**NUTRIXYL ANTI-DANDRUFF- salicylic acid liquid**
**APOLLO HEALTH AND BEAUTY CARE**

*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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**DRUG FACTS**

**ACTIVE INGREDIENT**

SALICYLIC ACID 2%

**PURPOSE**

ANTI-DANDRUFF/ANTI-SEBORRHEIC DERMATITIS/ANTI-PSORIASIS

**USES**

CONTROLS THE SYMPTOMS OF DANDRUFF, SEBORRHEIC DERMATITIS, AND PSORIASIS

**WARNINGS**

FOR EXTERNAL USE ONLY

**ASK A DOCTOR BEFORE USE IF YOU HAVE**

A CONDITION THAT COVERS A LARGE AREA OF THE BODY

WHEN USING THIS PRODUCT

AVOID CONTACT WITH EYES. IF CONTACT OCCURS, RINSE EYES THOROUGHLY WITH WATER. FOR USE ON COLOR TREATED OR PERMED HAIR, RINSE THOROUGHLY

*STOP USE AND ASK A DOCTOR IF*

CONDITION WORSENS OR DOES NOT IMPROVE AFTER REGULAR USE OF THIS PRODUCT AS DIRECTED

**KEEP OUT OF REACH OF CHILDREN**

IN CASE OF ACCIDENTAL INGESTION, GET MEDICAL HELP OR CONTACT A POISON CONTROL CENTER IMMEDIATELY

**DIRECTIONS**

- FOR BEST RESULTS, USE TWICE A WEEK OR AS DIRECTED BY A DOCTOR
- WET HAIR, MASSAGE ONTO SCALP FOR SEVERAL MINUTES, AND THEN RINSE
- REPEAT IF DESIRED

**INACTIVE INGREDIENTS**

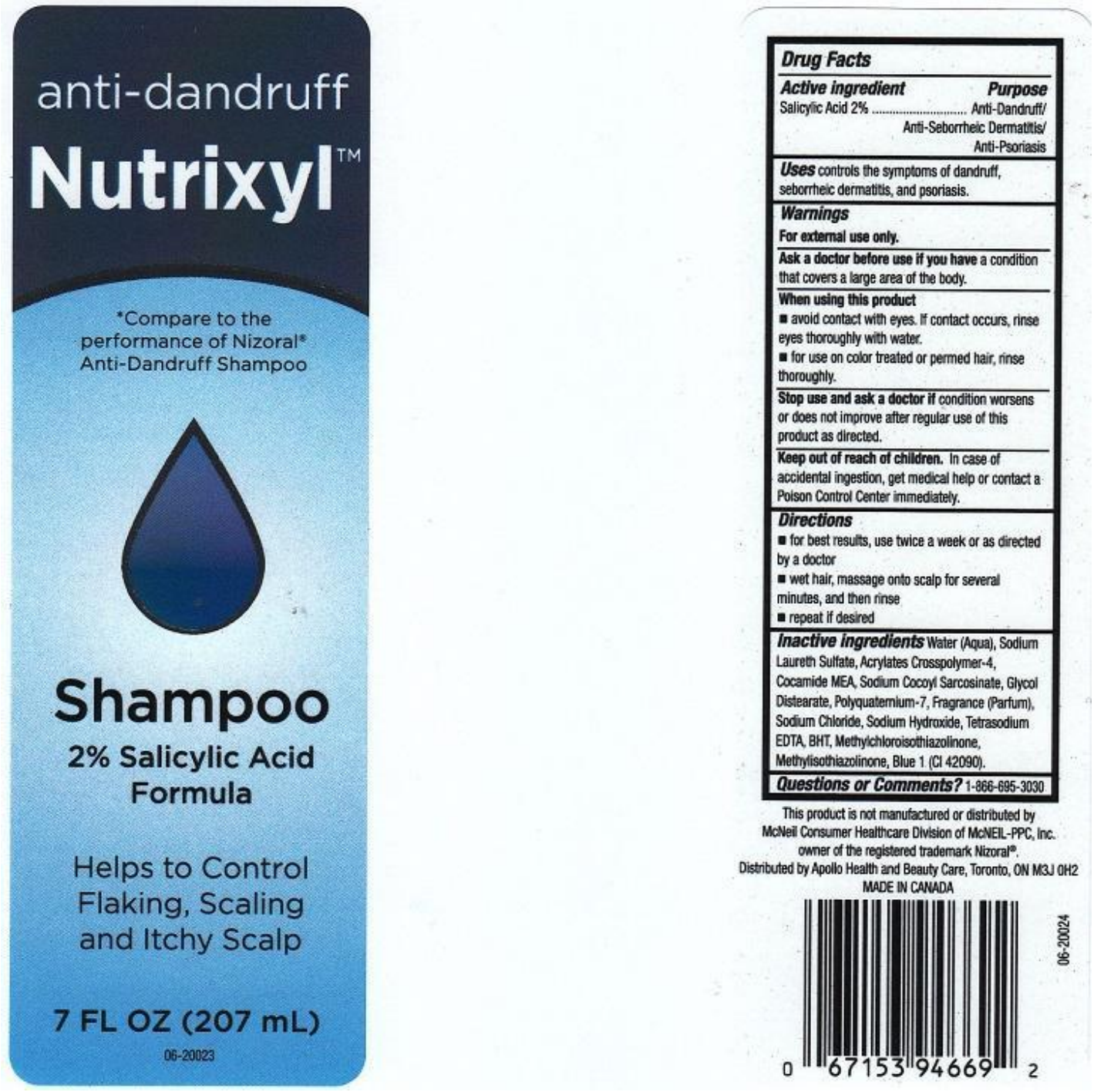
WATER (AQUA), SODIUM LAURETH SULFATE, ACRYLATES CROSSPOLYMER-4, COCAMIDE MEA, SODIUM COCOYL SARCOSINATE, GLYCOL DISTEARATE, POLYQUATERNIUM-7, FRAGRANCE (PARFUM), SODIUM CHLORIDE, SODIUM HYDROXIDE, TETRASODIUM EDTA, BHT, METHYLCHLOROISOTHIAZOLINONE,

