

**MEDULLA OSSIS SUIS- sus scrofa bone marrow pellet
BOIRON**

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.

MEDULLA OSSIS SUIS

MEDULLA OSSIS SUIS HPUS

8X

Arthritic pain in spine

Arthritic pain in spine

STOP USE IF SYMPTOMS PERSIST FOR MORE THAN 3 DAYS OR WORSEN

ASK A HEALTH PROFESSIONAL BEFORE USE

KEEP OUT OF REACH OF CHILDREN

5 PELLETS 3 TIMES A DAY UNTIL SYMPTOMS ARE RELIEVED

DO NOT USE IF PELLETT DISPENSER IS BROKEN.

CONTAINS APPROX. 80 PELLETS

LACTOSE, SUCROSE

1-800-BOIRON-1

Newtown Square, PA 19073-3267

info@boironusa.com

These "Uses" have not been evaluated by the FDA.

Medulla Ossis Suis

8 x

BOIRON

Drug Facts: Active ingredient listed above. Use for symptoms listed below. **Warnings:** ■ Stop use and ask a doctor if symptoms persist for more than 3 days or worsen. ■ If pregnant or breast-feeding, ask a health professional before use. ■ Keep out of reach of children. **Directions:** 5 pellets 3 times a day until symptoms are relieved. **Other information:** Do not use if pellet dispenser seal is broken. contains approx. 80 pellets ■ **Inactive ingredients:** lactose, sucrose. *These "Uses" have not been evaluated by the FDA.

Lot: Exp:
Arthritic pain in spine*

HOMEOPATHIC MEDICINE ■ Made in France
1-800-808-8100 ■ Newtown Square, PA 19073 ■ info@boironusa.com



US

MEDULLA OSSIS SUIS

sus scrofa bone marrow pellet

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:0220-3329
Route of Administration	ORAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
SUS SCROFA BONE MARROW (UNII: VP2CN2G7Y8) (SUS SCROFA BONE MARROW - UNII:VP2CN2G7Y8)	SUS SCROFA BONE MARROW	8 [hp_X] in 8 [hp_X]

Inactive Ingredients

Ingredient Name	Strength
SUCROSE (UNII: C151H8M554)	
LACTOSE, UNSPECIFIED FORM (UNII: J2B2A4N98G)	

Product Characteristics

Color	white	Score	
Shape	ROUND	Size	4mm
Flavor		Imprint Code	

Contains**Packaging**

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:0220-3329-41	8 [hp_X] in 1 TUBE; Type 0: Not a Combination Product	01/08/2018	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
unapproved homeopathic		01/08/2018	

Labeler - BOIRON (282560473)**Registrant** - BOIRON, INC. (014892269)**Establishment**

Name	Address	ID/FEI	Business Operations
BOIRON		282560473	manufacture(0220-3329)

Revised: 11/2023

BOIRON