**ADULT LONG LASTING COUGH RELIEF- dextromethorphan hbr capsule, liquid filled**
**Shield Pharmaceuticals Corp**

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**Adult Long Lasting Cough Relief**

***Drug Facts***

***Active ingredient (in each softgel)***

Dextromethorphan HBr, USP 15 mg

***Purpose***

Cough suppressant

***Uses***

temporarily relieves cough due to minor throat and bronchial irritation as may occur with a cold.

***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.

**Ask a doctor before use if you have**
• a cough that occurs with too much phlegm (mucus)
• a cough that lasts or is chronic as occurs with smoking, asthma, or emphysema

**Stop use and ask a doctor if**cough lasts for more than 7 days, comes back, or is accompanied by fever, rash, or persistent headache. 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***

do not take more than 8 softgels in any 24-hour period
this adult product is not intended for use in children under 12 years of age

| | |
|---|---|
| adults and children 12 years and over | take 2 softgels every 6 to 8 hours, as needed |

| | |
|---|---|
| children under 12 years | do not use |

### *Other information*

- store at controlled room temperature 20-25ºC (68-77ºF)
- avoid excessive heat above 40°C (104°F)
- protect from light

### *Questions or comments?*

**1-800-373-6981** (toll-free)

### *Inactive ingredients*

FD&C blue no. 1, FD&C red no. 40, FD&C yellow 6, gelatin, glycerin, polyethylene glycol 400, povidone (pvp k 30), propylene glycol, purified water, sorbitan, sorbitol solution*.
*may contain this inactive ingredient

Distributed by:
**Shield Pharmaceuticals Corp.**
Ronkonkoma, NY 11779

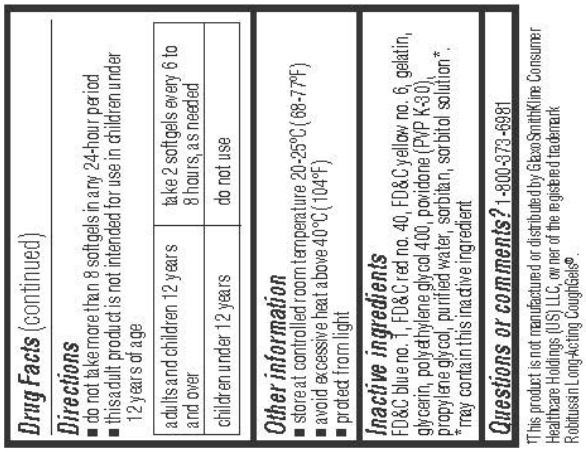
### PRINCIPAL DISPLAY PANEL

ValuRx

Compare to the active ingredient in Robitussin® Long-Acting Coughgels®†

Up to 8 Hr. Relief

Adult Long-Lasting
Cough Relief
Dextromethorphan HBr
(Cough Suppressant)



**Drug Facts** (continued)

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- do not take more than 8 softgels in any 24-hour period
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- protect from light

**Inactive ingredients**
FD&C blue no. 1, FD&C red no. 40, FD&C yellow no. 6, gelatin, glycerin, polyethylene glycol 400, povidone (PVP K-30), propylene glycol, purified water, sorbitan, sorbitol solution*.
*may contain this inactive ingredient

**Questions or comments?** 1-800-879-6981

†This product is not manufactured or distributed by GlaxoSmithKline Consumer Healthcare Holdings (US) LLC, owner of the registered trademark Robitussin Long-Acting CoughGels®.

# ADULT LONG LASTING COUGH RELIEF

dextromethorphan hbr capsule, liquid filled

## Product Information

| | | | |
|---|---|---|---|
| **Product Type** | HUMAN OTC DRUG | **Item Code (Source)** | NDC:83059-0116 |
| **Route of Administration** | ORAL | | |

## Active Ingredient/Active Moiety

| Ingredient Name | Basis of Strength | Strength |
|---|---|---|
| **DEXTROMETHORPHAN HYDROBROMIDE** (UNII: 9D2RTI9KYH) (DEXTROMETHORPHAN - UNII:7355X3ROTS) | DEXTROMETHORPHAN HYDROBROMIDE | 15 mg |

## Inactive Ingredients

| Ingredient Name | Strength |
|---|---|
| **FD&C BLUE NO. 1** (UNII: H3R47K3TBD) | |
| **FD&C RED NO. 40** (UNII: WZB9127XOA) | |
| **FD&C YELLOW NO. 6** (UNII: H77VEI93A8) | |
| **GELATIN** (UNII: 2G86QN327L) | |
| **GLYCERIN** (UNII: PDC6A3C0OX) | |
| **POLYETHYLENE GLYCOL 400** (UNII: B697894SGQ) | |
| **POVIDONE** (UNII: FZ989GH94E) | |
| **PROPYLENE GLYCOL** (UNII: 6DC9Q167V3) | |
| **SORBITAN** (UNII: 6O92ICV9RU) | |
| **SORBITOL SOLUTION** (UNII: 8KW3E207O2) | |
| **WATER** (UNII: 059QF0KO0R) | |

## Product Characteristics

| | | | |
|---|---|---|---|
| **Color** | pink | **Score** | no score |
| **Shape** | CAPSULE | **Size** | 10mm |
| **Flavor** | | **Imprint Code** | DX |
| **Contains** | | | |

| Packaging | | | | |
|---|---|---|---|---|
| # | Item Code | Package Description | Marketing Start Date | Marketing End Date |
| 1 | NDC:83059-0116-6 | 60 in 1 PACKAGE; Type 0: Not a Combination Product | 12/04/2027 | |

## Marketing Information

| Marketing Category | Application Number or Monograph Citation | Marketing Start Date | Marketing End Date |
|---|---|---|---|
| OTC Monograph Drug | M012 | 12/04/2025 | |

## Labeler - Shield Pharmaceuticals Corp (118724924)

Revised: 12/2025 Shield Pharmaceuticals Corp