AX PHARMACEUTICAL CORP- cyclophosphamide powder
AX Pharmaceutical Corp

Caution: For manufacturing, processing, or repacking. Use according to practitioner's prescription. Federal law prohibits dispensing without prescription.

Keep tightly closed in a cool place

AX Pharmaceutical Corp

Toll free: 1 866 305 0566

Cyclophosphamide , USP

Retest date: Dec 09, 2020

Original Reference: 20171210

NDC Number: 62157-514-01

Relabelled by: AX Pharmaceutical Corp

500g

CAS: 6055-19-2

Lot: D006-17L10SH

100 West Beaver Creek Road, Unit 12, Richmond Hill, ON Canada L4B 1H4 Fax: 416 852 1618

Toxic if swallowed. May cause cancer. Causes skin irritation. Causes serious eye damage. May cause genetic defects. May damage fertility or the unborn child. Causes damage to organs (bone marrow) through prolonged or repeated exposure.

Obtain special instructions before use. Do not handle until all safety precautions have been read and understood. Wash thoroughly after handling. Wear protective gloves/protective clothing/eye protection/face protection. If swallowed: Immediately call a poison centre/doctor. Rinse mouth. Do not induce vomiting. If on skin (or hair) or water: Take off immediately all contaminated clothing and wash it before reuse. If in eyes: Rinse cautiously with water for several minutes. Remove contact lenses, if present and easy to do. Continue rinsing. Immediately call a poison centre/doctor. If exposed or concerned: Get medical advice/attention. If medical advice/attention is needed, have the product or the product's name and the Dispose of contents/container in accordance with local/regional/national/international regulations.

Danger

AX PHARMACEUTICAL CORP

cyclophosphamide powder

Product Information

Product Type	BULK INGREDIENT	**Item Code (Source)**	NDC:62157-514
Route of Administration	NOT APPLICABLE		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
CYCLOPHOSPHAMIDE (UNII: 8N3DW7272P) (CYCLOPHOSPHAMIDE ANHYDROUS - UNII:6UXW23996M)	CYCLOPHOSPHAMIDE ANHYDROUS	495 g in 500 g

Inactive Ingredients

Ingredient Name	Strength
WATER (UNII: 059QF0KO0R)	

Product Characteristics

Color	white	**Score**	
Shape		**Size**	
Flavor		**Imprint Code**	
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:62157-514-01	500 g in 1 JAR	01/08/2018	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
bulk ingredient for animal drug compounding		01/08/2018	

Labeler - AX Pharmaceutical Corp (202924858)

Establishment

Name	Address	ID/FEI	Business Operations
AX Pharmaceutical Corp		202924858	repack(62157-514) , relabel(62157-514)