K OXIDERM OP POST PROCEDURE SKIN CARE BRUISE- magnesium ascorbyl 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.

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 PHOSPHATE (UNII: 0R822556M5) (ASCORBIC ACID - UNII:PQ6CK8PD0R)	MAGNESIUM ASCORBYL PHOSPHATE	0.5 g in 100 mL

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

Packaging				
#	**Item Code**	**Package Description**	**Marketing Start Date**	**Marketing End Date**
1	NDC: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