Sodium Salicyl 80% WSP powder for use in drinking water/milk for cattle (calves) and pigs

Authorised

- Sodium salicylate

Product identification

Medicine name:
Sodium Salicyl 80% WSP powder for use in drinking water/milk for cattle (calves) and pigs

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:
Powder for use in drinking water/milk

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

- **Cattle (calf)**
  - Meat and offal. 0 day

- **Pig**
  - Meat and offal. 0 day

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QN02BA04

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

Authorisation status:
Valid

Authorised in:
Netherlands

Available in:
Netherlands

Package description:
Securitainer: white polypropylene cylindrical container provided with a low-density polyethylene lid. The securitainer contains 1 kg of product.
2,5 kg bucket. Bucket: polypropylene bucket provided with a polypropylene lid.
5 kg bucket. Bucket: polypropylene bucket provided with a polypropylene lid.
1 kg bucket. Bucket: polypropylene bucket provided with a polypropylene lid.
1 kg composite can. Composite can: container consisting of PET/aluminium/adhesive/paper, with a PET/aluminium tear-off membrane and a HDPE lid.

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:
12/10/2009

Manufacturing sites for batch release:
Dopharma B.V.

Responsible authority:
Medicines Evaluation Board

Authorisation number:
REG NL 104033

Date of authorisation status change:
12/10/2009

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

Combined File of all Documents

English (PDF)
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