### MUCUS RELIEF DM MAX MAXIMUM STRENGTH- dextromethorphan hbr, guaifenesin liquid TOP CARE (Topco Associates LLC)

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## **Drug Facts**

## Active ingredients (in each 20 mL)

Dextromethorphan HBr 20 mg

Guaifenesin 400 mg

## Purposes

Cough suppressant

Expectorant

## Uses

- helps loosen phlegm (mucus) and thin bronchial secretions to rid the bronchial passageways of botherosme mucus and make coughs more productive
- temporarily relieves:
  - cough due to minor throat and bronchial irritation as may occur with the common cold or inhaled irritants
  - the intensity of coughing
  - 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.

## Ask a doctor before use if you have

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

## When using this product,

## do not use more than directed

## Stop use and ask a doctor if

cough lasts more than 7 days, 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 (1800-222-1222) right away.

## Directions

- do not take more than 6 doses in any 24-hour period
- measure only with dosing cup provided. Do not use any other dosing device.
- keep dosing cup with products
- mL = milliliter
- dose as follows or as directed by a doctor
- adults and children 12 years of age and older: 20 mL every 4 hours
- children under 12 years of age: do not use

## Other information

- each 20 mL contains: sodium 20 mg
- store between 20-25°C (68-77°F). Do not refrigerate

## Inactive ingredients

anhydrous citric acid, disodium EDTA, FD&C red #40, flavor, glycerin, propyl gallate, propylene glycol, purified water, sodium benzoate, sodium citrate, sorbitol, sucralose, xanthan gum

## **Questions or comments?**

call toll free 1-888-423-0139

## **Principal Display Panel**

COMPARE TO MAXIMUM STRENGTH MUCINEX® FAST-MAX® DM MAX ACTIVE INGREDIENTS\*

MAXIMUM STRENGTH

## **Mucus Relief DM Max**

DEXTROMETHORPHAN HBr 20 mg COUGH SUPPRESSANT

GUAIFENESIN 400 mg EXPECTORANT

Controls Cough

- Relieves Chest Congestion
- Thins & Loosens Mucus
- 4-Hour Dosing

FOR AGES 12 +

FL OZ (mL)

\*This product is not manufactured or distributed by Reckitt Benckiser, distributor of Maximum Strength Mucinex® Fast-Max® DM Max.

# TAMPER EVIDENT: DO NOT USE IF PRINTED SAFETY SEAL AROUND OR UNDER CAP IS BROKEN OR MISSING.

DISTRIBUTED BY TOPCO ASSOCIATES LLC

ELK GROVE VILLAGE, IL 60007

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www.topcarebrand.com

## Package Label



NDC 36800-605-06

COMPARE TO MAXIMUM STRENGTH MUCINEX® FAST-MAX® DM MAX ACTIVE INGREDIENTS\*

# MAXIMUM STRENGTH Mucus Relief DM Max

DEXTROMETHORPHAN HBr 20 mg COUGH SUPPRESSANT GUAIFENESIN 400 mg EXPECTORANT

- Controls Cough
- Relieves Chest Congestion
- Thins & Loosens Mucus
- 4-Hour Dosing

PLD-C409A LB005239

## 6 FL 0Z (177 mL) FOR AGES 12+

### Drug Facts (continued)

### 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

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

### When using this product, do not use more than directed.

Stop use and ask a doctor if cough lasts more than 7 days, comes back, or occurs with fever, rash, or headache that lasts. These could be signs of a serious condition.

### TAMPER EVIDENT: DO NOT USE IF PRINTED SAFETY SEAL AROUND OR UNDER CAP IS BROKEN OR MISSING.

This product is not manufactured or distributed by Reckitt Benckiser, distributor of Maximum Strength Mucinex® Fast-Max® DM Max.





DISTRIBUTED BY TOPCO ASSOCIATES LLC ELK GROVE VILLAGE, IL 60007 ©TOPCO PLVA0521 QUESTIONS? 1-888-423-0139 topcare@topco.com www.topcarebrand.com



Scan here for more information or call 1-888-423-0139

PEEL CORNER FOR DRUG FACTS

## Drug Facts (continued)

**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 (1-800-222-1222) right away.

### Directions

- do not take more than 6 doses in any 24-hour period
- measure only with dosing cup provided.
   Do not use any other dosing device.
- keep dosing cup with product
   mL = milliliter
- dose as follows or as directed by a doctor
- adults and children 12 years of age and older: 20 mL every 4 hours
- children under 12 years of age: do not use

### Other information

- each 20 mL contains: sodium 20 mg
- store between 20-25°C (68-77°F). Do not refrigerate.

