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Prazivetin 500 mg/g - Premix for medicated feeding stuff

Authorised

- Praziquantel

Product identification

Medicine name:
Prazivetin 500 mg/g - Premix for medicated feeding stuff

Active substance:
Praziquantel

Target species:
Gilthead

Route of administration:
In-feed use

Product details

Active substance and strength:
Praziquantel

Presentation_strength:500 mg Reference:Ph.Eur. Index:0

Pharmaceutical form:
Premix for medicated feeding stuff

Withdrawal period by route of administration:

In-feed use:

- **Gilthead**
  - Not applicable. 120 day 120 degree days

Anatomical therapeutic chemical veterinary (ATCvet) codes:

QP52AA01

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)

Available 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:Bag (PE), Package_size:1 bag, Content:20 kg

Packaging:Bag (PE), Package_size:8 bags, Content:2 kg

Additional information

Entitlement type:

Marketing Authorisation

Legal basis of product authorisation:

Full application - known active substance (Article 8 of Regulation (EU) 2019/6)

Marketing authorisation holder:
Vethellas S.A.

Marketing authorisation date:
23/04/2025

Manufacturing sites for batch release:
Vethellas S.A.

Responsible authority:
European Commission

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

Date of authorisation status change:
23/04/2025

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: 13/03/2026
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ema-puar-v6247-prazivetin-initial-en.pdf