

**Keep out of reach of children
FOR ANIMAL TREATMENT ONLY**

Bovilis[®] BVD

INACTIVATED VACCINE FOR PROTECTION AGAINST INFECTION WITH BOVINE VIRAL DIARRHOEA VIRUS (BVDV) IN CATTLE AND THE FOETUS.

For intramuscular injection.

50mL & 100mL

Read entire label before use.

DESCRIPTION

BOVILIS BVD is an inactivated vaccine containing 50 Elisa Units per dose of cytopathogenic BVD virus strain C86. The virus is grown in cell cultures and is inactivated with betapropiolactone. The antigen is adsorbed onto an aluminium salts adjuvant. The vaccine contains methyl parahydroxybenzoate as a preservative.

INDICATIONS

BOVILIS BVD is an inactivated vaccine for the active immunization of cattle, four months of age and onwards, for protection against infection with bovine viral diarrhoea virus and protection against foetal infection with Bovine Viral Diarrhoea Virus (BVDV). In healthy animals no clinical reactions due to the vaccination will be observed. A slight swelling may be felt at the site of vaccination during 1 to 2 weeks.

DOSAGE AND ADMINISTRATION

Each animal should be given 2mL of vaccine by intramuscular injection.

RECOMMENDED VACCINATION PROGRAMME

BOVILIS BVD should be given to cattle from four months of age onwards.

For the primary vaccination the animals should be given a single dose (2mL) followed by a booster (2mL) 4 weeks later.

After these primary vaccinations, to maintain immunity, revaccinate all cattle annually with a single dose (2mL) approximately 4 weeks before each pregnancy.

If primary vaccinations are given to calves at less than 8 months of age, they should be given a booster dose (2mL) approximately 4 weeks before the first pregnancy. If primary vaccinations are given to animals of breeding age, they should be given at approximately 2 months before the first pregnancy followed by a booster (2mL) 4 weeks later.

WITHHOLDING PERIOD

Nil.

STORAGE CONDITIONS

Store at 2 to 8°C. Do not freeze.

ADDITIONAL INFORMATION

- Vaccinate healthy cattle only.
- Allow the vaccine to reach ambient temperature (15-25°C) before use.
- Shake the bottle well before use.
- Use sterile injection equipment.
- Do not mix with other vaccines.
- There is no special precaution necessary when given during pregnancy and lactation.

Prescription Animal Remedy (**P.A.R**) Class I. For use only under the authority or prescription of a veterinarian.

Registered pursuant to the ACVM Act 1997, No. A8237.
See www.nzfsa.govt.nz/acvm/ for registration conditions

Registered to:
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