Changes to legislation: Regulation (EC) No 1223/2009 of the European Parliament and of the Council is up to date with all changes known to be in force on or before 19 March 2024. There are changes that may be brought into force at a future date. Changes that have been made appear in the content and are referenced with annotations. (See end of Document for details) View outstanding changes

Regulation (EC) No 1223/2009 of the European Parliament and of the Council of 30 November 2009 on cosmetic products (recast) (Text with EEA relevance)

CHAPTER I

SCOPE, DEFINITIONS

Article 1 Article 2	Scope and objective Definitions	
		CHAPTER II

SAFETY, RESPONSIBILITY, FREE MOVEMENT

Article 3	Safety
Article 4	Responsible person
Article 5	Obligations of responsible persons
Article 5A	Obligations of responsible persons established in Northern
	Ireland
Article 6	Obligations of distributors
Article 7	Identification within the supply chain
Article 8	Good manufacturing practice
Article 9	Free movement

CHAPTER III

SAFETY ASSESSMENT, PRODUCT INFORMATION FILE, NOTIFICATION

Article 10	Safety assessment
Article 11	Product information file
Article 12	Sampling and analysis
Article 13	Notification

CHAPTER IV

RESTRICTIONS FOR CERTAIN SUBSTANCES

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CHAPTER V

ANIMAL TESTING

Article 18 Animal testing

Article 37 Article 38

Article 39

Article 40

Repeal

Signature

Transitional provisions

Entry into force and date of application

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CHAPTER VI

	CONSUMER INFORMATION	
Article 19 Article 20 Article 21	Labelling Product claims Access to information for the public	
CHAPTER VII		
MARKET SURVEILLANCE		
Article 22 Article 23 Article 24	In-market control Communication of serious undesirable effects Information on substances	
CHAPTER VIII		
	NON-COMPLIANCE, SAFEGUARD CLAUSE	
Article 25 Article 26 Article 27 Article 28	Non-compliance by the responsible person Non-compliance by distributors Safeguard clause Good administrative practices	
	CHAPTER IX	
	ADMINISTRATIVE COOPERATION	
Article 29	Cooperation between competent authorities	
	CHAPTER 10	
POWERS AND FURTHER DUTIES OF THE SECRETARY OF STATE		
Article 30 Article 31 Article 32 Article 33 Article 34	Power to amend Articles Power to amend the annexes Procedure for making regulations Further duties of the Secretary of State Competent authorities, poison control centres or assimilated entities Annual report on animal testing	
Article 36 Article 37	Formal objection against harmonised standards Penalties	

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ANNEX I COSMETIC PRODUCT SAFETY REPORT

PART A -

Cosmetic product safety information

- 1. Quantitative and qualitative composition of the cosmetic product
- 2. Physical/chěmical characteristics and stability of the cosmetic product
- 3. Microbiological quality
- 4. Impurities, traces, information about the packaging material
- 5. Normal and reasonably foreseeable use
- 6. Exposure to the cosmetic product
- 7. Exposure to the substances
- 8. Toxicological profile of the substances
- 9. Undesirable effects and serious undesirable effects
- 10. Information on the cosmetic product

PART B -

Cosmetic product safety assessment

- 1. Assessment conclusion
- 2. Labelled warnings and instructions of use
- 3. Reasoning
- Assessor's credentials and approval of part B 4.

Preamble to Annexes II to VI

For the purposes of the Annexes II to VI: 'Rinse-off...

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ANNEX II

LIST OF SUBSTANCES PROHIBITED IN COSMETIC PRODUCTS

ANNEX III

LIST OF SUBSTANCES WHICH COSMETIC PRODUCTS MUST NOT CONTAIN EXCEPT SUBJECT TO THE RESTRICTIONS LAID DOWN

ANNEX IV

LIST OF COLORANTS ALLOWED IN COSMETIC PRODUCTS

Preamble

ANNEX V

LIST OF PRESERVATIVES ALLOWED IN COSMETIC PRODUCTS

Preamble

- 1. For the purposes of this list:
- 2. All finished products containing substances in this Annex and which...

ANNEX VI

LIST OF UV FILTERS ALLOWED IN COSMETIC PRODUCTS

ANNEX VII

SYMBOLS USED ON PACKAGING/CONTAINER

- Reference to enclosed or attached information 1.
- 2. Period-after-opening
- 3. Date of minimum durability

ANNEX VIII

LIST OF VALIDATED ALTERNATIVE METHODS TO ANIMAL TESTING

This Annex lists the alternative methods validated by the European...

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ANNEX IX

PART A

Repealed Directive with its successive amendments

PART B

List of time-limits for transposition into national law and application

ANNEX X
CORRELATION TABLE

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- (1) OJ C 27, 3.2.2009, p. 34.
- (2) Opinion of the European Parliament of 24 March 2009 (not yet published in the Official Journal) and Council Decision of 20 November 2009.
- (**3**) OJ L 262, 27.9.1976, p. 169.
- (4) OJ L 396, 30.12.2006, p. 1.
- **(5)** OJ L 192, 11.7.1987, p. 49.
- (**6**) OJ L 196, 2.8.2003, p. 7.
- (7) OJ L 157, 30.4.2004, p. 45.
- (8) OJ L 241, 10.9.2008, p. 21.
- **(9)** OJ L 353, 31.12.2008, p. 1.
- (10) OJ L 358, 18.12.1986, p. 1.
- (11) OJ L 149, 11.6.2005, p. 22.
- (12) OJ L 184, 17.7.1999, p. 23.

Changes to legislation:

Regulation (EC) No 1223/2009 of the European Parliament and of the Council is up to date with all changes known to be in force on or before 19 March 2024. There are changes that may be brought into force at a future date. Changes that have been made appear in the content and are referenced with annotations.

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Changes and effects yet to be applied to:

- Annex 3 TABL Text repeal by EUR 2017/1410 Regulation (This amendment by the EU not applied to legislation.gov.uk because it is brought into force after IP completion day.)
- Annex 3 words inserted by S.I. 2023/764 Sch. 1
- Annex 3 table words inserted by S.I. 2023/836 reg. 3(2)Sch. 1

Changes and effects yet to be applied to the whole legislation item and associated provisions

Art. 2(1)(g) words substituted by S.I. 2019/696 Sch. 34 para. 3(c)(i) (This amendment not applied to legislation.gov.uk. Sch. 34 para. 3(c)(i) substituted immediately before IP completion day by virtue of S.I. 2020/1460, reg. 1(4), Sch. 3 para. 23(2)(a))