PROPOLIS NASAL PLUS- plus nasal liquid
PROPOLIS NASAL- nasal liquid
SUNSCREEN SPF50- non nano zinc oxide cream
SUNSCREEN FOR FACE SPF50- non nano zinc oxide cream
AFTERSUN- non nano zinc oxide cream
DIAPER RASH CREAM- non nano zinc oxide cream
SUNSCREEN FOR KIDS SPF50- non nano zinc oxide cream
S.O.S CREAM INTENSIVE MOISTURIZING SKIN PROTECTANT- non nano zinc oxide cream
ACNE CONTROL SERUM- salicylic acid liquid

SBS BILIMSEL BIO COZUMLER SANAYI VE TICARET ANONIM SIRKETI PINAR SUBESI

Active Ingredient(s)

Alcohol 80% v/v. Purpose: Antiseptic

Purpose

Antiseptic, Hand Sanitizer

Use

Hand Sanitizer to help reduce bacteria that potentially can cause disease. For use when soap and water are not available.

Warnings

For external use only. Flammable. Keep away from heat or flame

Do not use

- in children less than 2 months of age
- on open skin wounds

When using this product keep out of eyes, ears, and mouth. In case of contact with eyes, rinse eyes thoroughly with water.

Stop use and ask a doctor if irritation or rash occurs. These may be signs of a serious condition.

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

Stop use and ask a doctor if irritation or rash occurs. These may be signs of a serious condition.

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

Directions

- Place enough product on hands to cover all surfaces. Rub hands together until dry.
- Supervise children under 6 years of age when using this product to avoid swallowing.

Other information

- Store between 15-30C (59-86F)
- Avoid freezing and excessive heat above 40C (104F)

Inactive Ingredients

Acne Control Serum

30 ml NDC: 82123-009



ACNE CONTROL SERUM

Clears acne and help prevent new breakouts

Hydrating

- Salicylic Acid
- ✓ Bee Venom
- → Anatolian Propolis (A.P.E.®)
- Hyaluronic Acid
- √ Tea Tree Oil

1fl oz. (30 ml)

Drug Facts

Active Ingredient

Salicylic acid 0.5%

Purpose

.. Acne treatment

Uses

- · Clears acne blemishes and allows skin to heal
- Helps prevent new acne blemishes from forming

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.

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 applying 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 a doctor
- if bothersome dryness or peeling occurs, reduce application to once a day or every other day

Inactive Ingredients: aqua, glycerin, rosa multiflora fruit extract, actinidia polygama fruit extract, beta vulgaris (beet) root extract, propanediol (and) xylitol (and) caprylic acid, hydroxyethyl cellulose, anatolian propolis extract, melaleuca alternifolia leaf oil, hyaluronic acid, bee venom

NDC 82123-008

Lot:

Product of Türkiye



90853 **"** 00350



Mineral Sunscreen Cream formulated with non-nano mineral zinc to deliver effective UVA/UVB protection for you and your loved ones. Produced with (MicNo® technology) novel and superiormineral-based broadspectrum UVA-UVB filter.

Drug Facts

Helps protect the skin against the harmful effects of the sun. Produced with natural ingredients. Suitable for sensitive skin. Sheer Tinted formula. No white cast.

Active Ingredients

Purpose

Non-nano Zinc Oxide 25%

Sunscreen

Uses: ■ Helps prevent sunburn ■ If used with other sun protection measures (see Directions), it helps lower the risk of skin cancer and early skin aging caused by the sun.

Warning; For external use only.

Do not use on open wounds, cuts, or when skin is open.

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

Stop use and ask a doctor if rash occurs.

Directions: ■ Apply liberally 15 minutes before sun exposure. ■ Children under 6 months old: Ask a doctor. Reapply: ■ After 80 minutes of swimming or sweating. ■ Immediately after towel drying. ■ At least every 2 hours.

Other Information Protect the product in its container from excessive heat and direct sun. Water resistant (80 minutes).

Inactive Ingredients: Aqua, Coco-Caprylate/Caprate, Dicaprylyl Carbonate, Olus Oil, Glycerin, Aloe Barbadensis Leaf Juice, Butyrospermum Parkii (Shea) Butter, Panthenol (Vitamin B5) Glyceryl Citrate/Lactate/Linoleate/Oleate, Propolis Extract, Tocopheryl Acetate (Vitamin E), Argania Spinosa Fruit Extract, Sodium Stearoyl Glutamate, Magnesium Sulfate.

FREE OF x Parabens x Petrolatum x Dyes x Oxybenzone x Octinoxate x Titanium Dioxide

ANATOLIAN PROPOLIS

Propolis is a natural bee product that bees collect from plants to protect their hives from microbes. It is a very rich natural antioxidant. We source our propolis from pristine Anatolian mountains, one of the most biodiverse regions in the world. Therefore Anatolian Propolis has a very special phenolic profile.

