CORETEX PROFESSIONAL OUTDOOR SKIN PROTECTION WALLET- coretex professional outdoor skin protection wallet
CoreTex Products Inc

CoreTex Professional Outdoor Skin Protection Wallet
65753-514

Hand Sanitizer
Active Ingredients:

SD Alcohol 40-B 62.5%

Hand Sanitizer
Purpose

Antimicrobial

Hand Sanitizer
Uses:

Hand Sanitizer to help reduce bacteria on the skin.

Hand Sanitizer
Warnings

Flammable. Keep away from heat or flame.

For external use only.

When using this product

do not use in or near the eyes. In case of contact, rinse eyes thoroughly with water.

Stop use and ask a doctor

if irritation or rash appears and lasts.

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

Hand Sanitizer
Directions:

Put enough product in your palm to cover hands and rub hands together briskly until dry.

Hand Sanitizer

Inactive Ingredients:

actinidia chinesis (kiwi) extract, aloe barbadenis leaf juice, aminomethyl propanol, brassica oleracea italica (broccoli) extract, cannabis sativa (hemp) seed extract, carbomer citrullus vulgaris (watermelon) fruit extract, euterpe oleracea fruit extract, fragaria vesca (strawberry) fruit extract, glycerin, helianthus annus (sunfloer) seed oil, hippophae rhamnoides (sea buckthorn) fruit extract, lycium barbarum (goji) extract, myrciaria dubia (camu camu) fruit extract, propanediol, propylene glycol, punica granatum extract, rubus idaeus seed extract, tocopherol acetate (vitamin E acetate), vaccinium angustifolium (blueberry) extract, vaccinium macrocarpon (cranberry) fruit extract, vitis vinifera (grape) seed extra, water

Hand Sanitizer
Other Information:

Store below 110 ºF (43 º), Read the Safety Data Sheet for this Product. You may obtain an SDS from our website: www.coretexproducts.com or Call: 1-877-684-5774

SunX 30
Active Ingredients:

Avobenzone 3.0%

Homosalate 7.5%

Octisalate 5.0%

Octocrylene 5.0%

SunX 30
Purpose

Sunscreen

Sunscreen

Sunscreen

Sunscreen

SunX 30
Uses:

- helps prevent sunburn.
- higher SPF gives more sunburn protection.
- retains SPF after 80 minutes of activity in the water or sweating.
- provides high protection against sunburn.

SunX 30
Warnings

For external use only

SunX 30
Do Not Use
- on damaged or broken skin

When using this product
- keep out of the eyes
- rinse with water to remove

Stop use and ask a doctor if
- rash or irritation develops and lasts

Keep out of reach of children.

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

SunX 30
Directions:
- apply liberally and evenly 15 minutes before sun exposure
- reapply:
- after 80 minutes of swimming or sweating
- after towel drying, swimming, or
- at least every 2 hours
- children under 6 months of age: Ask a doctor

SunX 30
Other Information:
- protect this product from excessive heat or direct sun.

Sun alert: Limiting sun exposure, wearing protective clothing, and using sunscreens may reduce the risk of skin aging, skin cancer, and other harmful effects of the sun.

SunX 30
Inactive Ingredients:

acrylates/C10-30 alkyl acrylate crosspolymer, aloe barbadensis leaf juice, C12-15 alkyl benzoate, carbomer, disodium EDTA, ethylexyiglycerin, hydroxypropyl methylcellulose, phenoxyethanol, polyethylene, polysorbate 20, propylene glycol, sodium hydroxide, sorbitan oleate, theobroma cacao (cocoa) seed butter, tocopheryl acetate, water

SunX 30
Questions?

Call: 1-877-684-5774

SunX 50 Lip Balm
Active Ingredients:

Avobenzone...3.0%

Homosalate...3.0%

Octinoxate...7.5%

Octisalate...5.0%

Petrolatum...40.0%

SunX 50 Lip Balm
Purpose

Sunscreen

Sunscreen

Sunscreen

Sunscreen

Skin Protectant

SunX 50 Lip Balm
Uses:

Helps protect against sunburn and chapped lips.

SunX 50 Lip Balm
Warnings

Sun Alert: Limiting sun exposure, wearing protective clothing, and using sunscreens may reduce the risks of skin aging, skin cancer, and other harmful effects of the sun.

