

# 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

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**Active substance:**

Prednisolone acetate

Lidocaine hydrochloride monohydrate

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**Target species:**

Horse

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**Route of administration:**

Cutaneous use

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## 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)

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**Pharmaceutical form:**

Gel

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**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.

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**Anatomical therapeutic chemical veterinary (ATCvet) codes:**

QM02AX99

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**Legal status of supply:**

Veterinary medicinal product subject to veterinary prescription

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**Authorisation status:**

Valid

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**Authorised in:**

Finland

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**Available in:**

Finland

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**Package description:**

Box of one 125 ml Brown Type III glass bottle with a dosing pump made of high-density polyethylene / polypropylene and a dip tube made of low-density polyethylene and polypropylene. Polypropylene screw-fit cap.

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## Additional information

**Entitlement type:**

Marketing Authorisation

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**Legal basis of product authorisation:**

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

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**Marketing authorisation holder:**

Audevard

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**Marketing authorisation date:**

10/06/2020

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**Manufacturing sites for batch release:**

Dopharma France

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**Responsible authority:**

Finnish Medicines Agency

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**Authorisation number:**

36229

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**Date of authorisation status change:**

10/06/2020

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**Reference member state:**

France

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**Procedure number:**

FR/V/0344/001

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**Concerned member states:**

Austria Denmark Finland Germany Ireland Luxembourg Netherlands Norway  
Poland Portugal Sweden

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To consult adverse reactions on veterinary medicinal products please go to  
[www.adrreports.eu/vet](http://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

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