PENICHILLIN- benzocaine cream Sambria Pharmaceuticals, 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.

Drug Facts

Active ingredient

Benzocaine 20%

Purpose

External Analgesic

Uses

For temporary relief of pain and itching due to minor skin irritation.

Warnings

For external use only. Avoid contact with eyes. Do not use in large quantities, particularly over raw surfaces or blistered areas.

Stop use and ask doctor if condition worsens or if symptoms persist for more than 7 days or clear up and occur again within a few days.

Keep out of reach of children. If product is swallowed, get medical help or contact a poison control center immediately.

Keep out of reach of children

If swallowed, get medical help or contact a Poison Control Center right away.

Directions

For adults and children two-years or older: Apply to affected area not more than 3 or 4 times daily.

Children under 2 years of age: consult a physician. Apply in a circular motion for 30 to 60 seconds.

Inactive ingredients

Aqua (Deionized Water), Arnica Montana Flower Extract, C13-14 Isoparaffin, Chrondroitin Sulfate, Emu Oil, Ethoxydiglycol, Ethylhexyglycerin, Glucosamine Sulfate, Isopropyl Palmitate, Laureth-7, Melaleuca Alternifolia (Tea Tree) Oil, Methylsulfonylmethane (MSM), Phenoxyethanol, Polyacrylamide, Propylene Glycol,StearicAcid, Triethanolamine

Other information

Protect this product from excessive heat and direct sun. Never tested on animals, just late clients.

Product label

Designed to help DRUG FACTS when tequila fails. ACTIVE INGREDIENTS: PURPOSE: Benzocaine 20% External Analgesic USES: For temporary relief of pain and itching No need for prayers due to minor skin irritation. to prevent or reduce WARNINGS: For external use only. Avoid contact with eyes. Do not use in large tattoo, permanent quantities, particularly over raw surfaces or blistered areas. Stop use and ask doctor if make up, sugaring or condition worsens or if symptoms persist for more than 7 days or clear up and occur again waxing pain. within a few days. Keep out of reach of children. If product is swallowed, get medical help or contact a poison control center Individually wrapped immediately. to maximize DIRECTIONS FOR USE: For adults and children two-years or older: ingredient freshness Apply to affected area not more than 3 or 4 times daily. and potency. Children under 2 years of age: consult a physician. Apply in a circular motion for 30 to 60 seconds. **INACTIVE INGREDIENTS:** Agua (Deionized) Water), Arnica Montana Flower Extract, C13-14 Isoparaffin, Chrondroitin Sulfate, Emu Oil, Ethoxydiglycol, Ethylhexyglycerin, Glucosamine Sulfate, Isopropyl Palmitate, Laureth-7, Melaleuca Alternifolia (Tea Tree) Oil, Methylsulfonylmethane (MSM), Phenoxyethanol, Polyacrylamide, Propylene Glycol, StearicAcid,Triethanolamine **OTHER INFORMATION:** Protect this product from excessive heat and direct sun. Never tested on animals, just late clients. QUESTIONS OR COMMENTS: FDA Registered: NDC No. ######## hello@ladypeng.com PENICHILLIN Distributed by Lady Peng, LLC Portland OR, 97206 MADE IN THE USA **BENZOCAINE 20%** 3ml / .10 fl. oz.

PENICHILLIN benzocaine cream **Product Information** HUMAN OTC DRUG NDC:54723-012 **Product Type** Item Code (Source) **Route of Administration** TOPICAL **Active Ingredient/Active Moiety Basis of Strength** Strength Ingredient Name BENZOCAINE (UNII: U3RSY48/W5) (BENZOCAINE - UNII:U3RSY48/W5) BENZOCAINE 20 g in 100 mL

| Inactive Ingre | dients | | |
|-----------------------|-----------------------------------------------------|-------------------------|-----------------------|
| | Ingredient Name | | Strength |
| C13-14 ISOPARAF | FIN (UNII: E4F12ROE70) | | |
| CHONDROITIN SU | LFATE (BOVINE) (UNII: 6IC1M3OG5Z) | | |
| EMU OIL (UNII: 344 | 821WD61) | | |
| DIETHYLENE GLYC | OL MONOETHYL ETHER (UNII: A1A1I8X02B) | | |
| THYLHEXYLGLYC | ERIN (UNII: 147D247K3P) | | |
| GLUCOSAMINE SU | LFATE (UNII: 1FW7WLR731) | | |
| SOPROPYL PALM | TATE (UNII: 8CRQ2TH63M) | | |
| AURETH-7 (UNII: 2 | 295S6G8201) | | |
| TEA TREE OIL (UNI | I: VIF565UC2G) | | |
| DIMETHYL SULFO | NE (UNII: 9H4PO4Z4FT) | | |
| PHENOXYETHANO | L (UNII: HIE492ZZ3T) | | |
| POLYACRYLAMIDE | (10000 MW) (UNII: E2KR9C9V2I) | | |
| PROPYLENE GLYC | OL (UNII: 6DC9Q167V3) | | |
| STEARIC ACID (UN | II: 4ELV7Z65AP) | | |
| ROLAMINE (UNII: | 9O3K93S3TK) | | |
| NATER (UNII: 059Q | F0KO0R) | | |
| ARNICA MONTANA | FLOWER (UNII: OZ0E5Y15PZ) | | |
| | | | |
| | | | |
| Packaging | | | |
| # Item Code | Package Description | Marketing Start Date | Marketing End Date |
| NDC:54723-012- 01 | 3 mL in 1 PACKET; Type 0: Not a Combination Product | 03/31/2023 | |
| | | | |
| Marketing | Information | | |
| | Application Number or Monograph | Marketing Start | Marketing End |
| Marketing | Citation | Date | Date |
| Marketing Category | Citation | | |

Labeler - Sambria Pharmaceuticals, LLC (078676259)

Revised: 8/2023

Sambria Pharmaceuticals, LLC