DISTRIBUTED BY: SBS AMERICAS INC. 42 Broadway Suite 12-403 New York, NY 10004

www.beeandyou.com - info@beeandyou.com

(i) (ii) **J*/beeandyounatural**

This investment is partly financed with funding by the European Union



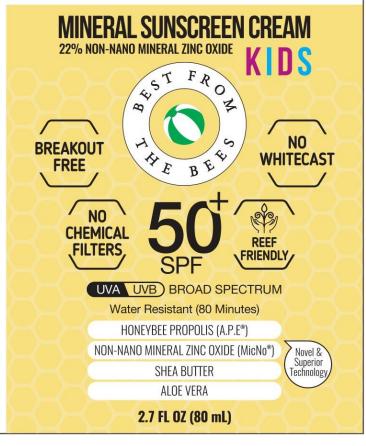




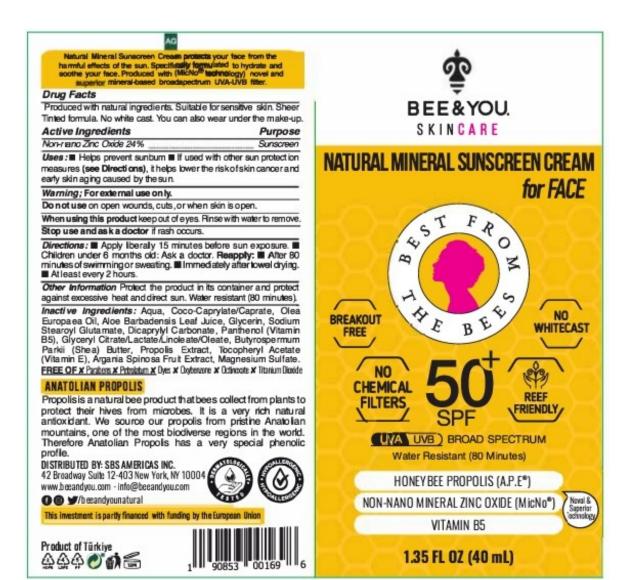
Product of Türkiye







Sunscreen for face SPF50



Aftersun



rich

Novel &

Superior

Technology

Diaper Rash Cream





Natural baby diaper cream for babies to prevent and heal rashes and soothe the skin

Drug Facts

Active Ingredients

Purpose

Non-nano Zinc Oxide 22%--

Skin Protectant

Uses

- Helps treat and prevent diaper rash
- Protects chafed skin due to diaper rash and helps seal out wetness

Warning;

For external use only.

When using this product • do not get into eyes

Stop use and ask a doctor if • condition worsen • symptoms last more than 7 days or clear up and occur within a few days.

Keep out of reach of children. If swallowed, get medical help or contact a Poison Control Center immediately

Directions

- Change wet or soiled diapers promptly
- Cleanse the diaper area and allow to dry
- Apply cream liberally as often as necessary with each diaper change especially at bedtime or any time exposure to wet diapers.

Other Information

- Store between 20° to 25°C (68° to 77°F)
- Do not use if safety seal is broken

Inactive Ingredients

Butyrospermum Parkii Butter, Olus Oil, Simmondsia Chinensis Seed Oil, Panthenol, Coco-Caprylate / Caprate, Aloe Barbadensis Leaf Juice, Glycerin, Polyglyceryl-2 Dipolyhydroxystearate, Polyglyceryl-3 Diisostearate, Jojoba Esters, Helianthus Annuus Seed Wax, Aqua, Propolis Extract, Diisostearoyl Polyglyceryl-3 Dimer / Dilinoleate, Acacia Decurrens Flower Wax, Cera Alba, Polyglyceryl-4 Diisostearate / Polyhydroxystearate / Sebacate. Caprylic / Capric Triglyceride.

82123-005

Propolis Nasal Plus





Drug Facts

Active Ingredient

Purpose

Oxymetazoline hydrochloride 0.05%.

... nasal decongestant

Uses

- temporarily relieves nasal congestion due to:
 - common cold • hay fever (allergic rhinitis)
- other upper respiratory allergies shrinks swollen nasal membranes so
- you can breathe more freely promotes nasal and sinus drainage
- temporarily relieves sinus congestion and pressure

Warnings

Ask a doctor before use if you have

- heart disease
- high blood pressure
- thyroid diesase
- diabetes
- trouble urinating due to an enlarged prostate gland

When using this product

- do not exceed recommended dosage
- do not use this product for more than 3 days. use only as directed. Frequent or prolonged use may cause nasal congestion to recur or worsen.
- temporary discomfort such as burning, stinging, sneezing or an increase in nasal discharge may occur
- use of this container by more than one person may spread infection

Stop use and ask doctor if symptoms persist.

If pregnant or breast-feeding, ask a health professional before use.

Drug Facts (continued)

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

Directions

- adults and children 6 to under 12 years of age (with adult supervision): 2 or 3 sprays in each nostril not more often than every 10 to 12 hours. Do not exceed 2 doses in any 24-hour period.
- children under 6 years of age: ask a doctor

Shake well before use

to spray, insert the nozzle into the nostril and depress the pump completely while breathing in through the nose.

