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:
Finland

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:
9/04/2019

Manufacturing sites for batch release:
Bela-Pharm GmbH & Co. KG

Responsible authority:
Finnish Medicines Agency

Authorisation number:
34211

Date of authorisation status change:
9/04/2019

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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