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NOTE

From: General Secretariat of the Council
To: Delegations

Subject: Proposal for a Regulation of the European Parliament and of the Council amending Regulations (EC) No 1272/2008, (EC) No 1223/2009 and (EU) 2019/1009 as regards simplification of certain requirements and procedures for chemical products
- Initial 4-column table

Delegations will find in the Annex, for information, the initial four-column table on the above-mentioned proposal.

Proposal for a REGULATION OF THE EUROPEAN PARLIAMENT AND OF THE COUNCIL amending Regulations (EC) No 1272/2008, (EC) No 1223/2009 and (EU) 2019/1009 as regards simplification of certain requirements and procedures for chemical products (Text with EEA relevance)

2025/0531(COD)

30-04-2026

	CLEAN	Commission Proposal	vs.EC	EP Mandate	vs.EC	Council Mandate	vs.EC	Draft Agreement
Formula								
1		2025/0531 (COD)		2025/0531 (COD)		2025/0531 (COD)		
Document Stage								
2		Proposal for a		Proposal for a		Proposal for a		
Document Type								
3		REGULATION OF THE EUROPEAN PARLIAMENT AND OF THE COUNCIL		REGULATION OF THE EUROPEAN PARLIAMENT AND OF THE COUNCIL		REGULATION OF THE EUROPEAN PARLIAMENT AND OF THE COUNCIL		
Document Purpose								
4		amending Regulations (EC) No 1272/2008, (EC) No 1223/2009 and (EU) 2019/1009 as regards simplification of certain requirements and procedures for chemical products		amending Regulations (EC) No 1272/2008, (EC) No 1223/2009 and (EU) 2019/1009 as regards simplification of certain requirements and procedures for chemical products		amending Regulations (EC) No 1272/2008, (EC) No 1223/2009 and (EU) 2019/1009 as regards simplification of certain requirements and procedures for chemical products		
EEA Relevance								
5		(Text with EEA relevance)		(Text with EEA relevance)		(Text with EEA relevance)		
Formula								

	CLEAN	Commission Proposal	VS.EC	EP Mandate	VS.EC	Council Mandate	VS.EC	Draft Agreement
6	THE EUROPEAN PARLIAMENT AND THE COUNCIL OF THE EUROPEAN UNION,	THE EUROPEAN PARLIAMENT AND THE COUNCIL OF THE EUROPEAN UNION,	VS.EC	THE EUROPEAN PARLIAMENT AND THE COUNCIL OF THE EUROPEAN UNION,	VS.EC	THE EUROPEAN PARLIAMENT AND THE COUNCIL OF THE EUROPEAN UNION,	VS.EC	
Citation 1								
7	Having regard to the Treaty on the Functioning of the European Union, and in particular Article 114 thereof,	Having regard to the Treaty on the Functioning of the European Union, and in particular Article 114 thereof,				Having regard to the Treaty on the Functioning of the European Union, and in particular Article 114 thereof,		
Citation 2								
8	Having regard to the proposal from the European Commission,	Having regard to the proposal from the European Commission,		Having regard to the proposal from the European Commission,		Having regard to the proposal from the European Commission,		
Citation 3								
9	After transmission of the draft legislative act to the national Parliaments,	After transmission of the draft legislative act to the national Parliaments,		After transmission of the draft legislative act to the national Parliaments,		After transmission of the draft legislative act to the national Parliaments,		
Citation 4								
10	Having regard to the opinion of the European Economic and Social Committee ¹ , <u>I. OJ C [...], [...], p. [...].</u>	Having regard to the opinion of the European Economic and Social Committee ¹ , <u>I. OJ C [...], [...], p. [...].</u>		Having regard to the opinion of the European Economic and Social Committee ¹ , <u>I. OJ C [...], [...], p. [...].</u>		Having regard to the opinion of the European Economic and Social Committee ¹ , <u>I. OJ C [...], [...], p. [...].</u>		
Citation 5								
11	Acting in accordance with the ordinary legislative procedure,	Acting in accordance with the ordinary legislative procedure,		Acting in accordance with the ordinary legislative procedure,		Acting in accordance with the ordinary legislative procedure,		
Formula								
12	Whereas:	Whereas:		Whereas:		Whereas:		

	CLEAN	Commission Proposal	VS.EC	EP Mandate	VS.EC	Council Mandate	VS.EC	Draft Agreement
Recital 1		(1) High quality and safety requirements for products on the Single Market ensure a high level of protection of human health and the environment and contribute to a fair and sustainable economy. In international competition, the reputation of high-grade products manufactured in the Union can create an advantage for Union companies.	VS.EC	(1) High quality and safety requirements for products on the Single Market ensure a high level of protection of human health and the environment and contribute to a fair and sustainable economy. In international competition, the reputation of high-grade products manufactured in the Union can create an advantage for Union companies.	VS.EC	(1) High quality and safety requirements for products on the Single Market ensure a high level of protection of human health and the environment and contribute to a fair and sustainable economy. In international competition, the reputation of high-grade products manufactured in the Union can create an advantage for Union companies.		
13								
Recital 2		(2) The findings of the 2024 Draghi report ¹ indicated that the increasing number and complexity of rules risks limiting room for manoeuvre for Union businesses and preventing them from remaining competitive. Against this background, certain procedures and requirements laid down in Regulations (EC) No 1272/2008 ² , (EC) No 1223/2009 ³ and (EU) 2019/1009 ⁴ of the European Parliament and of the Council should be simplified and unnecessary regulatory burdens should be removed, while maintaining the same level of	VS.EC	(2) The findings of the 2024 Draghi report ¹ indicated that the increasing number and complexity of rules risks limiting room for manoeuvre for Union businesses and preventing them from remaining competitive. Against this background, certain procedures and requirements laid down in Regulations (EC) No 1272/2008 ² , (EC) No 1223/2009 ³ and (EU) 2019/1009 ⁴ of the European Parliament and of the Council should be simplified and unnecessary <u>administrative and</u> regulatory burdens should be removed, while maintaining the <u>same high</u> level of <u>consumer</u>	VS.EC	(2) The findings of the 2024 Draghi report ¹ indicated that the increasing number and complexity of rules risks limiting room for manoeuvre for Union businesses and preventing them from remaining competitive. Against this background, certain procedures and requirements laid down in Regulations (EC) No 1272/2008 ² , (EC) No 1223/2009 ³ and (EU) 2019/1009 ⁴ of the European Parliament and of the Council should be simplified and unnecessary regulatory burdens should be removed, while maintaining the same level of		
14								

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<p>protection of human health and of the environment.</p> <p>1. 2024 report by Mario Draghi on the future of European competitiveness: https://commission.europa.eu/topics/eu-competitiveness/draghi-report_en#paragraph_47059</p> <p>2. Regulation (EC) No 1272/2008 of the European Parliament and of the Council of 16 December 2008 on classification, labelling and packaging of substances and mixtures, amending and repealing Directives 67/548/EEC and 1999/45/EC, and amending Regulation (EC) No 1907/2006 (OJ L 353, 31.12.2008, p. 1, ELI: http://data.europa.eu/eli/reg/2008/1272/oj)</p> <p>3. Regulation (EC) No 1223/2009 of the European Parliament and of the Council of 30 November 2009 on cosmetic products (recast) (OJ L 342, 22.12.2009, p. 59, ELI: http://data.europa.eu/eli/reg/2009/1223/oj)</p> <p>4. Regulation (EU) 2019/1009 of the European Parliament and of the Council of 5 June 2019 laying down rules on the making available on the market of EU fertilising products and amending Regulations (EC) No 1069/2009 and (EC) No 1107/2009 and repealing Regulation (EC) No 2003/2003 (OJ L 170, 25.6.2019, p. 1, ELI: http://data.europa.eu/eli/reg/2019/1009/oj)</p>	<p>protection of human health and of the environment.</p> <p>1. 2024 report by Mario Draghi on the future of European competitiveness: https://commission.europa.eu/topics/eu-competitiveness/draghi-report_en#paragraph_47059</p> <p>2. Regulation (EC) No 1272/2008 of the European Parliament and of the Council of 16 December 2008 on classification, labelling and packaging of substances and mixtures, amending and repealing Directives 67/548/EEC and 1999/45/EC, and amending Regulation (EC) No 1907/2006 (OJ L 353, 31.12.2008, p. 1, ELI: http://data.europa.eu/eli/reg/2008/1272/oj)</p> <p>3. Regulation (EC) No 1223/2009 of the European Parliament and of the Council of 30 November 2009 on cosmetic products (recast) (OJ L 342, 22.12.2009, p. 59, ELI: http://data.europa.eu/eli/reg/2009/1223/oj)</p> <p>4. Regulation (EU) 2019/1009 of the European Parliament and of the Council of 5 June 2019 laying down rules on the making available on the market of EU fertilising products and amending Regulations (EC) No 1069/2009 and (EC) No 1107/2009 and repealing Regulation (EC) No 2003/2003 (OJ L 170, 25.6.2019, p. 1, ELI: http://data.europa.eu/eli/reg/2019/1009/oj)</p>	<p>protection, protection of human health and of the environment.</p> <p>1. 2024 report by Mario Draghi on the future of European competitiveness: https://commission.europa.eu/topics/eu-competitiveness/draghi-report_en#paragraph_47059</p> <p>2. Regulation (EC) No 1272/2008 of the European Parliament and of the Council of 16 December 2008 on classification, labelling and packaging of substances and mixtures, amending and repealing Directives 67/548/EEC and 1999/45/EC, and amending Regulation (EC) No 1907/2006 (OJ L 353, 31.12.2008, p. 1, ELI: http://data.europa.eu/eli/reg/2008/1272/oj)</p> <p>3. Regulation (EC) No 1223/2009 of the European Parliament and of the Council of 30 November 2009 on cosmetic products (recast) (OJ L 342, 22.12.2009, p. 59, ELI: http://data.europa.eu/eli/reg/2009/1223/oj)</p> <p>4. Regulation (EU) 2019/1009 of the European Parliament and of the Council of 5 June 2019 laying down rules on the making available on the market of EU fertilising products and amending Regulations (EC) No 1069/2009 and (EC) No 1107/2009 and repealing Regulation (EC) No 2003/2003 (OJ L 170, 25.6.2019, p. 1, ELI: http://data.europa.eu/eli/reg/2019/1009/oj)</p>	<p>protection of human health and of the environment.</p> <p>1. 2024 report by Mario Draghi on the future of European competitiveness: https://commission.europa.eu/topics/eu-competitiveness/draghi-report_en#paragraph_47059</p> <p>2. Regulation (EC) No 1272/2008 of the European Parliament and of the Council of 16 December 2008 on classification, labelling and packaging of substances and mixtures, amending and repealing Directives 67/548/EEC and 1999/45/EC, and amending Regulation (EC) No 1907/2006 (OJ L 353, 31.12.2008, p. 1, ELI: http://data.europa.eu/eli/reg/2008/1272/oj)</p> <p>3. Regulation (EC) No 1223/2009 of the European Parliament and of the Council of 30 November 2009 on cosmetic products (recast) (OJ L 342, 22.12.2009, p. 59, ELI: http://data.europa.eu/eli/reg/2009/1223/oj)</p> <p>4. Regulation (EU) 2019/1009 of the European Parliament and of the Council of 5 June 2019 laying down rules on the making available on the market of EU fertilising products and amending Regulations (EC) No 1069/2009 and (EC) No 1107/2009 and repealing Regulation (EC) No 2003/2003 (OJ L 170, 25.6.2019, p. 1, ELI: http://data.europa.eu/eli/reg/2019/1009/oj)</p>	
Recital 2a				

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14a		(2a) <u>Fertilising products covered by Regulation (EU) 2019/1009 directly affect farmers' production costs, incomes, safety and competitiveness, while regulatory requirements and administrative burdens on producers are transmitted along the supply chain. As competitiveness of the Union depends, inter alia, on its capacity to innovate, the regulatory framework should support innovation and technical progress by remaining proportionate and avoiding unnecessary burdens, while ensuring the objectives of that Regulation are achieved.</u>					
Recital 3 15	(3) In line with the Commission's objective to promote the 'digital by default' principle to support digital transformations and in order to facilitate communication between economic operators and national authorities responsible for enforcement, the indication of a digital contact on the label of hazardous substances and mixtures is necessary to enhance the	(3) In line with the Commission's objective to promote the 'digital by default' principle to support digital transformations and in order to facilitate communication between economic operators <u>suppliers and individuals and between suppliers</u> and national authorities responsible for enforcement, the indication of a digital contact on the label of hazardous substances	(3) In line with the Commission's objective to promote the 'digital by default' principle to support digital transformations and in order to facilitate communication between economic operators <u>suppliers</u> and national authorities responsible for enforcement , the indication of a digital contact on the label of hazardous substances and mixtures is necessary to enhance the	(3) In line with the Commission's objective to promote the 'digital by default' principle to support digital transformations and in order to facilitate communication between economic operators <u>suppliers</u> and national authorities responsible for enforcement , the indication of a digital contact on the label of hazardous substances and mixtures is necessary to enhance the			

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<p>effectiveness of official controls and enforcement and to expedite the process of detecting substances and mixtures that do not comply with the requirements of Regulation (EC) No 1272/2008. Currently, suppliers are required to indicate their address and telephone number on the label of the packaging of hazardous substances or mixtures, but this is not always sufficient to ensure that authorities responsible for enforcement can establish rapid contact. It is therefore necessary to require suppliers to provide a digital contact, which could be any up-to-date and accessible online communication channel with the supplier.</p>	<p>and mixtures is necessary to <u>enhance contributes to enhancing</u> the effectiveness of official controls, <u>traceability, accountability</u> and enforcement and to expedite the process of detecting substances and mixtures that do not comply with the requirements of Regulation (EC) No 1272/2008, <u>as well as to ensure consumers have a quick and direct contact in case of an emergency or accident</u>. Currently, suppliers are required to indicate their address and telephone number on the label of the packaging of hazardous substances or mixtures, but this is not always sufficient to ensure that authorities responsible for enforcement can establish rapid contact. It is therefore necessary to require suppliers to <u>also</u> provide a digital contact, which could <u>should</u> be any up-to-date, <u>easily and freely</u> accessible online communication channel with the supplier, <u>which allows for the storage of information on a durable medium</u>. <u>Where a digital contact is provided, the telephone number</u></p>	<p>effectiveness of official controls and enforcement and to expedite the process of detecting substances and mixtures that do not comply with the requirements of Regulation (EC) No 1272/2008. Currently, suppliers are required to indicate their address and telephone number on the label of the packaging of hazardous substances or mixtures, but this is not always sufficient to ensure that authorities responsible for enforcement can establish rapid contact. It is therefore necessary to require suppliers to also provide a digital contact, which could be any up-to-date and accessible online communication channel with the supplier. To reduce required label space, suppliers should be permitted to provide this telephone number on the digital label alone or through the digital contact. For example, where the digital contact is an email address, suppliers should be allowed to provide the telephone number in an auto-response, or where the digital contact is a webpage, suppliers should be allowed to provide the telephone</p>	<p>effectiveness of official controls and enforcement and to expedite the process of detecting substances and mixtures that do not comply with the requirements of Regulation (EC) No 1272/2008. Currently, suppliers are required to indicate their address and telephone number on the label of the packaging of hazardous substances or mixtures, but this is not always sufficient to ensure that authorities responsible for enforcement can establish rapid contact. It is therefore necessary to require suppliers to also provide a digital contact, which could be any up-to-date and accessible online communication channel with the supplier. To reduce required label space, suppliers should be permitted to provide this telephone number on the digital label alone or through the digital contact. For example, where the digital contact is an email address, suppliers should be allowed to provide the telephone number in an auto-response, or where the digital contact is a webpage, suppliers should be allowed to provide the telephone</p>	

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		<p><u>can be made directly available through that digital contact.</u></p>	<p>number by clearly stating it on the webpage. To allow suppliers sufficient time to adapt, the requirement to provide a digital contact should first be applied to the placing on the market of substances and mixtures three years after the entry into force of this Regulation. Products placed on the market within the three-year deadline should be permitted to remain on the market without relabelling for a further two years after the passing of the three-year deadline. During the transitional period, the existing rules will continue to apply, including the requirement to use provide a telephone number. Suppliers should also be able to apply the new rules following the entry into force of this Regulation and be allowed to provide the telephone number on digital labels or through the digital contact.</p>	
Recital 3a		<p><u>(3a) The digital contact should allow consumers and authorities to contact suppliers directly and</u></p>		
15a				

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		<p><u>swiftly, and should be accessible free of charge, and without the need to provide any personal data, download or use an application or an obligation to register solely for the purpose of contacting the supplier. Such digital contact could include, for example, an email address or a contact form on a website. However, it should not be understood as encompassing automatic replies to queries, chatbots or fax numbers. The term 'digital contact', similarly to the term 'electronic address' in Regulation (EU) 2023/988 of the European Parliament and of the Council, should be understood in a technologically neutral manner, capable of evolving with future technological developments, and cover all forms of direct digital communication.</u></p>					
Recital 4							
16	<p>(4) Regulation (EC) No 1272/2008 laid down the exemptions from labelling and packaging requirements for packaging of specific shapes, forms or sizes. Those exemptions</p>	<p>(4) Regulation (EC) No 1272/2008 laid down the exemptions from labelling and packaging requirements for packaging of specific shapes, forms or sizes. Those exemptions</p>	<p>(4) Regulation (EC) No 1272/2008 laid down the exemptions from labelling and packaging requirements for packaging of specific shapes, forms or sizes. Those exemptions</p>				

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	can only be triggered if all required label elements do not fit on the outer packaging or on a tie-on tag. To simplify the use of those exemptions, it is appropriate to allow the existing exemptions to be applied to smaller packages without the need to prove the impossibility of using the outer packaging or tie-on tag.	can only be triggered if all required label elements do not fit on the outer packaging or on a tie-on tag. To simplify the use of those exemptions, it is appropriate to allow the existing exemptions to be applied to smaller packages without the need to prove the impossibility of using the outer packaging or tie-on tag.	can only be triggered if all required label elements do not fit on the outer packaging or on a tie-on tag. To simplify the use of those exemptions, it is appropriate to allow the existing exemptions to be applied to smaller packages without the need to prove the impossibility of using the outer packaging or tie-on tag.	
Recital 5				
17	<p>(5) Regulation (EU) 2024/2865 of the Council¹ introduced exemptions from labelling and packaging requirements for packages containing less than 10 ml. However, further simplifications are needed with regard to the derogation in cases where the 10 ml derogation is applied.</p> <p>¹ Regulation (EU) 2024/2865 of the European Parliament and of the Council of 23 October 2024 amending Regulation</p>	<p>(5) Regulation (EU) 2024/2865 of the Council⁴ introduced exemptions from labelling and packaging requirements for packages containing less than 10 ml. <u>It introduced a possibility to omit label elements from such inner packaging under certain conditions.</u> However, further <u>simplifications are needed with regard to the application of this derogation in cases where these packages are subject to the supplementary hazard statement EUH-208—it is also necessary to simplify these provisions and</u> clarify <u>cases that require labelling elements to be presented on the</u></p>	<p>(5) Regulation (EU) 2024/2865 of the Council¹ introduced exemptions from labelling and packaging requirements for packages containing less than 10 ml. It introduced a possibility to omit label elements from such inner packaging under certain conditions. However, further simplifications are needed with regard to the application of this derogation in cases where these packages are subject to the supplementary hazard statement EUH-208—it is also necessary to simplify these provisions and clarify cases that require labelling elements to be</p>	

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	<p>(EC) No 1272/2008 on classification, labelling and packaging of substances and mixtures (OJ L, 2024/2865, 20.11.2024, ELI: http://data.europa.eu/eli/reg/2024/2865/oj)</p>	<p>requirements for inner and outer packaging in cases where the 10 ml derogation is applied allowing <u>for these elements to be fully omitted. In addition, the Commission should carry out an assessment on whether further specific reductions of mandatory label elements should apply to packages between 10 and 125 ml.</u></p> <p>4. Regulation (EU) 2024/2865 of the European Parliament and of the Council of 23 October 2024 amending Regulation (EC) No 1272/2008 on classification, labelling and packaging of substances and mixtures (OJ L, 2024/2865, 20.11.2024, ELI: http://data.europa.eu/eli/reg/2024/2865/oj)</p>	<p>presented on the requirements for inner and outer packaging and cases where the 10 ml derogation is applied allowing for these elements to be fully omitted.</p> <p><u>1. Regulation (EU) 2024/2865 of the European Parliament and of the Council of 23 October 2024 amending Regulation (EC) No 1272/2008 on classification, labelling and packaging of substances and mixtures (OJ L, 2024/2865, 20.11.2024, ELI: http://data.europa.eu/eli/reg/2024/2865/oj)</u></p>	
Recital 6				
18	<p>(6) In order to provide the flexibility for suppliers of substances and mixtures, to create equal conditions for small and medium-sized enterprises who often outsource label printing services and to facilitate the preparation and production of fold-out labels, which is significantly longer than the production of the standard labels, it is necessary to remove a fixed</p>	<p>(6) <u>With regard to the updating of labels in case of new or more severe self-classification, suppliers should inform their direct downstream users about the results of the new evaluation without undue delay.</u> In order to provide <u>the flexibility for sufficient time for all</u> suppliers of substances and mixtures, to create equal conditions in particular, for small and medium-sized enterprises who</p>	<p>(6) In order to provide the flexibility for suppliers of substances and mixtures, to create equal conditions for small and medium-sized enterprises who often outsource label printing services and to facilitate the preparation and production of fold-out labels, which is significantly longer than the production of the standard labels, it is necessary to remove extend</p>	

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<p>six months relabelling deadline and to require the labels to be changed without undue delay after new data was obtained by or communicated to a supplier.</p>	<p>often outsource label printing services and to facilitate the preparation and <u>the</u> production of fold-out labels, which is significantly longer than the production of the standard labels, it is necessary to remove <u>extend</u> <u>the</u> fixed six months relabelling deadline ###to eighteen months. <u>while continuing</u> to require the labels to be changed without undue delay after new data was obtained by or communicated to a supplier.</p>	<p>the fixed six months relabelling deadline ###to twelve months, while continuing to require the labels to be changed without undue delay after new data was obtained by or communicated to a supplier.</p>		
Recital 7	<p>(7) Regulation (EU) 2024/2865 laid down rules on mandatory requirements for label formatting. New information¹ pointed to excessive administrative burden and costs, associated with these requirements. To balance the need for label information to be clearly understood by consumers with the need to reduce market barriers and burden for industry², it is necessary to simplify the current formatting obligations without reducing the level of protection of human health and the</p>	<p>(7) Regulation (EU) 2024/2865 laid down rules on mandatory requirements for label formatting. New information⁴ pointed to excessive administrative burden and costs, associated with these requirements. To balance the need for label information to be clearly understood by consumers with the need to reduce market barriers and <u>unjustified</u> burden for industry^{2,8}, <u>in particular for small and medium-sized enterprises</u>, it is necessary to simplify the current formatting obligations without</p>	<p>(7) Regulation (EU) 2024/2865 laid down rules on mandatory requirements for label formatting. New information¹ pointed to excessive administrative burden and costs, associated with these requirements. To balance the need for label information to be clearly understood by consumers with the need to reduce market barriers and burden for industry², it is necessary to simplify the current formatting obligations without reducing the level of protection of human health and the</p>	

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	<p>environment. Economic operators and enforcement authorities must remain responsible for ensuring that the labels are legible in accordance with the legal requirements.</p> <p>1. Detailed analysis of costs associated with new formatting requirements is provided in the Staff Working Document Accompanying the document Proposal for a Regulation of the European Parliament and of the Council amending Regulations (EC) No 1272/2008, (EC) No 1223/2009 and (EU) 2019/1009 as regards simplification of certain requirements and procedures for chemical products, SWD(2025) 531, p. 14.</p> <p>2. As outlined in the Communication from the Commission to the European Parliament, the Council, the European Economic and Social Committee and the Committee of the Regions The Single Market: our European home market in an uncertain world, A Strategy for making the Single Market simple, seamless and strong, COM(2025) 500 final, p. 10, available at: https://single-market-economy.ec.europa.eu/document/download/492c78d0-7447-4a16-b53f-1cead54bc649_en?filename=Communicati on%20Single%20Market%20Strategy.pdf.</p>	<p>reducing the level of protection of human health and the environment. <i>Economic operators and enforcement authorities</i> <u>The new measures should maintain a high level of consumer protection and ensure the proper functioning of the internal market. Suppliers</u> must remain responsible for ensuring that the labels are legible in accordance with the legal requirements.</p> <p><i>1. Detailed analysis of costs associated with new formatting requirements is provided in the Staff Working Document Accompanying the document Proposal for a Regulation of the European Parliament and of the Council amending Regulations (EC) No 1272/2008, (EC) No 1223/2009 and (EU) 2019/1009 as regards simplification of certain requirements and procedures for chemical products, SWD(2025) 531, p. 14.</i></p> <p><i>2. As outlined in the Communication from the Commission to the European Parliament, the Council, the European Economic and Social Committee and the Committee of the Regions The Single Market: our European home market in an uncertain world, A Strategy for making the Single Market simple, seamless and strong, COM(2025) 500 final, p. 10, available at: https://single-market-economy.ec.europa.eu/document/download/492c78d0-7447-4a16-b53f-1cead54bc649_en?filename=Communicati on%20Single%20Market%20Strategy.pdf.</i></p>	<p>environment. Economic operators and enforcement Suppliers must remain responsible for ensuring that the labels are legible in accordance with the legal requirements.</p> <p>1. Detailed analysis of costs associated with new formatting requirements is provided in the Staff Working Document Accompanying the document Proposal for a Regulation of the European Parliament and of the Council amending Regulations (EC) No 1272/2008, (EC) No 1223/2009 and (EU) 2019/1009 as regards simplification of certain requirements and procedures for chemical products, SWD(2025) 531, p. 14.</p> <p>2. As outlined in the Communication from the Commission to the European Parliament, the Council, the European Economic and Social Committee and the Committee of the Regions The Single Market: our European home market in an uncertain world, A Strategy for making the Single Market simple, seamless and strong, COM(2025) 500 final, p. 10, available at: https://single-market-economy.ec.europa.eu/document/download/492c78d0-7447-4a16-b53f-1cead54bc649_en?filename=Communicati on%20Single%20Market%20Strategy.pdf.</p>				

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Recital 7a			text%20-%20Single%20Market%20Strategy.pdf					
19a		<p><u>(7a) The label serves as a primary and often single source of hazard information and safe-use guidance readily available to consumers, while professional and industrial users are also informed about the hazards associated with a particular substance or mixture prior to use through safety data sheets and safety training. Therefore, it is indispensable that a label is easily readable not only under normal conditions, but also in exceptional circumstances such as accidents.</u></p>						
Recital 7b		<p><u>(7b) To ensure that a label has an appropriate degree of readability, it should at least have a clear contrast of the text of the label to the background, a suitable typeface, an appropriately sized font, appropriate line and letter spacing, overall label design and other relevant formatting elements. In particular, for</u></p>						
19b								

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		<p><u>substances or mixtures made available on the market for the general public, the label elements referred to in Article 17(1) should use a font size where the x-height is equal to or greater than 1.2 mm. However, when the contents of the package do not exceed 125 ml, the label elements referred to in Article 17(1) might use a font size where the x-height is equal to or greater than 0.9 mm. The European Chemicals Agency (ECHA) should update its guidance on formatting of labels and include clear examples of what constitutes acceptable and unacceptable examples of label formatting. The guidance should consider labelling formats from other relevant Union laws and take account of best practice on "accessible design" and accepted standards for readability, including acceptable as well as unacceptable colour combinations to ensure contrast. The guidance should assist enforcement by relevant national authorities, be developed in line with Better Regulation principles and informed by appropriate</u></p>		

CLEAN	Commission Proposal	VS.EC EP Mandate	VS.EC Council Mandate	VS.EC Draft Agreement
Recital 8	<p>(8) To alleviate the burden on industry and to improve the free circulation of substances and mixtures in the internal market it is appropriate to amend Regulation (EC) No 1272/2008 as regards the rules on advertisements and distance offers, taking advantages of existing provisions in other Union legislation with the same objectives. In this regard, requirements for advertisements and distance offers should be limited to products placed on the market for the general public, as Regulation (EC) No 1907/2006¹ already provides clear obligations on information flows in supply chains for substances and mixtures.</p> <p>1. Regulation (EC) No 1907/2006 of the European Parliament and of the Council of 18 December 2006 concerning the Registration, Evaluation, Authorisation</p>	<p><u>stakeholder consultation, support competitiveness by reducing unnecessary burdens, provide certainty for suppliers, and ensure fair competition on the internal market.</u></p>		
20	<p>(8) To alleviate the burden on industry and to improve the free circulation of substances and mixtures in the internal market it is appropriate to amend Regulation (EC) No 1272/2008 as regards the rules on advertisements and distance <u>sales</u> offers, taking advantages of existing provisions in other Union legislation with the same objectives. In this regard, requirements for advertisements and distance <u>sales</u> offers should be limited to products placed on the market for the general public, as Regulation (EC) No 1907/2006² already provides clear obligations on information flows in supply chains for substances and mixtures.</p> <p>1. Regulation (EC) No 1907/2006 of the European Parliament and of the Council of 18 December 2006 concerning the Registration, Evaluation, Authorisation</p>	<p>(8) To alleviate the burden on industry and to improve the free circulation of substances and mixtures in the internal market it is appropriate to amend Regulation (EC) No 1272/2008 as regards the rules on advertisements and distance <u>sales</u> offers, taking advantages of existing provisions in other Union legislation with the same objectives. In this regard, requirements for advertisements and distance <u>sales</u> offers should be limited to products placed on the market for the general public, as Regulation (EC) No 1907/2006¹ already provides clear obligations on information flows in supply chains for substances and mixtures for professional and industrial users through the safety data sheet, which must be provided no later than the date on which the</p>	<p>(8) To alleviate the burden on industry and to improve the free circulation of substances and mixtures in the internal market it is appropriate to amend Regulation (EC) No 1272/2008 as regards the rules on advertisements and distance sales offers, taking advantages of existing provisions in other Union legislation with the same objectives. In this regard, certain requirements for advertisements and distance sales offers should be limited to products placed on the market for the general public, as Regulation (EC) No 1907/2006¹ already provides clear obligations on information flows in supply chains for substances and mixtures for professional and industrial users through the safety data sheet, which must be provided no later than the date on which the</p>	

CLEAN	Commission Proposal	VS,EC	EP Mandate	VS,EC	Council Mandate	VS,EC	Draft Agreement
	<p>and Restriction of Chemicals (REACH), establishing a European Chemicals Agency, amending Directive 1999/45/EC and repealing Council Regulation (EEC) No 793/93 and Commission Regulation (EC) No 1488/94 as well as Council Directive 76/769/EEC and Commission Directives 91/155/EEC, 93/67/EEC, 93/105/EC and 2000/21/EC (OJ L 396, 30.12.2006, p. 1, ELI: http://data.europa.eu/eli/reg/2006/1907/oj)</p>	<p>and Restriction of Chemicals (REACH), establishing a European Chemicals Agency, amending Directive 1999/45/EC and repealing Council Regulation (EEC) No 793/93 and Commission Regulation (EC) No 1488/94 as well as Council Directive 76/769/EEC and Commission Directives 91/155/EEC, 93/67/EEC, 93/105/EC and 2000/21/EC (OJ L 396, 30.12.2006, p. 1, ELI: http://data.europa.eu/eli/reg/2006/1907/oj)</p>	<p>substance or a mixture is first supplied.</p> <p>1. Regulation (EC) No 1907/2006 of the European Parliament and of the Council of 18 December 2006 concerning the Registration, Evaluation, Authorisation and Restriction of Chemicals (REACH), establishing a European Chemicals Agency, amending Directive 1999/45/EC and repealing Council Regulation (EEC) No 793/93 and Commission Regulation (EC) No 1488/94 as well as Council Directive 76/769/EEC and Commission Directives 91/155/EEC, 93/67/EEC, 93/105/EC and 2000/21/EC (OJ L 396, 30.12.2006, p. 1, ELI: http://data.europa.eu/eli/reg/2006/1907/oj)</p>				
20a		<p><u>(8a) Professional and industrial users are generally better informed about the hazards associated with substances and mixtures than consumers. It is therefore appropriate to differentiate information requirements with respect to advertisement depending on whether products are intended for professional and industrial use or for consumers or directly made available to them. When assessing whether an</u></p>					