PEEL CORNER FOR MORE DRUG FACTS

Drug Facts

### Active ingredients Purposes (in each 20 mL) Dextromethorphan HBr 20 mg.....

.....Cough suppressant Guaifenesin 400 mg.....Expectorant

### Uses

- helps loosen phlegm (mucus) and thin bronchial secretions to rid the bronchial passageways of bothersome mucus and make coughs more productive
- temporarily relieves
- cough due to minor throat and bronchial irritation as may occur with the common cold or inhaled irritants
   the intensity of coughing
- the impulse to cough to help you get to sleep

## Drug Facts (continued)

### Inactive ingredients

citric acid, disodium EDTA, FD&C red #40, flavor, glycerin, propyl gallate, propylene glycol, purified water, sodium benzoate, sodium citrate, sorbitol, sucralose, xanthan gum

Questions or comments? Call toll free 1-888-423-0139

**TOPCARE HEALTH Mucus Relief DM Maximum Strength** 

# MUCUS RELIEF DM MAX MAXIMUM STRENGTH

dextromethorphan hbr, guaifenesin liquid

| Information         Product Type       HUMAN OTC DRUG       NDC:36800-605         Route of Administration       ORAL       NDC:36800-605         Active Ingredient/Active Moiety       DexTROMETHORPHAN HYDROBROMIDE (UNII: 902RT19KYH)       DexTROMETHORPHAN 20 mg in 20 mL         DEXTROMETHORPHAN HYDROBROMIDE (UNII: 902RT19KYH)       DEXTROMETHORPHAN 20 mg in 20 mL         GUAIFENESIN (UNII: 495W7451VQ) (GUAIFENESIN - UNII:495W7451VQ)       GUAIFENESIN (UNII: 495W7451VQ) (GUAIFENESIN - UNII:495W7451VQ)       GUAIFENESIN (UNII: 7474703751)         Somuto Ciftaic Actio (UNII: 8741703PSL)       Strength         Advise model (UNII: 60450N7792)       Somuto Enter CACID (UNII: 8741703PSL)       Somuto Ciftaic Actio (UNII: 60450N7792)       Somuto Ciftaic Actio (UNII: 8741703PSL)       Somuto Ciftaic Actio (UNII: 8741703PSL)       Somuto Ciftaic Actio (UNII: 8045NN792)       Somuto Ciftaic Actio (UNII: 8045NN792)       Somuto Ciftaic Actio (UNII: 8741703PSL)       Somuto Ciftaic Actio (UNII: 80450N278)       Somuto Ciftaic Actio (UNII: 80450N278)       Somuto Ciftaic Actio (UNII: 80250N278)       Somuto Ciftaic Actio (UNII: 80450N26)       Somuto Ciftaic Actio (UNII: 80250N278)       Somuto Ciftaic Actio (UNII: 80250N278) <th colspa<="" th=""><th></th><th></th><th></th><th></th><th></th><th></th><th></th></th>  | <th></th> <th></th> <th></th> <th></th> <th></th> <th></th> <th></th> |                         |                            |         |              |               |     |  |
|--|---|-------------------------|----------------------------|---------|--------------|---------------|-----|--|
| Route of Administration       ORAL         Active Ingredient/Active Moiety       Basis of Strength       Strength         Ingredient Name       Basis of Strength       Strength         DEXTROMETHORPHAN HYDROBROMIDE (UNII: 902RTI9KYH)       DEXTROMETHORPHAN 20 mg       in 20 mL         GUAIFENESIN (UNII: 495W7451VQ) (GUAIFENESIN - UNII:495W7451VQ)       GUAIFENESIN       400 mg         Inactive Ingredients       Ingredient Name       Strength         Inactive Ingredients       Ingredient Name       Strength         Inactive