File downloaded on 2026-11-09
Source URL: https://medicines.health.europa.eu/veterinary/en/600000993725

Lexylan 180 mg/ml - Suspension for injection

Authorised

- Cefalexin

Product identification

Medicine name:
Lexylan 180 mg/ml - Suspension for injection

Active substance:
Cefalexin

Target species:
Cattle
Dog
Cat

Route of administration:
Intramuscular use
Subcutaneous use

Product details

Active substance and strength:
Cefalexin
180.00 milligram(s) / 1.00 millilitre(s)

Pharmaceutical form:

Suspension for injection

Withdrawal period by route of administration:

Intramuscular use:

- **Cattle**
  - Meat and offal. 12 day 12 Days
  - Milk. 0 hour Zero Hours

Subcutaneous use:

- **Cattle**
  - Meat and offal. 12 day 12 Days
  - Milk. 0 hour Zero Hours

Anatomical therapeutic chemical veterinary (ATCvet) codes:

QJ01DB01

Legal status of supply:

Veterinary medicinal product subject to veterinary prescription

---

Authorisation status:

Valid

Authorised in:

Austria , Belgium , Bulgaria , Croatia , Cyprus , Czechia , Denmark , Estonia , Finland , France , Germany , Greece , Hungary , Iceland , Ireland , Italy , Latvia , Liechtenstein , Lithuania , Luxembourg , Malta , Netherlands , Norway , Poland , Portugal , Romania , Slovakia , Slovenia , Spain , Sweden , United Kingdom (Northern Ireland)

Available in:

Austria , Belgium , Denmark , France , Germany , Ireland , Italy , Luxembourg , Netherlands , Spain

---

Package description:

Packaging:Glass Type II, Package_size:1 vial, Content:250 ml
Packaging:Glass type II, Package_size:1 vial, Content:100 ml

Additional information

Entitlement type:
Marketing Authorisation

Legal basis of product authorisation:
Generic application (Article 18 of Regulation (EU) 2019/6)

Marketing authorisation holder:
Emdoka

Marketing authorisation date:
8/04/2024

Manufacturing sites for batch release:
Wirtschaftsgenossenschaft deutscher Tieraerzte eG

Responsible authority:
European Commission

Authorisation number:
This information is not available for this product.

Date of authorisation status change:
8/04/2024

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

Documents

Combined File of all Documents

English (PDF)
Published on: 13/03/2026
Download

ema-puar-v6103-lexylan-en.pdf