

**GEMBOOXT ECZEMA RELIEF CREAM- hydrocortisone cream**  
**Guangdong Quadrant Ecological Technology Co., Ltd.**

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**76986-019 Gemboox ECZEMA RELIEF CREAM**

Hydrocortisone (1%)

Anti-Itch

Temporarily relieves the itching and discomfort associated with minor skin irritations, rashes, and inflammation, and due to eczema.  
Other uses of this product should only be under the advice and supervision of a doctor.

For external use only.

Do not use if allergic to any ingredients in this product.  
Do not use for the treatment of diaper rash.  
Do not use more than directed unless told to do so by a doctor.

When using this product, avoid contact with eyes, nose and mouth. If contact occurs, rinse thoroughly with water.

Ask a doctor before use if pregnant or nursing

Stop using and ask a doctor if condition worsens, symptoms persist for more than 7 days or clear up and occur again within a few days, and do not begin use of any other hydrocortisone product unless you have asked a doctor.

Keep out of reach of children. In case of accidental ingestion, get medical help or contact a poison control center immediately.

Adults and children 2 years of age and older: apply a thin layer to affected area not more than 3 to 4 times daily.

Children under 2 years of age: ask a doctor.

WATER  
CETYL ALCOHOL  
STEARYL ALCOHOL  
GLYCERYL STEARATE  
GLYCERIN  
MINERAL OIL  
CETEARETH-25  
STEARIC ACID  
ETHYLHEXYL PALMITATE  
CAMPHOR  
MENTHOLUM  
CENTELLA ASIATICA EXTRACT  
AVENA SATIVA (OAT) KERNEL EXTRACT  
ROSA RUGOSA FLOWER EXTRACT  
LACTOBACILLUS FERMENT  
BUTYLENE GLYCOL  
DIMETHICONE

GLYCERYL CAPRYLATE  
HYDROXYACETOPHENONE  
ETHYLHEXYLGLYCERIN  
1,2-HEXANEDIOL  
TRIETHANOLAMINE  
ALLANTOIN  
ACRYLATES/C10-30 ALKYL ACRYLATE CROSSPOLYMER  
XANTHAN GUM  
BUTYLENE GLYCOL  
PENTYLENE GLYCOL  
HYDROXYPHENYL PROPAMIDOBENZOIC ACID  
ASCORBYL PALMITATE  
POTASSIUM LAURYL PHOSPHATE  
DIPOTASSIUM PHOSPHATE



## GEMBOOX ECZEMA RELIEF CREAM

hydrocortisone cream

Product Information			
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:76986-019
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety		
Ingredient Name	Basis of Strength	Strength
<b>HYDROCORTISONE</b> (UNII: W4X0X7BPJ) (HYDROCORTISONE - UNII:W4X0X7BPJ)	HYDROCORTISONE	0.01 g in 1 g

Inactive Ingredients	
Ingredient Name	Strength
<b>WATER</b> (UNII: 059QF0KO0R)	
<b>CETEARETH-25</b> (UNII: 8FA93U5T67)	
<b>DIMETHICONE</b> (UNII: 92RU3N3Y1O)	
<b>DIBASIC POTASSIUM PHOSPHATE</b> (UNII: CI71S98N1Z)	
<b>HYDROXYPHENYL PROPAMIDOBENZOIC ACID</b> (UNII: 25KRT26H77)	
<b>BUTYLENE GLYCOL</b> (UNII: 3XUS85K0RA)	
<b>STEARYL ALCOHOL</b> (UNII: 2KR89I4H1Y)	
<b>XANTHAN GUM</b> (UNII: TTV12P4NEE)	
<b>STEARIC ACID</b> (UNII: 4ELV7Z65AP)	
<b>CAMPHOR (SYNTHETIC)</b> (UNII: 5TJD82A1ET)	
<b>ROSA RUGOSA FLOWER</b> (UNII: 969IQC5YJU)	
<b>ASCORBYL PALMITATE</b> (UNII: QN83US2B0N)	
<b>POTASSIUM LAURYL PHOSPHATE</b> (UNII: C4QT53N4MK)	
<b>TRIETHANOLAMINE</b> (UNII: 9O3K93S3TK)	
<b>CETYL ALCOHOL</b> (UNII: 936JST6JCN)	
<b>ALLANTOIN</b> (UNII: 344S277G0Z)	
<b>OAT</b> (UNII: Z6J799EAJK)	
<b>GLYCERYL CAPRYLATE</b> (UNII: TM2TZD4G4A)	
<b>1,2-HEXANEDIOL</b> (UNII: TR046Y3K1G)	
<b>ETHYLHEXYL PALMITATE</b> (UNII: 2865993309)	
<b>MINERAL OIL</b> (UNII: T5L8T28FGP)	
<b>MENTHOL</b> (UNII: L7T10EIP3A)	
<b>CENTELLA ASIATICA TRITERPENOIDS</b> (UNII: 4YS74Q4G4J)	
<b>HYDROXYACETOPHENONE</b> (UNII: G1L3HT4CMH)	
<b>PENTYLENE GLYCOL</b> (UNII: 50C1307PZG)	
<b>GLYCERYL STEARATE</b> (UNII: 230OU9XXE4)	
<b>ACRYLATES/C10-30 ALKYL ACRYLATE CROSSPOLYMER (60000 MPA.S AT 0.5%)</b> (UNII: YY2H MJ9NZF)	
<b>GLYCERIN</b> (UNII: PDC6A3C0OX)	
<b>LACTOBACILLUS REUTERI F275 STRAIN</b> (UNII: 3K209OP33H)	
<b>ETHYLHEXYLGLYCERIN</b> (UNII: 147D247K3P)	

Packaging				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:76986-019-01	2 in 1 BOX	12/19/2025	
1		15 g in 1 BOTTLE; Type 0: Not a Combination Product		
2	NDC:76986-019-02	20 in 1 BOX	12/19/2025	
2		1.5 g in 1 BAG; Type 0: Not a Combination Product		

## Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M016	12/19/2025	

**Labeler** - Guangdong Quadrant Ecological Technology Co., Ltd. (554532634)

## Establishment

Name	Address	ID/FEI	Business Operations
Guangdong Quadrant Ecological Technology Co., Ltd.		554532634	manufacture(76986-019)

Revised: 12/2025

Guangdong Quadrant Ecological Technology Co., Ltd.