DEXAMETAZONA FP, soluție injectabilă

Authorised

- Dexamethasone

Product identification

Medicine name:
DEXAMETAZONA FP, soluție injectabilă

Active substance:
Dexamethasone

Target species:
Cattle
Horse
Goat
Pig
Dog
Cat

Route of administration:
Intravenous use
Subcutaneous use
Intramuscular use
Intraarticular use

Product details

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

Pharmaceutical form:
Solution for injection

Withdrawal period by route of administration:
Intravenous use:

- **Cattle**
  - Meat and offal. 28 day
  - Milk. 7 day
- **Horse**
  - Meat and offal. 28 day
- **Goat**
  - Meat and offal. 28 day
  - Milk. 7 day
- **Pig**
  - Meat and offal. 28 day

Subcutaneous use:

- **Cattle**
  - Meat and offal. 28 day
  - Milk. 7 day
-

Horse

- Meat and offal. 28 day

•

Goat

- Meat and offal. 28 day
- Milk. 7 day

•

Pig

- Meat and offal. 28 day

Intramuscular use:

•

Cattle

- Meat and offal. 28 day
- Milk. 7 day

•

Horse

- Meat and offal. 28 day

•

Goat

- Meat and offal. 28 day
- Milk. 7 day

•

Pig

- Meat and offal. 28 day

Intraarticular use:

•

Cattle

- Meat and offal. 28 day
- Milk. 7 day

- **Horse**
  - Meat and offal. 28 day
- **Goat**
  - Meat and offal. 28 day
  - Milk. 7 day
- **Pig**
  - Meat and offal. 28 day

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QH02AB02

Legal status of supply:
This information is not available for this product.

Authorisation status:
Valid

Authorised in:
Romania

Available in:
Romania

Package description:
Available only in Romanian
Available only in Romanian
Available only in Romanian
Available only in Romanian
Available only in Romanian
Available only in Romanian
Available only in Romanian
Available only in Romanian

Additional information

Entitlement type:
Marketing Authorisation

Legal basis of product authorisation:
Complete application (stand-alone) - Directive No 2001/82/EC

Marketing authorisation holder:
Pasteur Filiala Filipesti S.A.

Marketing authorisation date:
7/12/2005

Manufacturing sites for batch release:
Pasteur Filiala Filipesti S.A.

Responsible authority:
Institute For Control Of Biological Products And Veterinary Medicines

Authorisation number:
110214

Date of authorisation status change:
6/02/2022

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

Documents

Summary of Product Characteristics

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