CURACEF DUO, 50 MG/ML + 150 MG/ML, SUSPENSION FOR INJECTION FOR CATTLE

Authorised

- Ceftiofur hydrochloride
- Ketoprofen

Product identification

Medicine name:
CURACEF DUO, 50 MG/ML + 150 MG/ML, SUSPENSION FOR INJECTION FOR CATTLE

Active substance:
Ceftiofur hydrochloride

Ketoprofen

Target species:
Cattle

Route of administration:
Intramuscular use

Product details

Active substance and strength:
Ceftiofur hydrochloride
53.50 milligram(s) / 1.00 millilitre(s)

Ketoprofen
150.00 milligram(s) / 1.00 millilitre(s)

Pharmaceutical form:
Suspension for injection

Withdrawal period by route of administration:
Intramuscular use:

- **Cattle**
  - Meat and offal. 8 day
  - Milk. 0 day

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QJ01DD99

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

Authorisation status:
Valid

Authorised in:
Cyprus

Package description:
Available only in French
Available only in French
Available only in French
Available only in French
Available only in French
Available only in French

Additional information

Entitlement type:
Marketing Authorisation

Legal basis of product authorisation:
Fixed combination application (Article 13b of Directive No 2001/82/EC)

Marketing authorisation holder:
Virbac

Marketing authorisation date:
2/10/2014

Manufacturing sites for batch release:
Virbac

Responsible authority:
Veterinary Services, Ministry Of Agriculture, Natural Resources And Environment

Authorisation number:
CY00485V

Date of authorisation status change:
23/05/2019

Reference member state:
France

Procedure number:
FR/V/0258/001

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

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

Documents

Combined File of all Documents

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