File downloaded on 2026-11-18
Source URL: https://medicines.health.europa.eu/veterinary/en/600000014299

ALPHA JECT micro 6 emulsion for injection for atlantic salmon

Authorised

- Aeromonas salmonicida, strain AL2017, Inactivated
- Vibrio anguillarum, serotype O1, strain AL 112, Inactivated
- Vibrio anguillarum, serotype O2a, strain AL 104, Inactivated
- Aliivibrio salmonicida, strain AL 1134, Inactivated
- Moritella viscosa, strain AL 266, Inactivated
- Infectious pancreatic necrosis virus, serotype Sp, strain AL V103, Inactivated

Product identification

Medicine name:
ALPHA JECT micro 6 emulsion for injection for atlantic salmon

Active substance:
Aeromonas salmonicida, strain AL2017, Inactivated

Vibrio anguillarum, serotype O1, strain AL 112, Inactivated

Vibrio anguillarum, serotype O2a, strain AL 104, Inactivated

Aliivibrio salmonicida, strain AL 1134, Inactivated

Moritella viscosa, strain AL 266, Inactivated

Infectious pancreatic necrosis virus, serotype Sp, strain AL V103, Inactivated

Target species:
Atlantic salmon

Route of administration:
Intraperitoneal use

Product details

Active substance and strength:
Aeromonas salmonicida, strain AL2017, Inactivated
70.00 Relative Percentage Survival / 0.05 millilitre(s)

Vibrio anguillarum, serotype O1, strain AL 112, Inactivated
75.00 Relative Percentage Survival / 0.05 millilitre(s)

Vibrio anguillarum, serotype O2a, strain AL 104, Inactivated
75.00 Relative Percentage Survival / 0.05 millilitre(s)

Aliivibrio salmonicida, strain AL 1134, Inactivated
90.00 Relative Percentage Survival / 0.05 millilitre(s)

Moritella viscosa, strain AL 266, Inactivated
60.00 Relative Percentage Survival / 0.05 millilitre(s)

Infectious pancreatic necrosis virus, serotype Sp, strain AL V103, Inactivated
0.28 antigen unit(s) / 0.05 millilitre(s)

Pharmaceutical form:
Emulsion for injection

Withdrawal period by route of administration:
Intraperitoneal use:

- **Atlantic salmon**
  - Meat. 0 degree day

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QI10AL02

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

Authorisation status:

Valid

Authorised in:
Ireland

Available in:
Ireland

Package description:
250 ml bag
500 ml bag
10x500 ml bag

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:
Pharmaq AS

Marketing authorisation date:
15/03/2019

Manufacturing sites for batch release:
Pharmaq AS

Responsible authority:
Health Products Regulatory Authority

Authorisation number:
VPA10804/005/001

Date of authorisation status change:
15/03/2019

Reference member state:

Norway

Procedure number:
NO/V/0008/001

Concerned member states:
Ireland 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

Package Leaflet

Labelling

Summary of Product Characteristics