ALPHA JECT 3000 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

Product identification

Medicine name:
ALPHA JECT 3000 injeksjonsvæske, emulsjon til atlantisk laks
ALPHA JECT 3000 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

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 / 1.00 Dose

Vibrio anguillarum, serotype O1, strain AL 112, Inactivated
75.00 Relative Percentage Survival / 1.00 Dose

Vibrio anguillarum, serotype O2a, strain AL 104, Inactivated
75.00 Relative Percentage Survival / 1.00 Dose

Pharmaceutical form:
Emulsion for injection

Withdrawal period by route of administration:
Intraperitoneal use:

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

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QI10AB02

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

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Authorisation status:
Valid

Authorised in:
Norway

Available in:
Norway

Package description:
UVO injection bags made of a multilayer plastic foil with ethylene vinyl acetate as the product contact layer. Thegiving port is closed with a bromobutyl based rubber stopper.Pack size: 500 ml

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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:
24/11/2008

Manufacturing sites for batch release:
Pharmaq AS

Responsible authority:
Norwegian Medical Products Agency

Authorisation number:
07-5472

Date of authorisation status change:
1/10/2012

Reference member state:
Ireland

Procedure number:
IE/V/0219/001

Concerned member states:
Denmark Finland Iceland Norway

To consult adverse reactions on veterinary medicinal products please go to www.adrreports.eu/vet

Documents

Summary of Product Characteristics

English (PDF)
Published on: 4/05/2025
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Combined File of all Documents

Package Leaflet

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