PARTICULARS TO APPEAR ON THE OUTER PACKAGE
{Cardboard box}

1. NAME OF THE VETERINARY MEDICINAL PRODUCT

Nobivac Solvent

2. STATEMENT OF ACTIVE SUBSTANCES

Each 1 ml dose contains sterile phosphate buffered solvent.

3. PACKAGE SIZE

10 x 1 ml
50 x 1 ml

4. TARGET SPECIES

Cats and dogs.

5. INDICATIONS

Solvent for reconstitution of lyophilisates in the Nobivac vaccine range authorised for use with Nobivac Solvent.

6. ROUTES OF ADMINISTRATION

Read the instructions supplied with the vaccine carefully before using this solvent.

7. WITHDRAWAL PERIODS

8. EXPIRY DATE

Exp. {mm/yyyy}
Once broached, use immediately.

9. SPECIAL STORAGE PRECAUTIONS

Do not store above 25 °C.
Do not freeze.
Protect from light.
Keep the vials in the outer carton.

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

Intervet International B.V.

14. MARKETING AUTHORISATION NUMBERS

Vm 06376/4126

15. BATCH NUMBER

Lot {number}

MINIMUM PARTICULARS TO APPEAR ON SMALL IMMEDIATE PACKAGING UNITS {Glass vial }

1. NAME OF THE VETERINARY MEDICINAL PRODUCT

Nobivac Solvent
– sterile buffered solution

2. QUANTITATIVE PARTICULARS OF THE ACTIVE SUBSTANCES

1 dose
1 ml

3. BATCH NUMBER

Lot {number}

4. EXPIRY DATE

Exp. {mm/yyyy}

PARTICULARS TO APPEAR ON THE PACKAGE LEAFLET:

PACKAGE LEAFLET

1. Name of the veterinary medicinal product

Nobivac Solvent

2. Composition

Each 1 ml dose contains:

Excipients:
Potassium dihydrogen phosphate
Disodium phosphate
Water for injections

A clear, colourless solution.

3. Target species

Cats and dogs.

4. Indications for use

Solvent for reconstitution of lyophilisates in the Nobivac vaccine range authorised for use with Nobivac Solvent.

5. Contraindications

Any contraindications specified for the vaccine product for which the Nobivac Solvent is used for reconstitution will apply.

6. Special warnings

Special precautions for safe use in the target species:
No special precautions are required for handling the solvent, however any recommendations specified for the vaccine product for which Nobivac Solvent is used for reconstitution will apply.

Special precautions to be taken by the person administering the veterinary medicinal product to animals:
In case of accidental self-injection, seek medical advice immediately and show the package leaflet or the label to the physician.
After reconstitution: Any recommendations specified for the vaccine product for which Nobivac Solvent is used for reconstitution will apply.

Pregnancy and lactation:
Any recommendations specified for the vaccine product for which Nobivac Solvent is used for reconstitution will apply.

Interaction with other medicinal products and other forms of interaction:
Nobivac Solvent should only be used as a solvent for the reconstitution of lyophilisates in the Nobivac vaccine range authorised to be used with Nobivac Solvent.
No information is available on the safety and efficacy when used with any other veterinary medicinal product.
Any recommendations specified for the vaccine product for which Nobivac Solvent is used for reconstitution will apply.

Overdose:
Any recommendations specified for the vaccine product for which Nobivac Solvent is used for reconstitution will apply.

Major incompatibilities:
For use only with lyophilisates in the Nobivac vaccine product range which are authorised to be reconstituted with Nobivac Solvent.
Do not mix with any other veterinary medicinal product.

7. Adverse events

Any adverse events specified for the vaccine product for which Nobivac Solvent is used for reconstitution will apply.

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 the local representative of the marketing authorisation holder 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

Read the instructions supplied with the vaccine product carefully before using Nobivac Solvent.
The contents of one vial (1 ml) of Nobivac Solvent should be transferred aseptically into the vial of freeze-dried vaccine immediately prior to use.
Care should be taken to ensure that the freeze-dried powder plug has fully dissolved.

9. Advice on correct administration

Sterile equipment should be used for administration. Avoid contamination of vaccine with traces of chemical sterilising agents. Do not use chemicals such as disinfectant or spirit to disinfect the skin prior to inoculation.

10. Withdrawal periods

Not applicable.

11. Special storage precautions

Keep out of the sight and reach of children.

Do not store above 25 °C.
Do not freeze.
Protect from light.
Keep the vials in the outer carton.

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

Shelf life after first opening the immediate packaging: use immediately.

12. Special precautions for disposal

Medicines should not be disposed of via wastewater.

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.

Solvent only: Any unused veterinary medicinal product or waste materials derived from such veterinary medicinal product should be disposed of in accordance with local requirements.
After reconstitution: Any recommendations specified for the vaccine for which Nobivac Solvent is used will apply.

13. Classification of veterinary medicinal products

Veterinary medicinal product subject to prescription.

14. Marketing authorisation numbers and pack sizes

Vm 06376/4126

Pack sizes:
Cardboard boxes with 10 x 1 ml or 50 x 1 ml vials.
Not all pack sizes may be marketed.

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 and manufacturer responsible for batch release:
Intervet International B.V.
Wim de Körverstraat 35
5831 AN Boxmeer
The Netherlands

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

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

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

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

17. Other information

POM-V

Solvent authorised for reconstitution of:
Nobivac DHP, Nobivac DHPPi, Nobivac Parvo-C, Nobivac Pi, Nobivac Ducat, Nobivac Tricat Trio.

Gavin Hall

Approved 22 February 2026