Inactive ingredients

Anatolian propolis extract, purified water, eucalyptus oil, glycerin, sea salt

Propolis Nasal



Drug Facts

Active Ingredient

Oxymetazoline hydrochloride 0.05%.....

decongestant

Purpose

Uses

- temporarily relieves nasal congestion due to:
 - common cold hay fever (allergic rhinitis)
- other upper respiratory allergiesshrinks swollen nasal membranes so
- shrinks swollen nasal membranes so you can breathe more freely
- promotes nasal and sinus drainage
- temporarily relieves sinus congestion and pressure

Warnings

Ask a doctor before use if you have

- heart disease
- high blood pressure
- thyroid diesase
- diabetes
- trouble urinating due to an enlarged prostate gland

When using this product

- do not exceed recommended dosage
- do not use this product for more than 3 days. use only as directed. Frequent or prolonged use may cause nasal congestion to recur or worsen.
- temporary discomfort such as burning, stinging, sneezing or an increase in nasal discharge may occur
- use of this container by more than one person may spread infection

Stop use and ask doctor if symptoms persist.

If pregnant or breast-feeding, ask a health professional before use.

Drug Facts (continued)

Keep out of reach of children.

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

Directions

- adults and children 6 to under 12 years of age (with adult supervision): 2 or 3 sprays in each nostril not more often than every 10 to 12 hours. Do not exceed 2 doses in any 24-hour period.
- children under 6 years of age: ask a doctor

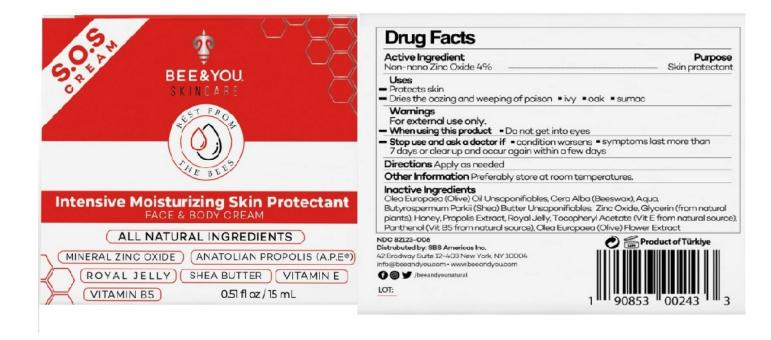
Shake well before use

to spray, insert the nozzle into the nostril and depress the pump completely while breathing in through the nose.

Inactive ingredients

Anatolian propolis extract, purified water, eucalyptus oil, glycerin, sea salt

SOS Cream



SOS Cream

Sunscreen SPF50



Mineral Sunscreen Cream formulated with non-nano mineral zinc to deliver effective UVA/UVB protection for you and your loved ones. Produced with (MicNo® technology) novel and superiormineral-based broadspectrum UVA-UVB filter.

Drug Facts

Helps protect the skin against the harmful effects of the sun. Produced with natural ingredients. Suitable for sensitive skin. Sheer Tinted formula.

No white cast. Active Ingredients

Non-nano Zinc Oxide 22% ...

Sunscreen

Uses: ■ Helps prevent sunburn ■ If used with other sun protection measures (see Directions), it helps lower the risk of skin cancer and early skin aging caused by the sun

Warning; For external use only.

Do not use on open wounds, cuts, or when skin is open.

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

Stop use and ask a doctor if rash occurs.

Directions: ■ Apply liberally 15 minutes before sun exposure. ■ Children under 6 months old: Ask a doctor. Reapply: ■ After 80 minutes of swimming or sweating. ■ Immediately after towel drying. ■ At least every 2 hours.

Other Information Protect the product in its container from excessive heat and direct sun. Water resistant (80 minutes).

Inactive Ingredients: Aqua, Coco-Caprylate/Caprate, Glycerin, Simmontsia Chinensis (Jojoba) Seed Oil, Aloe Barbadensis Leaf Juice, Glyceryl Citrate/Lactate/Linoleate/Oleate, Sodium Stearoyl Glutamate, Butyrospermum Parkii (Shea) Butter, Dicaprylyl Carbonate, Tocopheryl Acetate (Vitamin E), Panthenol (Vitamin B5), Propolis Extract, Argania Spinosa Fruit Extract, Magnesium Sulfate.

FREE OF X Parabens X Petrolatum X Dyes X Oxybenzone X Octinoxate X Titanium Dioxide

ANATOLIAN PROPOLIS

Propolis is a natural bee product that bees collect from plants to protect their hives from microbes. It is a very rich natural antioxidant. We source our propolis from pristine Anatolian mountains, one of the most biodiverse regions in the world. Therefore Anatolian Propolis has a very special phenolic profile.

