#### AMAZON BASIC CARE CHILDRENS COUGH DM- dextromethorphan polistirex suspension Praxis, LLC

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### Amazon Children's Cough DM Drug Facts

### Active ingredient (in each 5 mL)

Dextromethorphan polistirex equivalent to 30 mg dextromethorphan hydrobromide

#### Purpose

Cough suppressant

#### Uses

temporarily relieves

- cough due to minor throat and bronchial irritation as may occur with the common cold or inhaled irritants
- the impulse to cough to help you get to sleep

#### Warnings

#### Do not use

if you are now 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 prescription drug contains an MAOI, ask a doctor or pharmacist before taking this product.

**Allergy Alert:**Contains sodium metabisulfite, a sulfite that may cause allergic-type reactions.

#### Ask a doctor before use if you have

- chronic cough that lasts as occurs with smoking, asthma or emphysema
- cough that occurs with too much phlegm (mucus)

#### Stop use and ask a doctor if

- side effects occur. You may report side effects to FDA at 1-800-FDA-1088.
- cough lasts more than 7 days, cough comes back, or occurs with fever, rash or headache that
- lasts. These could be signs of a serious condition.

#### 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 (1-800-222-1222).

### Directions

- shake bottle well before use
- measure only with dosing cup provided
- do not use dosing cup with other products
- dose as follows or as directed by doctor

10 mL every 12 hours, not to
exceed 20 mL in 24 hours
5 mL every 12 hours, not to
exceed 10 mL in 24 hours
2.5 mL every 12 hours, not to
exceed 5 mL in 24 hours
do not use

## Other information

- each 5 mL contains: sodium 5 mg
- store at 20° to 25°C (68° to 77°F)
- dosing cup provided

### Inactive ingredients

D&C Red #30 aluminum lake, D&C Yellow #10 aluminum lake, glycerin, high fructose corn syrup, methylparaben, natural and artificial orange flavor, polysorbate 80, polyvinyl acetate, povidone, propylparaben, purified water, sodium metabisulfite, sodium polystyrene sulfonate, sucrose, tartaric acid, tragacanth gum, triacetin, xanthan gum

### Questions or comments?

1-800-719-9260

### Package/Label Principal Display Panel

Age 4+

Compare to Children's Delsym <sup>®</sup>active ingredient

Children's Cough DM

Dextromethorphan Polistirex

Extended-Release Oral Suspension

Cough Suppressant

12 Hour Cough Relief

# Contains sodium metabisulfite, a sulfite that may cause allergic-type reactions

Day or Night

For Children & Adults

Alcohol-free

- Orange Flavored Liquid
- Dosing Cup Included
- 5 FL OZ (148 mL)



	aometrorpha	an polistirex s	suspension					
Pr	oduct Infor	mation						
Pr	oduct Type		HUMAN OTC DRUG Item Code (Source)			NDC:59368-253		
Ro	ute of Admini	stration	ORAL					
Ac	tive Ingredi	ent/Active	Moiety					
		Ingre	edient Name			Basis of Stren		Strengt
DEXTROMETHORPHAN HYDROI (DEXTROMETHORPHAN - UNII:7355						DEXTROMETHORPHAN HYDROBROMIDE		30 mg in 5 mL
In	active Ingre	dients						
			Ingredient Name				St	rength
GĽ	YCERIN (UNII: PE	DC6A3C0OX)						
110	GH FRUCTOSE C		JNII: XY6UN3QB6S)					
ME	THYLPARABEN	(UNII: A2I8C7HI	9Т)					
<b>°</b> 0	LYSORBATE 80	(UNII: 60ZP392	ZG8H)					
<b>°</b> 0	VIDONE, UNSPE	ECIFIED (UNII:	Z989GH94E)					
PR	OPYLPARABEN	(UNII: Z8IX2SC1	.OH)					
	<b>TER</b> (UNII: 059Q							
	DIUM METABIS							
			<b>ATE</b> (UNII: 1699G8679Z)					
	CROSE (UNII: C1							
	RTARIC ACID (U							
	AGACANTH (UNI							
	IACETIN (UNII: X		•					
	NTHAN GUM (UI		·					
	C YELLOW NO.							
	C RED NO. 30 (							
PO	LYVINYL ACETA	<b>(UNII: 32K4</b> )	972 K2U)					
Pr	oduct Chara	acteristics						
Co	lor		orange	Score				
Shape				Size				
Flavor			ORANGE	Imprint Code				
Contains								
D-	ickaging							
r d	ickaying				<b>P4 1</b>	ating Chart		time Furl
#	ltem Code	Pa	ckage Description		Marl	ceting Start Date		ting End ate
	NDC:59368-253-					021		

-	Product								
Marketing Information									
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date						
ANDA	ANDA091135	07/20/2021							

Labeler - Praxis, LLC (016329513)

Establishment					
Name	Address	ID/FEI	Business Operations		
Praxis, LLC		016329513	manufacture(59368-253) , label(59368-253) , pack(59368-253)		

Revised: 1/2023

Praxis, LLC