CLEAN	Commission Proposal	VS,EC EP Mandate	VS,EC Council Mandate	VS,EC Draft Agreement
		<p><u>advertisement is targeted at consumers or made available to them, competent authorities should take into account objective elements, including whether it is clearly indicated that the substance or mixture is intended exclusively for professional use, as well as the context in which the advertisement or offer is made available, such as trade fairs, trade publications or digital platforms addressed to professional users. Similar provisions already exist under Union law.</u></p>		
Recital 9	<p>(9) Before the amendments introduced by Regulation (EU) 2024/2865, Regulation (EC) No 1272/2008 required the advertisements for hazardous mixtures, for which a member of the general public is allowed to conclude a contract for purchase without first having sight of the label, to mention the type or types of hazards indicated on the label, and required advertisements for substances to mention the hazard classes or hazard categories</p>	<p>(9) Before the amendments introduced by Regulation (EU) 2024/2865, Regulation (EC) No 1272/2008 required the advertisements for hazardous mixtures, for which a member of the general public is allowed to conclude a contract for purchase without first having sight of the label, to mention the type or types of hazards indicated on the label, and required advertisements for substances to mention the hazard classes or hazard categories</p>	<p>(9) Before the amendments introduced by Regulation (EU) 2024/2865, Regulation (EC) No 1272/2008 required the advertisements for hazardous mixtures, for which a member of the general public is allowed to conclude a contract for purchase without first having sight of the label, to mention the type or types of hazards indicated on the label, and required advertisements for substances to mention the hazard classes or hazard categories</p>	
21	<p>(9) Before the amendments introduced by Regulation (EU) 2024/2865, Regulation (EC) No 1272/2008 required the advertisements for hazardous mixtures, for which a member of the general public is allowed to conclude a contract for purchase without first having sight of the label, to mention the type or types of hazards indicated on the label, and required advertisements for substances to mention the hazard classes or hazard categories</p>	<p>(9) Before the amendments introduced by Regulation (EU) 2024/2865, Regulation (EC) No 1272/2008 required the advertisements for hazardous mixtures, for which a member of the general public is allowed to conclude a contract for purchase without first having sight of the label, to mention the type or types of hazards indicated on the label, and required advertisements for substances to mention the hazard classes or hazard categories</p>	<p>(9) Before the amendments introduced by Regulation (EU) 2024/2865, Regulation (EC) No 1272/2008 required the advertisements for hazardous mixtures, for which a member of the general public is allowed to conclude a contract for purchase without first having sight of the label, to mention the type or types of hazards indicated on the label, and required advertisements for substances to mention the hazard classes or hazard categories</p>	

CLEAN	Commission Proposal	VS.EC	EP Mandate	VS.EC	Council Mandate	VS.EC	Draft Agreement
	<p>concerned. Regulation (EU) 2024/2865 introduced a new requirement for all distance sales of hazardous substances and mixtures to include all labelling information in the offer, thus ensuring that the buyer is always informed about the hazards before buying the product. That Regulation also expanded requirements for advertisements, requiring them to indicate the hazard pictograms, signal words, hazard statements and supplemental statements and, in addition, to invite general public to follow the information on the product label. Advertisements are means of promoting the sale or use of chemical products, and at the moment of sale the label on the packaging of the substance or mixture or the label information in the distance offer provides full information about the hazards associated with that substance or mixture. It would therefore be appropriate to require advertisements to invite customers to read the label and product information before use, but not to</p>	<p>concerned. Regulation (EU) 2024/2865 introduced a new requirement for all distance sales of hazardous substances and mixtures to include all labelling information in the offer, thus ensuring that the buyer is always informed about the hazards before buying the product. That Regulation also expanded requirements for advertisements, requiring them to indicate the hazard pictograms, signal words, hazard statements and supplemental statements and, in addition, to invite general public to follow the information on the product label. Advertisements are means of promoting the sale or use of chemical products, and at the moment of sale the label on the packaging of the substance or mixture or the label information in the distance offer provides full information about the hazards associated with that substance or mixture. It would therefore be appropriate to require advertisements to invite customers to read the label and product information before use, <u>and to include either the signal word or</u></p>	<p>concerned. Regulation (EU) 2024/2865 introduced a new requirement for all distance sales of hazardous substances and mixtures to include all labelling information in the offer, thus ensuring that the buyer is always informed about the hazards before buying the product. That Regulation also expanded requirements for advertisements, requiring them to indicate the hazard pictograms, signal words, hazard statements and supplemental statements and, in addition, to invite general public to follow the information on the product label. Advertisements are means of promoting the sale or use of chemical products, and at the moment of sale the label on the packaging of the substance or mixture or the label information in the distance offer provides full information about the hazards associated with that substance or mixture. It would therefore be appropriate to require advertisements to invite customers to read the label and product information before use or to include the pictogram, but not to</p>				

CLEAN	Commission Proposal	VS,EC EP Mandate	VS,EC Council Mandate	VS,EC Draft Agreement
	<p>duplicate the hazard information from the label.</p>	<p><u>the pictogram</u> but not to duplicate the hazard information from the label. <u>This approach ensures that advertisements remain sufficiently informative in alerting potential users to the hazardous nature of the product without imposing requirements that would make advertisements cluttered, difficult to interpret or commercially impractical. Given the many different forms of advertisements, suppliers should be provided a degree of flexibility on how this information should be conveyed in advertisements.</u></p>	<p>duplicate the hazard information from the label. Given the many different forms of advertisements, suppliers should be provided a degree of flexibility on how this information should be conveyed in advertisements.</p>	
Recital 10	<p>(10) As Regulation (EC) No 1107/2009 of the European Parliament and of the Council¹ and Regulation (EU) No 528/2012 of the European Parliament and of the Council² require advertisements for authorised plant protection products and biocidal products to use the statement ‘Always read the label and product information before use’, it would be appropriate for use the same requirement for hazardous advertisements of hazardous</p>	<p>(10) As Regulation (EC) No 1107/2009 of the European Parliament and of the Council¹⁰ and Regulation (EU) No 528/2012 of the European Parliament and of the Council¹¹ require advertisements for authorised plant protection products and biocidal products to use the statement ‘Always read the label and product information before use’, it would be appropriate <u>for the advertisers to use the same requirement for</u></p>	<p>(10) As Regulation (EC) No 1107/2009 of the European Parliament and of the Council¹ and Regulation (EU) No 528/2012 of the European Parliament and of the Council² require advertisements for authorised plant protection products and biocidal products to use the statement ‘Always read the label and product information before use’, it would be appropriate to allow advertisers to use a similar statementuse the same</p>	
22				

	CLEAN	Commission Proposal	VS.EC	EP Mandate	VS.EC	Council Mandate	VS.EC	Draft Agreement
	<p>substances and mixtures to ensure consistency, especially in cases where advertised hazardous substances and mixtures are also authorised plant protection products or biocidal products.</p> <p>1. Regulation (EC) No 1107/2009 of the European Parliament and of the Council of 21 October 2009 concerning the placing of plant protection products on the market and repealing Council Directives 79/117/EEC and 91/414/EEC (OJ L 309, 24.11.2009, p. 1, ELI: http://data.europa.eu/eli/reg/2009/1107/oj)</p> <p>2. Regulation (EU) No 528/2012 of the European Parliament and of the Council of 22 May 2012 concerning the making available on the market and use of biocidal products (OJ L 167, 27.6.2012, p. 1, ELI: http://data.europa.eu/eli/reg/2012/528/oj).</p>	<p>advertisements of hazardous substances and mixtures to ensure consistency, especially in cases where advertised hazardous substances and mixtures are also authorised plant protection products or biocidal products.</p> <p>1. Regulation (EC) No 1107/2009 of the European Parliament and of the Council of 21 October 2009 concerning the placing of plant protection products on the market and repealing Council Directives 79/117/EEC and 91/414/EEC (OJ L 309, 24.11.2009, p. 1, ELI: http://data.europa.eu/eli/reg/2009/1107/oj)</p> <p>2. Regulation (EU) No 528/2012 of the European Parliament and of the Council of 22 May 2012 concerning the making available on the market and use of biocidal products (OJ L 167, 27.6.2012, p. 1, ELI: http://data.europa.eu/eli/reg/2012/528/oj)</p>	<p>advertisements of hazardous substances and mixtures to ensure consistency, especially in cases where advertised hazardous substances and mixtures are also authorised plant protection products or biocidal products.</p> <p>1. Regulation (EC) No 1107/2009 of the European Parliament and of the Council of 21 October 2009 concerning the placing of plant protection products on the market and repealing Council Directives 79/117/EEC and 91/414/EEC (OJ L 309, 24.11.2009, p. 1, ELI: http://data.europa.eu/eli/reg/2009/1107/oj)</p> <p>2. Regulation (EU) No 528/2012 of the European Parliament and of the Council of 22 May 2012 concerning the making available on the market and use of biocidal products (OJ L 167, 27.6.2012, p. 1, ELI: http://data.europa.eu/eli/reg/2012/528/oj).</p>	<p>requirement for advertisements of hazardous substances and mixtures to ensure consistency, especially in cases where advertised hazardous substances and mixtures are also authorised plant protection products or biocidal products.</p> <p>1. Regulation (EC) No 1107/2009 of the European Parliament and of the Council of 21 October 2009 concerning the placing of plant protection products on the market and repealing Council Directives 79/117/EEC and 91/414/EEC (OJ L 309, 24.11.2009, p. 1, ELI: http://data.europa.eu/eli/reg/2009/1107/oj)</p> <p>2. Regulation (EU) No 528/2012 of the European Parliament and of the Council of 22 May 2012 concerning the making available on the market and use of biocidal products (OJ L 167, 27.6.2012, p. 1, ELI: http://data.europa.eu/eli/reg/2012/528/oj).</p>	Draft Agreement			
Recital 11								
23	<p>(11) Regulation (EU) 2024/2865 introduced specific provisions for the labelling of fuels supplied at fuelling stations. To remove unnecessary burdens on businesses, without undermining the protection of human health and the environment, it is necessary to</p>	<p>(11) Regulation (EU) 2024/2865 introduced specific provisions for the labelling of fuels supplied at fuelling stations. To remove unnecessary burdens on businesses, without undermining the protection of human health and the environment, it is necessary to</p>	<p>(11) Regulation (EU) 2024/2865 introduced specific provisions for the labelling of fuels supplied at fuelling stations. To remove unnecessary burdens on businesses, without undermining the protection of human health and the environment, it is necessary to</p>	<p>(11) Regulation (EU) 2024/2865 introduced specific provisions for the labelling of fuels supplied at fuelling stations. To remove unnecessary burdens on businesses, without undermining the protection of human health and the environment, it is necessary to</p>	<p>(11) Regulation (EU) 2024/2865 introduced specific provisions for the labelling of fuels supplied at fuelling stations. To remove unnecessary burdens on businesses, without undermining the protection of human health and the environment, it is necessary to</p>	<p>(11) Regulation (EU) 2024/2865 introduced specific provisions for the labelling of fuels supplied at fuelling stations. To remove unnecessary burdens on businesses, without undermining the protection of human health and the environment, it is necessary to</p>	<p>(11) Regulation (EU) 2024/2865 introduced specific provisions for the labelling of fuels supplied at fuelling stations. To remove unnecessary burdens on businesses, without undermining the protection of human health and the environment, it is necessary to</p>	<p>(11) Regulation (EU) 2024/2865 introduced specific provisions for the labelling of fuels supplied at fuelling stations. To remove unnecessary burdens on businesses, without undermining the protection of human health and the environment, it is necessary to</p>

CLEAN	Commission Proposal	VS.EC EP Mandate	VS.EC Council Mandate	VS.EC Draft Agreement
	clarify in Annex II of Regulation (EC) No 1272/2008 which of the label elements required by Article 17(1) of that Regulation are not needed on the pump.	clarify in Annex II of Regulation (EC) No 1272/2008 which of the label elements required by Article 17(1) of that Regulation are not needed on the pump.	clarify in Annex II of Regulation (EC) No 1272/2008 which of the label elements required by Article 17(1) of that Regulation are not needed on the pump.	
Recital 12	<p>(12) Regulation (EU) 2024/2865 introduced the possibility to include certain labelling elements in the digital label only. To ensure broader use of technology and to allow a simpler and more flexible approach to labelling, suppliers should be allowed to place contact details of any additional suppliers on the digital label only. Inclusion of the digital contact would also be appropriate where contact details of additional suppliers are provided in the digital label.</p>	<p>(12) Regulation (EU) 2024/2865 introduced the possibility to include certain labelling elements in the digital label only. To ensure broader use of technology and to allow a simpler and more flexible approach to labelling, suppliers should be allowed to place contact details of any additional suppliers on the digital label only. Inclusion of the digital contact would also be appropriate where contact details of additional suppliers are provided in the digital label. <u><i>In order to ensure the possibility for rapid contact which is essential in certain situations such as in cases of emergency, the presence of a digital contact should not exclude the provision of a telephone number.</i></u></p>	<p>(12) Regulation (EU) 2024/2865 introduced the possibility to include certain labelling elements in the digital label only. To ensure broader use of technology and to allow a simpler and more flexible approach to labelling, suppliers should be allowed to place contact details of any additional suppliers on the digital label only. Inclusion of the digital contact would also be appropriate where contact details of additional suppliers are provided in the digital label.</p>	
Recital 12a				

	CLEAN	Commission Proposal	VS,EC	EP Mandate	VS,EC	Council Mandate	VS,EC	Draft Agreement
24a			VS,EC	<u>(12a) Inkjet cartridges (≤150 ml) (supplied in outer packaging and designed to be installed in a printer by a consumer or professional user) have very limited usable surface area for labelling and cannot benefit from the fold out label option enabling multilanguage solutions. In this case suppliers should be permitted to reduce hazard label information under certain conditions.</u>				
Recital 13								
25		(13) To ensure that suppliers of substances and mixtures have time to adapt to the new rules on classification, labelling and packaging, the application of some provisions of this Regulation should be deferred. Substances and mixtures which are already placed on the market before the end of that deferral period, should not be required to be reclassified or relabelled in accordance with this Regulation, to avoid an additional burden on suppliers of substances and mixtures.		(13) To ensure that suppliers of substances and mixtures have time to adapt to the new rules on classification, labelling and packaging, the application of some provisions of this Regulation should be deferred. Substances and mixtures which are already placed on the market before the end of that deferral period, should not be required to be reclassified or relabelled in accordance with this Regulation, to avoid an additional burden on suppliers of substances and mixtures.		(13) To ensure that suppliers of substances and mixtures have time to adapt to the new rules on classification, labelling and packaging, the application of some provisions of this Regulation should be deferred. Substances and mixtures which are already placed on the market before the end of that deferral period, should not be required to be reclassified or relabelled in accordance with this Regulation, to avoid an additional burden on suppliers of substances and mixtures.		
Recital 14								

	CLEAN	Commission Proposal	vs.EC	EP Mandate	vs.EC	Council Mandate	vs.EC	Draft Agreement
26	(14) In line with the transitional provisions of Regulation (EC) No 1272/2008, suppliers should have the possibility of applying the new classification, labelling and packaging provisions introduced by this Regulation on a voluntary basis before the date of the deferred application of these provisions.	(14) In line with the transitional provisions of Regulation (EC) No 1272/2008, suppliers should have the possibility of applying the new classification, labelling and packaging provisions introduced by this Regulation on a voluntary basis before the date of the deferred application of these provisions <u>and national authorities should encourage them to do so.</u>		(14) In line with the transitional provisions of Regulation (EC) No 1272/2008, suppliers should have the possibility of applying the new classification, labelling and packaging provisions introduced by this Regulation on a voluntary basis before the date of the deferred application of these provisions.				
Recital 15								
27	(15) In accordance with Regulation (EC) No 1223/2009, colorants, preservatives and UV filters may only be used in cosmetic products if they are listed in Annexes IV to VI to that Regulation. To ensure legal certainty for economic operators, a procedure should be provided specifying the different steps in the process for a substance to be included into the respective Annex for the purposes of adapting the Annexes to technical and scientific progress in accordance with Article 31(2) of Regulation (EC) No 1223/2009.	(15) In accordance with Regulation (EC) No 1223/2009, colorants, preservatives and UV filters may only be used in cosmetic products if they are listed in Annexes IV to VI to that Regulation. To ensure legal certainty for economic operators, a procedure should be provided specifying the different steps in the process for a substance to be included into the respective Annex for the purposes of adapting the Annexes to technical and scientific progress in accordance with Article 31(2) of Regulation (EC) No 1223/2009.		(15) In accordance with Regulation (EC) No 1223/2009, colorants, preservatives and UV filters may only be used in cosmetic products if they are listed in Annexes IV to VI to that Regulation. To ensure legal certainty for economic operators, a procedure should be provided specifying the different steps in the process for a substance to be included into the respective Annex for the purposes of adapting the Annexes to technical and scientific progress in accordance with Article 31(2) of Regulation (EC) No 1223/2009.		(15) In accordance with Regulation (EC) No 1223/2009, colorants, preservatives and UV filters may only be used in cosmetic products if they are listed in Annexes IV to VI to that Regulation. To ensure legal certainty for economic operators, a procedure should be provided specifying the different steps in the process for a substance to be included into the respective Annex for the purposes of adapting the Annexes to technical and scientific progress in accordance with Article 31(2) of Regulation (EC) No 1223/2009.		

CLEAN	Commission Proposal	VS.EC	EP Mandate	VS.EC	Council Mandate	VS.EC	Draft Agreement
Recital 16	(16) Regulation (EC) No 1223/2009 provides the possibility to use in cosmetic products substances classified as CMR under Part 3 of Annex VI to Regulation (EC) No 1272/2008 under certain conditions. A request for derogation should be submitted to the Commission at the latest three months after the entry into force of the respective changes to Part 3 of Annex VI to Regulation (EC) No 1272/2008. As the relevant opinion of the Committee for Risk Assessment proposing the substances for the harmonised classification is made publicly available several months before the Commission follows with the regulatory measure, such deadline is sufficient.	VS.EC	(16) Regulation (EC) No 1223/2009 provides the possibility to use in cosmetic products substances classified as CMR under Part 3 of Annex VI to Regulation (EC) No 1272/2008 under certain conditions. A request for derogation should be submitted to the Commission at the latest three months after the entry into force of the respective changes to Part 3 of Annex VI to Regulation (EC) No 1272/2008. As the relevant opinion of the Committee for Risk Assessment proposing the substances for the harmonised classification is made publicly available several months before the Commission follows with the regulatory measure, such deadline is sufficient.	VS.EC	(16) Regulation (EC) No 1223/2009 provides the possibility to use in cosmetic products substances classified as CMR under Part 3 of Annex VI to Regulation (EC) No 1272/2008 under certain conditions. A request for derogation should be submitted to the Commission at the latest three months after the entry into force of the respective changes to Part 3 of Annex VI to Regulation (EC) No 1272/2008. As the relevant opinion of the Committee for Risk Assessment proposing the substances for the harmonised classification is made publicly available several months before the Commission follows with the regulatory measure, such deadline is sufficient.	VS.EC	
28							
Recital 17	(17) The conditions allowing for exemptions from the ban of use of such substances in cosmetic products should be streamlined, and their scope should be set out in more detail. In addition, compliance with food safety	VS.EC	(17) The conditions allowing for exemptions from the ban of use of such substances in cosmetic products should be streamlined <u>without lowering the high level of human health and safety and consumer protection</u> , and their	VS.EC	(17) The conditions allowing for exemptions from the ban of use of such substances in cosmetic products should be streamlined, and their scope should be set out in more detail. In addition, compliance with food safety	VS.EC	
29							

CLEAN	Commission Proposal	V.S.EC	EP Mandate	V.S.EC	Council Mandate	V.S.EC	Draft Agreement
	<p>requirements is not compatible with the scientific and technical developments that allow the development of new substances for use in cosmetic products that are not used or found in food. The compliance with food safety requirements does not enhance the safety of cosmetic products as both categories of products are inherently different. It is, therefore, appropriate to abolish this condition.</p>	<p>scope should be set out in more detail. In addition, compliance with food safety requirements is not compatible with the scientific and technical developments that allow the development of new substances for use in cosmetic products that are not used or found in food. The compliance with food safety requirements does not <u>necessarily</u> enhance the safety of cosmetic products as both categories of products are inherently different. It is, therefore, appropriate to abolish this condition.</p>	<p>requirements is not compatible with the scientific and technical developments that allow the development of new substances for use in cosmetic products that are not used or found in food. The compliance with food safety requirements does not enhance the safety of cosmetic products as both categories of products are inherently different. It is, therefore, appropriate to abolish this condition.</p>				
30	<p>(18) Furthermore, elements to be considered under the availability of suitable alternatives condition should be specified. In particular, it should be provided that the use of alternative substance should result in reduced overall risk to human health and the environment and the substance should provide an equivalent or similar function in a cosmetic product, be available on the market in sufficient quantities, so that it can be technically feasible</p>	<p>(18) Furthermore, <u>the</u> elements to be considered under the availability of suitable alternatives condition should be specified<u>outlined</u>. In particular, it should be provided that the use of <u>any</u> alternative substance should result in reduced overall risk to <u>replace the classified substance is safe for</u> human health and the environment and the substance. <u>The alternative</u> should provide an equivalent or similar function in a cosmetic product, and comparable</p>	<p>(18) Furthermore, elements to be considered under the availability of suitable alternatives condition should be specified<u>outlined</u>. In particular, it should be provided that the use of <u>an</u> alternative substance, <u>a combination of substances, or, where relevant, an alternative technology that replaces the need for the substance</u>, should result in reduced overall risk to be safe for human health and the environment and the substance.</p>				

Recital 18

CLEAN	Commission Proposal	VS.EC EP Mandate	VS.EC Council Mandate	VS.EC Draft Agreement
	<p>and economically viable for businesses and especially for SMEs. In addition, access to the substance should not be restricted by patents or raw material restrictions. It should also be possible to consider the economic aspects, such as costs of reformulation and comparative contribution to overall production costs, as relevant factors in the analysis of the suitability of alternatives.</p>	<p><u>level of efficacy and performance and should</u> be available on the market in sufficient quantities, so that it can be <u>likely to be available in sufficient quantities to meet current demand and has the demonstrated potential to meet expected demands in a reasonable timeframe. Its use should be technically feasible and economically viable feasible</u> for businesses and especially for SMEs. In addition, access to the substance should not be restricted by patents or raw material restrictions. It should also be possible to consider the to sustained production. To assess economic feasibility, the economic aspects, such as costs of reformulation and comparative contribution to overall production costs, as relevant factors in the analysis of the suitability of alternatives, <u>can be considered.</u></p>	<p>The alternative should provide an equivalent or similar a function of a similar product and level of efficacy comparable to the classified substance, be available on the market in sufficient quantities, or likely to be available within a reasonable timeframe, so that it can be technically feasible and economically viable feasible also for businesses and especially for SMEs. In addition, access to the substance should not be restricted by patents or raw material restrictions. It should also be possible to consider the to assess economic feasibility, economic aspects, such as costs of reformulation and comparative contribution to overall production costs, as relevant factors in the analysis of the suitability of alternatives, should be considered.</p>	
Recital 19	(19) In addition, in order to streamline the derogation procedure, the condition that the derogation request is made for a particular use of a product	(19) In addition, in order to streamline the derogation procedure, the condition that the derogation request is made for a particular use of a product	(19) In addition, in order to streamline the derogation procedure, the condition that the derogation request is made for a particular use of a product	
31	(19) In addition, in order to streamline the derogation procedure, the condition that the derogation request is made for a particular use of a product	(19) In addition, in order to streamline the derogation procedure, the condition that the derogation request is made for a particular use of a product	(19) In addition, in order to streamline the derogation procedure, the condition that the derogation request is made for a particular use of a product	

CLEAN	Commission Proposal	VS.EC EP Mandate	VS.EC Council Mandate	VS.EC Draft Agreement
	<p>category with a known exposure should become part of the SCCS assessment criterion. Currently, the scientific committee is already assessing the safety of the substance considering its hazard properties and exposure, (i.e., namely specific use in particular product category), therefore, a separate criterion is redundant.</p>	<p>category with a known exposure should become part of the SCCS assessment criterion. Currently, the scientific committee is already assessing the safety of the substance considering its hazard properties and exposure, (ie, namely specific use in particular product category), therefore, a separate criterion is redundant.</p>	<p>category with a known exposure should become part of the SCCS assessment criterion. Currently, the scientific committee is already assessing the safety of the substance considering its hazard properties and exposure, (i.e., namely specific use in particular product category), therefore, a separate criterion is redundant.</p>	
Recital 20	<p>(20) Due account should be taken of the specific exposure of cosmetic products, which are mainly placed in contact with the external parts of human body (for example epidermis, hair system, nails, external genital organs) and that they are not ingested, inhaled, injected or implanted into the human body. The prohibition triggered by Article 15 of Regulation (EC) No 1223/2009 should cover the substances with CMR harmonised classification under the Regulation (EC) No 1272/2008, where the CMR hazards are not assigned to specific routes of exposure or when they are assigned explicitly to the dermal route of exposure.</p>	<p>(20) Due account should be taken of the specific exposure of cosmetic products, which are mainly placed in contact with the external parts of human body (for example epidermis, hair system, nails, external genital organs) and that they are not ingested, inhaled, injected or implanted into the human body. The prohibition triggered by Article 15 of Regulation (EC) No 1223/2009 should cover the substances with CMR harmonised classification under the Regulation (EC) No 1272/2008, where the CMR hazards are not assigned to specific routes of exposure or when they are assigned explicitly to the dermal route of exposure.</p>	<p>(20) Due account should be taken of the specific exposure of cosmetic products, which are mainly placed in contact with the external parts of human body (for example epidermis, hair system, nails, external genital organs) and that they are not ingested, inhaled, injected or implanted into the human body. The prohibition triggered by Article 15 of Regulation (EC) No 1223/2009 should cover the substances with CMR harmonised classification under the Regulation (EC) No 1272/2008, where the CMR hazards are not assigned to specific routes of exposure or when they are assigned explicitly to the dermal route of exposure.</p>	
32	<p>(20) Due account should be taken of the specific exposure of cosmetic products, which are mainly placed in contact with the external parts of human body (for example epidermis, hair system, nails, external genital organs) and that they are not ingested, inhaled, injected or implanted into the human body. The prohibition triggered by Article 15 of Regulation (EC) No 1223/2009 should cover the substances with CMR harmonised classification under the Regulation (EC) No 1272/2008, where the CMR hazards are not assigned to specific routes of exposure or when they are assigned explicitly to the dermal route of exposure.</p>	<p>(20) Due account should be taken of the specific exposure of cosmetic products, which are mainly placed in contact with the external parts of human body (for example epidermis, hair system, nails, external genital organs) and that they are not ingested, inhaled, injected or implanted into the human body. The prohibition triggered by Article 15 of Regulation (EC) No 1223/2009 should cover the substances with CMR harmonised classification under the Regulation (EC) No 1272/2008, where the CMR hazards are not assigned to specific routes of exposure or when they are assigned explicitly to the dermal route of exposure.</p>	<p>(20) Due account should be taken of the specific exposure of cosmetic products, which are mainly placed in contact with the external parts of human body (for example epidermis, hair system, nails, external genital organs) and that they are not ingested, inhaled, injected or implanted into the human body. The prohibition triggered by Article 15 of Regulation (EC) No 1223/2009 should cover the substances with CMR harmonised classification under the Regulation (EC) No 1272/2008, where the CMR hazards are not assigned to specific routes of exposure or when they are assigned explicitly to the dermal route of exposure.</p>	

CLEAN	Commission Proposal	VS.EC	EP Mandate	VS.EC	Council Mandate	VS.EC	Draft Agreement
	<p>Where the CMR classification of a substance is only associated with oral or inhalation routes of exposure, its use in cosmetic products does not result in the same level of risk for end-users, since oral and inhalation exposure are incidental (for example, cosmetic products used on lips, teeth or mucous membranes of the oral cavity or cosmetic products used in spray are not intended to be ingested or inhaled). Therefore, such substances should not be subject to a prohibition under Article 15 of Regulation (EC) No 1223/2009. However, the fact that a substance used in oral or sprayable cosmetic products is classified as CMR due to its oral or inhalation route of exposure, may raise concerns for human health. In such cases, the Commission should mandate the SCCS to assess the safety of such substances when used in cosmetic products and is to follow up with the appropriate regulatory measures in accordance with Article 31(1) of Regulation (EC) No 1223/2009.</p>	<p>Where the CMR classification of a substance is only associated with oral or inhalation routes of exposure, its use in cosmetic products does not result in the same level of risk for end-users, since oral and inhalation exposure are incidental (for example, cosmetic products used on lips, teeth or mucous membranes of the oral cavity or cosmetic products used in spray are not intended to be ingested or inhaled). Therefore, such substances should not be subject to a prohibition under Article 15 of Regulation (EC) No 1223/2009. However, the fact that a substance used in oral or sprayable cosmetic products is classified as CMR due to its oral or inhalation route of exposure, may raise concerns for human health. In such cases, the Commission should mandate the SCCS to assess the safety of such substances when used in cosmetic products and is to follow up with the appropriate regulatory measures in accordance with Article 31(1) of Regulation (EC) No 1223/2009.</p>	<p>Where the CMR classification of a substance is only associated with oral or inhalation routes of exposure, its use in cosmetic products does not result in the same level of risk for end-users, since oral and inhalation exposure are incidental (for example, cosmetic products used on lips, teeth or mucous membranes of the oral cavity or cosmetic products used in spray are not intended to be ingested or inhaled). Therefore, such substances should not be subject to a prohibition under Article 15 of Regulation (EC) No 1223/2009. However, the fact that a substance used in oral or sprayable cosmetic products is classified as CMR due to its oral or inhalation route of exposure, may raise concerns for human health. In such cases, the Commission should mandate the SCCS to assess the safety of such substances when used in cosmetic products and is to follow up with the appropriate regulatory measures in accordance with Article 31(1) of Regulation (EC) No 1223/2009.</p>				
Recital 21							

CLEAN	Commission Proposal	VS,EC	EP Mandate	VS,EC	Council Mandate	VS,EC	Draft Agreement
<p>(21) Often a substance can also be a constituent of natural complex substances, for example essential oils. In such cases, the prohibition of use in cosmetic products under Article 15 of Regulation (EC) No 1223/2009 is relevant only to the substance as it appears in Part 3 of Annex VI to the Regulation (EC) No 1272/2008. This means that natural complex substances that contain a CMR classified constituent are not subject to the prohibition, except if that natural complex substance is itself listed as CMR substance of category 1A, 1B or 2 in Part 3 of Annex VI to the Regulation (EC) No 1272/2008. Nevertheless, since the harmonised classification of a constituent may raise concerns as to the safety of the natural complex substances when used in cosmetic products, the Commission should mandate the SCCS to assess the impact of such constituent on the safety of natural complex substances, if a safety concern arises, and is to follow up with the appropriate regulatory measures in accordance with</p>	<p>(21) Often a substance can also be a <u>Substances containing more than one constituent which are extracted from plants or plant parts and which are not chemically modified as defined in Article 3, point (40), of Regulation (EC) No 1907/2006, are often referred to as</u> natural complex substances, for example essential oils. In such cases, or <u>'NCS'. They might contain one or more constituents classified as CMR substances, while the overall natural complex substance is not itself classified as a CMR substance. A high level of consumer protection must apply to such substances and should be based on scientific knowledge and the actual conditions of exposure arising from their use in cosmetic products.</u> The prohibition of use in cosmetic products under Article 15 of Regulation (EC) No 1223/2009 is relevant only applies to the substance as it appears in Part 3 of Annex VI to the Regulation (EC) No 1272/2008. His means that Natural complex substances that contain a CMR classified</p>	<p>(21) Often a substance can also be a <u>Article 5 of Regulation (EC) No 1272/2008 provides a specific rule for identification and examination of available information when evaluating the hazardous properties of substances containing more than one constituent which are extracted from plants or plant parts and which are not chemically modified as defined in Article 3, point (40), of Regulation (EC) No 1907/2006.</u> Such substances, often referred to as 'of Natural complex substances', or 'NCS' may contain one or more substances classified as CMR substances; for example essential oils. In such cases, while the overall Natural complex substance is not itself classified as a CMR substance. The prohibition of use in cosmetic products under Article 15 of Regulation (EC) No 1223/2009 is relevant only applies to the substance as it appears in Part 3 of Annex VI to the Regulation (EC) No 1272/2008. This means that Natural complex substances that contain a CMR classified</p>					

CLEAN	Commission Proposal	VS.EC EP Mandate	VS.EC Council Mandate	VS.EC Draft Agreement
Article 31(1) of Regulation (EC) No 1223/2009.	<p>constituent are not subject to the prohibition, except if that <u>where the natural complex substance is itself listed as classified as a</u> CMR substance of category IA, IB or 2 in Part 3 of Annex VI to the Regulation (EC) No 1272/2008. Nevertheless, since the harmonised classification of a constituent may raise concerns <u>concerns as to the about the use and</u> safety of the natural complex substances when used in cosmetic products, the Commission should <u>without delay</u> mandate the SCCS to assess the impact of such constituent on the safety of natural complex substances, if a safety concern arises, and is to follow up with the appropriate such constituents and, where necessary, take regulatory measures in accordance with <u>assessment of such constituent on the safety of natural complex substances, if a safety concern arises, and is to follow up with the appropriate such constituents and, where necessary, take regulatory measures in accordance with</u> <u>the assessment of the SCCS should be timely to ensure predictability for the industry.</u></p>	<p>constituent are not subject to the prohibition, except if that <u>where the natural complex substance is itself listed as classified as a</u> CMR substance of category IA, IB or 2 in Part 3 of Annex VI to the Regulation (EC) No 1272/2008. Nevertheless, since the harmonised classification of a constituent may raise concerns <u>concerns as to the about the use and</u> safety of the natural complex substances when used in cosmetic products, the Commission should <u>without delay</u> mandate the SCCS to assess the impact of such constituent on the safety of natural complex substances, if a safety concern arises, and is to follow up with the appropriate such constituents and, where necessary, take regulatory measures in accordance with <u>assessment of such constituent on the safety of natural complex substances, if a safety concern arises, and is to follow up with the appropriate such constituents and, where necessary, take regulatory measures in accordance with</u> <u>the assessment of the SCCS should be timely to ensure predictability for the industry.</u></p>	<p>constituent are not subject to the prohibition, except if that Natural complex substance is itself listed as CMR substance of category IA, IB or 2 in Part 3 of Annex VI to the Regulation (EC) No 1272/2008. Nevertheless, since the harmonised classification of When a constituent may raise concerns as to the safety of the of a Natural complex substances when used in cosmetic products substance is classified as a CMR (categories 1A, 1B or 2), this raises concerns about its safe use in cosmetics. In such cases, the Commission should mandatestystematically request the SCCS to assess the impact of such constituent on the safety of natural complex substances, if a safety concern arises, and is to follow up with the appropriate such constituents and, where necessary, take regulatory measures in accordance with action under Article 31(1) of Regulation (EC) No 1223/2009.</p>	
Recital 22				
34	(22) When a substance is prohibited or restricted from the use in cosmetic products, the	(22) When a substance is prohibited or restricted from the use in cosmetic products, the	(22) When a substance is prohibited or restricted from the use in cosmetic products, the	