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★ beeandyounatural



Product of Türkiye







PROPOLIS NASAL PLUS

plus nasal liquid

Product Information

Product Type HUMAN OTC DRUG Item Code (Source) NDC:82123-010

NASAL **Route of Administration**

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
	OXYMETAZ OLINE HYDROCHLORIDE	0.015 mg in 30 mL

Inactive Ingredients	
Ingredient Name	Strength
EUCALYPTUS OIL (UNII: 2R040NI662)	

WATER (UNII: 059QF0KO0R)	
SEA SALT (UNII: 87GE52P74G)	
PROPOLIS WAX (UNII: 6Y8XYV2NOF)	
GLYCERIN (UNII: PDC6A3C0OX)	

Packaging			
# Item Code	Package Description	Marketing Start Date	Marketing End Date
NDC:82123- 010-31	1 in 1 BOX	10/03/2023	
NDC:82123- 010-30	30 mL in 1 BOTTLE, GLASS; Type 0: Not a Combination Product		

Marketing Information			
Marketing Application Number or Monograph Marketing Start Marketing End Category Citation Date Date			
OTC Monograph Drug	M012	10/03/2023	

PROPOLIS NASAL

nasal liquid

Product Information			
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:82123-009
Route of Administration	NASAL		

Active Ingredient/Active Moiety				
Ingredient Name	Basis of Strength	Strength		
OXYMETAZOLINE HYDROCHLORIDE (UNII: K89MJ0S5VY) (OXYMETAZOLINE - UNII:8VLN5B44ZY)	OXYMETAZ OLINE HYDROCHLORIDE	0.015 mg in 30 mL		

Inactive Ingredients			
Ingredient Name	Strength		
EUCALYPTUS OIL (UNII: 2R04ONI662)			
SEA SALT (UNII: 87GE52P74G)			
GLYCERIN (UNII: PDC6A3C0OX)			
PROPOLIS WAX (UNII: 6Y8XYV2NOF)			
WATER (UNII: 059QF0KO0R)			

Packaging			
# Item Code	Package Description	Marketing Start Date	Marketing End Date
NDC-02122			

	NDC:82123-	1 in 1 BOX	10/02/2023	
1	NDC:82123- 009-30	30 mL in 1 BOTTLE, GLASS; Type 0: Not a Combination Product		

Marketing Information			
Marketing Application Number or Monograph Marketing Start Marketing End Category Citation Date Date			
OTC Monograph Drug	M012	10/02/2023	

SUNSCREEN SPF50

non nano zinc oxide cream

Product Information			
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:82123-011

Route of Administration TOPICAL

Active Ingredient/Active Moiety			
Ingredient Name	Basis of Strength	Strength	
ZINC OXIDE (UNII: SOI2LOH54Z) (ZINC OXIDE - UNII:SOI2LOH54Z)	ZINC OXIDE	12 mg in 80 mL	

Inactive Ingredients		
Ingredient Name	Strength	
WATER (UNII: 059QF0KO0R)		
BUTYROSPERMUM PARKII (SHEA) BUTTER UNSAPONIFIABLES (UNII: 0C9AC7D6XU)		
JOJOBA OIL (UNII: 724GKU717M)		
.ALPHATOCOPHEROL ACETATE (UNII: 9E8X80D2L0)		
PROPOLIS WAX (UNII: 6Y8XYV2NOF)		
PANTHENOL (UNII: WV9CM0O67Z)		
COCO-CAPRYLATE/CAPRATE (UNII: 8D9H4QU99H)		
MAGNESIUM SULFATE ANHYDROUS (UNII: ML30MJ2U7I)		
ALOE VERA LEAF (UNII: ZY81Z83H0X)		
ARGAN OIL (UNII: 4V59G5UW9X)		
DICAPRYLYL CARBONATE (UNII: 609A3V1SUA)		
SODIUM STEAROYL GLUTAMATE (UNII: 65A9F4P024)		
GLYCERYL CITRATE (UNII: 4987GT719I)		
GLYCERIN (UNII: PDC6A3C0OX)		

Product Characteristics			
Color	Score		
Shape	Size		
Flavor	Imprint Code		
Contains			

ı	Packaging			
;	# Item Code	Package Description	Marketing Start Date	Marketing End Date
	NDC:82123-011-	1 in 1 TUBE	06/28/2021	
	NDC:82123-011-	80 mL in 1 TUBE; Type 0: Not a Combination Product		



Natural Mineral Sunscreen Cream protects your face from the harmful effects of the sun. Specifically formulated to hydrate and soothe your face.

Drug Facts

Helps protect the skin against the harmful effects of the sun. Produced with natural ingredients, Suitable for sensitive skin. Sheer Tinted formula. No white cast. You can also wear under the make-up.

Active Ingredients Purpose
Non-nano Zinc Oxide 15% Sunscreen

Uses: ■ Helps prevent sunburn ■ If used with other sun protection measures (see Directions), it helps lower the risk of skin cancer and early skin aging caused by the sun.

Warning; For external use only.

Do not use on open wounds, cuts, or when skin is open.

When using this product keep out of eyes. Rinse with water to remove. Stop use and ask a doctor if rash occurs.

Directions: ■ Apply liberally 15 minutes before sun exposure. ■ Children under 6 months old: Ask a doctor. Reapply: ■ After 80 minutes of swimming or sweating. ■ Immediately after towel drying. ■ At least every 2 hours.

Other Information Protect the product in its container and protect against excessive heat and direct sun.

Inactive Ingredients: Aqua, Coco-Caprylate/Caprate, Olea Europaea Oil, Aloe Barbadensis Leaf Juice, Glycerin, Sodium Stearoyl Glutamate, Dicaprylyl Carbonate, Panthenol (Vitamin B5), Glyceryl Citrate/Lactate/Linoleate/Oleate, Butyrospermum Parkii (Shea) Butter, Propolis Extract, Tocopheryl Acetate (Vitamin E), Argania Spinosa Fruit Extract, Magnesium Sulfate.

