

## **NATURAL ACNE SOLUTIONS 3 STEP REGIMEN- salicylic acid**

### **Burt's Bees**

*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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### **Active Ingredients**

SALICYLIC ACID 1.0%

### **Purpose**

Acne Treatment

PENETRATES PORES TO CONTROL ACNE.

HELPS PREVENT THE DEVELOPMENT OF NEW BLEMISHES

### **WARNINGS**

FOR EXTERNAL USE ONLY. USING OTHER TOPICAL ACNE MEDICATION AT THE SAME TIME OR IMMEDIATELY FOLLOWING USE OF THIS PRODUCT MAY INCREASE DRYNESS OR IRRITATION OF THE SKIN. IF THIS OCCURS, ONLY ONE MEDICATION SHOULD BE USED UNLESS DIRECTED BY A DOCTOR.

**Keep out of reach of children** IF SWALLOWED, GET MEDICAL HELP OR CONTACT A POISON CONTROL CENTER RIGHT AWAY.

### **DIRECTIONS**

APPLY TO FACE IN THE MORNING AND EVENING AFTER CLEANSING. IF BOTHERSOME DRYNESS OCCURS, REDUCE TO ONCE A DAY OR EVERY OTHER DAY.

### **INACTIVE INGREDIENTS**

WATER, HELIANTHUS ANNUUS (SUNFLOWER) SEED OIL, GLYCERIN, STEARIC ACID, SUCROSE POLYSTEARATE, SALIX NIGRA (WILLOW) BARK EXTRACT, BEESWAX, CYMBOPOGON SCHOENANTHUS EXTRACT, HAMAMELIS VIRGINIANA (WITCH HAZEL) LEAF EXTRACT, HUMULUS LUPULUS (HOPS) EXTRACT, EQUISETUM HIEMALE EXTRACT, HYDRASTIS CANADENSIS (GOLDENSEAL) EXTRACT, EPILOBIUM FLEISCHERI EXTRACT, LECITHIN, CHONDRUS CRISPUS (CARAGEENAN), FRAGRANCE, GLUCOSE, XANTHAN GUM, SUCROSE, SUCROSE STEARATE, POTASSIUM SORBATE, SODIUM CHLORIDE, GLUCOSE OXIDASE, LACTOPEROXIDASE, PHENOXYETHANOL.

### **QUESTIONS?**

**800-849-7112** OR [WWW.BURTSBEEES.COM](http://WWW.BURTSBEEES.COM)

### **ACTIVE INGREDIENTS**

SALICYLIC ACID 1.0%

### **PURPOSE**

## ACNE TREATMENT

### USES

PENETRATES PORES TO CONTROL ACNE.

HELPS PREVENT THE DEVELOPMENT OF NEW BLEMISHES.

### WARNINGS

FOR EXTERNAL USE ONLY. USING OTHER TOPICAL ACNE MEDICATION AT THE SAME TIME OR IMMEDIATELY FOLLOWING USE OF THIS PRODUCT MAY INCREASE DRYNESS OR IRRITATION OF THE SKIN. IF THIS OCCURS, ONLY ONE MEDICATION SHOULD BE USED UNLESS DIRECTED BY A DOCTOR.

**Keep out of reach of children.** IF SWALLOWED, GET MEDICAL HELP OR CONTACT A POISON CONTROL CENTER RIGHT AWAY.

### DIRECTIONS

WASH FACE IN THE MORNING AND EVENING AVOIDING THE EYE AREA. IF CONTACT OCCURS, RINSE THOROUGHLY WITH WATER.

### QUESTIONS?

**800-849-7112** OR [WWW.BURTSBEES.COM](http://WWW.BURTSBEES.COM)

### INACTIVE INGREDIENTS

WATER, DECYL GLUCOSIDE, LAURYL GLUCOSIDE, SUCROSE LAURATE, SALIX NIGRA (WILLOW) BARK EXTRACT, COCO-BETAINE, BETAINE, BETA VULGARIS (BEET) ROOT EXTRACT, CENOTHERA BIENNIS (EVENING PRIMROSE) LEAF EXTRACT, SODIUM COCOYL HYDROLYZED SOY PROTEIN, HYDROXYPROPYLTRIMONIUM HONEY, GLYCERIN, GLYCERYL OLEATE, COCO GLUCOSIDE, FRAGRANCE, GLUCOSE, ALCOHOL DENAT., GLUCOSE OXIDASE, LACTOPEROXIDASE, CITRIC ACID, SODIUM CHLORIDE, PHENOXYETHANOL.

Salicylic Acid 0.75%

Acne Treatment

### Uses

- Penetrates pores to control acne

### Warnings

For external use only. Using other topical acne medication at the same time or immediately following use of this product may increase dryness or irritation of the skin. If this occurs, only one medication should be used unless directed by a doctor.

