PAIN RELIEF- lidocaine, menthol cream
HUMN Pharmaceuticals Inc

Pain Relief Cream TPR ^{20}

Active ingredients

Lidocaine Hydrochloride 4.0% (w/w)

Menthol 1.0% (w/w)

Purpose

Anesthetic

Analgesic, anesthetic, and antipruritic

Uses

Temporary relief of:

• pain

• itching associated with:

■ minor burns ■ sunburn ■ minor cuts ■ scrapes ■ insect bites ■ minor skin irritations

Warnings

For external use only.

Do not use

• in large quantities • over large areas of the body • over raw surfaces • over blistered areas

Ask a doctor before use

if child is under 2 years of age, and use only as directed

When using this product

• avoid contact with eyes; if this happens, rinse thoroughly with water

Stop use and ask a doctor

• condition worsens • if symptoms persist for more than 7 days

• clear up and occur again within a few days • you experience: pain, swelling, or blistering

Keep out of reach of children. If swallowed, get

medical help or contact a Poison Control Center right away.

Directions

• adults and children 2 years of age and older: apply to affected area not more than 3 to 4 times daily.

Other information

- do not use if seal is broken
- store at 60-85° F

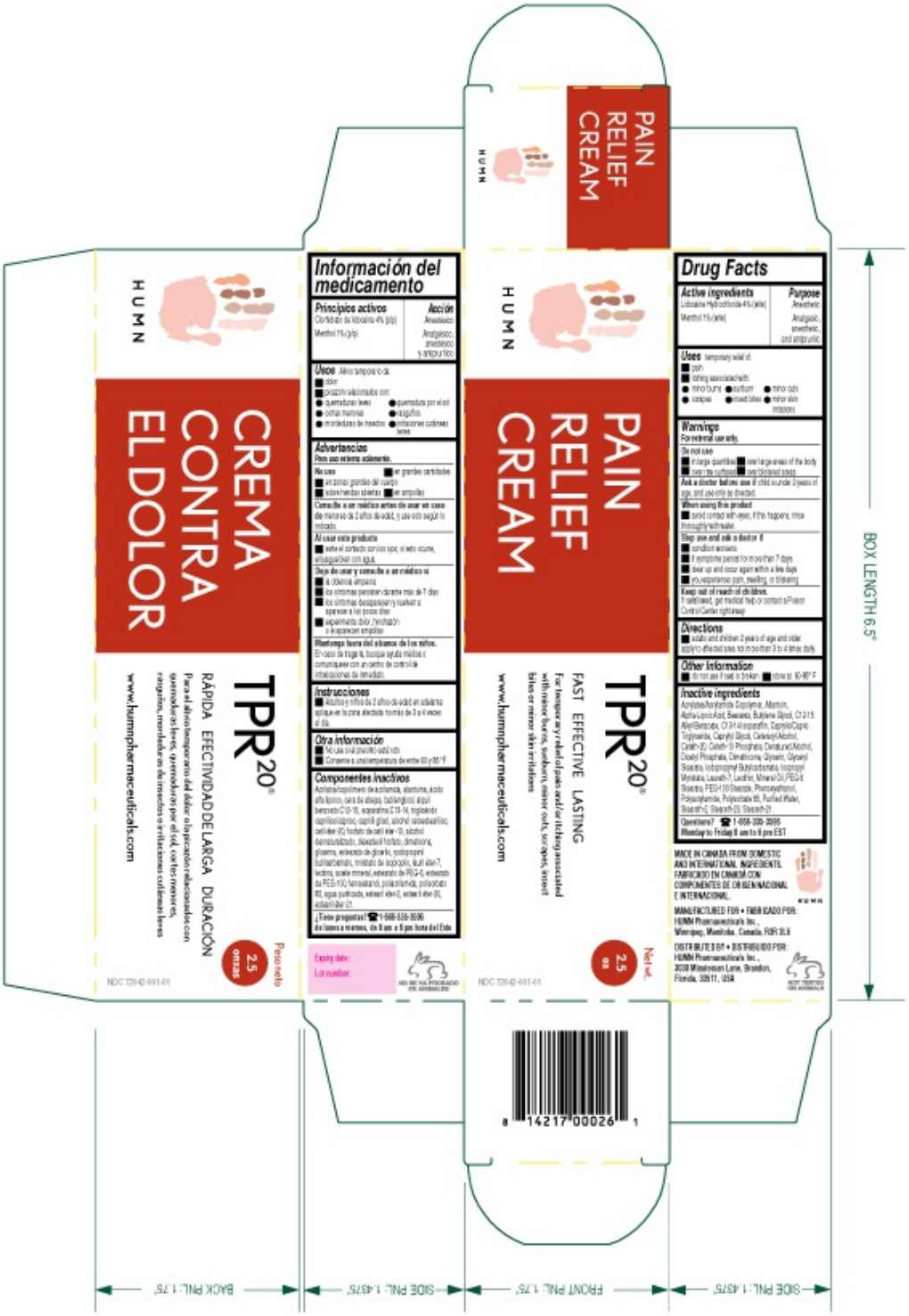
Inactive ingredients

Acrylates/Acrylamide Copolymer, Allantoin, Alpha-Lipoic Acid, Beeswax, Butylene Glycol, C12-15 Alkyl Benzoate, C13-14 Isoparaffin, Caprylic/Capric Triglyceride, Caprylyl Glycol, Ceteraryl Alcohol, Ceteth-20, Ceteth-10 Phosphate, Denatured Alcohol, Dicetyl Phosphate, Dimethicone, Glycerin, Glyceryl Stearate, Iodopropynyl Butylcarbamate, Isopropyl Myristate, Laureth-7, Lecithin, Mineral Oil, PEG-6 Stearate, PEG-100 Stearate, Phenoxyethanol, Polyacrylamide, Polysorbate 85, Purified Water, Steareth-2, Steareth-20, Steareth-21.

Questions?

1-866-335-3596 Monday to Friday 8 am to 6 pm EST .

TPR20 Pain Relief Label



PAIN RELIEF

lidocaine, menthol cream

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:72042-001
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
