

FATROXIMIN endomammario asciutta 100 mg/5 ml pomata intramammaria per bovine e bufale

Authorised

- Rifaximin

Product identification

Medicine name:

FATROXIMIN endomammario asciutta 100 mg/5 ml pomata intramammaria per bovine e bufale

Active substance:

Rifaximin

Target species:

Cattle (cow)

Buffalo (female)

Route of administration:

Intramammary use

Product details

Active substance and strength:

Rifaximin

100.00 milligram(s) / 1.00 Syringe

Pharmaceutical form:

Intramammary emulsion

Withdrawal period by route of administration:

Intramammary use:

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Cattle (cow)

- Milk. 0 day

Non utilizzare a scopo alimentare le mammelle degli animali trattati

- Meat and offal. 0 day

Non utilizzare a scopo alimentare le mammelle degli animali trattati

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Buffalo (female)

- Milk. 0 day

Non utilizzare a scopo alimentare le mammelle degli animali trattati

- Meat and offal. 0 day

Non utilizzare a scopo alimentare le mammelle degli animali trattati

Anatomical therapeutic chemical veterinary (ATCvet) codes:

QJ51XX01

Legal status of supply:

Veterinary medicinal product subject to veterinary prescription

Authorisation status:

Valid

Authorised in:

Italy

Package description:

Available only in Italian

Available only in [Italian](#)

Available only in [Italian](#)

Additional information

Entitlement type:

Marketing Authorisation

Legal basis of product authorisation:

Full application (Article 12(3) of Directive No 2001/82/EC)

Marketing authorisation holder:

Fatro S.p.A.

Marketing authorisation date:

16/10/1987

Manufacturing sites for batch release:

Fatro S.p.A.

Responsible authority:

Ministry Of Health

Authorisation number:

100077

Date of authorisation status change:

16/10/2007

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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