

SPINAL SARCODE- cartilago suis, cerebellum (suis), discus intervertebralis (suis), medulla ossis suis spray
Nutritional Specialties, Inc.

Disclaimer: This homeopathic product has not been evaluated by the Food and Drug Administration for safety or efficacy. FDA is not aware of scientific evidence to support homeopathy as effective.

DRUG FACTS:

ACTIVE INGREDIENTS:

Cartilago Suis 6X, 12X, 30X, 60X, 100X, Cerebellum 6X, 12X, 30X, 60X, 100X, Discus Intervertebralis 6X, 12X, 30X, 60X, 100X, Medulla Ossis Suis 6X, 12X, 30X, 60X, 100X.

PURPOSE:

Provides specific sarcode support for the spinal nerves.†

†Claims based on traditional homeopathic practice, not accepted medical evidence. Not FDA evaluated.

WARNINGS:

Professional Use Only

If pregnant or breast-feeding, ask a health professional before use.

In case of overdose, get medical help or contact a Poison Control Center right away.

If condition worsens, seek medical attention.

KEEP OUT OF REACH OF CHILDREN

Do not use if tamper evident seal is broken or missing.

Store in a cool place after opening

KEEP OUT OF REACH OF CHILDREN:

In case of overdose, get medical help or contact a Poison Control Center right away.

DIRECTIONS:

Adults: 2 to 3 sprays orally, three times daily. Children under twelve one half adult dosage. Do not take within 15 minutes of consuming food, beverage or brushing teeth. Consult a physician for use in children under 12 years of age.

INDICATIONS:

Provides specific sarcode support for the spinal nerves.†

†Claims based on traditional homeopathic practice, not accepted medical evidence. Not FDA evaluated.

INACTIVE INGREDIENTS:

Alcohol USP 20%, Purified Water USP.

QUESTIONS:

MANUFACTURED EXCLUSIVELY FOR

NUTRITIONAL SPECIALTIES, INC.

PO BOX 97227

PITTSBURGH, PA 15229

www.phpltd.com

PACKAGE LABEL DISPLAY:

Professional

Health Products

HOMEOPATHIC

NDC 83027-0082-1

SPINAL

SARCODE

2 FL. OZ (60 ml)

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**SPINAL
SARCODE**

HSD(reorder code)

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SPINAL SARCODE

cartilago suis, cerebellum (suis), discus intervertebralis (suis), medulla ossis suis spray

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:83027-0082
Route of Administration	ORAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
SUS SCROFA CARTILAGE (UNII: 73ECW5WG2F) (SUS SCROFA CARTILAGE - UNII:73ECW5WG2F)	SUS SCROFA CARTILAGE	6 [hp_X] in 1 mL
SUS SCROFA CEREBELLUM (UNII: 49NGK53TPQ) (SUS SCROFA CEREBELLUM - UNII:49NGK53TPQ)	SUS SCROFA CEREBELLUM	6 [hp_X] in 1 mL
SUS SCROFA INTERVERTEBRAL DISC (UNII: OJ17O2WTSM) (SUS SCROFA INTERVERTEBRAL DISC - UNII:OJ17O2WTSM)	SUS SCROFA INTERVERTEBRAL DISC	6 [hp_X] in 1 mL
SUS SCROFA BONE MARROW (UNII: VP2CN2G7Y8) (SUS SCROFA BONE MARROW - UNII:VP2CN2G7Y8)	SUS SCROFA BONE MARROW	6 [hp_X] in 1 mL

Inactive Ingredients

Ingredient Name	Strength
WATER (UNII: 059QF0KO0R)	
ALCOHOL (UNII: 3K9958V90M)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:83027-0082-1	60 mL in 1 BOTTLE, SPRAY; Type 0: Not a Combination Product	06/27/2023	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
unapproved homeopathic		06/27/2023	

Labeler - Nutritional Specialties, Inc. (032744609)

Revised: 6/2023

Nutritional Specialties, Inc.