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Uniferon 200 mg/ml Solution for Injection for pigs (piglets)

Authorised

- Iron dextran

Product identification

Medicine name:
Uniferon 200 mg/ml Solution for Injection for pigs (piglets)

Active substance:
Iron dextran

Target species:
Pig

Route of administration:
Intramuscular use

Product details

Active substance and strength:
Iron dextran
551.33 milligram(s) / 1.00 millilitre(s)

Pharmaceutical form:
Solution for injection

Withdrawal period by route of administration:

Intramuscular use:

- **Pig**
  - Meat and offal. 0 day

Anatomical therapeutic chemical veterinary (ATCvet) codes:

QB03AC

Legal status of supply:

Veterinary medicinal product not subject to veterinary prescription

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Authorisation status:

Valid

Authorised in:

Ireland

Available in:

Ireland

Package description:

Pack sizes of 5 x 100 ml multidose collapsible polyethylene vials. Vials are closed by rubberstoppers and aluminium caps. They are packed in transparent foil.The packaging may contain an injector.

Pack sizes of 10 x 100 ml multidose collapsible polyethylene vials. Vials are closed by rubberstoppers and aluminium caps. They are packed in transparent foil.The packaging may contain an injector.

Pack sizes of 12 x 200 ml multidose collapsible polyethylene vials. Vials are closed by rubberstoppers and aluminium caps. They are packed in transparent foil.The packaging may contain an injector.

Pack sizes of 12 x 200 ml multidose collapsible polyethylene vials. Vials are closed by rubberstoppers and aluminium caps. They are packed in aluminium foil.The packaging may contain an injector.

Pack sizes of 10 x 100 ml multidose collapsible polyethylene vials. Vials are closed by rubberstoppers and aluminium caps. They are packed in aluminium foil.The packaging may contain an injector.

Pack sizes of 5 x 100 ml multidose collapsible polyethylene vials. Vials are closed by rubberstoppers and aluminium caps. They are packed in aluminium foil.The packaging may contain an injector.

Additional information

Entitlement type:
Marketing Authorisation

Legal basis of product authorisation:
Complete application (stand-alone) - Council Directive 81/851/EEC

Marketing authorisation holder:
Pharmacosmos A/S

Marketing authorisation date:
3/03/2006

Manufacturing sites for batch release:
Pharmacosmos A/S

Responsible authority:
Health Products Regulatory Authority

Authorisation number:
VPA10794/001/001

Date of authorisation status change:
3/03/2006

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

Documents

Summary of Product Characteristics