

Commission Regulation (EC) No 523/2008 of 11 June 2008 amending Annexes VIII, X and XI to Regulation (EC) No 1774/2002 of the European Parliament and of the Council as regards the import of blood products for the manufacture of technical products (Text with EEA relevance)

- Article 1 Annexes VIII, X and XI to Regulation (EC) No 1774/2002...
- Article 2 Consignments of blood products accompanied by health certificates completed and...
- Article 3 This Regulation shall enter into force and apply on the...
Signature

ANNEX

The Annexes to Regulation (EC) No 1774/2002 are amended as...
In Annex VIII, Chapter IV is replaced by the following:...

Changes to legislation: Commission Regulation (EC) No 523/2008 is up to date with all changes known to be in force on or before 23 August 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](#)

- (1) [OJ L 273, 10.10.2002, p. 1](#). Regulation as last amended by Commission Regulation (EC) No 437/2008 ([OJ L 132, 22.5.2008, p. 7](#)).
- (2) [OJ L 125, 23.5.1996, p. 3](#). Directive as amended by Directive 2003/74/EC of the European Parliament and of the Council ([OJ L 262, 14.10.2003, p. 17](#)).

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Changes and effects yet to be applied to :

- Regulation implicit repeal by [EUR 2009/1069](#) Regulation