Regulation (EC) No 1069/2009 of the European Parliament and of the Council of 21 October 2009 laying down health rules as regards animal byproducts and derived products not intended for human consumption and repealing Regulation (EC) No 1774/2002 (Animal by-products Regulation)

TITLE II U.K.

OBLIGATIONS OF OPERATORS

CHAPTER II U.K.

Placing on the market

Section 3 U.K.

Derived products regulated by certain other [F1 retained EU law]

Article 33 U.K.

Placing on the market

Operators may place on the market the following derived products:

- (a) cosmetic products as defined in [F1Article 2(1)(a) of Regulation 1223/2009/EC];
- (b) active implantable medical devices as defined in Article 1(2)(c) of Directive 90/385/ EEC;
- (c) medical devices as defined in Article 1(2)(a) of Directive 93/42/EEC;
- (d) in vitro diagnostic medical devices as defined in Article 1(2)(b) of Directive 98/79/EC;
- (e) veterinary medicinal products as defined in Article 1(2) of Directive 2001/82/EC;
- (f) medicinal products as defined in Article 1(2) of Directive 2001/83/EC.

Textual Amendments

F1 Words in Art. 33(a) substituted (E.W.S.) (31.12.2020) by The Animals, Aquatic Animal Health, Invasive Alien Species, Plant Propagating Material and Seeds (Amendment) (EU Exit) Regulations 2020 (S.I. 2020/1388), regs. 1(2)(c), **12(17)**

Changes to legislation:

There are currently no known outstanding effects for the Regulation (EC) No 1069/2009 of the European Parliament and of the Council, Article 33.