HIPRALONA ENRO - I

Not authorised

- Enrofloxacin

Product identification

Medicine name:
ХИПРАЛОНА ЕНРО - И
HIPRALONA ENRO - I

Active substance:
Enrofloxacin

Target species:
Cattle (calf)
Pig

Route of administration:
Intramuscular use

Product details

Active substance and strength:
Enrofloxacin
50.00 milligram(s) / 1.00 millilitre(s)

Pharmaceutical form:
Solution for injection

Withdrawal period by route of administration:
Intramuscular use:

- **Cattle (calf)**
  - Meat and offal. 12 day

Не се разрешава за употреба при животни, чието мляко е предназначено за човешка консумацияСлед подкожно инжектиране: Месо и вътрешни органи: 12 дни

  - Meat and offal. 5 day

След подкожно инжектиране: Месо и вътрешни органи: 12 дни. Не се разрешава за употреба при животни, чието мляко е предназначено за човешка консумация

- **Pig**
  - Meat and offal. 13 day

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QJ01MA90

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

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Authorisation status:
Surrendered

Authorised in:
Bulgaria

Package description:
Available only in Bulgarian

Additional information

Entitlement type:

Marketing Authorisation

Legal basis of product authorisation:
Full application - Known active substance (Article 12(3) of Directive No 2001/82/EC)

Marketing authorisation holder:
Laboratorios Hipra S.A.

Marketing authorisation date:
20/11/2007

Manufacturing sites for batch release:
Laboratorios Hipra S.A.

Responsible authority:
Bulgarian Food Safety Authority

Authorisation number:
0022-1934-21.01.2013

Date of authorisation status change:
30/03/2025

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 and Labelling

This document does not exist in this language (English). You can find it in another language below.