

# 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

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**Active substance:**

*Aeromonas salmonicida*, strain AL2017, Inactivated  
*Vibrio anguillarum*, serotype O1, strain AL 112, Inactivated  
*Vibrio anguillarum*, serotype O2a, strain AL 104, Inactivated

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**Target species:**

Atlantic salmon

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**Route of administration:**

Intraperitoneal use

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## 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

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**Pharmaceutical form:**

Emulsion for injection

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**Withdrawal period by route of administration:**

**Intraperitoneal use:**

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**Atlantic salmon**

- Meat and offal. 0 degree day

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**Anatomical therapeutic chemical veterinary (ATCvet) codes:**

QI10AB02

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**Legal status of supply:**

Veterinary medicinal product subject to veterinary prescription

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**Authorisation status:**

Valid

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**Authorised in:**

Norway

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**Available in:**

Norway

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**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

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## Additional information

**Entitlement type:**

Marketing Authorisation

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**Legal basis of product authorisation:**

Full application (Article 12(3) of Directive No 2001/82/EC)

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**Marketing authorisation holder:**

Pharmaq AS

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**Marketing authorisation date:**

24/11/2008

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**Manufacturing sites for batch release:**

Pharmaq AS

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**Responsible authority:**

Norwegian Medical Products Agency

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**Authorisation number:**

07-5472

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**Date of authorisation status change:**

1/10/2012

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**Reference member state:**

Ireland

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**Procedure number:**

IE/V/0219/001

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**Concerned member states:**

Denmark Finland Iceland Norway

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To consult adverse reactions on veterinary medicinal products please go to [www.adrreports.eu/vet](http://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

This document does not exist in this language (English). You can find it in another language below.