Depo-Medrone V 40 mg/ml Suspension for Injection

Authorised

- Methylprednisolone acetate

Product identification

Medicine name:
Depo-Medrone V 40 mg/ml Suspension for Injection

Active substance:
Methylprednisolone acetate

Target species:
Dog
Horse (non food-producing)
Cat

Route of administration:
Intraarticular use
Intramuscular use

Product details

Active substance and strength:
Methylprednisolone acetate
40.00 milligram(s) / 1.00 millilitre(s)

Pharmaceutical form:
Suspension for injection

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QH02AB04

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

Authorisation status:
Valid

Authorised in:
Ireland

Available in:
Ireland

Package description:
5 ml Type I glass vial with butyl rubber bung and aluminium overseal containing a white sterile aqueous suspension for injection.

Additional information

Entitlement type:
Marketing Authorisation

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

Marketing authorisation holder:
Zoetis Belgium S.A.

Marketing authorisation date:
6/01/2014

Manufacturing sites for batch release:
Pfizer Manufacturing Belgium

Responsible authority:

Health Products Regulatory Authority

Authorisation number:
VPA10387/021/001

Date of authorisation status change:
6/01/2014

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

Documents

Summary of Product Characteristics