Equisolon 300 mg Sachet - Oral powder

Authorised

- Prednisolone

Product identification

Medicine name:
Equisolon 300 mg Sachet - Oral powder

Active substance:
Prednisolone

Target species:
Horse

Route of administration:
Oral use

Product details

Active substance and strength:
Prednisolone
300.00 milligram(s) / 1.00 Sachet

Pharmaceutical form:
Oral powder

Withdrawal period by route of administration:

Oral use:

- **Horse**
  - Meat and offal. 10 day

10 days (Not authorised for use in mares producing milk for human consumption)

Anatomical therapeutic chemical veterinary (ATCvet) codes:

QH02AB06

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:

Austria , Belgium , Croatia , Denmark , Finland , France , Germany , Ireland , Italy , Netherlands , Norway , Poland , Portugal , Slovenia , Spain , Sweden

Package description:

Packaging:Sachet (PET/PE/alu/PE/LD-LLDPE), Package_size:10 sachets of 9 g

Additional information

Entitlement type:

Marketing Authorisation

Legal basis of product authorisation:

Full application - Known active substance (Article 12(3) of Directive No 2001/82/EC)

Marketing authorisation holder:

Le Vet. B.V.

Marketing authorisation date:
12/03/2014

Manufacturing sites for batch release:
Lelypharma B.V.

Responsible authority:
European Commission

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

Date of authorisation status change:
12/03/2014

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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