OHEAL SHINGLES PAIN RELIEF CREAM- shingles pain relief cream cream
Dr.luke Healthcare 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.

83176-004
OHEAL SHINGLES PAIN RELIEF CREAM

Active Ingredient

MENTHOL, UNSPECIFIED FORM 3%

Purpose

Topical analgesic

Use

For the treatment of symptomsof shingles, hives, and rashes.

Warnings

For external use only.Avoid contact with eyes, flush with water immediately if it getsinto your eyes.Keep out of reach of children.

Do not use

On broken skin
On eyes area

WHEN USING SECTION

For external use only.Avoid contact with eyes, flush with water immediately if it getsinto your eyes.Keep out of reach of children.

STOP USE section

If you are allergic to this product.

Directions

Apply to the affected areas 2-3 times daily.
Children should use it under the supervision of adults.

Inactive ingredients

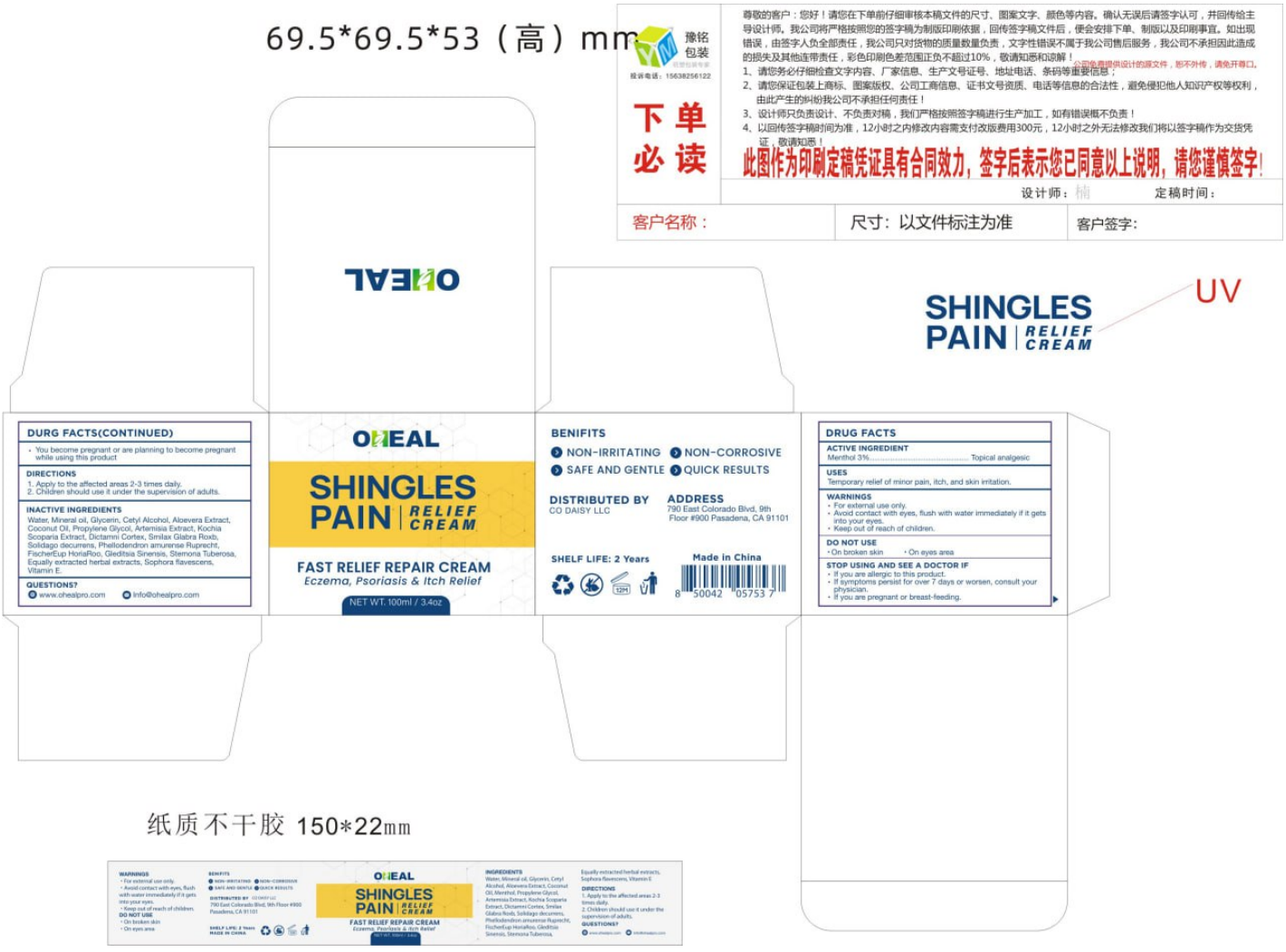
Water, Mineral oil, Glycerin, Cetyl Alcohol, Aloevera Extract,Coconut Oil, Propylene Glycol, Artemisia Extract, K chiaSoparia seeds , Dictamni Cortex, Smilax Glabra Roxb.Solidago decurrens, Phellodendron amurense Ruprecht.Euphorbia Fischeriana Root,Gleditsia Sinensis, StemonaTuberosa,Equally extracted herbalextracts,Sophora flavescensVitamin E.

QUESTIONS

www.ohealpro.com
Info@ohealpro.com

Keep out of reach of children.

83176-004-01



OHEAL SHINGLES PAIN RELIEF CREAM

shingles pain relief cream cream

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:83176-004

Route of Administration	TOPICAL

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
MENTHOL (UNII: L7T10EIP3A) (MENTHOL - UNII:L7T10EIP3A)	MENTHOL	3 mg in 100 mL

Inactive Ingredients

Ingredient Name	Strength
WATER (UNII: 059QF0KO0R)	
MINERAL OIL (UNII: T5L8T28FGP)	
STEMONA TUBEROSA WHOLE (UNII: 9373ZOT316)	
ALOE (UNII: V5VD430YW9)	
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)	
EUPHORBIA FISCHERIANA ROOT (UNII: R4DRG4762Y)	
SOPHORA FLAVESCENS WHOLE (UNII: X8KX602M5L)	
GLYCERIN (UNII: PDC6A3C0OX)	
ARTEMISIA PRINCEPS WHOLE (UNII: 2849G48V9J)	
COCONUT OIL (UNII: Q9L0O73W7L)	
SOLIDAGO DECURRENS WHOLE (UNII: Y026RJV99T)	
CETYL ALCOHOL (UNII: 936JST6JCN)	
BASSIA SCOPARIA FRUIT (UNII: 04W97Z676Y)	
SMILAX LANCEIFOLIA WHOLE (UNII: WBP9EYZ1XE)	
PHELLODENDRON AMURENSE WHOLE (UNII: 3OZN5NGR9L)	
GLEDITSIA SINENSIS WHOLE (UNII: FS3UB95UTG)	
DICTAMNUS DASYCARPUS ROOT (UNII: 6153LEN214)	
EQUISETUM ARVENSE BRANCH (UNII: 1L0VKZ185E)	
.ALPHA.-TOCOPHEROL (UNII: H4N855PNZ1)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:83176-004-01	100 mL in 1 BOTTLE; Type 0: Not a Combination Product	03/10/2023	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part348	03/10/2023	

Labeler - Dr.luke Healthcare LLC (118868014)

Establishment

Name	Address	ID/FEI	Business Operations
Dr.luke Healthcare LLC		118868014	manufacture(83176-004)

Revised: 3/2023