KAYDIA PATCH- kaydia patch patch
STRONG CURRENT ENTERPRISES LIMITED

Disclaimer: This drug has not been found by FDA to be safe and effective, and this labeling has not been approved by FDA. For further information about unapproved drugs, click here.

Kaydia Patch

Active Ingredients

CBD 0.15%, Camphor 3%, Mentholum 1%, Capsicum Annuum Extract 0.12%

Purpose

Uses

Warning

For external use only, do not ingest
Do not use on wounds,rash or damaged skin
Do not use if you have skin allergy
Not suitable for expectant mothers

when using this product

Use only as directed
Apply on clean and dry skin only
Avoid area near eye, mouth or other mucous opening
Do not use with combination with a heating device
Do not apply more than 1 large patch at a time

Stop using this product and consult a doctor if:Condition worsens
Allergic reaction like rash, itching and other skin irritation developed
Keep out of reach of children
If ingested, seek medical/poison control attention immediately

Directions

1.Remove backing film
2.Apply patch on dry and clean skin
3.Remove and discard patch after 8-10 hours

Other information

Store in dry, cool place
Avoid direct sunlight exposure
Seal bag after opening to keep product fresh

Inactive Ingredients

Water 56.53%, glycerin 25%, ammonium polyacrylate 9%, alcohol 3%, sodium acrylate 1.5%, 2,4-Imidazolidinedione 0.2%, disodium edta 0.1%, tartaric acid 0.25%, aluminum glycinate 0.15%

Package Label - Principal Display Panel

KAYDIA PATCH

kaydia patch patch

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:75140-002
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
CAMPHOR (NATURAL) (UNII: N20HL7Q941) (CAMPHOR (NATURAL) - UNII:N20HL7Q941)	CAMPHOR (NATURAL)	3 g in 100 g
LEVOMENTHOL (UNII: BZ1R15MTK7) (LEVOMENTHOL - UNII:BZ1R15MTK7)	LEVOMENTHOL	1 g in 100 g
CANNABIDIOL (UNII: 19GBJ60SN5) (CANNABIDIOL - UNII:19GBJ60SN5)	CANNABIDIOL	0.15 g in 100 g

Inactive Ingredients

Ingredient Name	Strength
WATER (UNII: 059QF0KO0R)	
TARTARIC ACID (UNII: W4888I119H)	
DIHYDROXYALUMINUM AMINOACETATE (UNII: DO250MG0W6)	
ALCOHOL (UNII: 3K9958V90M)	
HYDANTOIN (UNII: I6208298TA)	
GLYCERIN (UNII: PDC6A3C0OX)	
SODIUM ACRYLATE (UNII: 7C98FKB43H)	
DISODIUM EDTA-COPPER CU-64 (UNII: 4J875U5U11)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:75140-002-01	166 g in 1 BOX; Type 0: Not a Combination Product	05/02/2020	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
unapproved drug other		05/02/2020	

Labeler - STRONG CURRENT ENTERPRISES LIMITED (685811978)

Registrant - STRONG CURRENT ENTERPRISES LIMITED (685811978)

Establishment

Name	Address	ID/FEI	Business Operations
STRONG CURRENT ENTERPRISES LIMITED		685811978	manufacture(75140-002)

Revised: 5/2020 STRONG CURRENT ENTERPRISES LIMITED