File downloaded on 2026-11-20
Source URL: https://medicines.health.europa.eu/veterinary/en/600000097476

VITATOX emulsione iniettabile per cani e gatti

Authorised

- Glucose
- Sorbitol
- Nicotinamide
- Ascorbic acid
- Thiamine
- Cholecalciferol
- Retinol

Product identification

Medicine name:
VITATOX emulsione iniettabile per cani e gatti

Active substance:
Glucose

Sorbitol

Nicotinamide

Ascorbic acid

Thiamine

Cholecalciferol

Retinol

Target species:
Dog

Cat

Route of administration:
Subcutaneous use
Intramuscular use
Intravascular use
Intraperitoneal use

Product details

Active substance and strength:
Glucose
50.00 milligram(s) / 1.00 millilitre(s)

Sorbitol
50.00 milligram(s) / 1.00 millilitre(s)

Nicotinamide
1.20 milligram(s) / 1.00 millilitre(s)

Ascorbic acid
8.00 milligram(s) / 1.00 millilitre(s)

Thiamine
0.36 milligram(s) / 1.00 millilitre(s)

Cholecalciferol
2000.00 international unit(s) / 1.00 millilitre(s)

Retinol
400.00 international unit(s) / 1.00 millilitre(s)

Pharmaceutical form:
Emulsion for injection

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QA11AB

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

Authorisation status:
Valid

Authorised in:
Italy

Package description:
Available only in Italian

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:
Fatro S.p.A.

Marketing authorisation date:
27/02/1968

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

Responsible authority:
Ministry Of Health

Authorisation number:
101799

Date of authorisation status change:
1/01/2009

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

Documents

Combined File of all Documents

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