CLEAN	Commission Proposal	V.S.EC EP Mandate	V.S.EC Council Mandate	V.S.EC Draft Agreement
	<p>manufacturers, importers, distributors and responsible persons should be given appropriate time to take necessary measures to reformulate and relabel their products, withdraw from the distribution and destroy the unsold products not complying with the new requirements. Therefore, periods of 12 months for placing and 24 months for making available on the market of cosmetic products containing the substance concerned following the entry into force of the respective amendments to Regulation (EC) No 1223/2009 should be provided.</p>	<p>manufacturers, importers, distributors and responsible persons should be given appropriate time to take necessary measures to reformulate, <u>test</u> and relabel their products, withdraw from the distribution and destroy the unsold products not complying with the new requirements. Therefore, periods of 12 <u>Accordingly, if no derogation request was submitted, a period of 6</u> months for placing and <u>24</u> 15 months for making available on the market of cosmetic products containing the substance concerned following the entry into force of the respective amendments to Regulation (EC) No 1223/2009 should be provided. <u>In case a request for derogation has been submitted and the SCCS has found that the substance is not safe, those deadlines should be shortened to a period of 3 months for placing and 12 months for making available on the market. In case a request for derogation has been submitted and the SCCS has found that the substance is safe, but the request has been refused due to the</u></p>	<p>manufacturers, importers, distributors and responsible persons should be given appropriate time to take necessary measures. To enable manufacturers of cosmetic to reformulate and relabel their products to secure the suitable alternative substances, to carry out tests as part of reformulation process and to proceed with the safety assessment of the final products and their re-labelling and mandatory notifications, they should be provided with appropriate adjustment period. During this period distributors and retailers continue to receive cosmetic products containing the substance subject to regulatory measures. Therefore, they need additional time to withdraw such products from the distribution chain and to destroy the unsold products not complying with the new requirements. Therefore, periods of 12 These actions impact the existing contractual arrangements and requires substantial logistical efforts. Accordingly, a period of</p>	

CLEAN	Commission Proposal	VS.EC EP Mandate	VS.EC Council Mandate	VS.EC Draft Agreement
		<p><u>availability of suitable alternatives, those deadlines should be extended to a period of 24 months for placing and 48 months for making available on the market, under the condition that an up-to-date Cosmetic Product Safety Report (CPSR) remains available at all times.</u></p>	<p>6 months for placing and 24 months for making available on the market of cosmetic products containing the substance concerned following the entry into force of the respective amendments to Regulation (EC) No 1223/2009 should be provided.</p>	
Recital 23	<p>(23) To reduce compliance and administrative burden on businesses active in the cosmetic sector, only one notification of the cosmetic products should be required before placing them on the Union market. The conditions of such notification should apply to a non-discriminatory way to cosmetic products containing nanomaterials and to those cosmetic products which do not contain them. To maintain vigilance on nanomaterials, it should be required that the specific information on nanomaterials used in a cosmetic product is provided in the cosmetic product safety report so that it can be consulted by the competent authorities where the concerns over the potential risk</p>	<p>(23) To reduce compliance and administrative burden on businesses active in the cosmetic sector, only one notification of the cosmetic products <u>to the Commission</u> should be required before placing them on the Union market. The conditions of such notification should apply in a non-discriminatory way to cosmetic products containing nanomaterials and to those cosmetic products which do not contain them. To maintain vigilance on nanomaterials <u>in cosmetic products</u>, it should be required that <u>this notification includes the identification of the nanomaterial including its chemical name (IUPAC) and</u></p>	<p>(23) To reduce compliance and administrative burden on businesses active in the cosmetic sector, only one notification of the cosmetic products should only be required before placing them on the Union market. The conditions of such notification should apply in a non-discriminatory way to cosmetic products containing nanomaterials and to those cosmetic products which do not contain them. To maintain vigilance on nanomaterials, it should be required that the specific information on nanomaterials used in a cosmetic product should continue to be provided by the</p>	

CLEAN	Commission Proposal	VS,EC EP Mandate	VS,EC Council Mandate	VS,EC Draft Agreement
	<p>to human health arise from the use of a particular nanomaterial</p>	<p><u>other descriptors as specified in point 2 of the Preamble to Annexes II to VI, and the specification of the nanomaterial including size of particles, physical and chemical properties, intended to be placed on the market per year. This is necessary so that a safety assessment can be requested by the Commission in case of concerns. It should also be explicitly required that</u>he specific information on nanomaterials used in a cosmetic product is provided in the Cosmetic <u>Products Notification Portal (CPNP) notification and in the cosmetic product safety report</u> so that <u>that are both accessible to the competent authorities and</u> can be consulted by the competent authorities, where the concerns over the potential risk to human health arise from the use of a particular nanomaterial <u>in cosmetic products.</u></p>	<p>responsible persons through the existing process before the product safety report so that it can be consulted by the competent authorities where the concerns over the potential risk to human health arise from the use of a particular nanomaterial is placed on the market.</p>	
Recital 24				
36	<p>(24) In accordance with Regulation (EU) 2019/1020¹, the Commission has developed an information and communication</p>	<p>(24) In accordance with Regulation (EU) 2019/1020¹, the Commission has developed an information and communication</p>	<p>(24) In accordance with Regulation (EU) 2019/1020¹, the Commission has developed an information and communication</p>	

CLEAN	Commission Proposal	VS.EC EP Mandate	VS.EC Council Mandate	VS.EC Draft Agreement
	<p>system for the collection, processing and storing of information on issues relating to the enforcement of Union product legislation, including Regulation (EC) No 1223/2009. In practice this system has replaced the reporting obligation laid down in Article 22 of Regulation (EC) No 1223/2009 requiring the Member States to regularly submit the review and assessment of their surveillance activities to the Commission and other Member States. This reporting obligation should therefore be abolished.</p> <p>1. Regulation (EU) 2019/1020 of the European Parliament and of the Council of 20 June 2019 on market surveillance and compliance of products and amending Directive 2004/42/EC and Regulations (EC) No 765/2008 and (EU) No 305/2011 (OJ L 169, 25.6.2019, p. 1, ELI: http://data.europa.eu/eli/reg/2019/1020/oj)</p>	<p>system for the collection, processing and storing of information on issues relating to the enforcement of Union product legislation, including Regulation (EC) No 1223/2009. In practice this system has replaced the reporting obligation laid down in Article 22 of Regulation (EC) No 1223/2009 requiring the Member States to regularly submit the review and assessment of their surveillance activities to the Commission and other Member States. This reporting obligation should therefore be abolished.</p> <p>1. Regulation (EU) 2019/1020 of the European Parliament and of the Council of 20 June 2019 on market surveillance and compliance of products and amending Directive 2004/42/EC and Regulations (EC) No 765/2008 and (EU) No 305/2011 (OJ L 169, 25.6.2019, p. 1, ELI: http://data.europa.eu/eli/reg/2019/1020/oj)</p>	<p>system for the collection, processing and storing of information on issues relating to the enforcement of Union product legislation, including Regulation (EC) No 1223/2009. In practice this system has replaced the reporting obligation laid down in Article 22 of Regulation (EC) No 1223/2009 requiring the Member States to regularly submit the review and assessment of their surveillance activities to the Commission and other Member States. This reporting obligation should therefore be abolished.</p> <p>1. Regulation (EU) 2019/1020 of the European Parliament and of the Council of 20 June 2019 on market surveillance and compliance of products and amending Directive 2004/42/EC and Regulations (EC) No 765/2008 and (EU) No 305/2011 (OJ L 169, 25.6.2019, p. 1, ELI: http://data.europa.eu/eli/reg/2019/1020/oj)</p>	
Recital 25				
37	<p>(25) Cosmetics are globally traded goods. It is therefore important that the ingredient names present on their labels reflect the current state of scientific and technological</p>	<p>(25) Cosmetics are globally traded goods-and it is therefore important<u>essential</u> that the ingredient names present on their labels reflect the current state of scientific and technological</p>	<p>(25) Cosmetics are globally traded goods-and it is therefore important<u>essential</u> that the ingredient names present on their labels reflect the current state of scientific and technological</p>	

CLEAN	Commission Proposal	VS.EC	EP Mandate	VS.EC	Council Mandate	VS.EC	Draft Agreement
	<p>development. The use of internationally recognised cosmetic ingredient' names is an important factor promoting transparency and facilitating cross-border trade in cosmetics. This Regulation should enable internationally recognised names to be used on the labelling of cosmetic products without any additional regulatory action from the Commission. As a glossary of common ingredient names adopted by the Commission would slow down the process of uptake of the new names, the provision requiring the Commission to adopt such a glossary should be abolished.</p>	<p>development <i>in a timely manner</i>. The use of internationally recognised <i>nomenclature, such as the International Nomenclature of Cosmetic ingredient' names is an important factor promoting <u>Ingredients</u> transparency for consumers, ensures consistency across jurisdictions, and facilitates cross-border trade in cosmetics. INCI names are maintained by the Personal Care Products Council (PCPC) as an international industry standard and are widely recognised by regulators and stakeholders worldwide.</i> This Regulation should enable <i>the direct use of</i> internationally recognised names to be used on the labelling of cosmetic products without any additional regulatory action from the Commission. <i>Where a common ingredient name is not available in INCI, other generally accepted nomenclature should be used, for example names established in recognised international chemical or</i></p>	<p>development in a timely manner. The use of internationally recognised nomenclature, such as the International Nomenclature of Cosmetic ingredient' names is an important factor promoting <u>Ingredients</u> transparency for consumers, ensures consistency across jurisdictions, and facilitates cross-border trade in cosmetics. INCI names are maintained by the Personal Care Products Council (PCPC) as an international industry standard and are widely recognised by regulators and stakeholders worldwide. This Regulation should enable the direct use of internationally recognised namesnomenclature, to be used on the labelling of cosmetic products without any additionalfurther regulatory action from the Commission. Where a common ingredient name is not available in INCI, other generally accepted nomenclature should be used, for example names established in recognised international chemical or</p>				

CLEAN	Commission Proposal	VS.EC EP Mandate	VS.EC Council Mandate	VS.EC Draft Agreement
		<p><u>references, or in other authoritative sources commonly relied upon by industry and regulators. This approach ensures flexibility, avoids unnecessary administrative burden, and guarantees that ingredient names used on cosmetic product labelling remain up to date, internationally coherent, and easily understandable to consumers.</u> As a glossary of common ingredient names adopted by the Commission would slow down the process of uptake of the new names, the provision requiring the Commission to adopt such a glossary should be abolished.</p>	<p>pharmacopoeia references, or in other authoritative sources commonly relied upon by industry and regulators. This approach ensures flexibility, avoids unnecessary administrative burden, and guarantees that ingredient names used on cosmetic product labelling remain up to date, internationally coherent, and easily understandable to consumers. As a glossary of common ingredient names adopted by the Commission would slow down the process of uptake of the new names, the provision requiring the Commission to adopt such a glossary should be abolished. The Commission is encouraged to facilitate access to the internationally recognised nomenclature through digital tools, such as the CosIng ingredients (CosIng) database.</p>	
Recital 25a				
37a		<p><u>(25a) In order to ensure a high level of protection of human health, all operators placing cosmetic products on the Union market, whether offline or online,</u></p>		

CLEAN	Commission Proposal	VS,EC EP Mandate	VS,EC Council Mandate	VS,EC Draft Agreement
		<p><u>should be subject to equivalent obligations and effective enforcement. This is particularly important given the growing sale of cosmetics via online marketplaces, including products originating from third countries that are not subject to the same health and safety requirements. Therefore, it is necessary to require certain labelling information referred to in Article 19 to be clearly and visibly indicated in case of distance sales, including via online marketplaces. This requirement will simplify enforcement of Regulation (EC) No 1223/2009 and thereby contribute to fair competition and a high level of protection of human health. Furthermore, post-market surveillance should be strengthened, notably for online sales and imports, and cosmetic products identified as non-compliant through the Union rapid alert system (Safety Gate) should not be listed or offered for sale. To this end, online platforms should verify the identity of the responsible person before</u></p>		

CLEAN	Commission Proposal	VS,EC EP Mandate	VS,EC Council Mandate	VS,EC Draft Agreement
		<p><u>allowing products to be placed on the Union market, and the online sale, offering for sale and promotion, including via social media, of banned cosmetics, in particular mercury-added products, should be explicitly prohibited.</u></p>		
Recital 26	<p>(26) In line with the Commission's objective to rationalise and simplify reporting requirements and to promote the 'digital by default' principle to support digital transformations, economic operators dealing with EU fertilising products in accordance with Regulation (EU) 2019/1009 should provide a digital contact through which they can be reached, draw up the EU declaration of conformity in electronic form and make it accessible via an internet address or data carrier, and provide authorities, upon request, with all relevant information and documentation in electronic form. Documents and correspondence to and from notified bodies related to conformity assessments of EU</p>	<p>(26) In line with the Commission's objective to rationalise and simplify reporting requirements and to promote the 'digital by default' principle to support digital transformations, economic operators dealing with EU fertilising products in accordance with Regulation (EU) 2019/1009 should provide a digital contact through which they can be reached, draw up the EU declaration <u>authorities and end-users so as to adequately answer any queries</u> from those, draw up the EU declaration of conformity in electronic form and make it accessible via an internet address or data carrier, and provide <u>competent</u> authorities, upon request, with all relevant information and documentation <u>in</u></p>	<p>(26) In line with the Commission's objective to rationalise and simplify reporting requirements and to promote the 'digital by default' principle to support digital transformations, economic operators dealing with EU fertilising products in accordance with Regulation (EU) 2019/1009 should provide a digital contact through which they can be reached, draw up the EU declaration of conformity in electronic form and make it accessible via an internet address or data carrier, and provide authorities, upon request, with all relevant information and documentation in electronic form. Documents and correspondence to and from notified bodies related to conformity assessments of EU</p>	
38				

CLEAN	Commission Proposal	VS.EC EP Mandate	VS.EC Council Mandate	VS.EC Draft Agreement
	<p>fertilising products should also be provided in electronic form. Where a digital label is used, manufacturers should use the same data carrier used for the digital label to provide access to the EU declaration of conformity, to avoid the presence of multiple data carriers on the same product. Where a Digital Product Passport is required for EU fertilising products under other EU legislation, the digital labelling information and the EU declaration of conformity should be provided in that Digital Product Passport.</p>	<p><u>a swift manner</u> in electronic form. Documents and correspondence to and from notified bodies related to conformity assessments of EU fertilising products should also be provided in electronic form <u>in a swift manner</u>. Where a digital label is used, manufacturers should use the same data carrier used for the digital label to provide access to the EU declaration of conformity, to avoid the presence of multiple data carriers on the same product. Where a Digital Product Passport is required for EU fertilising products under other EU legislation, the digital labelling information and the EU declaration of conformity should be provided in that Digital Product Passport.</p>	<p>fertilising products should also be provided in electronic form. Where a digital label is used, manufacturers should use the same data carrier used for the digital label to provide access to the EU declaration of conformity, to avoid the presence of multiple data carriers on the same product. Where a Digital Product Passport is required for EU fertilising products under other EU legislation, the digital labelling information and the EU declaration of conformity should be provided in that Digital Product Passport.</p>	
Recital 26a		<p><u>(26a) Fertiliser products covered by Regulation (EU) 2019/1009 are subject to regulatory requirements and administrative burdens which, when increased, may widen the gap between agricultural production costs in the Union and those in third countries. The</u></p>		
38a				

CLEAN	Commission Proposal	VS,EC EP Mandate	VS,EC Council Mandate	VS,EC Draft Agreement
		<p><u>revision of this Regulation should therefore provide for regulatory simplification for EU fertiliser producers and benefit EU farmers making it easier for operators to enter the EU market and operate beyond national markets to the benefit of the Single Market. Therefore, it should not undermine a high level of consumer, health, environmental protection and risk management. Furthermore, easing sector-specific requirements for fertilising products should not lower the levels of control on these products when they are entering the market, nor should it raise concerns regarding traceability, safety, and reputational impacts.</u></p>		
Recital 26b		<p><u>(26b) Digitalisation of declarations of conformity and technical product information may offer certain advantages. Due regard should be given to cybersecurity, effective and swift enforcement oversight, the availability and interoperability of digital infrastructure, the</u></p>		
38b				

CLEAN	Commission Proposal	VS.EC	EP Mandate	VS.EC	Council Mandate	VS.EC	Draft Agreement
			<u>potential costs of introducing and operating such systems and the diversity of economic operators and national systems.</u>				
Recital 27	<p>(27) Under Regulation (EU) 2019/1009, only micro-organisms listed on a positive list in Annex II to that Regulation may be used as component material in microbial plant biostimulants. The Commission is empowered to add new micro-organisms or strains of micro-organisms to that list after an assessment concluding that none of the strains presents a risk to human, animal or plant health, to safety or to the environment and that it ensures agronomic efficiency. Given the large number of micro-organisms on the market, the assessment and subsequent inclusion of new micro-organisms or strains of micro-organisms to the positive list are lagging scientific progress. The current mechanism slows down the development of microbial plant biostimulants and delays farmers' access to those innovative fertilising products which may stimulate plant</p>	(27)	<p>Under Regulation (EU) 2019/1009, only micro-organisms listed on a positive list in Annex II to that Regulation may be used as component material in microbial plant biostimulants. The Commission is empowered to add new micro-organisms or strains of micro-organisms to that list after an assessment concluding that none of the strains presents a risk to human, animal or plant health, to safety or to the environment and that it ensures agronomic efficiency. Given the large number of micro-organisms on the market, the assessment and subsequent inclusion of new micro-organisms or strains of micro-organisms to the positive list are lagging scientific progress. The current mechanism slows down the development of microbial plant biostimulants and delays farmers' access to those innovative fertilising products which may stimulate plant</p>	(27)	<p>Under Regulation (EU) 2019/1009, only micro-organisms listed on a positive list in Annex II to that Regulation may be used as component material in microbial plant biostimulants. The Commission is empowered to add new micro-organisms or strains of micro-organisms to that list after an assessment concluding that none of the strains presents a risk to human, animal or plant health, to safety or to the environment and that it ensures agronomic efficiency. Given the large number of micro-organisms on the market, the assessment and subsequent inclusion of new micro-organisms or strains of micro-organisms to the positive list are lagging scientific progress. The current mechanism slows down the development of microbial plant biostimulants and delays farmers' access to those innovative fertilising products which may stimulate plant</p>		
	39						

CLEAN	Commission Proposal	VS,EC EP Mandate	VS,EC Council Mandate	VS,EC Draft Agreement
	<p>nutrition processes and thereby reduce the use of traditional fertilisers.</p>	<p>nutrition processes and thereby reduce. <u>A proactive monitoring of the scientific developments and updating of the list of micro-organisms or strains of micro-organisms, as well as of the relevant criteria and methodology is necessary on a continuous basis. Furthermore, all updates should be implemented swiftly. At the same time, the regulatory framework should ensure that the use of traditional and natural fertilisers remains a viable and accessible option, recognising their role alongside microbial solutions, so as not to undermine competitiveness or agricultural production within the Union.</u></p>	<p>nutrition processes and thereby reduce the use of traditional fertilisers.</p>	
Recital 27a		<p><u>(27a) Given the rapid pace of innovation in agricultural biotechnology, it is important that existing regulatory procedures, including the updating of Annexes by the Commission, are applied in a timely and science-based manner, making full use of the possibilities already provided for under Regulation (EU) 2019/1009. This should facilitate</u></p>		
39a				

CLEAN	Commission Proposal	VS,EC	EP Mandate	VS,EC	Council Mandate	VS,EC	Draft Agreement
			<u><i>the timely assessment of new strains, while ensuring that only those meeting Union safety requirements are allowed on the market.</i></u>				
Recital 28	(28) In order to accelerate the assessment of micro-organisms and to open the single market for more microbial plant biostimulants, the power to adopt acts in accordance with Article 290 of the Treaty on the Functioning of the European Union should be delegated to the Commission in respect of Annex II, Part II, component material category (CMC) 7, to Regulation (EU) 2019/1009 to allow the Commission to introduce general criteria and a methodology for the assessment of micro-organisms. Those criteria and the methodology should allow manufacturers and notified bodies to demonstrate and verify that micro-organisms used in microbial plant biostimulants, other than those listed in CMC 7, do not present a risk to human, animal or plant health, to safety or to the	(28) <u><i>The wider use of microbial plant biostimulants can improve nutrient-use efficiency and soil health, thereby fostering the development of sustainable while highly productive agriculture.</i></u> In order to accelerate the assessment of micro-organisms and to open the single market for more microbial plant biostimulants, the power to adopt acts in accordance with Article 290 of the Treaty on the Functioning of the European Union should be delegated to the Commission in respect of Annex II, Part II, component material category (CMC) 7, to Regulation (EU) 2019/1009 to allow the Commission to introduce general criteria and a methodology for the assessment of micro-organisms. Those criteria and the methodology should allow manufacturers to demonstrate and notified bodies to demonstrate verify that micro-organisms used in microbial plant biostimulants, other than those listed in CMC 7, do not present a risk to human, animal or plant	(28) <u><i>The wider use of microbial plant biostimulants can improve nutrient-use efficiency and soil health, thereby fostering the development of sustainable while highly productive agriculture.</i></u> In order to accelerate the assessment of micro-organisms and to open the single market for more microbial plant biostimulants, the power to adopt acts in accordance with Article 290 of the Treaty on the Functioning of the European Union should be delegated to the Commission in respect of Annex II, Part II, component material category (CMC) 7, to Regulation (EU) 2019/1009 to allow the Commission to introduce general criteria and a methodology for the assessment of micro-organisms. Those criteria and the methodology should allow manufacturers to demonstrate and notified bodies to demonstrate verify that micro-organisms used in microbial plant biostimulants, other than those listed in CMC 7, do not present a risk to human, animal or plant	(28) In order to accelerate the assessment of micro-organisms and to open the single market for more microbial plant biostimulants, the power to adopt acts in accordance with Article 290 of the Treaty on the Functioning of the European Union should be delegated to the Commission in respect of Annex II, Part II, component material category (CMC) 7, to Regulation (EU) 2019/1009 to allow the Commission to introduce general criteria and a methodology for the assessment of micro-organisms. Those criteria and the methodology should allow manufacturers to demonstrate and notified bodies to demonstrate verify that micro-organisms used in microbial plant biostimulants, other than those listed in CMC 7, do not present a risk to human, animal or plant			
40			Those criteria and the methodology should <u><i>reflect the most recent scientific</i></u>				

CLEAN	Commission Proposal	VS,EC EP Mandate	VS,EC Council Mandate	VS,EC Draft Agreement
<p>environment and ensure agronomic efficiency. In order to refine and validate the criteria and methodology to be introduced, it is of particular importance that the Commission carry out appropriate consultations during its preparatory work, including at expert level, and that those consultations be conducted in accordance with the principles laid down in the Interinstitutional Agreement of 13 April 2016 on Better Law-Making¹. In particular, to ensure equal participation in the preparation of delegated acts, the European Parliament and the Council receive all documents at the same time as Member States' experts, and their experts systematically have access to meetings of Commission expert groups dealing with the preparation of delegated acts.</p> <p>1. OJ L 123, 12.5.2016, p. 1, ELI: http://data.europa.eu/eli/agree_interinstit/2016/512/oj.</p>	<p><u>developments and</u> allow manufacturers and notified bodies, to demonstrate and verify that micro-organisms used in microbial plant biostimulants, other than those listed in CMC 7, do not present a risk to human, animal or plant health, to safety or to the environment and ensure agronomic efficiency <u>while taking into account EU competitiveness and production. Verifying compliance of micro-organisms with the criteria and methodology set under this Regulation requires specific competences, scientific and technical knowledge from the conformity assessment bodies. Those competences should be therefore scrutinized by the national bodies in charge of accreditation and notification, as well as aspects such as impartiality and professional integrity.</u> In order to refine and validate the criteria and methodology to be introduced, it is of particular importance that the Commission <u>involve the relevant scientific bodies, in particular the European Food Safety Agency,</u></p>	<p>health, to safety or to the environment and ensure agronomic efficiency. This assessment requires special expertise for which conformity assessment bodies will need to be specifically accredited and notified. The national bodies responsible for accreditation and notification should carefully verify the conformity assessment bodies' technical and scientific competence in the assessment of micro-organisms and that they meet the stringent requirements for notified bodies set out by the Regulation, including independence, objectivity, impartiality and professional integrity. In order to refine and validate the criteria and methodology to be introduced, it is of particular importance that the Commission should invite a scientific body, either EFSA or the JRC, to review and contribute and carry out appropriate consultations during its preparatory work, including at expert level, and that those consultations be conducted in accordance with the principles laid</p>		

CLEAN	Commission Proposal	VS,EC EP Mandate	VS,EC Council Mandate	VS,EC Draft Agreement
	<p><i>the JRC and the ECHA, to review and contribute, and also</i> carry out appropriate consultations during its preparatory work, including at expert level, and that. <i>It is essential that the Commission preserves strong interinstitutional cooperation, while engaging in</i> those consultations, <i>which should</i> be conducted in accordance with the principles laid down in the Interinstitutional Agreement of 13 April 2016 on Better Law-Making¹. In particular, to ensure equal participation in the preparation of delegated acts, the European Parliament and the Council receive all documents at the same time as Member States' experts, and their experts systematically have access to meetings of Commission expert groups dealing with the preparation of delegated acts.</p> <p>1. OJ L 123, 12.5.2016, p. 1, ELI: http://data.europa.eu/eli/agree_interinstit/2016/512/oj</p>	<p><i>the JRC and the ECHA, to review and contribute, and also</i> carry out appropriate consultations during its preparatory work, including at expert level, and that. <i>It is essential that the Commission preserves strong interinstitutional cooperation, while engaging in</i> those consultations, <i>which should</i> be conducted in accordance with the principles laid down in the Interinstitutional Agreement of 13 April 2016 on Better Law-Making¹. In particular, to ensure equal participation in the preparation of delegated acts, the European Parliament and the Council receive all documents at the same time as Member States' experts, and their experts systematically have access to meetings of Commission expert groups dealing with the preparation of delegated acts.</p> <p>1. OJ L 123, 12.5.2016, p. 1, ELI: http://data.europa.eu/eli/agree_interinstit/2016/512/oj</p>	<p>down in the Interinstitutional Agreement of 13 April 2016 on Better Law-Making¹. In particular, to ensure equal participation in the preparation of delegated acts, the European Parliament and the Council receive all documents at the same time as Member States' experts, and their experts systematically have access to meetings of Commission expert groups dealing with the preparation of delegated acts.</p> <p>1. OJ L 123, 12.5.2016, p. 1, ELI: http://data.europa.eu/eli/agree_interinstit/2016/512/oj</p>	
Recital 29				
41	(29) Where the Commission makes use of its empowerment to amend the component material	(29) Where the Commission makes use of its empowerment to amend the component material	(29) Where the Commission makes use of its empowerment to amend the component material	

CLEAN	Commission Proposal	VS.EC EP Mandate	VS.EC Council Mandate	VS.EC Draft Agreement
<p>categories in Annex II to Regulation (EU) 2019/1009, it may currently only do so via separate delegated acts in respect of each component material category. Considering the need to introduce additional materials to the various component material categories in the future and the constant technical and scientific progress in the fertilising product sector, there is a frequent need to amend the different component material categories. In some cases, for instance where a new raw material may be allowed in multiple CMCs, the Commission would introduce the same change in all relevant CMCs, each of them covered by a different delegated act. To speed up the adoption of the respective delegated acts, the Commission should be allowed to amend several component material categories by one delegated act.</p>	<p>categories in Annex II to Regulation (EU) 2019/1009, it may currently only do so via separate delegated acts in respect of each component material category. Considering the need to introduce additional materials to the various component material categories in the future and the constant technical and scientific progress in the fertilising product sector, there is a frequent need to amend the different component material categories. In some cases, for instance where a new raw material may be allowed in multiple CMCs, the Commission would introduce the same change in all relevant CMCs, each of them covered by a different delegated act. To speed up the adoption of the respective delegated acts, the Commission should be allowed to amend several component material categories by one delegated act.</p>	<p>categories in Annex II to Regulation (EU) 2019/1009, it may currently only do so via separate delegated acts in respect of each component material category. Considering the need to introduce additional materials to the various component material categories in the future and the constant technical and scientific progress in the fertilising product sector, there is a frequent need to amend the different component material categories. In some cases, for instance where a new raw material may be allowed, or similar requirements be introduced or amended in multiple CMCs, the Commission would introduce the same change in all relevant CMCs, each of them covered by a need to make these changes by adopting different delegated acts. To speed up the adoption of the respective delegated acts development of Regulation (EU) 2019/1009 in such cases, the Commission should be allowed to amend several component material categories by make all related amendments in one delegated act.</p>		

CLEAN	Commission Proposal	VS,EC	EP Mandate	VS,EC	Council Mandate	VS,EC	Draft Agreement
Recital 29a		<p>(29a) <u>Regulation (EU) 2019/1009 aims to facilitate the placing on the internal market and free movement of safe fertilising products, while supporting the recycling of nutrients and the circular use of raw materials and ensuring a high level of protection for human, animal and plant health and the environment. The effective application thereof depends, among other things, on the recognition of end points for constituent materials derived from animal by-products within the meaning of Regulation (EC) No 1069/2009. Delegated Regulation (EU) 2023/1605 has established certain end points for animal by-products intended for fertiliser applications. However, the processing parameters and risk mitigation measures included therein are largely derived from frameworks for feed hygiene and the prevention of feed fraud and do not always reflect the distinct exposure pathways and risk profiles of fertiliser products applied to the soil and water.</u></p>					
41a							

CLEAN	Commission Proposal	VS.EC	EP Mandate	VS.EC	Council Mandate	VS.EC	Draft Agreement
Recital 30	<p>(30) Chemical substances, on their own or in mixtures, if manufactured or imported in quantities above 1 tonne per company per year, need to be registered in accordance with Regulation (EC) No 1907/2006, with information requirements depending on the actual volume. Regulation (EU) 2019/1009, going beyond the requirements of Regulation (EC) No 1907/2006, requires that all substances used in an EU fertilising products, regardless of the quantity in which they are manufactured or imported, are registered, as a minimum, with the information requirements set out by Regulation (EC) No 1907/2006 for substances manufactured or imported in quantities of 10 to 100 tonnes per company per year, together with a chemical safety report covering their use in a fertilising product, in accordance with Article 14 of that Regulation. Those extensive information requirements might prevent manufacturers, especially small and medium-sized enterprises, from using substances</p>	<p>(30) Chemical <u>Regulation (EU) 2019/1009 introduced additional registration requirements for substances, on their own or in mixtures, if manufactured or imported in EU fertilising products, going beyond those set out in Regulation (EC) No 1907/2006. In order to ensure proportionality while maintaining a high level of protection of human health and the environment, it is appropriate to align the registration requirements for substances used in EU fertilising products with those set out in Regulation (EC) No 1907/2006, taking into account the relevant tonnage thresholds and information requirements. At the same time, for substances with particularly hazardous properties, including those classified under Regulation (EC) No 1272/2008 as carcinogenic, mutagenic, toxic for reproduction, endocrine disrupting or persistent, bioaccumulative and toxic, specific information requirements should apply even where such</u></p>	<p>(30) Chemical substances, on their own or in mixtures, if manufactured or imported in quantities above 1 tonne per company per year, need to be registered in accordance with Regulation (EC) No 1907/2006, with information requirements depending on the actual volume. Regulation (EU) 2019/1009, going beyond the requirements of Regulation (EC) No 1907/2006, requires that all substances used in an EU fertilising products, regardless of the quantity in which they are manufactured or imported, are registered, as a minimum, with the information requirements set out by Regulation (EC) No 1907/2006 for substances manufactured or imported in quantities of 10 to 100 tonnes per company per year, together with a chemical safety report covering their use in a fertilising product, in accordance with Article 14 of that Regulation. Those extensive information requirements might prevent manufacturers, especially small and medium-sized enterprises, from using substances</p>				

CLEAN	Commission Proposal	VS.EC EP Mandate	VS.EC Council Mandate	VS.EC Draft Agreement
	<p>that are not yet registered according to those requirements or force them to place their products only on national markets according to national rules. For the sake of proportionality, and considering the general obligation of manufactures and importers of substances and EU fertilising products under Regulation (EC) No 1907/2006 and Regulation (EU) 2019/1009, respectively, to ensure the safety of the products that they place on the market, registration of substances used in EU fertilising products should only follow the requirements, including the relevant gradations, set out in Regulation (EC) No 1907/2006.</p>	<p><i>substances are used in low quantities above 1 tonne per company per year, need to be registered in fertilising products.</i> <i><u>This approach ensures that safety-relevant information is available, while avoiding unnecessary burdens for substances presenting lower risks.</u></i> <i><u>The information provided should be based on available data, including the use of alternative methods and adaptations in</u></i> accordance with Regulation (EC) No 1907/2006, <i>with information requirements depending on the actual volume and should, wherever relevant and in last resort, avoid unnecessary testing, in particular on vertebrate animals.</i> <i>Regulation (EU) 2019/1009, going beyond the requirements of Regulation (EC) No 1907/2006, requires that all substances used in an EU fertilising products, regardless of the quantity in which they are manufactured or imported, are registered, as a minimum, with the information requirements set out by Regulation (EC) No 1907/2006 for substances manufactured or</i></p>	<p>that are not yet registered according to those requirements or force them to place their products only on national markets according to national rules. For the sake of proportionality, and considering the general obligation of manufactures and importers of substances and EU fertilising products under Regulation (EC) No 1907/2006 and Regulation (EU) 2019/1009, respectively, to ensure the safety of the products that they place on the market, registration of substances used in EU fertilising products should only follow the requirements, including the relevant gradations, set out in Regulation (EC) No 1907/2006.</p>	

