

X- orlistat powder
AX Pharmaceutical Corp



AX Pharmaceutical

Orlistat

Retest date: March 31, 2018

Original Reference #: 601160309

NDC#: 62157-394-01

CAS #: 96829-58-2

Relabelled by: AX Pharmaceutical

500g

Lot #: B336-16D23SH

Toll free: 1.866.3050566

Do not breathe dust/ fume/ gas/
mist/ vapours/ spray, avoid
release to the environment,
wear protective gloves. Wear
respiratory protection. Immedi-
ately call a POISON CENTER or
doctor/ physician.

Caution: For prescription compounding use only.
Use according to practitioner's prescription.
Federal law prohibits dispensing without prescrip-
tion.
Keep the product at temperature 2° C ~ 8° C.

100 West Beaver Creek Road, Unit 12, Richmond Hill, ON Canada L4B 1H4 Fax: 416 8521618

orlistat powder

Product Information

Product Type	BULK INGREDIENT	Item Code (Source)	NDC:62157-394
Route of Administration	NOT APPLICABLE		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
ORLISTAT (UNII: 95M8R751W8) (ORLISTAT - UNII:95M8R751W8)	ORLISTAT	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:62157-394-01	500 g in 1 VIAL	03/01/2017	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
bulk ingredient for animal drug compounding		03/01/2017	

Labeler - AX Pharmaceutical Corp (202924858)

Establishment

Name	Address	ID/FEI	Business Operations
AX Pharmaceutical Corp		202924858	pack(62157-394) , relabel(62157-394)