Equilis StrepE (--) - Lyophilisate and solvent for suspension for injection

Authorised

- Streptococcus equi subsp. equi, strain TW928, Live

Product identification

Medicine name:
Equilis StrepE (--) - Lyophilisate and solvent for suspension for injection

Active substance:
Streptococcus equi subsp. equi, strain TW928, Live

Target species:
Horse

Route of administration:
Submucosal use

Product details

Active substance and strength:
Streptococcus equi subsp. equi, strain TW928, Live
Presentation_strength:$10^{9.0}$ to $10^{9.4}$ CFU Reference:HSE Index:0

Pharmaceutical form:

Lyophilisate and solvent for suspension for injection

Withdrawal period by route of administration:

Submucosal use:

- **Horse**
  - Not applicable. 0 day Zero days

Anatomical therapeutic chemical veterinary (ATCvet) codes:

QI05AE

Legal status of supply:

Veterinary medicinal product subject to veterinary prescription

Authorisation status:

Valid

Authorised in:

Austria , Belgium , Bulgaria , Croatia , Cyprus , Czechia , Denmark , Estonia , Finland , France , Germany , Greece , Hungary , Iceland , Ireland , Italy , Latvia , Liechtenstein , Lithuania , Luxembourg , Malta , Netherlands , Norway , Poland , Portugal , Romania , Slovakia , Slovenia , Spain , Sweden , United Kingdom (Northern Ireland)

Available in:

Belgium , France , Netherlands

Package description:

Packaging:Lyophilisate: vial (glass); solvent: vial (glass), Package_size:10 vials lyophilisate, 10 vials solvent, 10 applicators, 10 syringes with needle, Content:Powder: 1 dose; solvent 0.5 ml

Additional information

Entitlement type:

Marketing Authorisation

Legal basis of product authorisation:

Complete application (stand-alone) - Council Directive 81/851/EEC

Marketing authorisation holder:
Intervet International B.V.

Marketing authorisation date:
7/05/2004

Manufacturing sites for batch release:
Intervet International B.V.

Responsible authority:
European Commission

Authorisation number:
This information is not available for this product.

Date of authorisation status change:
7/05/2004

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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