

PARVOKAN LYOPHILISATE AND SUSPENSION FOR SUSPENSION FOR INJECTION

Authorised

- Derzsy's disease virus, strain H, Live
- Muscovy duck parvovirus, strain GM, Inactivated

Product identification

Medicine name:

PARVOKAN LYOPHILISATE AND SUSPENSION FOR SUSPENSION FOR INJECTION

Active substance:

Derzsy's disease virus, strain H, Live

Muscovy duck parvovirus, strain GM, Inactivated

Target species:

Duck

Route of administration:

Subcutaneous use

Product details

Active substance and strength:

Derzsy's disease virus, strain H, Live

2.50 log₁₀ 50% cell culture infectious dose / 1.00 Dose

Muscovy duck parvovirus, strain GM, Inactivated
1.50 log₁₀ serum neutralising unit(s) / 1.00 Dose

Pharmaceutical form:

Lyophilisate and suspension for suspension for injection

Withdrawal period by route of administration:

Subcutaneous use:

-

Duck

- All relevant tissues. 0 day

Anatomical therapeutic chemical veterinary (ATCvet) codes:

QI01BH01

Legal status of supply:

Veterinary medicinal product subject to veterinary prescription

Authorisation status:

Valid

Authorised in:

Italy

Package description:

Box of one 500-dose bottle of suspension and one 500-dose bottle of freeze-dried pellet

Box of ten 1,500-dose bottles of suspension and thirty 500-dose bottles of freeze-dried pellet

Box of ten 500-dose bottles of suspension and ten 500-dose bottles of freeze-dried pellet

Box of one 1,500-dose bottle of suspension and three 500-dose bottles of freeze-dried pellet

Additional information

Entitlement type:

Marketing Authorisation

Legal basis of product authorisation:

Full application - New active substance (Article 12(3) of Directive No 2001/82/EC)

Marketing authorisation holder:

Boehringer Ingelheim Animal Health Italia S.p.A. In Breve Boehringer Ingelheim Ah It S.p.A.

Marketing authorisation date:

6/05/2009

Manufacturing sites for batch release:

Boehringer Ingelheim Animal Health France

Responsible authority:

Ministry Of Health

Authorisation number:

104060

Date of authorisation status change:

6/05/2009

Reference member state:

France

Procedure number:

FR/V/0144/001

Concerned member states:

Germany Italy

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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