

DICLOFENAC SODIUM- diclofenac sodium powder

AX Pharmaceutical Corp

Diclofenac sodium

Diclofenac sodium

Caution: For pharmacy compounding only. Use according to practitioner's prescription. Federal law prohibits dispensing without prescription. Preserve in tight, light-resistant containers at controlled room temperature.



AX Pharmaceutical Corp

Diclofenac Sodium, USP

Retest Date: Jun 16, 2024

500g

Original Reference: 301200617-6

NDC: 73377-113-01

CAS: 15307-79-6

Repackaged by: AX Pharmaceutical Corp

Lot: F438-20F17P2

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

Toll free: 1 866 305 0566

Harmful if swallowed. Suspected of damaging fertility or the unborn child. Causes damage to organs (gastrointestinal, oral) through prolonged or repeated exposure. Toxic to aquatic life with long lasting effects.



Obtain special instructions before use. Do not handle until all safety precautions have been read and understood. Do not breathe dust/fume/gas/vapor/spray. Wash (hands) thoroughly after handling. Avoid release to the environment. Wear protective gloves/protective clothing/eye protection/face protection. IF SWALLOWED: Call a POISON CENTER or doctor/physician if you feel unwell. If exposed or concerned: Get medical attention/advice. Get medical attention/advice if you feel unwell. Rinse mouth. Collect spillage.

Danger

DICLOFENAC SODIUM

diclofenac sodium powder

Product Information

Product Type	BULK INGREDIENT	Item Code (Source)	NDC:73377-113
Route of Administration	NOT APPLICABLE		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
DICLOFENAC SODIUM (UNII: QTG126297Q) (DICLOFENAC SODIUM - UNII:QTG126297Q)	DICLOFENAC 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:73377-113-01	500 g in 1 JAR	06/02/2021	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
bulk ingredient for animal drug compounding		06/02/2021	

Labeler - AX Pharmaceutical Corp (204011316)

Establishment

Name	Address	ID/FEI	Business Operations
AX Pharmaceutical Corp		204011316	repack(73377-113) , relabel(73377-113)

Revised: 6/2021

AX Pharmaceutical Corp