

TYLOSIN TARTRATE- tylosin tartrate powder
AMERICAN PHARMACEUTICAL INGREDIENTS LLC

Tylosin Tartrate



Tylosin Tartrate EP (VET ONLY)

Qty: 5 GM Lot: 000000
 CAS#: 1405-54-5 NDC#: 58597-8569-2
 Mfg Date: MM/DD/YY Exp Date: MM/DD/YY

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

FDA: 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

TYLOSIN TARTRATE

tylosin tartrate powder

Product Information

Product Type	BULK INGREDIENT	Item Code (Source)	NDC:58597-8569
Route of Administration	NOT APPLICABLE		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
Tylosin Tartrate (UNII: 5P4625C51T) (Tylosin Tartrate - UNII:5P4625C51T)	Tylosin Tartrate	1 g in 1 g

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:58597-8569-2	5 g in 1 BOTTLE		
2	NDC:58597-8569-4	25 g in 1 BOTTLE		
3	NDC:58597-8569-6	100 g in 1 BOTTLE		
4	NDC:58597-8569-7	500 g in 1 BOTTLE		
5	NDC:58597-8569-8	1000 g in 1 BOTTLE		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
bulk ingredient for animal drug compounding		05/02/2014	

Labeler - AMERICAN PHARMACEUTICAL INGREDIENTS LLC (078793641)

Revised: 5/2014

AMERICAN PHARMACEUTICAL INGREDIENTS LLC