

**END-ITCH FOR ECZEMA- end-itch for eczema cream**  
**Crystal Connections, LLC**

*Disclaimer: Most OTC drugs are not reviewed and approved by FDA, however they may be marketed if they comply with applicable regulations and policies. FDA has not evaluated whether this product complies.*

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**END-ITCH FOR ECZEMA**

**OTC - ACTIVE INGREDIENT**

Colloidal Oatmeal 37.5%

**INACTIVE INGREDIENTS**

Almond Oil, Beeswax, Jojoba Oil, Lanolin,  
Rosemary, Vitamin E Oil

**OTC - KEEP OUT OF REACH OF CHILDREN SECTION**

If swallowed get medical help or contact  
a poison control center immediately.

**WARNINGS**

For external use only  
Do not use on open cuts  
When using this product do not get into eyes  
Do not ingest

**OTC - PURPOSE SECTION**

Temporarily relieves itching from eczema.

**OTC - STOP USE SECTION**

Condition worsens  
Sensitivity develops or increases  
Symptoms last more than 7 days or  
clear up and occur again in a few days

**INDICATIONS & USAGE**

Temporarily relieves itching from eczema.

**DOSAGE & ADMINISTRATION**

Apply to affected area  
Rub in circular motion  
Apply as needed

**PACKAGE LABEL.PRINCIPAL DISPLAY PANEL**

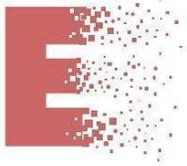
NDC 60858-315-03

**END-ITCH**  
(Colloidal Oatmeal)  
Skin Protectant

**For Eczema**

Temporary Relief Due To  
Dry Skin from Eczema

No steroids or cortizone



MADE IN THE USA • ALL NATURAL  
Net Contents: 3.2 oz (94ml)

**END-ITCH FOR ECZEMA**

end-itch for eczema cream

**Product Information**

<b>Product Type</b>	HUMAN OTC DRUG	<b>Item Code (Source)</b>	NDC:60858-315
<b>Route of Administration</b>	TOPICAL		

**Active Ingredient/Active Moiety**

Ingredient Name	Basis of Strength	Strength
OATMEAL (UNII: 8 PI54V663Y) (OATMEAL - UNII:8 PI54V663Y)	OATMEAL	0.24 g in 1 mL

**Inactive Ingredients**

Ingredient Name	Strength
ALMOND OIL (UNII: 66 YXD4DKO9)	
YELLOW WAX (UNII: 2ZA36H0S2V)	
JOJOBA OIL (UNII: 724GKU717M)	
lanolin (UNII: 7EV65EAW6H)	

rosemary (UNII: IJ67X351P9)

ALPHA-TOCOPHEROL (UNII: H4N855PNZ1)

### Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:60858-315-03	94 mL in 1 BOTTLE; Type 0: Not a Combination Product	05/15/2015	

### Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph final	part347	05/15/2015	

**Labeler** - Crystal Connections, LLC (016582119)

### Establishment

Name	Address	ID/FEI	Business Operations
American Pharmaceuticals and Cosmetics, Inc.		038023805	manufacture(60858-315)

Revised: 5/2015

Crystal Connections, LLC