

belfer 100 mg/ml Solution for injection for horses, cattle, pigs, sheep, goats and dogs

Authorised

- IRON(III)-HYDROXIDE DEXTRAN COMPLEX

Product identification

Medicine name:

belfer 100 mg/ml Solution for injection for horses, cattle, pigs, sheep, goats and dogs

Active substance:

IRON(III)-HYDROXIDE DEXTRAN COMPLEX

Target species:

Cattle

Pig

Dog

Goat

Sheep

Horse (suckling foal)

Route of administration:

Intravenous use

Intramuscular use

Product details

Active substance and strength:

IRON(III)-HYDROXIDE DEXTRAN COMPLEX

333.33 milligram(s) / 1.00 millilitre(s)

Pharmaceutical form:

Solution for injection

Withdrawal period by route of administration:

Intravenous use:

-

Cattle

- Milk. 0 hour
- Meat and offal. 0 day

-

Pig

- Meat and offal. 0 day

Intramuscular use:

-

Cattle

- Milk. 0 hour
- Meat and offal. 0 day

-

Goat

- Milk. 0 hour
- Meat and offal. 0 day

-

Sheep

- Milk. 0 hour
- Meat and offal. 0 day

-

Pig

- Meat and offal. 0 day

-

Horse (suckling foal)

- Meat and offal. 0 day

- Milk. 0 hour

Anatomical therapeutic chemical veterinary (ATCvet) codes:

QB03AC

Legal status of supply:

Veterinary medicinal product subject to veterinary prescription

Authorisation status:

Valid

Authorised in:

Latvia

Package description:

(ID1) 100 millilitre(s): unspecified outer container with 1 Vial (brown glass) with 100 millilitre(s), closed with Stopfen (bromobutyl rubber)

(ID2) 600 millilitre(s): unspecified outer container with 6 Vial (brown glass) each with 100 millilitre(s), closed with Stopfen (bromobutyl rubber)

(ID3) 1200 millilitre(s): unspecified outer container with 12 Vial (brown glass) each with 100 millilitre(s), closed with Stopfen (bromobutyl rubber)

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:

Bela-Pharm GmbH & Co. KG

Marketing authorisation date:

3/01/2018

Manufacturing sites for batch release:

Bela-Pharm GmbH & Co. KG

Responsible authority:

Food And Veterinary Service

Authorisation number:

V/DCP/17/0050

Date of authorisation status change:

3/01/2018

Reference member state:

Germany

Procedure number:

DE/V/0167/001

Concerned member states:

Austria Denmark Estonia Finland Greece Hungary Iceland Latvia Lithuania
Poland Portugal Romania Slovenia Spain Sweden

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

Documents

Summary of Product Characteristics

English (RTF)

Published on: 12/01/2026

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Package Leaflet

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Labelling

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

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