Hepizovac (--) - Suspension for injection

Authorised

- Epizootic haemorrhagic disease virus, serotype 8, strain EHDV8 SPA 2022/LCV_03 LCV Cod.:O78, Inactivated

Product identification

Medicine name:
Hepizovac (--) - Suspension for injection

Active substance:
Epizootic haemorrhagic disease virus, serotype 8, strain EHDV8 SPA 2022/LCV_03 LCV Cod.:O78, Inactivated

Target species:
Cattle

Route of administration:
Subcutaneous use

Product details

Active substance and strength:
Epizootic haemorrhagic disease virus, serotype 8, strain EHDV8 SPA 2022/LCV_03 LCV Cod.:O78, Inactivated

Presentation_strength:4 x $10^{5.5}$ $CCID_{50}$ Reference:In-house Index:0

Pharmaceutical form:

Suspension for injection

Withdrawal period by route of administration:

Subcutaneous use:

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

Anatomical therapeutic chemical veterinary (ATCvet) codes:

QI02AA

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:Bottle (HDPE), Package_size:1 bottle, Content:100 mL

Packaging:Bottle (HDPE), Package_size:1 bottle, Content:252 mL

Packaging:Bottle (HDPE), Package_size:1 bottle, Content:52 mL

Additional information

Entitlement type:

Marketing Authorisation

Legal basis of product authorisation:

Applications in exceptional circumstances (Article 25 of Regulation (EU) 2019/6)

Marketing authorisation holder:
CZ Vaccines S.A.U.

Marketing authorisation date:
23/04/2025

Manufacturing sites for batch release:
CZ Vaccines S.A.U.

Responsible authority:
European Commission

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

Date of authorisation status change:
23/04/2025

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: 18/06/2025
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