

SERTRALINE HCL- sertraline hcl powder

AX Pharmaceutical Corp

Sertraline Hcl

Sertraline Hcl

Complies with USP Organic Impurities Procedure 2.
Caution: For pharmacy compounding only. For veterinary use only. Use according to practitioner's prescription. Federal law prohibits dispensing without prescription.
Preserve in tight, light-resistant containers at a temperature below 40°C.



AX Pharmaceutical Corp

Sertraline HCl, USP

Expiry Date: May 14, 2025

100g

Original Reference: DK70-2205251

NDC: 73377-134-01

CAS: 79559-97-0

Relabelled by: AX Pharmaceutical Corp

Lot: H285-22E15SH

Original Manufacturer: Shangyu Jingxin Pharmaceutical Co., Ltd.
No. 31 Weisan Road, Hangzhou Bay Shangyu Economic and Technological Development Area,
Shangyu, 312369, Zhejiang Province, China

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

Toll free: 1 866 305 0566

Harmful if swallowed. Very toxic to aquatic life.



Wash skin thoroughly after handling. Do not eat, drink or smoke when using this product. Avoid release to the environment. IF SWALLOWED: Call a POISON CENTER/doctor if you feel unwell. Rinse mouth. Collect spillage. Dispose of contents/ container to an approved waste disposal plant.

Warning

Complies with USP Organic Impurities Procedure 2.
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Preserve in tight, light-resistant containers at a temperature below 40°C.



AX Pharmaceutical Corp

Sertraline HCl, USP

Expiry Date: May 14, 2025

50g

Original Reference: DK70-2205251

NDC: 73377-134-02

CAS: 79559-97-0

Relabelled by: AX Pharmaceutical Corp

Lot: H285-22E15SH

Original Manufacturer: Shangyu Jingxin Pharmaceutical Co., Ltd.
No. 31 Weisan Road, Hangzhou Bay Shangyu Economic and Technological Development Area,
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Warning

SERTRALINE HCL

sertraline hcl powder

Product Information

Product Type

Item Code (Source)

NDC:73377-134

Route of Administration

NOT APPLICABLE

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
SERTRALINE HYDROCHLORIDE (UNII: UTI8907Y6X) (SERTRALINE - UNII: QUC7NX6VMB)	SERTRALINE	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:73377-134-01	100 g in 1 JAR		
2	NDC:73377-134-02	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		07/19/2022	

Labeler - AX Pharmaceutical Corp (204011316)

Establishment

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

Establishment

Name	Address	ID/FEI	Business Operations
Shangyu Jingxin Pharmaceutical Co., Ltd.		527982528	api manufacture

Revised: 10/2024

AX Pharmaceutical Corp