



SAFETY DATA SHEET

Creation Date: 17-APR-2018

Revision Date: 03-DEC-2018

Version: 003

Page 1 of 5

1. IDENTIFICATION OF THE SUBSTANCE OR MIXTURE AND THE MANUFACTURER

Product Identifier ANTHRAX SPORE VACCINE, Nonencapsulated Live Culture
Trade Name None
Recommended Use Veterinary Vaccine
Manufacturer Colorado Serum Company
4950 York Street – P.O. Box 16428
Denver, CO 80216
Emergency Phone No. 303-295-7527

2. HAZARDS IDENTIFICATION

Hazard Classification Not classified
Hazard Statements Not classified
Signal Word Not classified
Other Hazards May cause allergic reaction with skin exposure
Unknown acute toxicity None known
Precautionary Statements This is an animal vaccine for veterinary use. Use only as recommended.

3. COMPOSITION / INFORMATION ON INGREDIENTS

CATEGORY	COMMON NAME	CAS NO.	%
CHEMICAL NAME			
Sodium Chloride (NaCl)	Salt	7647-14-5	*
Glycerine (C ₃ H ₈ O ₃)	Glycerol, (CP)	56-81-5	*
Saponin (C ₅₈ H ₉₄ O ₂₇)	N/A	8047-15-2	*
Dihydrogen Monoxide (H ₂ O)	Purified Water	7789-20-0	*
MIXTURE			
Anthrax Spore Vaccine, Nonencapsulated Live Culture	Anthrax spore Vaccine	N/A	*

*The components are not hazardous or are below required disclosure limits. The percent of each ingredient is proprietary.

SAFETY DATA SHEET

ANTHRAX SPORE VACCINE, Nonencapsulated Live Culture

Colorado Serum Company

Creation Date: 17-APR-2018

Revision Date: 03-DEC-2018

Version: 003

Page 2 of 5

4. FIRST AID MEASURES

Necessary First-Aid Instructions: Anthrax Spore Vaccine, Nonencapsulated Live Culture

This strain of *Bacillus anthracis* (Sterne 34F2) is not known to be pathogenic for humans. However, it is advised that it be handled carefully to avoid human exposure.

In the event of human exposure, contact a physician for medical advice /attention regarding any exposure as indicated:

Accidental injection:	Contact a physician. Local reaction at injection site (swelling) may occur. Infection at injection site may occur from contaminated equipment.
Eye Contact:	Immediately flush eyes with water for at least 10 minutes.
Skin Contact:	Wash skin with soap and water.
Ingestion:	Contact a physician.
Inhalation:	Contact a physician.

Important Symptoms or Effects - Acute or Delayed: Exposure has potential for allergic reaction.

Recommendations for Immediate Care and Special Treatment: Contact a physician if injected or ingested. Treat symptomatically.

General Information: This is an animal vaccine. NOT FOR HUMAN USE.

5. FIRE FIGHTING MEASURES

Suitable Extinguishing Equipment: Water, Carbon dioxide, Dry Chemical

Unsuitable Extinguishing Equipment: Not known

Specific Hazards Arising from Substance or Mixture: Avoid causing aerosol.

Fire Fighting: Special Protective Equipment: Firefighters: Wear appropriate personal protective equipment as conditions warrant, which may include self-contained breathing apparatus. Avoid aerosols and environmental release.

6. ACCIDENTAL RELEASE MEASURES

Personal Precautions: Use as directed. Avoid contact with exposed skin and eyes and clothing. It is an aqueous suspension not expected to support combustion.

Protective Equipment: Use personal protective clothing, gloves and eye/face shield as indicated.

Emergency Procedures: None

Containment Methods: Isolate a spill and apply hypochlorite solution (10 % bleach) or equivalent disinfectant.

Clean Up Procedures: Utilize absorbent material to soak up spill. Precautionary recommendations: burn, incinerate, autoclave or boil materials which contacted the spill. For disposal, see Section 13 of this SDS. Avoid release to the environment.

