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ALAMYCIN 32,1 mg/mL δερματικό εκνέφωμα, διάλυμα για πρόβατα

Authorised

- Oxytetracycline hydrochloride

Product identification

Medicine name:
ALAMYCIN 32,1 mg/mL δερματικό εκνέφωμα, διάλυμα για πρόβατα

Active substance:
Oxytetracycline hydrochloride

Target species:
Sheep

Route of administration:
Topical use

Product details

Active substance and strength:
Oxytetracycline hydrochloride
35.80 milligram(s) / 1.00 gram(s)

Pharmaceutical form:
Cutaneous spray, solution

Withdrawal period by route of administration:

Topical use:

- **Sheep**
  - Meat and offal. 0 day
  - Milk. 0 hour

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QD06AA03

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:
Chellafarm Vet A.E.

Marketing authorisation date:
10/12/1996

Manufacturing sites for batch release:

Norbrook Laboratories (Ireland) Limited

Responsible authority:
National Organization For Medicines

Authorisation number:
75237/10-08-2021/K-0053002

Date of authorisation status change:
29/12/2021

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