File downloaded on 2026-11-26
**Source URL:** https://medicines.health.europa.eu/veterinary/en/600000033173

# Salmoporc, Lyophilisate and Solvent for Suspension for Injection for Pigs

Ima dovoljenje za promet

- Salmonella enterica, subsp. enterica, serovar Typhimurium, strain 421/125 (histidine-adenine auxotrophic), Live

## Informacije o zdravilu

**Ime zdravila:**
Salmoporc, Lyophilisate and Solvent for Suspension for Injection for Pigs

**Učinkovina:**
Na voljo samo v English

**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

**Pot uporabe:**
Peroralna uporaba
Subkutana uporaba

# Podatki o zdravilu

**Učinkovina / Jakost:**
Na voljo samo v English
8.00 log10 colony forming unit(s) / 1.00 millilitre(s)

---

**Farmacevtska oblika:**
Liofilizat za suspenzijo za injiciranje

---

**Karenca glede na pot uporabe:**
**Peroralna uporaba:**

- **Pig**
  - Meat and offal. 6 week — six weeks after the second vaccination

**Subkutana uporaba:**

- **Pig**
  - Meat and offal. 6 week — six weeks after the second vaccination

---

**Anatomsko-terapevtsko-kemična veterinarska oznaka klasifikacija zdravil, ki se uporabljajo v veterini (ATCvet):**
QI09AE02

---

**Pravni status za dobavo/izdajo zdravila:**
Zdravilo, ki se izdaja na veterinarski recept

---

**Status dovoljenja za promet z zdravilom:**
Valid

---

**Ima dovoljenje za promet v:**
Na voljo samo v Estonian English French Lithuanian Portuguese Swedish Icelandic Norwegian

---

**Na voljo v:**
United Kingdom (Northern Ireland)

---

**Opis ovojnine zdravila:**
Na voljo samo v English
Na voljo samo v English

## Dodatne informacije

**Vrsta dovoljenja:**
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:**
Ceva Sante Animale

**Datum dovoljenja za promet z zdravilom:**
18/04/2019

**Proizvodna mesta, odgovorna za sproščanje serij:**
Ceva-Phylaxia Zrt.
IDT Biologika GmbH

**Pristojni organ:**
The Veterinary Medicines Directorate

**Številka dovoljenja za promet z zdravilom:**
Vm 14966/3050

**Datum spremembe statusa dovoljenja za promet:**
27/06/2024

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

**Številka postopka:**
NL/V/0247/001

**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

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

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

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

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 Lithuanian Dutch Portuguese Romanian Slovak Swedish Icelandic Norwegian

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

Na voljo samo v Estonian English French Lithuanian Portuguese Swedish Icelandic Norwegian

---

Za poročila o domnevnih neželenih učinkih zdravil za uporabo v veterinaski medicini obiščite stran: www.adrreports.eu/vet