FREE OF x Parabens x Petrolatum x Dyes x Oxybenzone x Octinoxate x Titanium Dioxide

Marketing Information			
Application Number or Monograph Citation	Marketing Start Date	Marketing End Date	
1020	06/28/2021		
	Application Number or Monograph Citation	Application Number or Monograph Marketing Start Citation Date	

SUNSCREEN FOR FACE SPF50

non nano zinc oxide cream

Product Information			
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:82123-013
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety			
Ingredient Name	Basis of Strength	Strength	
ZINC OXIDE (UNII: SOI2LOH54Z) (ZINC OXIDE - UNII:SOI2LOH54Z)	ZINC OXIDE	6 mg in 40 mL	

Inactive Ingredients		
Ingredient Name	Strength	
COCO-CAPRYLATE/CAPRATE (UNII: 8D9H4QU99H)		
OLEA EUROPAEA (OLIVE) OIL UNSAPONIFIABLES (UNII: XO45V955LT)		
ALOE VERA LEAF (UNII: ZY81Z83H0X)		
DICAPRYLYL CARBONATE (UNII: 609A3V1SUA)		
PANTHENOL (UNII: WV9CM0O67Z)		
BUTYROSPERMUM PARKII (SHEA) BUTTER UNSAPONIFIABLES (UNII: 0C9AC7D6XU)		
MAGNESIUM SULFATE ANHYDROUS (UNII: ML30MJ2U7I)		
GLYCERYL CITRATE (UNII: 4987GT719I)		
PROPOLIS WAX (UNII: 6Y8XYV2NOF)		
.ALPHATOCOPHEROL ACETATE (UNII: 9E8X80D2L0)		
WATER (UNII: 059QF0KO0R)		
GLYCERIN (UNII: PDC6A3C0OX)		
SODIUM STEAROYL GLUTAMATE (UNII: 65A9F4P024)		
ARGAN OIL (UNII: 4V59G5UW9X)		

Product Characteristics	
Color	Score
Shape	Size
Flavor	Imprint Code
Contains	

P	ackaging			
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:82123-013- 41	1 in 1 TUBE	06/28/2021	
1	NDC:82123-013- 40	40 mL in 1 TUBE; Type 0: Not a Combination Product		



Natural Mineral Sunscreen Cream protects your face from the harmful effects of the sun. Specifically formulated to hydrate and soothe your face.

Drug Facts

Helps protect the skin against the harmful effects of the sun. Produced with natural ingredients. Suitable for sensitive skin. Sheer Tinted formula. No white cast. You can also wear under the make-up.

Active Ingredients Purpose

Non-nano Zinc Oxide 15%

Sunscreen

Uses: ■ Helps prevent sunburn ■ If used with other sun protection measures (see Directions), it helps lower the risk of skin cancer and early skin aging caused by the sun.

Warning; For external use only.

Do not use on open wounds, cuts, or when skin is open.

When using this product keep out of eyes. Rinse with water to remove. Stop use and ask a doctor if rash occurs.

Directions: ■ Apply liberally 15 minutes before sun exposure. ■ Children under 6 months old: Ask a doctor. Reapply: ■ After 80 minutes of swimming or sweating. ■ Immediately after towel drying. ■ At least every 2 hours.

Other Information Protect the product in its container and protect against excessive heat and direct sun.

Inactive Ingredients: Aqua, Coco-Caprylate/Caprate, Olea Europaea Oli, Aloe Barbadensis Leaf Julice, Glycerin, Sodium Stearoyl Glutamate, Dicaprylyl Carbonate, Panthenol (Vitamin B5), Glyceryl Citrate/Lactate/Linoleate/Oleate, Butyrospermum Parkii (Shea) Butter, Propolis Extract, Tocopheryl Acetate (Vitamin E), Arqania Spinosa Fruit Extract, Magnesium Sulfate.

FREE OF x Parabens x Petrolatum x Dyes x Oxybenzone x Octinoxate x Titanium Dioxide

Marketing Information			
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M020	06/28/2021	

AFTERSUN

non nano zinc oxide cream

Product Information

Product Type HUMAN OTC DRUG Item Code (Source) NDC:82123-004

Route of Administration TOPICAL

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
ZINC OXIDE (UNII: SOI2LOH54Z) (ZINC OXIDE - UNII:SOI2LOH54Z)	ZINC OXIDE	1.6 mg in 80 mL

Inactive Ingredients Ingredient Name PROPOLIS WAX (UNII: 6Y8XYV2NOF) Strength

SODIUM STEAROYL GLUTAMATE (UNII: 65A9F4P024)

PHYSALIS ANGULATA (UNII: W4TKW9D5GG)
ALOE VERA LEAF (UNII: ZY81Z83H0X)

GLYCERYL CITRATE (UNII: 4987GT719I)

WHITE WAX (UNII: 7G1J5DA97F)

.ALPHA.-TOCOPHEROL ACETATE (UNII: 9E8X80D2L0)

GLYCERIN (UNII: PDC6A3C0OX)

