Nobivac NXT HCPCh (--) + (--) + (--) + (--) - Lyophilisate and solvent for suspension for injection

Authorised

- Chlamydia felis, strain Baker, Live
- Feline panleucopenia virus, strain MW-1, Live
- Feline calicivirus, strain F9, Live
- Felid herpesvirus 1, strain G2620A, Live

Product identification

Medicine name:
Nobivac NXT HCPCh (--) + (--) + (--) + (--) - Lyophilisate and solvent for suspension for injection

Active substance:
Chlamydia felis, strain Baker, Live

Feline panleucopenia virus, strain MW-1, Live

Feline calicivirus, strain F9, Live

Felid herpesvirus 1, strain G2620A, Live

Target species:
Cat

Route of administration:
Subcutaneous use

Product details

Active substance and strength:

Chlamydia felis, strain Baker, Live

Feline panleucopenia virus, strain MW-1, Live

Feline calicivirus, strain F9, Live

Felid herpesvirus 1, strain G2620A, Live

Pharmaceutical form:

Lyophilisate and solvent for suspension for injection

Anatomical therapeutic chemical veterinary (ATCvet) codes:

QI06AF01

Legal status of supply:

Veterinary medicinal product subject to veterinary prescription

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:Vial (glass), Package_size:Lyophilisate: 5 vials + solvent: 5 vials, Content:Lyophilisate: 1 dose + solvent: 0.5 ml

Packaging:Vial (glass), Package_size:Lyophilisate: 25 vials + solvent: 25 vials, Content:Lyophilisate: 1 dose + solvent: 0.5 ml

Additional information

Entitlement type:

Marketing Authorisation

Legal basis of product authorisation:
Full application - known active substance (Article 8 of Regulation (EU) 2019/6)

Marketing authorisation holder:
Intervet International B.V.

Marketing authorisation date:
2/07/2026

Manufacturing sites for batch release:
Intervet International B.V.

Responsible authority:
European Commission

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

Date of authorisation status change:
2/07/2026

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

Documents

Combined File of all Documents

English (PDF)
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