SAFETY DATA SHEET

ANTHRAX SPORE VACCINE, Nonencapsulated Live Culture

Colorado Serum Company

Creation Date: 17-APR-2018

Revision Date: 03-DEC-2018

Version: 003

Page 3 of 5

7. HANDLING AND STORAGE

Precautions for Safe Handling: Avoid contact with skin, eyes and clothing. Use only with adequate ventilation. Wash hands thoroughly after handling. Avoid breathing mists or aerosols of this product. Use appropriate personal protective equipment and avoid accidental injection.

Storage Conditions: Store only in unopened containers. Store at 2 to 8° C.

8. EXPOSURE CONTROLS / PERSONAL PROTECTION

OSHA Permissible Exposure Limits (PEL): Please refer to other sections herein regarding Anthrax Spore vaccine, the active live component of this vaccine. Exposure limits for other component(s) are not noted.

American Conference of Governmental Industrial Hygienists (ACGIH): Not known

Threshold Limit Values (TLV): Not known

Engineering Controls: Not known

Personal Protective Measures: Safety glasses with side shields, or face shield, rubber/latex/vinyl gloves. Wash hands after handling. Disinfect any equipment in contact with vaccine.

9. PHYSICAL AND CHEMICAL PROPERTIES

Appearance/Physical State:	Colorless to light yellow or tan liquid in sealed container
Odor:	No data available
Odor Threshold:	No data available
Vapor Density:	No data available
PH:	No data available
Melting/Freezing Points:	No data available
Solubility:	No data available
Initial Boiling Point & Range:	No data available
Flash Point:	No data available
Evaporation Rate:	No data available
Flammability (solid, gas):	No data available
Flammability/Explosive limits:	No data available
Vapor Pressure/Vapor Density:	No data available
Relative Density:	No data available
Solubility:	No data available
Partition Coefficient: n-octanol/water	No data available
Auto-ignition Temperature:	No data available
Decomposition Temperature:	No data available
Viscosity:	No data available

SAFETY DATA SHEET

ANTHRAX SPORE VACCINE, Nonencapsulated Live Culture

Colorado Serum Company

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Revision Date: 03-DEC-2018

Version: 003

Page 4 of 5

10. STABILITY AND REACTIVITY

Reactivity: Product is stable and non-reactive when stored according to label instructions. No other data is available.

Chemical Stability: Product is stable under normal conditions.

Oxidizing Properties: No data available

Conditions to Avoid: Store at 2 to 8° C. Prolonged exposure to higher temperatures may adversely affect potency. Do not freeze.

Incompatible Materials: Do not combine with any other material. Use only as directed.

Hazardous Decomposition Products: None expected

11. TOXICOLOGICAL INFORMATION

Likely Routes of Exposure: Accidental exposure is most likely to be topical or needle stick.

Physical, Chemical and Toxicological Symptoms: No data available

Delayed and Immediate Effects of Exposure: No data available

Acute Toxicity Estimates: No data available

Carcinogenic Properties: No data available

12. ECOLOGICAL INFORMATION

Avoid product being introduced into the environment.

13. DISPOSAL CONSIDERATIONS

Follow disposal instructions on the product label.

14. TRANSPORT INFORMATION

Not required information

15. REGULATORY INFORMATION

Not required information

SAFETY DATA SHEET

ANTHRAX SPORE VACCINE, Nonencapsulated Live Culture

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Revision Date: 03-DEC-18

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Page 5 of 5

16. OTHER INFORMATION

SDS Code: Not applicable

SDS Preparation Date: 17-APR-2018

Latest Revision Date: 03-DEC-18

Reasons for Revision: Not relevant

Data Sources: Information included in this SDS may have been gathered from confidential internal sources, raw material suppliers, or from published literature.

Prepared by: Colorado Serum Company Regulatory Department

Colorado Serum Company believes that the information contained in this Safety Data Sheet is accurate and provided in good faith. It is without warranty of any kind, expressed or implied. If data for a hazard is not included in this document, there is no known information at this time.

END OF SAFETY DATA SHEET