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 PELLET DISPENSER IS BROKEN.

CONTAINS APPROX. 80 PELLETS

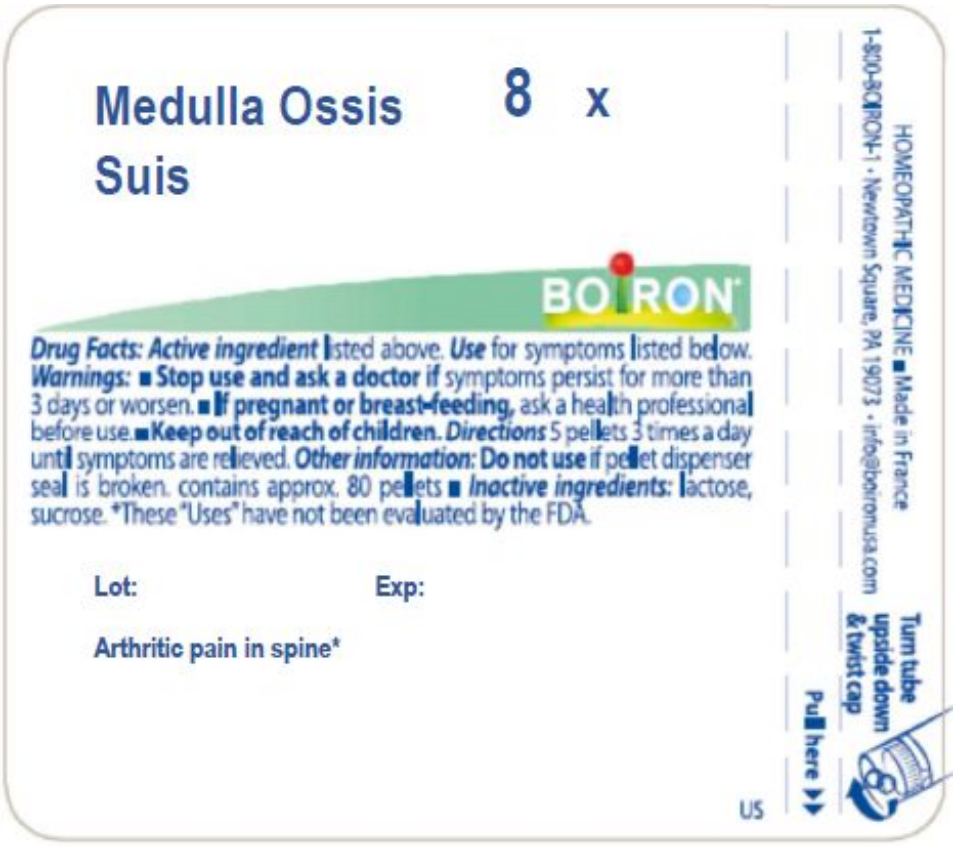
LACTOSE, SUCROSE

1-800-BOIRON-1

Newtown Square, PA 19073-3267

info@boironusa.com

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



MEDULLA OSSIS SUIS

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