

Footvax, Injekční emulze

Authorised

- Dichelobacter nodosus, serogroup A, strain 6, Inactivated
- Dichelobacter nodosus, serotype B1, strain 44, Inactivated
- Dichelobacter nodosus, serotype B2, strain 58, Inactivated
- Dichelobacter nodosus, serogroup D, strain 16, Inactivated
- Dichelobacter nodosus, serogroup C, strain 8, 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
- Dichelobacter nodosus, serogroup I, strain 109, Inactivated

Product identification

Medicine name:

Footvax, Injekční emulze

Active substance:

Dichelobacter nodosus, serogroup A, strain 6, Inactivated
Dichelobacter nodosus, serotype B1, strain 44, Inactivated
Dichelobacter nodosus, serotype B2, strain 58, Inactivated
Dichelobacter nodosus, serogroup D, strain 16, Inactivated
Dichelobacter nodosus, serogroup C, strain 8, 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

Dichelobacter nodosus, serogroup I, strain 109, Inactivated

Target species:

Sheep

Route of administration:

Subcutaneous use

Product details

Active substance and strength:

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 D, strain 16, 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 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)

Dichelobacter nodosus, serogroup I, strain 109, Inactivated

500000000.00 cells / 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 hour

Anatomical therapeutic chemical veterinary (ATCvet) codes:

QI04AB03

Legal status of supply:

Veterinary medicinal product subject to veterinary prescription

Authorisation status:

Valid

Authorised in:

Czechia

Available in:

Czechia

Package description:

Available only in [Czech](#)

Available only in [Czech](#)

Available only in [Czech](#)

Additional information

Entitlement type:

Marketing Authorisation

Legal basis of product authorisation:

Full application (Article 12(3) of Directive No 2001/82/EC)

Marketing authorisation holder:

Intervet International B.V.

Marketing authorisation date:

21/05/2001

Manufacturing sites for batch release:

Intervet International B.V.

Responsible authority:

Institute For State Control Of Veterinary Biologicals And Medicaments

Authorisation number:

97/023/01-C

Date of authorisation status change:

22/09/2011

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

Documents

Summary of Product Characteristics

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

Package Leaflet

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

Labelling

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