

# Oxybee 39.4 mg/ml - Powder and solution for bee-hive dispersion

Authorised

- Oxalic acid dihydrate

## Product identification

**Medicine name:**

Oxybee 39.4 mg/ml - Powder and solution for bee-hive dispersion

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**Active substance:**

Oxalic acid dihydrate

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**Target species:**

Honey bee

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**Route of administration:**

In-hive use

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## Product details

**Active substance and strength:**

Oxalic acid dihydrate

34.90 milligram(s) / 1.00 Bottle

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**Pharmaceutical form:**

Powder and solution for bee-hive dispersion

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**Withdrawal period by route of administration:****In-hive use:**

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**Honey bee**

- Honey. 0 day  
Zero days

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

QP53AG03

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**Legal status of supply:**

Veterinary medicinal product not subject to veterinary prescription

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**Authorisation status:**

Valid

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**Authorised in:**

Austria , Belgium , Bulgaria , Croatia , Cyprus , Czechia , Denmark , Estonia , Finland , France , Germany , Greece , Hungary , Iceland , Ireland , Italy , Latvia , Liechtenstein , Lithuania , Luxembourg , Malta , Netherlands , Norway , Poland , Portugal , Romania , Slovakia , Slovenia , Spain , Sweden , United Kingdom (Northern Ireland)

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**Package description:**

Packaging: Bottle (HDPE); sachet (LDPE/Alu/LDPE/clay coated paper), Package\_size: 1 bottle + 1 sachet, Content: 375 g (500 ml bottle); 125 g sachet

Packaging: Bottle (HDPE); sachet (LDPE/Alu/LDPE/clay coated paper), Package\_size: 1 bottle + 2 sachets, Content: 750 g (1000 ml bottle); 125 g sachet

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## Additional information

**Entitlement type:**

Marketing Authorisation

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**Legal basis of product authorisation:**

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

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**Marketing authorisation holder:**

Dany Bienenwohl GmbH

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**Marketing authorisation date:**

1/02/2018

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**Manufacturing sites for batch release:**

Wirtschaftsgenossenschaft deutscher Tieraerzte eG

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**Responsible authority:**

European Commission

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**Authorisation number:**

This information is not available for this product.

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**Date of authorisation status change:**

1/02/2018

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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

English (PDF)

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