PARTICULARS TO APPEAR ON THE OUTER PACKAGE Cardboard box (12 x 25 ml)

1. NAME OF THE VETERINARY MEDICINAL PRODUCT

Bovilis Huskvac oral suspension

2. STATEMENT OF ACTIVE SUBSTANCES

Each 25 ml dose contains:
1000 – 2000 viable 3rd stage irradiated *Dictyocaulus viviparus* larvae

3. PACKAGE SIZE

12 doses (12 x 25 ml)

4. TARGET SPECIES

Cattle.

5. INDICATIONS

6. ROUTES OF ADMINISTRATION

Oral use.
Shake well before use.

7. WITHDRAWAL PERIODS

Withdrawal period: Zero days.

8. EXPIRY DATE

Exp. {dd/mm/yy}

Following text included in red/white circle on front panel

Notice:
Short shelf life vaccine.
Please check the expiry date before use.

9. SPECIAL STORAGE PRECAUTIONS

Following text included in red/white circle on front panel

Store in a refrigerator between 2 °C – 8 °C.
Do not freeze.

10. THE WORDS “READ THE PACKAGE LEAFLET BEFORE USE”

Read the package leaflet before use.

11. THE WORDS “FOR ANIMAL TREATMENT ONLY”

For animal treatment only.

12. THE WORDS “KEEP OUT OF THE SIGHT AND REACH OF CHILDREN”

Keep out of the sight and reach of children.

13. NAME OF THE MARKETING AUTHORISATION HOLDER

MA Holder:
Intervet International B.V.

Distributor in Northern Ireland:
Intervet Ireland Ltd.

14. MARKETING AUTHORISATION NUMBERS

Vm 06376/4137

15. BATCH NUMBER

Lot {number}

MINIMUM PARTICULARS TO APPEAR ON SMALL IMMEDIATE PACKAGING UNITS
Glass Bottle Label

1. NAME OF THE VETERINARY MEDICINAL PRODUCT

Bovilis Huskvac

2. QUANTITATIVE PARTICULARS OF THE ACTIVE SUBSTANCES

Each 25 ml dose contains:
1000 - 2000 viable 3rd stage irradiated *Dictyocaulus viviparus* larvae

3. BATCH NUMBER

Lot {number}

4. EXPIRY DATE

Exp. {dd/mm/yy}

PARTICULARS TO APPEAR ON THE PACKAGE LEAFLET:

PACKAGE LEAFLET

1. Name of the veterinary medicinal product

Bovilis Huskvac oral suspension for cattle

2. Composition

Each 25 ml dose contains:

Active substances:
Dictyocaulus viviparus 3rd stage irradiated larvae: ≥ 1000 viable larvae, ≤ 2000 viable larvae

Clear aqueous suspension.

3. Target species

Cattle.

4. Indications for use

For the active immunisation of cattle to reduce clinical signs and lesions of parasitic bronchitis attributable to *Dictyocaulus viviparus* (lungworm).

Onset of immunity: 2 weeks after completion of the basic vaccination scheme.

Duration of immunity: Lungworm immunity is maintained from season to season by the exposure to lungworm larvae, which in most cases occurs from the grazing of normal pastures. If such exposure does not occur, a single booster dose is required prior to each season's turnout.

5. Contraindications

Do not use in calves showing any signs of illness, particularly those exhibiting any signs of respiratory distress.

6. Special warnings

Special warnings:
Vaccinate healthy animals only.

Routine vaccination of housed or suckled young stock prior to exposure to field lungworm challenge will help protect the calves and help reduce the levels of pasture contamination with lungworm larvae. However, owing to the ability of lungworm larvae to survive on pasture, calfhood vaccination programmes to control lungworm infection can only be successful if all susceptible calves are vaccinated in the spring before exposure to natural field infection occurs at turnout or weaning.

A good immune response is reliant on the reaction of an immunogenic agent and a fully competent immune system. Immunogenicity of the vaccine antigen will be reduced by poor storage or inappropriate administration. Immunocompetence of the animal may be compromised by a variety of factors including poor health, nutritional status, genetic factors, concurrent drug therapy and stress.

Special precautions for safe use in the target species:
Following vaccination, vaccinated stock should not be mixed with unvaccinated stock or allowed to graze on pastures recently used by unvaccinated stock until 2 weeks after the second dose of this vaccine.

Note: For optimum benefit it is important that the calf is exposed to pasture carrying some infection after this time, as this low-level exposure enhances the immunity induced by vaccination with this vaccine.

Special precautions to be taken by the person administering the veterinary medicinal product to animals:
Wash hands after use.

