# Fluboral 200 mg/ml, suspension for use in drinking water for pigs and chickens

Pooblaščeno

- Flubendazole

## Product identification

**Ime zdravila:**
Fluboral 200 mg/ml, suspension for use in drinking water for pigs and chickens
Fluboral 200 mg/ml suspension til anvendelse i drikkevand

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**Učinkovina:**
Na voljo samo v English

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**Ciljne živalske vrste:**
Na voljo samo v Bulgarian Spanish Czech Danish German Estonian Greek English French Italian Latvian Lithuanian Hungarian Dutch Romanian Finnish Swedish Icelandic Norwegian
Na voljo samo v Bulgarian Spanish Czech Danish German Estonian Greek English French Italian Latvian Lithuanian Hungarian Dutch Romanian Finnish Swedish Icelandic Norwegian

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**Pot uporabe:**
Dajanje v vodo za pitje

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## Product details

**Učinkovina / Jakost:**

Na voljo samo v English
200.00 milligram(s) / 1.00 millilitre(s)

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**Farmacevtska oblika:**

Suspenzija za dajanje v vodo za pitje

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**Withdrawal period by route of administration:**

**Dajanje v vodo za pitje:**

- **Chicken**
  - Eggs. 0 day
  - Meat and offal. 2 day
- **Pig**
  - Meat and offal. 4 day 1 mg / kg for 5 days.
  - Meat and offal. 5 day 2.5 mg / kg for 2 days.

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**Anatomsko-terapevtsko-kemična veterinarska oznaka (ATCvet):**

QP52AC12

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**Pravni status za dobavo / izdajo zdravila:**

Zdravilo, ki se izdaja na veterinarski recept

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**Status dovoljenja:**

Valid

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**Authorised in:**

Na voljo samo v Spanish Czech German Estonian English French Italian Dutch Portuguese Slovak Swedish Icelandic Norwegian

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**Opis ovojnine:**

Na voljo samo v English

Na voljo samo v English

Na voljo samo v English

## Additional information

**Entitlement type:**
Na voljo samo v English French Croatian Italian Latvian Finnish Swedish Icelandic Norwegian

**Pravna podlaga za izdajo dovoljenja za promet z zdravilom:**
Na voljo samo v English Italian Latvian Norwegian

**Imetnik dovoljenja za promet z zdravilom:**
Dechra Regulatory B.V.

**Marketing authorisation date:**
27/01/2023

**Proizvodna mesta, odgovorna za sproščanje serij:**
Genera d.d.

**Pristojni organ:**
Danish Medicines Agency

**Številka dovoljenja :**
67072

**Datum spremembe statusa dovoljenja:**
27/01/2023

**Referenčna država članica:**
Na voljo samo v Spanish Czech German Estonian English French Italian Dutch Portuguese Slovak Swedish Icelandic Norwegian

**Številka postopka:**
IE/V/0664/001

**Zadevne države članice:**

Na voljo samo v Spanish Czech German Estonian English French Italian Dutch Portuguese Slovak Swedish Icelandic Norwegian
Na voljo samo v Spanish Czech German Estonian English French Italian Dutch Portuguese Slovak Swedish Icelandic Norwegian
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Za poročila o domnevnih neželenih učinkih zdravil za uporabo v veterinaski medicini obiščite stran: www.adrreports.eu/vet

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