

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



Drug Facts

Active ingredient	Purpose
Chloroxylenol 0.30%	Antibacterial hand soap

Use for handwashing to decrease bacteria on the skin

Warnings
For external use only.
KEEP OUT OF REACH OF CHILDREN. If in eyes, may cause temporary discomfort; flush thoroughly with water. If swallowed, get medical help or contact Poison Control Center right away.

Directions •Wet hands and forearms. Apply 5mL or palmful to hands and forearms. Scrub thoroughly for 30 sec. Rinse.

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

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P&G
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Empty Before Recycling
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PLASTIC BOTTLE
20187921

Contains no phosphate.
Patents: www.pg.com/patents
Distr. by
PROCTER & GAMBLE
CINCINNATI, OHIO 45202

0 30772 00179 0

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	02/01/2024

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	M003	07/07/2014	

Labeler - The Procter & Gamble Manufacturing Company (004238200)