DG BODY TONIC- salicylic acid liquid
DOLGENCORP INC

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

SALICYLIC ACID 2.0%

PURPOSE

ACNE TREATMENT

USES

FOR THE TREATMENT OF ACNE

WARNINGS

FOR EXTERNAL USE ONLY

- FLAMMABLE
- KEEP AWAY FROM FIRE OR HEAT SOURCE

ASK DOCTOR OR PHARMACIST BEFORE USE IF YOU ARE

USING OTHER TOPICAL ACNE MEDICATIONS AT THE SAME TIME OR IMMEDIATELY FOLLOWING USE OF THIS PRODUCT. THIS MAY INCREASE DRYNESS OR IRRITATION OF THE SKIN. iF THIS OCCURS, ONLY ONE MEDICATION SHOULD BE USED UNLESS OTHERWISE DIRECTED BY A DOCTOR

WHEN USING THIS PRODUCT

AVOID CONTACT WITH EYES. IF CONTACT OCCURS, RINSE THOROUGHLY WITH WATER

STOP USING THIS PRODUCT AND ASK A DOCTOR IF

IRRITATION OR REDNESS DEVELOPS AND LASTS

KEEP OUT OF REACH OF CHILDREN

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

DIRECTIONS

- CLEANSE SKIN THOROUGHLY
- MOISTEN A COTTON BALL WITH PRODUCT AND COVER AFFECTED AREA ONE TO THREE TIMES DAILY
- DUE TO EXCESSIVE DRYING OF THE SKIN, START WITH ONE APPLICATION DAILY, INCREASING TO 2 OR 3 IF NEEDED OR AS DIRECTED BY A PHYSICIAN
- IF PEELING OCCURS, REDUCE USE TO ONCE A DAY OR EVERY OTHER DAY

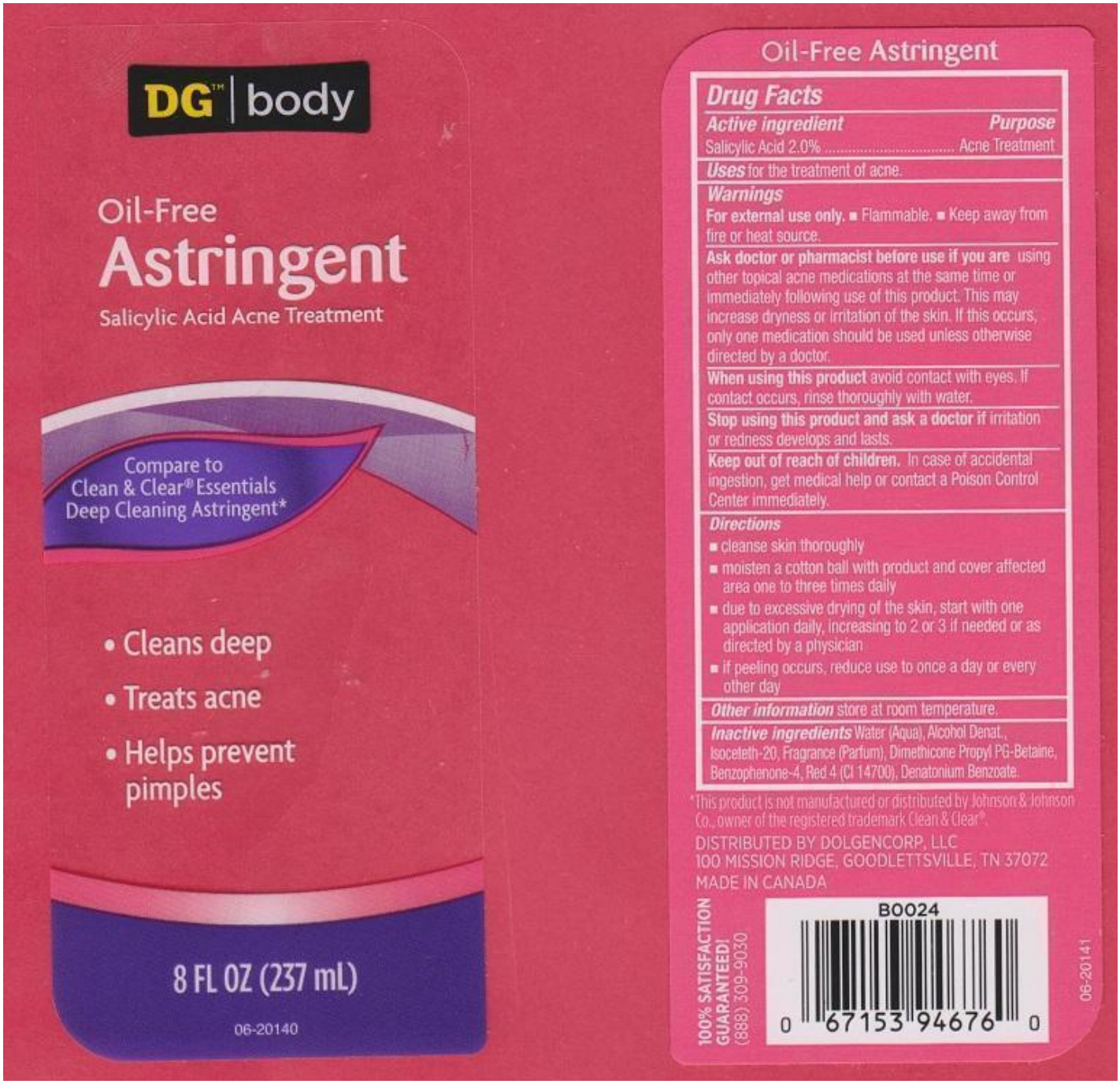
OTHER INFORMATION

STORE AT ROOM TEMPERATURE

INACTIVE INGREDIENTS

WATER (AQUA), ALCOHOL DENAT., ISOCETETH-20, FRAGRANCE (PARFUM), DIMETHICONE PROPYL PG-BETAINE, BENZOPHENONE-4, RED 4 (CI 14700), DENATONIUM BENZOATE

LABEL COPY



DG BODY TONIC
salicylic acid liquid

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:55910-101
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)	
ALCOHOL (UNII: 3K9958V90M)	
ISOCETETH-20 (UNII: O020065R7Z)	
SULISOBENZONE (UNII: 1W6L629B4K)	
FD&C RED NO. 40 (UNII: WZB9127XOA)	
DENATONIUM BENZOATE (UNII: 4YK5Z54AT2)	

Packaging				
#	**Item Code**	**Package Description**	**Marketing Start Date**	**Marketing End Date**
1	NDC:55910-101-08	237 mL in 1 BOTTLE, PLASTIC		

Marketing Information			
Marketing Category	**Application Number or Monograph Citation**	**Marketing Start Date**	**Marketing End Date**
OTC monograph final	part333D	03/02/2015	

Labeler - DOLGENCORP INC (068331990)

Registrant - APOLLO HEALTH AND BEAUTY CARE (201901209)

Establishment			
Name	**Address**	**ID/FEI**	**Business Operations**
APOLLO HEALTH AND BEAUTY CARE		201901209	manufacture(55910-101)

Revised: 3/2015 DOLGENCORP INC