

**ERYTHROMYCIN ETHYLSUCCINATE- erythromycin ethylsuccinate powder**  
**AX Pharmaceutical Corp**

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**Erythromycin Ethylsuccinate**

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Caution: For pharmacy compounding only. For veterinary use only. Use according to practitioner's prescription. Federal law prohibits dispensing without prescription.  
 Keep tightly closed at a controlled room temperature.



**AX Pharmaceutical Corp**

**Erythromycin Ethylsuccinate, USP**

Toll free: 1 866 305 0566

**Retest date:** Apr 11, 2022

**200g**

**Original Reference:** C01-201804013

**NDC:** 62157-864-01

**CA S:** 1264-62-6

**Relabelled by:** AX Pharmaceutical Corp

Original manufacturer: Wuhan Longke Biotechnology Co., Ltd.  
 No.199, Shendun rd, Wuhan, Hubei 430000, China

**Lot:** E235-18D12SH

May cause an allergic skin reaction. May cause dizziness or lightheadedness or breathing difficulties if inhaled. May be harmful if inhaled. May cause respiratory tract irritation. May be harmful if swallowed through skin. May cause dizziness if inhaled. May cause eye irritation. May be harmful if inhaled.  
 Avoid breathing about head (use full length of spray). Wear protective gloves. If symptoms persist or worsen, call a POISON CENTER or doctor/physician. If swallowed, more persistent headache, if not breathing, give artificial respiration. In case of skin contact, wash with soap and plenty of water. In case of eye contact, wash thoroughly with water. If inhaled, if possible, have person breathe mouth-to-mouth for an unobstructed passage. Have mouth with water. Consult a physician.



Danger

100 West Beaver Creek Road, Unit 12, Richmond Hill, ON Canada L4B 1H4 Fax: 416 852 1618

**ERYTHROMYCIN ETHYLSUCCINATE**

erythromycin ethylsuccinate powder

**Product Information**

<b>Product Type</b>	BULK INGREDIENT	<b>Item Code (Source)</b>	NDC:62157-864
<b>Route of Administration</b>	NOT APPLICABLE		

**Active Ingredient/Active Moiety**

Ingredient Name	Basis of Strength	Strength
ERYTHROMYCIN ETHYLSUCCINATE (UNII: 1014KSJ86F) (ERYTHROMYCIN - UNII:63937KV33D)	ERYTHROMYCIN	1 g in 1 g

**Product Characteristics**

<b>Color</b>	white	<b>Score</b>	
<b>Shape</b>		<b>Size</b>	
<b>Flavor</b>		<b>Imprint Code</b>	
<b>Contains</b>			

**Packaging**

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:62157-864-01	200 g in 1 JAR	06/14/2019	

## Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
bulk ingredient for animal drug compounding		06/14/2019	

**Labeler** - AX Pharmaceutical Corp (202924858)

## Establishment

Name	Address	ID/FEI	Business Operations
AX Pharmaceutical Corp		202924858	relabel(62157-864) , repack(62157-864)

Revised: 6/2019

AX Pharmaceutical Corp