

Ubrostar Red 100 mg / 280 mg / 100 mg, Intramammary Suspension for Cattle

Authorised

- Penethamate hydriodide
- Framycetin sulfate
- Benethamine penicillin

Product identification

Medicine name:

Ubrostar Red 100 mg / 280 mg / 100 mg Intramammary Suspension for cattle
Ubrostar Red 100 mg / 280 mg / 100 mg, Intramammary Suspension for Cattle

Active substance:

Penethamate hydriodide
Framycetin sulfate
Benethamine penicillin

Target species:

Cattle

Route of administration:

Intramammary use

Product details

Active substance and strength:

Penethamate hydriodide
100.00 milligram(s) / 1.00 Syringe

Framycetin sulfate
100.00 milligram(s) / 1.00 Syringe

Benethamine penicillin
280.00 milligram(s) / 1.00 Syringe

Pharmaceutical form:

Intramammary suspension

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

- Meat and offal. 10 day
 - Milk. 37 day
-

Anatomical therapeutic chemical veterinary (ATCvet) codes:

QJ51RC25

Legal status of supply:

Veterinary medicinal product subject to veterinary prescription

Authorisation status:

Valid

Authorised in:

United Kingdom (Northern Ireland)

Package description:

Cardboard box or plastic container containing 20 single use intramammary syringes and 20 teat wipes (containing isopropanol 70%). Each 4.5 g syringe (cylinder with piston and cap, all made of low density polyethylene) contains 5 ml intramammary suspension.

Cardboard box or plastic container containing 60 single use intramammary syringes and 60 teat wipes (containing isopropanol 70%). Each 4.5 g syringe (cylinder with piston and cap, all made of low density polyethylene) contains 5 ml intramammary suspension.

Cardboard box or plastic container containing 120 single use intramammary syringes and 120 teat wipes (containing isopropanol 70%). Each 4.5 g syringe (cylinder with piston and cap, all made of low density polyethylene) contains 5 ml intramammary suspension.

Additional information

Entitlement type:

Marketing Authorisation

Legal basis of product authorisation:

Generic application (Article 13(1) of Directive No 2001/82/EC)

Marketing authorisation holder:

Boehringer Ingelheim Animal Health UK Limited

Marketing authorisation date:

13/12/2011

Manufacturing sites for batch release:

Lohmann Pharma Herstellung GmbH

Haupt Pharma Latina S.r.l.

Responsible authority:

The Veterinary Medicines Directorate

Authorisation number:

Vm 08327/4307

Date of authorisation status change:

29/08/2022

Reference member state:

Ireland

Procedure number:

IE/V/0271/001

Concerned member states:

Austria Belgium Bulgaria Cyprus Czechia Estonia France Germany Greece
Hungary Latvia Lithuania Netherlands Poland Portugal Romania Slovakia
Slovenia Spain United Kingdom (Northern Ireland)

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

Documents

Summary of Product Characteristics

Source URL: <https://medicines.health.europa.eu/veterinary/600000052260>