

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 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

Authorisation status:

Valid

Authorised in:

Denmark

Available in:

Denmark

Package description:

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

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:

7/07/2011

Manufacturing sites for batch release:

Pharmaq AS

Responsible authority:

Danish Medicines Agency

Authorisation number:

47944

Date of authorisation status change:

7/07/2011

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

Updated on: 13/03/2026

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Combined File of all Documents