CHILDRENS MULTI-SYMPTOM COLD AND FEVER LIQUID - acetaminophen, dextromethorphan hbr, guaifenesin, phenylephrine hcl liquid
Wal-Mart Stores, 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.

Equate Children's Multi-Symptom Cold & Fever Liquid

ACTIVE INGREDIENTS (in each 10 mL)

Acetaminophen 325 mg

Dextromethorphan HBr 10 mg

Guaifenesin 200 mg

Phenyephrine HCl 5 mg

PURPOSE

Pain reliever / fever reducer

Cough Suppressant

Expectorant

Nasal decongestant

USE(S)

temporarily relieves these common cold and flu symptoms:

- nasal congestion
- stuffy nose
- cough due to minor throat and bronchial irritation
- the intensity of coughing
- the impulse to cough to help your child get to sleep
- minor aches and pains
- sore throat
- headache
- temporarily reduces fever
- helps loosen phlegm (mucus) and thin bronchial secretions to drain bronchial tubes and make coughs more productive

WARNINGS

Liver warning: This product contains acetaminophen. Severe liver damage may occur if your child takes

- more than 5 doses in 24 hous, which is the maximum daily amount
- with other drugs containing acetaminophen.

Allergy alert: acetaminophen may cause severe skin reactions. Symptoms may include:

- skin reddening

- blisters
- rash

If a skin reaction occurs, stop use and seek medical help right away

Sore throat warning: if sore throat is severe, persists for more than 2 days, is accompanied or followed by fever, headache, rash, nausea or vomiting, consult a doctor promptly.

DO NOT USE

- **with any other drug containing acetaminophen** (prescription or nonprescription). If you are not sure whether a drug contains acetaminophen, ask a doctor or pharmacist.
- in a child who is taking a prescription monoamine oxidase inhibitor (MAOI) (certain drugs for depression, psychiatric, or emotional conditions, or Parkinson's disease), or for 2 weeks after stopping the MAOI drug. If you do not know if your child's prescription drug contains an MAOI, ask a doctor or pharmacist before giving this product.

ASK A DOCTOR BEFORE USE IF THE CHILD HAS

- liver disease
- thyroid disease
- diabetes
- heart disease
- high blood pressure
- a breathing problem such as chronic bronchitis
- persistent or chronic cough such as occurs with asthma
- cough that occurs with too much phlegm (mucus)

ASK A DOCTOR OR PHARMACIST BEFORE USE IF

- your child is taking the blood thinning drug warfarin

WHEN USING THIS PRODUCT

do not use more than directed

STOP USE AND ASK DOCTOR IF

- nervousness, dizziness or sleeplessness occur
- pain, nasal congestion or cough gets worse or lasts more than 5 days
- fever gets worse or lasts more than 3 days
- redness or swelling is present
- new symptoms occur
- cough comes back, or occurs with rash or headache that lasts. These could be signs of a serious condition.

KEEP OUT OF REACH OF CHILDREN

Overdose warning: Taking more than the recommended dose (overdose) may cause liver damage. In case of overdose, get medical help or contact a Poison Control Center right away. Quick medical attention is critical even if you do not notice any signs or symptoms.

DIRECTIONS

- **this product does not contain directions or complete warnings for adult use**
- **do not give more than directed (see Overdose warning)**
- **shake well before use**
- do not give more than 5 doses in any 24-hour period
- if needed, repeat dose every 4 hours while symptoms last
- do not give more than 5 days unless directed by a doctor
- measure only with dosing cup provided
- do not use dosing cup with other products
- dose as follows or as directed by a doctor
- **Children 6 to under 12 years of age:** 10 mL in dosing cup provided.
- mL = milliliter
- **Children under 6 years of age:** do not use.

OTHER INFORMATION

- **each 10 mL contains:** sodium 10 mg
- store between 15-30°C (59-86°F)
- do not refrigerate
- dosing cup provided

INACTIVE INGREDIENTS

citric acid anhydrous, edetate disodium, FD&C blue #1, FD&C red #40, flavors, glycerin, propylene glycol, propyl gallate, purified water, sodium benzoate, sodium citrate, sorbitol, sucralose, xanthan gum.

