FLATULEX (17+1,67)MG/ML ΕΝΕΣΙΜΟ ΔΙΑΛΥΜΑ

Authorised

- TOCOPHERYL ACETATE
- SODIUM SELENITE ANHYDROUS

Product identification

Medicine name:
FLATULEX (17+1,67)MG/ML ΕΝΕΣΙΜΟ ΔΙΑΛΥΜΑ

Active substance:
TOCOPHERYL ACETATE
SODIUM SELENITE ANHYDROUS

Target species:
Pig
Horse
Cattle
Sheep
Goat

Route of administration:
Intramuscular use

Product details

Active substance and strength:

TOCOPHERYL ACETATE
17.00 milligram(s) / 1.00 millilitre(s)

SODIUM SELENITE ANHYDROUS
1.67 milligram(s) / 1.00 millilitre(s)

Pharmaceutical form:
Solution for injection

Withdrawal period by route of administration:
Intramuscular use:

- **Pig**
  - Meat and offal. 28 day
- **Horse**
  - Meat and offal. 28 day
  - Milk. 0 day
- **Cattle**
  - Meat and offal. 28 day
  - Milk. 0 day
- **Sheep**
  - Meat and offal. 28 day
  - Milk. 0 day
- **Goat**
  - Meat and offal. 28 day
  - Milk. 0 day

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QA12CE99

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

Authorisation status:
Valid

Authorised in:
Greece

Available in:
Greece

Package description:
Available only in Greek
Available only in Greek

Additional information

Entitlement type:
Marketing Authorisation

Legal basis of product authorisation:
Complete application (stand-alone)

Marketing authorisation holder:
PROVET S.A.

Marketing authorisation date:
9/09/1996

Manufacturing sites for batch release:
PROVET S.A.

Responsible authority:
National Organization For Medicines

Authorisation number:
31805/09-04-2021/K-0018301

Date of authorisation status change:
10/10/2021

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