

DR. DREAM ADVANCED FACIAL TREATMENT TONER - witch hazel liquid

Dr. Dream Inc

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.

Drug Facts

witch hazel

bis-PEG-18 methyl ether dimethyl silane, butylene glycol, panthenol, sodium PCA, caprylyl glycol, torrefied nutmeg extract, cornus kousa extract, rosa multiflora fruit extract, chloranthus glaber extract, musa basjoo extract, musa basjoo extract, rh-oligopeptide-1, acetyl hexapeptide-8, palmitoyl oligopeptide, palmitoyl tetrapeptide-7, polysorbate 80, glycerin, caprylhydroxamic acid, adenosine, ammonium acryloyldimethyltaurate, hydroxyethylcellulose, pentaerythritol, cellulose gum, sodium chondroitin sulfate, lavandula angustifolia oil, cymbopogon schoenanthus oil, abies sibirica oil, citrus aurantium dulcis peel oil, amyris balsamifera bark oil, salvia sclarea oil, lecithin, sodium phosphate, sodium hyaluronate, rh-polypeptide-60, rh-polypeptide-9, rh-polypeptide-11, rh-polypeptide-2, rh-polypeptide-1, rh-polypeptide-10, copper tripeptide-1, palmitoyl pentapeptide-4, glycine, glutamine, lysine, leucine, methionine, valine, serine, cysteine, cystine, asparagine, alanine, arginine, ornithine, isoleucine, tyrosine, threonine, tryptophan, phenylalanine, proline, histidine

anti wrinkle

keep out of reach of the children

after cleaning up your face with clean water apply proper amount to your skin and massage a few minutes

When using this product

- keep out of eyes, ears, and mouth. If contact occurs, rinse with plenty of cold water right away and contact a physician. If swallowing, drink plenty of water and contact a physician

for external use only



DR. DREAM ADVANCED FACIAL TREATMENT TONER

witch hazel liquid

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:530 77-100 1
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
WITCH HAZEL (UNII: 10 114J0 U34) (WITCH HAZEL - UNII: 10 114J0 U34)	WITCH HAZEL	0.1 mg in 100 mL

Inactive Ingredients

Ingredient Name	Strength
BIS-PEG-18 METHYL ETHER DIMETHYL SILANE (UNII: OEB4R3WW9C)	
BUTYLENE GLYCOL (UNII: 3XUS85K0RA)	
PANTHENOL (UNII: WV9CM0O67Z)	
SODIUM PYRROLIDONE CARBOXYLATE (UNII: 469OTG57A2)	
CAPRYLYL GLYCOL (UNII: 00YIU5438U)	
TORREYA NUCIFERA WHOLE (UNII: 91Z68135ZV)	
ROSA DAMASCENA FLOWER OIL (UNII: 18920M3T13)	
MUSA BASJOO WHOLE (UNII: V3AYC3JL9Y)	
ACETYL HEXAPEPTIDE-8 (UNII: L4EL31FWIL)	
PALMITOYL OLIGOPEPTIDE (UNII: HO4ZT5S86C)	
PALMITOYL TETRAPEPTIDE-7 (UNII: Q41S464P1R)	
POLYSORBATE 80 (UNII: 6OZP39ZG8H)	
GLYCERIN (UNII: PDC6A3C0OX)	
CAPRYLHYDROXAMIC ACID (UNII: UPY805K99W)	
ADENOSINE (UNII: K72T3FS567)	
AMMONIUM ACRYLOYLDIMETHYLTAURATE/VP COPOLYMER (UNII: W59H9296ZG)	
NEOPENTYL GLYCOL (UNII: Q180HXD6S5)	
CARBOXYMETHYLCELLULOSE SODIUM (UNII: K679OBS311)	
SODIUM CHONDROITIN SULFATE (PORCINE; 5500 MW) (UNII: H5BJH23Z9A)	
LAVANDULA ANGUSTIFOLIA FLOWERING TOP (UNII: 9YT4B71U8P)	
CYMBOPOGON SCHOENANTHUS OIL (UNII: XE7K568ILO)	
ABIES SIBIRICA LEAF OIL (UNII: XRY0V4VZKZ)	
ORANGE (UNII: 5EVU04N5QU)	
AMYRIS BALSAMIFERA OIL (UNII: I1BJ961J2E)	
CLARY SAGE OIL (UNII: 87L0D4U3M0)	
LECITHIN, SOYBEAN (UNII: 1DI56QDM62)	
SODIUM PHOSPHATE (UNII: SE337SVY37)	
HYALURONATE SODIUM (UNII: YSE9PPT4TH)	
THIOREDOXIN (UNII: XJZ418133Z)	
PREZATIDE COPPER (UNII: 6BJQ43T119)	
PALMITOYL PENTAPEPTIDE-4 (UNII: KK181SM5JG)	
GLYCINE (UNII: TE7660XO1C)	
GLUTAMINE (UNII: 0RH81L854J)	
LYSINE (UNII: K3Z4F929H6)	
LEUCINE (UNII: GMW67QNF9C)	
METHIONINE (UNII: AE28F7PNPL)	
VALINE (UNII: HG18B9YRS7)	
SERINE (UNII: 452VLY9402)	
CYSTEINE (UNII: K848JZ4886)	
CYSTINE (UNII: 48TCX9A1VT)	
ASPARAGINE (UNII: 5Z33R5TKO7)	
ALANINE (UNII: OF5P57N2ZX)	
ARGININE (UNII: 94ZLA3W45F)	
ORNITHINE (UNII: E524N2IXA3)	
ISOLEUCINE (UNII: 04Y7590D77)	
TYROSINE (UNII: 42HK56048U)	
THREONINE (UNII: 2ZD004190S)	

TRYPTOPHAN (UNII: 8DUH1N11BX)	
PHENYLALANINE (UNII: 47E5O17Y3R)	
PROLINE (UNII: 9DLQ4CIU6V)	
HISTIDINE (UNII: 4QD397987E)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:53077-1001-1	100 mL in 1 BOTTLE, GLASS		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph final	part347	08/24/2012	

Labeler - Dr. Dream Inc (557821161)

Registrant - Dr. Dream Inc (557821161)

Establishment

Name	Address	ID/FEI	Business Operations
New & New Co., Ltd		557821160	manufacture(53077-1001)

Revised: 8/2012

Dr. Dream Inc