Do not use

Do not use on damaged or broken skin.

Stop use and ask a doctor if
- rash occurs

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

Keep out of reach of children.

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

SunX 50 Lip Balm
Directions:
- Apply liberally before sun exposure and as needed

- Children under 6 months of age: **Ask a doctor before use.**

SunX 50 Lip Balm
Other information:

protect the product in this container from excessive heat and direct sun.

SunX 50 Lip Balm
Inactive Ingredients:

C12-15 alkyl benzoate, caprylic/capric triglyceride, mineral oil, ozokerite, phenyl trimethicone, tocopherol.

Anti-Itch
Active Ingredients:

Camphor 0.1%

Diphenhydramine hydrochloride 2%

Zinc acetate 1%

Anti-Itch
Purpose

Extrenal analgesic

Antihistamine

Skin protectant

Anti-Itch
Uses:
- For the temporary relief of itching and pain associated with minor skin irritations and rashes due to insect bites, poison ivy, poison oak, poison sumac.
- dries the oozing and weeping of poison ivy, poison oak, poison sumac.

Anti-Itch
Warnings

For external use only

Do Not Use
- on chicken pox, blisters or on extensive areas of the skin
- with any drugs containing diphenhydramine while using this product.

When using this product
- keep out of eyes.

Stop use and ask a doctor if

- conditions worsen or if symptoms persist for more than 7 days or clear up and occur again within a few days.

Keep out of reach of children

If swallowed get medical help or contact Poison Control center right away.

Anti-Itch
Directions:

- Adults and children 12 years of age and older, apply to affected area not more than 3 to 4 times daily, or as directed bu a doctor.

Anti-Itch
Other Information:

- protect this product from excessive heat and direct sun.

Anti-Itch
Inactive Ingredients:

citric acid, diazolidinyl urea, glycerin, hydroxypropyl methylcellulose, methylparaben, propylene glycol, propylparaben, SD alcohol 40, sodium citrate, water (aqua).

Anti-Itch
Questions?

Call: 1-877-684-5774

Sting X
Active Ingredients:

Benzocaine 6%

Sting X
Purpose

Topical Analgesic

Sting X
Use:

For temporary pain relief from insect bites and stings

Sting X
Warnings

For external use only

Do not use

- in or near eyes
- over large areas of the body
- over raw or blistered areas

Stop use and ask a doctor if

- conditions worsen or persist for more than 7 days or clear up and occur again within a few days

Keep out of reach of children.

If swallowed get medical help or contact Poison Control center right away.

Sting X
Directions:

Apply to affected area not more than 3 to 4 times daily, for adults and children 2 years of age or older.

Sting X
Inactive Ingredients:

SD alcohol 40, water (aqua), glycerin, allantoin

Sting X
Other Information:

Made in USA for CoreTex Products, Inc.

Bakersfield, CA 93308

www.CoreTexProducts.com (877)684-5774

Outdoor Professional Kit and Refill Kit Contents

1 ea – Sun X SPF 30 Broad Spectrum Lotion Pouch

1 ea – Sun X SPF 30 Broad Spectrum Multi-Pack Pouch w/Towelette
1 ea – Sun X SPF 30 Lip Balm
2 ea – Bug X 30 Insect Repellent Towelette
2 ea – Ivy X Pre-Contact Barrier Towelette
2 ea – Ivy X Post-Contact Cleanser Towelette
2 ea – Burn X Lite Cooling Gel Pouch
1 ea – Anti-Itch Gel Pouch
1 ea – Sting X Pain Relief Pad
1 ea – Hand Sanitizer Gel Pouch

