Commission Regulation (EC) No 2007/2006 of 22 December 2006 implementing Regulation (EC) No 1774/2002 of the European Parliament and of the Council as regards the importation and transit of certain intermediate products derived from Category 3 material intended for technical uses in medical devices, in vitro diagnostics and laboratory reagents and amending that Regulation (Text with EEA relevance) (repealed)

COMMISSION REGULATION (EC) No 2007/2006

of 22 December 2006

implementing Regulation (EC) No 1774/2002 of the European Parliament and of the Council as regards the importation and transit of certain intermediate products derived from Category 3 material intended for technical uses in medical devices, in vitro diagnostics and laboratory reagents and amending that Regulation

(Text with EEA relevance) (repealed)

F1 . . . . . .

Textual Amendments

F1 Repealed by Commission Regulation (EU) No 142/2011 of 25 February 2011 implementing Regulation (EC) No 1069/2009 of the European Parliament and of the Council laying down health rules as regards animal by-products and derived products not intended for human consumption and implementing Council Directive 97/78/EC as regards certain samples and items exempt from veterinary checks at the border under that Directive Text with EEA relevance.

Status:
Point in time view as at 03/03/2011.

Changes to legislation:
There are currently no known outstanding effects for the Commission Regulation (EC) No 2007/2006 (repealed).