

Orbenin EDC 600 mg/ 3,6 g intramammális szuszpenzió szarvasmarhák részére

Authorised

- Cloxacillin

Product identification

Medicine name:

Orbenin EDC 600 mg/ 3,6 g intramammális szuszpenzió szarvasmarhák részére

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 solution

Withdrawal period by route of administration:

Intramammary use:

-

Cattle

- Meat and offal. 49 day

Anatomical therapeutic chemical veterinary (ATCvet) codes:

QJ51CF02

Legal status of supply:

Veterinary medicinal product subject to veterinary prescription

Authorisation status:

Valid

Authorised in:

Hungary

Package description:

Available only in [Hungarian](#)

Available only in [Hungarian](#)

Available only in [Hungarian](#)

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 Hungary Kft.

Marketing authorisation date:

16/04/1998

Manufacturing sites for batch release:

Haupt Pharma Latina S.r.l.

Responsible authority:

Directorate Of Veterinary Medicinal Products

Authorisation number:

This information is not available for this product.

Date of authorisation status change:

16/04/1998

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.