

**ACTINEL DM PEDIATRIC- dextromethorphan hydrobromide, guaifenesin,
phenylephrine hydrochloride liquid
ACTIPHARMA, LLC**

ACTINEL® DM Pediatric

Drug Facts

Active ingredients (in each 5 mL teaspoonful)

Dextromethorphan HBr 5 mg

Guaifenesin 75 mg

Phenylephrine HCl 2.5 mg

Purposes

Cough suppressant

Expectorant

Nasal decongestant

Uses

- helps loosen phlegm (mucus) and thin bronchial secretions to make coughs more productive
- temporarily relieves:
 - nasal congestion due to the common cold
 - cough due to minor throat and bronchial irritation associated with a cold or inhaled irritants

Warnings

Do not use 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 your child has

- heart disease • high blood pressure • thyroid disease
- diabetes • cough accompanied by excessive phlegm (mucus)
- persistent or chronic cough such occurs with asthma

When using this product

- **do not exceed recommended dosage**

Stop use and ask a doctor if

- your child gets nervous, dizzy, or sleepless
- symptoms do not improve within 7 days or are accompanied by fever
- cough persists for more than 1 week, tends to recur, or is accompanied by fever, rash, or a persistent headache. These could be signs of a serious condition.

Keep out of reach of children. In case of accidental overdose, seek professional assistance or contact a poison control center immediately.

Directions

- do not exceed 6 doses in any 24-hour period
- mL = milliliter, tsp = teaspoon

children 6 to under 12 years of age	10 mL (2 tsp) every 4 hours
children 2 to under 6 years of age	5 mL (1 tsp) every 4 hours
children under 2 years of age	consult a doctor

Other information

- tamper evident feature: Do not use if inner seal is torn, cut, or opened
- store at controlled room temperature 15°- 30°C (59° - 86°F)
- avoid excessive heat or humidity.

Inactive ingredients

citric acid, flavor, glycerin, methylparaben, propylene glycol, propylparaben, purified water, sodium citrate, sucralose

Questions or comments?

1.787.608.0882

You may also report serious side effects to this phone number.

Call weekdays from 8AM to 4PM AST

Alcohol, Dye & Sugar FREE

Contains the same active ingredients as Tusnel® DM Pediatric*

ActiPharma

COMMITTED TO HEALTH AND WELL-BEING

WWW.ACTIPHARMA.NET

Manufactured for Actipharma

San Juan, PR 00917.

Manufactured in USA with imported ingredients

* Tusnel® DM Pediatric is a registered trademark of Llorens Pharmaceutical. This product is not manufactured, distributed or marketed by Llorens Pharmaceutical.

Packaging

NDC 63102-130-16

ACTINEL® DM Pediatric

**Dextromethorphan HBr
COUGH SUPPRESSANT
Guaifenesin
EXPECTORANT
Phenylephrine HCl
NASAL DECONGESTANT**

Alcohol, Dye & Sugar FREE

Contains the same active ingredients as Tusnel® DM Pediatric*

Strawberry Banana Flavor

16 fl oz (1 pt.) 473 mL

Drug Facts

Active ingredients (in each 5 mL teaspoonful) **Purposes**

Dextromethorphan HBr 5 mg..... Cough suppressant

Guaifenesin 75 mg.....Expectorant

Phenylephrine HCl 2.5 mg..... Nasal decongestant

Uses

- helps loosen phlegm (mucus) and thin bronchial secretions to make coughs more productive
- temporarily relieves:
 - nasal congestion due to the common cold
 - cough due to minor throat and bronchial irritation associated with a cold or inhaled irritants

Warnings

Do not use 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 your child has

- heart disease ■ high blood pressure ■ thyroid disease
- diabetes ■ cough accompanied by excessive phlegm (mucus)
- persistent or chronic cough such occurs with asthma

When using this product

- do not exceed recommended dosage

Stop use and ask a doctor if

- your child gets nervous, dizzy, or sleepless
- symptoms do not improve within 7 days or are accompanied by fever
- cough persists for more than 1 week, tends to recur, or is accompanied by fever, rash or a persistent headache. These could be signs of a serious condition.

Drug Facts (continued)

Keep out of reach of children. In case of accidental overdose, seek professional assistance or contact a poison control center immediately.

Directions ■ do not exceed 6 doses in any 24-hour period

- mL = milliliter, tsp = teaspoon

children 6 to under 12 years of age	10 mL (2 tsp) every 4 hours
children 2 to under 6 years of age	5 mL (1 tsp) every 4 hours
children under 2 years of age	consult a doctor

Other information

- tamper evident feature: Do not use if inner seal is torn, cut, or opened
- store at controlled room temperature 15°-30°C (59°-86°F)
- avoid excessive heat or humidity

Inactive ingredients

citric acid, flavor, glycerin, methylparaben, propylene glycol, propylparaben, purified water, sodium citrate, sucralose

Questions or comments?

1.787.608.0882

You may also report serious side effects to this phone number.

Call weekdays from 8AM to 4PM AST

Manufactured for ActiPharma
San Juan, PR 00917
www.actipharma.net
**Manufactured in USA
with imported ingredients**

Rev. 12/25

3 63102 13016 8

ACTINEL DM PEDIATRIC

dextromethorphan hydrobromide, guaifenesin, phenylephrine hydrochloride liquid

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:63102-130
Route of Administration	ORAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
DEXTROMETHORPHAN HYDROBROMIDE (UNII: 9D2RTI9KYH) (DEXTROMETHORPHAN - UNII:7355X3ROTS)	DEXTROMETHORPHAN HYDROBROMIDE	5 mg in 5 mL
GUAIFENESIN (UNII: 495W7451VQ) (GUAIFENESIN - UNII:495W7451VQ)	GUAIFENESIN	75 mg in 5 mL
PHENYLEPHRINE HYDROCHLORIDE (UNII: 04JA59TNSJ) (PHENYLEPHRINE - UNII:1WS297W6MV)	PHENYLEPHRINE HYDROCHLORIDE	2.5 mg in 5 mL

Inactive Ingredients

Ingredient Name	Strength
CITRIC ACID MONOHYDRATE (UNII: 2968PHW8QP)	
GLYCERIN (UNII: PDC6A3C0OX)	
METHYLPARABEN (UNII: A2I8C7HI9T)	
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)	
PROPYLPARABEN (UNII: Z8IX2SC1OH)	
WATER (UNII: 059QF0KO0R)	
SODIUM CITRATE, UNSPECIFIED FORM (UNII: 1Q73Q2JULR)	
SUCRALOSE (UNII: 96K6UQ3ZD4)	

Product Characteristics

Color		Score	
Shape		Size	
Flavor	STRAWBERRY (Strawberry Banana)	Imprint Code	
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:63102-130-16	473 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	02/12/2026	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M012	02/12/2026	

Labeler - ACTIPHARMA, LLC (079340948)

Revised: 1/2026

ACTIPHARMA, LLC