

Uniferon 200 mg/ml solution for injection

Authorised

- Iron dextran
- Iron dextran

Product identification

Medicine name:

Uniferon 200 mg/ml solution for injection

Active substance:

Iron dextran

Iron dextran

Target species:

Pig

Route of administration:

Subcutaneous use

Intramuscular use

Product details

Active substance and strength:

Iron dextran

560.00 milligram(s) / 1.00 millilitre(s)

Iron dextran
560.00 milligram(s) / 1.00 millilitre(s)

Pharmaceutical form:

Solution for injection

Anatomical therapeutic chemical veterinary (ATCvet) codes:

QB03AC

Legal status of supply:

Veterinary medicinal product subject to veterinary prescription

Authorisation status:

Valid

Authorised in:

Portugal

Available in:

Portugal

Package description:

12 x 200 ml vials (LDPE) in a carton box
12 x 100 ml vials (LDPE) in a carton box
12 x 100 ml vials (glass) in a carton box
12 x 100 ml vials (HDPE) in a carton box
20 x 100 ml vials (glass) in a carton box
5 x 100 ml vials (glass) in a carton box
20 x 100 ml vials (LDPE) in a carton box
20 x 100 ml vials (HDPE) in a carton box
5 x 100 ml vials (LDPE) in a carton box
5 x 100 ml vials (HDPE) in a carton box

Additional information

Entitlement type:

Marketing Authorisation

Legal basis of product authorisation:

Legal basis not covered by Directive 2001/82/EC

Marketing authorisation holder:

Pharmacosmos A/S

Marketing authorisation date:

15/11/2010

Manufacturing sites for batch release:

Pharmacosmos A/S

Responsible authority:

Directorate General For Food And Veterinary

Authorisation number:

289/01/10RFVPT

Date of authorisation status change:

17/07/2026

Reference member state:

Denmark

Procedure number:

DK/V/0114/001

Concerned member states:

Czechia Estonia Finland France Germany Iceland Italy Latvia Lithuania
Netherlands Norway Poland Portugal Romania Slovakia Spain

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

Documents

Combined File of all Documents

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