

File downloaded on 2026-06-24

Source URL: <https://medicines.health.europa.eu/veterinary/en/600000035207>

Prednicortone vet. 20 mg tablets for dogs

Not
authorised

- Prednisolone

Product identification

Medicine name:

Prednicortone vet. 20 mg tablets for dogs

Active substance:

Prednisolone

Target species:

Dog

Cat

Route of administration:

Oral use

Product details

Active substance and strength:

Prednisolone

20.00 milligram(s) / 1.00 Piece

Pharmaceutical form:

Tablet

Anatomical therapeutic chemical veterinary (ATCvet) codes:

QH02AB06

Legal status of supply:

Veterinary medicinal product subject to veterinary prescription

Authorisation status:

Surrendered

Authorised in:

Germany

Package description:

Cardboard box of 50 Aluminium - PVC/PE/PVDC blisters of 10 tablets.
Cardboard box of 5 Aluminium - PVC/PE/PVDC blisters of 10 tablets.
Cardboard box of 4 Aluminium - PVC/PE/PVDC blisters of 10 tablets.
Cardboard box of 3 Aluminium - PVC/PE/PVDC blisters of 10 tablets.
Cardboard box of 25 Aluminium - PVC/PE/PVDC blisters of 10 tablets.
Cardboard box of 2 Aluminium - PVC/PE/PVDC blisters of 10 tablets.
Cardboard box of 15 Aluminium - PVC/PE/PVDC blisters of 10 tablets.
Cardboard box of 10 Aluminium - PVC/PE/PVDC blisters of 10 tablets.
Cardboard box of 7 Aluminium - PVC/PE/PVDC blisters of 10 tablets.
Cardboard box of 1 Aluminium - PVC/PE/PVDC blister of 10 tablets.
Cardboard box of 9 Aluminium - PVC/PE/PVDC blisters of 10 tablets.
Cardboard box of 6 Aluminium - PVC/PE/PVDC blisters of 10 tablets.
Cardboard box of 8 Aluminium - PVC/PE/PVDC blisters of 10 tablets.

Additional information

Entitlement type:

Marketing Authorisation

Legal basis of product authorisation:

Generic application (Article 13(1) of Directive No 2001/82/EC)

Marketing authorisation holder:

CP-Pharma Handelsgesellschaft mbH

Marketing authorisation date:

22/07/2015

Manufacturing sites for batch release:

Lelypharma B.V.

Responsible authority:

Federal Office Of Consumer Protection And Food Safety

Authorisation number:

402147.01.00

Date of authorisation status change:

25/07/2023

Reference member state:

Netherlands

Procedure number:

NL/V/0190/002

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