# K OXIDERM OP POST PROCEDURE SKIN CARE BRUISE- magnesium as corbyl phosphate cream EZEKIELCOSMETIC CO., LTD

Disclaimer: This drug has not been found by FDA to be safe and effective, and this labeling has not been approved by FDA. For further information about unapproved drugs, click here.

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#### **Drug Facts**

Magnesium Ascorbyl Phosphate

water, propylene glycol, alcohol etc

Reduces Discoloration on skin

keep out of reach of the children

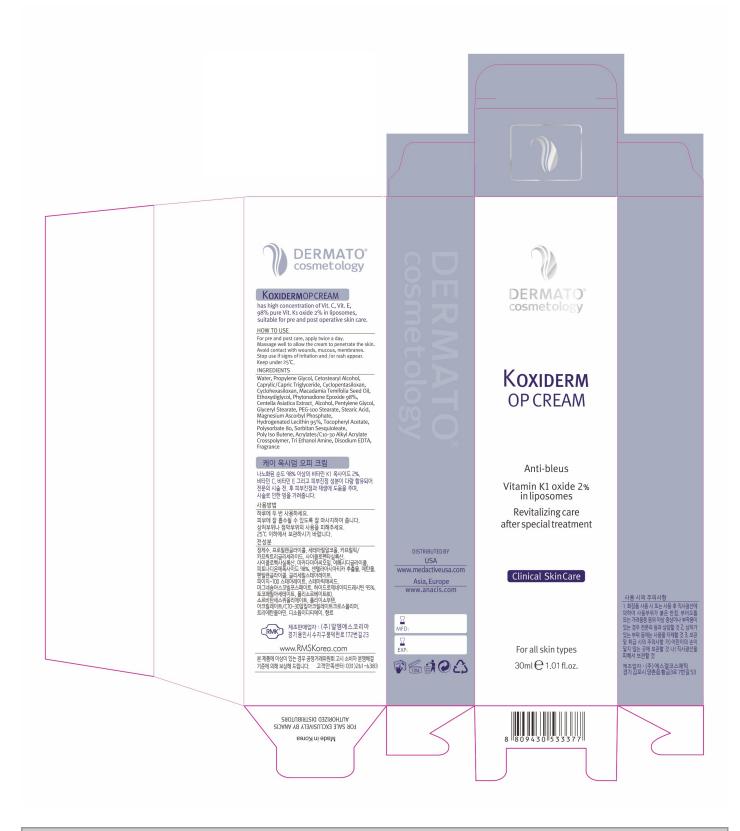
for 4~5 days before cosmetic application, apply thinly twice a day, morning and night, after cleansing the skin

apply K oxiderm OP cream right after cosmetic application

 $K\ oxiderm\ OP\ cream\ is\ recommended\ for\ 10\ to\ 15\ days\ as\ preparatory\ skin\ care\ before\ and\ after\ surgical\ and\ medical\ cosmetic\ procedure$ 

- 1. If the following symptoms occur after product use, stop using the product immediately and consult a dermatologist (continuous use can exacerbate the symptoms).
- 1) Occurrence of red spots, swelling, itchiness, and other skin irritation
- 2) If the symptoms above occur after the application area is exposed to direct sunlight
- 2. Do not use on open wounds, eczema, and other skin irritations
- 3. Precaution for Storage and Handling
- 1) Close the lid after use
- 2) Keep out of reach of infants and children
- 3) Do not to store in a place with high/low temperature and exposed to direct sunlight
- 4. Use as avoiding eye areas.

for external use only



#### K OXIDERM OP POST PROCEDURE SKIN CARE BRUISE

magnesium ascorbyl phosphate cream

Product Information			
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:71588-0011
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety		
Ingredient Name	Basis of Strength	Strength
MAGNESIUM ASCORBYL PHO SPHATE (UNII: 0 R8 22556 M5) (ASCORBIC ACID - UNII: PQ6 CK8 PD0 R)	MAGNESIUM ASCORBYL PHOSPHATE	0.5 g in 100 mL

Inactive Ingredients		
Ingredient Name	Strength	
WATER (UNII: 059QF0KO0R)		
ALCOHOL (UNII: 3K9958V90M)		
MAGNESIUM PHO SPHIDE (UNII: Q846538H9E)		

ı	Pac	ckaging			
l	#	# Item Code Package Description		<b>Marketing Start Date</b>	<b>Marketing End Date</b>
l	1 N	DC:71588-0011-1	30 mL in 1 TUBE; Type 0: Not a Combination Product	08/17/2017	

Marketing Information				
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date	
unapproved drug other		08/17/2017		

### Labeler - EZEKIELCOSMETIC CO., LTD (689851966)

## Registrant - EZEKIELCOSMETIC CO., LTD (689851966)

Establishment				
Name	Address	ID/FEI	Business Operations	
EZEKIELCOSMETIC CO., LTD		689851966	label(71588-0011), manufacture(71588-0011), pack(71588-0011)	

Revised: 8/2017 EZEKIELCOSMETIC CO., LTD