PANTHENOL (UNII: WV9CM0O67Z)

WATER (UNII: 059QF0K0OR)

BUTYROSPERMUM PARKII (SHEA) BUTTER UNSAPONIFIABLES (UNII: 0C9AC7D6XU)

COCO-CAPRYLATE/CAPRATE (UNII: 8D9H4QU99H)

SESAME OIL (UNII: QX10HYY4QV)

XANTHAN GUM (UNII: TTV12P4NEE)

ARGAN OIL (UNII: 4V59G5UW9X)

MAGNESIUM SULFATE ANHYDROUS (UNII: ML30MJ2U7I)

Product Characteristics			
Color	Score		
Shape	Size		
Flavor	Imprint Code		
Contains			

ı	Packaging				
	#	Item Code	Package Description	Marketing Start Date	Marketing End Date
	1	NDC:82123-004- 81	1 in 1 TUBE	06/28/2021	
	1	NDC:82123-004- 80	80 mL in 1 TUBE; Type 0: Not a Combination Product		



Natural After Sun Lotion to soothe and moisturize your skin after sun exposure.

Drug Facts

Helps protect the skin against the harmful effects of the sun. Produced with natural ingredients. Suitable for sensitive skin. Sheer Tinted formula. No white cast.

Active Ingredients	Purpose	
Non-nano Zinc Oxide 2%	 Sunscreen	

Uses: ■ Helps prevent sunburn ■ If used with other sun protection measures (see Directions), it helps lower the risk of skin cancer and early skin aging caused by the sun.

Warning; For external use only.

Do not use on open wounds, cuts, or when skin is open.

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

Stop use and ask a doctor if rash occurs.

Directions: ■ Apply liberally to all areas of the skin for soothing relief.
■ Gently massage into affacted areas ■ Reapply as frequently as you like
■ For best results, apply right after sunburn occurs.

Other Information Protect the product in its container from excessive heat and direct sun

Inactive Ingredients: Aqua, Glycerin, Coco-Caprylate/Caprate, Butyrospermum Parkii (Shea) Butter, Sesamum Indicum Seed Oil, Aloe Barbadensis Leaf Juice, Glyceryl Citrate/Lactate/Linoleate/Oleate, Sodium Stearoyl Glutamate, Panthenol (Vitamin BS), Cera Alba, Tocopheryl Acetate (Vitamin E), Argania Spinosa Fruit Extract, Physalis Angulata Extract, Xanthan Gum, Propolis Extract, Magnesium Sulfate.

FREE OF x Parabens x Petrolatum x Dyes x Oxybenzone x Octinoxate x Titanium Dioxide

Marketing Information			
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M020	06/28/2021	

DIAPER RASH CREAM

non nano zinc oxide cream

Product Information

Product Type HUMAN OTC DRUG Item Code (Source) NDC:82123-005

Route of Administration TOPICAL

Active Ingredient/Active Moiety				
Ingredient Name	Basis of Strength	Strength		
ZINC OXIDE (UNII: SOI2LOH54Z) (ZINC OXIDE - UNII:SOI2LOH54Z)	ZINC OXIDE	11 mg in 50 mg		

Inactive Ingredients	
Ingredient Name	Strength
BUTYROSPERMUM PARKII (SHEA) BUTTER UNSAPONIFIABLES (UNII: 0C9AC7D6XU)	
HYDROLYZED JOJOBA ESTERS (POTASSIUM SALTS) (UNII: CH428W5O62)	
WHITE WAX (UNII: 7G1J5DA97F)	
POLYGLYCERYL-3 OLEATE (UNII: XRQ165498B)	
PANTHENOL (UNII: WV9CM0O67Z)	
COCO-CAPRYLATE/CAPRATE (UNII: 8D9H4QU99H)	
ALOE VERA LEAF (UNII: ZY81Z83H0X)	
GLYCERIN (UNII: PDC6A3C0OX)	
POLYGLYCERYL-2 DIPOLYHYDROXYSTEARATE (UNII: 9229XJ4V12)	
POLYGLYCERYL-3 DIISOSTEARATE (UNII: 46P231IQV8)	
CAPRYLIC/CAPRIC/LAURIC TRIGLYCERIDE (UNII: FJ1H6M2JG9)	
.ALPHATOCOPHEROL ACETATE (UNII: 9E8X80D2L0)	
JOJOBA OIL (UNII: 724GKU717M)	
HELIANTHUS ANNUUS SEED WAX (UNII: 42DG15CHXV)	
POLYGLYCERYL-4 ISOSTEARATE (UNII: 820DPX33S7)	
POLYGLYCERIN-3 (UNII: 4A0NCJ6RD6)	
PALM OIL (UNII: 5QUO05548Z)	
PALM KERNEL OIL (UNII: B0S90M0233)	
COCONUT OIL (UNII: Q9L0O73W7L)	
CANOLA OIL (UNII: 331KBJ17RK)	

Product Characteristics	
Color	Score
Shape	Size
Flavor	Imprint Code

Contains

P	Packaging			
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:82123-005- 51	1 in 1 TUBE	06/28/2021	
1	NDC:82123-005- 50	50 mg in 1 TUBE; Type 0: Not a Combination Product		



Natural baby diaper cream for babies to prevent and heal rashes and soothe the skin

Drug Facts

Active Ingredients

Purpose

Non-nano Zinc Oxide 22%----- Skin Protectant

Uses

- Helps treat and prevent diaper rash
- Protects chafed skin due to diaper rash and helps seal out wetness

Warning;

For external use only.

