Source URL: https://medicines.health.europa.eu/veterinary/en/600000028013

ByeMite 500 mg/ml concentrate for spraying emulsion for laying hens

Authorised

Phoxim

Product identification

Medicine name:

BYEMITE 500 MG/ML CONCENTRATE FOR SPRAYING EMULSION FOR LAYING HENS ByeMite 500 mg/ml concentrate for spraying emulsion for laying hens

Active substance:

Phoxim

Target species:

Chicken (layer hen)

Route of administration:

Cutaneous use

Product details

Active substance and strength:

Phoxim

500.00 milligram(s) / 1.00 millilitre(s)

Pharmaceutical form:

Concentrate for spray emulsion

Withdrawal period by route of administration:

Cutaneous use:

Chicken (layer hen)

- Meat and offal. 25 day

Remove eggs before treatment. Discard eggs laid during and on the same day after the treatment. Meat and offal: 25 days after the second treatment.

- Eggs. 12 hour

Remove eggs before treatment. Discard eggs laid during and on the same day after the treatment.

Anatomical therapeutic chemical veterinary (ATCvet) codes:

OP53AF01

Legal status of supply:

Veterinary medicinal product subject to veterinary prescription

Authorisation status:

Valid

Authorised in:

Ireland

Available in:

Ireland

Package description:

Available only in French

Available only in French

Available only in French

Additional information

Entitlement type:

Marketing Authorisation

Legal basis of product authorisation:

Generic application (Article 13(1) of Directive No 2001/82/EC)

Marketing authorisation holder:

Elanco GmbH

Marketing authorisation date:

20/02/2009

Manufacturing sites for batch release:

KVP Pharma+Veterinaer Produkte GmbH

Responsible authority:

Health Products Regulatory Authority

Authorisation number:

VPA22020/049/001

Date of authorisation status change:

20/02/2009

Reference member state:

France

Procedure number:

FR/V/0196/001

Concerned member states:

Austria Belgium Bulgaria Cyprus Czechia Finland Greece Hungary Iceland Ireland Italy Luxembourg Netherlands Portugal Romania Slovakia Slovenia Sweden

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