

## STATUTORY INSTRUMENTS

# 2013 No. 2033

## The Veterinary Medicines Regulations 2013

### PART 2

#### Authorised veterinary medicinal products

#### Exemptions **E+W+S**

**15.**—(1) These Regulations do not apply to an inactivated autogenous vaccine that is manufactured, on the instructions of a veterinary surgeon, from pathogens or antigens obtained from an animal and used for the treatment of that animal.

(2) Schedule 1 and Part 1 of Schedule 2 do not apply in relation to an inactivated autogenous vaccine that is—

- (a) manufactured by a person and in premises authorised in accordance with Part 2 of Schedule 2, on the instructions of a veterinary surgeon, from pathogens or antigens obtained from an animal; and
- (b) used for the treatment of—
  - (i) other animals on the same site;
  - (ii) animals intended to be sent to those premises; or
  - (iii) animals on a site that receives animals from those premises.

(3) Schedule 1 and Part 1 of Schedule 2 do not apply in relation to—

- (a) blood or blood constituents from a blood bank authorised in accordance with [<sup>F1</sup>Part 2] of Schedule 2;
- (b) a product manufactured for administration under the cascade by a person and in premises authorised in accordance with [<sup>F2</sup>Part 2] of Schedule 2; or
- (c) <sup>F3</sup>... stem cell products for use as an autologous treatment for [<sup>F4</sup>non-food producing animals] from an <sup>F3</sup>... collection centre authorised in accordance with [<sup>F5</sup>Part 2] of Schedule 2.

(4) Schedule 6 (exemptions for small pet animals) has effect.

#### Extent Information

- E1** This version of this provision extends to England and Wales and Scotland only; a separate version has been created for Northern Ireland only

#### Textual Amendments

- F1** Words in [reg. 15\(3\)\(a\)](#) substituted (E.W.S.) (17.5.2024) by [The Veterinary Medicines \(Amendment etc.\) Regulations 2024 \(S.I. 2024/567\)](#), [regs. 1\(1\)](#), [10\(a\)](#)
- F2** Words in [reg. 15\(3\)\(b\)](#) substituted (E.W.S.) (17.5.2024) by [The Veterinary Medicines \(Amendment etc.\) Regulations 2024 \(S.I. 2024/567\)](#), [regs. 1\(1\)](#), [10\(b\)](#)

*Status: There are multiple versions of this provision on screen. These apply to different geographical extents. Skip to: E+W+S - England, Wales and Scotland extent N.I. - Northern Ireland extent*  
*Changes to legislation: There are currently no known outstanding effects for the The Veterinary Medicines Regulations 2013, Section 15. (See end of Document for details)*

- F3** Word in reg. 15(3)(c) omitted (E.W.S.) (17.5.2024) by virtue of [The Veterinary Medicines \(Amendment etc.\) Regulations 2024 \(S.I. 2024/567\)](#), regs. 1(1), **10(c)(i)**
- F4** Words in reg. 15(3)(c) substituted (E.W.S.) (17.5.2024) by [The Veterinary Medicines \(Amendment etc.\) Regulations 2024 \(S.I. 2024/567\)](#), regs. 1(1), **10(c)(ii)**
- F5** Words in reg. 15(3)(c) substituted (E.W.S.) (17.5.2024) by [The Veterinary Medicines \(Amendment etc.\) Regulations 2024 \(S.I. 2024/567\)](#), regs. 1(1), **10(c)(iii)**

## Exemptions **N.I.**

15.—(1) These Regulations do not apply to an inactivated autogenous vaccine that is manufactured, on the instructions of a veterinary surgeon, from pathogens or antigens obtained from an animal and used for the treatment of that animal.

(2) Schedule 1 and Part 1 of Schedule 2 do not apply in relation to an inactivated autogenous vaccine that is—

- (a) manufactured by a person and in premises authorised in accordance with Part 2 of Schedule 2, on the instructions of a veterinary surgeon, from pathogens or antigens obtained from an animal; and
- (b) used for the treatment of—
  - (i) other animals on the same site;
  - (ii) animals intended to be sent to those premises; or
  - (iii) animals on a site that receives animals from those premises.

(3) Schedule 1 and Part 1 of Schedule 2 do not apply in relation to—

- (a) blood or blood constituents from a blood bank authorised in accordance with Part 3 of Schedule 2;
- (b) a product manufactured for administration under the cascade by a person and in premises authorised in accordance with Part 4 of Schedule 2; or
- (c) equine stem cell products for use as an autologous treatment for horses from an equine collection centre authorised in accordance with Part 5 of Schedule 2.

(4) Schedule 6 (exemptions for small pet animals) has effect.

## Extent Information

- E2** This version of this provision extends to Northern Ireland only; a separate version has been created for England and Wales and Scotland only

**Status:**

There are multiple versions of this provision on screen. These apply to different geographical extents.

**Skip to:**

- E+W+S - England, Wales and Scotland extent
- N.I. - Northern Ireland extent

**Changes to legislation:**

There are currently no known outstanding effects for the The Veterinary Medicines Regulations 2013, Section 15.