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Footvax Emulsion for injection

Authorised

- Dichelobacter nodosus, serogroup I, strain 109, Inactivated
- Dichelobacter nodosus, serogroup A, strain 6, Inactivated
- Dichelobacter nodosus, serotype B1, strain 44, Inactivated
- Dichelobacter nodosus, serotype B2, strain 58, Inactivated
- Dichelobacter nodosus, serogroup C, strain 8, Inactivated
- Dichelobacter nodosus, serogroup D, strain 16, Inactivated
- Dichelobacter nodosus, serogroup E, strain 5, Inactivated
- Dichelobacter nodosus, serogroup F, strain 66, Inactivated
- Dichelobacter nodosus, serogroup G, strain 52, Inactivated
- Dichelobacter nodosus, serogroup H, strain 340, Inactivated

Product identification

Medicine name:
Footvax Emulsion for injection

Active substance:
Dichelobacter nodosus, serogroup I, strain 109, Inactivated
Dichelobacter nodosus, serogroup A, strain 6, Inactivated
Dichelobacter nodosus, serotype B1, strain 44, Inactivated
Dichelobacter nodosus, serotype B2, strain 58, Inactivated
Dichelobacter nodosus, serogroup C, strain 8, Inactivated
Dichelobacter nodosus, serogroup D, strain 16, Inactivated
Dichelobacter nodosus, serogroup E, strain 5, Inactivated
Dichelobacter nodosus, serogroup F, strain 66, Inactivated

Dichelobacter nodosus, serogroup G, strain 52, Inactivated

Dichelobacter nodosus, serogroup H, strain 340, Inactivated

Target species:
Sheep

Route of administration:
Subcutaneous use

Product details

Active substance and strength:
Dichelobacter nodosus, serogroup I, strain 109, Inactivated
500000000.00 cells / 1.00 millilitre(s)

Dichelobacter nodosus, serogroup A, strain 6, Inactivated
10.00 microgram(s) / 1.00 millilitre(s)

Dichelobacter nodosus, serotype B1, strain 44, Inactivated
10.00 microgram(s) / 1.00 millilitre(s)

Dichelobacter nodosus, serotype B2, strain 58, Inactivated
10.00 microgram(s) / 1.00 millilitre(s)

Dichelobacter nodosus, serogroup C, strain 8, Inactivated
10.00 microgram(s) / 1.00 millilitre(s)

Dichelobacter nodosus, serogroup D, strain 16, Inactivated
10.00 microgram(s) / 1.00 millilitre(s)

Dichelobacter nodosus, serogroup E, strain 5, Inactivated
10.00 microgram(s) / 1.00 millilitre(s)

Dichelobacter nodosus, serogroup F, strain 66, Inactivated
10.00 microgram(s) / 1.00 millilitre(s)

Dichelobacter nodosus, serogroup G, strain 52, Inactivated
10.00 microgram(s) / 1.00 millilitre(s)

Dichelobacter nodosus, serogroup H, strain 340, Inactivated
10.00 microgram(s) / 1.00 millilitre(s)

Pharmaceutical form:
Emulsion for injection

Withdrawal period by route of administration:

Subcutaneous use:

- **Sheep**
  - Meat and offal. 0 day
  - Milk. 0 day

Anatomical therapeutic chemical veterinary (ATCvet) codes:
QI04AB03

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

Authorisation status:
Valid

Authorised in:
Germany

Available in:
Germany

Package description:
(ID1) 20 millilitre(s): Box (Cardboard) with 1 Bottle (Low Density PolyEthylene) with 20 millilitre(s)
(ID2) 50 millilitre(s): Box (Cardboard) with 1 Bottle (Low Density PolyEthylene) with 50 millilitre(s)

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:
Intervet Deutschland GmbH

Marketing authorisation date:
24/09/1996

Manufacturing sites for batch release:
Intervet International B.V.

Responsible authority:
Paul-Ehrlich-Institut

Authorisation number:
66a/95

Date of authorisation status change:
29/03/2007

Reference member state:
Germany

Procedure number:
DE/V/0220/001

Concerned member states:
Austria

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

Documents

Combined File of all Documents

English (PDF)

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