

TADALAFIL- tadalafil powder
AX Pharmaceutical Corp

Tadalafil

Tadalafil

Caution: For pharmacy compounding only. For veterinary use only. Use according to practitioner's prescription. Federal law prohibits dispensing without prescription.
 Preserve in a well-closed container at controlled room temperature.



AX Pharmaceutical Corp

Tadalafil, USP

Retest date: Apr 30, 2029

Original Reference: D5383-24-009

NDC: 73377-299-01

Repackaged by: AX Pharmaceutical Corp

Original Manufacturer: Zhejiang Huahai Pharmaceutical Co., Ltd.
 Coastal Industrial Zone, Duqiao, Linhai, Zhejiang 317016, China (CHN)

100 Tesma Way, Unit 8, Concord, ON Canada L4K 0J9 Fax: 416 352 1618

50g

CAS: 171596-29-5

Lot: J123-24E05P10

Toll free: 1 866 305 0566

Causes skin irritation. Causes serious eye irritation.
 May cause respiratory irritation.
 Avoid breathing dust/ fumes/ gas/ mist/ vapours/ spray. Wash skin thoroughly after handling. Use only outdoors or in a well-ventilated area. Wear protective gloves. IF ON SKIN: Wash with plenty of water. IF INHALED: Remove person to fresh air and keep comfortable for breathing. Call a POISON CENTER/doctor if you feel unwell. IF IN EYES: Rinse cautiously with water for several minutes. Remove contact lenses, if present and easy to do. Continue rinsing. IF skin irritation occurs: Get medical advice/attention. If eye irritation persists: Get medical advice/attention. Take off contaminated clothing and wash it before reuse. Store in a well-ventilated place. Keep container tightly closed. Store locked up. Dispose of contents/ container to an approved waste disposal plant.



Warning

TADALAFIL

tadalafil powder

Product Information

Product Type		Item Code (Source)	NDC:73377-299
Route of Administration	NOT APPLICABLE		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
Tadalafil (UNII: 742SXX0ICT) (TADALAFIL - UNII:742SXX0ICT)	Tadalafil	1 g in 1 g

Product Characteristics

Color	white	Score	
Shape		Size	
Flavor		Imprint Code	
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:73377-299-01	50 g in 1 JAR		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
bulk ingredient for animal drug compounding		10/30/2024	

Labeler - AX Pharmaceutical Corp (204011316)

Establishment

Name	Address	ID/FEI	Business Operations
AX Pharmaceutical Corp		204011316	analysis, pack, label, relabel, repack

Revised: 10/2024

AX Pharmaceutical Corp