METHYLISOTHIAZOLINONE, BLUE 1 (CI 42090)

**QUESTIONS OR COMMENTS?**

1-866-695-3030

**LABEL COPY**




**NUTRIXYL ANTI-DANDRUFF**

salicylic acid liquid

**Product Information**

| | | | |
|---|---|---|---|
| **Product Type** | HUMAN OTC DRUG | **Item Code (Source)** | NDC:63148-450 |
| **Route of Administration** | TOPICAL | | |

## Active Ingredient/Active Moiety

| Ingredient Name | Basis of Strength | Strength |
|---|---|---|
| **SALICYLIC ACID** (UNII: O414PZ4LPZ) (SALICYLIC ACID - UNII:O414PZ4LPZ) | SALICYLIC ACID | 20 mg in 1 mL |

## Inactive Ingredients

| Ingredient Name | Strength |
|---|---|
| **WATER** (UNII: 059QF0KO0R) | |
| **SODIUM LAURETH SULFATE** (UNII: BPV390UAP0) | |
| **CARBOMER COPOLYMER TYPE A (ALLYL PENTAERYTHRITOL CROSSLINKED)** (UNII: 71DD5V995L) | |
| **COCO MONOETHANOLAMIDE** (UNII: C80684146D) | |
| **SODIUM COCOYL SARCOSINATE** (UNII: 1R9DUY89CZ) | |
| **GLYCOL DISTEARATE** (UNII: 13W7MDN21W) | |
| **POLYQUATERNIUM-7 (70/30 ACRYLAMIDE/DADMAC; 1600000 MW)** (UNII: 0L414VCS5Y) | |
| **SODIUM CHLORIDE** (UNII: 451W47IQ8X) | |
| **SODIUM HYDROXIDE** (UNII: 55X04QC32I) | |
| **EDETATE SODIUM** (UNII: MP1J8420LU) | |
| **BUTYLATED HYDROXYTOLUENE** (UNII: 1P9D0Z171K) | |
| **METHYLCHLOROISOTHIAZOLINONE** (UNII: DEL7T5QRPN) | |
| **METHYLISOTHIAZOLINONE** (UNII: 229D0E1QFA) | |
| **FD&C BLUE NO. 1** (UNII: H3R47K3TBD) | |

## Packaging

| # | Item Code | Package Description | Marketing Start Date | Marketing End Date |
|---|---|---|---|---|
| **1** | NDC:63148-450-07 | 207 mL in 1 BOTTLE, PLASTIC | | |

## Marketing Information

| Marketing Category | Application Number or Monograph Citation | Marketing Start Date | Marketing End Date |
|---|---|---|---|
| OTC monograph final | part358H | 09/18/2014 | |

## Labeler - APOLLO HEALTH AND BEAUTY CARE (201901209)

## Registrant - APOLLO HEALTH AND BEAUTY CARE (201901209)

## Establishment

| Name | Address | ID/FEI | Business Operations |
|---|---|---|---|
| APOLLO HEALTH AND BEAUTY CARE | | 201901209 | manufacture(63148-450) |

Revised: 9/2014 APOLLO HEALTH AND BEAUTY CARE