

Lenzelta, Suspension for injection

Authorised

- Escherichia coli, serotype O111, strain J5, Inactivated
- Staphylococcus aureus, strain DSM 4910, Inactivated

Product identification

Medicine name:

Lenzelta, Suspension for injection

LENZELTA SUSPENSION INJECTABLE POUR BOVINS

Active substance:

Escherichia coli, serotype O111, strain J5, Inactivated

Staphylococcus aureus, strain DSM 4910, Inactivated

Target species:

Cattle

Route of administration:

Intramuscular use

Product details

Active substance and strength:

Escherichia coli, serotype O111, strain J5, Inactivated

1.00 relative potency / 1.00 Dose

Staphylococcus aureus, strain DSM 4910, Inactivated

1.00 relative potency / 1.00 Dose

Pharmaceutical form:

Suspension for injection

Withdrawal period by route of administration:**Intramuscular use:**

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Cattle

- Meat and offal. 0 day

- Milk. 0 hour

Anatomical therapeutic chemical veterinary (ATCvet) codes:

QI02AB17

Legal status of supply:

Veterinary medicinal product subject to veterinary prescription

Authorisation status:

Valid

Authorised in:

France

Package description:

Glass Vial 10 x 5.0 Dose obal- plastová krabička

Plastic (HDPE) Vial 10 x 5.0 Dose obal- plastová krabička

Glass Vial 1 x 5.0 Dose obal- papírová krabička

Plastic (HDPE) Vial 1 x 5.0 Dose obal- papírová krabička

Glass Vial 1 x 25.0 Dose obal- papírová krabička

Plastic (HDPE) Vial 1 x 25.0 Dose obal- papírová krabička

Glass Vial 1 x 50.0 Dose obal- papírová krabička

Plastic (HDPE) Vial 1 x 50.0 Dose obal- papírová krabička

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:

Boehringer Ingelheim Vetmedica GmbH

Marketing authorisation date:

17/11/2025

Manufacturing sites for batch release:

Bioveta a.s.

Responsible authority:

French Agency For Food, Environmental And Occupational Health & Safety

Authorisation number:

FR/V/1783815 5/2025

Date of authorisation status change:

17/11/2025

Reference member state:

Czechia

Procedure number:

CZ/V/0204/001

Concerned member states:

Austria Belgium Denmark Finland France Germany Greece Ireland Italy
Luxembourg Netherlands Norway Portugal Spain Sweden
United Kingdom (Northern Ireland)

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 and Labelling

This document does not exist in this language (English). You can find it in another language below.