

# Aquavac PD7 vet. injeksjonsvæske, emulsjon

Not  
authorised

- *Moritella viscosa*, Inactivated
- *Vibrio anguillarum*, serotype O2a, Inactivated
- *Vibrio anguillarum*, serotype O1, Inactivated
- *Aliivibrio salmonicida*, Inactivated
- *Aeromonas salmonicida*, subsp. *salmonicida*, Inactivated
- Infectious pancreatic necrosis virus, serotype Sp, Inactivated
- Salmon pancreas disease virus, strain F93-125, Inactivated

## Product identification

### **Medicine name:**

Aquavac PD7 vet. injeksjonsvæske, emulsjon

### **Active substance:**

*Moritella viscosa*, Inactivated

*Vibrio anguillarum*, serotype O2a, Inactivated

*Vibrio anguillarum*, serotype O1, Inactivated

*Aliivibrio salmonicida*, Inactivated

*Aeromonas salmonicida*, subsp. *salmonicida*, Inactivated

Infectious pancreatic necrosis virus, serotype Sp, Inactivated

Salmon pancreas disease virus, strain F93-125, Inactivated

**Target species:**

Atlantic salmon

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**Route of administration:**

Intraperitoneal use

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## Product details

**Active substance and strength:**

Moritella viscosa, Inactivated

5.80 log<sub>2</sub> enzyme-linked immunosorbent assay unit(s) / 0.10 millilitre(s)

Vibrio anguillarum, serotype O2a, Inactivated

75.00 Relative Percentage Survival / 0.10 millilitre(s)

Vibrio anguillarum, serotype O1, Inactivated

75.00 Relative Percentage Survival / 0.10 millilitre(s)

Aliivibrio salmonicida, Inactivated

75.00 Relative Percentage Survival / 0.10 millilitre(s)

Aeromonas salmonicida, subsp. salmonicida, Inactivated

10.70 log<sub>2</sub> enzyme-linked immunosorbent assay unit(s) / 0.10 millilitre(s)

Infectious pancreatic necrosis virus, serotype Sp, Inactivated

1.50 enzyme-linked immunosorbent assay unit / 0.10 millilitre(s)

Salmon pancreas disease virus, strain F93-125, Inactivated

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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:**

QI10AL05

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**Legal status of supply:**

Veterinary medicinal product subject to veterinary prescription

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**Authorisation status:**

Surrendered

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**Authorised in:**

Norway

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**Package description:**

Available only in Norwegian

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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:**

Intervet International B.V.

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**Marketing authorisation date:**

3/02/2015

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**Manufacturing sites for batch release:**

Intervet International B.V.

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**Responsible authority:**

Norwegian Medical Products Agency

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**Authorisation number:**

13-9717

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**Date of authorisation status change:**

3/06/2025

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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

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.