CLEAN	Commission Proposal	VS.EC EP Mandate	VS.EC Council Mandate	VS.EC Draft Agreement
		<p>imported in quantities of 10 to 100 tonnes per company per year, together with a Chemical safety report covering their use in a fertilising product, in accordance with Article 14 of that Regulation. Those extensive information requirements might prevent manufacturers, especially small and medium-sized enterprises, from using substances that are not yet registered according to those requirements or force them to place their products only on national markets according to national rules. For the sake of proportionality, and considering the general obligation of manufactures and importers of substances and EU assessments <u>should be limited to relevant exposure scenarios related to the agronomic use of fertilising products under Regulation (EC) No 1907/2006 and Regulation (EU) 2019/1009, respectively, to ensure the safety of the products that they place on the and the environment. Such an approach ensures a balanced framework that supports innovation and market access, while safeguarding</u></p>		

CLEAN	Commission Proposal	VS,EC	EP Mandate	VS,EC	Council Mandate	VS,EC	Draft Agreement
			<i>a high level of protection and legal certainty for operators; registration of substances used in EU fertilising products should only follow the requirements, including the relevant gradations, set out in Regulation (EC) No 1907/2006.</i>				
Recital 31							
43	(31) To ensure a smooth and effective transition, to minimise disruptions, and to provide a reasonable timeframe for economic operators and authorities to adjust to the new requirements, the application of the amendments to Regulation (EU) 2019/1009 concerning digitalisation should be deferred.	(31)	To ensure a smooth and effective transition, to minimise disruptions, and to provide a reasonable timeframe for economic operators and authorities to adjust to the new requirements, the application of the amendments to Regulation (EU) 2019/1009 concerning digitalisation should be deferred.	(31)	To ensure a smooth and effective transition, to minimise disruptions, and to provide a reasonable timeframe for economic operators and authorities to adjust to the new requirements, the application of the amendments to Regulation (EU) 2019/1009 concerning digitalisation should be deferred.		
Recital 32							
44	(32) In order to enable economic operators to supply stock of products that have been placed on the market before the date of application of the amendments to Regulation (EU) 2019/1009 concerning digitalisation, it is necessary to provide for reasonable transitional arrangements that do not impede	(32)	In order to enable economic operators to supply stock of products that have been placed on the market before the date of application of the amendments to Regulation (EU) 2019/1009 concerning digitalisation, it is necessary to provide for reasonable transitional arrangements that do not impede	(32)	In order to enable economic operators to supply stock of products that have been placed on the market before the date of application of the amendments to Regulation (EU) 2019/1009 concerning digitalisation, it is necessary to provide for reasonable transitional arrangements that do not impede		

	CLEAN	Commission Proposal	vs.EC	EP Mandate	vs.EC	Council Mandate	vs.EC	Draft Agreement
		the making available on the market of products that have been placed on the market in accordance with that Regulation in its version applicable before that date.	vs.EC	the making available on the market of products that have been placed on the market in accordance with that Regulation in its version applicable before that date.	vs.EC	the making available on the market of products that have been placed on the market in accordance with that Regulation in its version applicable before that date.		
Recital 33								
45	(33) Regulations (EC) No 1272/2008, (EC) No 1223/2009 and (EU) 2019/1009 should therefore be amended accordingly,	(33) Regulations (EC) No 1272/2008, (EC) No 1223/2009 and (EU) 2019/1009 should therefore be amended accordingly,	vs.EC	(33) Regulations (EC) No 1272/2008, (EC) No 1223/2009 and (EU) 2019/1009 should therefore be amended accordingly,	vs.EC	(33) Regulations (EC) No 1272/2008, (EC) No 1223/2009 and (EU) 2019/1009 should therefore be amended accordingly,		
Formula								
46	HAVE ADOPTED THIS REGULATION:	HAVE ADOPTED THIS REGULATION:		HAVE ADOPTED THIS REGULATION:		HAVE ADOPTED THIS REGULATION:		
Article 1								
47	Article 1 Amendments to Regulation (EC) No 1272/2008	Article 1 Amendments to Regulation (EC) No 1272/2008		Article 1 Amendments to Regulation (EC) No 1272/2008		Article 1 Amendments to Regulation (EC) No 1272/2008		
Article 1, first paragraph								
48	Regulation (EC) No 1272/2008 is amended as follows:	Regulation (EC) No 1272/2008 is amended as follows:		Regulation (EC) No 1272/2008 is amended as follows:		Regulation (EC) No 1272/2008 is amended as follows:		
Article 1, first paragraph, point (1)								
49	(1) in Article 2, the following point is added:	(1) in Article 2, the following point is added:		(1) in Article 2, the following point is added:		(1) in Article 2, the following point is added:		
Article 1, first paragraph, point (1), amending provision, numbered paragraph (42)								
50	,	,		,		,		

CLEAN	Commission Proposal	VS,EC	EP Mandate	VS,EC	Council Mandate	VS,EC	Draft Agreement
42.	“digital contact” means any up-to-date and accessible online communication channel through which a supplier can be reached or engaged without the need to register or to download an application.;	42.	“digital contact” means any up-to-date <u>easily and freely</u> accessible online communication channel <u>such as email addresses or a weblink</u> through which a supplier can be reached or engaged without the need to register or to download <u>or use</u> an application.;	42.	“digital contact” means any up-to-date and freely accessible online communication channel such as email addresses through which a supplier can be reached or engaged without the need to register or to download an application or use additional applications specific to the supplier. ;		
Article 1, first paragraph, point (1), amending provision, numbered paragraph (42a)							
50a		<u>42a. (1a) In article 5, the following paragraph is inserted:</u> <u>‘3a. By 18 months from the date of the entering into force of this Amending Regulation, the Commission shall carry out an assessment on whether further specific reductions of mandatory label elements should apply to packages between 10 and 125 ml.’</u>					
Article 1, first paragraph, point (2)							
51	(2) in Article 17(1), point (a) is replaced by the following:	(2) in Article 17(1) is <u>amended as follows:</u> <u>(a)</u> point (a) is replaced by the following:	(2) in Article 17(1), point (a) is replaced by the following:				
Article 1, first paragraph, point (2), amending provision, numbered paragraph (a)							
52							

	CLEAN	Commission Proposal	VS.EC	EP Mandate	VS.EC	Council Mandate	VS.EC	Draft Agreement
	(a) the name, address and digital contact of the suppliers.;	(a) the name, address, telephone number and digital contact of the supplier supplier .;				(a) the name, address, and digital contact of the supplier supplier(s) .;		
Article 1, first paragraph, point (2), amending provision, numbered paragraph (aa)								
52a				<u>2a</u> in Article 17(1) the following subparagraph is added: <u>'The telephone number in point (a) of the first subparagraph may be omitted from the label if such telephone number is directly available through the digital contact.'</u>				
Article 1, first paragraph, point (3)								
53	(3) in Article 25(6), the third subparagraph is replaced by the following:	(3) in Article 25(6), the third subparagraph is replaced by the following:				(3) in Article 25(6), the third subparagraph is replaced by the following:		
Article 1, first paragraph, point (3), amending provision, first paragraph								
54	The label shall also include the product identifier referred to in Article 18 and the name, address and digital contact of the supplier of the mixture.;	The label shall also include the product identifier referred to in Article 18 and the name, address and digital contact of the <u>supplier(s)</u> of the mixture <u>and the telephone number, unless this telephone number is directly available through the digital contact. Without prejudice to the deadline established in Article</u>				The label shall also include the product identifier referred to in Article 18 and the name, address, and digital contact of the supplier(s) of the mixture, and the telephone number, unless this telephone number is immediately available through the digital contact. ;		

CLEAN	Commission Proposal	VS,EC	EP Mandate	VS,EC	Council Mandate	VS,EC	Draft Agreement
			<u>61(8), the inclusion or change to the digital contact may be added or updated at any time or during the supplier's regular label update cycles;</u>				
Article 1, first paragraph, point (4)							
55	(4) in Article 29, paragraph 2 is replaced by the following:	(4) in Article 29, paragraph 2 is paragraphs 1 and 2 are replaced by the following:		(4) in Article 29, paragraph 2 is replaced by the following:			
Article 1, first paragraph, point (4a)							
55b		<u>1. Where the packaging of a substance or a mixture is either in such a shape or form or is so small that it is impossible to meet the requirements laid down in of Article 31 for a label in the languages of the Member State in which the substance or mixture is placed on the market, the label elements set out in Article 17(1) shall be provided in accordance with section 1.5.1 of Annex I.;</u> "					
Article 1, first paragraph, point (4), amending provision, numbered paragraph (2)							
56	The label elements set out in Article 17(1) may be reduced in accordance with the rules set out in section 1.5.2 of Annex I.;	2. The label elements set out in Article 17(1) may be reduced in accordance with the rules set out in section 1.5.2 of Annex I.		2. Where the packaging of a substance or a mixture is either in such a shape or form or is so small that it is impossible to			

CLEAN	Commission Proposal	VS.EC	EP Mandate	VS.EC	Council Mandate	VS.EC	Draft Agreement
		<p><i>where: a) the content of the packaging of a substance or a mixture does not exceed the quantities indicated in section 1.5.2. of Annex I; and b) the packaging is either in such a shape or form or is too small in size to allow for a full reference to all the elements referred to in Article 31 in all the languages of the Member State in which the substance or mixture is placed on the market.”</i></p>	<p>meet the requirements of Article 31 for a label in the languages of the Member State in which the substance or mixture is placed on the market, the label elements set out in Article 17(1) information may be reduced in accordance with the rules set out in section 1.5.2 of Annex I.;</p>				
56a	Article 1, first paragraph, point (4a), amending provision, numbered paragraph (2a)	<p><i>4a. In article 29, the following paragraph is inserted: 2a. By way of derogation from Article 17(1) and Article 25(6), the label elements of ink cartridges may be reduced in accordance with the rules set out in 1.5.2.5a of Annex I. For the purpose of this paragraph, ‘ink cartridge’ means a replaceable unit that holds ink and which must be inserted into a printer during printing.</i></p>					
	Article 1, first paragraph, point (5)						

	CLEAN	Commission Proposal	VS,EC	EP Mandate	VS,EC	Council Mandate	VS,EC	Draft Agreement
57	(5) in Article 30, paragraph 1 is replaced by the following:	(5) in Article 30, paragraph 1 is replaced by the following:	VS,EC	(5) in Article 30, paragraph 1 is replaced by the following:	VS,EC	(5) in Article 30, paragraph 1 is replaced by the following:	VS,EC	
	Article 1, first paragraph, point (5), amending provision, numbered paragraph (1)	Article 1, first paragraph, point (5), amending provision, numbered paragraph (1)						
58	1. In the event of a change regarding the classification or labelling of a substance or a mixture, which results in the addition of a new hazard class or in a more severe classification, or which requires new supplemental information on the label in accordance with Article 25, the supplier of that substance or that mixture shall ensure that the label is updated without undue delay after the results of the new evaluation referred to in Article 15(4) are obtained by, or communicated to, that supplier.;	1. In the event of a change regarding the classification or labelling of a substance or a mixture, which results in the addition of a new hazard class or in a more severe classification, or which requires new supplemental information on the label in accordance with Article 25, the supplier of that substance or that mixture shall ensure that the label is updated without undue delay and in any event no later than eighteen months after the results of the new evaluation referred to in Article 15(4) are obtained by, or communicated to, that supplier. <i>With a view to complete the changes to the labelling without undue delay, suppliers shall cooperate in accordance with Article 4(9) and inform their direct downstream users about the results of the new evaluation as referred to in subparagraph 1 in accordance with the applicable requirements of Regulation (EC)</i>		1. In the event of a change regarding the classification or labelling of a substance or a mixture, which results in the addition of a new hazard class or in a more severe classification, or which requires new supplemental information on the label in accordance with Article 25, the supplier of that substance or that mixture shall ensure that the label is updated without undue delay and in any event no later than 12 months after the results of the new evaluation referred to in Article 15(4) are obtained by, or communicated to, that supplier.;		1. In the event of a change regarding the classification or labelling of a substance or a mixture, which results in the addition of a new hazard class or in a more severe classification, or which requires new supplemental information on the label in accordance with Article 25, the supplier of that substance or that mixture shall ensure that the label is updated without undue delay and in any event no later than 12 months after the results of the new evaluation referred to in Article 15(4) are obtained by, or communicated to, that supplier.;		

CLEAN	Commission Proposal	VS,EC	EP Mandate	VS,EC	Council Mandate	VS,EC	Draft Agreement
		1907/2006. Suppliers may apply the new or updated classifications and adapt the labelling accordingly on a voluntary basis before the expiry of the eighteen months period, in order to ensure a high level of protection of human health and the environment and to provide sufficient flexibility;					
59	Article 1, first paragraph, point (6) (6) in Article 31, paragraph 3 is replaced by the following:	(6) in Article 31, paragraph 3 is replaced by the following:	(6) in Article 31, paragraph 3 is replaced by the following:	(6) in Article 31, paragraph 3 is replaced by the following:			
60	Article 1, first paragraph, point (6), amending provision, numbered paragraph (3) 3. The label elements referred to in Article 17(1) shall be clearly and indelibly marked. They shall stand out clearly from the background and they shall be of such a size and be spaced in such a way as to be easily read;	3. The label elements referred to in Article 17(1) shall be clearly and indelibly marked. They shall stand out clearly from the background and they shall be of such a size and be spaced in such a way as to be easily read. They shall be formatted in accordance with section 1.2.1 of Annex I;	3. The label elements referred to in Article 17(1) shall be clearly and indelibly marked. They shall stand out clearly from the background and they shall be of such a size and be spaced in such a way as to be easily read;	3. The label elements referred to in Article 17(1) shall be clearly and indelibly marked. They shall stand out clearly from the background and they shall be of such a size and be spaced in such a way as to be easily read;			
	Article 1, first paragraph, point (7)						

CLEAN	Commission Proposal	VS.EC	EP Mandate	VS.EC	Council Mandate	VS.EC	Draft Agreement
61	(7) Article 48 is replaced by the following:	(7)	Article 48 is replaced by the following:	(7)	Article 48 is replaced by the following:		
	Article 1, first paragraph, point (7), amending provision, first paragraph						
62	Article 48		Article 48		Article 48		
	Article 1, first paragraph, point (7), amending provision, second paragraph						
63	Advertisement		Advertisement		Advertisement		
	Article 1, first paragraph, point (7), amending provision, numbered paragraph (1)						
64	1. Any advertisement to the general public for a substance or a mixture classified as hazardous or referred to in Part 2 of Annex II shall include the sentence: 'Always read the label and product information before use.'	1.	Any advertisement to the general public for a substance or a mixture classified as hazardous or referred to in Part 2 of Annex II shall <u>always</u> include the sentence: 'Always read the label and product information before use.'; <u>and shall also include one of the following:</u> <u>a) the applicable hazard pictogram(s); or b) the relevant signal word in accordance with Article 20.</u>	1.	Any advertisement to the general public for When advertising a substance or a mixture classified as hazardous or referred to in Part 2 of Annex II, the advertisement shall include the label and product information before use. only be made if the advertisement includes at least one of any of the following:		
	Article 1, first paragraph, point (7), amending provision, numbered paragraph (1a)						
64a		<u>1a.</u>	<u>The first subparagraph shall not apply to advertisements to professional users for substances or mixtures intended for use in the course of their industrial or professional activities, provided that the</u>				

CLEAN	Commission Proposal	VS,EC	EP Mandate	VS,EC	Council Mandate	VS,EC	Draft Agreement
			<i>advertisement is not targeted at or directly made available to the general public.</i>				
Article 1, first paragraph, point (7), amending provision, numbered paragraph (2), first subparagraph							
65	2. Any advertisement for a substance or a mixture classified as hazardous shall not contain statements that are not allowed to appear on the label or packaging of that substance or mixture in accordance with Article 25(4). ⁷	2. Any advertisement for a substance or a mixture classified as hazardous shall not contain statements that are not allowed to appear on the label or packaging of that substance or mixture in accordance with Article 25(4). ⁷	2. Any advertisement for a substance or a mixture classified as hazardous shall not contain statements that are not allowed to appear on the label or packaging of that substance or mixture in accordance with Article 25(4). ⁷	2. Any advertisement for a substance or a mixture classified as hazardous shall not contain statements that are not allowed to appear on the label or packaging of that substance or mixture in accordance with Article 25(4). ⁷	2. Any advertisement for a substance or a mixture classified as hazardous, the advertisement shall only be made if the advertisement does not contain statements that are not allowed to appear on the label or packaging of that substance or mixture in accordance with Article 25(4). ⁷		
Article 1, first paragraph, point (7), amending provision, numbered paragraph (2), second subparagraph							
66	;	;	;	;	;	;	
Article 1, first paragraph, point (8)							
67	(8) Article 48a is replaced by the following:	(8) Article 48a is replaced by the following:	(8) Article 48a is replaced by the following:	(8) Article 48a is replaced by the following:	(8) Article 48a is replaced by the following:	(8) Article 48a is replaced by the following:	
Article 1, first paragraph, point (8), amending provision, first paragraph							
68	Article 48a	Article 48a	Article 48a	Article 48a	Article 48a	Article 48a	
Article 1, first paragraph, point (8), amending provision, second paragraph							
69	Distance sales offers	Distance sales offers	Distance sales offers	Distance sales offers	Distance sales offers	Distance sales offers	
Article 1, first paragraph, point (8), amending provision, third paragraph							

	CLEAN	Commission Proposal	VS,EC	EP Mandate	VS,EC	Council Mandate	VS,EC	Draft Agreement
70	When substances or mixtures are placed on the market for the general public through distance sales, the offer shall clearly and visibly indicate the label elements referred to in Article 17.'	<p><u>1.</u> When substances or mixtures are placed on the market for the general public through distance sales, the offer shall clearly and visibly indicate the label elements referred to in Article 17;</p> <p><u>2. Paragraph 1 also applies to distance sales offers to professional users for substances or mixtures intended for use in the course of their industrial or professional activities, if the offer allows a member of the general public to conclude a distance contract as defined in Article 2, point (7) of Directive 2011/83/EU;'</u></p>	VS,EC	<p><u>1.</u> When substances or mixtures are placed on the market for the general public through distance sales, the offer shall clearly and visibly indicate the label elements referred to in Article 17.'</p>	VS,EC			
	Article 1, first paragraph, point (8), amending provision, fourth paragraph							
71	;	;		;		;		,
	Article 1, first paragraph, point (9)							
72	(9) Article 61 is amended as follows:	(9) Article 61 is amended as follows:		(9) Article 61 is amended as follows:	(9) Article 61 is amended as follows:	(9) Article 61 is amended as follows:		
	Article 1, first paragraph, point (9)(a)							
73	(a) paragraph 8 is replaced by the following:	(a) paragraph 8 is replaced by the following:		(a) paragraph 8 is replaced by the following:	(a) paragraph 8 is replaced by the following:	(a) paragraph 8 is replaced by the following:		
	Article 1, first paragraph, point (9)(a), amending provision, numbered paragraph (8), first subparagraph							
74	,	,		,		,		,

CLEAN	Commission Proposal	VS.EC EP Mandate	VS.EC Council Mandate	VS.EC Draft Agreement
<p>8. Substances and mixtures which have been classified, labelled and packaged in accordance with Article 18(3) as applicable on 9 December 2024 and which were placed on the market before 1 January 2027 shall not be required to be classified, labelled and packaged in accordance with this Regulation as amended by Regulation (EU) 2024/2865 of the European Parliament and of the Council until 1 January 2029.’</p>	<p>8. Substances and mixtures which have been classified, labelled and packaged in accordance with Article 18(3) as applicable on 9 December 2024 and which were placed on the market before 1 January 2027 shall not be required to be classified, labelled and packaged in accordance with this Regulation as amended by Regulation (EU) 2024/2865 of the European Parliament and of the Council until 1 January 2029. <u>Substances and mixtures which have been classified, labelled and packaged in accordance with Article 31(3) and section 1.2.1 of Annex I as applicable on 9 December 2024 and which were placed on the market before 1 January 2028 shall not be required to be classified, labelled and packaged in accordance with this Regulation as amended by Regulation (EU) 2024/2865 of the European Parliament and of the Council until 1 January 2030.’</u></p>	<p>8. Substances and mixtures which have been classified, labelled and packaged in accordance with Article 18(3), Article 31(3) and section 1.2.1 of Annex I as applicable on 9 December 2024 and which were placed on the market before 1 January 2027 shall not be required to be classified, labelled and packaged in accordance with this Regulation as amended by Regulation (EU) 2024/2865 of the European Parliament and of the Council until 1 January 2029.’</p>		
<p>75</p>	<p>Article 1, first paragraph, point (9)(a), amending provision, numbered paragraph (8), second subparagraph</p>			

CLEAN	Commission Proposal	VS.EC	EP Mandate	VS.EC	Council Mandate	VS.EC	Draft Agreement
	Article 1, first paragraph, point (9)(b)						
76	(b) the following paragraph is added:	(b) the following paragraph is added:	(b) the following paragraph is added:	(b) the following paragraph is added:			
	Article 1, first paragraph, point (9)(b), amending provision, numbered paragraph (9), first subparagraph						
77	9. Substances and mixtures which have been labelled in accordance with Article 17(1), Article 25(6) and section 1.5.1.2 and section 1.6 of Annex I as applicable on [OP: please insert the date of the day before the date of entry into force of this Regulation] and which were placed on the market before [OP: please insert 36 months after entry into force of this Regulation] shall not be required to be labelled in accordance with this Regulation as amended by [OP: please add reference to this Regulation] until [OP: please insert 60 months after entry into force of this Regulation].'	9. Substances and mixtures which have been labelled in accordance with Article 17(1), Article 25(6) and section 1.5.1.2 and section 1.6 of Annex I as applicable on [OP: please insert the date of the day before the date of entry into force of this Regulation] and which were placed on the market before [OP: please insert 36 months after entry into force of this Regulation] shall not be required to be labelled in accordance with this Regulation as amended by [OP: please add reference to this Regulation] until [OP: please insert 60 months after entry into force of this Regulation].'	9. Substances and mixtures which have been labelled in accordance with Article 17(1), Article 25(6) and section 1.5.1.2 and section 1.6 of Annex I as applicable on [OP: please insert the date of the day before the date of entry into force of this Regulation] and which were placed on the market before [OP: please insert 36 months after entry into force of this Regulation] shall not be required to be labelled in accordance with this Regulation as amended by [OP: please add reference to this Regulation] until [OP: please insert 60 months after entry into force of this Regulation].'	9. Substances and mixtures which have been labelled in accordance with Article 17(1), Article 25(6) and section 1.5.1.2 and section 1.6 of Annex I as applicable on [OP: please insert the date of the day before the date of entry into force of this Regulation] and which were placed on the market before [OP: please insert 36 months after entry into force of this Regulation] shall not be required to be labelled in accordance with this Regulation as amended by [OP: please add reference to this Regulation] until [OP: please insert 60 months after entry into force of this Regulation].'-			
	Article 1, first paragraph, point (9)(b), amending provision, numbered paragraph (9), second subparagraph						
78	;	;	;	;			

CLEAN	Commission Proposal	VS,EC	EP Mandate	VS,EC	Council Mandate	VS,EC	Draft Agreement
Article 1, first paragraph, point (10)							
79	(10) Annexes I and II are amended in accordance with Annex I to this Regulation.	(10)	Annexes I and II are amended in accordance with Annex I to this Regulation.	(10)	Annexes I and II are amended in accordance with Annex I to this Regulation.		
Article 2							
80	Article 2 Amendments to Regulation (EC) No 1223/2009	Article 2 Amendments to Regulation (EC) No 1223/2009	Article 2 Amendments to Regulation (EC) No 1223/2009	Article 2 Amendments to Regulation (EC) No 1223/2009	Article 2 Amendments to Regulation (EC) No 1223/2009		
Article 2, first paragraph							
81	Regulation (EC) No 1223/2009 is amended as follows:	Regulation (EC) No 1223/2009 is amended as follows:	Regulation (EC) No 1223/2009 is amended as follows:	Regulation (EC) No 1223/2009 is amended as follows:	Regulation (EC) No 1223/2009 is amended as follows:		
Article 2, first paragraph, point (1)							
82	(1) The following Article is inserted:	(1) The following Article is inserted:	(1) The following Article is inserted:	(1) The following Article is inserted:	(1) The following Article is inserted:		
Article 2, first paragraph, point (1), amending provision, first paragraph							
83	Article 14a	Article 14a	Article 14a	Article 14a	Article 14a		
Article 2, first paragraph, point (1), amending provision, second paragraph							
84	Requests for inclusion of substances used as colorants, preservatives or UV-filters in Annexes IV, V or VI	Requests for inclusion of substances used as colorants, preservatives or UV-filters in Annexes IV, V or VI	Requests for inclusion of substances used as colorants, preservatives or UV-filters in Annexes IV, V or VI	Requests for inclusion of substances used as colorants, preservatives or UV-filters in Annexes IV, V or VI	Requests for inclusion of substances used as colorants, preservatives or UV-filters in Annexes IV, V or VI		
Article 2, first paragraph, point (1), amending provision, numbered paragraph (1)							
85	1. A request for a substance to be used as a colorant, preservative or UV-filter to be included in Annex IV, Annex V or	1. A request for a substance to be used as a colorant, preservative or UV-filter to be included in Annex IV, Annex V or	1. A request for a substance to be used as a colorant, preservative or UV-filter to be included in Annex IV, Annex V or	1. A request for a substance to be used as a colorant, preservative or UV-filter to be included in Annex IV, Annex V or	1. A request for a substance to be used as a colorant, preservative or UV-filter to be included in Annex IV, Annex V or		

	CLEAN	Commission Proposal	VS.EC	EP Mandate	VS.EC	Council Mandate	VS.EC	Draft Agreement
		Annex VI, as applicable, may be submitted to the Commission. It shall be accompanied by scientific evidence and documentation showing that due to the latest technical and scientific progress, the substance is safe for use in cosmetic products.	VS.EC	Annex VI, as applicable, may be submitted to the Commission. It shall be accompanied by scientific evidence and documentation showing that due to the latest technical and scientific progress, the substance is safe for use in cosmetic products.	VS.EC	Annex VI, as applicable, may be submitted to the Commission. It shall be accompanied by scientific evidence and documentation showing that due to the latest technical and scientific progress, the substance is safe for use in cosmetic products.	VS.EC	
		Article 2, first paragraph, point (1), amending provision, numbered paragraph (2)						
86		2. After receiving the request referred to in paragraph 1, the Commission shall seek an opinion of the SCCS on the safety of the substance for use in cosmetic products without undue delay.		2. After receiving the request referred to in paragraph 1, the Commission shall seek a request <u>the</u> opinion of the SCCS on the safety of the substance for use in cosmetic products without undue delay.		2. After receiving the request referred to in paragraph 1, the Commission shall seek an opinion request the SCCS to give its opinion on the safety of the substance for use in cosmetic products without undue delay.		
		Article 2, first paragraph, point (1), amending provision, numbered paragraph (3)						
87		3. The SCCS shall transmit its opinion to the Commission within 12 months after receiving the request from the Commission referred to in paragraph 2. The Commission may extend that deadline if additional evidence is required.'		3. The SCCS shall transmit its opinion to the Commission within 12 months after receiving the request from the Commission referred to in paragraph 2. The Commission may extend that deadline if additional evidence is required.'		3. The SCCS shall transmit its opinion to the Commission within 12 months after receiving the request from the Commission referred to in paragraph 2. The Commission may extend that deadline if additional evidence is required.'		
		Article 2, first paragraph, point (1), amending provision, numbered paragraph (3), second subparagraph						
88		;		;		;		

CLEAN	Commission Proposal	VS.EC	EP Mandate	VS.EC	Council Mandate	VS.EC	Draft Agreement
	Article 2, first paragraph, point (2)						
89	(2) Article 15 is amended as follows:	(2) Article 15 is amended as follows:	(2) Article 15 is amended as follows:	(2) Article 15 is amended as follows:	(2) Article 15 is amended as follows:		
	Article 2, first paragraph, point (2)(a)						
90	(a) paragraph 2 is amended as follows:	(a) paragraph 2 is amended as follows:	(a) paragraph 2 is amended as follows:	(a) paragraph 2 is amended as follows:	(a) paragraph 2 is amended as follows:		
	Article 2, first paragraph, point (2)(a)(i)						
91	(i) the second subparagraph is replaced by the following:	(i) the second subparagraph is replaced by the following:	(i) the second subparagraph is replaced by the following:	(i) the second subparagraph is replaced by the following:	(i) the second subparagraph is replaced by the following:		
	Article 2, first paragraph, point (2)(a)(i), amending provision, numbered paragraph (2)						
92	2. However, such substances may be used in cosmetic products if a derogation request is submitted to the Commission at the latest three months after the date of entry into force of the amendments to Part 3 of Annex VI to Regulation (EC) No 1272/2008 classifying the substance as CMR substance of category 1A or 1B. The Commission shall grant the derogation where all of the following conditions are fulfilled:	2. However, such substances may be used in cosmetic products <u>exceptionally</u> , if a derogation request is submitted to the Commission at the latest three months after the date of entry into force of the amendments to Part 3 of Annex VI to Regulation (EC) No 1272/2008 classifying the substance as CMR substance of category 1A or 1B, <u>and the Commission grants the derogation from the general prohibition laid out in subparagraph 1.</u> The Commission shall <u>may</u> grant the derogation where all of the following conditions are fulfilled:	2. However, such substances may be used in cosmetic products if a derogation request is submitted to the Commission at the latest three months after the date of entry into force of the amendments to Part 3 of Annex VI to Regulation (EC) No 1272/2008 classifying the substance as CMR substance of category 1A or 1B. The Commission shall grant the derogation where all of the following conditions are fulfilled:	2. However, such substances may be used in cosmetic products if a derogation request is submitted to the Commission at the latest three months after the date of entry into force of the amendments to Part 3 of Annex VI to Regulation (EC) No 1272/2008 classifying the substance as CMR substance of category 1A or 1B. The Commission shall grant the derogation where all of the following conditions are fulfilled:	2. However, such substances may be used in cosmetic products if a derogation request is submitted to the Commission at the latest three months after the date of entry into force of the amendments to Part 3 of Annex VI to Regulation (EC) No 1272/2008 classifying the substance as CMR substance of category 1A or 1B. The Commission shall grant the derogation where all of the following conditions are fulfilled:		

CLEAN	Commission Proposal	VS,EC	EP Mandate	VS,EC	Council Mandate	VS,EC	Draft Agreement
	Article 2, first paragraph, point (2)(a)(i), amending provision, numbered paragraph (2), point (a)						
93	(a) there are no suitable alternative substances available as documented in an analysis of alternatives;	(a) there are no suitable alternative substances available as documented in an analysis of alternatives;	(a) there are no suitable alternative substances available as documented in an analysis of alternatives;	(a) there are no suitable alternative substances available as documented in an analysis of alternatives;	(a) there are no suitable alternative substances available as documented in an analysis of alternatives;		
	Article 2, first paragraph, point (2)(a)(i), amending provision, numbered paragraph (2), point (b)						
94	(b) the substances have been evaluated and found safe by the SCCS for a particular use of the cosmetic product category, considering exposure to those products, overall exposure from sources other than cosmetics and of vulnerable population groups..	(b) the substances have been evaluated and found safe by the SCCS for one or more particular use of the uses of one or more cosmetic product category; categories considering exposure to those products; overall exposure from the uses in those products categories as well as from sources other than cosmetics and of vulnerable population groups.;	(b) the substances have been evaluated and found safe by the SCCS for one or more particular use of the uses of one or more cosmetic product category; categories considering exposure to those products; overall exposure from the uses in those products categories as well as from sources other than cosmetics and of vulnerable population groups.;	(b) the substances have been evaluated and found safe by the SCCS for a particular use of the cosmetic product category, considering exposure to those products, overall exposure from sources other than cosmetics and of vulnerable population groups.:-	(b) the substances have been evaluated and found safe by the SCCS for a particular use of the cosmetic product category, considering exposure to those products, overall exposure from sources other than cosmetics and of vulnerable population groups.:-		
	Article 2, first paragraph, point (2)(a)(ii)						
95	(ii) the third subparagraph is replaced by the following:	(ii) the third subparagraph is replaced by the following:	(ii) the third subparagraph is replaced by the following:	(ii) the third subparagraph is replaced by the following:	(ii) the third subparagraph is replaced by the following:		
	Article 2, first paragraph, point (2)(a)(ii), amending provision, first paragraph						
96	For the purpose of the second subparagraph, point (a), a substance shall be considered a	For the purpose of the second subparagraph, point (a), a substance shall be considered a	For the purpose of the second subparagraph, point (a), a substance shall be considered a	For the purpose of the second subparagraph, point (a), a substance shall be considered a	For the purpose of the second subparagraph, point (a), a substance, a combination of substances, or where relevant,		

