

VITA SOLUBALL- ascorbic acid, zinc oxide powder
Silexn Technology 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

Ascorbic acid 200mg, Zinc oxide 40mg per 1000mg

silicon dioxide

Antioxidation, Antiinflammatory on the skin

KEEP OUT OF REACH OF THE CHILDREN

Sprinkle on the affected area of the skin and rub it.

Once a day - below 100mg per day

Stop using it immediately if there are signs of skin rashes or hypersensitivity.

for topical use only

<i>Drug Facts</i>
<u>Active Ingredients</u> Ascorbic acid, Zinc oxide
<u>Uses</u> <ul style="list-style-type: none"> ■ Antioxidation, <u>Antiinflammatory</u> on the skin
<u>Warning</u> For topical use only When using this product <ul style="list-style-type: none"> ■ Do not use it on open wounds and sensitive skin ■ Stop use and ask a doctor if there are any abnormal symptoms or side effect such as red spots, swelling or itching during or immediately after use ■ Keep out of reach of children
<u>Directions</u> <ul style="list-style-type: none"> ■ Sprinkle on the affected area of the skin and rub it. ■ Once a day - below 100mg per day
<u>Other Information</u> - read the directions and warnings before use - avoid freezing and excessive heat above 40 degree C (<u>104 degree F</u>)
<u>Inactive Ingredients</u> Silicon dioxide
[MANUFACTURER] [DISTRIBUTOR]

VITA SOLUBALL			
ascorbic acid, zinc oxide powder			
Product Information			
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:84020-0001
Route of Administration	TOPICAL		
Active Ingredient/Active Moiety			
		Base of	

Ingredient Name	Basis of Strength	Strength
ZINC OXIDE (UNII: SOI2LOH54Z) (ZINC OXIDE - UNII:SOI2LOH54Z)	ZINC OXIDE	40 mg in 1000 mg
ASCORBIC ACID (UNII: PQ6CK8PD0R) (ASCORBIC ACID - UNII:PQ6CK8PD0R)	ASCORBIC ACID	200 mg in 1000 mg

Inactive Ingredients

Ingredient Name	Strength
BUTYLENE GLYCOL (UNII: 3XUS85K0RA)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:84020-0001-1	10000 mg in 1 BOTTLE; Type 0: Not a Combination Product	01/24/2024	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
unapproved drug other		01/24/2024	

Labeler - Silexn Technology Co.,Ltd. (987655536)

Registrant - Silexn Technology Co.,Ltd. (987655536)

Establishment

Name	Address	ID/FEI	Business Operations
Silexn Technology Co.,Ltd.		987655536	manufacture(84020-0001)

Revised: 1/2024

Silexn Technology Co.,Ltd.