PREDNISOLONA FATRO

Authorised

- Prednisolone sodium phosphate

Product identification

Medicine name:
PREDNISOLONA FATRO

Active substance:
Prednisolone sodium phosphate

Target species:
Dog
Cat

Route of administration:
Intraarticular use
Intramuscular use

Product details

Active substance and strength:
Prednisolone sodium phosphate
33.75 milligram(s) / 1.00 millilitre(s)

Pharmaceutical form:
Solution for injection

Withdrawal period by route of administration:

Intraarticular use:

- **Dog**

Intramuscular use:

- **Dog**
- **Cat**

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QH02AB06

Legal status of supply:
Veterinary medicinal product subject to veterinary prescription

Authorisation status:
Valid

Authorised in:
Spain

Package description:
Available only in Spanish
Available only in Spanish

Additional information

Entitlement type:
Marketing Authorisation

Legal basis of product authorisation:
Full application - Known active substance (Article 12(3) of Directive No 2001/82/EC)

Marketing authorisation holder:
Fatro Iberica S.L.

Marketing authorisation date:
15/07/1992

Manufacturing sites for batch release:
Fatro S.p.A.

Responsible authority:
Spanish Agency For Medicines And Health Products

Authorisation number:
475 ESP

Date of authorisation status change:
1/01/2019

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

Documents

Labelling

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Package Leaflet

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Summary of Product Characteristics

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