BI-FISHVAX

Authorised

- Yersinia ruckeri, Inactivated
- Vibrio anguillarum, Inactivated

Product identification

Medicine name:
BI-FISHVAX

Active substance:
Yersinia ruckeri, Inactivated

Vibrio anguillarum, Inactivated

Target species:
Trout

Route of administration:
Water-borne use
Intraperitoneal use

Product details

Active substance and strength:
Yersinia ruckeri, Inactivated
70.00 Relative Percentage Survival / 1.00 millilitre(s)

Vibrio anguillarum, Inactivated
70.00 Relative Percentage Survival / 1.00 millilitre(s)

Pharmaceutical form:
Suspension for injection

Withdrawal period by route of administration:

Water-borne use:

- **Trout**
  - Meat and offal. 0 day

Intraperitoneal use:

- **Trout**
  - Meat and offal. 0 day

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QI10BB01

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

Authorisation status:
Valid

Authorised in:
Italy

Package description:
Available only in Italian
Available only in Italian

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:
Fatro S.p.A.

Marketing authorisation date:
31/05/1993

Manufacturing sites for batch release:
Fatro S.p.A.

Responsible authority:
Ministry Of Health

Authorisation number:
This information is not available for this product.

Date of authorisation status change:
31/05/1993

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

Documents

Combined File of all Documents

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