# MICOSPECTONE 166,5 mg + 333,5 mg prášok na perorálny roztok

Geautoriseerd

- Lincomycin hydrochloride
- Streptomycin hydrochloride

## Product identification

**Naam van het geneesmiddel:**
MICOSPECTONE 166,5 mg + 333,5 mg prášok na perorálny roztok

**Werkzame stof:**
Alleen beschikbaar in English
Alleen beschikbaar in English

**Doeldiersoort(en):**
Biggen
Kip

**Toedieningsweg:**
Alleen beschikbaar in Bulgarian Spanish Czech Danish German Estonian Greek English French Irish Croatian Italian Latvian Polish Portuguese Slovak Slovenian Finnish Swedish Icelandic Norwegian

## Product details

**Werkzame stof / Sterkte:**
Alleen beschikbaar in English

205.23 milligram(s) / 1.00 gram(s)

Alleen beschikbaar in English
497.11 milligram(s) / 1.00 gram(s)

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**Farmaceutische vorm:**

Poeder voor drank

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**Withdrawal period by route of administration:**

**In drinking water use:**

- **Biggen**

  - Meat and offal. 0 day

Meat and offal zero days. Animals must not be killed for human consumption during treatment

- **Kip**

  - Meat and offal. 2 day

The medicinal product is not authorized for use in birds producing eggs for human consumption, including replacement chickens intended for the production of eggs for human consumption. Animals must not be killed for human consumption during treatment.

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**Anatomisch therapeutisch chemische veterinaire classificatie (ATCvet code):**

QJ01FF52

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**Afleverstatus:**

Alleen beschikbaar in Czech Estonian English French Italian Latvian Portuguese Slovenian Finnish Swedish Icelandic Norwegian

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**Status toelating:**

Valid

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**Authorised in:**

Slowakije

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**Package description:**

Alleen beschikbaar in Slovak

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## Additional information

**Entitlement type:**
Alleen beschikbaar in English French Croatian Italian Latvian Finnish Swedish Icelandic Norwegian

**Rechtsgrondslag productvergunning:**
Alleen beschikbaar in English French Italian Latvian Norwegian

**Handelsvergunninghouder:**
Fatro S.p.A.

**Marketing authorisation date:**
21/12/1999

**Productielocaties partijvrijgifte:**
Fatro S.p.A.

**Verantwoordelijke instantie:**
USKVBL

**Toelatingsnummer:**
96/128/99-S

**Wijzigingsdatum status toelating:**
21/12/1999

To consult adverse reactions on veterinary medicinal products please go to www.adrreports.eu/vet

## Documents

Combined File of all Documents

Dit document bestaat niet in deze taal (Nederlands). Je kunt het hieronder vinden in een andere taal.

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