

Medrate Solubile ad us.vet. 125 mg Pulver und Lösungsmittel zur Herstellung einer Injektionslösung für Hunde

Authorised

- Methylprednisolone sodium succinate

Product identification

Medicine name:

Medrate Solubile ad us.vet. 125 mg Pulver und Lösungsmittel zur Herstellung einer Injektionslösung für Hunde

Active substance:

Methylprednisolone sodium succinate

Target species:

Dog

Route of administration:

Intramuscular use

Intravenous use

Product details

Active substance and strength:

Methylprednisolone sodium succinate
82.88 milligram(s) / 1.00 millilitre(s)

Pharmaceutical form:

Powder and solvent for solution 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:

Germany

Available in:

Germany

Package description:

Available only in German

Additional information

Entitlement type:

Marketing Authorisation

Legal basis of product authorisation:

Legal basis not covered by Directive 2001/82/EC

Marketing authorisation holder:

Zoetis Deutschland GmbH

Marketing authorisation date:

2/08/1999

Manufacturing sites for batch release:

Pfizer Manufacturing Belgium

Responsible authority:

Federal Office Of Consumer Protection And Food Safety

Authorisation number:

400128.00.00

Date of authorisation status change:

28/11/2005

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

Documents

Package Leaflet

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

Summary of Product Characteristics

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