LEVOPLIX 10%, pulbere orală

Valtuutettu

- Levamisole hydrochloride

Product identification

Lääkkeen nimi:
LEVOPLIX 10%, pulbere orală

Vaikuttava aine:
Saatavissa vain English

Kohde-eläinlaji(t):
Saatavissa vain Bulgarian Spanish Czech Danish German Estonian Greek English French Italian Latvian Lithuanian Hungarian Dutch Romanian Slovenian Swedish Icelandic Norwegian
Saatavissa vain Bulgarian Spanish Czech Danish German Estonian Greek English French Italian Latvian Lithuanian Hungarian Dutch Romanian Swedish Icelandic Norwegian
Saatavissa vain Bulgarian Spanish Czech Danish German Estonian Greek English French Italian Latvian Lithuanian Hungarian Dutch Romanian Swedish Icelandic Norwegian
Saatavissa vain Bulgarian Spanish Danish German Estonian Greek English French Italian Latvian Lithuanian Hungarian Dutch Romanian Swedish Icelandic Norwegian

Antoreitti:
Juomaveteen sekoitettuna

Product details

Vaikuttava aine / Vahvuus:

Saatavissa vain English
100.00 milligram(s) / 1.00 gram(s)

Lääkemuoto:

Jauhe

Withdrawal period by route of administration:

Juomaveteen sekoitettuna:

- **Cattle**

  - Meat and offal. 28 day

Nu este autorizată utilizarea la animalele al căror lapte este destinat consumului uman.

- **Sheep**

  - Meat and offal. 28 day

Nu este autorizată utilizarea la animalele al căror lapte este destinat consumului uman.

- **Pig**

  - Meat and offal. 28 day

- **Chicken (hen)**

  - Meat and offal. 28 day

Nu este autorizată utilizarea la păsările ale căror ouă sunt destinate consumului uman.

Anatomis-terapeuttis-kemiallinen koodi (ATCvet):

QP52AE01

Toimituksen oikeudellinen asema:

Tätä tietoa ei ole saatavana tästä valmisteesta.

Myyntiluvan tila:

Valid

Authorised in:

Saatavissa vain Spanish Czech German Estonian English French Italian Dutch Portuguese Romanian Slovak Swedish Icelandic Norwegian

Pakkauksen kuvaus:
Saatavissa vain Romanian
Saatavissa vain Romanian
Saatavissa vain Romanian
Saatavissa vain Romanian
Saatavissa vain Romanian

Additional information

Entitlement type:
Marketing Authorisation

Myyntiluvan oikeusperusta:
Saatavissa vain English

Myyntiluvan haltija:
Erfar S.A.

Marketing authorisation date:
28/09/2006

Erän vapauttamisesta vastaavat valmistuspaikat:
Erfar S.A.

Vastuullinen viranomainen:
Institute For Control Of Biological Products And Veterinary Medicines

Myyntiluvan numero:
160178

Myyntiluvan tilan muutoksen päivämäärä:
14/06/2016

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

Documents

Valmisteyhteenveto

Tätä asiakirjaa ei ole tällä kielellä (suomi). Löydät sen toisesta kielestä alla.
