

GUDAIR Emulsion for injection for sheep and goats

Authorised

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

Product identification

Medicine name:

GUDAIR Emulsion for injection for sheep and goats

Active substance:

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

Target species:

Sheep

Goat

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:

-

Sheep

- Meat and offal. 0 day

-

Goat

- Meat and offal. 0 day

Anatomical therapeutic chemical veterinary (ATCvet) codes:

QI03AB01

Legal status of supply:

Veterinary medicinal product subject to veterinary prescription

Authorisation status:

Valid

Authorised in:

Portugal

Available in:

Portugal

Package description:

box containing 1 vial of 30 ml (30 doses)

Additional information

Entitlement type:

Marketing Authorisation

Legal basis of product authorisation:

Full application - Known active substance (Article 12(3) of Directive No 2001/82/EC)

Marketing authorisation holder:

CZ Vaccines S.A.U.

Marketing authorisation date:

23/12/2020

Manufacturing sites for batch release:

CZ Vaccines S.A.U.

Responsible authority:

Directorate General For Food And Veterinary

Authorisation number:

1392/01/20RIVPT

Date of authorisation status change:

26/08/2022

Reference member state:

Spain

Procedure number:

ES/V/0210/001

Concerned member states:

France Iceland Norway Portugal

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

Documents

Combined File of all Documents

English (PDF)

Published on: 13/03/2026

Download

Summary of Product Characteristics

Package Leaflet

Labelling