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

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:
QP53AF01

Legal status of supply:
Veterinary medicinal product subject to veterinary prescription

Authorisation status:
Valid

Authorised in:
Italy

Available in:
Italy

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 Italia S.p.A.

Marketing authorisation date:
25/05/2009

Manufacturing sites for batch release:
KVP Pharma+Veterinaer Produkte GmbH

Responsible authority:
Ministry Of Health

Authorisation number:
104053

Date of authorisation status change:
25/05/2009

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

Documents

Combined File of all Documents

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