PARTICULARS TO APPEAR ON THE OUTER PACKAGE
Carton box - 500 ml

1. NAME OF THE VETERINARY MEDICINAL PRODUCT

Nobilis REO Inac emulsion for injection

2. STATEMENT OF ACTIVE SUBSTANCES

Each dose (0.5 ml) contains:
Avian reovirus strains 1733 and 2408, inactivated inducing ≥ 7.4 $\log_2$ ELISA units

3. PACKAGE SIZE

1000 doses

4. TARGET SPECIES

Chickens (for reproduction).

5. INDICATIONS

6. ROUTES OF ADMINISTRATION

Intramuscular or subcutaneous use.

7. WITHDRAWAL PERIODS

Withdrawal period: Zero days.

8. EXPIRY DATE

Exp. {mm/yyyy}
Once broached use within 3 hours.

9. SPECIAL STORAGE PRECAUTIONS

Store in a refrigerator.
Do not freeze.
Protect from light.
Allow to reach ambient temperature before use (15 °C - 25 °C)
Shake vigorously before and periodically during use.

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.

14. MARKETING AUTHORISATION NUMBERS

Vm 06376/4133

15. BATCH NUMBER

Lot {number}

PARTICULARS TO APPEAR ON THE IMMEDIATE PACKAGE
PET or glass 500 ml

1. NAME OF THE VETERINARY MEDICINAL PRODUCT

Nobilis REO Inac emulsion for injection

2. STATEMENT OF ACTIVE SUBSTANCES

One dose (0.5 ml) contains:
Avian reovirus strains 1733 and 2408, inactivated inducing ≥ 7.4 $\log_2$ ELISA units

1000 doses

3. TARGET SPECIES

Chickens (for reproduction)

4. ROUTES OF ADMINISTRATION

Intramuscular or subcutaneous use.
Read the package leaflet before use.

5. WITHDRAWAL PERIODS

Withdrawal period: Zero days.

6. EXPIRY DATE

Exp. {mm/yyyy}

Once broached use within 3 hours.

7. SPECIAL STORAGE PRECAUTIONS

Store in a refrigerator.
Do not freeze.
Protect from light.
Allow to reach ambient temperature before use (15 °C - 25 °C). Shake vigorously before and periodically during use.

8. NAME OF THE MARKETING AUTHORISATION HOLDER

Intervet International B.V.

9. BATCH NUMBER

Lot {number}

PARTICULARS TO APPEAR ON THE PACKAGE LEAFLET:

PACKAGE LEAFLET

1. Name of the veterinary medicinal product

Nobilis REO Inac emulsion for injection for chickens

2. Composition

Each dose (0.5 ml) contains:

Active substance:
Avian reovirus strains 1733 and 2408, inactivated inducing ≥ 7.4 $\log_2$ ELISA units
Adjuvant:
Liquid paraffin

Homogenous and white to nearly white emulsion.

3. Target species

Chickens (for reproduction).

4. Indications for use

For the active immunisation of parent birds for the passive immunisation of their progeny, to reduce mortality and clinical signs of disease caused by avian reoviruses.

Onset of immunity: Active immunity develops in the parent within 4 weeks, and off spring born at any stage of the subsequent laying period will have passive immunity against reovirus infections.
Duration of immunity: Offspring born to a vaccinated parent at any stage of the subsequent laying period will have passive immunity against reovirus infections for protection during the susceptible period in the early phase of life.

5. Contraindications

None.

6. Special warnings

Special warnings:
Vaccinate healthy birds only.

Special precautions for safe use in the target species:
Not applicable.

Special precautions to be taken by the person administering the veterinary medicinal product to animals:

To the user:
This veterinary medicinal product contains mineral oil. Accidental injection/self-injection may result in severe pain and swelling, particularly if injected into a joint or finger, and in rare cases could result in the loss of the affected finger if prompt medical attention is not given. If you are accidentally injected with this veterinary medicinal product, seek prompt medical advice even if only a very small amount is injected and take the package leaflet with you. If pain persists for more than 12 hours after medical examination, seek medical advice again.

To the physician:
This veterinary medicinal product contains mineral oil. Even if small amounts have been injected, accidental injection with this veterinary medicinal product can cause intense swelling, which may, for example, result in ischaemic necrosis and even the loss of a digit. Expert, PROMPT, surgical attention is required and may necessitate early incision and irrigation of the injected area, especially where there is involvement of finger pulp or tendon.

Laying birds:
The safety of the veterinary medicinal product has not been established during lay. Do not use in birds in lay and within 4 weeks before the start of the laying period.

Interaction with other medicinal products and other forms of interaction:
Safety and efficacy data are available which demonstrate that this vaccine can be administered on the same day but not mixed with other Intervet inactivated oil emulsion vaccines, containing the avian infectious bronchitis (IB), Gumboro disease (GD), Newcastle disease (ND), turkey rhinotracheitis (TRT) and/or egg drop syndrome (EDS) antigens.

No information is available on the safety and efficacy of this vaccine when used with any other veterinary medicinal product except the products mentioned above. 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.

Overdose:
No symptoms were observed from a double dose other than those described in section 3.6.

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

7. Adverse events

Chickens (for reproduction):

Very common (>1 animal / 10 animals treated):	Injection site swelling[1].

[1] A slight transient swelling may be observed which resolves within 3 weeks and may be palpable at the site of vaccination in 50% of vaccinated birds.

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

Intramuscular or subcutaneous use.
The veterinary medicinal product should be given to birds around 10–20 weeks of age but not less than 4 weeks before the expected onset of lay.

Dose: 0.5 ml vaccine per bird by intramuscular injection into the thigh or chest muscle or by subcutaneous injection into the back of the neck, using a medium sized needle (20 g x ½”). For an optimal response in birds not primed by field virus, two vaccinations should be given approximately 6 weeks apart.

9. Advice on correct administration

Allow to reach ambient temperature before use (15 °C – 25 °C).

The vaccine may occasionally separate into two layers on storage. This in no way affects its potency, but the vaccine should be shaken vigorously before and during use to ensure a good emulsification.

An automatic injection system, incorporating a means to prevent back flushing and hence possible contamination of the vaccine, should be used to administer the vaccine.

Ensure that vaccination equipment is clean and sterile before use. Do not use vaccination equipment with rubber parts as the excipient may damage certain types of rubber.

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.
Protect from light.
Do not use this veterinary medicinal product after the expiry date which is stated on the carton. The expiry date refers to the last day of that month.
Keep the bottle in the outer carton.

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.

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

13. Classification of veterinary medicinal products

Veterinary medicinal product subject to prescription.

14. Marketing authorisation numbers and pack sizes

Vm 06376/4133

Pack size: Carton box containing 1 x bottle of 500 ml (1000 doses).

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
The Netherlands

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

Gavin Hall
Approved: 08 March 2026