

ALENDRONATE SODIUM- alendronate sodium powder

AX Pharmaceutical Corp

Alendronate Sodium

Alendronate Sodium

Caution: For pharmacy compounding only. For veterinary use only. Use according to practitioner's prescription. Federal law prohibits dispensing without prescription.
 Preserve in well-closed containers. Store at room temperature, excursion allowed between 15°C ~ 30°C.



AX Pharmaceutical Corp

Alendronate Sodium, USP

Expiry date: Jun 30, 2027

Original Reference: 22AS013

NDC: 73377-173-01

Relabelled by: AX Pharmaceutical Corp

Original Manufacturer: Cadila Pharmaceuticals Limited
 Plot No. 294, GIDC Industrial Estate, Ankleshwar, Gujarat 393002, India

100g

CA S: 121268-17-5

Lot: H281-22G01SH

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

Toll free: 1 866 305 0566

Harmful if swallowed. Causes severe skin irritation. Causes serious eye damage. May cause respiratory irritation. May cause damage to bones, stomach through prolonged or repeated exposure.



Danger

Obtain special instructions before use. Do not handle until all safety precautions have been read and understood. Do not breathe dust/fume/gas/mist/vapor/spray. Wash hands thoroughly after handling. Wear protective gloves/protective clothing/eye protection/face protection. IF SWALLOWED: Call a POISON CENTER or doctor/physician if you feel unwell. IF ON SKIN: Wash with plenty of soap and water. IF INHALED: Remove victim to fresh air and keep at rest in a position comfortable for breathing. IF IN EYES: Rinse cautiously with water for several minutes. Remove contact lenses if present and easy to do. Continue rinsing. IF ON CLOTHING: Get medical advice/attention. Immediately call a POISON CENTER or doctor/physician. Get medical attention if you feel unwell. Rinse mouth. If skin irritation occurs, get medical advice/attention. Take off contaminated clothing and wash it before reuse.

ALENDRONATE SODIUM

alendronate sodium powder

Product Information

Product Type		Item Code (Source)	NDC:73377-173
Route of Administration	NOT APPLICABLE		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
ALENDRONATE SODIUM (UNII: 2UY4M2U3RA) (ALENDRONIC ACID - UNII:X1J18R4W8P)	ALENDRONATE 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:73377-173-01	100 g in 1 JAR		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
bulk ingredient for animal drug compounding		11/09/2022	

Labeler - AX Pharmaceutical Corp (204011316)

Establishment

Name	Address	ID/FEI	Business Operations
AX Pharmaceutical Corp		204011316	relabel, repack

Establishment

Name	Address	ID/FEI	Business Operations
Cadila Pharmaceuticals Limited		916130698	api manufacture

Revised: 11/2022

AX Pharmaceutical Corp