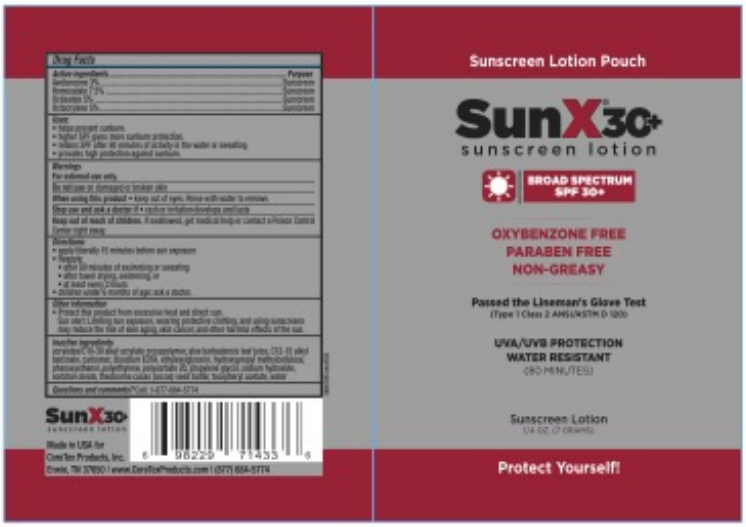
Professional Outdoor Kit Label

Hand Sanitizer
Package Label

Hand Sanitizer Pouch

SunX 30 Label

Sun X Pouch



Black

PMS 187

Pouch Size: 2.5 x 3.5"

SunX 30 Multipack with Towelette Label


Sunscreen Lotion Pouch
SunX 30+
sunscreen lotion
BROAD SPECTRUM SPF 30+
OXYBENZONE FREE
PARABEN FREE
NON-GREASY
Passed the Lineman's Glove Test
(Type 1 Class 2 ANSI/ASTM D 120)
UVA/UVB PROTECTION
WATER RESISTANT
(80 MINUTES)
Sunscreen Lotion
1/4 OZ. (7 GRAMS)
Protect Yourself!
Utility Towelette
SunX
utility towelette
DRY & SUPER ABSORBENT
Multi-Purpose | Convenient | 8X5 Towelette
Single-Use Package | Dry Utility Towelette
Prepare Yourself!

Front


Sunscreen Lotion Pouch
Drug Facts
Active ingredients Purpose
Avobenzone 3% Sunscreen
Homosalate 7.5% Sunscreen
Octisalate 5% Sunscreen
Octocrylene 5% Sunscreen
Uses:
• helps prevent sunburn.
• higher SPF gives more sunburn protection.
• retains SPF after 80 minutes of activity in the water or sweating.
• provides high protection against sunburn.
Warnings
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 use and ask a doctor if • rash or irritation develops and lasts
Keep out of reach of children. If swallowed, get medical help or contact a Poison Control Center right away.
Directions:
• apply liberally 15 minutes before sun exposure
• Reapply:
• after 80 minutes of swimming or sweating
• after towel drying, swimming, or
• at least every 2 hours
• children under 6 months of age: ask a doctor.
Other information
• Protect this product from excessive heat and direct sun.
Sun alert: Limiting sun exposure, wearing protective clothing, and using sunscreens may reduce the risk of skin aging, skin cancer, and other harmful effects of the sun.
Inactive ingredients
acrylates/C10-30 alkyl acrylate crosspolymer, aloe barbadensis leaf juice, C12-15 alkyl benzoate, carbomer, disodium EDTA, ethylexyiglycerin, hydroxypropyl methylcellulose, phenoxyethanol, polyethylene, polysorbate 20, propylene glycol, sodium hydroxide, sorbitan oleate, theobroma cacao (cocoa) seed butter, tocopheryl acetate, water
Questions and comments? Call: 1-877-684-5774
Made in USA | Mfg. for CoreTex Products, Inc. | Erwin, TN 37650 | SKU#71443 rev.1222
Utility Towelette
SunX
utility towelette
SUN X UTILITY TOWELETTE IS A MULTI-PURPOSE, CONVENIENT, DRY TOWELETTE IN A SINGLE DOSE FOIL PACK. IT IS GENTLE ON THE SKIN, FACE AND HANDS. IT IS NOT PRE-MOISTENED AND IS INTENDED TO ABSORB SWEAT DIRT, SUNSCREEN AND/OR OTHER SUBSTANCES.
PLEASE DISCARD RESPONSIBLY.
DRY & SUPER ABSORBENT
CoreTex
6 98229 71443 5
Made in USA | Mfg. for CoreTex Products, Inc.
Erwin, TN 37650 | www.CoreTexProducts.com | (877) 684-5774

Back

SunX 50 Lip Balm Label

Anti-Itch Label

Anti-Itch Gel Pouch

Drug Facts (continued)

Use: For the temporary relief of itching and pain associated with minor cuts, minor skin irritations and rashes due to insect bites, poison ivy, poison oak, and poison sumac.

