WONDERCEF, 1g 50 mg/ml polvere e solvente per soluzione iniettabile per equidi non destinati alla produzione di alimenti per il consumo umano (NDPA)

Authorised

- Ceftiofur

Product identification

Medicine name:
WONDERCEF, 1g 50 mg/ml polvere e solvente per soluzione iniettabile per equidi non destinati alla produzione di alimenti per il consumo umano (NDPA)

Active substance:
Ceftiofur

Target species:
Horse (non food-producing)

Route of administration:
Intramuscular use

Product details

Active substance and strength:
Ceftiofur
1060.00 milligram(s) / 1.00 gram(s)

Pharmaceutical form:
Lyophilisate and solvent for solution for injection

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QJ01DD90

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

Authorisation status:
Valid

Authorised in:
Italy

Package description:
Available only in Italian

Additional information

Entitlement type:
Marketing Authorisation

Legal basis of product authorisation:
Generic application (Article 13(1) of Directive No 2001/82/EC)

Marketing authorisation holder:
Fatro S.p.A.

Marketing authorisation date:
11/06/2008

Manufacturing sites for batch release:
Fatro S.p.A.

Responsible authority:
Ministry Of Health

Authorisation number:
This information is not available for this product.

Date of authorisation status change:
18/12/2012

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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