

AX PHARMACEUTICAL CORP- cetirizine hcl powder
AX Pharmaceutical Corp

Caution: For manufacturing, processing or repackaging. Use according to practitioner's prescription. Federal law prohibits dispensing without prescription.
Keep tightly closed at controlled room temperature.



AX Pharmaceutical Corp

Cetirizine Hydrochloride, EP

Retest date: June 05, 2022

500g

Original Reference: BCTDZ17008

NDC: 62157-568-01

CAS: 83881-52-1

Relabelled by: AX Pharmaceutical Corp

Lot: D087-17F06SH

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

Toll free: 1 866 305 0566

Harmful, if swallowed. May cause drowsiness or dizziness.
Dispose of contents/container in accordance with local/regional/national/international regulations.



Warning

Do not breathe dust/fume/gas/mist/vapour/spray, avoid release to the environment. Wear protective gloves. Wear respiratory protection. Immediately call a poison centre or doctor/physician.

Product Information

Product Type	BULK INGREDIENT	Item Code (Source)	NDC:62157-568
Route of Administration	NOT APPLICABLE		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
CETIRIZINE HYDROCHLORIDE (UNII: 64O047KTOA) (CETIRIZINE - UNII:YO7261ME24)	CETIRIZINE HYDROCHLORIDE	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:62157-568-01	500 g in 1 JAR	04/27/2018	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
bulk ingredient for animal drug compounding		04/27/2018	

Labeler - AX Pharmaceutical Corp (202924858)**Establishment**

Name	Address	ID/FEI	Business Operations
AX Pharmaceutical Corp		202924858	repack(62157-568) , relabel(62157-568)