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Kenostart Spray en Dip 3 mg/g Cutaneous spray, solution@Dip solution

Authorised

- Iodine

Product identification

Medicine name:

Kenostart Spray en Dip 3 mg/g Cutaneous spray, solution@Dip solution

Active substance:

Iodine

Target species:

Cattle

Route of administration:

Dipping

Nebulisation use

Product details

Active substance and strength:

Iodine

3.00 milligram(s) / 1.00 gram(s)

Pharmaceutical form:

Teat dip/spray solution

Withdrawal period by route of administration:**Dipping:**

-

Cattle

- Meat and offal. 0 day
- Milk. 0 day

Nebulisation use:

-

Cattle

- Meat and offal. 0 day
 - Milk. 0 day
-

Anatomical therapeutic chemical veterinary (ATCvet) codes:

QD08AG03

Legal status of supply:

Veterinary medicinal product subject to veterinary prescription except for some pack sizes

Authorisation status:

Valid

Authorised in:

France

Available in:

France

Package description:

Kenostart Spray en Dip 3 mg/g dip sol./cut. spray sol. 200 l
Kenostart Spray en Dip 3 mg/g dip sol./cut. spray sol. 60 l
Kenostart Spray en Dip 3 mg/g dip sol./cut. spray sol. 25 l
Kenostart Spray en Dip 3 mg/g dip sol./cut. spray sol. 20 l

Kenostart Spray en Dip 3 mg/g dip sol./cut. spray sol. 10 l
Kenostart Spray en Dip 3 mg/g dip sol./cut. spray sol. 5 l
Kenostart Spray en Dip 3 mg/g dip sol./cut. spray sol. 1 l

Additional information

Entitlement type:

Marketing Authorisation

Legal basis of product authorisation:

Full application (Article 12(3) of Directive No 2001/82/EC)

Marketing authorisation holder:

Cid Lines

Marketing authorisation date:

18/10/2007

Manufacturing sites for batch release:

Cid Lines

Responsible authority:

French Agency For Food, Environmental And Occupational Health & Safety

Authorisation number:

FR/V/6708751 6/2007

Date of authorisation status change:

18/10/2012

Reference member state:

Belgium

Procedure number:

BE/V/0042/001

Concerned member states:

Austria Cyprus Estonia France Germany Greece Ireland Italy Netherlands
Poland Portugal 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

English (PDF)

Published on: 13/03/2026

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Labelling