CLEAN	Commission Proposal	VS.EC	EP Mandate	VS.EC	Council Mandate	VS.EC	Draft Agreement
	suitable alternative if it fulfils all of the following conditions:		suitable alternative if it fulfils all of the following conditions:		an alternative technology, that replaces the need for the substance shall be considered a suitable alternative if it fulfils all of the following conditions:		
97	Article 2, first paragraph, point (2)(a)(ii), amending provision, first paragraph, point (a) (a) its use in cosmetic products results in reduced overall risk to human health and the environment;	(a) its use in cosmetic products <i>is safe and</i> results in reduced <i>reduction of</i> overall risk to human health, <i>when assessed against the substance it is intended to replace and the environment</i> ;	(a) its use in cosmetic products results in reduced overall risk to human health and the environment;				
98	Article 2, first paragraph, point (2)(a)(ii), amending provision, first paragraph, point (b) (b) it provides an equivalent function to the classified substance, in a finished cosmetic product with a similar effect and the same level of efficacy;	(b) it provides an equivalent function to the classified substance, in a finished cosmetic product with a similar <i>comparable</i> effect and the same level of efficacy <i>and performance</i> ;	(b) it provides an equivalent function to the classified substance, in a finished cosmetic product with a similar <i>comparable</i> effect and the same level of efficacy;				
99	Article 2, first paragraph, point (2)(a)(ii), amending provision, first paragraph, point (c) (c) is technically feasible and economically viable;	(c) is technically feasible and economically viable <i>feasible</i> <u>provided costs and supply conditions allow sustained production</u> ;	(c) is technically feasible and economically viable <i>feasible</i> ;				
100	Article 2, first paragraph, point (2)(a)(ii), amending provision, first paragraph, point (d) (d) it is not restricted, not protected by exclusive rights, and	(d) it is not restricted, not protected by exclusive rights , and	(d) it is not restricted, not protected by exclusive rights, and				

	CLEAN	Commission Proposal	VS.EC	EP Mandate	VS.EC	Council Mandate	VS.EC	Draft Agreement
		is available on the market at scale, in quantities large enough to meet current and expected demand.'		is <u>either</u> available on the market at scale; and in quantities large enough to meet current and expected demand in a reasonable timeframe. '		is available on the market at scale, in quantities large enough to meet current and expected demand.'		
	Article 2, first paragraph, point (2)(a)(iii), amending provision, first paragraph, point (da)							
100a			(<u>ia</u>) <u>The following subparagraph is inserted after the third subparagraph:</u> ' <u>The Commission shall consult relevant stakeholders for the purpose of the second subparagraph, point (a) and for the purpose of the third subparagraph.</u> '					
	Article 2, first paragraph, point (2)(a)(iii), amending provision, second paragraph							
101	;	;	;	;	;	;	;	
	Article 2, first paragraph, point (2)(a)(iii)							
102	(iii) the following subparagraph is inserted after the fourth subparagraph:	(iii) the following subparagraph is inserted after the fourth subparagraph:	(iii) the following subparagraph is inserted after the fourth subparagraph:	(iii) the following subparagraph is inserted after the fourth subparagraph:	(iii) the following subparagraph is inserted after the fourth subparagraph:	(iii) the following subparagraph is inserted after the fourth subparagraph:	(iii) the following subparagraph is inserted after the fourth subparagraph:	
	Article 2, first paragraph, point (2)(a)(iii), amending provision, first paragraph							
103	The deadline laid down in the fourth subparagraph of this	The deadline laid down in the fourth subparagraph of this	The deadline laid down in the fourth subparagraph of this	The deadline laid down in the fourth subparagraph of this	The deadline laid down in the fourth subparagraph of this	The deadline laid down in the fourth subparagraph of this	The deadline laid down in the fourth subparagraph of this	

	CLEAN	Commission Proposal	VS,EC	EP Mandate	VS,EC	Council Mandate	VS,EC	Draft Agreement
	paragraph shall start on the date of entry into application of the relevant amendments to Part 3 of Annex VI to Regulation (EC) No 1272/2008 classifying the substance concerned as CMR substance of category 1A, or 1B.;	paragraph shall start on the date of entry into application force of the relevant amendments to Part 3 of Annex VI to Regulation (EC) No 1272/2008 classifying the substance concerned as CMR substance of category 1A, or 1B.;	VS,EC	EP Mandate	VS,EC	Council Mandate	VS,EC	Draft Agreement
Article 2, first paragraph, point (2)(a)(iii), amending provision, first paragraph a								
103a				<u>5a. Where a derogation request referred to in the second subparagraph of paragraph 2 has been submitted for CMR substances of category 1A, or 1B, this deadline may, where relevant, be extended by twelve months.</u>				
Article 2, first paragraph, point (2)(b)								
104	(b) the following paragraphs 5, 6 and 7 are added:	(b) the following paragraphs 5, 6 and 7 are added:			(b) the following paragraphs 5, 6 and 7 are added:	(b) the following paragraphs 5, 6 and 7 are added:		
Article 2, first paragraph, point (2)(b), amending provision, numbered paragraph (5)								
105	5. The prohibition referred to in paragraphs 1 and 2 of this Article shall not apply to a substance where the oral or inhalation route of exposure for CMR harmonised classification has been explicitly indicated in the	5. The prohibition referred to in paragraphs 1 and 2 of this Article shall not apply to a substance where the oral or inhalation route of exposure for CMR harmonised classification has been explicitly indicated in the			5. The prohibition referred to in paragraphs 1 and 2 of this Article shall not apply to a substance where the oral or inhalation route of exposure for CMR harmonised classification has been explicitly indicated in the	5. The prohibition referred to in paragraphs 1 and 2 of this Article shall not apply to a substance where the oral or inhalation route of exposure for CMR harmonised classification has been explicitly indicated in the		

CLEAN	Commission Proposal	V.S.EC	EP Mandate	V.S.EC	Council Mandate	V.S.EC	Draft Agreement
	<p>'Hazard statement Code(s)' column under the 'Classification' in Part 3 of Annex VI to Regulation (EC) No 1272/2008. If a potential risk to human health arises from cosmetic products containing such substance due to incidental ingestion or inhalation, the Commission shall request an SCCS opinion on the safety of the substance concerned in those specific product types without undue delay.</p>	<p>'Hazard statement Code(s)' column under the 'Classification' in Part 3 of Annex VI to Regulation (EC) No 1272/2008. If a potential risk to human health arises from cosmetic products containing such substance due to incidental ingestion or inhalation, the Commission shall request an SCCS opinion on the safety of the substance concerned in those specific product types without undue delay: deleted</p>	<p>'Hazard statement Code(s)' column under the 'Classification' in Part 3 of Annex VI to Regulation (EC) No 1272/2008. If a potential risk to human health arises from cosmetic products containing such substance due to incidental ingestion or inhalation, the Commission shall request an SCCS opinion on the safety of the substance concerned in those specific product types without undue delay.</p>				
106	<p>Article 2, first paragraph, point (2)(b), amending paragraph (6), first subparagraph</p> <p>6. The prohibition referred to in paragraphs 1 and 2 of this Article shall not apply to a substance extracted from plants or plant parts and not chemically modified as defined in Article 3, point (40), of Regulation (EC) No 1907/2006, containing more than one constituent, at least one of which has been classified as a CMR substance of category 1A, 1B, or 2 under Part 3 of Annex VI to Regulation (EC) No 1272/2008. If a potential risk to human health arises from the use of such substance in cosmetic products, the Commission shall seek an</p>	<p>6. The prohibition referred to in paragraphs 1 and 2 of this Article shall not apply to a <u>substances</u> extracted from plants or plant parts and not chemically modified as defined in Article 3, point (40), of Regulation (EC) No 1907/2006, containing more than one constituent, at least one of which has been classified as a CMR substance of category 1A, 1B, or 2 under Part 3 of Annex VI to Regulation (EC) No 1272/2008. <u>As a potential risk to human health arises from the presence of such constituent classified as CMR category 1A, 1B or 2 in</u></p>	<p>6. The prohibition referred to in paragraphs 1 and 2 of this Article shall not apply to a <u>substances</u> extracted from plants or plant parts and not chemically modified as defined in Article 3, point (40), of Regulation (EC) No 1907/2006, containing more than one constituent, at least one of which has been classified as a CMR substance of category 1A, 1B, or 2 under Part 3 of Annex VI to Regulation (EC) No 1272/2008. <u>As a potential risk to human health arises from the use of such substance in cosmetic products of a substance containing a</u></p>				

CLEAN	Commission Proposal	VS.EC	EP Mandate	VS.EC	Council Mandate	VS.EC	Draft Agreement
	<p>opinion of the SCCS on the safety of that substance for its use in cosmetic products without undue delay.</p>	<p><u>such substances in cosmetic products, the Commission shall without delay request an opinion of the SCCS on the safety of that constituent for its presence in cosmetic products. The SCCS shall deliver its opinion within 12 months of the Commission's request. The Commission may extend that deadline by six months if additional evidence is required. The SCCS shall deliver its final opinion within six months of submission of additional data. The opinion of the SCCS shall be made publicly available. Taking into account the opinion of the SCCS, and where a potential risk to human health arises from the use of such substance referred to in cosmetic products, the Commission shall seek an opinion of the SCCS on the safety of that substance for its use the first subparagraph in cosmetic products containing a constituent classified as CMR category IA, IB or 2 for human health, the Commission shall, without undue delay, amend the Annexes to this Regulation. For the purposes of this paragraph,</u></p>	<p>constituent classified as CMR category IA, IB or 2, the Commission shall seek an opinion of the SCCS on the safety of that substance for its use in cosmetic products without undue delay request the SCCS to give its opinion on the safety of the CMR-constituents present in the substance for a particular use of the cosmetic product category, as part of the exposure assessment.</p>				

CLEAN	Commission Proposal	VS,EC	EP Mandate	VS,EC	Council Mandate	VS,EC	Draft Agreement
			<u><i>“plants” means living or dead organisms from the kingdoms Plantae and Fungi, including algae, lichens and yeasts.</i></u>				
Article 2, first paragraph, point (2)(b), amending provision, numbered paragraph (6), second subparagraph							
107	For the purpose of this paragraph, ‘plants’ means living or dead organisms from the kingdoms Plantae and Fungi, and includes algae, lichens and yeasts.		For the purpose of this paragraph, ‘plants’ means living or dead organisms from the kingdoms Plantae and Fungi, and includes algae, lichens and yeasts.		For the purpose of this paragraph, ‘plants’ means living or dead organisms from the kingdoms Plantae and Fungi, and includes algae, lichens and yeasts.		
Article 2, first paragraph, point (2)(b), amending provision, numbered paragraph (7), first subparagraph							
108	7. Cosmetic products containing a substance classified as a CMR substance of category 1A, 1B, or 2 under Part 3 of Annex VI to Regulation (EC) No 1272/2008 prohibited from use in cosmetic products or such substance not compliant with a restriction may continue to be placed on the market for 12 months and be made available on the market for 24 months after the entry into force of the relevant amendments to the relevant Annexes to this Regulation.’		7. <u><i>Cosmetic products containing a substance classified as a CMR substance of category 1A, 1B, or 2 under Part 3 of Annex VI to Regulation (EC) No 1272/2008 prohibited from use in cosmetic products and for which no derogation request was submitted in accordance with paragraph 2 or such substance is not compliant with a restriction may continue to be placed on the market for 6 months, and be available on the market for 15 months after the entry into force of the relevant amendments to this Regulation. Cosmetic products containing a substance classified</i></u>		7. Cosmetic products containing a substance classified as a CMR substance of category 1A, 1B, or 2 under Part 3 of Annex VI to Regulation (EC) No 1272/2008 prohibited from use in cosmetic products or such substance not compliant with a restriction may continue to be placed on the market for 12 6 months and be made available on the market for 24 12 months after the entry into force of the relevant amendments to the relevant Annexes to this Regulation.-		

CLEAN	Commission Proposal	VS,EC EP Mandate	VS,EC Council Mandate	VS,EC Draft Agreement
		<p><u>as a CMR substance of category IA, IB, or 2 under Part 3 of Annex VI to Regulation (EC) No 1272/2008 prohibited from use in cosmetic products and for which a derogation request was submitted in accordance with paragraph 2, but not granted due to safety concerns by the SCCS or such substance not compliant with a restriction may continue to be placed on the market for 3 months, and be available on the market for 12 months after the entry into force of the relevant amendments to the relevant Annexes to this Regulation.</u></p> <p>Cosmetic products containing a substance classified as a CMR substance of category 1A, 1B, or 2 under Part 3 of Annex VI to Regulation (EC) No 1272/2008 prohibited from use in cosmetic products or such substance not compliant with a restriction <i>and for which a derogation request was submitted in accordance with paragraph 2, but not granted due to the availability of a suitable alternative</i>, may continue to be placed on the market for 124 months and be made available on</p>		

CLEAN	Commission Proposal	VS,EC	EP Mandate	VS,EC	Council Mandate	VS,EC	Draft Agreement
			the market for 24 48 months after the entry into force of the relevant amendments to the relevant Annexes to this Regulation, <u>and where needed updated the cosmetic product safety report (CPSR) remains available.</u> ?				
	Article 2, first paragraph, point (2)(b), amending provision, numbered paragraph (7), second subparagraph						
109	;	;	;	;	;	;	
	Article 2, first paragraph, point (3)						
110	(3) In Article 16, paragraphs 3 and 7 are deleted;	(3) In Article 16, paragraphs 3 and 7 are deleted ; <u>is replaced by the following:</u> <u>3. In addition to the notification under Article 13, cosmetic products containing nanomaterials shall be notified to the Commission by the responsible person by electronic means prior to being placed on the market. The first subparagraphs shall not apply to cosmetic products containing nanomaterials that are in conformity with the requirements set out in Annex III. The information notified to the</u>	(3) In Article 16, paragraphs 3 and 7 are deleted ; <u>is replaced by the following:</u> <u>3. In addition to the notification under Article 13, cosmetic products containing nanomaterials shall be notified to the Commission by the responsible person by electronic means prior to being placed on the market. The first subparagraphs shall not apply to cosmetic products containing nanomaterials that are in conformity with the requirements set out in Annex III. The information notified to the</u>	(3) In Article 16, paragraphs 3 and 7 are deleted paragraph 3 is replaced by the following;			

CLEAN	Commission Proposal	VS,EC	EP Mandate	VS,EC	Council Mandate	VS,EC	Draft Agreement
		<p><u>Commission shall contain at least/</u> <u>(a) the identification of the nanomaterial including its chemical name (IUPAC) and other descriptors as specified in point 2 of the Preamble to Annexes II to VI, and</u> <u>(b) the specification of the nanomaterial including size of particles, physical and chemical properties.</u> <u>The responsible person may designate another legal or natural person by written mandate for the notification of nanomaterials and shall inform the Commission thereof. The Commission shall provide a reference number for the submission of, which may substitute the information to be notified in case of the same nanomaterial used in different products.</u></p>					
Article 2, first paragraph, point (4)							
111	(4) In Article 19, paragraph 6 is replaced by the following:	(4) In Article 19, paragraph 6 is replaced by the following:	(4) In Article 19, paragraph 6 is replaced by the following:	(4) In Article 19, paragraph 6 is replaced by the following:			
Article 2, first paragraph, point (4), amending provision, numbered paragraph (6), first subparagraph							
112	‘	‘	‘	‘			

CLEAN	Commission Proposal	VS.EC	EP Mandate	VS.EC	Council Mandate	VS.EC	Draft Agreement
6.	The information mentioned in paragraph 1, point (g) shall be expressed by using the common ingredient name in accordance with the internationally recognised nomenclature and in the absence of a common ingredient name, a term as contained in a generally accepted nomenclature shall be used.'	6.	The information mentioned in paragraph 1, point (g) shall be expressed by using the common ingredient name in accordance with the internationally recognised nomenclature and in the absence of a common ingredient name, a term as contained in a generally accepted nomenclature shall be used.'	6.	The information mentioned in paragraph 1, point (g) shall be expressed by using the common ingredient name in accordance with the internationally recognised nomenclature and in the absence of a common ingredient name, a term as contained in a generally accepted nomenclature shall be used.'		
Article 2, first paragraph, point (4), amending provision, numbered paragraph (1), first subparagraph a							
112a		<u>4a</u>	<u>In Article 19, the following paragraph is added:</u> <u>'6a. When cosmetic products are made available on the market through distance sales, the offer shall clearly and visibly indicate the information referred to in paragraph 1.'</u>				
Article 2, first paragraph, point (4), amending provision, numbered paragraph (6), second subparagraph							
113	;	;	;	;	;	;	;
Article 2, first paragraph, point (5)							
114	(5) In Article 22, fourth subparagraph, the second sentence is deleted;	(5)	In Article 22, fourth subparagraph, the second sentence is deleted;	(5)	In Article 22, fourth subparagraph, the second sentence is deleted;		
Article 2, first paragraph, point (6)							

	CLEAN	Commission Proposal	VS,EC	EP Mandate	VS,EC	Council Mandate	VS,EC	Draft Agreement
115	(6)	Article 33 is deleted;	(6)	Article 33 is deleted;	(6)	Article 33 is deleted;	(6)	Article 33 is deleted;
Article 2, first paragraph, point (7)								
116	(7)	Annex I is amended in accordance with Annex II to this Regulation;	(7)	Annex I is amended in accordance with Annex II to this Regulation;	(7)	Annex I is amended in accordance with Annex II to this Regulation;		
Article 2, first paragraph, point (8)								
117	(8)	Annexes II to VI are amended in accordance with Annex III this Regulation.	(8)	Annexes II to VI are amended in accordance with Annex III this Regulation.	(8)	Annexes II to VI are amended in accordance with Annex III to this Regulation.		
Article 3								
118	Article 3	Amendments to Regulation (EU) 2019/1009	Article 3	Amendments to Regulation (EU) 2019/1009	Article 3	Amendments to Regulation (EU) 2019/1009		
Article 3, first paragraph								
119	Regulation (EU) 2019/1009 is amended as follows:	Regulation (EU) 2019/1009 is amended as follows:	Regulation (EU) 2019/1009 is amended as follows:	Regulation (EU) 2019/1009 is amended as follows:	Regulation (EU) 2019/1009 is amended as follows:	Regulation (EU) 2019/1009 is amended as follows:		
Article 3, first paragraph, point (1)								
120	(1)	in Article 2, the following point (15a) is inserted:	(1)	in Article 2, the following point (15a) is inserted:	(1)	in Article 2, the following point (15a) is inserted:		
Article 3, first paragraph, point (1), amending provision, numbered paragraph (15a), first subparagraph								
121	(15a)	‘digital contact’ means any up-to-date and accessible online communication channel through which economic operators can be reached or engaged without	(15a)	‘digital contact’ means any up-to-date and <u>freely</u> accessible online communication channel through which <u>an</u> economic operator <u>operator</u> can be reached or engaged <u>contacted</u>	(15a)	‘digital contact’ means any up-to-date and freely accessible online communication channel such as email addresses through which economic operators can be reached or		

	CLEAN	Commission Proposal	VS,EC	EP Mandate	VS,EC	Council Mandate	VS,EC	Draft Agreement
		the need to register or to download an application;		without the need to register or to download an application or use <u>additional applications specific to the economic operator;</u>		engaged contacted without the need to register or to download an application or use additional applications specific to the economic operator;		
		Article 3, first paragraph, point (1), amending provision, numbered paragraph (15a), second subparagraph						
122		;	;	;	;	;		
		Article 3, first paragraph, point (2)						
123		(2) Article 6 is amended as follows:	(2) Article 6 is amended as follows:	(2) Article 6 is amended as follows:	(2) Article 6 is amended as follows:			
		Article 3, first paragraph, point (2)(a)						
124		(a) paragraph 2 is amended as follows:	(a) paragraph 2 is amended as follows:	(a) paragraph 2 is amended as follows:	(a) paragraph 2 is amended as follows:			
		Article 3, first paragraph, point (2)(a)(i)						
125		(i) the second subparagraph is replaced by the following:	(i) the second subparagraph is replaced by the following:	(i) the second subparagraph is replaced by the following:	(i) the second subparagraph is replaced by the following:			
		Article 3, first paragraph, point (2)(a)(i), amending provision, first paragraph						
126		Where compliance of an EU fertilising product with the applicable requirements laid down in this Regulation has been demonstrated by that conformity assessment procedure, manufacturers shall draw up an EU declaration of conformity, in	Where compliance of an EU fertilising product with the applicable requirements laid down in this Regulation has been demonstrated by that conformity assessment procedure, manufacturers shall draw up an EU declaration of conformity, in	Where compliance of an EU fertilising product with the applicable requirements laid down in this Regulation has been demonstrated by that conformity assessment procedure, manufacturers shall draw up an EU declaration of conformity, in	Where compliance of an EU fertilising product with the applicable requirements laid down in this Regulation has been demonstrated by that conformity assessment procedure, manufacturers shall draw up an EU declaration of conformity, in			

	CLEAN	Commission Proposal	VS.EC	EP Mandate	VS.EC	Council Mandate	VS.EC	Draft Agreement
		electronic form, and affix the CE marking.;		electronic form, and affix the CE marking.;		electronic form, and affix the CE marking.;		
Article 3, first paragraph, point (2)(a)(ii)								
127	(ii) the following subparagraph is added:	(ii) the following subparagraph is added:				(ii) the following subparagraph is added:		
Article 3, first paragraph, point (2)(a)(ii), amending provision, first paragraph								
128	Manufacturers shall ensure that the EU fertilising product is accompanied by the internet address or data carrier through which the EU declaration of conformity can be accessed.;	Manufacturers shall ensure that the EU fertilising product is accompanied by the internet address or data carrier through which the EU declaration of conformity can be <u>directly</u> accessed.;		Manufacturers shall ensure that the EU fertilising product is accompanied by the internet address or data carrier through which the EU declaration of conformity can be accessed.;		Manufacturers shall ensure that the EU fertilising product is accompanied by the internet address or data carrier through which the EU declaration of conformity can be accessed.;		
Article 3, first paragraph, point (2)(b)								
129	(b) in paragraph 3, the second subparagraph is replaced by the following:	(b) in paragraph 3, the second subparagraph is replaced by the following:		(b) in paragraph 3, the second subparagraph is replaced by the following:		(b) in paragraph 3, the second subparagraph is replaced by the following:		
Article 3, first paragraph, point (2)(b), amending provision, first paragraph								
130	On request, manufacturers shall make the EU declaration of conformity available to other economic operators in electronic form.;	On request, manufacturers shall make the EU declaration of conformity available to other economic operators in electronic form.;		On request, manufacturers shall make the EU declaration of conformity available to other economic operators in electronic form.;		On request, manufacturers shall make the EU declaration of conformity available to other economic operators in electronic form.;		

CLEAN	Commission Proposal	VS.EC	EP Mandate	VS.EC	Council Mandate	VS.EC	Draft Agreement
	,		,		,		
Article 3, first paragraph, point (2)(c)							
131	(c) in paragraph 6, first subparagraph, the first and second sentences are replaced by the following:	(c) in paragraph 6, first subparagraph, the first and second sentences are replaced by the following:	(c) in paragraph 6, first subparagraph, the first and second sentences are replaced by the following:	(c) in paragraph 6, first subparagraph, the first and second sentences are replaced by the following:			
Article 3, first paragraph, point (2)(c), amending provision, first paragraph							
132	Manufacturers shall indicate on the packaging of the EU fertilising product their name, registered trade name or registered trademark as well as their postal address and digital contact or, where the EU fertilising product is supplied without packaging, in a document accompanying the EU fertilising product. The postal address and digital contact shall indicate a single point through which the manufacturer can be reached.²	Manufacturers shall indicate on the packaging of the EU fertilising product their name, registered trade name or registered trademark as well as their postal address and digital contact or, where the EU fertilising product is supplied without packaging, in a document accompanying the EU fertilising product. The postal address and digital contact shall indicate a single point through which the manufacturer can be reached.²	Manufacturers shall indicate on the packaging of the EU fertilising product their name, registered trade name or registered trademark as well as their postal address and digital contact or, where the EU fertilising product is supplied without packaging, in a document accompanying the EU fertilising product. The postal address and digital contact shall indicate a single point through which the manufacturer can be reached.²	Manufacturers shall indicate on the packaging of the EU fertilising product their name, registered trade name or registered trademark as well as their postal address and digital contact or, where the EU fertilising product is supplied without packaging, in a document accompanying the EU fertilising product. The postal address and digital contact shall indicate a single point through which the manufacturer can be reached.²			
Article 3, first paragraph, point (2)(c), amending provision, second paragraph							
133	;	;	;	;	;		
Article 3, first paragraph, point (2)(d)							

	CLEAN	Commission Proposal	vs.EC	EP Mandate	vs.EC	Council Mandate	vs.EC	Draft Agreement
134		(d) in paragraph 9, the first sentence is replaced by the following:		(d) in paragraph 9, the first sentence is replaced by the following:		(d) in paragraph 9, the first sentence is replaced by the following:		
Article 3, first paragraph, point (2)(d), amending provision, first paragraph								
135		Manufacturers shall, further to a reasoned request from a competent national authority, provide it, in electronic form, with all the information and documentation necessary to demonstrate the conformity of the EU fertilising product with this Regulation, in a language which can be easily understood by that authority.?		Manufacturers shall, further to a reasoned request from a competent national authority, provide it, in electronic form, with all the information and documentation necessary to demonstrate the conformity of the EU fertilising product with this Regulation, in a language which can be easily understood by that authority.?		Manufacturers shall, further to a reasoned request from a competent national authority, provide it, in electronic form, with all the information and documentation necessary to demonstrate the conformity of the EU fertilising product with this Regulation, in a language which can be easily understood by that authority.?		
Article 3, first paragraph, point (2)(d), amending provision, second paragraph								
136		;		;		;		
Article 3, first paragraph, point (3)								
137		(3) in Article 7(2), point (b) is replaced by the following:		(3) in Article 7(2), point (b) is replaced by the following:		(3) in Article 7(2), point (b) is replaced by the following:		
Article 3, first paragraph, point (3), amending provision, numbered paragraph (b), first subparagraph								
138		(b) further to a reasoned request from a competent national authority, provide that authority, in electronic form, with all the information and documentation		(b) further to a reasoned request from a competent national authority, provide that authority, in electronic form, with all the information and documentation		(b) further to a reasoned request from a competent national authority, provide that authority, in electronic form, with all the information and documentation		

CLEAN	Commission Proposal	VS,EC	EP Mandate	VS,EC	Council Mandate	VS,EC	Draft Agreement
	necessary to demonstrate the conformity of an EU fertilising product;		necessary to demonstrate the conformity of an EU fertilising product;		necessary to demonstrate the conformity of an EU fertilising product;²		
	Article 3, first paragraph, point (3), amending provision, numbered paragraph (b), second subparagraph						
139	;	;	;	;	;	;	
	Article 3, first paragraph, point (4)						
140	(4) Article 8 is amended as follows:	(4)	Article 8 is amended as follows:	(4)	Article 8 is amended as follows:		
	Article 3, first paragraph, point (4)(a)						
141	(a) in paragraph 2, first subparagraph, the second sentence is replaced by the following:	(a)	in paragraph 2, first subparagraph, the second sentence is replaced by the following:	(a)	in paragraph 2, first subparagraph, the second sentence is replaced by the following:		
	Article 3, first paragraph, point (4)(a), amending provision, first paragraph						
142	They shall ensure that the manufacturer has drawn up the technical documentation, that the EU fertilising product is accompanied by the internet address or data carrier through which the EU declaration of conformity can be accessed and, where appropriate, by other required documents, and that the manufacturer has complied with the requirements set out in Article 6(5) and (6).²	They shall ensure that the manufacturer has drawn up the technical documentation, that the EU fertilising product is accompanied by the internet address or data carrier through which the EU declaration of conformity can be accessed and, where appropriate, by other required documents, and that the manufacturer has complied with the requirements set out in Article 6(5) and (6).²	They shall ensure that the manufacturer has drawn up the technical documentation, that the EU fertilising product is accompanied by the internet address or data carrier through which the EU declaration of conformity can be <i>directly</i> accessed and, where appropriate, by other required documents, and that the manufacturer has complied with the requirements set out in Article 6(5) and (6).²	They shall ensure that the manufacturer has drawn up the technical documentation, that the EU fertilising product is accompanied by the internet address or data carrier through which the EU declaration of conformity can be accessed and, where appropriate, by other required documents, and that the manufacturer has complied with the requirements set out in Article 6(5) and (6).²			

CLEAN	Commission Proposal	VS.EC	EP Mandate	VS.EC	Council Mandate	VS.EC	Draft Agreement
Article 3, first paragraph, point (4)(a), amending provision, second paragraph							
143	;	;	;	;	;	;	
Article 3, first paragraph, point (4)(b)							
144	(b) in paragraph 3, the first sentence is replaced by the following:	(b) in paragraph 3, the first sentence is replaced by the following:	(b) in paragraph 3, the first sentence is replaced by the following:	(b) in paragraph 3, the first sentence is replaced by the following:	(b) in paragraph 3, the first sentence is replaced by the following:		
Article 3, first paragraph, point (4)(b), amending provision, first paragraph							
145	Importers shall indicate their name, registered trade name or their postal address and digital contact on the packaging of the EU fertilising product or, where the EU fertilising product is supplied without packaging, in a document accompanying the EU fertilising product. ²	Importers shall indicate their name, registered trade name or their postal address and digital contact on the packaging of the EU fertilising product or, where the EU fertilising product is supplied without packaging, in a document accompanying the EU fertilising product. ²	Importers shall indicate their name, registered trade name or their postal address and digital contact on the packaging of the EU fertilising product or, where the EU fertilising product is supplied without packaging, in a document accompanying the EU fertilising product. ²	Importers shall indicate their name, registered trade name or their postal address and digital contact on the packaging of the EU fertilising product or, where the EU fertilising product is supplied without packaging, in a document accompanying the EU fertilising product. ²	Importers shall indicate their name, registered trade name or their postal address and digital contact on the packaging of the EU fertilising product or, where the EU fertilising product is supplied without packaging, in a document accompanying the EU fertilising product. ²		
Article 3, first paragraph, point (4)(b), amending provision, second paragraph							
146	;	;	;	;	;	;	
Article 3, first paragraph, point (4)(c)							
147	(c) paragraph 8 is replaced by the following:	(c) paragraph 8 is replaced by the following:	(c) paragraph 8 is replaced by the following:	(c) paragraph 8 is replaced by the following:	(c) paragraph 8 is replaced by the following:	(c) paragraph 8 is replaced by the following:	
Article 3, first paragraph, point (4)(c), amending provision, numbered paragraph (8), first subparagraph							

	CLEAN	Commission Proposal	VS.EC	EP Mandate	VS.EC	Council Mandate	VS.EC	Draft Agreement
148		8. Importers shall, for 5 years after the EU fertilising product has been placed on the market, keep the EU declaration of conformity at the disposal of the market surveillance authorities and ensure that the technical documentation can be made available to those authorities, upon request.	8. Importers shall, for 5 years after the EU fertilising product has been placed on the market, keep the EU declaration of conformity at the disposal of the market surveillance authorities and ensure that the technical documentation can be made available <u>in a swift manner</u> to those authorities, upon request.	8. Importers shall, for 5 years after the EU fertilising product has been placed on the market, keep the EU declaration of conformity at the disposal of the market surveillance authorities and ensure that the technical documentation can be made available to those authorities, upon request.				
Article 3, first paragraph, point (4)(c), amending provision, numbered paragraph (8), second subparagraph								
149		On request, importers shall make the EU declaration of conformity available to other economic operators in electronic form. ²	On request, importers shall make the EU declaration of conformity available to other economic operators in electronic form <u>in a swift manner</u> .	On request, importers shall make the EU declaration of conformity available to other economic operators in electronic form. ²				
Article 3, first paragraph, point (4)(c), amending provision, numbered paragraph (8), third subparagraph								
150		;	;	;				
Article 3, first paragraph, point (4)(d)								
151		(d) in paragraph 9, the first sentence is replaced by the following:	(d) in paragraph 9, the first sentence is replaced by the following:	(d) in paragraph 9, the first sentence is replaced by the following:				
Article 3, first paragraph, point (4)(d), amending provision, first paragraph								
152		Importers shall, further to a reasoned request from a competent	Importers shall, further to a reasoned request from a competent	Importers shall, further to a reasoned request from a competent				