Ingredients       Ingredient Name       Strength         ANHYDROUS CITRIC ACID (UNII: 8741703PSL)       EDETATE DISODIUM (UNII: 7FL091C86K)       Fernethin         PROPYL GALLATE (UNII: 012302)ULR)       Sobilum BENZOATE (UNII: 012302)ULR)       Sobilum BENZOATE (UNII: 012302)ULR)       Sobilum SenzoATE (UNII: 012302)ULR)         Sobilum CITRATE (UNII: 012302)ULR)       Sobilum CITRATE (UNII: 107302)ULR)       Sobilum CITRATE (UNII: 107302)ULR)       Sobilum CITRATE (UNII: 107302)ULR)         Sobilum CITRATE (UNII: 107302)ULR)       Sobilum CITRATE (UNII: 107302)ULR)       Sobilum CITRATE (UNII: 107302)ULR)       Marketing Start         PROPYLENE GLYCOL (UNII: 10020167V3)       Marketing Start       Marketing End         Succasion       Quarketing Information       Marketing Start       Marketing End       Date  | Product Info  | rmation                 |                            |         |              |               |     |  |
| Active Ingredient/Active Moiety<br>Ingredient Name<br>Basis of Strength<br>Strengt<br>DEXTROMETHORPHAN - UNII:7355X3ROTS)<br>DEXTROMETHORPHAN - UNII:7355X3ROTS)<br>GUAIFENESIN (UNII: 495W7451VQ) (GUAIFENESIN - UNII:495W7451VQ)<br>GUAIFENESIN (UNII: 7FL91C66K)<br>PROPYL GALTATE (UNII: 8045W7V92)<br>SODIUM BENZOATE (UNII: 01245FE5EU)<br>SODIUM CITRATE (UNII: 0123Q2)ULR)<br>SORBITOL (UNII: 506T60A25R)<br>PROC NED A0. 40 (UNII: WZ B9127X0A)<br>GLYCERIN (UNII: 9DC6A3C00X)<br>PROPYLENE GLYCOL (UNII: 60C90167V3)<br>WATER (UNII: 90590FKOOR)<br>SUCRALOSE (UNII: 9690FKOOR)<br>SUCRALOSE (UNII: 9690FKOOR)<br>SUCRALOSE (UNII: 97T1LE N LASTIC: Type 0: Not a<br>1 NDC:36800-<br>1 NDC:36800-<br>1 NDC:36800-<br>1 NDC:36800-<br>Combination Product<br>Marketing Information<br>Marketing Information<br>Marketing Application Number or Monograph<br>Marketing Start Marketing Ent<br>Date   | Product Type  |                         | HUMAN OTC DRUG Item Cod    |         | de (Source)  | NDC:36800-605 |     |  |
| Ingredient Name         Basis of Strength         Strength           DEXTROMETHORPHAN HYDROBROMIDE (UNII: 9D2RT19KYH)         DEXTROMETHORPHAN HYDROBROMIDE (UNII: 9D2RT19KYH)         DEXTROMETHORPHAN 20 mg in 20 mL           GUAIFENESIN (UNII: 495W7451VQ) (GUAIFENESIN - UNII:495W7451VQ)         GUAIFENESIN         400 mg in 20 mL           Inactive Ingredients         Ingredient Name         Strength           ANHYDROUS CITRIC ACID (UNII: XF417D3PSL)         EDETATE DISODIUM (UNII: 7FL991C86K)         Foregreen and the strength           PROPYL GALLATE (UNII: 0245FESEU)         Sobalum Citrate (UNII: 0245FESEU)         Sobalum Citrate (UNII: 0245FESEU)           SODIUM CITRATE (UNII: 0245FESEU)         Sobalum Citrate (UNII: 0245FESEU)         Sobalum Citrate (UNII: 0245FESEU)           SOBUM CITRATE (UNII: 0245FESEU)         Sobalum Citrate (UNII: 0245FESEU)         Sobalum Citrate (UNII: 0245FESEU)           SOBUM CITRATE (UNII: 0245FESEU)         Sobalum Citrate (UNII: 0245FESEU)         Sobalum Citrate (UNII: 0245FESEU)           SOBUM CITRATE (UNII: 1073Q2JULR)         Sobalum Citrate (UNII: 0545FESEU)         Sobalum Citrate (UNII: 0545FESEU)           SOBUM CITRATE (UNIII: 96460Q32 DA)         SUCRALOSE (UNII: 96660Q32 DA)         SUCRALOSE (UNII: 96660Q32 DA)           SUCRALOSE (UNII: 96660Q32 DA)         SUCRALOSE (UNII: 96660Q32 DA)         SUCRALOSE (UNII: 96660Q32 DA)           1         NDC:36800-         177 mL in 1 BOTTLE, PLASTIC; Type   | Route of Admin  | Administration ORAL     |                            |         |              |               |     |  |
