**ADVANCED RELIEF A AND D- petrolatum ointment**
**Ultra Seal Corporation**

*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.*

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Active Ingredient (in each gram): White Petrolatum 93.5%

Purpose-Skin Protectant

Uses-Temporarily protects minor cuts, scrapes, and burns. Temporarily protects and helps chapped or cracked skin and lips. Helps protect lips from drying effects of wind and cold weather.

Directions: Apply as needed

Keep out of reach of children. If swallowed, get medical help or contact a Poison Control Center right away.

For external use only. Do not use in eyes, on deep puncture wounds, animal bites, or serious burns or for more than 1 week unless directed by a doctor.

Stop use and ask a doctor if: A rash or allergic reaction develops. Condition worsens, persists, or recurs.

Inactive Ingredients: Corn Oil, Light Mineral Oil, Vitamin A Palmitate, Vitamin D

NDC# 42213-375-05

ADVANCED RELIEF™

THE SCIENCE OF FAST RELIEF.

# A & D OINTMENT

## Skin Protectant

**NET WT. 5.0 grams**

Ultra Seal Corporation, 521 Main St.
New Paltz, NY 12561 • 845-255-2490

Tamper Evident. Do not use if packet is torn, cut or opened.

## Drug Facts

***Active Ingredient*** ***Purpose***

(in each gram)
White Petrolatum
93.5% ..... Skin Protectant

### *Uses:*

- temporarily protects minor cuts, scrapes and burns
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▶

## *Drug Facts* (continued)

### *Warnings:*

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If ingested, contact a Poison Control Center right away.

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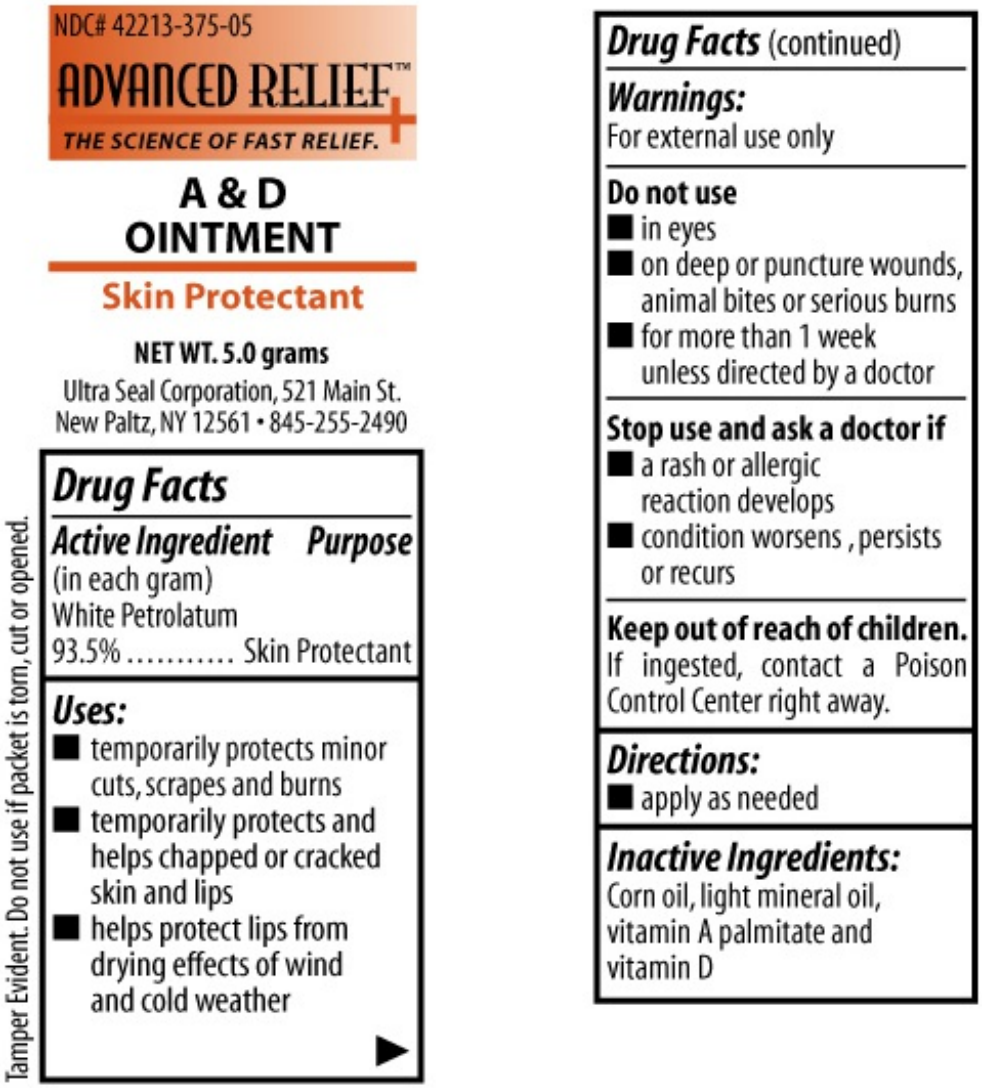


# ADVANCED RELIEF A AND D

petrolatum ointment

| Product Information | | | |
|---|---|---|---|
| **Product Type** | HUMAN OTC DRUG | **Item Code (Source)** | NDC:42213-375 |
| **Route of Administration** | TOPICAL | | |

| Active Ingredient/Active Moiety | | |
|---|---|---|
| **Ingredient Name** | **Basis of Strength** | **Strength** |
| **PETROLATUM** (UNII: 4T6H12BN9U) (PETROLATUM - UNII:4T6H12BN9U) | PETROLATUM | 937.5 mg in 1 g |

| Inactive Ingredients |
|---|

| Ingredient Name | Strength |
|---|---|
| **MINERAL OIL** (UNII: T5L8T28FGP) | |
| **CORN OIL** (UNII: 8470G57WFM) | |
| **VITAMIN A PALMITATE** (UNII: 1D1K0N0VVC) | |
| **ERGOCALCIFEROL** (UNII: VS041H42XC) | |

## Packaging

| # | Item Code | Package Description | Marketing Start Date | Marketing End Date |
|---|---|---|---|---|
| **1** | NDC:42213-375-47 | 144 in 1 CARTON | 12/01/2011 | |
| **1** | NDC:42213-375-05 | 5 g in 1 PACKET; Type 0: Not a Combination Product | | |

## Marketing Information

| Marketing Category | Application Number or Monograph Citation | Marketing Start Date | Marketing End Date |
|---|---|---|---|
| OTC monograph final | part347 | 12/01/2011 | |

## Labeler - Ultra Seal Corporation (085752004)

## Registrant - ULTRAtab Laboratories, Inc. (151051757)

## Establishment

| Name | Address | ID/FEI | Business Operations |
|---|---|---|---|
| Ultraseal Corporation | | 085752004 | pack(42213-375) |

## Establishment

| Name | Address | ID/FEI | Business Operations |
|---|---|---|---|
| Ultratab Laboratories, Inc. | | 151051757 | manufacture(42213-375) |

Revised: 1/2020 Ultra Seal Corporation