Dexalin

Authorised

- Dexamethasone sodium phosphate

Product identification

Medicine name:
Dexalin

Active substance:
Dexamethasone sodium phosphate

Target species:
Dog
Cat

Route of administration:
Intraarticular use
Intramuscular use
Intravenous use

Product details

Active substance and strength:
Dexamethasone sodium phosphate
2.00 milligram(s) / 1.00 millilitre(s)

Pharmaceutical form:
Solution for injection

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
Available only in Dutch

Additional information

Entitlement type:
Marketing Authorisation

Legal basis of product authorisation:
Legal basis not covered by Directive 2001/82/EC

Marketing authorisation holder:
Kepro B.V.

Marketing authorisation date:
12/11/1993

Manufacturing sites for batch release:
Kepro B.V.

Responsible authority:
Medicines Evaluation Board

Authorisation number:
REG NL 2483

Date of authorisation status change:
2/12/2016

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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