Vetbromide

Authorised

- Potassium bromide

Product identification

Medicine name:
Vetbromide
Vetbromide, 600 mg tabletid koertele

Active substance:
Potassium bromide

Target species:
Dog

Route of administration:
Oral use

Product details

Active substance and strength:
Potassium bromide
600.00 milligram(s) / 1.00 Piece

Pharmaceutical form:
Tablet

Withdrawal period by route of administration:

Oral use:

- **Dog**

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QN03AX91

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

Authorisation status:
Valid

Authorised in:
Estonia

Available in:
Estonia

Package description:
Cardboard box containing 60 tablets (four PVC/PVDC/Aluminium blisters with 15 tablets each)
Cardboard box containing 120 tablets (eight PVC/PVDC/Aluminium blisters with 15 tablets each)

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:
Domes Pharma

Marketing authorisation date:
22/02/2021

Manufacturing sites for batch release:
EUROPHARTECH

Responsible authority:

State Agency Of Medicines

Authorisation number:
2281

Date of authorisation status change:
22/02/2021

Reference member state:
Netherlands

Procedure number:
NL/V/0346/001

Concerned member states:
Austria Belgium Bulgaria Croatia Cyprus Czechia Denmark Estonia Finland Germany Greece Hungary Ireland Italy Latvia Lithuania Luxembourg Malta Norway Poland Portugal Romania Slovakia Slovenia 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

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