	CLEAN	Commission Proposal	VS,EC	EP Mandate	VS,EC	Council Mandate	VS,EC	Draft Agreement
		national authority, provide it, in electronic form, with all the information and documentation necessary to demonstrate the conformity of the EU fertilising product with this Regulation in a language which can be easily understood by that authority.’	VS,EC	national authority, provide it, in electronic form, with all the information and documentation necessary to demonstrate the conformity of the EU fertilising product with this Regulation in a language which can be easily understood by that authority.’	VS,EC	national authority, provide it, in electronic form, with all the information and documentation necessary to demonstrate the conformity of the EU fertilising product with this Regulation in a language which can be easily understood by that authority.’-		
		Article 3, first paragraph, point (4)(d), amending provision, second paragraph						
153	;	;	;	;	;	;	;	;
		Article 3, first paragraph, point (5)						
154	(5) follows:	Article 9 is amended as follows:	(5) follows:	Article 9 is amended as follows:	(5) follows:	Article 9 is amended as follows:		
		Article 3, first paragraph, point (5)(a)						
155	(a) follows:	in paragraph 2, the first subparagraph is replaced by the following:	(a) follows:	in paragraph 2, the first subparagraph is replaced by the following:	(a) follows:	in paragraph 2, the first subparagraph is replaced by the following:		
		Article 3, first paragraph, point (5)(a), amending provision, first paragraph						
156	Before making an EU fertilising product available on the market, distributors shall verify that it is accompanied by the internet address or data carrier through which the EU declaration of conformity can be accessed and, where appropriate, by other	Before making an EU fertilising product available on the market, distributors shall verify that it is accompanied by the internet address or data carrier through which the EU declaration of conformity can be accessed and, where appropriate, by other	Before making an EU fertilising product available on the market, distributors shall verify that it is accompanied by the internet address or data carrier through which the EU declaration of conformity can be accessed and, where appropriate, by other	Before making an EU fertilising product available on the market, distributors shall verify that it is accompanied by the internet address or data carrier through which the EU declaration of conformity can be accessed and, where appropriate, by other	Before making an EU fertilising product available on the market, distributors shall verify that it is accompanied by the internet address or data carrier through which the EU declaration of conformity can be accessed and, where appropriate, by other	Before making an EU fertilising product available on the market, distributors shall verify that it is accompanied by the internet address or data carrier through which the EU declaration of conformity can be accessed and, where appropriate, by other		

CLEAN	Commission Proposal	VS.EC	EP Mandate	VS.EC	Council Mandate	VS.EC	Draft Agreement
	required documents, including the information referred to in Article 6(7) or Article 8(4) provided in the manner specified therein, in a language which can be easily understood by end-users in the Member State in which the EU fertilising product is to be made available on the market, and that the manufacturer and the importer have complied with the requirements set out in Article 6(5) and (6) and Article 8(3) respectively.’	VS.EC	by other required documents, including the information referred to in Article 6(7) or Article 8(4) provided in the manner specified therein, in a language which can be easily understood by end-users in the Member State in which the EU fertilising product is to be made available on the market, and that the manufacturer and the importer have complied with the requirements set out in Article 6(5) and (6) and Article 8(3) respectively.’	VS.EC	required documents, including the information referred to in Article 6(7) or Article 8(4) provided in the manner specified therein, in a language which can be easily understood by end-users in the Member State in which the EU fertilising product is to be made available on the market, and that the manufacturer and the importer have complied with the requirements set out in Article 6(5) and (6) and Article 8(3) respectively.’		
	Article 3, first paragraph, point (5)(a), amending provision, second paragraph						
157	;	;	;	;	;		
	Article 3, first paragraph, point (5)(b)						
158	(b) in paragraph 5, the first sentence is replaced by the following:	(b) in paragraph 5, the first sentence is replaced by the following:	(b) in paragraph 5, the first sentence is replaced by the following:	(b) in paragraph 5, the first sentence is replaced by the following:			
	Article 3, first paragraph, point (5)(b), amending provision, first paragraph						
159	‘ Distributors shall, further to a reasoned request from a competent national authority, provide it, in electronic form, with all the information and documentation necessary to demonstrate the	‘ Distributors shall, further to a reasoned request from a competent national authority, provide it, in electronic form, with all the information and documentation necessary to demonstrate the	‘ Distributors shall, further to a reasoned request from a competent national authority, provide it, in electronic form, with all the information and documentation necessary to demonstrate the	‘ Distributors shall, further to a reasoned request from a competent national authority, provide it, in electronic form, with all the information and documentation necessary to demonstrate the	‘ Distributors shall, further to a reasoned request from a competent national authority, provide it, in electronic form, with all the information and documentation necessary to demonstrate the		

	CLEAN	Commission Proposal	VS,EC	EP Mandate	VS,EC	Council Mandate	VS,EC	Draft Agreement
		conformity of the EU fertilising product with this Regulation.'		conformity of the EU fertilising product with this Regulation.'		conformity of the EU fertilising product with this Regulation.-'		
		Article 3, first paragraph, point (5)(b), amending provision, second paragraph						
160		;		;		;		
		Article 3, first paragraph, point (6)						
161		(6) Article 15 is amended as follows:	(6)	Article 15 is amended as follows:	(6)	Article 15 is amended as follows:		
		Article 3, first paragraph, point (6)(a)						
162		(a) paragraph 2 is replaced by the following:	(a)	paragraph 2 is replaced by the following:	(a)	paragraph 2 is replaced by the following:		
		Article 3, first paragraph, point (6)(a), amending provision, numbered paragraph (2)						
163	2.	Records and correspondence relating to the conformity assessment procedures shall be drawn up, in electronic form, in an official language of the Member State where the notified body carrying out the procedures is established or in a language accepted by that body.	2.	Records and correspondence relating to the conformity assessment procedures shall be drawn up, in electronic form, in an official language of the Member State where the notified body carrying out the procedures is established or in a language accepted by that body.	2.	Records and correspondence relating to the conformity assessment procedures shall be drawn up, in electronic form, in an official language of the Member State where the notified body carrying out the procedures is established or in a language accepted by that body.		
		Article 3, first paragraph, point (6)(b)						
164	(b)	the following paragraph 3 is added:	(b)	the following paragraph 3 is added:	(b)	the following paragraph 3 is added:		

CLEAN	Commission Proposal	VS,EC	EP Mandate	VS,EC	Council Mandate	VS,EC	Draft Agreement
	Article 3, first paragraph, point (6)(b), amending provision, numbered paragraph (3), first subparagraph						
165	3. The manufacturer shall provide the notified body carrying out the conformity assessment procedure with all the information and documentation relating to conformity assessment procedures in electronic form.²	3.	The manufacturer shall provide the notified body carrying out the conformity assessment procedure with all the information and documentation relating to conformity assessment procedures in electronic form.²	3.	The manufacturer shall provide the notified body carrying out the conformity assessment procedure with all the information and documentation relating to conformity assessment procedures in electronic form.²		
	Article 3, first paragraph, point (6)(b), amending provision, numbered paragraph (3), second subparagraph						
166	;	;	;	;	;	;	
	Article 3, first paragraph, point (7)						
167	(7) in Article 16, the following paragraphs 5 and 6 are added:	(7)	in Article 16, the following paragraphs 5 and 6 are added:	(7)	in Article 16, the following paragraphs 5 and 6 are added:		
	Article 3, first paragraph, point (7), amending provision, numbered paragraph (5), first subparagraph						
168	5. The EU declaration of conformity shall be provided in a machine-readable and open format as defined in Article 2, points (13) and (14), of Directive (EU) 2019/1024 of the European Parliament and of the Council* and meet the requirements for digital labels set out in Article 11b(4), points (a) to (d).	5.	The EU declaration of conformity shall be provided in a machine-readable and open -format as defined in Article 2, points (13) and (14), of Directive (EU) 2019/1024 of the European Parliament and of the Council* and meet the requirements for digital labels set out in Article	5.	The EU declaration of conformity shall be provided in a machine-readable and open format as defined in Article 2, points (13) and (14), of Directive (EU) 2019/1024 of the European Parliament and of the Council* and meet the requirements for digital labels set out in Article 11b(4), points (a) to (d).		

CLEAN	Commission Proposal	VS.EC	EP Mandate	VS.EC	Council Mandate	VS.EC	Draft Agreement
			11b(4) , points (a) to (d) <u>through which it can be directly accessed.</u>				
	Article 3, first paragraph, point (7), amending provision, numbered paragraph (5), second subparagraph		Where a data carrier is used for providing access to the EU declaration of conformity, it should accompany the product in accordance with Article 11b(5) and be based on one of the electronic technical solutions which economic operators can use for providing the digital label established on the basis of Article 42(9). Economic operators providing the EU declaration of conformity using a data carrier shall not track, analyse or use any usage information for purposes other than what is absolutely necessary for providing the relevant information digitally.		Where a data carrier is used for providing access to the EU declaration of conformity, it should accompany the product in accordance with Article 11b(5) and be based on one of the electronic technical solutions which economic operators can use for providing the digital label established on the basis of Article 42(9). Economic operators providing the EU declaration of conformity using a data carrier shall not track, analyse or use any usage information for purposes other than what is absolutely necessary for providing the relevant information digitally.		
169							
	Article 3, first paragraph, point (7), amending provision, numbered paragraph (5), third subparagraph		Where a digital label is used in accordance with Article 11a, the data carrier used for the digital label shall also provide access to the EU declaration of conformity.		Where a digital label is used in accordance with Article 11a, the data carrier used for the digital label shall also provide access to the EU declaration of conformity.		
170							
	Article 3, first paragraph, point (7), amending provision, numbered paragraph (5), fourth subparagraph		*Directive (EU) 2019/1024 of the European Parliament and of the		*Directive (EU) 2019/1024 of the European Parliament and of the		
171							

	CLEAN	Commission Proposal	VS,EC	EP Mandate	VS,EC	Council Mandate	VS,EC	Draft Agreement
		Council of 20 June 2019 on open data and the re-use of public sector information, (OJ L 172, 26.06.2019, p. 56, ELI: http://data.europa.eu/eli/dir/2019/1024/oj).	VS,EC	Council of 20 June 2019 on open data and the re-use of public sector information, (OJ L 172, 26.06.2019, p. 56, ELI: http://data.europa.eu/eli/dir/2019/1024/oj).	VS,EC	Council of 20 June 2019 on open data and the re-use of public sector information, (OJ L 172, 26.06.2019, p. 56, ELI: http://data.europa.eu/eli/dir/2019/1024/oj).		
	Article 3, first paragraph, point (7), amending provision, numbered paragraph (6), first subparagraph							
172	6. Where other Union legislation applicable to EU fertilising products requires the economic operator to include the information that the product complies with the requirements set out in that legislation in a digital product passport or to upload the EU declaration of conformity in a digital product passport, the information set out in Annex V to be included in the EU declaration of conformity and any digital labelling information in accordance with Article 11b, if applicable, shall be provided only in that digital product passport.'	6. Where other Union legislation applicable to EU fertilising products requires the economic operator to include the information that the product complies with the requirements set out in that legislation in a digital product passport or to upload the EU declaration of conformity in a digital product passport, the information set out in Annex V to be included in the EU declaration of conformity and any digital labelling information in accordance with Article 11b, if applicable, shall be provided only in that digital product passport.'	VS,EC	6. Where other Union legislation applicable to EU fertilising products requires the economic operator to include the information that the product complies with the requirements set out in that legislation in a digital product passport or to upload the EU declaration of conformity in a digital product passport, the information set out in Annex V to be included in the EU declaration of conformity and any digital labelling information in accordance with Article 11b, if applicable, shall be provided only in that digital product passport.'	VS,EC	6. Where other Union legislation applicable to EU fertilising products requires the economic operator to include the information that the product complies with the requirements set out in that legislation in a digital product passport or to upload the EU declaration of conformity in a digital product passport, the information set out in Annex V to be included in the EU declaration of conformity and any digital labelling information in accordance with Article 11b, if applicable, shall be provided only in that digital product passport.'		
	Article 3, first paragraph, point (7), amending provision, numbered paragraph (6), second subparagraph							
173	;	;	VS,EC	;	VS,EC	;		
	Article 3, first paragraph, point (8)							

	CLEAN	Commission Proposal	VS.EC	EP Mandate	VS.EC	Council Mandate	VS.EC	Draft Agreement
174		(8) in Article 41(1), point (c) is replaced by the following:	(8) in Article 41(1), point (c) is replaced by the following:	(8) in Article 41(1), point (c) is replaced by the following:	(8) in Article 41(1), point (c) is replaced by the following:			
175		(c) the EU declaration of conformity has not been drawn up or has not been drawn up correctly, or the EU fertilising product is not accompanied by the internet address or data carrier through which the EU declaration of conformity can be accessed.	(c) the EU declaration of conformity has not been drawn up or has not been drawn up correctly, or the EU fertilising product is not accompanied by the internet address or data carrier through which the EU declaration of conformity can be accessed.	(c) the EU declaration of conformity has not been drawn up or has not been drawn up correctly, or the EU fertilising product is not accompanied by the internet address or data carrier through which the EU declaration of conformity can be accessed.	(c) the EU declaration of conformity has not been drawn up or has not been drawn up correctly, or the EU fertilising product is not accompanied by the internet address or data carrier through which the EU declaration of conformity can be accessed.			
176		(9) Article 42 is amended as follows:	(9) Article 42 is amended as follows:	(9) Article 42 is amended as follows:	(9) Article 42 is amended as follows:			
177		(a) in paragraph 4, the introductory statement is replaced by the following:	(a) in paragraph 4, the introductory statement is replaced by the following:	(a) in paragraph 4, the introductory statement is replaced by the following:	(a) in paragraph 4, the introductory statement is replaced by the following:			
178		The Commission may adopt delegated acts pursuant to paragraph 1 amending Annex II to add new micro-organisms or strains of micro-organisms, or additional processing methods to	The Commission may adopt delegated acts pursuant to paragraph 1 amending Annex II to add new micro-organisms or strains of micro-organisms, or additional processing methods to	The Commission may adopt delegated acts pursuant to paragraph 1 amending Annex II to add new micro-organisms or strains of micro-organisms, or additional processing methods to	The Commission may adopt delegated acts pursuant to paragraph 1 amending Annex II to add new micro-organisms or strains of micro-organisms, or additional processing methods to			

	CLEAN	Commission Proposal	VS.EC EP Mandate	VS.EC Council Mandate	VS.EC Draft Agreement
	the component material category for such organisms after having verified which strains of the additional micro-organism fulfil the criteria in paragraph 1, point (b), on the basis of the following data:²	the component material category for such organisms after having verified which strains of the additional micro-organism fulfil the criteria in paragraph 1, point (b), on the basis of the following data:²	the component material category for such organisms after having verified which strains of the additional micro-organism fulfil the criteria in paragraph 1, point (b), on the basis of the following data:²	the component material category for such organisms after having verified which strains of the additional micro-organism fulfil the criteria in paragraph 1, point (b), on the basis of the following data:²	
	Article 3, first paragraph, point (9)(a), amending provision, second paragraph				
179	;	;	;	;	;
	Article 3, first paragraph, point (9)(b)				
180	(b) the following paragraph 4a is inserted:	(b) the following paragraph 4a is inserted:	(b) the following paragraph 4a is inserted:	(b) the following paragraph 4a is inserted:	
	Article 3, first paragraph, point (9)(b), amending provision, numbered paragraph (4a), first subparagraph				
181	4a. The Commission may also adopt delegated acts pursuant to paragraph 1 amending Annex II to set out criteria and a methodology for the assessment of micro-organisms other than those listed in Annex II, which, if compliance with those criteria is demonstrated in the conformity assessment of the EU fertilising product in accordance with that methodology, may be used as component material in EU fertilising products. The criteria and methodology shall	4a. The Commission may also adopt delegated acts pursuant to paragraph 1 amending Annex II to set out criteria and a methodology for the assessment of micro-organisms other than those listed in Annex II, which, if the notifying bodies verify the compliance with those criteria in the conformity assessment of the EU fertilising product in accordance with that methodology, may be used as component material in EU fertilising products.	4a. The Commission may also adopt delegated acts pursuant to paragraph 1 amending Annex II to set out criteria and a methodology for the assessment of micro-organisms other than those listed in Annex II, which, if a manufacturer demonstrates the compliance with those criteria in the conformity assessment of the EU fertilising product in accordance with that methodology, may be used as component material in EU	4a. The Commission may also adopt delegated acts pursuant to paragraph 1 amending Annex II to set out criteria and a methodology for the assessment of micro-organisms other than those listed in Annex II, which, if a manufacturer demonstrates the compliance with those criteria in the conformity assessment of the EU fertilising product in accordance with that methodology, may be used as component material in EU	

CLEAN	Commission Proposal	VS,EC	EP Mandate	VS,EC	Council Mandate	VS,EC	Draft Agreement
	allow for verification that the micro-organisms fulfil the criteria in paragraph 1, point (b), and provide, as a minimum, for the consideration of the following elements:	VS,EC	component material in EU fertilising products. The criteria and methodology shall allow for verify that the micro-organisms fulfil the criteria in paragraph 1, point (b), and provide, as a minimum, for the consideration of the following elements:	VS,EC	fertilising products. The criteria and methodology shall allow for verify that the micro-organisms fulfil the criteria in paragraph 1, point (b), and provide, as a minimum, for the consideration of the following elements:		
	Article 3, first paragraph, point (9)(b), amending provision, numbered paragraph (4a), first subparagraph a						
181a							
	Article 3, first paragraph, point (9)(b), amending provision, numbered paragraph (4a), first subparagraph, point (a)						
182	(a) scientific literature reporting about safe production, conservation and use of the micro-organism;	VS,EC	(a) scientific literature reporting about safe production, conservation and use of the micro-organism;	VS,EC	(a) scientific literature reporting about safe production, conservation and use of the micro-organism;		
	Article 3, first paragraph, point (9)(b), amending provision, numbered paragraph (4a), first subparagraph, point (b)						
183	(b) taxonomic relation of the micro-organism to micro-organisms species fulfilling the requirements for a Qualified Presumption of Safety as established by the European Food Safety Authority;	VS,EC	(b) taxonomic relation of the micro-organism to micro-organisms species fulfilling the requirements for a Qualified Presumption of Safety as established by the European Food Safety Authority;	VS,EC	(b) taxonomic relation of the micro-organism to micro-organisms species fulfilling the requirements for a Qualified Presumption of Safety as established by the European Food Safety Authority;		
	Article 3, first paragraph, point (9)(b), amending provision, numbered paragraph (4a), first subparagraph, point (c)						
184	(c) information on the production process of the micro-organism, including, where relevant, the composition of the	VS,EC	(c) information on the production process of the micro-organism, including, where relevant, the composition of the	VS,EC	(c) information on the production process of the micro-organism, including, where relevant, the composition of the		

	CLEAN	Commission Proposal	VS.EC	EP Mandate	VS.EC	Council Mandate	VS.EC	Draft Agreement
		cultivation medium, processing methods such as spray drying, fluid-bed drying, static drying, centrifugation, deactivation by heat, filtration and grinding;		cultivation medium, processing methods such as spray drying, fluid-bed drying, static drying, centrifugation, deactivation by heat, filtration and grinding;		cultivation medium, processing methods such as spray drying, fluid-bed drying, static drying, centrifugation, deactivation by heat, filtration and grinding;		
	Article 3, first paragraph, point (9)(b), amending provision, numbered paragraph (4a), first subparagraph, point (d)							
185	(d) information on the identity and residue levels of residual intermediates, toxins or microbial metabolites in the component material;	(d) information on the identity and residue levels of residual intermediates, toxins or microbial metabolites in the component material;		(d) information on the identity and residue levels of residual intermediates, toxins or microbial metabolites in the component material;		(d) information on the identity and residue levels of residual intermediates, toxins or microbial metabolites in the component material;		
	Article 3, first paragraph, point (9)(b), amending provision, numbered paragraph (4a), first subparagraph, point (e)							
186	(e) natural occurrence, survival and mobility in the environment;	(e) natural occurrence, survival and mobility in the environment;		(e) natural occurrence, survival and mobility in the environment;		(e) natural occurrence, survival and mobility in the environment;		
	Article 3, first paragraph, point (9)(b), amending provision, numbered paragraph (4a), first subparagraph, point (f)							
187	(f) susceptibility to all relevant antimicrobial agents as defined in the Annex, Introduction to Part B, point (ii)(28), to Commission Regulation (EU) No 283/2013*, with the exception of intrinsic resistance.	(f) susceptibility to all relevant antimicrobial agents as defined in the Annex, Introduction to Part B, point (ii)(28), to Commission Regulation (EU) No 283/2013*, with the exception of intrinsic resistance.		(f) susceptibility to all relevant antimicrobial agents as defined in the Annex, Introduction to Part B, point (ii)(28), to Commission Regulation (EU) No 283/2013*, with the exception of intrinsic resistance.		(f) susceptibility to all relevant antimicrobial agents as defined in the Annex, Introduction to Part B, point (ii)(28), to Commission Regulation (EU) No 283/2013*, with the exception of intrinsic resistance.		
	Article 3, first paragraph, point (9)(b), amending provision, numbered paragraph (4a), second subparagraph							
188	*Commission Regulation (EU) No 283/2013 of 1 March 2013 setting out the data requirements for active substances, in accordance with Regulation (EC) No	*Commission Regulation (EU) No 283/2013 of 1 March 2013 setting out the data requirements for active substances, in accordance with Regulation (EC) No		*Commission Regulation (EU) No 283/2013 of 1 March 2013 setting out the data requirements for active substances, in accordance with Regulation (EC) No		*Commission Regulation (EU) No 283/2013 of 1 March 2013 setting out the data requirements for active substances, in accordance with Regulation (EC) No		

CLEAN	Commission Proposal	VS,EC	EP Mandate	VS,EC	Council Mandate	VS,EC	Draft Agreement
	1107/2009 of the European Parliament and of the Council concerning the placing of plant protection products on the market (OJ L 93, 3.4.2013, p. 1, ELI: http://data.europa.eu/eli/reg/2013/283/oj).	1107/2009 of the European Parliament and of the Council concerning the placing of plant protection products on the market (OJ L 93, 3.4.2013, p. 1, ELI: http://data.europa.eu/eli/reg/2013/283/oj).	1107/2009 of the European Parliament and of the Council concerning the placing of plant protection products on the market (OJ L 93, 3.4.2013, p. 1, ELI: http://data.europa.eu/eli/reg/2013/283/oj).	1107/2009 of the European Parliament and of the Council concerning the placing of plant protection products on the market (OJ L 93, 3.4.2013, p. 1, ELI: http://data.europa.eu/eli/reg/2013/283/oj).	1107/2009 of the European Parliament and of the Council concerning the placing of plant protection products on the market (OJ L 93, 3.4.2013, p. 1, ELI: http://data.europa.eu/eli/reg/2013/283/oj).		
	Article 3, first paragraph, point (9)(b), amending provision, numbered paragraph (4a), third subparagraph						
189	;	;	;	;	;		
	Article 3, first paragraph, point (9)(ba)						
189a		<u>(ba) The following paragraph 4aa is inserted:</u>					
	Article 3, first paragraph, point (9)(ba), amending provision, point (a)						
189b		<u>4aa The Commission may also adopt delegated acts pursuant to paragraph 1 amending Annex II to establish general criteria and a methodology for the assessment of materials and processing methods other than those already listed in Annex II, excluding micro-organisms, which may be used as component materials in EU fertilising products where compliance with those criteria is demonstrated in the conformity assessment. The criteria and</u>					

CLEAN	Commission Proposal	VS,EC	EP Mandate	VS,EC	Council Mandate	VS,EC	Draft Agreement
			<u>methodology shall, as a minimum, provide for the consideration of scientific or technical information supporting safe sourcing, processing and use of the material.</u>				
Article 3, first paragraph, point (9)(bb)							
189c			<u>(bb) The following paragraph 4ab is inserted:</u>				
Article 3, first paragraph, point (9)(bb), amending provision, point (a)							
189d			<u>4ab By [12 months after the date of entry into force of this amending Regulation] the Commission shall adopt a delegated act to establish the criteria and methodology and upon assessment to create a corresponding new category for materials and processing methods other than those already listed in Annex II, excluding micro-organisms, which may be used as component materials in EU fertilising products provided that compliance with those criteria is demonstrated in the conformity assessment. The criteria and methodology shall aim to ensure safe sourcing, processing and use of the material. This new</u>				

CLEAN	Commission Proposal	VS,EC	EP Mandate	VS,EC	Council Mandate	VS,EC	Draft Agreement
			<u>category shall be established without prejudice to the application of other component material categories set out in Annex II.</u>				
Article 3, first paragraph, point (9)(bc)							
189e			<u>(bc) The following paragraph 5a is inserted:</u>				
Article 3, first paragraph, point (9)(bc), amending provision, point (a)							
189f			<u>5a For the purposes of this Regulation, products derived from animal by-products that are used solely as component materials in EU fertilising products may be made available on the market only where they originate from animal by-products or derived products that have reached an end point in the manufacturing chain in accordance with Article 5(2) of Regulation (EC) No 1069/2009. When determining such end points for derived products intended for use in EU fertilising products, the Commission shall ensure that: a) the processing and safety criteria, as well as any necessary risk-mitigation</u>				

CLEAN	Commission Proposal	VS,EC EP Mandate	VS,EC Council Mandate	VS,EC Draft Agreement
		<p><u>measures, are proportionate and sufficient to ensure a high level of protection for human health and the environment, specifically relevant to fertilising uses;</u></p> <p><u>b) those criteria and measures are appropriate to the risks arising from the application of fertilising products to soil and water; and</u></p> <p><u>c) those criteria and measures are aligned with the requirements set out in Article 42(5) of this Regulation.</u></p> <p><u>Where existing delegated or implementing acts adopted under Regulation (EC) No 1069/2009 do not meet those conditions, the Commission shall, where appropriate and in accordance with the procedures laid down in that Regulation, review them and, where necessary, amend them to ensure such consistency. For that purpose, the Commission shall, where relevant, request scientific opinions from the European Food Safety Authority to assess risks specific to fertilising uses, taking into account exposure pathways that are distinct from those associated with feed or other uses.</u></p>		

CLEAN	Commission Proposal	VS,EC	EP Mandate	VS,EC	Council Mandate	VS,EC	Draft Agreement
Article 3, first paragraph, point (9)(bc), amending provision, point (b)		VS,EC	EP Mandate	VS,EC	Council Mandate	VS,EC	Draft Agreement
189g			<u>(bb) The following paragraph is added: '8a.'</u>				
Article 3, first paragraph, point (9)(bc), amending provision, point (c)							
189h			<u>(8a) The Commission shall adopt the first delegated act pursuant to paragraph 4a by ... [18 months after the entry into force of this Regulation].</u>				
Article 3, first paragraph, point (10)							
190	(10) Article 43 is deleted;	(10)	Article 43 is deleted;	(10)	In Article 43, the following paragraph is addedis deleted;		
Article 3, first paragraph, point (10a)							
190a			<u>(10a) the following Article 49a is inserted:</u>				
Article 3, first paragraph, point (10a), amending provision, first paragraph							
190b			<u>Review</u> <u>The Commission shall periodically assess whether the requirements governing the treatment of materials intended for use in fertilising products remain appropriate and, where necessary, adapt them in light of scientific and technical advances, taking into account national</u>				

CLEAN	Commission Proposal	VS,EC	EP Mandate	VS,EC	Council Mandate	VS,EC	Draft Agreement
			<u>practices, and the objectives of Regulation (EU) 2019/1009. This review shall be carried out for the first time no later than two years after the date of application of this amending Regulation.'</u>				
Article 3, first paragraph, point (10b)							
190c			<u>(10b) the following Article 49aa is inserted:</u>				
Article 3, first paragraph, point (10b), amending provision, first paragraph							
190d			<u>Report</u> <u>By [insert date: 12 months after entry into force], the Commission shall submit a report to the European Parliament and the Council assessing the requirements for fertilising product blends where the primary component is one or more growing media belonging to PFC 4.</u> <u>The report shall, in particular, evaluate the conditions for adjusting nutrient content, pH value or biological activity, updated labelling requirements, and appropriate conformity assessment procedures for such blends and it may accompany,</u>				

CLEAN	Commission Proposal	vs.EC	EP Mandate	vs.EC	Council Mandate	vs.EC	Draft Agreement
			<i>where appropriate, amendments to this Regulation.</i>				
Article 3, first paragraph, point (11)							
191	(11) Annexes I, II and IV to Regulation (EU) 2019/1009 are amended in accordance with Annex IV to this Regulation.	(11)	Annexes I, II and IV to Regulation (EU) 2019/1009 are amended in accordance with Annex IV to this Regulation.	(11)	Annexes I, II and IV to Regulation (EU) 2019/1009 are amended in accordance with Annex IV to this Regulation.		
Article 4							
192	Article 4 Transitional provisions	Article 4 Transitional provisions	Article 4 Transitional provisions	Article 4 Transitional provisions	Article 4 Transitional provisions		
Article 4(1), first subparagraph							
193	1. By way of derogation from section 1.5.2.4.1 and section 1.5.2.4.2 of Annex I to Regulation (EC) No 1272/2008 as applicable on 9 December 2024 substances and mixtures may until 30 June 2026 be classified, labelled and packaged in accordance with Regulation (EC) No 1272/2008 as amended by points (5), (6) and (7) of Annex I to this Regulation.	1.	By way of derogation from section 1.5.2.4.1 and section 1.5.2.4.2 of Annex I to Regulation (EC) No 1272/2008 as applicable on 9 December 2024 substances and mixtures may until 30 June 2026 be classified, labelled and packaged in accordance with Regulation (EC) No 1272/2008 as amended by points (5), (6) and (7) of Annex I to this Regulation.	1.	By way of derogation from section 1.5.2.4.1 and section 1.5.2.4.2 of Annex I to Regulation (EC) No 1272/2008 as applicable on 9 December 2024 substances and mixtures may until 30 June 2026 be classified, labelled and packaged in accordance with Regulation (EC) No 1272/2008 as amended by points (5), (6) and (7) of Annex I to this Regulation.		
Article 4(1), second subparagraph							
194	By way of derogation from Article 30 and Article 48 of Regulation (EC) No 1272/2008 and Part 5 of Annex II to Regulation (EC) No 1272/2008 as applicable on 9 December 2024, substances and	By way of derogation from Article 30 and Article 48 of Regulation (EC) No 1272/2008 and Part 5 of Annex II to Regulation (EC) No 1272/2008 as applicable on 9 December 2024, substances and	By way of derogation from Article 30 and Article 48 of Regulation (EC) No 1272/2008 and Part 5 of Annex II to Regulation (EC) No 1272/2008 as applicable on 9 December 2024, substances and	By way of derogation from Article 30 and Article 48 of Regulation (EC) No 1272/2008 and Part 5 of Annex II to Regulation (EC) No 1272/2008 as applicable on 9 December 2024, substances and	By way of derogation from Article 30 and Article 48 of Regulation (EC) No 1272/2008 and Part 5 of Annex II to Regulation (EC) No 1272/2008 as applicable on 9 December 2024, substances and		

CLEAN	Commission Proposal	VS.EC EP Mandate	VS.EC Council Mandate	VS.EC Draft Agreement
	<p>mixtures may until 31 December 2027 be classified, labelled and packaged in accordance with Regulation (EC) No 1272/2008 as amended by Article 1, points (5), (7) and (8) of this Regulation and point (9) of Annex I to this Regulation.</p>	<p>mixtures may until 31 December 2027 be classified, labelled and packaged in accordance with Regulation (EC) No 1272/2008 as amended by Article 1, points (5), (7) and (8) of this Regulation and point (9) of Annex I to this Regulation.</p>	<p>mixtures may until 31 December 2027 be classified, labelled and packaged in accordance with Regulation (EC) No 1272/2008 as amended by Article 1, points (5), (7) and (8) of this Regulation and point (9) of Annex I to this Regulation.</p>	
Article 4(1), third subparagraph				
195	<p>By way of derogation from Article 17(1), Article 25(6) of Regulation (EC) No 1272/2008, section 1.5.1.2 and section 1.6 of Annex I to Regulation (EC) No 1272/2008 as applicable on [OP: please insert the date of the day before the date of entry into force of this Regulation], substances and mixtures may until [OP: please insert the date of the last day of the month following 35 months after the date of entry into force of this Regulation] be classified, labelled and packaged in accordance with Regulation (EC) No 1272/2008 as amended by Article 1, points (2) and (3) of this Regulation and points (3) and (8) of Annex I to this Regulation.</p>	<p>By way of derogation from Article 17(1), Article 25(6) of Regulation (EC) No 1272/2008, section 1.5.1.2 and section 1.6 of Annex I to Regulation (EC) No 1272/2008 as applicable on [OP: please insert the date of the day before the date of entry into force of this Regulation], substances and mixtures may until [OP: please insert the date of the last day of the month following 35 months after the date of entry into force of this Regulation] be classified, labelled and packaged in accordance with Regulation (EC) No 1272/2008 as amended by Article 1, points (2) and (3) of this Regulation and points (3) and (8) of Annex I to this Regulation.</p>	<p>By way of derogation from Article 17(1), Article 25(6) of Regulation (EC) No 1272/2008, section 1.5.1.2 and section 1.6 of Annex I to Regulation (EC) No 1272/2008 as applicable on [OP: please insert the date of the day before the date of entry into force of this Regulation], substances and mixtures may until [OP: please insert the date of the last day of the month following 35 months after the date of entry into force of this Regulation] be classified, labelled and packaged in accordance with Regulation (EC) No 1272/2008 as amended by Article 1, points (2) and (3) of this Regulation and points (3) and (8) of Annex I to this Regulation.</p>	
Article 4(2)				

	CLEAN	Commission Proposal	VS.EC	EP Mandate	VS.EC	Council Mandate	VS.EC	Draft Agreement
196		2. Member States shall not impede the making available on the market of products which were placed on the market in accordance with Regulation (EU) 2019/1009 before [OP: please insert 24 months after entry into force of this amending Regulation]).	VS.EC	2. Member States shall not impede the making available on the market of products which were placed on the market in accordance with Regulation (EU) 2019/1009 before [OP: please insert 24 months after entry into force of this amending Regulation]).	VS.EC	2. Member States shall not impede the making available on the market of products which were placed on the market in accordance with Regulation (EU) 2019/1009 before [OP: please insert 24 months after entry into force of this amending Regulation]).		
Article 5								
197		Article 5 Entry into force and application		Article 5 Entry into force and application		Article 5 Entry into force and application		
Article 5(1)								
198		1. This Regulation shall enter into force on the twentieth day following that of its publication in the Official Journal of the European Union.		1. This Regulation shall enter into force on the twentieth day following that of its publication in the Official Journal of the European Union.		1. This Regulation shall enter into force on the twentieth day following that of its publication in the Official Journal of the European Union.		
Article 5(2)								
199		2. Points (4) to (7) of Annex I shall apply from 1 July 2026.		2. Points (4) to (7) of Annex I shall apply from 1 July 2026.		2. Points (4) to (7) of Annex I shall apply from 1 July 2026.		
Article 5(3)								
200		3. Article 1, points (5) to (8) and points (1), (2) and (9) of Annex I shall apply from 1 January 2028.		3. Article 1, points (5) to (8) and points (1), (2) and (9) of Annex I shall apply from 1 January 2028.		3. Article 1, points (5) to (8) and points (1), (2) and (9) of Annex I shall apply from 1 January 2028.		
Article 5(4)								

	CLEAN	Commission Proposal	VS,EC	EP Mandate	VS,EC	Council Mandate	VS,EC	Draft Agreement
201		4. Article 1, points (1), (2) and (3), points (3) and (8) of Annex I shall apply from [OP: please insert the date of 36 months after the entry into force of this Regulation)]	VS,EC	4. Article 1, points (1), (2) and (3), points (3) and (8) of Annex I shall apply from [OP: please insert the date of 36 months after the entry into force of this Regulation)]	VS,EC	4. Article 1, points (1), (2) and (3), points (3) and (8) of Annex I shall apply from [OP: please insert the date of 36 months after the entry into force of this Regulation)]		
	Article 5(5)							
202		5. Article 2, point (1) to (8) shall apply from [OP: please insert the date of entry into force of this Regulation)]	VS,EC	5. Article 2, point (1) to (8) shall apply from [OP: please insert the date of entry into force of this Regulation)]	VS,EC	5. Article 2, point (1) to (8) shall apply from [OP: please insert the date of entry into force of this Regulation)]		
	Article 5(6)							
203		6. Article 3, point (1) to (8), and Annex IV, point (1) and (3), shall apply from [OP: please insert 24 months after entry into force of this Regulation].	VS,EC	6. Article 3, point (1) to (8), and Annex IV, point (1) and (3), shall apply from [OP: please insert 24 months after entry into force of this Regulation].	VS,EC	6. Article 3, point (1) to (8), and Annex IV, point (1) and (3), shall apply from [OP: please insert 24 months after entry into force of this Regulation].		
	Article 5, seventh paragraph							
204		This Regulation shall be binding in its entirety and directly applicable in all Member States.	VS,EC	This Regulation shall be binding in its entirety and directly applicable in all Member States.	VS,EC	This Regulation shall be binding in its entirety and directly applicable in all Member States.		
	Formula							
205		Done at Strasbourg,	VS,EC	Done at Strasbourg,	VS,EC	Done at Strasbourg,		
	Formula							
206		For the European Parliament	VS,EC	For the European Parliament	VS,EC	For the European Parliament		
	Formula							
207		[...]	VS,EC	[...]	VS,EC	[...]		

