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# Ubrostar Red 100 mg / 280 mg / 100 mg Intramammary Suspension for cattle

Authorised

- Framycetin sulfate
- Benethamine penicillin
- Penethamate hydriodide

## Product identification

**Medicine name:**
Ubrostar Red 100 mg / 280 mg / 100 mg Intramammary Suspension for cattle

**Active substance:**
Framycetin sulfate

Benethamine penicillin

Penethamate hydriodide

**Target species:**
Cattle

**Route of administration:**
Intramammary use

## Product details

**Active substance and strength:**

Framycetin sulfate
100.00 milligram(s) / 1.00 Syringe

Benethamine penicillin
280.00 milligram(s) / 1.00 Syringe

Penethamate hydriodide
100.00 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:**
Poland

**Available in:**
Poland

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

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

**Marketing authorisation date:**
12/12/2011

**Manufacturing sites for batch release:**
Lohmann Pharma Herstellung GmbH
Haupt Pharma Latina S.r.l.

**Responsible authority:**
Office For Registration Of Medicinal Products Medical Devices And Biocidal Products

**Authorisation number:**
2161

**Date of authorisation status change:**
12/12/2011

**Reference member state:**
Ireland

**Procedure number:**
IE/V/0271/001

**Concerned member states:**
Belgium Bulgaria Czechia France Hungary Netherlands Poland Romania Slovakia Slovenia United Kingdom (Northern Ireland)

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

## Documents

Summary of Product Characteristics

English (PDF)
Published on: 23/08/2024
Updated on: 14/03/2026
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Package Leaflet

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