

**FAMILY DOLLAR DAYTIME AND NIGHTTIME COMBO PACK COLD AND FLU-
acetaminophen, dextromethorphan hbr, phenylephrine hydrochloride, and
doxylamine succinate
FAMILY DOLLAR SERVICES 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.

**Family Dollar Daytime & Nighttime Combo Pack Cold & Flu
FAMILY DOLLAR DAYTIME COLD & FLU RELIEF**

Active ingredients (in each 15 mL)

Acetaminophen 325 mg
Dextromethorphan HBr 10 mg
Phenylephrine HCl 5 mg

Purpose

Pain reliever/Fever reducer
Cough suppressant
Nasal decongestant

Uses

temporarily relieves common cold/flu symptoms:

- nasal congestion
- cough due to minor throat & bronchial irritation
- minor aches & pains
- headache
- fever
- sore throat

Warnings

If pregnant or breast-feeding, ask a health professional before use.

Keep out of reach of children. In case of overdose, get medical help or contact a Poison Control Center right away. Quick medical attention is critical for adults as well as for children even if you do not notice any signs or symptoms.

Directions

- take only as directed
- Only use the dose cup provided

- do not exceed 4 doses per 24 hrs

adults & children 12 yrs & over	30 mL every 4 hours
Children 6 to under 12 yrs	15 mL every 4 hours
children 4 to under 12 yrs	ask a doctor
children under 4 yrs	do not use

Other information

each 15 mL contains:

- sodium 42 mg
- store at room temperature
- Do not refrigerate.

Inactive ingredients

Anhydrous citric acid, disodium edetate, FD&C Yellow No. 6, flavor, glycerin, menthol, potassium citrate, propyl gallate, purified water, sodium benzoate, sodium chloride, sorbitol, sucralose, xanthan gum

Questions?

1-866-467-2748

FAMILY DOLLAR NIGHTTIME COLD & FLU RELIEF

Active ingredients (in each 30 mL)

Acetaminophen 650 mg
 Dextromethorphan HBr 30 mg
 Doxylamine succinate 12.5 mg

Purpose

Pain reliever/Fever reducer
 Cough suppressant
 Antihistamine

Uses

temporarily relieves common cold/flu symptoms:

- cough due to minor throat & bronchial irritation
- minor aches & pains
- headache
- fever
- sore throat

- runny nose & sneezing

Warnings

If pregnant or breast-feeding, ask a health professional before use.

Keep out of reach of children. In case of overdose, get medical help or contact a Poison Control Center right away. Quick medical attention is critical for adults as well as for children even if you do not notice any signs or symptoms.

Directions

- take only as directed
- Only use the dose cup provided
- do not exceed 4 doses per 24 hrs

adults & children 12 yrs & over	30 mL every 4 hours
children 4 to under 12 yrs	ask a doctor
children under 4 yrs	do not use

Other information

- **each 30 mL contains:** sodium 16 mg
- Store at room temperature
- Do not refrigerate

Inactive ingredients

Citric acid, D&C Yellow No. 10, disodium edetate, FD&C Green No.3, FD&C Yellow no 6, flavor, propyl gallate, propylene glycol, purified water, sodium benzoate, sodium citrate, sorbitol, sucralose

Questions?

1-866-467-2748

Distributed By:

PRINCIPAL DISPLAY PANEL - Kit Carton

Compare to the active ingredients in Vicks®DayQuil™ & NyQuil™ Cold & Flu*

NDC# 55319-808-16

DayTime

Cold & Flu

Relief

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:55319-808-16	1 in 1 PACKAGE; Type 0: Not a Combination Product	08/21/2023	

Quantity of Parts

Part #	Package Quantity	Total Product Quantity
Part 1	1 BOTTLE	236 mL
Part 2	1 BOTTLE	236 mL

Part 1 of 2

FD DAYTIME COLD AND FLU RELIEF

acetaminophen, dextromethorphan hydrobromide, and phenylephrine hydrochloride liquid

