

Nobilis®

NEWCAVAC

VACCINE AGAINST NEWCASTLE DISEASE IN POULTRY

**ACTIVE CONSTITUENTS**

Inactivated Newcastle Disease Virus inducing at least 4 log₂ HI units per 1/50 dose or at least 50 PD₅₀ units per dose. Virus is ND clone 30 (Beaudette's La Sota strain).

Nobilis NEWCAVAC contains one immunogenic strain of Newcastle disease virus. The virus has been grown on embryonated eggs and is inactivated with formaldehyde. Subsequently it has been suspended in the aqueous phase of an oil adjuvant emulsion.

The vaccine is recommended for the vaccination of layers and breeders for protection against sickness and death caused by Newcastle disease in early – mid lay.

DIRECTIONS FOR USE**Restraints**

Do not mix with other vaccines.

Precautions

Vaccinate healthy birds only.

Side effects**Vaccination reactions**

In healthy birds no clinical reaction to vaccination will be observed. For some weeks after vaccination a slight swelling may be felt at the site of vaccination. Local tissue reactions may occur.

Dosage and administration**Shake the bottle well before use.**

Broached vials should be used within 3 hours.

Before using the vaccine allow it to reach ambient temperature (between 15 °C and 25 °C). Use sterile injection equipment only and use the entire contents when first opened within 3 hours. Each bird should be given 0.5 mL of vaccine intramuscularly in the thigh or chest muscle or subcutaneously into the lower part of the neck.

Recommended vaccination program

Nobilis NEWCAVAC should be given to birds around 14-18 weeks of age, but not less than 4 weeks before the expected onset of lay. For an optimal effect, the birds must be primed with live vaccine against Newcastle disease.

WITHHOLDING PERIODS

Zero (0) days.

FIRST AID

If poisoning occurs, contact a doctor or Poisons Information Centre. Phone Australia 131126.

ADDITIONAL USER SAFETY INFORMATION**Take care to avoid self-injection.**

This product contains a mineral oil and is an irritant. In the event of accidental self-administration, it can cause significant pain and swelling at the injection site, perhaps also involving the draining lymph nodes. Medical or surgical intervention may be required, especially if the site of injection involves a finger joint or tendon sheath. Contact a doctor as soon as possible, even if only a very small amount is injected, and take this package leaflet/carton with you. Allow the wound to bleed freely and do not squeeze or interfere with the injection site to avoid spread of the vaccine.

Advice to the medical practitioner

This product contains mineral oil. Even if small amounts of this product have been accidentally self-administered, it can cause intense swelling and a persistent granulomatous inflammatory reaction. If injected into a finger joint or tendon sheath, the product may track along the tendon. The swelling and inflammation may compromise blood supply and result in necrosis that, in rare cases, may lead to the loss of a digit.

Following appropriate immediate local cleansing, corticosteroids may be considered to decrease the severity of any local reaction. Ascertain the patient's tetanus immunisation status, and provide booster or primary series, as appropriate.

In some cases of accidental self-injection, PROMPT surgical attention may be required. The wound should be incised and irrigated to remove the vaccine, especially where there is involvement of finger pulp or tendon. Complete curettage or total excision of the lesion may be required for chronic granulomatous reactions. Meticulous technique is required to stop inadvertent spread of the product.

Additional information is listed in the safety data sheet.

DISPOSAL

Dispose of empty container by wrapping with paper and putting in garbage. Discarded needles/sharps should be placed in a designated and appropriately labelled 'sharps' container.

STORAGE

KEEP OUT OF REACH OF CHILDREN. Store between 2 °C and 8 °C (refrigerate, do not freeze).

Warranty

Intervet Australia Pty Limited (IAPL), trading as MSD Animal Health, warrants that this product is of merchantable quality and fit for its intended purpose. IAPL's liability for any loss, including consequential losses or injury caused by act or omission, including negligent acts or omissions, by IAPL or its agent, is limited to replacing or repairing the product at the option of IAPL. If possible, a sample of any product causing concern should be retained or delivered to IAPL within 30 days for a scientific examination.

Intervet Australia Pty Limited
(trading as MSD Animal Health)
91-105 Harpin Street,
Bendigo East VIC 3550
Phone: 1800 033 461

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