| Ingredient Name         Basis of Strength         Strength           DEXTROMETHORPHAN HYDROBROMIDE (UNII: 9D2RT19KYH)         DEXTROMETHORPHAN HYDROBROMIDE (UNII: 9D2RT19KYH)         DEXTROMETHORPHAN 20 mg in 20 mL           GUAIFENESIN (UNII: 495W7451VQ) (GUAIFENESIN - UNII:495W7451VQ)         GUAIFENESIN         400 mg in 20 mL           Inactive Ingredients         Ingredient Name         Strength           ANHYDROUS CITRIC ACID (UNII: XF417D3PSL)         EDETATE DISODIUM (UNII: 7FL991C86K)         Foregreen and the strength           PROPYL GALLATE (UNII: 0245FESEU)         Sobalum Citrate (UNII: 0245FESEU)         Sobalum Citrate (UNII: 0245FESEU)           SODIUM CITRATE (UNII: 0245FESEU)         Sobalum Citrate (UNII: 0245FESEU)         Sobalum Citrate (UNII: 0245FESEU)           SOBUM CITRATE (UNII: 0245FESEU)         Sobalum Citrate (UNII: 0245FESEU)         Sobalum Citrate (UNII: 0245FESEU)           SOBUM CITRATE (UNII: 0245FESEU)         Sobalum Citrate (UNII: 0245FESEU)         Sobalum Citrate (UNII: 0245FESEU)           SOBUM CITRATE (UNII: 1073Q2JULR)         Sobalum Citrate (UNII: 0545FESEU)         Sobalum Citrate (UNII: 0545FESEU)           SOBUM CITRATE (UNIII: 96460Q32 DA)         SUCRALOSE (UNII: 96660Q32 DA)         SUCRALOSE (UNII: 96660Q32 DA)           SUCRALOSE (UNII: 96660Q32 DA)         SUCRALOSE (UNII: 96660Q32 DA)         SUCRALOSE (UNII: 96660Q32 DA)           1         NDC:36800-         177 mL in 1 BOTTLE, PLASTIC; Type   |   |                         |                            |         |              |               |     |  |
| DEXTROMETHORPHAN HYDROBROMIDE (UNII: 9D2RTI9KYH)<br>(DEXTROMETHORPHAN - UNII: 7355X3ROTS)       DEXTROMETHORPHAN - UNII: 495W7451VQ)       20 mg<br>in 20 mL<br>400 mg<br>in 20 mL         GUAIFENESIN (UNII: 495W7451VQ) (GUAIFENESIN - UNII: 495W7451VQ)       GUAIFENESIN       400 mg<br>in 20 mL         Inactive Ingredients       Strength         ANHYDROUS CITRIC ACID (UNII: XF417D3P5L)       E         EDETATE DISODIUM (UNII: 7FL091C86K)       F         PROPYL GALATE (UNII: 8D45NN7V92)       SODIUM ENZOATE (UNII: 0)245FE5EU)         SODIUM EINZOATE (UNII: 0)245FE5EU)       SODIUM CITRATE (UNII: 0)245FE5EU)         SODBITOL (UNII: 506T60A25R)       F         FDAC RED NO. 40 (UNII: WZ B9127XOA)       GLYCERIN (UNII: 906K0QR)         SUCRALOSE (UNII: 809Q167V3)       F         WATER (UNII: 059Q760K0R)       SUCRALOSE (UNII: 96K6UQ3ZD4)         SUCRALOSE (UNII: 96K6UQ3ZD4)       SUCRALOSE (UNII: 96K6UQ3ZD4)         XANTHAN GUM (UNII: TTV12P4NEE)       SUCRALOSE (UNII: 96K6UQ3ZD4)         1       NDC:36800-       177 mL in 1 BOTTLE, PLASTIC; Type 0: Not a       04/30/2016         1       NDC:36800-       177 mL in 1 BOTTLE, PLASTIC; Type 0: Not a       04/30/2016         1       NDC:36800-       177 mL in 1 BOTTLE, PLASTIC; Type 0: Not a       04/30/2016   | Active Ingred   | ient/Active             | Moiety                     |         |              |               |     |  |
