

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 Scope and objective
- Article 2 Definitions

CHAPTER II

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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 03 September 2023. 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

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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/chemical 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
4. Assessor's credentials and approval of part B

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

1. Reference to enclosed or attached information
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Repealed Directive with its successive amendments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:

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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))