	CLEAN	Commission Proposal	VS.EC	EP Mandate	VS.EC	Council Mandate	VS.EC	Draft Agreement
Formula								
208	The President	The President	The President	The President	The President	The President		
Formula								
209	For the Council	For the Council	For the Council	For the Council	For the Council	For the Council		
Formula								
210	[...]	[...]	[...]	[...]	[...]	[...]		
ANNEX I								
211	ANNEX I	ANNEX I	ANNEX I	ANNEX I	ANNEX I	ANNEX I		
ANNEX I, first paragraph								
212	Annexes I and II to Regulation (EC) No 1272/2008 are amended as follows:	Annexes I and II to Regulation (EC) No 1272/2008 are amended as follows:	Annexes I and II to Regulation (EC) No 1272/2008 are amended as follows:	Annexes I and II to Regulation (EC) No 1272/2008 are amended as follows:	Annexes I and II to Regulation (EC) No 1272/2008 are amended as follows:	Annexes I and II to Regulation (EC) No 1272/2008 are amended as follows:		
ANNEX I, second paragraph								
213	(1) in Annex I, section 1.2.1.4 is replaced by the following:	(1) in Annex I, section 1.2.1.4 is replaced by the following:	(1) in Annex I, section 1.2.1.4 is replaced by the following:	(1) in Annex I, section 1.2.1.4 is replaced by the following:	(1) in Annex I, section 1.2.1.4 is replaced by the following:	(1) in Annex I, section 1.2.1.4 is replaced by the following:		
ANNEX I, second paragraph, amending provision, numbered paragraph (1.2.1.4)								
214	‘ 1.2.1.4.	‘ 1.2.1.4.	‘ 1.2.1.4.	‘ 1.2.1.4.	‘ 1.2.1.4.	‘ 1.2.1.4.		
ANNEX I, second paragraph, amending provision, numbered paragraph (1.2.1.4), first subparagraph								
215	1.2.1.4. The dimensions of the label and of each pictogram shall be as follows:	1.2.1.4. The dimensions of the label and of each pictogram shall be as follows:	1.2.1.4. The dimensions of the label and of each pictogram shall be as follows:	1.2.1.4. The dimensions of the label and of each pictogram shall be as follows:	1.2.1.4. The dimensions of the label and of each pictogram shall be as follows:	1.2.1.4. The dimensions of the label and of each pictogram shall be as follows:		
ANNEX I, second paragraph, amending provision, numbered paragraph (1.2.1.4), second subparagraph								
216	Table 1.3	Table 1.3	Table 1.3	Table 1.3	Table 1.3	Table 1.3		
ANNEX I, second paragraph, amending provision, numbered paragraph (1.2.1.4), third subparagraph								

	CLEAN	Commission Proposal	VS,EC	EP Mandate	VS,EC	Council Mandate	VS,EC	Draft Agreement
217	Minimum dimensions of labels and pictograms	Minimum dimensions of labels and pictograms	VS,EC	Minimum dimensions of labels and pictograms	VS,EC	Minimum dimensions of labels and pictograms	VS,EC	
	ANNEX I, second paragraph, amending provision, numbered paragraph (1.2.1.4), Table							
218	Table	Table				Table		
	ANNEX I, second paragraph, amending provision, numbered paragraph (1.2.1.4), fourth subparagraph							
219	' ; ,	' ; ,		' ; ,		' ; ,		
	ANNEX I, 2 paragraph							
220	(2) in Annex I, section 1.2.1.5 is deleted;	(2) in Annex I, section 1.2.1.5 is deleted <u>replaced by the following: The text on the label shall be legible. For the purpose of this section, a label shall be considered legible if the physical appearance of information, by means of which the information is visually accessible and which is determined by various elements, inter alia, font size, letter spacing, spacing between lines, stroke width, type colour, typeface, width-height ratio of the letters, the surface of the material and significant contrast between the print and the background. It shall have at least the following characteristics: (a) printed in a contrasting colour compared to the background; (b) a single</u>		(2) in Annex I, section 1.2.1.5 is deleted <u>replaced by the following:</u>		(2) in Annex I, section 1.2.1.5 is deleted <u>replaced by the following:</u>		

CLEAN	Commission Proposal	VS,EC EP Mandate	VS,EC Council Mandate	VS,EC Draft Agreement
		<p><u>typeface that is easily legible and without serifs shall be used; (c) appropriate letter spacing for the selected typeface to be easily legible; (d) appropriate line spacing for the selected typeface to be easily readable and to ensure that lines of text do not overlap; (e) an appropriate font size with regard to the size of the label and the required label elements and the intended user.</u></p> <p><u>For substances or mixtures made available on the market for the general public, the label elements referred to in Article 17(1) shall use a font size where the x-height is equal to or greater than 1.2 mm. However, where the contents of the package do not exceed 125 ml, the label elements referred to in Article 17(1) may use a font size where the x-height is equal to or greater than 0.9 mm. These characteristics shall be further clarified in the ECHA guidance on legibility and label presentation to be developed under this Regulation.</u></p> <p><u>Compliance with this section shall be assessed on the basis of the overall legibility of the label</u></p>		

CLEAN	Commission Proposal	VS.EC	EP Mandate	VS.EC	Council Mandate	VS.EC	Draft Agreement
			<i>and applied in a manner compatible with the physical characteristics and technical limitations of the packaging, including size as well as the intended user.</i>				
ANNEX I, 3 paragraph							
221	(3) in Annex I, section 1.5.1.2 is replaced by the following:	(3) in Annex I, section 1.5.1.2 is replaced by the following:	(3) in Annex I, section 1.5.1.2 is replaced by the following:	(3) in Annex I, section 1.5.1.2 is replaced by the following:	(3) in Annex I, section 1.5.1.2 is replaced by the following:		
ANNEX I, 3 paragraph, amending provision, numbered paragraph (1.5.1.2)							
222	1.5.1.2. Where section 1.5.1.1 applies, the label on any inner packaging shall contain at least the hazard pictograms, the signal words, the product identifier referred to in Article 18(2) for substances or the trade name or designation referred to in Article 18(3), point (a) for mixtures, and the name and digital contact of the suppliers of the substance or mixture;	1.5.1.2. Where section 1.5.1.1 applies, the label on any inner packaging shall contain at least the hazard pictograms, the signal words, the product identifier referred to in Article 18(2) for substances or the trade name or designation referred to in Article 18(3), point (a) for mixtures, and the name, the digital contact <u>and the telephone number</u> of the suppliers of the substance or mixture: <u>unless this telephone number is directly available through the digital contact;</u>	1.5.1.2. Where section 1.5.1.1 applies, the label on any inner packaging shall contain at least the hazard pictograms, the signal words, the product identifier referred to in Article 18(2) for substances or the trade name or designation referred to in Article 18(3), point (a) for mixtures, and the name and digital contact of the suppliers of the substance or mixture, and the telephone number, unless this telephone number is immediately available through the digital contact;				
ANNEX I, 4 paragraph							

	CLEAN	Commission Proposal	VS.EC	EP Mandate	VS.EC	Council Mandate	VS.EC	Draft Agreement
223	(4) the heading of section 1.5.2.4 is replaced by the following:	(4) the heading of section 1.5.2.4 is replaced by the following:			(4) the heading of section 1.5.2.4 is replaced by the following:			
ANNEX I, 4 paragraph, amending provision, numbered paragraph (1.5.2.4)								
224	1.5.2.4. Labelling of packages where the contents do not exceed 10 ml ;	1.5.2.4. Labelling of packages where the contents do not exceed 10 ml ;			1.5.2.4. Labelling of packages where the contents do not exceed 10 ml ;			
ANNEX I, 5 paragraph								
225	(5) in Annex I, section 1.5.2.4.1 is replaced by the following:	(5) in Annex I, section 1.5.2.4.1 is replaced by the following:			(5) in Annex I, section 1.5.2.4.1 is replaced by the following:			
ANNEX I, 5 paragraph, amending provision, numbered paragraph (1.5.2.4.1)								
226	1.5.2.4.1. The label elements set out in Article 17 may be omitted from the inner packaging where the contents of the inner packaging do not exceed 10 ml, the outer packaging meets the requirements set out in Article 17(1) and any of the following applies:	1.5.2.4.1. The label elements set out in Article 17 may be omitted from the inner packaging where the contents of the inner packaging do not exceed 10 ml, the outer packaging meets the requirements set out in Article 17(1) and any of the following applies:			1.5.2.4.1. The label elements set out in Article 17(1) may be omitted from the inner packaging where the contents of the inner packaging do not exceed 10 ml, the outer packaging meets the requirements set out in Article 17(1) and any of the following applies:			
ANNEX I, 5 paragraph, amending provision, numbered paragraph (1.5.2.4.1), point (a)								
227	(a) the substance or mixture is placed on the market for scientific	(a) the substance or mixture is placed on the market for scientific			(a) the substance or mixture is placed on the market for scientific			

CLEAN	Commission Proposal	VS.EC	EP Mandate	VS.EC	Council Mandate	VS.EC	Draft Agreement
	research and development or quality control analysis;		research and development or quality control analysis;		research and development or quality control analysis;		
	ANNEX I, 5 paragraph, amending provision, numbered paragraph (1.5.2.4.1), point (b)						
228	(b) the substance or mixture requires labelling in accordance with Part 1 or 2 of Annex II, except for section 2.8 of Part 2 of Annex II, and is not classified in any of the following hazard classes and categories:	(b) the substance or mixture requires labelling in accordance with Part 1 or 2 of Annex II, <i>except for section 2.8 of Part 2 of Annex II</i> , and is not classified in any of the following hazard classes and categories:	(b) the substance or mixture requires labelling in accordance with Part 1 or 2 of Annex II, <i>except for section 2.8 of Part 2 of Annex II</i> , and is not classified in any of the following hazard classes and categories:	(b) the substance or mixture requires labelling in accordance with Part 1 or 2 of Annex II, <i>except for section 2.8 of Part 2 of Annex II</i> , and is not classified in any of the following hazard classes and categories:			
	ANNEX I, 5 paragraph, amending provision, numbered paragraph (1.5.2.4.1), point (b)(a)						
229	(a) acute toxicity, any category;	(a) acute toxicity, any category;	(a) acute toxicity, any category;	(a) acute toxicity, any category;	(a) acute toxicity, any category;		
	ANNEX I, 5 paragraph, amending provision, numbered paragraph (1.5.2.4.1), point (b)(b)						
230	(b) specific target organ toxicity – single exposure, categories 1 and 2;	(b) specific target organ toxicity – single exposure, categories 1 and 2;	(b) specific target organ toxicity – single exposure, categories 1 and 2;	(b) specific target organ toxicity – single exposure, categories 1 and 2;	(b) specific target organ toxicity – single exposure, categories 1 and 2;		
	ANNEX I, 5 paragraph, amending provision, numbered paragraph (1.5.2.4.1), point (b)(c)						
231	(c) specific target organ toxicity – repeated exposure, any category;	(c) specific target organ toxicity – repeated exposure, any category;	(c) specific target organ toxicity – repeated exposure, any category;	(c) specific target organ toxicity – repeated exposure, any category;	(c) specific target organ toxicity – repeated exposure, any category;		
	ANNEX I, 5 paragraph, amending provision, numbered paragraph (1.5.2.4.1), point (b)(d)						
232	(d) skin corrosion, category 1, any sub-category;	(d) skin corrosion, category 1, any sub-category;	(d) skin corrosion, category 1, any sub-category;	(d) skin corrosion, category 1, any sub-category;	(d) skin corrosion, category 1, any sub-category;		
	ANNEX I, 5 paragraph, amending provision, numbered paragraph (1.5.2.4.1), point (b)(e)						
233	(e) serious eye damage, category 1;	(e) serious eye damage, category 1;	(e) serious eye damage, category 1;	(e) serious eye damage, category 1;	(e) serious eye damage, category 1;		

CLEAN	Commission Proposal	VS,EC	EP Mandate	VS,EC	Council Mandate	VS,EC	Draft Agreement
	ANNEX I, 5 paragraph, amending provision, numbered paragraph (1.5.2.4.1), point (b)(f)						
234	(f) respiratory sensitisation, any category;	(f)	respiratory sensitisation, any category;	(f)	respiratory sensitisation, any category;	(f)	respiratory sensitisation, any category;
	ANNEX I, 5 paragraph, amending provision, numbered paragraph (1.5.2.4.1), point (b)(g)						
235	(g) aspiration hazard;	(g)	aspiration hazard;	(g)	aspiration hazard;	(g)	aspiration hazard;
	ANNEX I, 5 paragraph, amending provision, numbered paragraph (1.5.2.4.1), point (b)(h)						
236	(h) germ cell mutagenicity, any category;	(h)	germ cell mutagenicity, any category;	(h)	germ cell mutagenicity, any category;	(h)	germ cell mutagenicity, any category;
	ANNEX I, 5 paragraph, amending provision, numbered paragraph (1.5.2.4.1), point (b)(i)						
237	(i) carcinogenicity, any category;	(i)	carcinogenicity, any category;	(i)	carcinogenicity, any category;	(i)	carcinogenicity, any category;
	ANNEX I, 5 paragraph, amending provision, numbered paragraph (1.5.2.4.1), point (b)(j)						
238	(j) reproductive toxicity, any category;	(j)	reproductive toxicity, any category;	(j)	reproductive toxicity, any category;	(j)	reproductive toxicity, any category;
	ANNEX I, 5 paragraph, amending provision, numbered paragraph (1.5.2.4.1), point (b)(k)						
239	(k) endocrine disruption for human health, any category.;	(k)	endocrine disruption for human health, any category.;	(k)	endocrine disruption for human health, any category.;	(k)	endocrine disruption for human health, any category.;
	ANNEX I, 6 paragraph						
240	(6) in Annex I, section 1.5.2.4.2 is replaced by the following:	(6)	in Annex I, section 1.5.2.4.2 is replaced by the following:	(6)	in Annex I, section 1.5.2.4.2 is replaced by the following:	(6)	in Annex I, section 1.5.2.4.2 is replaced by the following:
	ANNEX I, 6 paragraph, amending provision, numbered paragraph (1.5.2.4.2)						
241	,	,	,	,	,	,	,

	CLEAN	Commission Proposal	VS,EC	EP Mandate	VS,EC	Council Mandate	VS,EC	Draft Agreement
	1.5.2.4.2. 1.5.2.4.1	Where section applies, the label on the inner packaging shall contain the product identifier referred to in Article 18(2) for substances or the trade name or designation referred to in Article 18(3), point (a) for mixtures, and, where applicable, the hazard pictograms ‘GHS01’, ‘GHS05’, ‘GHS06’ or ‘GHS08’.	1.5.2.4.2. 1.5.2.4.1	Where section applies, the label on the inner packaging shall contain the product identifier referred to in Article 18(2) for substances or the trade name or designation referred to in Article 18(3), point (a) for mixtures, and, where applicable, the hazard pictograms ‘GHS01’, ‘GHS05’, ‘GHS06’ or ‘GHS08’.	1.5.2.4.2. 1.5.2.4.1	Where section applies, the label on the inner packaging shall contain the product identifier referred to in Article 18(2) for substances or the trade name or designation referred to in Article 18(3), point (a) for mixtures, and, where applicable, the hazard pictograms ‘GHS01’, ‘GHS05’, ‘GHS06’ or ‘GHS08’.		
		Where more than two pictograms are assigned, ‘GHS06’ and ‘GHS08’ may take precedence over ‘GHS01’ and ‘GHS05’ ;	Where more than two pictograms are assigned, ‘GHS06’ and ‘GHS08’ may take precedence over ‘GHS01’ and ‘GHS05’ ;	Where more than two pictograms are assigned, ‘GHS06’ and ‘GHS08’ may take precedence over ‘GHS01’ and ‘GHS05’ ;	Where more than two pictograms are assigned, ‘GHS06’ and ‘GHS08’ may take precedence over ‘GHS01’ and ‘GHS05’ ;			
	ANNEX I, 7 paragraph							
242	(7) 1.5.2.4.3	in Annex I, section 1.5.2.4.3 is added:	(7) 1.5.2.4.3	in Annex I, section 1.5.2.4.3 is added:	(7) 1.5.2.4.3	in Annex I, section 1.5.2.4.3 is added:		
	ANNEX I, 7 paragraph, amending provision, numbered paragraph (1.5.2.4.3)							
243	1.5.2.4.3.	The label elements set out in Article 17(1) may be omitted from the package provided that the following conditions are met:	1.5.2.4.3.	The label elements set out in Article 17(1) may be omitted from the package provided that the following conditions are met:	1.5.2.4.3.	The label elements set out in Article 17(1) may be omitted from the package provided that the following conditions are met:		
	ANNEX I, 7 paragraph, amending provision, numbered paragraph (1.5.2.4.3), point (a)							
244	(a)	the contents of the package do not exceed 10 ml;	(a)	the contents of the package do not exceed 10 ml;	(a)	the contents of the package do not exceed 10 ml;		

	CLEAN	Commission Proposal	VS.EC	EP Mandate	VS.EC	Council Mandate	VS.EC	Draft Agreement
	ANNEX I, 7 paragraph, amending provision, numbered paragraph (1.5.2.4.3), point (b)							
245	(b) the substance or mixture does not require labelling in accordance with Part 1 or 2 of Annex II, except for section 2.8 of Part 2 of Annex II;	(b) the substance or mixture does not require labelling in accordance with Part 1 or 2 of Annex II, except for section 2.8 of Part 2 of Annex II;				(b) the substance or mixture does not require labelling in accordance with Part 1 or 2 of Annex II, except for section 2.8 of Part 2 of Annex II;		
	ANNEX I, 7 paragraph, amending provision, numbered paragraph (1.5.2.4.3), point (c)							
246	(c) the substance or mixture is not classified in any of the hazard classes and categories referred to in point (b) of section 1.5.2.4.1.;	(c) the substance or mixture is not classified in any of the hazard classes and categories referred to in point (b) of section 1.5.2.4.1.;				(c) the substance or mixture is not classified in any of the hazard classes and categories referred to in point (b) of section 1.5.2.4.1.;		
	ANNEX I, 7 paragraph a							
246a		<u>7a in Annex I, section 1.5.2.4.3a is added: 'Where section 1.5.2.4.3 applies, the label on the packaging shall contain the product identifier referred to in Article 18(2) for substances or the trade name or designation referred to in Article 18(3), point (a) for mixtures, and, where applicable, the hazard pictograms 'GHS01' or 'GHS05';'</u>						
	ANNEX I, 7 paragraph b							
246b		<u>(7 b) in Annex I, section 1.5.2.5a is added: Labelling of</u>						

CLEAN	Commission Proposal	VS.EC EP Mandate	VS.EC Council Mandate	VS.EC Draft Agreement
		<p><u>ink cartridges where the contents do not exceed 150 ml.</u></p> <p><u>1.5.2.5a1: The label elements required by Article 17 may be reduced in accordance with</u></p> <p><u>1.5.2.5a.2 and 1.5.2.5a.3 where:</u></p> <p><u>(a) the contents of the ink cartridge do not exceed 150 ml</u></p> <p><u>(b) the outer packaging complies with the provisions laid out in Article 17.; and</u></p> <p><u>(c) keep the hazard information on the outer packaging is kept with the printer</u></p> <p><u>1.5.2.5a.2: Where 1.5.2.5a.1 applies the label on the ink cartridge and any intermediate packaging shall contain at least:</u></p> <p><u>(a) the product identifier in accordance with Article 18(2) for substances and Article 18(3) for mixtures;</u></p> <p><u>(b) where applicable the pictograms;</u></p> <p><u>(c) where applicable the unique formula identifier;</u></p> <p><u>(d) name, registered name or trademark of the supplier; and</u></p> <p><u>(e) telephone number or digital contact</u></p> <p><u>1.5.2.5a.3: Where 1.5.2.5a.1 applies and the contents of the</u></p>		

CLEAN	Commission Proposal	VS,EC	EP Mandate	VS,EC	Council Mandate	VS,EC	Draft Agreement
		<p><u>ink cartridge do not exceed 30 ml the information required by 1.5.2.5a.1 may be further reduced so that the label on the ink cartridge shall contain at least:</u></p> <p><u>(a) the product identifier in accordance with Article 18(2) for substances and Article 18(3) for mixtures; and</u></p> <p><u>(b) where applicable the following pictograms: GHS01, GHS05, GHS06, GHS08, GHS07, GHS09, Where more than two pictograms are assigned GHS06 and GHS08 may take precedence over GHS01 and GHS05.</u></p>					
ANNEX I, 8 paragraph							
247	(8) in Annex I, section 1.6 is replaced by the following:	(8) in Annex I, section 1.6 is replaced by the following:				(8) (9) in Annex I, section 1.6 is replaced by the following:	
ANNEX I, 8 paragraph, amending provision, numbered paragraph (1.6)							
248	1.6. Label elements that may be provided on a digital label only	1.6. Label elements that may be provided on a digital label only	1.6. Label elements that may be provided on a digital label only	1.6. Label elements that may be provided on a digital label only			
ANNEX I, 8 paragraph, amending provision, numbered paragraph (1.6), point (a)							
249	(a) Supplemental information referred to in Article 25(3);	(a) Supplemental information referred to in Article 25(3);	(a) Supplemental information referred to in Article 25(3);	(a) Supplemental information referred to in Article 25(3);			
ANNEX I, 8 paragraph, amending provision, numbered paragraph (1.6), point (b)							
250	(b) Where more than one supplier is indicated on the label in	(b) Where more than one supplier is indicated on the label in	(b) Where more than one supplier is indicated on the label in	(b) Where more than one supplier is indicated on the label in			

	CLEAN	Commission Proposal	VS,EC	EP Mandate	VS,EC	Council Mandate	VS,EC	Draft Agreement
		accordance with Article 17(1), point (a), the name, address and digital contact of the suppliers may be provided on a digital label only, as long as the supplier referred to in Article 4(11) is indicated on the physical label.;	VS,EC	accordance with Article 17(1), point (a), the name, address and digital contact of the suppliers may be provided on a digital label only, as long as the supplier referred to in Article 4(11) is indicated on the physical label.;	VS,EC	accordance with Article 17(1), point (a), the name, address and digital contact of the suppliers may be provided on a digital label only, as long as the supplier referred to in Article 4(11) is indicated on the physical label.;		
	ANNEX I, 8 paragraph, amending provision, numbered paragraph (1.6), point (ba)							
250a				<i>(ba) The telephone number for the supplier(s) required according to Article 17(1), point (a), and where relevant the third subparagraph of Article 25(6) and section 1.5.1.2 of Annex I;</i>				
	ANNEX I, 9 paragraph							
251	(9)	in Annex II, Part 5 is replaced by the following:	(9)	in Annex II, Part 5 is replaced by the following:	☉(10)	in Annex II, Part 5 is replaced by the following:		
	ANNEX I, 9 paragraph, amending provision, first paragraph							
252		PART 5: HAZARDOUS SUBSTANCES AND MIXTURES TO WHICH ARTICLE 29(3) APPLIES		PART 5: HAZARDOUS SUBSTANCES AND MIXTURES TO WHICH ARTICLE 29(3) APPLIES		PART 5: HAZARDOUS SUBSTANCES AND MIXTURES TO WHICH ARTICLE 29(3) APPLIES		
	ANNEX I, 9 paragraph, amending provision, numbered paragraph (a)							
253	(a)	Ready mixed cement and concrete in the wet state shall be accompanied by a copy of the	(a)	Ready mixed cement and concrete in the wet state shall be accompanied by a copy of the	(a)	Ready mixed cement and concrete in the wet state shall be accompanied by a copy of the		

CLEAN	Commission Proposal	VS.EC	EP Mandate	VS.EC	Council Mandate	VS.EC	Draft Agreement
	label elements in accordance with Article 17.		label elements in accordance with Article 17.		label elements in accordance with Article 17.		
ANNEX I, 9 paragraph, amending provision, numbered paragraph (b)	(b) For a substance or a mixture supplied at a filling station and directly pumped into a receptacle that forms an integral part of a vehicle and from where the substance or mixture is normally not intended to be removed, the copy of the label elements referred to in Article 17, points (c) to (h) shall be provided on a visible place on the respective pump. The unique formula identifier referred to in Article 25(7) does not need to be provided.	(b) For a substance or a mixture supplied at a <u>filling fuel service</u> station and directly pumped into a receptacle that forms an integral part of a vehicle and from where the substance or mixture is normally not intended to be removed , the copy of the <u>following</u> label elements referred to in Article 17 <u>17.1</u> , points (c) to (h) shall be provided on a visible place on <u>or next to</u> the respective pump. The unique formula identifier referred to in Article 25(7) does not need to be provided.	(b) For a substance or a mixture supplied at a filling station and directly pumped into a receptacle that forms an integral part of a vehicle and from where the substance or mixture is normally not intended to be removed, the copy of the label elements referred to in Article 17, points (c) to (h) shall be provided on a visible place on the respective pump. The unique formula identifier referred to in Article 25(7) does not need to be provided.				
254							
ANNEX I, 9 paragraph, amending provision, numbered paragraph (c)	(c) When a vehicle fuel is supplied at a filling station through pumping into portable receptacles designed to be used for fuels, a copy of the label elements referred to in Article 17, points (c) to (h), shall be provided to be attached to the receptacle, unless the receptacle is already appropriately labelled. The unique formula	(c) When a vehicle fuel is supplied at a filling station through pumping into portable receptacles designed to be used for fuels, a copy of the label elements referred to in Article 17, points (c) to (h), shall be provided to be attached to the receptacle, unless the receptacle is already appropriately labelled. The unique formula	(c) When a vehicle fuel is supplied at a filling station through pumping into portable receptacles designed to be used for fuels, a copy of the label elements referred to in Article 17, points (c) to (h), shall be provided to be attached to the receptacle, unless the receptacle is already appropriately labelled. The unique formula				
255							

CLEAN	Commission Proposal	VS,EC	EP Mandate	VS,EC	Council Mandate	VS,EC	Draft Agreement
	identifier referred to in Article 25(7) does not need to be provided.. ,		identifier referred to in Article 25(7) does not need to be provided.. ,		identifier referred to in Article 25(7) does not need to be provided.: ,		
ANNEX II							
256	ANNEX II	ANNEX II		ANNEX II		ANNEX II	
ANNEX II, first paragraph							
257	In Annex I to Regulation (EC) No 1223/2009 Part A, point 2 is replaced by the following:	In Annex I to Regulation (EC) No 1223/2009 Part A, point 2 is replaced by the following:			In Annex I to Regulation (EC) No 1223/2009 Part A, point 2 is replaced by the following:		
ANNEX II, first paragraph, amending provision, numbered paragraph (2), first subparagraph							
258	2. Physical/chemical characteristics and stability of the cosmetic product	2. Physical/chemical characteristics and stability of the cosmetic product		2. Physical/chemical characteristics and stability of the cosmetic product	2. Physical/chemical characteristics and stability of the cosmetic product		
ANNEX II, first paragraph, amending provision, numbered paragraph (2), second subparagraph							
259	The physical and chemical characteristics of the substances or mixtures, as well as the cosmetic product.	The physical and chemical characteristics of the substances or mixtures, as well as the cosmetic product.		The physical and chemical characteristics of the substances or mixtures, as well as the cosmetic product.	The physical and chemical characteristics of the substances or mixtures, as well as the cosmetic product.		
ANNEX II, first paragraph, amending provision, numbered paragraph (2), third subparagraph							
260	The stability of the cosmetics product under reasonably foreseeable storage conditions.	The stability of the cosmetics product under reasonably foreseeable storage conditions.		The stability of the cosmetics product under reasonably foreseeable storage conditions.	The stability of the cosmetics product under reasonably foreseeable storage conditions.		
ANNEX II, first paragraph, amending provision, numbered paragraph (2), fourth subparagraph							

	CLEAN	Commission Proposal	VS,EC EP Mandate	VS,EC Council Mandate	VS,EC	Draft Agreement
261	The specification of the nanomaterial including its chemical name (IUPAC) and other descriptors as specified in point 2 of the preamble to Annexes II to VI, size of particles, physical and chemical properties.	The specification of the nanomaterial including its chemical name (IUPAC) and other descriptors as specified in point 2 of the preamble to Annexes II to VI, size of particles, physical and chemical properties.		The specification of the nanomaterial including its chemical name (IUPAC) and other descriptors as specified in point 2 of the preamble to Annexes II to VI, size of particles, physical and chemical properties.		
ANNEX II, first paragraph, amending provision, numbered paragraph (2), fifth subparagraph						
262	The safety data of the nanomaterial (including its toxicological profile and exposure conditions) relating to the category of cosmetic product, as used in such products.	The safety data of the nanomaterial (including its toxicological profile and exposure conditions) relating to the category of cosmetic product, as used in such products.		The safety data of the nanomaterial (including its toxicological profile and exposure conditions) relating to the category of cosmetic product, as used in such products.		
ANNEX III						
263	ANNEX III		ANNEX III		ANNEX III	
ANNEX III, first paragraph						
264	Annexes II to VI to Regulation (EC) No 1223/2009 are amended as follows:	Annexes II to VI to Regulation (EC) No 1223/2009 are amended as follows:		Annexes II to VI to Regulation (EC) No 1223/2009 are amended as follows:		
ANNEX III, second paragraph						
265	(1) in the preamble to Annexes II to VI, point 2, the fifth indent is replaced by the following:	(1) in the preamble to Annexes II to VI, point 2, the fifth indent is replaced by the following:		(1) in the preamble to Annexes II to VI, point 2, the fifth indent is replaced by the following:		
ANNEX III, second paragraph, amending provision, numbered paragraph						

	CLEAN	Commission Proposal	VS.EC	EP Mandate	VS.EC	Council Mandate	VS.EC	Draft Agreement
266		<p>the name included in the internationally recognised nomenclature.;</p>		<p>the name included in the internationally recognised nomenclature.;</p>		<p>the name included in the internationally recognised nomenclature.;</p>		
ANNEX III, 2 paragraph								
267		<p>in the heading of tables in Annexes III to VI the title 'Name of Common Ingredients Glossary' is replaced by 'Name in the Internationally Recognised Nomenclature'.</p>		<p>in the heading of tables in Annexes III to VI the title 'Name of Common Ingredients Glossary' is replaced by 'Name in the Internationally Recognised Nomenclature'.</p>		<p>in the heading of tables in Annexes III to VI the title 'Name of Common Ingredients Glossary' is replaced by 'Name in the Internationally Recognised Nomenclature'.</p>		
ANNEX IV								
268		ANNEX IV		ANNEX IV		ANNEX IV		
ANNEX IV, first paragraph								
269		Annexes I, II and IV to Regulation (EU) 2019/1009 are amended as follows:		Annexes I, II and IV to Regulation (EU) 2019/1009 are amended as follows:		Annexes I, II and IV to Regulation (EU) 2019/1009 are amended as follows:		
ANNEX IV, second paragraph								
270		<p>in Annex I, Part II, PFC 7: FERTILISING PRODUCT BLEND, point 4(c) is replaced by the following:</p>		<p>in Annex I, Part II, PFC 7: FERTILISING PRODUCT BLEND, point 4(c) is replaced by the following:</p>		<p>in Annex I, Part II, PFC 7: FERTILISING PRODUCT BLEND, point 4(c) is replaced by the following:</p>		
ANNEX IV, second paragraph, amending provision, numbered paragraph (c)								
271		<p>Article 8(8) (importers' obligation to keep the EU</p>		<p>Article 8(8) (importers' obligation to keep the EU</p>		<p>Article 8(8) (importers' obligation to keep the EU</p>		

