

Orbenin Extra Dry Cow 600 mg, intramammary suspension for cattle

Authorised

- Cloxacillin

Product identification

Medicine name:

Orbenin Extra Dry Cow 600 mg, интрамамарна супензия за говеда
Orbenin Extra Dry Cow 600 mg, intramammary suspension for cattle

Active substance:

Cloxacillin

Target species:

Cattle

Route of administration:

Intramammary use

Product details

Active substance and strength:

Cloxacillin

600.00 milligram(s) / 1.00 Syringe

Pharmaceutical form:

Intramammary suspension

Withdrawal period by route of administration:**Intramammary use:**

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Cattle

- Meat and offal. 28 day
- Milk. 96 hour

Мляко: 96 часа след отелване. Не се препоръчва при крави с по – къс от 42 дни сухостоен период.

Anatomical therapeutic chemical veterinary (ATCvet) codes:

QJ01CF02

Legal status of supply:

Veterinary medicinal product subject to veterinary prescription

Authorisation status:

Valid

Authorised in:

Bulgaria

Available in:

Bulgaria

Package description:

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:

Zoetis Belgium

Marketing authorisation date:

14/12/2006

Manufacturing sites for batch release:

Haupt Pharma Latina S.r.l.

Responsible authority:

Bulgarian Food Safety Authority

Authorisation number:

0022-1642

Date of authorisation status change:

14/12/2006

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

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.