

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

Authorised

- Penethamate hydriodide
- Benethamine penicillin
- Framycetin sulfate

## Product identification

**Medicine name:**

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

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**Active substance:**

Penethamate hydriodide  
Benethamine penicillin  
Framycetin sulfate

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**Target species:**

Cattle

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**Route of administration:**

Intramammary use

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## Product details

**Active substance and strength:**

Penethamate hydriodide  
100.00 milligram(s) / 1.00 Syringe  
Benethamine penicillin  
280.00 milligram(s) / 1.00 Syringe  
Framycetin sulfate  
100.00 milligram(s) / 1.00 Syringe

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**Pharmaceutical form:**

Intramammary suspension

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**Withdrawal period by route of administration:**

**Intramammary use:**

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**Cattle**

- Meat and offal. 10 day
  - Milk. 37 day
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**Anatomical therapeutic chemical veterinary (ATCvet) codes:**

QJ51RC25

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**Legal status of supply:**

Veterinary medicinal product subject to veterinary prescription

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**Authorisation status:**

Valid

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**Authorised in:**

Slovakia

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**Package description:**

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.

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 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.

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## Additional information

**Entitlement type:**

Marketing Authorisation

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**Legal basis of product authorisation:**

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

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**Marketing authorisation holder:**

Boehringer Ingelheim Vetmedica GmbH

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**Marketing authorisation date:**

2/12/2011

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**Manufacturing sites for batch release:**

Haupt Pharma Latina S.r.l.

Lohmann Pharma Herstellung GmbH

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**Responsible authority:**

Institute For State Control Of Veterinary Biologicals And Medicaments

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**Authorisation number:**

96/060/DC/11-S

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**Date of authorisation status change:**

2/12/2011

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**Reference member state:**

Ireland

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**Procedure number:**

IE/V/0271/001

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**Concerned member states:**

Belgium Bulgaria Czechia France Hungary Netherlands Poland Romania  
Slovakia Slovenia United Kingdom (Northern Ireland)

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To consult adverse reactions on veterinary medicinal products please go to  
[www.adrreports.eu/vet](http://www.adrreports.eu/vet)

## Documents

Combined File of all Documents

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Summary of Product Characteristics