

AX PHARMACEUTICAL CORP- ranitidine hcl powder
AX Pharmaceutical Corp

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Ranitidine HCl, USP

Retest date: Jan 16, 2020

1kg

Original Reference: RAA-1701025

CAS: 66357-59-3

NDC Number: 62157-515-01

Relabelled by: AX Pharmaceutical Corp

Lot: C345-17A17SH

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

Toll free: 1 866 305 0566

May cause an allergic skin reaction.
May cause allergy or asthma symptoms or breathing difficulties if inhaled.

Contaminated work clothing must not be allowed out of the workplace. Wear protective gloves in case of inadequate ventilation wear respiratory protection. If on skin: Wash with plenty of water. If skin irritation or rash occurs: Get medical advice/attention. Wash contaminated clothing before reuse. If inhaled: If breathing is difficult, remove person to fresh air and keep comfortable for breathing. If expending respiratory symptoms. Call a poison center/doctor.



Danger

AX PHARMACEUTICAL CORP

ranitidine hcl powder

Product Information			
Product Type	BULK INGREDIENT	Item Code (Source)	NDC:62157-515
Route of Administration	NOT APPLICABLE		

Active Ingredient/Active Moiety		
Ingredient Name	Basis of Strength	Strength
RANITIDINE HYDRO CHLO RIDE (UNII: BK76465IHM) (RANITIDINE - UNII:884KT10YB7)	RANITIDINE	.99 kg in 1 kg

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:62157-515-01	1 kg in 1 JAR	01/09/2017	

Marketing Information			
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
bulk ingredient for animal drug compounding		01/09/2017	

Labeler - AX Pharmaceutical Corp (202924858)

Establishment

Name	Address	ID/FEI	Business Operations
AX Pharmaceutical Corp		202924858	repack(62157-515) , relabel(62157-515)