AQUACOOL MULTI 500- menthol gel Pharmanuco

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

MENTHOL

Camphor, Carbomer, Eucalyptus Globulus Leaf Oil, Witch Hazel Extract, Arnica Montana Flower Extract, Aloe Barbadensis Leaf Extract, Acetyl Glucosamine, etc.

External Analgesic: Multi Action Recovery for Warm Up & Cool Down

keep out of reach of the children

- 1. Apply gel on to affected areas.
- 2. Massage until completely absorbed into skin.
- 3. Repeat as many as needed.
- * It is recommended to control the amount and times of use depending on your symptom.

for external use only

Avoid contact with eyes. Stop if irritation occurs. Avoid bandaging tightly. Keep out of reach of children. Ask your doctor before use if pregnant or breast feeding





AQUACOOL MULTI 500

menthol gel

Product Information

Product TypeHUMAN OTC DRUGItem Code (Source)NDC:70759-0012

Route of Administration TOPICAL

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
MENTHOL (UNII) I 7T10 FID3 A) (MENTHOL - UNII) I 7T10 FID3 A)	MENTHOI	3 g in 100 mI

Inactive Ingredients

	Ingredient Name	Strength
- 1		

EUCALYPTUS GLOBULUS LEAF (UNII: S546YLW6E6)

CAMPHO R, (-)- (UNII: 213N3S8275)

ARNICA MONTANA FLOWER (UNII: OZ0E5Y15PZ)	
WITCH HAZEL (UNII: 10 114J0 U34)	
ALOE VERA LEAF (UNII: ZY8 1Z8 3H0 X)	
N-ACETYLGLUCO SAMINE (UNII: V956696549)	

Packaging				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:70759-0012- 1	500 mL in 1 BOTTLE, PUMP; Type 0: Not a Combination Product	12/01/2017	

Marketing Information				
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date	
OTC monograph not final	part348	12/0 1/20 17		

Labeler - Pharmanuco (687825097)

Registrant - Pharmanuco (687825097)

Establishment			
Name	Address	ID/FEI	Business Operations
Pharmanuco		687825097	manufacture(70759-0012)

Revised: 12/2017 Pharmanuco