LUROCAINE

Authorised

• Lidocaine hydrochloride monohydrate

Product identification

Medicine name:

LUROCAINE

Active substance:

Lidocaine hydrochloride monohydrate

Target species:

Dog

Cat

Equid

Route of administration:

Intramuscular use

Retrobulbar use

Subcutaneous use

Paravertebral use

Ocular use

Intravenous use

Intraperitoneal use

Intraarticular use

Product details

Active substance and strength:

Pharmaceutical form:

Solution for injection

Withdrawal period by route of administration:

Intramuscular use:

•

Dog

•

Cat

•

Equid

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

Retrobulbar use:

•

Dog

•

Cat

Subcutaneous use:

•

Dog

•

Cat

Paravertebral use:

•

Equid

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

Ocular use:

•

Dog Cat **Equid** - Milk. 3 day - Meat and offal. 3 day **Intravenous use:** Dog Cat **Equid** - Milk. 3 day - Meat and offal. 3 day Intraperitoneal use: Dog Cat **Equid** - Milk. 3 day - Meat and offal. 3 day Intraarticular use: **Equid** - Milk. 3 day - Meat and offal. 3 day

Legal status of supply:

Veterinary medicinal product subject to veterinary prescription

Authorisation status:

Valid

Authorised in:

France

Package description:

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:

Vetoquinol S.A.

Marketing authorisation date:

21/12/1984

Manufacturing sites for batch release:

Vetoquinol Biowet Sp. z o.o.

VETOQUINOL

Responsible authority:

French Agency For Food, Environmental And Occupational Health & Safety

Authorisation number:

FR/V/9931286 6/1984

Date of authorisation status change:

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

Documents

Summary of Product Characteristics

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