

CROMOLYN SODIUM- cromolyn sodium powder

AX Pharmaceutical Corp

Cromolyn sodium

Cromolyn sodium

Caution: For pharmacy compounding only. For veterinary use only. Use according to practitioner's prescription. Federal law prohibits dispensing without prescription. Must be subjected to further processing during the preparation of sterile dosage forms. Preserve in tight containers, and store at controlled room temperature.



AX Pharmaceutical Corp

Cromolyn Sodium, USP

Expiry Date: Dec 15, 2023

100g

Original Reference: CP02-201201M

NDC: 73377-122-01

CAS: 15826-37-6

Relabelled by: AX Pharmaceutical Corp

Lot: G153-20L16SH

Original Manufacturer: Changzhou Tianhua Pharmaceutical Co., Ltd.
No. 3, Houyang Chemical Zone, Jintan., Changzhou, JIANGSU 213200, China

100 TesmaWay, Unit 8, Concord, ON Canada L4K 0J9 Fax: 416 352 1618

Toll free: 1 866 305 0566

Causes skin irritation. Causes serious eye irritation. May cause respiratory irritation. Avoid breathing dust/ fume/ gas/ mist/ vapour/ spray. Wash skin thoroughly after handling. Use only outdoors or in a well-ventilated area. Wear protective gloves/ eye protection/ face protection. IF ON SKIN: Wash with plenty of water. IF INHALED: Remove person to fresh air and keep comfortable if breathing. Call a POISON CENTER/doctor if you feel unwell. IF IN EYES: Rinse cautiously with water for several minutes. Remove contact lenses, if present and easy to do. Continuing if skin irritation occurs. Get medical advice/ attention. Take off contaminated clothing and wash it before reuse. Store in a well-ventilated place. Keep container tightly closed. Store locked up. Dispose of contents/container to an approved waste disposal plant.



Warning

CROMOLYN SODIUM

cromolyn sodium powder

Product Information

Product Type	BULK INGREDIENT	Item Code (Source)	NDC:73377-122
Route of Administration	NOT APPLICABLE		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
CROMOLYN SODIUM (UNII: Q2WXR1I0PK) (CROMOLYN SODIUM - UNII:Q2WXR1I0PK)	CROMOLYN SODIUM	1 g in 1 g

Product Characteristics

Color	white	Score	
Shape		Size	
Flavor		Imprint Code	
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:73377-122-01	100 g in 1 JAR	04/05/2021	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
bulk ingredient for animal drug compounding		04/05/2021	

Labeler - AX Pharmaceutical Corp (204011316)

Establishment

Name	Address	ID/FEI	Business Operations
AX Pharmaceutical Corp		204011316	repack(73377-122) , relabel(73377-122)

Revised: 4/2021

AX Pharmaceutical Corp