Pregnancy:
Can be used during pregnancy.

Interaction with other medicinal products and other forms of interaction:
No information is available on the safety and efficacy of this vaccine when used with any other veterinary medicinal product. A decision to use this vaccine before or after any other veterinary medicinal product therefore needs to be made on a case-by-case basis.
To ensure residual effects of long-acting anthelmintics and endectocides or sustained release bolus preparations do not interfere with the development of immunity following lungworm vaccination, avoid vaccination during the period of their activity, and do not use these veterinary medicinal products until 14 days after the second dose of this vaccine.

Overdose:
No symptoms other than those detailed in the section ‘Adverse events’ were observed following administration of twice the recommended dose.

Major incompatibilities:
Do not mix with any other veterinary medicinal product.

7. Adverse events

Cattle:

Very rare (<1 animal / 10 000 animals treated, including isolated reports):	Cough[1].

[1] May occur approximately 7 days after either dose. Transient episodes, which usually subside in a few days.

Reporting adverse events is important. It allows continuous safety monitoring of a product. If you notice any side effects, even those not already listed in this package leaflet, or you think that the medicine has not worked, please contact, in the first instance, your veterinarian. You can also report any adverse events to the marketing authorisation holder or its local representative using the contact details at the end of this leaflet, or via your national reporting system:
Website: https://www.gov.uk/report-veterinary-medicine-problem/animal-reacts-medicine
e-mail: adverse.events@vmd.gov.uk

8. Dosage for each species, routes and method of administration

Oral use.
One dose: 25 ml.

Vaccinate healthy animals of 8 weeks of age and older.

Vaccination regime

Basic vaccination scheme
Two doses at a dosage interval of approximately 4 weeks.

Revaccination
Lungworm immunity is maintained from season to season by the exposure to lungworm larvae, which in most cases occurs from the grazing of normal pastures after vaccination. Under these conditions of exposure, revaccination is generally not required.
A single dose of this vaccine prior to each season's turnout will boost immunity where such exposure has not occurred, e.g. extensive use of anthelmintics or if using reserved or clean pasture for a large part of the grazing season.

9. Advice on correct administration

Shake bottle well immediately before use.

10. Withdrawal periods

Zero days.

11. Special storage precautions

Keep out of the sight and reach of children.
Store in a refrigerator (2 °C – 8 °C).
Do not freeze.

Do not use this veterinary medicinal product after the expiry date which is stated on the label or carton after Exp.

12. Special precautions for disposal

Medicines should not be disposed of via wastewater or household waste.

Use take-back schemes for the disposal of any unused veterinary medicinal product or waste materials derived thereof in accordance with local requirements and with any applicable national collection systems. These measures should help to protect the environment.

Ask your veterinary surgeon or pharmacist how to dispose of medicines no longer required.

Dispose of waste material by boiling, incineration or immersion in an appropriate disinfectant approved for use by the competent authorities.

13. Classification of veterinary medicinal products

Veterinary medicinal product subject to prescription.

14. Marketing authorisation numbers and pack sizes

Vm 06376/4137

Pack size:
Cardboard box containing 12 x 25 ml (1 dose) bottles.

15. PID LINK (Do not print heading)

[The following statement must be included where reference to the European Union Product Database is included on the product information. This statement is relevant to both UK(GB) and UK(NI) products:]

Find more product information by searching for the 'Product Information Database' on www.gov.uk.

16. Contact details

Marketing authorisation holder:
Intervet International B.V.
Wim de Körverstraat 35
5831 AN Boxmeer
Netherlands

Manufacturer responsible for batch release[1]:
Intervet International B.V.
Wim de Körverstraat 35
5831 AN Boxmeer
Netherlands

MSD Animal Health UK Limited
Walton Manor, Walton
Milton Keynes
MK7 7AJ, United Kingdom

[1]The printed package leaflet will state the name and address of the manufacturer responsible for the release of the concerned batch only.

Local representative:
MSD Animal Health UK Limited
Walton Manor, Walton
Milton Keynes
MK7 7AJ, United Kingdom

Contact details to report suspected adverse events:
UK(GB)
MSD Animal Health UK Ltd.
Tel.: +44 (0)1908 685685

UK(NI)
Intervet Ireland Ltd.
Tel.: +353 (0)1 2970220

Distributor in Northern Ireland:
Intervet Ireland Ltd., Magna Drive, Magna Business Park, Citywest Road, Dublin 24, Ireland

For any information about this veterinary medicinal product, please contact the local representative of the marketing authorisation holder.

17. Other information

POM-V

Gavin Hall
Approved: 09 December 2025