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# ALPHA JECT micro 7 ILA i

Authorised

- Infectious salmon anaemia virus, Inactivated
- Infectious pancreatic necrosis virus, serotype Sp, Inactivated
- Moritella viscosa, Inactivated
- Aliivibrio salmonicida, Inactivated
- Vibrio anguillarum, serotype O2a, Inactivated
- Vibrio anguillarum, serotype O1, Inactivated
- Aeromonas salmonicida, subsp. salmonicida, Inactivated

## Product identification

**Medicine name:**
ALPHA JECT micro 7 ILA i

**Active substance:**
Infectious salmon anaemia virus, Inactivated

Infectious pancreatic necrosis virus, serotype Sp, Inactivated

Moritella viscosa, Inactivated

Aliivibrio salmonicida, Inactivated

Vibrio anguillarum, serotype O2a, Inactivated

Vibrio anguillarum, serotype O1, Inactivated

Aeromonas salmonicida, subsp. salmonicida, Inactivated

**Target species:**
Atlantic salmon

**Route of administration:**
Intraperitoneal use

## Product details

**Active substance and strength:**
Infectious salmon anaemia virus, Inactivated
60.00 Relative Percentage Survival / 0.05 millilitre(s)

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

Moritella viscosa, Inactivated
10.70 log2 enzyme-linked immunosorbent assay unit(s) / 0.05 millilitre(s)

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

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

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

Aeromonas salmonicida, subsp. salmonicida, Inactivated
12.60 log2 enzyme-linked immunosorbent assay unit(s) / 0.05 millilitre(s)

**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:**
QI10AL04

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

**Authorisation status:**
Valid

**Authorised in:**
Norway

**Available in:**
Norway

**Package description:**
Available only in Norwegian
Available only in Norwegian
Available only in Norwegian

## 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:**
12/02/2019

**Manufacturing sites for batch release:**
Pharmaq AS

**Responsible authority:**
Norwegian Medical Products Agency

**Authorisation number:**
17-11914

**Date of authorisation status change:**
12/02/2019

**Reference member state:**
Norway

**Procedure number:**
NO/V/0021/001

**Concerned member states:**
Iceland

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

## Documents

Summary of Product Characteristics

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

Package Leaflet

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

Labelling

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