

BIOFEL M PLUS

Authorised

- Microsporium canis, Inactivated

Product identification

Medicine name:

BIOFEL M PLUS

Active substance:

Microsporium canis, Inactivated

Target species:

Dog

Cat

Route of administration:

Intramuscular use

Subcutaneous use

Product details

Active substance and strength:

Microsporium canis, Inactivated

1.00 million colony forming units / 1.00 Dose

Pharmaceutical form:

Suspension for injection

Withdrawal period by route of administration:

Intramuscular use:

-

Dog

- Unspecified. 0 day

-

Cat

- Unspecified. 0 day

Subcutaneous use:

-

Cat

- Unspecified. 0 day

Anatomical therapeutic chemical veterinary (ATCvet) codes:

QI07AQ02

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](#)

Available only in [Italian](#)

Available only in [Italian](#)

Additional information

Entitlement type:

Marketing Authorisation

Legal basis of product authorisation:

Immunological veterinary medicinal product application (Article 13d of Directive No 2001/82/EC)

Marketing authorisation holder:

Bio 98 S.r.l.

Marketing authorisation date:

7/02/2017

Manufacturing sites for batch release:

Bioveta a.s.

Responsible authority:

Ministry Of Health

Authorisation number:

This information is not available for this product.

Date of authorisation status change:

7/02/2017

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

Documents

Combined File of all Documents

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