# UP AND UP DEEP CLEAN TONER- salicylic acid liquid TARGET CORPORATION

Disclaimer: Most OTC drugs are not reviewed and approved by FDA, however they may be marketed if they comply with applicable regulations and policies. FDA has not evaluated whether this product complies.

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#### **Up & Up Deep Clean Toner**

### **Active ingredient**

Salicylic Acid 0.5%

### **Purpose**

Acne Treatment

#### Uses

for the treatment of acne.

#### **Warnings**

For external use only

### When using this product

• skin irritation and dryness is more likely to occur if you use another topical acne medication at the same time. If irritation occurs only use one topical acne medication at a time. • avoid contact with eyes. If contact occurs, flush thoroughly with water.

### Keep out of reach of children.

If swallowed, get medical help or contact a Poison Control Center right away.

#### **Directions**

• clean the skin thoroughly before using this product • cover the entire affected area with a thin layer one to three times daily • because excessive drying of the skin may occur, start with one application daily, then gradually increase to two or three times daily if needed or as directed by doctor • if bothersome dryness or peeling occurs, reduce application to once a day or every other day.

#### Other information

• may stain some fabric

### **Inactive ingredients**

water, alcohol denat. (24.5 %), glycerin, isoceteth-20, PEG-32, sodium citrate, fragrance, propylene glycol, dimethicone propyl PG-betaine, benozphenone-4, algae extract, aloe barbadenis leaf juice, blue 1

Up & Up Deep Clean Toner

8 FL OZ (236.5 mL)

NDC 11676-936-11





#### **UP AND UP DEEP CLEAN TONER**

salicylic acid liquid

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Prod	uct	Inform	ation

WATER (UNII: 059QF0KO0R)

Product Type HUMAN OTC DRUG Item Code (Source) NDC:11673-936

Route of Administration TOPICAL

### **Active Ingredient/Active Moiety**

Ingredient Name Basis of Strength Strength

SALICYLIC ACID (UNII: O414PZ4LPZ) (SALICYLIC ACID - UNII:O414PZ4LPZ) SALICYLIC ACID 5 mg in 1 mL

Inactive Ingredients		
Ingredient Name	Strength	
ALOE VERA LEAF (UNII: ZY81Z83H0X)		
TRISODIUM CITRATE DIHYDRATE (UNII: B22547B95K)		
ALCOHOL (UNII: 3K9958V90M)		

GLYCERIN (UNII: PDC6A3C0OX)	
FD&C BLUE NO. 1 (UNII: H3R47K3TBD)	
POLYETHYLENE GLYCOL 1600 (UNII: 1212Z7S33A)	
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PHYMATOLITHON CALCAREUM (UNII: 6J1M3WA0ZK)	
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)	
SULISOBENZONE (UNII: 1W6L629B4K)	
ISOCETETH-20 (UNII: O020065R7Z)	
DIMETHICONE PROPYL PG-BETAINE (UNII: OB83A4S9K9)	

P	Packaging				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date	
1	NDC:11673- 936-11	236.5 mL in 1 BOTTLE; Type 0: Not a Combination Product	06/13/2014		

Marketing Information				
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date	
OTC monograph final	part333D	06/13/2014		

## Labeler - TARGET CORPORATION (006961700)

Revised: 10/2022 TARGET CORPORATION