

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

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