

Nobilis RT+IBmulti+ND+EDS emulsion for injection for chickens

Awtorizzat

- Newcastle disease virus, strain Clone 30, Inactivated
- Egg drop syndrome '76 virus, strain BC14, Inactivated
- Turkey rhinotracheitis virus, strain BUT1#8544, Inactivated
- Avian infectious bronchitis virus, type D274/D207, strain 249g, Inactivated
- Avian infectious bronchitis virus, type Massachusetts, strain M41, Inactivated

Product identification

Isem tal-mediċina:

Nobilis RT+IBmulti+ND+EDS emulsion for injection for chickens

Nobilis RT+IBmulti+ND+EDS, Injekční emulze

Sustanza attiva:

Disponibbli biss fi [English](#)

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Speċi li fuqhom ser jintuża l-prodott:

Disponibbli biss fi [Bulgarian](#) [Spanish](#) [Czech](#) [Danish](#) [German](#) [Estonian](#) [Greek](#) [English](#)

[French](#) [Italian](#) [Latvian](#) [Lithuanian](#) [Hungarian](#) [Dutch](#) [Romanian](#) [Finnish](#) [Swedish](#)

[Icelandic](#) [Norwegian](#)

Metodu ta' amministrazzjoni:

Użu għal ġol-muskoli

Product details

Sustanza attiva / Qawwa:

Disponibbli biss fi [English](#)

2.00 haemagglutination inhibiting unit(s) / 0.50 millilitre(s)

Disponibbli biss fi [English](#)

2.70 haemagglutination inhibiting unit(s) / 0.50 millilitre(s)

Disponibbli biss fi [English](#)

3.25 enzyme-linked immunosorbent assay unit / 0.50 millilitre(s)

Disponibbli biss fi [English](#)

2.00 virus neutralising unit(s) / 0.50 millilitre(s)

Disponibbli biss fi [English](#)

2.46 virus neutralising unit(s) / 0.50 millilitre(s)

Forma farmaċewtika:

Emulsjoni għall-injezzjoni

Withdrawal period by route of administration:

Użu għal ġol-muskoli:

• Chicken

- Meat and offal. 0 day

- Egg. 0 day

Kodiċi veterinarju anatomiku terapewtiku kimiku (ATCvet):

QI01AA18

Status legali tal-provvista:

Din l-informazzjoni mhijiex disponibbli għal dan il-prodott.

Status tal-awtorizzazzjoni:

Valid

Authorised in:

Disponibbli biss fi [Spanish](#) [Czech](#) [German](#) [Estonian](#) [English](#) [French](#) [Italian](#) [Latvian](#) [Dutch](#) [Portuguese](#) [Slovak](#) [Swedish](#) [Icelandic](#) [Norwegian](#)

Deskrizzjoni tal-pakkett:

Disponibbli biss fi [English](#)

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Additional information

Entitlement type:

Disponibbli biss fi [English](#) [French](#) [Croatian](#) [Italian](#) [Latvian](#) [Finnish](#) [Swedish](#) [Icelandic](#) [Norwegian](#)

Baži ġuridika tal-awtorizzazzjoni tal-prodott:

Disponibbli biss fi [English](#) [Italian](#) [Latvian](#) [Norwegian](#)

Id-detentur tal-awtorizzazzjoni għat-tqegħid fis-suq:

Intervet International B.V.

Marketing authorisation date:

Din l-informazzjoni mhijiex disponibbli għal dan il-prodott.

Siti ta' manifattura b'rilaxx tal-lott:

INTERVET INTERNATIONAL B.V.

Awtorità responsabbli:

Institute For State Control Of Veterinary Biologicals And Medicaments

Numru tal-awtorizzazzjoni:

97/127/04-C

Data tal-bidla fl-istatus tal-awtorizzazzjoni:

30/07/2004

Stat Membru ta' referenza:

Disponibbli biss fi [Spanish](#) [Czech](#) [German](#) [Estonian](#) [English](#) [French](#) [Italian](#) [Dutch](#) [Portuguese](#) [Slovak](#) [Swedish](#) [Icelandic](#) [Norwegian](#)

Numru tal-proċedura:

DE/V/0209/001

Stati Membri Kkonċernati:

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To consult adverse reactions on veterinary medicinal products please go to www.adrreports.eu/vet

Documents

Sommarju tal-karatteristiċi tal-prodott

Dan id-dokument ma jeżistix f'dan il-lingwa (Malti). Tista 'ssibha f'lingwa oħra hawn taħt.

Fuljett ta' tagħrif

Dan id-dokument ma jeżistix f'dan il-lingwa (Malti). Tista 'ssibha f'lingwa oħra hawn taħt.

Tikkettar

Dan id-dokument ma jeżistix f'dan il-lingwa (Malti). Tista 'ssibha f'lingwa oħra hawn taħt.

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