

PONAZURIL- ponazuril powder
AMERICAN PHARMACEUTICAL INGREDIENTS LLC

Ponazuril



Ponazuril (VET ONLY)

Qty: 10 GM Lot: 000000
 CAS#: 69004-04-2 NDC#: 58597-8452-3
 Mfg Date: MM/DD/YY Exp Date: MM/DD/YY

Produced under cGMP guidelines by American Pharmaceutical Ingredients, LLC, Waterford, Michigan USA

FD# 3010302040
 DEA: RA0481764
 C.S. Facility: 5015062541
 BDP Michigan Distributor: 5306004308

Federal Law Prohibits Dispensing Without a Prescription.

CAUTION/DIRECTION FOR USE:
 For Prescription compounding use only, by a licensed pharmacist, directed by a licensed physician's prescription in accordance with FDA Compliance Policy Guidelines pertaining to Custom Rx Pharmacy Compounding.

For more information please see MSDS & COA provided with this product.

Your State Law Prohibits wholesalers to distribute to your pharmacy without an active license.

Please keep a copy of our license authorizing us to distribute to you on file at all times.

Sales: (888) 405-7271
 Operations: (248) 522-6002
 Fax: (248) 493-5936
 Email: info@American-Pharmaceutical.com

Facility Address: 6650 Highland Road, Waterford, Michigan 48327 USA

PONAZURIL

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Product Information

Product Type	BULK INGREDIENT	Item Code (Source)	NDC:58597-8452
Route of Administration	NOT APPLICABLE		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
Ponazuril (UNII: JPW84AS66U) (Ponazuril - UNII:JPW84AS66U)	Ponazuril	1 g in 1 g

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:58597-8452-3	10 g in 1 BOTTLE		
2	NDC:58597-8452-4	25 g in 1 BOTTLE		
3	NDC:58597-8452-6	100 g in 1 BOTTLE		

Marketing Information

Marketing Category	Application Number or Monograph	Marketing Start	Marketing End
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Marketing Category	Citation	Date	Date
bulk ingredient for animal drug compounding		04/29/2014	

Labeler - AMERICAN PHARMACEUTICAL INGREDIENTS LLC (078793641)

Revised: 4/2014

AMERICAN PHARMACEUTICAL INGREDIENTS LLC