DIFTOSEC CT lyofilizát a rozpúšťadlo na injekčnú suspenziu pre kurčatá a morky

Authorised

- Fowlpox virus, strain DECP 25, Live

Product identification

Medicine name:
DIFTOSEC CT lyofilizát a rozpúšťadlo na injekčnú suspenziu pre kurčatá a morky

Active substance:
Fowlpox virus, strain DECP 25, Live

Target species:
Chicken (broiler)
Chicken (pullet future breeder)
Chicken (pullet for egg production, future layer)

Route of administration:
Skin scarification
Transdermal use

Product details

Active substance and strength:
Fowlpox virus, strain DECP 25, Live

3.00 log10 50% cell culture infectious dose / 0.01 millilitre(s)

Pharmaceutical form:

Lyophilisate and solvent for suspension for injection

Withdrawal period by route of administration:

Skin scarification:

- **Chicken (broiler)**
  - All relevant tissues. 0 day zero days
- **Chicken (pullet future breeder)**
  - All relevant tissues. 0 day zero days
- **Chicken (pullet for egg production, future layer)**
  - All relevant tissues. 0 day zero days

Transdermal use:

- **Chicken (broiler)**
  - All relevant tissues. 0 day zero days
- **Chicken (pullet future breeder)**
  - All relevant tissues. 0 day zero days
- **Chicken (pullet for egg production, future layer)**
  - All relevant tissues. 0 day zero days

Anatomical therapeutic chemical veterinary (ATCvet) codes:

QI01AD12

Legal status of supply:

Veterinary medicinal product subject to veterinary prescription

Authorisation status:
Valid

Authorised in:
Slovakia

Package description:
Available only in Slovak
Available only in Slovak
Available only in Slovak

Additional information

Entitlement type:
Marketing Authorisation

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

Marketing authorisation holder:
Boehringer Ingelheim Animal Health France

Marketing authorisation date:
22/05/1998

Manufacturing sites for batch release:
Boehringer Ingelheim Animal Health France

Responsible authority:
Institute For State Control Of Veterinary Biologicals And Medicaments

Authorisation number:
97/0019/98-S

Date of authorisation status change:
22/05/1998

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

Documents

Combined File of all Documents

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