

PENBEX

Authorised

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

Product identification

Medicine name:

ПЕНБЕКС
PENBEX

Active substance:

Benzylpenicillin procaine
Dihydrostreptomycin sulfate
Betamethasone
Chlorphenamine maleate
Procaine hydrochloride

Target species:

Cattle
Pig
Horse
Dog
Cat

Route of administration:

Intramuscular use

Product details

Active substance and strength:

Benzylpenicillin procaine

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

Dihydrostreptomycin sulfate

250.00 milligram(s) / 1.00 millilitre(s)

Betamethasone

0.50 milligram(s) / 1.00 millilitre(s)

Chlorphenamine maleate

8.00 milligram(s) / 1.00 millilitre(s)

Procaine hydrochloride

15.00 milligram(s) / 1.00 millilitre(s)

Pharmaceutical form:

Suspension for injection

Withdrawal period by route of administration:

Intramuscular use:

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Cattle

- Meat and offal. 30 day

- Milk. 3 day Мляко: 3 дни (6 издојавания)

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Pig

- Meat and offal. 30 day

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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

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

Package Leaflet and Labelling

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