CANNABIS SATIVA- cannabis sativa liquid Remedy Makers

Disclaimer: This homeopathic product has not been evaluated by the Food and Drug Administration for safety or efficacy. FDA is not aware of scientific evidence to support homeopathy as effective.

CANNABIS SATIVA 1X

(Cannabidiol) 100mg per mL Delta 9 THC < 0.3%

LESSENS PAIN, IMPROVES APPETITE, AND CAUSES SLEEP

WARNING:

"The FDA has not determined that this product is safe, effective and not misbranded for its intended use."

To be used according to standard homeopathic indications for self-limiting conditions such as those indicated above or as directed by a physician.

Keep this and all medication out of reach of children.

Warning:

Use only if cap and seal are unbroken. If symptoms persist for more than 3 days or worsen, discontinue (STOP) use and consult your physician. As with any drug, if you are pregnant or nursing (breast-feeding) a baby, seek the advise of a health professional before using this product. Store tightly closed in a cool area.

Directions:

Take 1mL by mouth or as directed by a physician.

Other information:

Contains approx. 30 doses.

Inactive Ingredients:

MCT Oil, Alcohol, and Strawberry Flavoring. Free from yeast, wheat, corn, and soy.

Questions or comments

(877) REM4YOU Fax (909) 594-4205 Pomona, CA 91768. USA







HOMEOPATHIC MEDICINE

CANNABIS SATIVA cannabis sativa liquid					
Product Information					
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:10191-6374		
Route of Administration	ORAL				
Active Ingredient/Active Moiety					

		Ingredient Name	Basis of Streng	gth Strengtl
	NNABIS SATIV NII:19GBJ60SN5	A FLOWERING TOP (UNII: 8X454SZ22D) (CANNABIDI)	OL CANNABIS SATIVA FLOWERING TOP	1 [hp_X] in 30 mL
In	active Ingr	edients		
	Strength			
AL				
ME				
Pa	ackaging			
#	ltem Code	Package Description	Marketing Start Date	Marketing En Date
	NDC:10191- 6374-1	30 mL in 1 BOTTLE, DROPPER; Type 0: Not a Combination Product	03/22/2023	
м	arkotina	Information		
	U			
	Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
	approved meopathic		03/22/2023	
	approved	Citation		Date

Labeler - Remedy Makers (018543582)

Registrant - Remedy Makers (018543582)

Revised: 3/2023

Remedy Makers