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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

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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: 14/03/2026
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Labelling

English (PDF)
Published on: 14/03/2026
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Package Leaflet

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