ZANIL SUSPENSION

Authorised

This information is not available for this product.

Product identification

Medicine name:

ZANIL SUSPENSION

Active substance:

This information is not available for this product.

Target species:

Cattle

Sheep

Route of administration:

Oral use

Product details

Active substance and strength:

This information is not available for this product.

Pharmaceutical form:

Oral suspension

Withdrawal period by route of administration:

Oral use:

- . Cattle
 - Meat and offal. 13 day

- Milk. 5 day
- . Sheep
 - Meat and offal. 14 day
 - Milk. 7 day

Anatomical therapeutic chemical veterinary (ATCvet) codes:

QP52AG06

Legal status of supply:

Veterinary medicinal product subject to veterinary prescription

Authorisation status:

Valid

Authorised in:

France

Package description:

Available only in French

Available only in French

Additional information

Entitlement type:

Marketing Authorisation

Legal basis of product authorisation:

Legal basis not covered by Directive 2001/82/EC

Marketing authorisation holder:

Intervet

Marketing authorisation date:

22/06/1979

Manufacturing sites for batch release:

Trirx Segre

Responsible authority:

French Agency For Food, Environmental And Occupational Health & Safety

Authorisation number:

FR/V/9273440 3/1979

Date of authorisation status change:

22/06/2009

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.

Source URL: https://medicines.health.europa.eu/veterinary/600000032136