

ICTHIOVAC-TM TENACIBACULOSIS RODABALLO SUSPENSION INYECTIONABLE PARA RODABALLO

Authorised

- Tenacibaculum maritimum, serotype O2, strain LPV 1.7, Inactivated

Product identification

Medicine name:

ICTHIOVAC-TM TENACIBACULOSIS RODABALLO SUSPENSION INYECTIONABLE PARA RODABALLO

Active substance:

Tenacibaculum maritimum, serotype O2, strain LPV 1.7, Inactivated

Target species:

Turbot

Route of administration:

Intraperitoneal use

Product details

Active substance and strength:

Tenacibaculum maritimum, serotype O2, strain LPV 1.7, Inactivated
75.00 Relative Percentage Survival / 1.00 Dose

Pharmaceutical form:

Suspension for injection

Withdrawal period by route of administration:**Intraperitoneal use:**

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Turbot

- Meat. 0 day

Anatomical therapeutic chemical veterinary (ATCvet) codes:

QI10D

Legal status of supply:

Veterinary medicinal product subject to veterinary prescription

Authorisation status:

Valid

Authorised in:

Spain

Package description:

Available only in Spanish

Additional information

Entitlement type:

Marketing Authorisation

Legal basis of product authorisation:

Full application - New active substance (Article 12(3) of Directive No 2001/82/EC)

Marketing authorisation holder:

Laboratorios Hipra S.A.

Marketing authorisation date:

31/05/2006

Manufacturing sites for batch release:

Laboratorios Hipra S.A.

Responsible authority:

Spanish Agency Of Medicines And Medical Devices

Authorisation number:

1691 ESP

Date of authorisation status change:

31/05/2006

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.

Labelling

This document does not exist in this language (English). You can find it in another language below.