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

Route of administration:
Intraperitoneal use

Product details

Active substance and strength:
Moritella viscosa, Inactivated
5.80 log2 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 log2 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

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:
QI10AL05

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

Authorisation status:
Surrendered

Authorised in:
Norway

Package description:
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:
Intervet International B.V.

Marketing authorisation date:
3/02/2015

Manufacturing sites for batch release:
Intervet International B.V.

Responsible authority:
Norwegian Medical Products Agency

Authorisation number:
13-9717

Date of authorisation status change:
3/06/2025

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.