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Fenylbutazon vet. 1 g Oralt pulver i dospåse

Authorised

- Phenylbutazone

Product identification

Medicine name:
Fenylbutazon vet. 1 g Oralt pulver i dospåse

Active substance:
Phenylbutazone

Target species:
Horse

Route of administration:
Oral use

Product details

Active substance and strength:
Phenylbutazone
1.00 gram(s) / 1.00 Sachet

Pharmaceutical form:
Oral powder in sachet

Withdrawal period by route of administration:

Oral use:

- **Horse**
  - All relevant tissues. no withdrawal period

Läkemedlet får endast användas till hästar som förklarats inte vara avsedda för livsmedelsproduktion, i enlighet med 2 kap 7 § Statens jordbruksverks föreskrifter (SJVFS 2009:84) om läkemedel och läkemedelsanvändning, D9.

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QM01AA01

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

Authorisation status:
Valid

Authorised in:
Sweden

Available in:
Sweden

Package description:
Available only in Swedish
Available only in Swedish

Additional information

Entitlement type:
Marketing Authorisation

Legal basis of product authorisation:
Legal basis not covered by Directive 2001/82/EC

Marketing authorisation holder:

Dechra Veterinary Products A/S

Marketing authorisation date:
15/12/1978

Manufacturing sites for batch release:
Genera d.d.

Responsible authority:
Swedish Medical Products Agency

Authorisation number:
9377

Date of authorisation status change:
15/12/1978

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

Documents

Package Leaflet

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Summary of Product Characteristics

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

Labelling

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