

AX PHARMACEUTICAL CORP- ponazuril powder
AX Pharmaceutical Corp

Caution: For manufacturing, processing, repack-
ing or compounding. Use according to practition-
er's prescription. Federal law prohibits dispensing
without prescription.
Keep tightly closed at controlled room temper-
ature.



AX Pharmaceutical Corp

Ponazuril

Retest date: May 17, 2020

500g

Original Reference: MK20180501

NDC: 62157-619-01

CAS: 69004-04-2

Relabelled by: AX Pharmaceutical Corp

Lot: D206-18E18SH

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

Toll free: 1 866 305 0566

Do not breathe dust/fume/gas/
mist/ vapours/ spray, avoid
release to the environment,
wear protective gloves. Wear
respiratory protection. In case
of fire, call the POISON CENTER or
doctor/physician.

AX PHARMACEUTICAL CORP

ponazuril powder

Product Information			
Product Type	BULK INGREDIENT	Item Code (Source)	NDC:62157-619
Route of Administration	NOT APPLICABLE		

Active Ingredient/Active Moiety		
Ingredient Name	Basis of Strength	Strength
PONAZURIL (UNII: JPW84AS66U) (PONAZURIL - UNII:JPW84AS66U)	PONAZURIL	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:62157-619-01	500 g in 1 JAR	07/24/2018	

Marketing Information			
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
bulk ingredient for animal drug compounding		07/24/2018	

Labeler - AX Pharmaceutical Corp (202924858)

Establishment

Name	Address	ID/FEI	Business Operations
AX Pharmaceutical Corp		202924858	repack(62157-619) , relabel(62157-619)