CLEAN	Commission Proposal	VS,EC	EP Mandate	VS,EC	Council Mandate	VS,EC	Draft Agreement
	declaration of conformity at the disposal of the market surveillance authorities); ,		declaration of conformity at the disposal of the market surveillance authorities); ,		declaration of conformity at the disposal of the market surveillance authorities); ,		
ANNEX IV, 2 paragraph							
272	(2) in Annex II, Part II, is amended as follows:	(2) in Annex II, Part II, is amended as follows:		(2) in Annex II, Part II, is amended as follows:			
ANNEX IV, 2 paragraph, point (a)							
273	(a) in CMC 1: VIRGIN MATERIAL SUBSTANCES AND MIXTURES, point 2 is deleted;	(a) in CMC 1: VIRGIN MATERIAL SUBSTANCES AND MIXTURES, point 2 is deleted <u>a) Substances classified in Part 3 of Annex VI to Regulation (EC) No 1272/2008 for the following hazard classes or categories:</u> <u>- Germ cell mutagenicity, category IA or IB;</u> <u>- Carcinogenicity, category IA or IB;</u> <u>- Reproductive toxicity, category IA or IB;</u> <u>- Specific target organ toxicity, repeated exposure, category I</u> <u>- Endocrine disruptor for human health, category I;</u> <u>- Endocrine disruptor for the environment, category I; and</u>		(a) in CMC 1: VIRGIN MATERIAL SUBSTANCES AND MIXTURES, point 2 is deleted ; replaced as follows:			

CLEAN	Commission Proposal	VS,EC EP Mandate	VS,EC Council Mandate	VS,EC Draft Agreement
		<p><u>- Persistent, bioaccumulative and toxic or very persistent and very bioaccumulative properties whose actual quantities placed on the market are lower than 10 tonnes per year, intentionally incorporated into the EU fertilising product, on their own or in a mixture, in a concentration equal or lower than the generic cut-off values set out in Article 11(3) of Regulation (EC) No 1272/2008, shall have been registered pursuant to Regulation (EC) No 1907/2006, with a dossier containing:</u></p> <p><u>(i) the information provided for by Annexes VI, VII and, insofar as relevant and available, Annex VIII to Regulation (EC) No 1907/2006, on the basis of available data, alternative methods pursuant to Article 13 and adaptations pursuant to Annex XI, and conducting new tests on vertebrate animals only as a last resort and where relevant, and</u></p> <p><u>(ii) a chemical safety report pursuant to Article 14 of Regulation (EC) No 1907/2006 strictly limited to the exposure</u></p>		

	CLEAN	Commission Proposal	VS,EC	EP Mandate	VS,EC	Council Mandate	VS,EC	Draft Agreement
				<p><u>scenarios related to the agronomic use and the environment, unless explicitly covered by one of the registration obligation exemptions provided for by Article 9 (PPORD) of Regulation (EC) No 1907/2006, used exclusively for scientific research and development, Annex IV to Regulation (EC) No 1907/2006 or by points 6, 7, 8, 9 or 10 9 (only for magnesia) of Annex V to that Regulation.</u></p> <p><u>b) Polymers are exempt from point a).;</u></p>				
		ANNEX IV, 2 paragraph, point (b)						
274	(b) in CMC 3: COMPOST, point 1(d) is replaced by the following:	(b) in CMC 3: COMPOST, point 1(d) is replaced by the following:		(b) in CMC 3: COMPOST, point 1(d) is replaced by the following:		(b) in CMC 3: COMPOST, point 1(d) is replaced by the following:		
	ANNEX IV, 2 paragraph, point (b), amending provision, numbered paragraph (d)							
275	(d) composting additives which are necessary to improve the process performance or the environmental performance of the composting process, provided that the total concentration of all additives does not exceed 5% of the total input material weight; or;	(d) composting additives which are necessary to improve the process performance or the environmental performance of the composting process, provided that the total concentration of all additives does not exceed 5% of the total input material weight; or;		(d) composting additives which are necessary to improve the process performance or the environmental performance of the composting process, provided that the total concentration of all additives does not exceed 5% of the total input material weight; or;		(d) composting additives which are necessary to improve the process performance or the environmental performance of the composting process, provided that the total concentration of all additives does not exceed 5% of the total input material weight; or;		

	CLEAN	Commission Proposal	VS.EC	EP Mandate	VS.EC	Council Mandate	VS.EC	Draft Agreement
		,		,		,		
ANNEX IV, 2 paragraph, point (c)								
276	(c) CMC 4: FRESH CROP DIGESTATE is amended as follows:	(c) CMC 4: FRESH CROP DIGESTATE is amended as follows:			(c) CMC 4: FRESH CROP DIGESTATE is amended as follows:			
ANNEX IV, 2 paragraph, point (c)(i)								
277	(i) point 1(b) is replaced by the following:	(i) point 1(b) is replaced by the following:			(i) point 1(b) is replaced by the following:			
ANNEX IV, 2 paragraph, point (c)(i), amending provision, numbered paragraph (b)								
278	(b) digestion additives which are needed to improve the process performance or the environmental performance of the digestion process, provided that the total concentration of all additives does not exceed 5% of the total input material weight; or;	(b) digestion additives which are needed to improve the process performance or the environmental performance of the digestion process, provided that the total concentration of all additives does not exceed 5% of the total input material weight; or;			(b) digestion additives which are needed to improve the process performance or the environmental performance of the digestion process, provided that the total concentration of all additives does not exceed 5% of the total input material weight; or;			
ANNEX IV, 2 paragraph, point (c)(ii)								
279	(ii) point 3d is replaced by the following:	(ii) point 3d is replaced by the following:			(ii) point 3d is replaced by the following:			
ANNEX IV, 2 paragraph, point (c)(ii), amending provision, numbered paragraph (3d)								
280	3d. Additives needed in the post processing of a digestate or a	3d. Additives needed in the post processing of a digestate or a			3d. Additives needed in the post processing of a digestate or a			

	CLEAN	Commission Proposal	VS.EC	EP Mandate	VS.EC	Council Mandate	VS.EC	Draft Agreement
		fraction in accordance with points 3a, 3b and 3c may be used provided that the concentration of the additives needed in each of the processes does not exceed 5 % of the weight of the digestate or fraction used as input in the respective process.;	VS.EC	fraction in accordance with points 3a, 3b and 3c may be used provided that the concentration of the additives needed in each of the processes does not exceed 5 % of the weight of the digestate or fraction used as input in the respective process.;	VS.EC	fraction in accordance with points 3a, 3b and 3c may be used provided that the concentration of the additives needed in each of the processes does not exceed 5 % of the weight of the digestate or fraction used as input in the respective process.;		
ANNEX IV, 2 paragraph, point (d)								
281	(d) CMC 5: DIGESTATE OTHER THAN FRESH CROP DIGESTATE is amended as follows	(d) CMC 5: DIGESTATE OTHER THAN FRESH CROP DIGESTATE is amended as follows	VS.EC	(d) CMC 5: DIGESTATE OTHER THAN FRESH CROP DIGESTATE is amended as follows	VS.EC	(d) CMC 5: DIGESTATE OTHER THAN FRESH CROP DIGESTATE is amended as follows:		
ANNEX IV, 2 paragraph, point (d)(i)								
282	(i) point 1(d) is replaced by the following:	(i) point 1(d) is replaced by the following:	VS.EC	(i) point 1(d) is replaced by the following:	VS.EC	(i) point 1(d) is replaced by the following:		
ANNEX IV, 2 paragraph, point (d)(i), amending provision, numbered paragraph (d)								
283	(d) digestion additives which are necessary to improve the process performance or the environmental performance of the digestion process, provided that the total concentration of all additives does not exceed 5% of the total input material weight; or;	(d) digestion additives which are necessary to improve the process performance or the environmental performance of the digestion process, provided that the total concentration of all additives does not exceed 5% of the total input material weight; or;	VS.EC	(d) digestion additives which are necessary to improve the process performance or the environmental performance of the digestion process, provided that the total concentration of all additives does not exceed 5% of the total input material weight; or;	VS.EC	(d) digestion additives which are necessary to improve the process performance or the environmental performance of the digestion process, provided that the total concentration of all additives does not exceed 5% of the total input material weight; or;		

	CLEAN	Commission Proposal	VS.EC	EP Mandate	VS.EC	Council Mandate	VS.EC	Draft Agreement
		ANNEX IV, 2 paragraph, point (d)(ii)						
284	(ii)	point 3d is replaced by the following:	(ii)	point 3d is replaced by the following:	(ii)	point 3d is replaced by the following:		
		ANNEX IV, 2 paragraph, point (d)(ii), amending provision, numbered paragraph (3d)						
285	3d.	Additives needed in the post processing of a digestate or a fraction in accordance with points 3a, 3b and 3c may be used, provided that the concentration of the additives needed in each of the processes does not exceed 5% of the weight of the digestate or fraction used as input in the respective process.;	3d.	Additives needed in the post processing of a digestate or a fraction in accordance with points 3a, 3b and 3c may be used, provided that the concentration of the additives needed in each of the processes does not exceed 5% of the weight of the digestate or fraction used as input in the respective process.;	3d.	Additives needed in the post processing of a digestate or a fraction in accordance with points 3a, 3b and 3c may be used, provided that the concentration of the additives needed in each of the processes does not exceed 5% of the weight of the digestate or fraction used as input in the respective process.;		
		ANNEX IV, 2 paragraph, point (e)						
286	(e)	in CMC 6: FOOD INDUSTRY BY-PRODUCTS, point 2 is deleted;	(e)	in CMC 6: FOOD INDUSTRY BY-PRODUCTS, point 2 is deleted;	(e)	in CMC 6: FOOD INDUSTRY BY-PRODUCTS, point 2 is deleted;		
		ANNEX IV, 2 paragraph, point (f)						
287	(f)	in CMC 8: NUTRIENT POLYMERS, point 1 is replaced by the following:	(f)	in CMC 8: NUTRIENT POLYMERS, point 1 is replaced by the following:	(f)	in CMC 8: NUTRIENT POLYMERS, point 1 is replaced by the following:		
		ANNEX IV, 2 paragraph, point (f), amending provision, numbered paragraph (1)						
288	1.	An EU fertilising product may contain polymers exclusively	1.	An EU fertilising product may contain polymers exclusively	1.	An EU fertilising product may contain polymers exclusively		

CLEAN	Commission Proposal	VS.EC	EP Mandate	VS.EC	Council Mandate	VS.EC	Draft Agreement
	made up of monomer substances complying with the criteria set out in point 1 of CMC 1, where the purpose of the polymerisation is to control the release of nutrients from one or more of the monomer substances.;	made up of monomer substances complying with the criteria set out in point 1 of CMC 1, where the purpose of the polymerisation is to control the release of nutrients from one or more of the monomer substances. <u>The polymer may additionally perform other functions, provided that this does not alter the safety characteristics of the product nor compromise the controlled-release function;</u>	made up of monomer substances complying with the criteria set out in point 1 of CMC 1, where the purpose of the polymerisation is to control the release of nutrients from one or more of the monomer substances.;				
ANNEX IV, 2 paragraph, point (g)	(g) in CMC 10: DERIVED PRODUCTS WITHIN THE MEANING OF REGULATION (EC) No 1069/2009, the table, point 1.3 is replaced by the following:	(g) in CMC 10: DERIVED PRODUCTS WITHIN THE MEANING OF REGULATION (EC) No 1069/2009, the table, point 1.3 is replaced by the following:	(g) in CMC 10: DERIVED PRODUCTS WITHIN THE MEANING OF REGULATION (EC) No 1069/2009, the table, point 1.3 is replaced by the following:				
289	ANNEX IV, 2 paragraph, point (g), amending provision, numbered paragraph (1.3)	1.3. Additives needed in the processing referred to in points 1.1 and 1.2 may be used, provided that the concentration of the additives needed in each of the processes does not exceed 5% of the weight of the processed manure or	1.3. Additives needed in the processing referred to in points 1.1 and 1.2 may be used, provided that the concentration of the additives needed in each of the processes does not exceed 5% of the weight of the processed manure or				
290							

	CLEAN	Commission Proposal	vs.EC	EP Mandate	vs.EC	Council Mandate	vs.EC	Draft Agreement
		fraction used as input in the respective process.;		fraction used as input in the respective process.;		fraction used as input in the respective process.;		
	ANNEX IV, 2 paragraph, point (h)							
291	(h) in CMC 11: BY-PRODUCTS WITHIN THE MEANING OF DIRECTIVE 2008/98/EC, point 2 is deleted;	(h) in CMC 11: BY-PRODUCTS WITHIN THE MEANING OF DIRECTIVE 2008/98/EC, point 2 is deleted;		(h) in CMC 11: BY-PRODUCTS WITHIN THE MEANING OF DIRECTIVE 2008/98/EC, point 2 is deleted;		(h) in CMC 11: BY-PRODUCTS WITHIN THE MEANING OF DIRECTIVE 2008/98/EC, point 2 is deleted;		
	ANNEX IV, 2 paragraph, point (i)							
292	(i) in CMC 12: PRECIPITATED PHOSPHATE SALTS AND DERIVATES, point 13 is deleted;	(i) in CMC 12: PRECIPITATED PHOSPHATE SALTS AND DERIVATES, point 13 is deleted;		(i) in CMC 12: PRECIPITATED PHOSPHATE SALTS AND DERIVATES, point 13 is deleted;		(i) in CMC 12: PRECIPITATED PHOSPHATE SALTS AND DERIVATES, point 13 is deleted;		
	ANNEX IV, 2 paragraph, point (j)							
293	(j) in CMC 13: THERMAL OXIDATION MATERIALS OR DERIVATES, point 8 is deleted;	(j) in CMC 13: THERMAL OXIDATION MATERIALS OR DERIVATES, point 8 is deleted;		(j) in CMC 13: THERMAL OXIDATION MATERIALS OR DERIVATES, point 8 is deleted;		(j) in CMC 13: THERMAL OXIDATION MATERIALS OR DERIVATES, point 8 is deleted;		
	ANNEX IV, 2 paragraph, point (k)							
294	(k) in CMC 14: PYROLYSIS AND GASIFICATION MATERIALS, point 7 is deleted;	(k) in CMC 14: PYROLYSIS AND GASIFICATION MATERIALS, point 7 is deleted;		(k) in CMC 14: PYROLYSIS AND GASIFICATION MATERIALS, point 7 is deleted;		(k) in CMC 14: PYROLYSIS AND GASIFICATION MATERIALS, point 7 is deleted;		
	ANNEX IV, 2 paragraph, point (l)							
295	(l) in CMC 15: RECOVERED HIGH PURITY MATERIALS, point 10 is deleted;	(l) in CMC 15: RECOVERED HIGH PURITY MATERIALS, point 10 is deleted;		(l) in CMC 15: RECOVERED HIGH PURITY MATERIALS, point 10 is deleted;		(l) in CMC 15: RECOVERED HIGH PURITY MATERIALS, point 10 is deleted;		
	ANNEX IV, 3 paragraph							

	CLEAN	Commission Proposal	VS.EC	EP Mandate	VS.EC	Council Mandate	VS.EC	Draft Agreement
296	(3) in Annex IV, Part II is amended as follows:	(3) in Annex IV, Part II is amended as follows:		(3) in Annex IV, Part II is amended as follows:		(3) in Annex IV, Part II is amended as follows:		
ANNEX IV, 3 paragraph, point (a)								
297	(a) MODULE A – INTERNAL PRODUCTION CONTROL is amended as follows:	(a) MODULE A – INTERNAL PRODUCTION CONTROL is amended as follows:		(a) MODULE A – INTERNAL PRODUCTION CONTROL is amended as follows:		(a) MODULE A – INTERNAL PRODUCTION CONTROL is amended as follows:		
ANNEX IV, 3 paragraph, point (a)(i)								
298	(i) in point 4.2, the first sentence is replaced by the following:	(i) in point 4.2, the first sentence is replaced by the following:		(i) in point 4.2, the first sentence is replaced by the following:		(i) in point 4.2, the first sentence is replaced by the following:		
ANNEX IV, 3 paragraph, point (a)(i), amending provision, first paragraph								
299	The manufacturer shall draw up an EU declaration of conformity for an EU fertilising product or type in electronic form and keep it together with the technical documentation at the disposal of the national authorities for 5 years after the EU fertilising product has been placed on the market.;	The manufacturer shall draw up an EU declaration of conformity for an EU fertilising product or type in electronic form and keep it together with the technical documentation at the disposal of the national authorities for 5 years after the EU fertilising product has been placed on the market.;		The manufacturer shall draw up an EU declaration of conformity for an EU fertilising product or type in electronic form and keep it together with the technical documentation at the disposal of the national authorities for 5 years after the EU fertilising product has been placed on the market.;		The manufacturer shall draw up an EU declaration of conformity for an EU fertilising product or type in electronic form and keep it together with the technical documentation at the disposal of the national authorities for 5 years after the EU fertilising product has been placed on the market.;		
ANNEX IV, 3 paragraph, point (a)(ii)								
300	(ii) point 4.3. is replaced by the following:	(ii) point 4.3. is replaced by the following:		(ii) point 4.3. is replaced by the following:		(ii) point 4.3. is replaced by the following:		
ANNEX IV, 3 paragraph, point (a)(ii), amending provision, numbered paragraph (4.3)								

	CLEAN	Commission Proposal	VS,EC EP Mandate	VS,EC Council Mandate	VS,EC	Draft Agreement
301		4.3. The manufacturer shall, further to a reasoned request from a competent national authority, provide it, in electronic form, with all the information and documentation necessary to demonstrate the conformity of the EU fertilising product with this Regulation, in a language which can be easily understood by that authority.;	4.3. The manufacturer shall, further to a reasoned request from a competent national authority, provide it, in electronic form, with all the information and documentation necessary to demonstrate the conformity of the EU fertilising product with this Regulation, in a language which can be easily understood by that authority.;	4.3. The manufacturer shall, further to a reasoned request from a competent national authority, provide it, in electronic form, with all the information and documentation necessary to demonstrate the conformity of the EU fertilising product with this Regulation, in a language which can be easily understood by that authority.;		
	ANNEX IV, 3 paragraph, point (b)					
302	(b) MODULE A1 - INTERNAL PRODUCTION CONTROL PLUS SUPERVISED PRODUCT TESTING is amended as follows:	(b) MODULE A1 - INTERNAL PRODUCTION CONTROL PLUS SUPERVISED PRODUCT TESTING is amended as follows:	(b) MODULE A1 - INTERNAL PRODUCTION CONTROL PLUS SUPERVISED PRODUCT TESTING is amended as follows:	(b) MODULE A1 - INTERNAL PRODUCTION CONTROL PLUS SUPERVISED PRODUCT TESTING is amended as follows:		
	ANNEX IV, 3 paragraph, point (b)(i)					
303	(i) point 2.2.(f) is replaced by the following:	(i) point 2.2.(f) is replaced by the following:	(i) point 2.2.(f) is replaced by the following:	(i) point 2.2.(f) is replaced by the following:		
	ANNEX IV, 3 paragraph, point (b)(i), amending provision, numbered paragraph (f)					
304	(f) the names, postal addresses and digital contacts of the sites, and of the operators of the sites, at which the product and	(f) the names, postal addresses and digital contacts of the sites, and of the operators of the sites, at which the product and	(f) the names, postal addresses and digital contacts of the sites, and of the operators of the sites, at which the product and	(f) the names, postal addresses and digital contacts of the sites, and of the operators of the sites, at which the product and		

	CLEAN	Commission Proposal	VS,EC	EP Mandate	VS,EC	Council Mandate	VS,EC	Draft Agreement
		its principal components were manufactured,;		its principal components were manufactured,;		its principal components were manufactured,;		
	ANNEX IV, 3 paragraph, point (b)(ii)							
305	(ii) in point 5.2., the first sentence is replaced by the following:	(ii) in point 5.2., the first sentence is replaced by the following:		(ii) in point 5.2., the first sentence is replaced by the following:		(ii) in point 5.2., the first sentence is replaced by the following:		
	ANNEX IV, 3 paragraph, point (b)(ii), amending provision, first paragraph							
306	The manufacturer shall draw up an EU declaration of conformity for an EU fertilising product type in electronic form and keep it together with the technical documentation at the disposal of the national authorities for 5 years after the EU fertilising product has been placed on the market.;	The manufacturer shall draw up an EU declaration of conformity for an EU fertilising product type in electronic form and keep it together with the technical documentation at the disposal of the national authorities for 5 years after the EU fertilising product has been placed on the market.;		The manufacturer shall draw up an EU declaration of conformity for an EU fertilising product type in electronic form and keep it together with the technical documentation at the disposal of the national authorities for 5 years after the EU fertilising product has been placed on the market.;		The manufacturer shall draw up an EU declaration of conformity for an EU fertilising product type in electronic form and keep it together with the technical documentation at the disposal of the national authorities for 5 years after the EU fertilising product has been placed on the market.;		
	ANNEX IV, 3 paragraph, point (b)(iii)							
307	(iii) point 5.3. is replaced by the following:	(iii) point 5.3. is replaced by the following:		(iii) point 5.3. is replaced by the following:		(iii) point 5.3. is replaced by the following:		
	ANNEX IV, 3 paragraph, point (b)(iii), amending provision, numbered paragraph (5.3)							
308	5.3. The manufacturer shall, further to a reasoned request from a competent national authority,	5.3. The manufacturer shall, further to a reasoned request from a competent national authority,		5.3. The manufacturer shall, further to a reasoned request from a competent national authority,		5.3. The manufacturer shall, further to a reasoned request from a competent national authority,		

CLEAN	Commission Proposal	VS.EC	EP Mandate	VS.EC	Council Mandate	VS.EC	Draft Agreement
	provide it, in electronic form, with all the information and documentation necessary to demonstrate the conformity of the EU fertilising product with this Regulation, in a language which can be easily understood by that authority.;	VS.EC	provide it, in electronic form, with all the information and documentation necessary to demonstrate the conformity of the EU fertilising product with this Regulation, in a language which can be easily understood by that authority.;	VS.EC	provide it, in electronic form, with all the information and documentation necessary to demonstrate the conformity of the EU fertilising product with this Regulation, in a language which can be easily understood by that authority.;	VS.EC	
ANNEX IV, 3 paragraph, point (c)							
309	(c) MODULE B – EU-TYPE EXAMINATION is amended as follows:	(c)	MODULE B – EU-TYPE EXAMINATION is amended as follows:	(c)	MODULE B – EU-TYPE EXAMINATION is amended as follows:		
ANNEX IV, 3 paragraph, point (c)(i)							
310	(i) point 3.2.(a) is replaced by the following:	(i)	point 3.2.(a) is replaced by the following:	(i)	point 3.2.(a) is replaced by the following:		
ANNEX IV, 3 paragraph, point (c)(i), amending provision, numbered paragraph (a)							
311	(a) the name, postal address and digital contact of the manufacturer and, if the application is lodged by the authorised representative, his or her name, postal address and digital contact as well.;	(a)	the name, postal address and digital contact of the manufacturer and, if the application is lodged by the authorised representative, his or her name, postal address and digital contact as well.;	(a)	the name, postal address and digital contact of the manufacturer and, if the application is lodged by the authorised representative, his or her name, postal address and digital contact as well.;		
ANNEX IV, 3 paragraph, point (c)(ii)							

	CLEAN	Commission Proposal	VS.EC	EP Mandate	VS.EC	Council Mandate	VS.EC	Draft Agreement
312	(ii) in point 6.1., the second sentence is replaced by the following:	(ii) in point 6.1., the second sentence is replaced by the following:		(ii) in point 6.1., the second sentence is replaced by the following:		(ii) in point 6.1., the second sentence is replaced by the following:		
ANNEX IV, 3 paragraph, point (c)(ii), amending provision, first paragraph								
313	The certificate shall contain the name, postal address and digital contact of the manufacturer, the conclusions of the examination, the conditions (if any) for its validity and the necessary data for identification of the approved type.;	The certificate shall contain the name, postal address and digital contact of the manufacturer, the conclusions of the examination, the conditions (if any) for its validity and the necessary data for identification of the approved type.;		The certificate shall contain the name, postal address and digital contact of the manufacturer, the conclusions of the examination, the conditions (if any) for its validity and the necessary data for identification of the approved type.;		The certificate shall contain the name, postal address and digital contact of the manufacturer, the conclusions of the examination, the conditions (if any) for its validity and the necessary data for identification of the approved type.;		
ANNEX IV, 3 paragraph, point (d)								
314	(d) MODULE C – CONFORMITY TO TYPE BASED ON INTERNAL PRODUCTION CONTROL is amended as follows:	(d) MODULE C – CONFORMITY TO TYPE BASED ON INTERNAL PRODUCTION CONTROL is amended as follows:		(d) MODULE C – CONFORMITY TO TYPE BASED ON INTERNAL PRODUCTION CONTROL is amended as follows:		(d) MODULE C – CONFORMITY TO TYPE BASED ON INTERNAL PRODUCTION CONTROL is amended as follows:		
ANNEX IV, 3 paragraph, point (d)(i)								
315	(i) in point 3.2., the first sentence is replaced by the following:	(i) in point 3.2., the first sentence is replaced by the following:		(i) in point 3.2., the first sentence is replaced by the following:		(i) in point 3.2., the first sentence is replaced by the following:		
ANNEX IV, 3 paragraph, point (d)(i), amending provision, first paragraph								
316	The manufacturer shall draw up an EU declaration of conformity for	The manufacturer shall draw up an EU declaration of conformity for		The manufacturer shall draw up an EU declaration of conformity for		The manufacturer shall draw up an EU declaration of conformity for		

	CLEAN	Commission Proposal	VS.EC EP Mandate	VS.EC Council Mandate	VS.EC Draft Agreement
		an EU fertilising product type in electronic form and keep it together with the technical documentation at the disposal of the national authorities for 5 years after the EU fertilising product has been placed on the market.;	an EU fertilising product type in electronic form and keep it together with the technical documentation at the disposal of the national authorities for 5 years after the EU fertilising product has been placed on the market.;	an EU fertilising product type in electronic form and keep it together with the technical documentation at the disposal of the national authorities for 5 years after the EU fertilising product has been placed on the market.;	
ANNEX IV, 3 paragraph, point (d)(ii)					
317	(ii) point 3.3. is replaced by the following:	(ii) point 3.3. is replaced by the following:	(ii) point 3.3. is replaced by the following:	(ii) point 3.3. is replaced by the following:	
ANNEX IV, 3 paragraph, point (d)(ii), amending provision, numbered paragraph (3.3)					
318	3.3. The manufacturer shall, further to a reasoned request from a competent national authority, provide it, in electronic form, with all the information and documentation necessary to demonstrate the conformity of the EU fertilising product with this Regulation, in a language which can be easily understood by that authority.;	3.3. The manufacturer shall, further to a reasoned request from a competent national authority, provide it, in electronic form, with all the information and documentation necessary to demonstrate the conformity of the EU fertilising product with this Regulation, in a language which can be easily understood by that authority.;	3.3. The manufacturer shall, further to a reasoned request from a competent national authority, provide it, in electronic form, with all the information and documentation necessary to demonstrate the conformity of the EU fertilising product with this Regulation, in a language which can be easily understood by that authority.;	3.3. The manufacturer shall, further to a reasoned request from a competent national authority, provide it, in electronic form, with all the information and documentation necessary to demonstrate the conformity of the EU fertilising product with this Regulation, in a language which can be easily understood by that authority.;	
ANNEX IV, 3 paragraph, point (e)					
319	(e) MODULE D1 - QUALITY ASSURANCE OF	(e) MODULE D1 - QUALITY ASSURANCE OF	(e) MODULE D1 - QUALITY ASSURANCE OF	(e) MODULE D1 - QUALITY ASSURANCE OF	

	CLEAN	Commission Proposal	VS,EC	EP Mandate	VS,EC	Council Mandate	VS,EC	Draft Agreement
		THE PRODUCTION PROCESS is amended as follows:		THE PRODUCTION PROCESS is amended as follows:		THE PRODUCTION PROCESS is amended as follows:		
	ANNEX IV, 3 paragraph, point (e)(i)							
320	(i) in point 5.2., the first indent is replaced by the following:	(i) in point 5.2., the first indent is replaced by the following:		(i) in point 5.2., the first indent is replaced by the following:		(i) in point 5.2., the first indent is replaced by the following:		
	ANNEX IV, 3 paragraph, point (e)(i), amending provision, first paragraph							
321	the name, postal address and digital contact of the manufacturer and, if the application is lodged by the authorised representative, his or her name, postal address and digital contact as well,;	the name, postal address and digital contact of the manufacturer and, if the application is lodged by the authorised representative, his or her name, postal address and digital contact as well,;		the name, postal address and digital contact of the manufacturer and, if the application is lodged by the authorised representative, his or her name, postal address and digital contact as well,;		the name, postal address and digital contact of the manufacturer and, if the application is lodged by the authorised representative, his or her name, postal address and digital contact as well,;		
	ANNEX IV, 3 paragraph, point (e)(ii)							
322	(ii) in point 7.2., the first sentence is replaced by the following:	(ii) in point 7.2., the first sentence is replaced by the following:		(ii) in point 7.2., the first sentence is replaced by the following:		(ii) in point 7.2., the first sentence is replaced by the following:		
	ANNEX IV, 3 paragraph, point (e)(ii), amending provision, first paragraph							
323	The manufacturer shall draw up an EU declaration of conformity for an EU fertilising product or type in electronic form and keep it together with the technical documentation at the disposal of the national authorities for 5 years	The manufacturer shall draw up an EU declaration of conformity for an EU fertilising product or type in electronic form and keep it together with the technical documentation at the disposal of the national authorities for 5 years		The manufacturer shall draw up an EU declaration of conformity for an EU fertilising product or type in electronic form and keep it together with the technical documentation at the disposal of the national authorities for 5 years		The manufacturer shall draw up an EU declaration of conformity for an EU fertilising product or type in electronic form and keep it together with the technical documentation at the disposal of the national authorities for 5 years		

	CLEAN	Commission Proposal	VS.EC	EP Mandate	VS.EC	Council Mandate	VS.EC	Draft Agreement
		after the EU fertilising product has been placed on the market.;	VS.EC	after the EU fertilising product has been placed on the market.;	VS.EC	after the EU fertilising product has been placed on the market.;	VS.EC	
	ANNEX IV, 3 paragraph, point (e)(iii)							
324		(iii) point 7.3. is replaced by the following:	(iii)	point 7.3. is replaced by the following:	(iii)	point 7.3. is replaced by the following:		
	ANNEX IV, 3 paragraph, point (e)(iii), amending provision, first paragraph							
325		The manufacturer shall, further to a reasoned request from a competent national authority, provide it, in electronic form, with all the information and documentation necessary to demonstrate the conformity of the EU fertilising product with this Regulation, in a language which can be easily understood by that authority.		The manufacturer shall, further to a reasoned request from a competent national authority, provide it, in electronic form, with all the information and documentation necessary to demonstrate the conformity of the EU fertilising product with this Regulation, in a language which can be easily understood by that authority.		The manufacturer shall, further to a reasoned request from a competent national authority, provide it, in electronic form, with all the information and documentation necessary to demonstrate the conformity of the EU fertilising product with this Regulation, in a language which can be easily understood by that authority.		

Commission Proposal Table

Capacity of the package	Dimensions of the label (in millimetres) for the information required by Article 17	Dimensions of each pictogram (in millimetres)
Not exceeding 3 litres:	If possible, at least 52×74	Not smaller than 10×10 If possible, at least 16×16
Greater than 3 litres but not exceeding 50 litres:	At least 74×105	At least 23×23
Greater than 50 litres but not exceeding 500 litres:	At least 105×148	At least 32×32
Greater than 500 litres:	At least 148×210	At least 46×46

EP Mandate Table

Capacity of the package	Dimensions of the label (in millimetres) for the information required by Article 17	Dimensions of each pictogram (in millimetres)
Not exceeding 3 litres:	If possible, at least 52×74	Not smaller than 10×10 If possible, at least 16×16
Greater than 3 litres but not exceeding 50 litres:	At least 74×105	At least 23×23
Greater than 50 litres but not exceeding 500 litres:	At least 105×148	At least 32×32
Greater than 500 litres:	At least 148×210	At least 46×46

Council Mandate Table

Capacity of the package	Dimensions of the label (in millimetres) for the information required by Article 17	Dimensions of each pictogram (in millimetres)
Not exceeding 3 litres:	If possible, at least 52×74	Not smaller than 10×10 If possible, at least 16×16
Greater than 3 litres but not exceeding 50 litres:	At least 74×105	At least 23×23
Greater than 50 litres but not exceeding 500 litres:	At least 105×148	At least 32×32
Greater than 500 litres:	At least 148×210	At least 46×46