When using this product · do not get into eyes

Stop use and ask a doctor if · condition worsen · symptoms last more than 7 days or clear up and occur within a few days.

Keep out of reach of children. If swallowed, get medical help or contact a Poison Control Center immediately

Directions

- Change wet or soiled diapers promptly
- Cleanse the diaper area and allow to dry
- Apply cream liberally as often as necessary with each diaper change especially at bedtime or any time exposure to wet diapers.

Other Information

Store between 20° to 25°C (68° to 77°F)

Do not use if safety seal is broken

Inactive Ingredients

Butyrospermum Parkii Butter, Olus Oil, Simmondsia Chinensis Seed Oil, Panthenol, Coco-Caprylate / Caprate, Aloe Barbadensis Leaf Juice, Glycerin, Polyglyceryl-2 Dipolyhydroxystearate, Polyglyceryl-3 Diisostearate, Jojoba Esters, Helianthus Annuus Seed Wax, Aqua, Propolis Extract, Diisostearoyl Polyglyceryl-3 Dimer / Dilinoleate, Acacia Decurrens Flower Wax, Cera Alba, Polyglyceryl-4 Diisostearate / Polyhydroxystearate / Sebacate, Caprylic / Capric Triglyceride, Polyglyceryl-3 Oleate, Tocopheryl Acetate, Polyglycerin-3

Marketing	Application Number or Monograph	Marketing Start	Marketing End
Category	Citation	Date	Date
OTC Monograph Drug	M016	06/28/2021	

SUNSCREEN FOR KIDS SPF50

non nano zinc oxide cream

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:82123-012
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Route of Administration TOPICAL

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
ZINC OXIDE (UNII: SOI2LOH547) (ZINC OXIDE - UNII: SOI2LOH547)	ZINC OXIDE	17.6 mg in 80 mg

Inactive Ingredients

Ingredient Name	Strength
MAGNESIUM SULFATE ANHYDROUS (UNII: ML30MJ2U7I)	
PALM KERNEL OIL (UNII: B0S90M0233)	
COCONUT OIL (UNII: Q9L0O73W7L)	
CANOLA OIL (UNII: 331KBJ17RK)	
WATER (UNII: 059QF0KO0R)	
COCO-CAPRYLATE/CAPRATE (UNII: 8D9H4QU99H)	
BUTYROSPERMUM PARKII (SHEA) BUTTER UNSAPONIFIABLES (UNII: 0C9AC7D6XU)	
ARGAN OIL (UNII: 4V59G5UW9X)	
GLYCERYL CITRATE (UNII: 4987GT719I)	
PROPOLIS WAX (UNII: 6Y8XYV2NOF)	
.ALPHATOCOPHEROL ACETATE (UNII: 9E8X80D2L0)	

PANTHENOL (UNII: WY9CM0067Z)

DICAPRYLYL CARBONATE (UNII: 609A3V1SUA)

ALOE VERA LEAF (UNII: ZY81Z83H0X)

PALM OIL (UNII: 5QU005548Z)

SODIUM STEAROYL GLUTAMATE (UNII: 65A9F4P024)

GLYCERIN (UNII: PDC6A3C0OX)

Product Characteristics				
Color				
Shape	Size			
Flavor	Imprint Code			
Contains				

P	Packaging			
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:82123-012- 81	1 in 1 TUBE	06/28/2021	
1	NDC:82123-012- 80	80 mg in 1 TUBE; Type 0: Not a Combination Product		



Natural Sunscreen Cream formulated with non-nano mineral zinc to deliver effective UVA/UVB protection for your kids and the whole family.

Drug Facts

Helps protect the skin against the harmful effects of the sun. Produced with natural ingredients. Suitable for sensitive skin. Sheer Tinted formula. No white cast.

Active Ingredients

Non-nano Zinc Oxide 22%

Uses: ■ Helps prevent sunburn ■ If used with other sun protection measures (see Directions), it helps lower the risk of skin cancer and early skin aging caused by the sun.

Warning; For external use only.

Do not use on open wounds, cuts, or when skin is open.

When using this product keep out of eyes. Rinse with water to remove. Stop use and ask a doctor if rash occurs.

Directions: ■ Apply liberally 15 minutes before sun exposure. ■ Children under 6 months old: Ask a doctor. Reapply: ■ After 80 minutes of swimming or sweating. ■ Immediately after towel drying. ■ At least every 2 hours.

Other Information Protect the product in its container from excessive heat and direct sun.

Inactive Ingredients: Aqua, Coco-Caprylate/Caprate, Dicaprylyl Carbonate, Olus Oil, Glycerin, Aloe Barbadensis Leaf Juice, Butyrospermum Parkii (Shea) Butter, Panthenol (Vitarnin BS) Glyceryl Citrate/Lactate/Linoleate/Oleate, Propolis Extract, Tocopheryl Acetate (Vitarnin E), Argania Spinosa Fruit Extract, Sodium Stearcyl Glutamate, Magnesium Sulfate.

