

Alfadexx UDA 2 mg/ml oplossing voor injectie voor paard, rund, geit, varken, hond en kat

Authorised

- Dexamethasone sodium phosphate

Product identification

Medicine name:

Alfadexx UDA 2 mg/ml oplossing voor injectie voor paard, rund, geit, varken, hond en kat

Active substance:

Dexamethasone sodium phosphate

Target species:

Cattle

Goat (adult female)

Pig

Horse

Dog

Cat

Route of administration:

Intramuscular use

Subcutaneous use

Product details

Active substance and strength:

Dexamethasone sodium phosphate

2.63 milligram(s) / 1.00 millilitre(s)

Pharmaceutical form:

Solution for injection

Withdrawal period by route of administration:

Intramuscular use:

-

Cattle

- Milk. 72 hour
- Meat and offal. 8 day

-

Goat (adult female)

- Milk. 72 hour
- Meat and offal. 8 day

-

Pig

- Meat and offal. 2 day

-

Horse

- Meat and offal. 8 day

Subcutaneous use:

-

Cattle

- Milk. 72 hour
- Meat and offal. 8 day

-

Goat (adult female)

- Milk. 72 hour
- Meat and offal. 8 day

-

Pig

- Meat and offal. 2 day

-

Horse

- 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:

Netherlands

Package description:

Available only in Dutch

Available only in Dutch

Additional information

Entitlement type:

Marketing Authorisation

Legal basis of product authorisation:

Generic application (Article 13(1) of Directive No 2001/82/EC)

Marketing authorisation holder:

Alfasan Nederland B.V.

Marketing authorisation date:

20/09/2021

Manufacturing sites for batch release:

Produlab Pharma Production B.V.

Alfasan Nederland B.V.

Responsible authority:

Medicines Evaluation Board

Authorisation number:

REG NL 128701

Date of authorisation status change:

20/09/2021

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

Documents

Combined File of all Documents

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