NEMAPROL SOLUTION BUVABLE

Authorised

- Amprolium hydrochloride

Product identification

Medicine name:
NEMAPROL SOLUTION BUVABLE

Active substance:
Amprolium hydrochloride

Target species:
Poultry

Route of administration:
Oral use

Product details

Active substance and strength:
Amprolium hydrochloride
120.00 milligram(s) / 1.00 millilitre(s)

Pharmaceutical form:
Oral solution

Withdrawal period by route of administration:

Oral use:

- **Poultry**
  - Meat and offal. 0 day
  - Eggs. 0 day

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QP51AX09

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

Authorisation status:
Valid

Authorised in:
France

Package description:
Available only in French
Available only in French

Additional information

Entitlement type:
Marketing Authorisation

Legal basis of product authorisation:
Legal basis not covered by Directive 2001/82/EC

Marketing authorisation holder:
Dopharma France

Marketing authorisation date:
17/02/1992

Manufacturing sites for batch release:
Dopharma France

Responsible authority:
French Agency For Food, Environmental And Occupational Health & Safety

Authorisation number:
FR/V/5336886 7/1992

Date of authorisation status change:
17/02/2012

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.