MOSQUITO BUG OFF- diethyltoluamide liquid Lydia Co., Ltd.

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.

Drug Facts

DIETHYLTOLUAMIDE

ET5HANOL, PURIFIED WATER, DIPROPYLENE GLYCOL, FRAGRANCE

mosquito repellents that prevent access to mosquitoes.

keep out of reach of the children

You can spray it on your skin or clothes.

do not use for the following people: infants under six months

do not use excessive amounts or for long periods of time beyond the required

do not use it in an enclosed area

for external use only

MOSQUITO OFF

IT'S NOT STICKY, IT'S CLEANER, IT'S MORE FREE TO DO OUTDOOR ACTIVITIES,

MOSQUITOES, MITES PREVENTION / SPRAY TYPE / MEDICINAL OUTLETS / PORTABLE

MOSQUITO-KIPERS, WHICH ARE MADE OF COLORLESS OR BROWN LIQUID, CAN BE USED TO MAKE MOSQUITOES, MOLES, ETC, AVOID MOSQUITO BITES WITH AN ANTI-ACCESS REPELLENTERY THAT PREVENTS ACCESS.

SPECIAL ADVANTAGES OF MOGI-KEEPER FLUID

- MOSQUITO CONTROL
 MOSQUITOES IN THE SUMMER, AND OTHER HAIR MITES.
- A CHILD'S RELIEF

 SAFE FOR HUMAN BODY WITH NATURAL INGREDIENTS
 AND MEDICATION CERTIFIED BY THE KOREA FOOD AND
 DRUG ADMINISTRATION
- THE SCENT OF NATURE EVEN IF YOU SPRAY IT ON YOUR SKIN OR CLOTHES, IT'S OKAY.
- 4 SPRAY TYPE

 EASY TO USE WITH SPRAY TYPE

 SPRINKLE ON NECK, ARMS, LEGS, ETC,

 EXCEPT EYES AND MOUTH,
- 5 SAFE MEDICAL SUPPLIES
 SAFE MEDICINES OUTLETS APPROVED BY
 THE FOOD AND DRUG ADMINISTRATION



MOSQUITO BUG OFF

diethyltoluamide liquid

Product	Intorm	ation

Product Type HUMAN OTC DRUG Item Code (Source) NDC:72988-0005

Route of Administration TOPICAL

Active Ingredient/Active Moiety

Ingredient Name	Strength	Strength
DIETHYLTOLUAMIDE (UNII: FB0C1XZV4Y) (DIETHYLTOLUAMIDE - UNII:FB0C1XZV4Y)	DIETHYLTOLUAMIDE	10 g in 100 mL

Inactive Ingredients		
Ingredient Name	Strength	
WATER (UNII: 059QF0KO0R)		
DIPROPYLENE GLYCOL (UNII: E107L85C40)		

l	Packaging				
	#	Item Code	Package Description	Marketing Start Date	Marketing End Date
	1	NDC:72988- 0005-1	60 mL in 1 BOTTLE; Type 0: Not a Combination Product	03/23/2019	

Marketing Information			
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
unapproved drug other		03/23/2019	

Labeler - Lydia Co., Ltd. (695735569)

Registrant - Lydia Co., Ltd. (695735569)

Establishment				
Name	Address	ID/FEI	Business Operations	
Aloe Vera Korea Co.,Ltd.		687773350	manufacture(72988-0005)	

Establishment				
Name	Address	ID/FEI	Business Operations	
Lydia Co., Ltd.		695735569	label(72988-0005)	

Revised: 6/2021 Lydia Co., Ltd.