Butazocare flavour 1g granules in sachet for horses and ponies

Authorised

- Phenylbutazone

Product identification

Medicine name:
Butazocare flavour 1g granules in sachet for horses and ponies

Active substance:
Phenylbutazone

Target species:
Horse (non food-producing)

Route of administration:
Oral use

Product details

Active substance and strength:
Phenylbutazone
1000.00 milligram(s) / 2.00 gram(s)

Pharmaceutical form:
Granules in sachet

Withdrawal period by route of administration:

Oral use:

- **Horse (non food-producing)**

  - Meat and offal. no withdrawal period

Not to be used in horses intended for human consumption. Treated horses may never be slaughtered for human consumption. The horse must have been declared as not intended for human consumption under national horse passport legislation.

  - Milk. no withdrawal period

Not to be used in horses intended for human consumption. Treated horses may never be slaughtered for human consumption. The horse must have been declared as not intended for human consumption under national horse passport legislation.

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QM01AA01

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

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Authorisation status:
Valid

Authorised in:
Austria

Package description:
(ID2) 200 gram(s): unspecified outer container with 100 Bag (paper; polyethylene; aluminium; polyethylene) each with 2 gram(s)
(ID1) 64 gram(s): unspecified outer container with 32 Bag (paper; polyethylene; aluminium; polyethylene) each with 2 gram(s)

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:
Ecuphar

Marketing authorisation date:
25/03/2019

Manufacturing sites for batch release:
Labo Smeets

Responsible authority:
Austrian Agency For Health And Food Safety

Authorisation number:
838820

Date of authorisation status change:
25/03/2019

Reference member state:
Germany

Procedure number:
DE/V/0332/001

Concerned member states:
Austria United Kingdom (Northern Ireland)

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

English (PDF)
Published on: 7/01/2026
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