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)

ANNEX

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

(**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).

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.
View outstanding changes

Changes and effects yet to be applied to :

– Regulation implicit repeal by EUR 2009/1069 Regulation