

File downloaded on 2026-05-22

Source URL: <https://medicines.health.europa.eu/veterinary/en/600000042921>

EQUEST PRAMOX 19.5 MG/G + 121.7 MG/G ORAL GEL FOR HORSES

Authorised

- Moxidectin
- Praziquantel

Product identification

Medicine name:

EQUEST PRAMOX 19.5 MG/G + 121.7 MG/G ORAL GEL FOR HORSES

Active substance:

Moxidectin

Praziquantel

Target species:

Horse (mare)

Horse

Route of administration:

Oral use

Product details

Active substance and strength:

Moxidectin

19.50 milligram(s) / 1.00 gram(s)

Praziquantel

121.70 milligram(s) / 1.00 gram(s)

Pharmaceutical form:

Oral gel

Withdrawal period by route of administration:

Oral use:

-

Horse (mare)

- Milk. no withdrawal period

Not permitted for use in lactating mares producing milk for human consumption.

-

Horse

- Meat and offal. 64 day

Anatomical therapeutic chemical veterinary (ATCvet) codes:

QP54AB52

Legal status of supply:

Veterinary medicinal product subject to veterinary prescription

Authorisation status:

Valid

Authorised in:

Belgium

Available in:

Belgium

Package description:

Box containing 1 individually boxed syringes

Box containing 20 syringes

Box containing 20 individually boxed syringes

Box containing 10 individually boxed syringes

Additional information

Entitlement type:

Marketing Authorisation

Legal basis of product authorisation:

Legal basis not covered by Directive 2001/82/EC

Marketing authorisation holder:

Zoetis Belgium

Marketing authorisation date:

16/08/2006

Manufacturing sites for batch release:

Zoetis Manufacturing & Research Spain S.L.

Responsible authority:

Federal Agency For Medicines And Health Products

Authorisation number:

BE-V285984

Date of authorisation status change:

16/08/2006

Reference member state:

France

Procedure number:

FR/V/0161/001

Concerned member states:

Austria Belgium Czechia Denmark Finland Germany Greece Hungary
Ireland Italy Lithuania Luxembourg Netherlands Norway Poland Portugal
Slovakia Spain Sweden United Kingdom (Northern Ireland)

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.

Combined File of all Documents