

AquaVac Relera Concentrate for Dip Suspension or Suspension for Injection for Rainbow Trout

Authorised

- *Yersinia ruckeri*, serotype O1, biotype 1, strain Hagerman, Inactivated
- *Yersinia ruckeri*, serotype O1, biotype 2 (EX5), strain SP/07/04, Inactivated

Product identification

Medicine name:

AquaVac Relera Concentrate for Dip Suspension or Suspension for Injection for Rainbow Trout

Active substance:

Yersinia ruckeri, serotype O1, biotype 1, strain Hagerman, Inactivated

Yersinia ruckeri, serotype O1, biotype 2 (EX5), strain SP/07/04, Inactivated

Target species:

Trout - Golden/Rainbow/Redband/Steelhead

Route of administration:

Dipping

Intraperitoneal use

Product details

Active substance and strength:

Yersinia ruckeri, serotype O1, biotype 1, strain Hagerman, Inactivated

75.00 Relative Percentage Survival / 1.00 millilitre(s)

Yersinia ruckeri, serotype O1, biotype 2 (EX5), strain SP/07/04, Inactivated

75.00 Relative Percentage Survival / 1.00 millilitre(s)

Pharmaceutical form:

Concentrate for dip suspension

Withdrawal period by route of administration:

Dipping:

-

Trout - Golden/Rainbow/Redband/Steelhead

- Meat. 0 degree day

Intraperitoneal use:

-

Trout - Golden/Rainbow/Redband/Steelhead

- Meat. 0 degree day

Anatomical therapeutic chemical veterinary (ATCvet) codes:

QI10BB03

Legal status of supply:

Veterinary medicinal product subject to veterinary prescription

Authorisation status:

Valid

Authorised in:

Spain

Package description:

bottle of 1 L

Additional information

Entitlement type:

Marketing Authorisation

Legal basis of product authorisation:

Full application - Known active substance (Article 12(3) of Directive No 2001/82/EC)

Marketing authorisation holder:

Merck Sharp & Dohme Animal Health S.L.

Marketing authorisation date:

27/07/2009

Manufacturing sites for batch release:

MSD Animal Health UK Limited

Merck Sharp & Dohme Animal Health S.L.

The Veterinary Medicines Directorate

Responsible authority:

Spanish Agency Of Medicines And Medical Devices

Authorisation number:

2054 ESP

Date of authorisation status change:

1/01/2023

Reference member state:

Spain

Procedure number:

ES/V/0309/001

Concerned member states:

Czechia Denmark Finland France Germany Italy Norway Portugal Slovakia

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: 5/04/2023

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Labelling

English (PDF)

Published on: 21/10/2024

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Package Leaflet

English (PDF)

Published on: 21/10/2024

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