

SODIUM SALICYL 80% WSP, powder for oral solution for cattle (calves) and pig

Authorised

- Sodium salicylate

Product identification

Medicine name:

SODIUM SALICYL 80% WSP, powder for oral solution for cattle (calves) and pig

Active substance:

Sodium salicylate

Target species:

Cattle (calf)

Pig

Route of administration:

In drinking water/milk use

Product details

Active substance and strength:

Sodium salicylate

800.00 milligram(s) / 1.00 gram(s)

Pharmaceutical form:

Oral powder

Withdrawal period by route of administration:**In drinking water/milk use:**

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Cattle (calf)

- Meat and offal. no withdrawal period zero days

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Pig

- Meat and offal. no withdrawal period zero days

Anatomical therapeutic chemical veterinary (ATCvet) codes:

QN02BA04

Legal status of supply:

Veterinary medicinal product subject to veterinary prescription

Authorisation status:

Valid

Authorised in:

France

Available in:

France

Package description:

1 kg composite can. Composite can: container consisting of PET/aluminium/adhesive/paper, with a PET/aluminium tear-off membrane and a HDPE lid.

1 kg bucket. Bucket: polypropylene bucket provided with a polypropylene lid.

5 kg bucket. Bucket: polypropylene bucket provided with a polypropylene lid.

2,5 kg bucket. Bucket: polypropylene bucket provided with a polypropylene lid.

Securitainer: white polypropylene cylindrical container provided with a low-density polyethylene lid. The securitainer contains 1 kg of product.

Additional information

Entitlement type:

Marketing Authorisation

Legal basis of product authorisation:

Generic application (Article 13(1) of Directive No 2001/82/EC)

Marketing authorisation holder:

Dopharma Research B.V.

Marketing authorisation date:

20/10/2009

Manufacturing sites for batch release:

Dopharma B.V.

Responsible authority:

French Agency For Food, Environmental And Occupational Health & Safety

Authorisation number:

FR/V/9392306 8/2009

Date of authorisation status change:

20/10/2014

Reference member state:

Netherlands

Procedure number:

NL/V/0133/001

Concerned member states:

Belgium Bulgaria Denmark Estonia France Germany Greece Hungary
Ireland Italy Latvia Lithuania Poland Portugal Romania Spain
United Kingdom (Northern Ireland)

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

Documents

Summary of Product Characteristics

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

Combined File of all Documents

NL V 133 001 DC Sodium salicyl 80 WSP PuAR_updated 082021.pdf