Dexasone 2 mg/ml Roztwór do wstrzykiwań

Authorised

- Dexamethasone sodium phosphate

Product identification

Medicine name:
Dexasone 2 mg/ml Roztwór do wstrzykiwań

Active substance:
Dexamethasone sodium phosphate

Target species:
Cattle
Cat
Horse
Dog

Route of administration:
Intramuscular use
Intravenous use

Product details

Active substance and strength:
Dexamethasone sodium phosphate
2.00 milligram(s) / 1.00 millilitre(s)

Pharmaceutical form:
Solution for injection

Withdrawal period by route of administration:

Intramuscular use:

- **Cattle**
  - Meat and offal. 21 day
  - Milk. 72 hour
- **Horse**
  - Meat and offal. 21 day
  - Milk. 72 hour

Intravenous use:

- **Cattle**
  - Meat and offal. 21 day
  - Milk. 72 hour
- **Horse**
  - Meat and offal. 21 day
  - Milk. 72 hour

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QH02AB02

Legal status of supply:
Veterinary medicinal product subject to veterinary prescription

Authorisation status:
Valid

Authorised in:
Poland

Available in:
Poland

Package description:
Available only in Polish

Additional information

Entitlement type:
Marketing Authorisation

Legal basis of product authorisation:
Well-established use application (Article 13a of Directive No 2001/82/EC)

Marketing authorisation holder:
Scanvet Poland Sp. z o.o.

Marketing authorisation date:
28/06/1994

Manufacturing sites for batch release:
Norbrook Laboratories Limited
Norbrook Manufacturing Limited

Responsible authority:
Office For Registration Of Medicinal Products Medical Devices And Biocidal Products

Authorisation number:
0041

Date of authorisation status change:
28/06/1994

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

Documents

Package Leaflet

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Labelling

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

Summary of Product Characteristics

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