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Gudair emulsie voor injectie voor schapen en geiten

Authorised

- Mycobacterium avium, subsp. paratuberculosis, strain 316F, Inactivated

Product identification

Medicine name:
Gudair emulsie voor injectie voor schapen en geiten

Active substance:
Mycobacterium avium, subsp. paratuberculosis, strain 316F, Inactivated

Target species:
Goat (adult female)
Sheep

Route of administration:
Subcutaneous use

Product details

Active substance and strength:
Mycobacterium avium, subsp. paratuberculosis, strain 316F, Inactivated
2.00 tuberculin unit(s) / 1.00 millilitre(s)

Pharmaceutical form:

Emulsion for injection

Withdrawal period by route of administration:

Subcutaneous use:

- **Goat (adult female)**
  - Milk. 0 day
  - Meat and offal. 0 day
- **Sheep**
  - Milk. 0 day
  - Meat and offal. 0 day

Anatomical therapeutic chemical veterinary (ATCvet) codes:

QI03AB01

QI04AB

Legal status of supply:

Veterinary medicinal product subject to veterinary prescription

Authorisation status:

Valid

Authorised in:

Netherlands

Available in:

Netherlands

Package description:

Available only in Dutch

Available only in Dutch

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:
CZ Vaccines S.A.U.

Marketing authorisation date:
11/09/2014

Manufacturing sites for batch release:
CZ Vaccines S.A.U.

Responsible authority:
Medicines Evaluation Board

Authorisation number:
REG NL 113936

Date of authorisation status change:
24/04/2019

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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