

CLOMIPRAMINE HCL- clomipramine hcl powder

AX Pharmaceutical Corp

Clomipramine HCl

Clomipramine HCl

Sample for testing only. Product is not for sale. Keep tightly closed at controlled room temperature, away from strong heat and light.



AX Pharmaceutical Corp

Clomipramine HCl, USP

SAMPLE FOR TESTING ONLY

Retest Date: Jan 31, 2031

100g

Original Reference: CLM/226011

CA S: 17321-77-6

NDC: 73377-196-04

Lot: L048-26B01SH

Relabelled by: AX Pharmaceutical Corp

Original Manufacturer: R.L. Fine Chem Private Limited
No IP-27-29, KIADB Industrial Area 1st Phase, Kudumalakunte Village, Chikkaballapur District Gowribidanur-Taluk, Chikkaballapur, Karnataka, 561208, India

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

Toll free: 1 866 305 0566

Hamful if swallowed. May cause drowsiness or dizziness. Causes damage to organs (Central nervous system, Cardiovascular). Very toxic to aquatic life with long lasting effects.



Danger

Do not breathe dust. Wash skin thoroughly after handling. Do not eat, drink or smoke when using this product. Use only outdoors or in a well-ventilated area. Avoid release to the environment. IF SWALLOWED: Call a POISON CENTER/ doctor if you feel unwell. Rinse mouth. IF INHALED: Remove person to fresh air and keep comfortable for breathing. Call a POISON CENTER/ doctor if you feel unwell. IF exposed or concerned: Call a POISON CENTER/ doctor. Collect spillage. Store in a well-ventilated place. Keep container tightly closed. Store locked up. Dispose of contents/ container to an approved waste disposal plant.

CLOMIPRAMINE HCL

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Product Information

Product Type		Item Code (Source)	NDC:73377-196
Route of Administration	NOT APPLICABLE		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
CLOMIPRAMINE HYDROCHLORIDE (UNII: 2LXW0L6GWJ) (CLOMIPRAMINE - UNII:NUV44L116D)	CLOMIPRAMINE HYDROCHLORIDE	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:73377-196-04	100 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		04/13/2026	

Labeler - AX Pharmaceutical Corp (204011316)

Establishment

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

Revised: 4/2026

AX Pharmaceutical Corp