

# 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

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**Active substance:**

Phoxim

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**Target species:**

Chicken (layer hen)

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**Route of administration:**

Cutaneous use

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## Product details

**Active substance and strength:**

Phoxim

500.00 milligram(s) / 1.00 millilitre(s)

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**Pharmaceutical form:**

Concentrate for spray emulsion

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**Withdrawal period by route of administration:**

**Cutaneous use:**

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**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.

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

QP53AF01

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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:**

Available only in [French](#)

Available only in [French](#)

Available only in [French](#)

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## Additional information

**Entitlement type:**

Marketing Authorisation

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**Legal basis of product authorisation:**

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

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**Marketing authorisation holder:**

Elanco GmbH

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**Marketing authorisation date:**

24/04/2009

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**Manufacturing sites for batch release:**

KVP Pharma+Veterinaer Produkte GmbH

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**Responsible authority:**

French Agency For Food, Environmental And Occupational Health & Safety

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**Authorisation number:**

FR/V/2385846 0/2009

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**Date of authorisation status change:**

24/04/2014

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**Reference member state:**

France

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**Procedure number:**

FR/V/0196/001

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**Concerned member states:**

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

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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.