

EKYFLOGYL 1.8 MG/ML + 8.7 MG/ML GEL FOR HORSES

Authorised

- Prednisolone acetate
- Lidocaine hydrochloride monohydrate

Product identification

Medicine name:

EKYFLOGYL 1.8 MG/ML + 8.7 MG/ML GEL FOR HORSES

Active substance:

Prednisolone acetate

Lidocaine hydrochloride monohydrate

Target species:

Horse

Route of administration:

Cutaneous use

Product details

Active substance and strength:

Prednisolone acetate

2.00 milligram(s) / 1.00 millilitre(s)

Lidocaine hydrochloride monohydrate

10.70 milligram(s) / 1.00 millilitre(s)

Pharmaceutical form:

Gel

Withdrawal period by route of administration:

Cutaneous use:

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Horse

- Meat and offal. 10 day
- Milk. no withdrawal period

Not authorised for use in animals producing milk for human consumption.

Anatomical therapeutic chemical veterinary (ATCvet) codes:

QM02AX99

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

Additional information

Entitlement type:

Marketing Authorisation

Legal basis of product authorisation:

Hybrid application (Article 13(3) of Directive No 2001/82/EC)

Marketing authorisation holder:

Audevard

Marketing authorisation date:

24/07/2019

Manufacturing sites for batch release:

Dopharma France

Responsible authority:

French Agency For Food, Environmental And Occupational Health & Safety

Authorisation number:

FR/V/0777108 2/2019

Date of authorisation status change:

4/05/2023

Reference member state:

France

Procedure number:

FR/V/0344/001

Concerned member states:

Austria Denmark Finland Germany Ireland Luxembourg Netherlands Norway
Poland Portugal Sweden

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.

Package Leaflet and Labelling

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

Combined File of all Documents

eu-puar-frv0344001-mr-rpe532-en.pdf