

# 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

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**Active substance:**

Derzsy's disease virus, strain H, Live

Muscovy duck parvovirus, strain GM, Inactivated

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**Target species:**

Duck

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**Route of administration:**

Subcutaneous use

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## Product details

**Active substance and strength:**

Derzsy's disease virus, strain H, Live

2.50 log<sub>10</sub> 50% cell culture infectious dose / 1.00 Dose

Muscovy duck parvovirus, strain GM, Inactivated  
1.50 log<sub>10</sub> serum neutralising unit(s) / 1.00 Dose

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**Pharmaceutical form:**

Lyophilisate and suspension for suspension for injection

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**Withdrawal period by route of administration:**

**Subcutaneous use:**

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**Duck**

- All relevant tissues. 0 day

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**Anatomical therapeutic chemical veterinary (ATCvet) codes:**

QI01BH01

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**Legal status of supply:**

Veterinary medicinal product subject to veterinary prescription

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**Authorisation status:**

Valid

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**Authorised in:**

France

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**Available in:**

France

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**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

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## Additional information

**Entitlement type:**

Marketing Authorisation

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**Legal basis of product authorisation:**

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

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**Marketing authorisation holder:**

Boehringer Ingelheim Animal Health France

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**Marketing authorisation date:**

16/05/2002

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**Manufacturing sites for batch release:**

Boehringer Ingelheim Animal Health France

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**Responsible authority:**

French Agency For Food, Environmental And Occupational Health & Safety

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**Authorisation number:**

FR/V/5823169 2/2002

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**Date of authorisation status change:**

16/05/2007

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**Reference member state:**

France

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**Procedure number:**

FR/V/0144/001

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**Concerned member states:**

Germany Italy

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To consult adverse reactions on veterinary medicinal products please go to [www.adrreports.eu/vet](http://www.adrreports.eu/vet)

## Documents

### Summary of Product Characteristics

This document does not exist in this language (English). You can find it in another language below.

### Package Leaflet and Labelling

This document does not exist in this language (English). You can find it in another language below.