Product Information

Item Code (Source)	NDC:55319-036
Route of Administration	ORAL

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
ACETAMINOPHEN (UNII: 362O9ITL9D) (ACETAMINOPHEN - UNII:362O9ITL9D)	ACETAMINOPHEN	325 mg in 15 mL
DEXTROMETHORPHAN HYDROBROMIDE (UNII: 9D2RTI9KYH) (DEXTROMETHORPHAN - UNII:7355X3ROTS)	DEXTROMETHORPHAN HYDROBROMIDE	10 mg in 15 mL
PHENYLEPHRINE HYDROCHLORIDE (UNII: 04JA59TNSJ) (PHENYLEPHRINE - UNII:1WS297W6MV)	PHENYLEPHRINE HYDROCHLORIDE	5 mg in 15 mL

Inactive Ingredients

Ingredient Name	Strength
ANHYDROUS CITRIC ACID (UNII: XF417D3PSL)	
EDETATE DISODIUM (UNII: 7FLD91C86K)	
FD&C YELLOW NO. 6 (UNII: H77VEI93A8)	
GLYCERIN (UNII: PDC6A3C0OX)	
MENTHOL, UNSPECIFIED FORM (UNII: L7T10EIP3A)	
POTASSIUM CITRATE (UNII: EE90ONI6FF)	
PROPYL GALLATE (UNII: 8D4SNN7V92)	
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)	
WATER (UNII: 059QF0KO0R)	
SODIUM BENZOATE (UNII: OJ245FE5EU)	
SODIUM CHLORIDE (UNII: 451W47IQ8X)	

SORBITOL (UNII: 506T60A25R)	
SUCRALOSE (UNII: 96K6UQ3ZD4)	
XANTHAN GUM (UNII: TTV12P4NEE)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:55319-036-08	236 mL in 1 BOTTLE; Type 0: Not a Combination Product		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph final	part341	08/21/2023	

Part 2 of 2

FD NIGHTTIME COLD AND FLU RELIEF

acetaminophen, doxylamine succinate, and dextromethorphan hydrobromide liquid

Product Information

Item Code (Source)	NDC:55319-096
Route of Administration	ORAL

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
ACETAMINOPHEN (UNII: 362O9ITL9D) (ACETAMINOPHEN - UNII:362O9ITL9D)	ACETAMINOPHEN	650 mg in 30 mL
DEXTROMETHORPHAN HYDROBROMIDE (UNII: 9D2RTI9KYH) (DEXTROMETHORPHAN - UNII:7355X3ROTS)	DEXTROMETHORPHAN HYDROBROMIDE	30 mg in 30 mL
DOXYLAMINE SUCCINATE (UNII: V9B19B5Y12) (DOXYLAMINE - UNII:95QB77JKPL)	DOXYLAMINE SUCCINATE	12.5 mg in 30 mL

Inactive Ingredients

Ingredient Name	Strength
ANHYDROUS CITRIC ACID (UNII: XF417D3PSL)	
D&C YELLOW NO. 10 (UNII: 35SW5USQ3G)	
EDETATE DISODIUM (UNII: 7FLD91C86K)	
FD&C GREEN NO. 3 (UNII: 3P3ONR6O1S)	
FD&C YELLOW NO. 6 (UNII: H77VEI93A8)	
PROPYL GALLATE (UNII: 8D4SNN7V92)	
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)	

WATER (UNII: 059QF0KO0R)	
SODIUM BENZOATE (UNII: OJ245FE5EU)	
SODIUM CITRATE, UNSPECIFIED FORM (UNII: 1Q73Q2JULR)	
SORBITOL (UNII: 506T60A25R)	
SUCRALOSE (UNII: 96K6UQ3ZD4)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:55319-096-08	236 mL in 1 BOTTLE; Type 0: Not a Combination Product		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph final	part341	08/21/2023	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph final	part341	08/21/2023	

Labeler - FAMILY DOLLAR SERVICES INC (024472631)

Revised: 9/2023

FAMILY DOLLAR SERVICES INC