Warnings: **For external use only.**

Do not use: • on chicken pox, blisters or on extensive areas of the skin. • With any other drugs containing **diphenhydramine** while using this products. • In the eyes. • more than directed.

Directions: Adults and children 2 years of age and older, apply to affected area not more than 3 to 4 times daily, or as directed by a doctor. For use on children under 2 years of age, consult a doctor.

Made in USA for CoreTex Products, Inc., Erwin, TN 37650 (877) 684-5774

SKU#26638 rev.1222

Anti-Itch

Dual Action Gel

Net Weight: 0.034 oz. (1 gram)

Drug Facts

Active Ingredients	***Purpose***
Camphor 0.1%	External analgestic
Diphenhydramine hydrochloride 2%	Antihistamine
Zinc acetate 1%	Skin protectant

Black

PMS 187

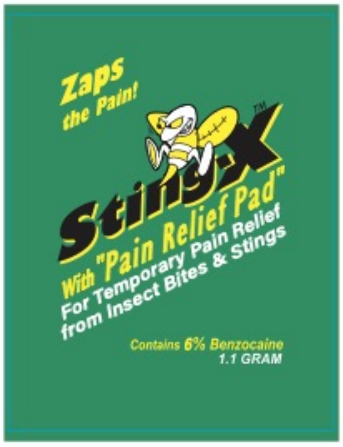
Sting X Label

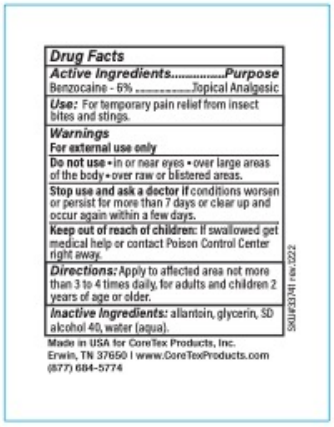
Sting X Pouch

PMS 339 C

PMS 3945 C

PMS Hexachrome Black





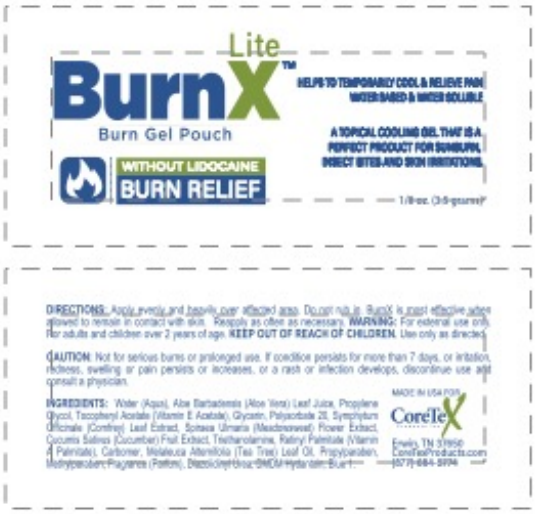
H2.5" x W2"