MENTHOL (UNII: L7T10EIP3A) (MENTHOL - UNII:L7T10EIP3A)	MENTHOL	1 mg in 100 mg
LIDOCAINE HYDROCHLORIDE ANHYDROUS (UNII: EC2CNF7XFP) (LIDOCAINE - UNII:98PI200987)	LIDOCAINE HYDROCHLORIDE ANHYDROUS	4 mg in 100 mg

Inactive Ingredients

Ingredient Name	Strength
CETETH-20 (UNII: I835H2IHHX)	
ALCOHOL (UNII: 3K9958V90M)	
ALUMINUM DICETYL PHOSPHATE (UNII: WMV3R5DS7O)	
GLYCERIN (UNII: PDC6A3C0OX)	
MINERAL OIL (UNII: T5L8T28FGP)	
PEG-6 STEARATE (UNII: 8LQC57C6B0)	
PEG-100 STEARATE (UNII: YD01N1999R)	
POLYACRYLAMIDE (10000 MW) (UNII: E2KR9C9V2I)	
STEARETH-2 (UNII: V56DFE46J5)	
STEARETH-20 (UNII: L0Q8IK9E08)	
CETETH-10 PHOSPHATE (UNII: 4E05O5N49G)	
PEG-120 GLYCERYL STEARATE (UNII: 6941286E4I)	
IODOPROPYNYL BUTYLCARBAMATE (UNII: 603P14DHEB)	
ISOPROPYL MYRISTATE (UNII: 0RE8K4LNJS)	
LAURETH-7 (UNII: Z95S6G8201)	
LECITHIN, SOYBEAN (UNII: 1DI56QDM62)	
PHENOXYETHANOL (UNII: HIE492ZZ3T)	
DIMETHICONE 100 (UNII: RO266O364U)	
POLYSORBATE 85 (UNII: A7F3N56197)	
WATER (UNII: 059QF0KO0R)	
CAPRYLYL GLYCOL (UNII: 00YIU5438U)	
CAPRYLIC/CAPRIC/LAURIC TRIGLYCERIDE (UNII: FJ1H6M2JG9)	
C13-14 ISOPARAFFIN (UNII: E4F12ROE70)	
ACRYLAMIDE (UNII: 20R035KLCI)	
CETYL ALCOHOL (UNII: 936JST6JCN)	
YELLOW WAX (UNII: 2ZA36H0S2V)	
BUTYLENE GLYCOL (UNII: 3XUS85K0RA)	
ALKYL (C12-15) BENZOATE (UNII: A9EJ3J61HQ)	
ALLANTOIN (UNII: 344S277G0Z)	
.ALPHA.-LIPOIC ACID (UNII: 73Y7P0K73Y)	
STEARETH-21 (UNII: 53J3F32P58)	

Product Characteristics

Color	white	**Score**	
Shape		**Size**	
Flavor		**Imprint Code**	
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:72042-001-01	1 in 1 CARTON	05/19/2018	
1		70873.8 mg in 1 TUBE; Type 0: Not a Combination Product		
2	NDC:72042-001-03	1 in 1 CARTON	05/19/2018	
2		28349.5 mg in 1 TUBE; Type 0: Not a Combination Product		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M017	05/19/2018	

Labeler - HUMN Pharmaceuticals Inc (245630272)

Registrant - Delta Pharma Inc (200161730)

Establishment

Name	Address	ID/FEI	Business Operations
Delta Pharma Inc.		200161730	manufacture(72042-001)

Revised: 1/2024 HUMN Pharmaceuticals Inc