PRACETAM 200 MG/ML SOLUTION FOR USE IN DRINKING WATER FOR PIGS

Authorised

- Paracetamol

Product identification

Medicine name:
PRACETAM 200 MG/ML SOLUTION FOR USE IN DRINKING WATER FOR PIGS

Active substance:
Paracetamol

Target species:
Pig

Route of administration:
Oral use

Product details

Active substance and strength:
Paracetamol
200.00 milligram(s) / 1.00 millilitre(s)

Pharmaceutical form:

Solution for use in drinking water

Withdrawal period by route of administration:

Oral use:

- **Pig**
  - Meat and offal. 0 day

Anatomical therapeutic chemical veterinary (ATCvet) codes:

QN02BE01

Legal status of supply:

Veterinary medicinal product subject to veterinary prescription

Authorisation status:

Valid

Authorised in:

Poland

Package description:

High density polyethylene bottle with high density polyethylene screwcap and polyethylene-aluminium-wax-paper-low density polyethylene seal (1 L bottle)
High density polyethylene bottle with high density polyethylene screwcap and polyethylene-PET-aluminium-wax-cardboard seal (10 L bottle)
High density polyethylene bottle with high density polyethylene screwcap and polyethylene-PET-aluminium-wax-cardboard seal (5 L bottle)
High density polyethylene bottle with high density polyethylene screwcap and polyethylene-PET-aluminium-wax-cardboard seal (2 L bottle)
High density polyethylene bottle with polypropylene screwcap and polyethylene seal (1 L bottle).
High density polyethylene bottle with polypropylene screwcap and polyethylene seal (5 L bottle).

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:
Ceva Animal Health Polska Sp. z o.o.

Marketing authorisation date:
16/11/2010

Manufacturing sites for batch release:
Ceva Sante Animale

Responsible authority:
Office For Registration Of Medicinal Products Medical Devices And Biocidal Products

Authorisation number:
2013

Date of authorisation status change:
16/11/2010

Reference member state:
France

Procedure number:
FR/V/0181/001

Concerned member states:
Austria Belgium Bulgaria Czechia Denmark Estonia Germany Hungary Italy Latvia Lithuania Netherlands Poland Portugal Romania Slovakia Spain United Kingdom (Northern Ireland)

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

Documents

Package Leaflet

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

Summary of Product Characteristics

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