Equip Rotavirus emulsion for injection for horses

Authorised

- Equine rotavirus A, type G3P12, strain H2, Inactivated

Product identification

Medicine name:
Equip Rotavirus emulsion for injection for horses

Active substance:
Equine rotavirus A, type G3P12, strain H2, Inactivated

Target species:
Horse

Route of administration:
Intramuscular use

Product details

Active substance and strength:
Equine rotavirus A, type G3P12, strain H2, Inactivated
74000000.00 fluorescent assay infectious dose 50% / 1.00 Dose

Pharmaceutical form:
Emulsion for injection

Withdrawal period by route of administration:

Intramuscular use:

- **Horse**
  - Meat and offal. 0 day

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QI05AA09

Legal status of supply:
Veterinary medicinal product subject to veterinary prescription

Authorisation status:
Valid

Authorised in:
Germany

Package description:
Single-dose type I glass syringes (Ph.Eur. 3.2.1) closed with bromobutyl rubber tips (Ph. Eur. 3.2.9).Syringes are supplied in cardboard packs of 3 units.
Single-dose type I glass syringes (Ph.Eur. 3.2.1) closed with bromobutyl rubber tips (Ph. Eur. 3.2.9).Syringes are supplied in cardboard packs of 10 units.
Single-dose type I glass syringes (Ph.Eur. 3.2.1) closed with bromobutyl rubber tips (Ph. Eur. 3.2.9).Syringes are supplied in cardboard packs of 20 units.
Single-dose type I glass syringes (Ph.Eur. 3.2.1) closed with bromobutyl rubber tips (Ph. Eur. 3.2.9).Syringes are supplied in cardboard packs of 40 units.

Additional information

Entitlement type:
Marketing Authorisation

Legal basis of product authorisation:
Full application (Article 12(3) of Directive No 2001/82/EC)

Marketing authorisation holder:

Zoetis Deutschland GmbH

Marketing authorisation date:
2/02/2012

Manufacturing sites for batch release:
Zoetis Belgium

Responsible authority:
Paul-Ehrlich-Institut

Authorisation number:
PEI.V.11552.01.1

Date of authorisation status change:
13/03/2017

Reference member state:
Ireland

Procedure number:
IE/V/0574/001

Concerned member states:
France Germany Italy Spain United Kingdom (Northern Ireland)

To consult adverse reactions on veterinary medicinal products please go to www.adrreports.eu/vet

Documents

Combined File of all Documents

English (PDF)
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