Burn X Label

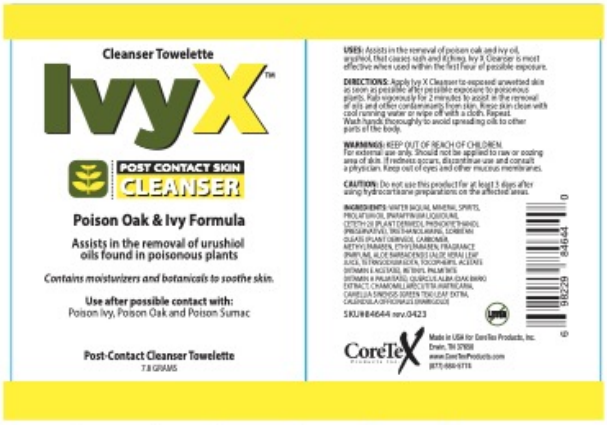
65753

Burn X Lite Pouch

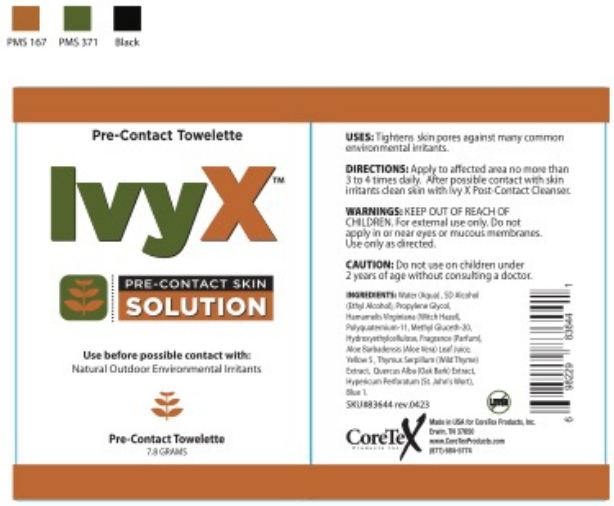
PMS 377 C PMS 286 C



IVY X Cleanser Label



IVY X Pre Label



CORETEX PROFESSIONAL OUTDOOR SKIN PROTECTION WALLET

coretex professional outdoor skin protection wallet kit

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:65753-514

Packaging				
#	**Item Code**	**Package Description**	**Marketing Start Date**	**Marketing End Date**
1	NDC:65753-514-36	1 in 1 KIT; Type 0: Not a Combination Product	02/24/2023	

Quantity of Parts		
Part #	**Package Quantity**	**Total Product Quantity**
Part 1	1 PACKET	44 mL
Part 2	1 POUCH	7 g
Part 3	14809258 APPLICATOR	65753105 mL
Part 4	1 POUCH	1 mL
Part 5	1 POUCH	1.1 mL
Part 6	1 POUCH	3.5 mL

Part 1 of 6

CORETEX SUN X SPF 30 NEW

avobenzone, homosalate, octisalate, octocrylene lotion

Product Information

Item Code (Source)	NDC:65753-110
Route of Administration	TOPICAL

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
HOMOSALATE (UNII: V06SV4M95S) (HOMOSALATE - UNII:V06SV4M95S)	HOMOSALATE	7.5 g in 100 mL
OCTISALATE (UNII: 4X49Y0596W) (OCTISALATE - UNII:4X49Y0596W)	OCTISALATE	5 g in 100 mL
AVOBENZONE (UNII: G63QQF2NOX) (AVOBENZONE - UNII:G63QQF2NOX)	AVOBENZONE	3 g in 100 mL
OCTOCRYLENE (UNII: 5A68WGF6WM) (OCTOCRYLENE - UNII:5A68WGF6WM)	OCTOCRYLENE	5 g in 100 mL

Inactive Ingredients

Ingredient Name	Strength
MEDIUM DENSITY POLYETHYLENE (UNII: 3W404QE89S)	
DISODIUM EDTA-COPPER (UNII: 6V475AX06U)	
THEOBROMA CACAO WHOLE (UNII: EB048G1S9J)	
ETHYLHEXYLGLYCERIN (UNII: 147D247K3P)	
POLYSORBATE 20 (UNII: 7T1F30V5YH)	
METHYLCELLULOSE, UNSPECIFIED (UNII: Z944H5SN0H)	
CARBOMER HOMOPOLYMER, UNSPECIFIED TYPE (UNII: 0A5MM307FC)	
SODIUM HYDROXIDE (UNII: 55X04QC32I)	
ACRYLATES CROSSPOLYMER-6 (UNII: 4GXD0Q3OS3)	
ALKYL (C12-15) BENZOATE (UNII: A9EJ3J61HQ)	
.ALPHA.-TOCOPHEROL ACETATE (UNII: 9E8X80D2L0)	
ALOE VERA LEAF (UNII: ZY81Z83H0X)	
WATER (UNII: 059QF0KO0R)	
PHENOXYETHANOL (UNII: HIE492ZZ3T)	
PEG-6 SORBITAN OLEATE (UNII: 58O7V09UCI)	
PROPYLENE GLYCOL PROPYL ETHER (UNII: 92KA3PYX0S)	

Product Characteristics

Color	white (Thick White Lotion)	**Score**	
Shape		**Size**	
Flavor		**Imprint Code**	
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:65753-110-37	44 mL in 1 PACKET; Type 0: Not a Combination Product		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M020	10/01/2023	

