

# HYOGEN EMULSION FOR INJECTION

Authorised

- *Mycoplasma hyopneumoniae*, strain 2940, Inactivated

## Product identification

**Medicine name:**

HYOGEN EMULSION FOR INJECTION

Hyogen emulsija injekcijām cūkām

**Active substance:**

*Mycoplasma hyopneumoniae*, strain 2940, Inactivated

**Target species:**

Pig (for fattening)

**Route of administration:**

Intramuscular use

## Product details

**Active substance and strength:**

*Mycoplasma hyopneumoniae*, strain 2940, Inactivated

328.00 enzyme-linked immunosorbent assay unit / 2.00 millilitre(s)

**Pharmaceutical form:**

Emulsion for injection

**Withdrawal period by route of administration:****Intramuscular use:**

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**Pig (for fattening)**

- All relevant tissues. 0 day

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

QI09AB13

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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:**

Latvia

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**Package description:**

Cardboard box of 1 bottle of 50 ml (1x25 doses)

Cardboard box of 5 bottles of 250 ml (5x125 doses)

Cardboard box of 1 bottle of 250 ml (1x125 doses)

Cardboard box of 5x200 ml in 5 bottles of 250 ml (5x100 doses)

Cardboard box of 200 ml in 1 bottle of 250 ml (1x100 doses)

Cardboard box of 5 bottles of 200 ml (5x100 doses)

Cardboard box of 1 bottle of 200 ml (1x100 doses)

Cardboard box of 5 bottles of 100 ml (5x50 doses)

Cardboard box of 1 bottle of 100 ml (1x50 doses)

Cardboard box of 5 bottles of 50 ml (5x25 doses)

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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:**

Ceva Sante Animale

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**Marketing authorisation date:**

27/04/2015

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**Manufacturing sites for batch release:**

Ceva-Phylaxia Zrt.

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**Responsible authority:**

Food And Veterinary Service

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**Authorisation number:**

V/DCP/15/0017

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**Date of authorisation status change:**

27/04/2015

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**Reference member state:**

France

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**Procedure number:**

FR/V/0278/001

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**Concerned member states:**

Austria Belgium Bulgaria Croatia Cyprus Czechia Denmark Estonia  
Germany Greece Hungary Ireland Italy Latvia Lithuania Netherlands Poland  
Portugal Romania Slovakia Slovenia Spain Sweden  
United Kingdom (Northern Ireland)

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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

Combined File of all Documents

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