Aquavac 6 Emulsion for Injection for Atlantic Salmon

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, Inactivated

Product identification

Medicine name:
Aquavac 6 Emulsion for Injection for Atlantic Salmon

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, Inactivated

Target species:
Atlantic salmon

Route of administration:
Intraperitoneal use

Product details

Active substance and strength:

Moritella viscosa, Inactivated
6.50 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
90.00 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, Inactivated
1.50 enzyme-linked immunosorbent assay unit / 0.10 millilitre(s)

Pharmaceutical form:

Emulsion for injection

Withdrawal period by route of administration:

Intraperitoneal use:

- **Atlantic salmon**
  - All relevant tissues. 0 degree day

Anatomical therapeutic chemical veterinary (ATCvet) codes:

QI10AL02

Legal status of supply:

Veterinary medicinal product subject to veterinary prescription

---

Authorisation status:

Surrendered

Authorised in:

United Kingdom (Northern Ireland)

Package description:
Bottles of polyethylene terephthalate (PET) closed with a rubber stopper and aluminium cap.

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:
24/05/2022

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

Responsible authority:
The Veterinary Medicines Directorate

Authorisation number:
Vm 06376/3021

Date of authorisation status change:
29/10/2025

Reference member state:
Norway

Procedure number:
NO/V/0019/001

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