ECTOPOR 2%W/V ΔΕΡΜΑΤΙΚΟ ΔΙΑΛΥΜΑ

Authorised

- Cypermethrin

Product identification

Medicine name:
ECTOPOR 2%W/V ΔΕΡΜΑΤΙΚΟ ΔΙΑΛΥΜΑ

Active substance:
Cypermethrin

Target species:
Goat
Cattle
Sheep

Route of administration:
Cutaneous use

Product details

Active substance and strength:
Cypermethrin
2.00 gram(s) / 100.00 millilitre(s)

Pharmaceutical form:

Cutaneous solution

Withdrawal period by route of administration:

Cutaneous use:

- **Goat**
  - Meat and offal. 4 day
  - Milk. 96 hour
- **Cattle**
  - Meat and offal. 4 day
  - Milk. 48 hour
- **Sheep**
  - Meat and offal. 4 day
  - Milk. 96 hour

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QP53AC08

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

Authorisation status:
Valid

Authorised in:
Greece

Package description:
Available only in Greek

Additional information

Entitlement type:

Marketing Authorisation

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

Marketing authorisation holder:
Premier Shukuroglou Hellas S.A. Emporias Ktiniatrikon Farmakon

Marketing authorisation date:
26/02/1990

Manufacturing sites for batch release:
Safapac Limited

Responsible authority:
National Organization For Medicines

Authorisation number:
17438/11-03-2013/K-0032601

Date of authorisation status change:
17/06/2024

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

Documents

Summary of Product Characteristics

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