DAWN ULTRA ANTIBACTERIAL HAND ORANGE SCENT- chloroxylenol soap
The Procter & Gamble Manufacturing Company

Dawn ® Ultra Antibacterial Hand

Orange Scent

Drug Facts

Active ingredient

Chloroxylenol 0.30%

Purpose

Antibacterial hand soap

Use

- for handwashing to decrease bacteria on the skin

Warnings

For external use only

Keep out of reach of children. In case of accidental ingestion, drink a glass of water to dilute. If eye contact occurs, rinse thoroughly with water.

Directions

- wet hands and forearms. Apply 5 ml or palmful to hands and forearms. Scrub thoroughly for 30 sec. and rinse.

Inactive ingredients

Water, sodium lauryl sulfate, lauramine oxide, alcohol denat., phenoxyethanol, tetrasodium glutamate diacetate, sodiumchloride, fragrance, deceth-8, C9-11 alketh-8, PPG-26, sodium hydroxide, yellow 5, red 33

Questions?

1-800-725-3296

Distributed by **PROCTER & GAMBLE,**
CINCINNATI, OH 45202

PRINCIPAL DISPLAY PANEL - 1.12L (1.18 QT) Bottle Label

DAWN ®
ULTRA

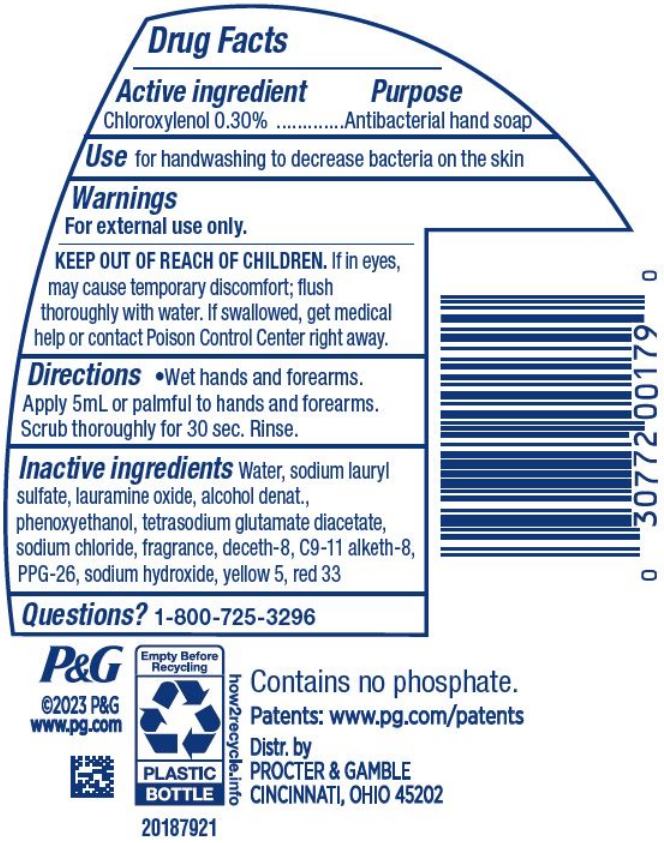
CHLOROXYLENOL ANTIBACTERIAL HAND SOAP

ORANGE SCENT

***VS DAWN NON-CONCENTRATED**

DISHWASHING LIQUID

1.12L (1.18 QT) 38 FL OZ



DAWN ULTRA ANTIBACTERIAL HAND ORANGE SCENT

chloroxylenol soap

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:37000-617
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
CHLOROXYLENOL (UNII: 0F32U78V2Q) (CHLOROXYLENOL - UNII:0F32U78V2Q)	CHLOROXYLENOL	0.3 g in 100 mL

Inactive Ingredients

Ingredient Name	Strength
TETRASODIUM GLUTAMATE DIACETATE (UNII: 5EHL50I4MY)	
DECETH-8 (UNII: 19FQ96EA8Q)	
ALCOHOL (UNII: 3K9958V90M)	
WATER (UNII: 059QF0KO0R)	
SODIUM LAURYL SULFATE (UNII: 368GB5141J)	
LAURAMINE OXIDE (UNII: 4F6FC4MI8W)	
PHENOXYETHANOL (UNII: HIE492ZZ3T)	
SODIUM CHLORIDE (UNII: 451W47IQ8X)	
PPG-26 (UNII: V86KZL3H2Z)	
FD&C YELLOW NO. 5 (UNII: I753WB2F1M)	
D&C RED NO. 33 (UNII: 9DBA0SBB0L)	
SODIUM HYDROXIDE (UNII: 55X04QC32I)	
C9-11 PARETH-8 (UNII: 80E6PSE1XL)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:37000-617-26	266 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	07/07/2014	01/10/2018
2	NDC:37000-617-59	591 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	07/07/2014	01/10/2018
3	NDC:37000-617-63	638 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	07/07/2014	12/12/2019
4	NDC:37000-617-70	709 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	07/07/2014	01/10/2018
5	NDC:37000-617-10	1010 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	07/07/2014	12/12/2019
6	NDC:37000-617-11	1120 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	07/07/2014	01/10/2018
7	NDC:37000-617-16	1660 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	07/07/2014	08/01/2024
8	NDC:37000-617-23	236 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	02/10/2017	02/01/2021
9	NDC:37000-617-53	532 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	02/10/2017	12/12/2019
10	NDC:37000-617-22	2210 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	02/10/2017	01/01/2022
11	NDC:37000-617-47	479 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	02/02/2018	12/01/2021
12	NDC:37000-617-57	573 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	02/02/2018	02/01/2025
13	NDC:37000-617-12	1210 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	02/02/2018	04/01/2021
14	NDC:37000-617-82	828 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	02/02/2018	
15	NDC:37000-617-20	207 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	04/02/2019	02/01/2024
16	NDC:37000-617-40	1200 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	04/02/2019	12/01/2023
17	NDC:37000-	192 mL in 1 BOTTLE, PLASTIC; Type 0: Not a	12/30/2021	09/01/2024

17	617-19	Combination Product	12/30/2021	09/01/2024
18	NDC:37000-617-38	1120 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	12/30/2021	
19	NDC:37000-617-21	221 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	12/01/2022	11/01/2025
20	NDC:37000-617-54	532 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	02/07/2023	10/01/2025
21	NDC:37000-617-65	650 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	02/07/2023	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	505G(a)(3)	07/07/2014	

Labeler - The Procter & Gamble Manufacturing Company (004238200)

Revised: 9/2024 The Procter & Gamble Manufacturing Company