**Keep out of reach of children.** If swallowed, get medical help and contact a Poison Control Center right away.

### Directions



1	NDC:26052-035-10	1 in 1 BOX; Type 0: Not a Combination Product	01/02/2014
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### Quantity of Parts

Part #	Package Quantity	Total Product Quantity
Part 1	0 BOTTLE, PLASTIC	1 mL in 100
Part 2	0 BOTTLE, WITH APPLICATOR	1 g in 100
Part 3	0 BOTTLE, PUMP	1 g in 100

### Part 1 of 3

## NAS PURIFYING CLEANSER

salicylic acid gel

### Product Information

Item Code (Source)	NDC:26052-010
Route of Administration	TOPICAL

### Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
SALICYLIC ACID (UNII: O414PZ4LPZ) (SALICYLIC ACID - UNII:O414PZ4LPZ)	SALICYLIC ACID	1 g in 100 mL

### Inactive Ingredients

Ingredient Name	Strength
GLYCERYL OLEATE (UNII: 4PC054V79P)	
DECYL GLUCOSIDE (UNII: Z17H97EA6Y)	
LAURYL GLUCOSIDE (UNII: 76LN7P7UCU)	
SUCROSE LAURATE (UNII: 05Q7CD0E49)	
COCO GLUCOSIDE (UNII: ICS790225B)	
GLUCOSE OXIDASE (UNII: 0T8392U5N1)	
GLYCERIN (UNII: PDC6A3C0OX)	
HYDROXYPROPYLTRIMONIUM CHLORIDE FRAGMENT (UNII: N30M4819DB)	
BETA VULGARIS (UNII: 4G174V5051)	
HYDROLYZED SOY PROTEIN (ENZYMATIC; 2000 MW) (UNII: 1394NXB9L6)	
ALCOHOL (UNII: 3K9958V90M)	
LACTOPEROXIDASE BOVINE (UNII: 01UIN5OC49)	
ANHYDROUS CITRIC ACID (UNII: XF417D3PSL)	
SODIUM CHLORIDE (UNII: 451W47IQ8X)	
PHENOXYETHANOL (UNII: HIE492ZZ3T)	
WATER (UNII: 059QF0K00R)	
SALIX NIGRA BARK (UNII: QU52J3A5B3)	
COCO-BETAINE (UNII: 03DH2IZ3FY)	
BETAINE (UNII: 3SCV180C9W)	
OENOTHERA BIENNIS LEAF (UNII: 153YBS895W)	

## Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:26052-010-02	145 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product		

## Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph final	part333D	02/01/2010	

## Part 2 of 3

### NAS TARGETED SPOT TREATMENT

salicylic acid liquid

## Product Information

Item Code (Source)	NDC:26052-011
Route of Administration	TOPICAL

## Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
SALICYLIC ACID (UNII: O414PZ4LPZ) (SALICYLIC ACID - UNII:O414PZ4LPZ)	SALICYLIC ACID	0.75 g in 100 g

## Inactive Ingredients

Ingredient Name	Strength
WATER (UNII: 059QF0KO0R)	
CALENDULA OFFICINALIS FLOWER (UNII: P0M7O4Y7YD)	
LEMON OIL (UNII: I9GRO824LL)	
BORAGO OFFICINALIS SEED (UNII: 2GXJ790US0)	
PARSLEY OIL (UNII: IXK9N7RJ7J)	
SALIX NIGRA BARK (UNII: QU52J3A5B3)	
TEA TREE OIL (UNII: VIF565UC2G)	
EUCALYPTUS GLOBULUS LEAF (UNII: S546YLW6E6)	
FENNEL OIL (UNII: 59AAO5F6HT)	
JUNIPER BERRY OIL (UNII: SZH16H44UY)	
ALCOHOL (UNII: 3K9958V90M)	
ACHILLEA MILLEFOLIUM (UNII: 2FXJ6SW4PK)	

## Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:26052-011-03	7.5 g in 1 BOTTLE, WITH APPLICATOR; Type 0: Not a Combination Product		

## Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph final	part333D	02/01/2010	

## Part 3 of 3

### NAS DAILY MOISTURIZING

salicylic acid lotion

## Product Information

Item Code (Source)	NDC:26052-012
Route of Administration	TOPICAL

## Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
SALICYLIC ACID (UNII: O414PZ4LPZ) (SALICYLIC ACID - UNII:O414PZ4LPZ)	SALICYLIC ACID	1 g in 100 g

## Inactive Ingredients

Ingredient Name	Strength
POTASSIUM SORBATE (UNII: 1VPU26JZZ4)	
HAMAMELIS VIRGINIANA LEAF (UNII: T07U1161SV)	
LEVOGLUCOSE (UNII: 02833ISA66)	
PHENOXYETHANOL (UNII: HIE492ZZ3T)	
WATER (UNII: 059QF0KO0R)	
EPILOBIUM FLEISCHERI WHOLE (UNII: 8E2KLS8J8K)	
SUNFLOWER OIL (UNII: 3W1JG795YI)	
GLYCERIN (UNII: PDC6A3C0OX)	
SALIX NIGRA BARK (UNII: QU52J3A5B3)	
WHITE WAX (UNII: 7G1J5DA97F)	
CYMOPOGON SCHOENANTHUS TOP (UNII: 9SJ1LW39W)	
HUMULUS LUPULUS WHOLE (UNII: 912A6Q1N4A)	
XANTHAN GUM (UNII: TTV12P4NEE)	
SUCROSE STEARATE (UNII: 274KW0O50M)	
SODIUM CHLORIDE (UNII: 451W47IQ8X)	
GLUCOSE OXIDASE (UNII: 0T8392U5N1)	
EQUISETUM HYEMALE (UNII: 59677RXH25)	
LECITHIN, SOYBEAN (UNII: 1DI56QDM62)	

CHONDRUS CRISPUS CARRAGEENAN (UNII: UE856F2T78)	
SUCROSE (UNII: C151H8M554)	
LACTOPEROXIDASE BOVINE (UNII: 01UIN5OC49)	
STEARIC ACID (UNII: 4ELV7Z65AP)	
HYDRASTIS CANADENSIS WHOLE (UNII: R763EBH88T)	

### Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:26052-012-01	55 g in 1 BOTTLE, PUMP; Type 0: Not a Combination Product		

### Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph final	part333D	01/02/2014	

### Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph final	part333D	01/02/2014	

**Labeler** - Burt's Bees (613480946)