

**MIRACLE LASER MASK- hyaluronate sodium cream**  
**TOAS 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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**72557-005\_Miracle Laser Mask**

Sodium Hyaluronate 0.005%

Moisturizing

Helps to keep skin moistness

Remove film and apply mask over cleansed face. Leave on for 15-30 minutes then remove.

For external use only.

Do not use on damaged or broken skin.

When using this product, keep out of eyes. Rinse with water to remove.

Stop using and ask a doctor if rash occurs.

Keep out of reach of the children. If product is swallowed, get medical help or contact a poison control center right away.

Water, Glycerin, Butylene Glycol, Propylene Glycol, Aloe Barbadensis Leaf Extract, Portulaca Oleracea Extract, Arginine, Carbomer, Polysorbate 80, Betaine, Allantoin, Methylparaben, Hydrolyzed Collagen, Sodium PCA, Disodium EDTA, Dipotassium Glycyrrhizate, Hamamelis Virginiana (Witchhazel) Water, Tocopheryl Acetate, Sodium Polyacrylate, Fragrance, Propylparaben, OligoPepTide-1



MIRACLE LASER MASK			
hyaluronate sodium cream			
<b>Product Information</b>			
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:72557-005
Route of Administration	TOPICAL		
<b>Active Ingredient/Active Moiety</b>			
Ingredient Name		Basis of Strength	Strength
HYALURONATE SODIUM (UNII: YSE9PPT4TH) (HYALURONIC ACID - UNII:S270N0TRQY)		HYALURONATE SODIUM	0.0014 g in 28 g
<b>Inactive Ingredients</b>			
Ingredient Name			Strength
WATER (UNII: 059QF0KO0R)			
GLYCERIN (UNII: PDC6A3C0OX)			
BUTYLENE GLYCOL (UNII: 3XUS85K0RA)			
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)			

<b>ALOE VERA LEAF</b> (UNII: ZY81Z83H0X)	
<b>ARGININE</b> (UNII: 94ZLA3W45F)	
<b>CARBOXYPOLYMETHYLENE</b> (UNII: 0A5MM307FC)	
<b>POLYSORBATE 80</b> (UNII: 6OZP39ZG8H)	
<b>BETAINE</b> (UNII: 3SCV180C9W)	
<b>ALLANTOIN</b> (UNII: 344S277G0Z)	

### Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:72557-005-02	10 in 1 CARTON	09/24/2018	
1	NDC:72557-005-01	28 g in 1 POUCH; Type 0: Not a Combination Product		

### Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
unapproved drug other		09/24/2018	

**Labeler** - TOAS Co., Ltd. (694485346)

**Registrant** - TOAS Co., Ltd. (694485346)

### Establishment

Name	Address	ID/FEI	Business Operations
TOAS Co., Ltd.		694485346	manufacture(72557-005)

Revised: 9/2018

TOAS Co., Ltd.