

Orbenin EDC, 600 mg intramammaarsuspensioo veistele

Authorised

- Cloxacillin

Product identification

Medicine name:

Orbenin EDC, 600 mg intramammaarsuspensioo veistele

Active substance:

Cloxacillin

Target species:

Cattle (dairy cow)

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:

-

Cattle (dairy cow)

- Meat and offal. 28 day

Ravi ajal loomi inимtoiduks tappa ei tohi. Loomi võib inимtoiduks tappa 28 päeva pärast ravimi manustamist.

- Milk. 96 hour

Mitte kasutada lehmadel, kelle kinnisjärgu pikkus on 42 päeva või vähem. Piima võib inимtoiduks kasutada 96 tundi pärast poegimist (kaks korda päevas lüpsmise puhul 8. lüpsikorral). Kui lehm poegib varem kui 42 päeva pärast ravimi manustamist, võib piima inимtoiduks kasutada 42 päeva pluss 96 tundi pärast ravimi manustamist.

Anatomical therapeutic chemical veterinary (ATCvet) codes:

QJ51CF02

Legal status of supply:

Veterinary medicinal product subject to veterinary prescription

Authorisation status:

Valid

Authorised in:

Estonia

Available in:

Estonia

Package description:

Available only in Estonian

Available only in Estonian

Available only in Estonian

Available only in Estonian

Additional information

Entitlement type:

Marketing Authorisation

Legal basis of product authorisation:

Full application (Article 12(3) of Directive No 2001/82/EC)

Marketing authorisation holder:

Zoetis Belgium

Marketing authorisation date:

6/04/2006

Manufacturing sites for batch release:

Haupt Pharma Latina S.r.l.

Responsible authority:

State Agency Of Medicines

Authorisation number:

1386

Date of authorisation status change:

6/04/2006

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

Documents

Summary of Product Characteristics

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