Novaderma 660 mg/g + 7.7 mg/g cutaneous paste for horses, cattle and sheep

Authorised

- Methyl salicylate
- Salicylic acid

Product identification

Medicine name:
Novaderma 660 mg/g + 7.7 mg/g cutaneous paste for horses, cattle and sheep

Active substance:
Methyl salicylate

Salicylic acid

Target species:
Cattle
Sheep
Horse

Route of administration:
Cutaneous use

Product details

Active substance and strength:

Methyl salicylate
7.70 milligram(s) / 1.00 gram(s)

Salicylic acid
660.00 milligram(s) / 1.00 gram(s)

Pharmaceutical form:
Cutaneous paste

Withdrawal period by route of administration:
Cutaneous use:

- **Cattle**
  - Milk. 24 hour
  - Meat and offal. 1 day
- **Sheep**
  - Milk. 24 hour
  - Meat and offal. 1 day
- **Horse**
  - Milk. 24 hour
  - Meat and offal. 1 day

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QD02AF

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

Authorisation status:
Valid

Authorised in:
France

Package description:

(ID1) 500 gram(s): Box (cardboard) with 1 Jar (polypropylene) with 500 gram(s), closed with Schraubdeckel (polypropylene)

Additional information

Entitlement type:
Marketing Authorisation

Legal basis of product authorisation:
Hybrid application – bioavailability studies cannot be used to demonstrate bioequivalence (Article 19(1)(b) of Regulation (EU) 2019/6)

Marketing authorisation holder:
Wirtschaftsgenossenschaft deutscher Tieraerzte eG

Marketing authorisation date:
16/04/2026

Manufacturing sites for batch release:
Wirtschaftsgenossenschaft deutscher Tieraerzte eG

Responsible authority:
French Agency For Food, Environmental And Occupational Health & Safety

Authorisation number:
FR/V/6781929 8/2026

Date of authorisation status change:
16/04/2026

Reference member state:
Germany

Procedure number:
DE/V/0349/001

Concerned member states:
Austria Belgium Czechia France Hungary Ireland Italy Luxembourg Netherlands Poland Portugal United Kingdom (Northern Ireland)

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

Documents

Combined File of all Documents

Summary of Product Characteristics

This document does not exist in this language (English). You can find it in another language below.

Package Leaflet and Labelling

This document does not exist in this language (English). You can find it in another language below.