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PENBEX

Authorised

- Procaine hydrochloride
- Chlorphenamine maleate
- Betamethasone
- Dihydrostreptomycin sulfate
- Benzylpenicillin procaine

Product identification

Medicine name:
PENBEX

Active substance:
Procaine hydrochloride
Chlorphenamine maleate
Betamethasone
Dihydrostreptomycin sulfate
Benzylpenicillin procaine

Target species:
Cattle
Pig
Horse
Dog
Cat

Route of administration:
Intramuscular use

Product details

Active substance and strength:

Procaine hydrochloride
15.00 milligram(s) / 1.00 millilitre(s)

Chlorphenamine maleate
8.00 milligram(s) / 1.00 millilitre(s)

Betamethasone
0.50 milligram(s) / 1.00 millilitre(s)

Dihydrostreptomycin sulfate
250.00 milligram(s) / 1.00 millilitre(s)

Benzylpenicillin procaine
200000.00 international unit(s) / 1.00 millilitre(s)

Pharmaceutical form:

Suspension for injection

Withdrawal period by route of administration:

Intramuscular use:

- **Cattle**
  - Meat and offal. 30 day
  - Milk. 3 day Мляко: 3 дни (6 издоявания
- **Pig**
  - Meat and offal. 30 day
- **Horse**
  - Meat and offal. no withdrawal period

Не се разрешава употребата при коне, чиито месо е предназначено за човешка консумация.

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QJ01RA01

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

Authorisation status:
Valid

Authorised in:
Bulgaria

Package description:
Available only in Bulgarian
Available only in Bulgarian
Available only in Bulgarian
Available only in Bulgarian

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:
Industrial Veterinaria S.A.

Marketing authorisation date:
17/06/2007

Manufacturing sites for batch release:
Industrial Veterinaria S.A.

Responsible authority:
Bulgarian Food Safety Authority

Authorisation number:

0022-1820-16.07.2012

Date of authorisation status change:
15/07/2012

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

Documents

Summary of Product Characteristics

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Package Leaflet and Labelling

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