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ENTEROSTREP SP

Authorised

- Dihydrostreptomycin sulfate

Product identification

Medicine name:
ENTEROSTREP SP

Active substance:
Dihydrostreptomycin sulfate

Target species:
Young rabbit

Route of administration:
In drinking water use

Product details

Active substance and strength:
Dihydrostreptomycin sulfate
625.90 milligram(s) / 1.00 gram(s)

Pharmaceutical form:
Powder for use in drinking water

Withdrawal period by route of administration:

In drinking water use:

- **Young rabbit**
  - Meat and offal. 8 day

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QJ01GA90

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

Authorisation status:
Valid

Authorised in:
Spain

Available in:
Spain

Package description:
bag of 500 g

Additional information

Entitlement type:
Marketing Authorisation

Legal basis of product authorisation:
Full application - Known active substance (Article 12(3) of Directive No 2001/82/EC)

Marketing authorisation holder:
S P Veterinaria S.A.

Marketing authorisation date:
30/07/2013

Manufacturing sites for batch release:

S P Veterinaria S.A.

Responsible authority:
Spanish Agency Of Medicines And Medical Devices

Authorisation number:
2859 ESP

Date of authorisation status change:
30/07/2013

Reference member state:
Spain

Procedure number:
ES/V/0285/001

Concerned member states:
Portugal

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

Documents

Summary of Product Characteristics

English (PDF)
Published on: 20/04/2026
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Package Leaflet

English (PDF)

Published on: 27/03/2026
Download

Labelling

English (PDF)
Published on: 27/03/2026
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Combined File of all Documents

es-puar-enterostrep-sp-es.pdf

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