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Nasym (--) - Lyophilisate and solvent for suspension for injection

Authorised

- Bovine respiratory syncytial virus, strain Lym-56, Live

Product identification

Medicine name:
Nasym (--) - Lyophilisate and solvent for suspension for injection

Active substance:
Bovine respiratory syncytial virus, strain Lym-56, Live

Target species:
Cattle

Route of administration:
Intramuscular use
Nasal use

Product details

Active substance and strength:
Bovine respiratory syncytial virus, strain Lym-56, Live

Presentation_strength:≥ $10^{5,1}$ $CCID_{50}$ Reference:Ph. Eur. 1177 Index:0

Pharmaceutical form:

This information is not available for this product.

Withdrawal period by route of administration:

Intramuscular use:

- **Cattle**
  - Not applicable. 0 day Zero days

Nasal use:

- **Cattle**
  - Not applicable. 0 day Zero days

Anatomical therapeutic chemical veterinary (ATCvet) codes:

QI02AD04

Legal status of supply:

Veterinary medicinal product subject to veterinary prescription

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Authorisation status:

Valid

Authorised in:

Austria , Belgium , Bulgaria , Croatia , Cyprus , Czechia , Denmark , Estonia , Finland , France , Germany , Greece , Hungary , Iceland , Ireland , Italy , Latvia , Liechtenstein , Lithuania , Luxembourg , Malta , Netherlands , Norway , Poland , Portugal , Romania , Slovakia , Slovenia , Spain , Sweden , United Kingdom (Northern Ireland)

Package description:

Packaging:Lyophilisate: vial (glass); solvent: vial (PET), Package_size:10 vials lyophilisate; 10 vials solvent, Content:5 doses lyophilisate; 10 ml solvent
Packaging:Lyophilisate: vial (glass); solvent: vial (PET), Package_size:10 vials lyophilisate; 10 vials solvent, Content:25 doses lyophilisate; 50 ml solvent
Packaging:Lyophilisate: vial (glass); solvent: vial (PET), Package_size:1 vial lyophilisate; 1 vial solvent, Content:25 doses lyophilisate; 50 ml solvent

Packaging:Lyophilisate: vial (glass); solvent: vial (PET), Package_size:1 vial lyophilisate, 1 vial solvent, Content:5 doses lyophilisate; 10 ml solvent
Packaging:Lyophilisate: vial (glass); solvent: vial (glass), Package_size:10 vials lyophilisate; 10 vials solvent, Content:1 dose lyophilisate; 2 ml solvent

Additional information

Entitlement type:
Marketing Authorisation

Legal basis of product authorisation:
Full application - Known active substance (Article 12(3) of Directive No 2001/82/EC)

Marketing authorisation holder:
Laboratorios Hipra S.A.

Marketing authorisation date:
29/07/2019

Manufacturing sites for batch release:
Laboratorios Hipra S.A.

Responsible authority:
European Commission

Authorisation number:
This information is not available for this product.

Date of authorisation status change:
29/07/2019

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

Documents

Combined File of all Documents

English (PDF)
Published on: 13/03/2026
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