Part 2 of 6

CORETEX SUN X SPF 30 MULTIPACK NEW

avobenzone, homosalate, octisalate, octocrylene lotion

Product Information

Item Code (Source)	NDC:65753-109
Route of Administration	TOPICAL

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
AVOBENZONE (UNII: G63QQF2NOX) (AVOBENZONE - UNII:G63QQF2NOX)	AVOBENZONE	3 g in 100 g
OCTOCRYLENE (UNII: 5A68WGF6WM) (OCTOCRYLENE - UNII:5A68WGF6WM)	OCTOCRYLENE	5 g in 100 g
OCTISALATE (UNII: 4X49Y0596W) (OCTISALATE - UNII:4X49Y0596W)	OCTISALATE	5 g in 100 g
HOMOSALATE (UNII: V06SV4M95S) (HOMOSALATE - UNII:V06SV4M95S)	HOMOSALATE	7.5 g in 100 g

Inactive Ingredients

Ingredient Name	Strength
DISODIUM EDTA-COPPER (UNII: 6V475AX06U)	
SODIUM HYDROXIDE (UNII: 55X04QC32I)	
ETHYLHEXYLGLYCERIN (UNII: 147D247K3P)	
POLYSORBATE 20 (UNII: 7T1F30V5YH)	
METHYLCELLULOSE, UNSPECIFIED (UNII: Z944H5SN0H)	
PROPYLENE GLYCOL PROPYL ETHER (UNII: 92KA3PYX0S)	
CARBOMER HOMOPOLYMER, UNSPECIFIED TYPE (UNII: 0A5MM307FC)	
ALOE VERA LEAF (UNII: ZY81Z83H0X)	
.ALPHA.-TOCOPHEROL ACETATE (UNII: 9E8X80D2L0)	
WATER (UNII: 059QF0KO0R)	
PHENOXYETHANOL (UNII: HIE492ZZ3T)	
PEG-6 SORBITAN OLEATE (UNII: 58O7V09UCI)	
ALKYL (C12-15) BENZOATE (UNII: A9EJ3J61HQ)	
MEDIUM DENSITY POLYETHYLENE (UNII: 3W404QE89S)	
ACRYLATES CROSSPOLYMER-6 (UNII: 4GXD0Q3OS3)	
THEOBROMA CACAO WHOLE (UNII: EB048G1S9J)	

Packaging

			Marketing Start	Marketing End

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:65753-109-39	7 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
OTC Monograph Drug	M020	10/01/2023	

Part 3 of 6

SUN X SPF 50 BROAD SPECTRUM SUNSCREEN LIP BALM

avobenzone, homosalate, octinoxate, octisalate, petrolatum lipstick

Product Information

Item Code (Source)	NDC:65753-108
Route of Administration	TOPICAL

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
OCTINOXATE (UNII: 4Y5P7MUD51) (OCTINOXATE - UNII:4Y5P7MUD51)	OCTINOXATE	7.5 g in 100 mL
HOMOSALATE (UNII: V06SV4M95S) (HOMOSALATE - UNII:V06SV4M95S)	HOMOSALATE	3 g in 100 mL
OCTISALATE (UNII: 4X49Y0596W) (OCTISALATE - UNII:4X49Y0596W)	OCTISALATE	5 g in 100 mL
AVOBENZONE (UNII: G63QQF2NOX) (AVOBENZONE - UNII:G63QQF2NOX)	AVOBENZONE	3 g in 100 mL
PETROLATUM (UNII: 4T6H12BN9U) (PETROLATUM - UNII:4T6H12BN9U)	PETROLATUM	40 g in 100 mL

Inactive Ingredients

Ingredient Name	Strength
PHENYL TRIMETHICONE (UNII: DR0K5NOJ4R)	
COCONUT OIL (UNII: Q9L0O73W7L)	
ALKYL (C12-15) BENZOATE (UNII: A9EJ3J61HQ)	
MINERAL OIL (UNII: T5L8T28FGP)	
WHITE WAX (UNII: 7G1J5DA97F)	
TOCOPHEROL (UNII: R0ZB2556P8)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:65753-108-39	4.44 mL in 1 APPLICATOR; Type 0: Not a Combination Product		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M020	02/24/2023	