FREE OF x Parabens x Petrolatum x Dyes x Oxybenzone x Octinoxate x Titanium Dioxide

Marketing Information				
Marketing Application Number or Monograph Marketing Start Marketing End Category Citation Date Date				
OTC Monograph Drug	M020	06/28/2021		

S.O.S CREAM INTENSIVE MOISTURIZING SKIN PROTECTANT

non nano zinc oxide cream

Product Information

ROYAL JELLY (UNII: L497I37F0C)

Product Type HUMAN OTC DRUG Item Code (Source) NDC:82123-007

Route of Administration TOPICAL

Active Ingredient/Active Moiety

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Ingredient Name	Basis of Strength	Strength	
ZINC OXIDE (UNII: SOI2LOH54Z) (ZINC OXIDE - UNII:SOI2LOH54Z)	ZINC OXIDE	0.6 mg in 15 mL	

Inactive Ingredients	
Ingredient Name	Strength
OLEA EUROPAEA FLOWER (UNII: 498M34P1VZ)	
WHITE WAX (UNII: 7G1J5DA97F)	
PANTHENOL (UNII: W/9CM0067Z)	
HONEY (UNII: Y9H1V576FH)	
PROPOLIS WAX (UNII: 6Y8XYV2NOF)	
BUTYROSPERMUM PARKII (SHEA) BUTTER UNSAPONIFIABLES (UNII: 0C9AC7D6XU)	
GLYCERIN (UNII: PDC6A3C0OX)	
OLEA EUROPAEA (OLIVE) OIL UNSAPONIFIABLES (UNII: XO45V955LT)	
WATER (UNII: 059QF0KO0R)	
.ALPHATOCOPHEROL ACETATE, D- (UNII: A7E6112E4N)	

Product Characteristics			
Color	Score		
Shape	Size		
Flavor	Imprint Code		
Contains			

P	Packaging				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date	
1	NDC:82123- 007-16	1 in 1 BOX	11/11/2021		
1	NDC:82123- 007-15	15 mL in 1 CONTAINER; Type 0: Not a Combination Product			





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LOT:

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Marketing Information

Marketing Application Number or Monograph Category Citation

ograph Marketing Start Date Marketing End Date

OTC Monograph Drug M016 11/11/2021

ACNE CONTROL SERUM

salicylic acid liquid

Product Information

Product Type HUMAN OTC DRUG Item Code (Source) NDC:82123-008

Route of Administration TOPICAL

Active Ingredient/Active Moiety

Ingredient Name Basis of Strength Strength

SALICYLIC ACID (UNII: O414PZ4LPZ) (SALICYLIC ACID - UNII:O414PZ4LPZ) SALICYLIC ACID 0.15 mg in 30 mL

Inactive Ingredients

Ingredient Name Strength

WATER (UNII: 059QF0KO0R)

APIS MELLIFERA VENOM (UNII: 760130881M)

HYDROXYETHYL CELLULOSE, UNSPECIFIED (UNII: T4V6TWG28D)

BEET (UNII: N487KM8COK)

CAPRYLIC ACID (UNII: OBL58JN025)

ROSA MULTIFLORA FRUIT (UNII: EZ5DSL4T27)	
HYALURONIC ACID (UNII: S270N0TRQY)	
ACTINIDIA POLYGAMA FRUIT (UNII: CJA97047JF)	
XYLITOL (UNII: VCQ006KQ1E)	
TEA TREE OIL (UNII: VIF565UC2G)	
GLYCERIN (UNII: PDC6A3C0OX)	
PROPANEDIOL (UNII: 5965N8W85T)	
PROPOLIS WAX (UNII: 6Y8XYV2NOF)	

Product Characteristics			
Color	Score		
Shape	Size		
Flavor	Imprint Code		
Contains			

P	Packaging			
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:82123- 008-31	1 in 1 BOX	10/11/2023	
1	NDC:82123- 008-30	30 mL in 1 BOTTLE, GLASS; Type 0: Not a Combination Product		



Marketing Information				
Marketing Application Number or Monograph Marketing Start Marketing End Category Citation Date Date				
OTC Monograph Drug	M006	09/29/2023		

Labeler - SBS BILIMSEL BIO COZUMLER SANAYI VE TICARET ANONIM SIRKETI PINAR SUBESI (533166749)

Registrant - SBS BILIMSEL BIO COZUMLER SANAYI VE TICARET ANONIM SIRKETI PINAR SUBESI (533166749)

Establishment

Name	Address	ID/FEI	Business Operations
SBS BILIMSEL BIO COZUMLER SANAYI VE TICARET ANONIM SIRKETI PINAR SUBESI		533166749	manufacture(82123-012, 82123-013, 82123-011, 82123-004, 82123-005, 82123-007, 82123-008, 82123-009, 82123-010)

Revised: 6/2023 SBS BILIMSEL BIO COZUMLER SANAYI VE TICARET ANONIM SIRKETI PINAR SUBESI