

AX PHARMACEUTICAL CORP- butaphosphan powder
AX Pharmaceutical Corp

Caution: For prescription compounding use only.
Use according to practitioner's prescription.
Federal law prohibits dispensing without prescription.
Keep tightly closed in a cool place



AX Pharmaceutical Corp

Butaphosphan

Retest date: Oct 29, 2019

500g

Original Reference: BTL-171101M

NDC Number: 62157-508-01

CAS: 17316-67-5

Relabelled by: AX Pharmaceutical Corp

Lot: C356-17J30SH

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

Toll free: 1 866 305 0566

Caution: Skin Irritation. Causes serious eye irritation.

May be harmful if inhaled. Contains dust which should not be inhaled and may irritate the respiratory tract. Avoid contact with clothing and do not be allowed out of the workplace. Wear protective gloves. If you breathe the dust, you may experience irritation of your eyes, nose and throat. If you experience any of these symptoms, stop using the product immediately and seek medical advice. Wash your face and hands thoroughly after use. Do not eat, drink or smoke while using this product. Use only in well-ventilated areas. Do not use if the container is damaged or if the contents are not as described. Keep this product away from children. Do not use if the container is damaged or if the contents are not as described. Keep this product away from children. Do not use if the container is damaged or if the contents are not as described. Keep this product away from children.



Warning

AX PHARMACEUTICAL CORP

butaphosphan powder

Product Information

Product Type	BULK INGREDIENT	Item Code (Source)	NDC:62157-508
Route of Administration	NOT APPLICABLE		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
BUTAFOSFAN (UNII: 3Q2LQ1149L) (BUTAFOSFAN - UNII:3Q2LQ1149L)	BUTAFOSFAN	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:62157-508-01	500 g in 1 JAR	12/15/2017	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
bulk ingredient for animal drug compounding		12/15/2017	

Labeler - AX Pharmaceutical Corp (202924858)

Revised: 12/2017

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