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Equipred 50 mg tablets for horses

Authorised

- Prednisolone

Product identification

Medicine name:
Equipred 50 mg tablets for horses

Active substance:
Prednisolone

Target species:
Horse

Route of administration:
Oral use

Product details

Active substance and strength:
Prednisolone
50.00 milligram(s) / 1.00 Tablet

Pharmaceutical form:
Tablet

Withdrawal period by route of administration:

Oral use:

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

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QH02AB06

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

Authorisation status:
Valid

Authorised in:
Denmark

Available in:
Denmark

Package description:
PVC/PVDC/Alu blisters containing 10 tablets, in cartons of 50 tablets
PVC/PVDC/Alu blisters containing 10 tablets, in cartons of 100 tablets
PVC/PVDC/Alu blisters containing 10 tablets, in cartons of 200 tablets

Additional information

Entitlement type:
Marketing Authorisation

Legal basis of product authorisation:
Hybrid application (Article 13(3) of Directive No 2001/82/EC)

Marketing authorisation holder:
CP-Pharma Handelsgesellschaft mbH

Marketing authorisation date:
28/10/2019

Manufacturing sites for batch release:
CP-Pharma Handelsgesellschaft mbH

Responsible authority:
Danish Medicines Agency

Authorisation number:
61900

Date of authorisation status change:
28/10/2019

Reference member state:
Netherlands

Procedure number:
NL/V/0276/001

Concerned member states:
Austria Belgium Denmark Finland France Germany Hungary Ireland Italy Spain Sweden United Kingdom (Northern Ireland)

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

Documents

Summary of Product Characteristics

This document does not exist in this language (English). You can find it in another language below.

Combined File of all Documents

PuAR Equipred NL_V_0276_001_DC fianl.pdf