DECONEX DMX- dextromethorphan hydrobromide, guaifenesin and phenylephrine hydrochloride tablet
Poly Pharmaceuticals

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.

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DECONEX DMX TABLETS

Drug Facts
Active ingredients (in each tablet)
Dextromethorphan HBr 15 mg
Guaifenesin 380 mg
Phenylephrine HCl 10 mg

Purpose
Cough Suppressant
Expectorant
Nasal Decongestant

Uses
Temporarily relieves these symptoms due to the common cold, hay fever (allergic rhinitis) or other upper respiratory allergies:

- cough due to minor throat and bronchial irritation
- helps loosen phlegm (mucus) and thin bronchial secretions to drain bronchial tubes and make coughs more productive
- nasal congestion
- reduces swelling of nasal passages

Warnings
Do not exceed recommended dosage.

Do not use this product
- 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.

Ask a doctor before use if you have
- a cough that lasts or is chronic such as occurs with smoking, asthma, chronic bronchitis, or emphysema
- a cough that occurs with too much phlegm (mucus)
- heart disease
- high blood pressure
- thyroid disease
- diabetes
- trouble urinating due to an enlarged prostate gland

Stop use and ask a doctor if
- nervousness, dizziness, or sleeplessness occur
- cough or nasal congestion persists for more than 1 week, tends to recur, or is accompanied by a fever, rash, or persistent headache. A persistent cough may be a sign of a serious condition.
- new symptoms occur

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.

Directions
Do not exceed recommended dosage.

| Adults and children 12 years of age and over: | 1 tablet every 4 hours, not to exceed 6 tablets in 24 hours |
| Children 6 to under 12 years of age: | 1/2 tablet every 4 hours, not to exceed 3 tablets in 24 hours |
| Children under 6 years of age: | Consult a physician. |

Other information
Store at 15°- 30° C (59°- 86° F).

Supplied in a tight, light-resistant container with a child-resistant cap.

Deconex DMX Tablets are orange, oblong, capsule-shaped, scored tablets, debossed "POLY" bisect "730" on one side and blank on the other side.

Inactive ingredients
Colloidal Silicon Dioxide, Croscarmellose Sodium, D&C Yellow #10 Aluminum Lake, FD&C Yellow #6 Aluminum Lake, Hypromellose, Maltodextrin, Povidone, Silicified Microcrystalline Cellulose, and Stearic Acid.

Questions? Comments?
Call 1-800-882-1041.

Manufactured for:
Poly Pharmaceuticals
Quitman, MS  39355
1(800) 882-1041

Rev: 12/11

Product Packaging
The packaging below represents the labeling currently used.
Principal display panel and side panel for 60 tablets label:

NDC 50991-730-60

DECONEX DMX
TABLETS

COUGH SUPPRESSANT · EXPECTORANT
NASAL DECONGESTANT

Each tablet contains:
Dextromethorphan HBr......................15 mg
Guaifenesin..................................380 mg
Phenylephrine HCl........................10 mg

Tamper evident by foil seal under cap.
Do not use if foil seal is broken or missing.

Distributed by:
Poly Pharmaceuticals
Quitman, MS  39355

60 tablets

Rev. 12/11
DECONEX DMX
dextromethorphan hydrobromide, guaifenesin, phenylephrine hydrochloride tablet

**Product Information**

<table>
<thead>
<tr>
<th>Product Type</th>
<th>HUMAN OTC DRUG</th>
<th>Item Code (Source)</th>
<th>NDC:50991-730</th>
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<tbody>
<tr>
<td>Route of Administration</td>
<td>ORAL</td>
<td>DEA Schedule</td>
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**Active Ingredient/Active Moiety**

<table>
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<tr>
<th>Ingredient Name</th>
<th>Basis of Strength</th>
<th>Strength</th>
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<tbody>
<tr>
<td>Dextromethorphan Hydrobromide (UNII: D2RT19KHYH) (Dextromethorphan - UNII:7355X3ROTS)</td>
<td>Dextromethorphan Hydrobromide</td>
<td>15 mg</td>
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<tr>
<td>Guaifenesin (UNII: 495W7451VQ) (Guaifenesin - UNII:495W7451VQ)</td>
<td>Guaifenesin</td>
<td>380 mg</td>
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<tr>
<td>Phenylephrine Hydrochloride (UNII: 041A59TNSJ) (Phenylephrine - UNII:1WS297W6MV)</td>
<td>Phenylephrine Hydrochloride</td>
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### Inactive Ingredients

<table>
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<th>Strength</th>
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<tr>
<td>Silicon Dioxide (UNII: ETJ7Z6XBU4)</td>
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<tr>
<td>Croscarmellose Sodium (UNII: M28OL1HH48)</td>
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<td>Hypromelloses (UNII: 3NXW29V3WO)</td>
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<td>Maltodextrin (UNII: 7CVR7L4A2D)</td>
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<tr>
<td>Povidone (UNII: FZ989GH94E)</td>
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<tr>
<td>Cellulose, Microcrystalline (UNII: OP1R32D61U)</td>
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<td>Stearic Acid (UNII: 4ELV7Z65AP)</td>
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### Product Characteristics

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<tr>
<th>Color</th>
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<tr>
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<tr>
<td>Flavor</td>
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<td>Imprint Code</td>
<td>POLY;730</td>
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<tr>
<td>Contains</td>
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### Packaging

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<tr>
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<th>Item Code</th>
<th>Package Description</th>
<th>Marketing Start Date</th>
<th>Marketing End Date</th>
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<tr>
<td>1</td>
<td>NDC:50991-730-60</td>
<td>60 in 1 BOTTLE</td>
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### Marketing Information

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<th>Application Number or Monograph Citation</th>
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<th>Marketing End Date</th>
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<tr>
<td>OTC monograph final</td>
<td>part341</td>
<td>02/17/2012</td>
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### Labeler

- Poly Pharmaceuticals (198449894)

### Registrant

- Great Southern Laboratories (056139553)

### Establishment

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<tr>
<th>Name</th>
<th>Address</th>
<th>ID/FEI</th>
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<tr>
<td>Great Southern Laboratories</td>
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<td>056139553</td>
<td>manufacture</td>
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Revised: 2/2012 Poly Pharmaceuticals