Part 4 of 6

CORETEX ANTI-ITCH GEL

camphor, diphenhydramine, zinc acetate gel

Product Information

Item Code (Source)	NDC:65753-400
Route of Administration	TOPICAL

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
CAMPHOR (SYNTHETIC) (UNII: 5TJD82A1ET) (CAMPHOR (SYNTHETIC) - UNII:5TJD82A1ET)	CAMPHOR (SYNTHETIC)	0.1 g in 100 mL
ZINC ACETATE (UNII: FM5526K07A) (ZINC CATION - UNII:13S1S8SF37)	ZINC ACETATE	1 g in 100 mL
DIPHENHYDRAMINE HYDROCHLORIDE (UNII: TC2D6JAD40) (DIPHENHYDRAMINE - UNII:8GTS82S83M)	DIPHENHYDRAMINE HYDROCHLORIDE	2 g in 100 mL

Inactive Ingredients

Ingredient Name	Strength
SODIUM CITRATE (UNII: 1Q73Q2JULR)	
METHYLPARABEN (UNII: A2I8C7HI9T)	
ANHYDROUS CITRIC ACID (UNII: XF417D3PSL)	
GLYCERIN (UNII: PDC6A3C0OX)	
DIAZOLIDINYL UREA (UNII: H5RIZ3MPW4)	
METHYLCELLULOSE (1500 CPS) (UNII: P0NTE48364)	
PROPYLPARABEN (UNII: Z8IX2SC1OH)	
ALCOHOL (UNII: 3K9958V90M)	
WATER (UNII: 059QF0KO0R)	
PROPYLENE GLYCOL 1-ALLYL ETHER (UNII: QRB8092KPK)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:65753-400-39	1 mL in 1 POUCH; Type 0: Not a Combination Product		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M017	11/01/2023	

Part 5 of 6

STINGX

benzocaine swab

Product Information

Item Code (Source)	NDC:65753-350
Route of Administration	TOPICAL

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
BENZOCAINE (UNII: U3RSY48JW5) (BENZOCAINE - UNII:U3RSY48JW5)	BENZOCAINE	6 g in 100 mL

Inactive Ingredients

Ingredient Name	Strength
WATER (UNII: 059QF0KO0R)	
ALLANTOIN (UNII: 344S277G0Z)	
GLYCERIN (UNII: PDC6A3C0OX)	
ALCOHOL (UNII: 3K9958V90M)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1		1.1 mL in 1 POUCH; Type 0: Not a Combination Product		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M017	11/25/2019	

Part 6 of 6

CORETEX ANTIBACTERIAL HAND SANITIZER

sd alcohol liquid

Product Information	
Item Code (Source)	NDC:65753-200
Route of Administration	TOPICAL

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
ALCOHOL (UNII: 3K9958V90M) (ALCOHOL - UNII:3K9958V90M)	ALCOHOL	62.5 mL in 100 mL

Inactive Ingredients	
Ingredient Name	**Strength**
1,3-PROPANEDIOL BIS(4-AMINOBENZOATE) (UNII: 8860R9ORQR)	
GLYCERIN (UNII: PDC6A3C0OX)	
ALPHA-TOCOPHEROL ACETATE (UNII: 9E8X80D2L0)	
WATER (UNII: 059QF0KO0R)	
ALOE VERA LEAF (UNII: ZY81Z83H0X)	
AMINOMETHYL PROPANEDIOL (UNII: CZ7BU4QZJZ)	
CARBOMER HOMOPOLYMER, UNSPECIFIED TYPE (UNII: 0A5MM307FC)	

Packaging				
#	**Item Code**	**Package Description**	**Marketing Start Date**	**Marketing End Date**
1		3.5 mL in 1 POUCH; Type 0: Not a Combination Product		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M004	11/25/2019	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M020	02/24/2023	

Labeler - CoreTex Products Inc (061944620)

Establishment

Name	Address	ID/FEI	Business Operations
Cosmetic Enterprises		017701475	manufacture(65753-400, 65753-350)

Establishment

Name	Address	ID/FEI	Business Operations
CoreTex Products Inc		061944620	pack(65753-514, 65753-200, 65753-108, 65753-110, 65753-109, 65753-350, 65753-400)

Establishment

Name	Address	ID/FEI	Business Operations
Pure Source		080354456	manufacture(65753-109, 65753-110)

Establishment

Name	Address	ID/FEI	Business Operations
Raining Rose		083819404	manufacture(65753-108)

Establishment

Name	Address	ID/FEI	Business Operations
HealthSpecialty		794053863	manufacture(65753-200)