

Bovilis Bovivac S suspension for injection for cattle

Authorised

- Salmonella enterica, subsp. enterica, serovar Dublin, strain S342/70, Inactivated
- Salmonella enterica, subsp. enterica, serovar Typhimurium, strain S341/70, Inactivated

Product identification

Medicine name:

Bovilis Bovivac S suspension for injection for cattle

Active substance:

Salmonella enterica, subsp. enterica, serovar Dublin, strain S342/70, Inactivated
Salmonella enterica, subsp. enterica, serovar Typhimurium, strain S341/70, Inactivated

Target species:

Cattle

Route of administration:

Subcutaneous use

Product details

Active substance and strength:

Salmonella enterica, subsp. enterica, serovar Dublin, strain S342/70, Inactivated

10000000.00 cells / 0.01 millilitre(s)

Salmonella enterica, subsp. enterica, serovar Typhimurium, strain S341/70,
Inactivated

10000000.00 cells / 0.01 millilitre(s)

Pharmaceutical form:

Suspension for injection

Withdrawal period by route of administration:

Subcutaneous use:

-

Cattle

- Meat and offal. 0 day

- Milk. 0 day

Anatomical therapeutic chemical veterinary (ATCvet) codes:

QI02AB

Legal status of supply:

Veterinary medicinal product not subject to veterinary prescription

Authorisation status:

Valid

Authorised in:

Ireland

Available in:

Ireland

Package description:

Polyethylene multidose bottles (50 ml) with rubber stoppers sealed with an aluminium crimp cap. The containers and closures comply with the Ph. Eur

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 (Ireland) Limited

Marketing authorisation date:

5/09/2001

Manufacturing sites for batch release:

MSD Animal Health UK Limited

Intervet International B.V.

Responsible authority:

Health Products Regulatory Authority

Authorisation number:

VPA10996/165/001

Date of authorisation status change:

5/09/2001

Reference member state:

Ireland

Procedure number:

IE/V/0436/001

Concerned member states:

United Kingdom (Northern Ireland)

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

Documents

Summary of Product Characteristics