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Pentium forte plus vet. injeksjonsvæske, emulsjon, vaksine til atlantisk laks

Not authorised

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

Product identification

Medicine name:
Pentium forte plus vet. injeksjonsvæske, emulsjon, vaksine til atlantisk laks

Active substance:
Infectious pancreatic necrosis virus, Inactivated

Aliivibrio salmonicida, Inactivated

Moritella viscosa, 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 pancreatic necrosis virus, Inactivated
1.00 enzyme-linked immunosorbent assay unit / 0.10 millilitre(s)

Aliivibrio salmonicida, Inactivated
90.00 Relative Percentage Survival / 0.10 millilitre(s)

Moritella viscosa, Inactivated
70.00 Relative Percentage Survival / 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)

Aeromonas salmonicida, subsp. salmonicida, Inactivated
70.00 Relative Percentage Survival / 0.10 millilitre(s)

Pharmaceutical form:
Emulsion for injection

Withdrawal period by route of administration:
Intraperitoneal use:

- **Atlantic salmon**
  - Fish meat. 0 degree day

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QI10AL02

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

Authorisation status:

Revoked

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:
10/07/2007

Manufacturing sites for batch release:
Lohmann Animal Health GmbH

Responsible authority:
Norwegian Medical Products Agency

Authorisation number:
07-4936

Date of authorisation status change:
28/07/2026

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.