Stelfonta 1 mg/ml - Solution for injection

Authorised

- Tigilanol tiglate

Product identification

Medicine name:
Stelfonta 1 mg/ml - Solution for injection

Active substance:
Tigilanol tiglate

Target species:
Dog

Route of administration:
Intratumoral use

Product details

Active substance and strength:
Tigilanol tiglate
1.00 milligram(s) / 1.00 Vial

Pharmaceutical form:
Solution for injection

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QL01XX91

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

Authorisation status:
Valid

Authorised in:
Austria , Belgium , Bulgaria , Croatia , Cyprus , Czechia , Denmark , Estonia , Finland , France , Germany , Greece , Hungary , Iceland , Ireland , Italy , Latvia , Liechtenstein , Lithuania , Luxembourg , Malta , Netherlands , Norway , Poland , Portugal , Romania , Slovakia , Slovenia , Spain , Sweden , United Kingdom (Northern Ireland)

Package description:
Packaging:Vial (glass), Package_size:1 vial, Content:2 ml

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:
QBiotics Netherlands B.V.

Marketing authorisation date:
15/01/2020

Manufacturing sites for batch release:
Virbac

Responsible authority:
European Commission

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

Date of authorisation status change:
15/01/2020

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

Documents

Combined File of all Documents

English (PDF)
Published on: 2/09/2022
Download

ema-puar-stelfonta-v-5018-par-en.pdf