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DEXAZONE SOLUTION INJECTABLE

Authorised

- Dexamethasone

Product identification

Medicine name:

DEXAZONE SOLUTION INJECTABLE

Active substance:

Dexamethasone

Target species:

Cattle

Pig

Cat

Horse

Horse (mare)

Goat

Dog

Route of administration:

Intramuscular use

Subcutaneous use

Periarticular use

Intravenous 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:

Intramuscular use:

-

Cattle

- Milk. 3 day

- Meat and offal. 8 day

-

Pig

- Meat and offal. 6 day

-

Horse

- Meat and offal. 6 day

-

Horse (mare)

- Milk. 3 day

-

Goat

- Milk. 3 day

- Meat and offal. 8 day

Subcutaneous use:

-

Cattle

- Milk. 3 day

- Meat and offal. 8 day

-

Pig

- Meat and offal. 6 day

-

Horse

- Meat and offal. 6 day

-

Horse (mare)

- Milk. 3 day

-

Goat

- Milk. 3 day

- Meat and offal. 8 day

Periarticular use:

-

Cattle

- Milk. 3 day

- Meat and offal. 8 day

-

Pig

- Meat and offal. 6 day

-

Horse

- Meat and offal. 6 day

-

Horse (mare)

- Milk. 3 day

-

Goat

- Milk. 3 day
- Meat and offal. 8 day

Intravenous use:

-

Cattle

- Milk. 3 day
- Meat and offal. 8 day

-

Pig

- Meat and offal. 6 day

-

Horse

- Meat and offal. 6 day

-

Horse (mare)

- Milk. 3 day

-

Goat

- Milk. 3 day
- Meat and offal. 8 day

Intraarticular use:

-

Cattle

- Milk. 3 day
- Meat and offal. 8 day

-

Pig

- Meat and offal. 6 day

•

Horse

- Meat and offal. 6 day

•

Horse (mare)

- Milk. 3 day

•

Goat

- Milk. 3 day

- Meat and offal. 8 day

Anatomical therapeutic chemical veterinary (ATCvet) codes:

QH02AB02

Legal status of supply:

Veterinary medicinal product subject to veterinary prescription

Authorisation status:

Valid

Authorised in:

France

Available in:

France

Package description:

Available only in [French](#)

Available only in [French](#)

Additional information

Entitlement type:

Marketing Authorisation

Legal basis of product authorisation:

Legal basis not covered by Directive 2001/82/EC

Marketing authorisation holder:

Virbac

Marketing authorisation date:

21/05/1984

Manufacturing sites for batch release:

Virbac

Responsible authority:

French Agency For Food, Environmental And Occupational Health & Safety

Authorisation number:

FR/V/6932577 5/1984

Date of authorisation status change:

21/05/2009

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

Documents

Summary of Product Characteristics

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