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Artuvetrin-Test Mijten/Insecten oplossing voor injectie voor honden

Authorised

- Aedes communis extract
- TYROPHAGUS PUTRESCENTIAE EXTRACT
- ACARUS SIRO EXTRACT
- DERMATOPHAGOIDES FARINAE EXTRACT
- DERMATOPHAGOIDES PTERONYSSINUS EXTRACT
- Musca domestica extract
- LEPIDOGLYPHUS DESTRUCTOR EXTRACT
- Ctenocephalides spp. extract

Product identification

Medicine name:
Artuvetrin-Test Mijten/Insecten oplossing voor injectie voor honden

Active substance:
Aedes communis extract

TYROPHAGUS PUTRESCENTIAE EXTRACT

ACARUS SIRO EXTRACT

DERMATOPHAGOIDES FARINAE EXTRACT

DERMATOPHAGOIDES PTERONYSSINUS EXTRACT

Musca domestica extract

LEPIDOGLYPHUS DESTRUCTOR EXTRACT

Ctenocephalides spp. extract

Target species:
Dog

Route of administration:
Intradermal use

Product details

Active substance and strength:
Aedes communis extract
100.00 noon unit / 1.00 millilitre(s)

TYROPHAGUS PUTRESCENTIAE EXTRACT
100.00 noon unit / 1.00 millilitre(s)

ACARUS SIRO EXTRACT
100.00 noon unit / 1.00 millilitre(s)

DERMATOPHAGOIDES FARINAE EXTRACT
100.00 noon unit / 1.00 millilitre(s)

DERMATOPHAGOIDES PTERONYSSINUS EXTRACT
100.00 noon unit / 1.00 millilitre(s)

Musca domestica extract
100.00 noon unit / 1.00 millilitre(s)

LEPIDOGLYPHUS DESTRUCTOR EXTRACT
100.00 noon unit / 1.00 millilitre(s)

Ctenocephalides spp. extract
10.00 noon unit / 1.00 millilitre(s)

Pharmaceutical form:
Solution for injection

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QV01AA

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

Authorisation status:
Valid

Authorised in:
Netherlands

Package description:
Available only in Dutch

Additional information

Entitlement type:
Marketing Authorisation

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

Marketing authorisation holder:
Nextmune B.V.

Marketing authorisation date:
21/11/1991

Manufacturing sites for batch release:
Nextmune B.V.

Responsible authority:
Medicines Evaluation Board

Authorisation number:
REG NL 1234

Date of authorisation status change:
21/11/1991

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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