QUESTIONS OR COMMENTS

1-888-287-1915

PRINCIPAL DISPLAY PANEL

NDC 49035-772-03

equate

Compare to Children's Mucinex®Multi-Symptom Cold & Fever active ingredients*

Children's Multi-Symptom Cold & Fever Liquid

Acetaminophen 325 mg
Pain reliever/Fever Reducer

Dextromethorphan HBr 10 mg

Cough Suppressant
Guaifenesin 200 mg

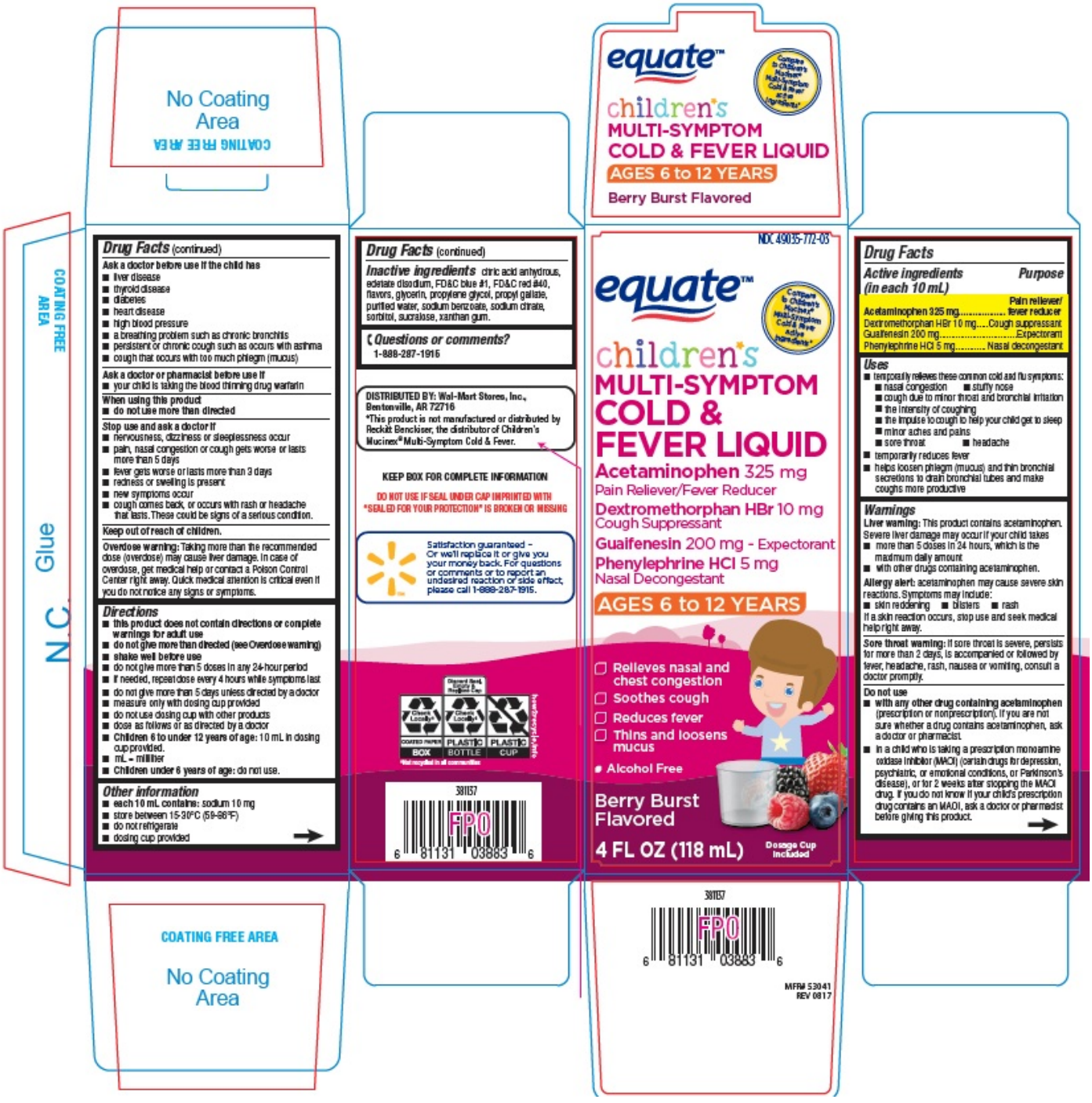
Expectorant
Phenylephrine HCl 5 mg

Nasal Decongestant

AGES 6 to 12 YEARS

- Relieves nasal and chest congestion
- Soothes Cough
- Reduces fever
- Thins and loosens Mucus
- Alcohol Free
- Berry Burst Flavored

4 FL OZ (118 mL)



CHILDRENS MULTI-SYMPTOM COLD AND FEVER LIQUID

acetaminophen, dextromethorphan hbr, guaifenesin, phenylephrine hcl liquid

Product Information

Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:49035-772
Route of Administration	ORAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
ACETAMINOPHEN (UNII: 362O9ITL9D) (ACETAMINOPHEN - UNII:362O9ITL9D)	ACETAMINOPHEN	325 mg in 10 mL
DEXTROMETHORPHAN HYDROBROMIDE (UNII: 9D2RTI9KYH) (DEXTROMETHORPHAN - UNII:7355X3ROTS)	DEXTROMETHORPHAN HYDROBROMIDE	10 mg in 10 mL
GUAIFENESIN (UNII: 495W7451VQ) (GUAIFENESIN - UNII:495W7451VQ)	GUAIFENESIN	200 mg in 10 mL
PHENYLEPHRINE HYDROCHLORIDE (UNII: 04JA59TNSJ) (PHENYLEPHRINE - UNII:1WS297W6MV)	PHENYLEPHRINE HYDROCHLORIDE	5 mg in 10 mL

Inactive Ingredients

Ingredient Name	Strength
ANHYDROUS CITRIC ACID (UNII: XF417D3PSL)	
EDETATE DISODIUM (UNII: 7FLD91C86K)	
FD&C BLUE NO. 1 (UNII: H3R47K3TBD)	
FD&C RED NO. 40 (UNII: WZB9127XOA)	
GLYCERIN (UNII: PDC6A3C0OX)	
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)	
PROPYL GALLATE (UNII: 8D4SNN7V92)	
WATER (UNII: 059QF0KO0R)	
SODIUM BENZOATE (UNII: OJ245FE5EU)	
SODIUM CITRATE, UNSPECIFIED FORM (UNII: 1Q73Q2JULR)	
SORBITOL (UNII: 506T60A25R)	
SUCRALOSE (UNII: 96K6UQ3ZD4)	
XANTHAN GUM (UNII: TTV12P4NEE)	

Product Characteristics

Color	BLUE	**Score**	
Shape		**Size**	
Flavor	BERRY	**Imprint Code**	
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:49035-772-03	1 in 1 CARTON	08/01/2017	
1		118 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	08/01/2017	

Labeler - Wal-Mart Stores, Inc. (051957769)

Revised: 1/2018 Wal-Mart Stores, Inc.