

S.E.T.-VAC -inaktywowany szczep Salmonella enteritidis, nie mniej niż 1×10^9 CFU,-inaktywowany szczep Salmonella typhimurium, nie mniej niż 1×10^9 CFU. Emulsja do wstrzykiwań

Authorised

- Salmonella enterica, subsp. enterica, serovar Enteritidis, Inactivated
- Salmonella enterica, subsp. enterica, serovar Typhimurium, Inactivated

Product identification

Medicine name:

S.E.T.-VAC -inaktywowany szczep Salmonella enteritidis, nie mniej niż 1×10^9 CFU,-inaktywowany szczep Salmonella typhimurium, nie mniej niż 1×10^9 CFU. Emulsja do wstrzykiwań

Active substance:

Salmonella enterica, subsp. enterica, serovar Enteritidis, Inactivated

Salmonella enterica, subsp. enterica, serovar Typhimurium, Inactivated

Target species:

Chicken (hen)

Route of administration:

Subcutaneous use

Product details

Active substance and strength:

Salmonella enterica, subsp. enterica, serovar Enteritidis, Inactivated
1000000000.00 Colony forming unit / 1.00 Colony forming unit

Salmonella enterica, subsp. enterica, serovar Typhimurium, Inactivated
1000000000.00 Colony forming unit / 1.00 Colony forming unit

Pharmaceutical form:

Emulsion for injection/infusion

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

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Chicken (hen)

- All relevant tissues. 0 day

Anatomical therapeutic chemical veterinary (ATCvet) codes:

QI01AB01

Legal status of supply:

Veterinary medicinal product subject to veterinary prescription

Authorisation status:

Valid

Authorised in:

Poland

Package description:

Available only in Polish

Available only in Polish

Additional information

Entitlement type:

Marketing Authorisation

Legal basis of product authorisation:

Full application (Article 12(3) of Directive No 2001/82/EC)

Marketing authorisation holder:

Fatro S.p.A.

Marketing authorisation date:

29/04/2004

Manufacturing sites for batch release:

Fatro S.p.A.

Responsible authority:

Office For Registration Of Medicinal Products Medical Devices And Biocidal Products

Authorisation number:

1477

Date of authorisation status change:

29/04/2004

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