CIRLUTONE- camphor ointment Herbal Health LLC

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.

Camphor 0.6%

Camphor 0.6%.....Topical Analgesic

For the temporary relief of pain associated with:

- sunburn
- minor cuts
- scrapes

For External Use only.

Flammable: lkeep away from fire or flame

When using this product

- avoid contact with eyes or mucous membrane
- use only as directed

OStop use and ask a doctor if

- condition worsens
- symptoms persist for more than 7 days
- symptoms clear up and occur again within a few days

If pregnant or breastfeeding, ask a healthcare professional before use.

IKeep out of reach of childrenI. If swallowed, get medical help or contact a Poison Control Center immediately.

- **Adults and children 2 years of age and older**: apply to affected area not more than 3 to 4 times daily.
- **Children under 2 yrs of ageC**: consult a doctor.
- Store in a cool & dry place
- Do not use if tamper evident seal is broken or missing

Angelica anemala, Emulsifying Wax NF, Fragrance, Frankincense Oil, Isopropyl alcohol, Ligusticum wallichii, Magnolia liliflora, Methylparaben, Mineral Oil, Petrolatum, Propylparaben, Purified Water, Sophora, Syzgium armaticum

Call 310-888-8878 Monday through Friday, 9am-5pm CST.

Herbal Health L.L.C. Made in the U.S.A. www.cirlutone.com

Exp. Date:

Lot No.:

Manufactured by:

Neutraderm, Inc.

20660 Nordhoff St.

Chatsworth, CA. 91311

info@cirlutone.com

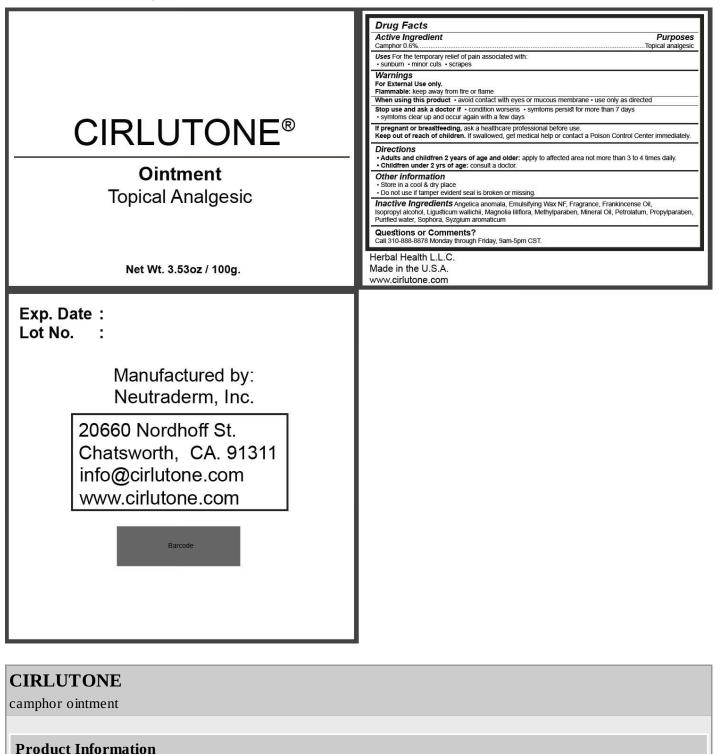
www.cirlutone.com

CIRLUTONE

Ointment

Topical Analgesic

INet Wt. 3.53oz/100g.



Product T ype

HUMAN OTC DRUG

NDC:71917-001

Route	of Administration	
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Active Ingredien	t/Active Moiety					
	Ingredient Name	Basis of St	rength Strength			
CAMPHOR (SYNTHE UNII:5TJD82A1ET)	TIC) (UNII: 5TJD82A1ET) (CAMPHOR (SYNTHETIC) -	CAMPHOR (SYNTHETIC)	0.6 g in 100 g			
Inactive Ingredie	ents					
Ingredient Name						
ANGELICA ANO MAL	A WHOLE (UNII: T1F4GB8MNL)					
WHITE WAX (UNII: 70	G1J5DA97F)					
FRANKINCENSE O IL	(UNII: 67ZYA5T02K)					
ISOPROPYL ALCOHOL (UNII: ND2M416302)						
LIGUSTICUM WALL	ICHII WHOLE (UNII: 8 Y1N3NX2DW)					
MAGNOLIA LILIIFL	ORA WHOLE (UNII: GDO2997SZZ)					
METHYLPARABEN (JNII: A218C7H19T)					
MINERAL OIL (UNII: T5L8T28FGP)						
PETROLATUM (UNII	4T6H12BN9U)					
PROPYLPARABEN (U	JNII: Z8 IX2SC1OH)					
WATER (UNII: 059QF	0KO0R)					
SOPHORA FLAVESC	ENS WHOLE (UNII: X8KX602M5L)					
SYZYGIUM AROMATICUM WHOLE (UNII: EY9 MMA0 P6 Y)						
Packaging						
# Item Code	Package Description	Marketing Start Date	Marketing End Date			
1 NDC:71917-001-35	100 g in 1 BOTTLE; Type 0: Not a Combination Product	12/06/2017				
Marketing Inf	ormation					
	wy Application Number or Menograph Citation	Marketing Start Date	Marketing End Date			
Marketing Catego	ry Application Number or Monograph Citation	Marketing Start Date	Marketing End Date			

Labeler - Herbal Health LLC (080764759)

Establishment							
Name	Address	ID/FEI	Business Operations				
Neutraderm, Inc.		146224444	manufacture(71917-001)				

Revised: 12/2017

Herbal Health LLC