

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:

-

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

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Labelling

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

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