| DEXTROMETHORPHAN HYDROBROMIDE (UNII: 902RT19KYH)       DEXTROMETHORPHAN - UNII: 7355X3R0TS)       20 mg<br>in 20 mL<br>HYDROBROMIDE       20 mg<br>in 20 mL         GUAIFENESIN (UNII: 495W7451VQ) (GUAIFENESIN - UNII:495W7451VQ)       GUAIFENESIN       400 mg<br>in 20 mL         Inactive Ingredients       Strength         ANHYDROUS CITRIC ACID (UNII: XF417D3P5L)       E         EDETATE DISODIUM (UNII: 7FL091C86K)       Strength         PROPYL GALATE (UNII: 8045NN7V92)       SODIUM BENZOATE (UNII: 01245FE5EU)         SODIUM TEATE (UNII: 01245FE5EU)       SORBITOL (UNII: 07302)ULR)         SORBITOL (UNII: 506T60A25R)       FDAC RED NO. 40 (UNII: WZ B9127XOA)         GLYCERIN (UNII: 9506GR0R)       SUCRALOSE (UNII: 96K6UQ3ZD4)         SUCRALOSE (UNII: 96K6UQ3ZD4)       SUCRALOSE (UNII: 96K6UQ3ZD4)         XANTHAN GUM (UNII: TTV12P4NEE)       Marketing Start<br>Combination Product  |   | Basis of Str            | ength Strei                | ngt     |              |               |     |  |
| Inactive Ingredients<br>Ingredient Name Strength<br>ANHYDROUS CITRIC ACID (UNII: XF417D3P5L)<br>EDETATE DISODIUM (UNII: 7FLD91C86K)<br>PROPYL GALATE (UNII: 8D4SNN7V92)<br>SODIUM CITRATE (UNII: 8D4SNN7V92)<br>SODIUM CITRATE (UNII: 10/342FE5EU)<br>SODIUM CITRATE (UNII: 10/342FE5EU)<br>SOCRALOSE (UNII: 9050F0K00R)<br>SUCRALOSE (UNII: 9050F0K00R)<br>SUCRALOSE (UNII: 9050F0K00R)<br>SUCRALOSE (UNII: 96K6U03Z D4)<br>XANTHAN GUM (UNII: TTV12P4NEE)<br>Packaging<br># Item Code Package Description Marketing Start Marketing End<br>Date<br>Marketing Information<br>Marketing Application Number or Monograph Marketing Start Marketing End<br>Date  |   | DEXTROMETHORP           | HAN 20 mg                  |         |              |               |     |  |
| Ingredient Name       Strength         ANHYDROUS CITRIC ACID (UNII: XF417D3PSL)       EDETATE DISODIUM (UNII: XF417D3PSL)       EDETATE DISODIUM (UNII: XF417D3PSL)         EDETATE DISODIUM (UNII: RFL091C86K)       PROPYL GALLATE (UNII: 0245FE5EU)       SOUDUM BENZOATE (UNII: 0245FE5EU)         SODIUM BENZOATE (UNII: 073Q2JULR)       SOBITOL (UNII: SOET60A25R)       SOBITOL (UNII: SOET60A25R)         FD&C RED NO. 40 (UNII: WZB9127XOA)       SOE  | GUAIFENESIN (UN   | II: 495W7451VQ          | ) (GUAIFENESIN - UNII:495W | 7451VQ) | GUAIFENES IN |               |     |  |
| Ingredient NameStrengthANHYDROUS CITRIC ACID (UNII: XF417D3P5L)EDETATE DISODIUM (UNII: XF417D3P5L)EDETATE DISODIUM (UNII: RFL091C86K)PROPYL GALLATE (UNII: 0245FE5EU)SODIUM BENZOATE (UNII: 0173Q2JULR)SORBITOL (UNII: S06T60A25R)FD&C RED NO. 40 (UNII: WZB9127XOA)GLYCERIN (UNII: DC633C00X)PROPYLENE GLYCOL (UNII: 6DC90167V3)WATER (UNII: 059QF0K00R)SUCRALOSE (UNII: 96K6UQ3ZD4)XANTHAN GUM (UNII: TTV12P4NEE)Marketing Start DateMarketing InformationMarketing InformationMarketing Start DateMarketing CategoryApplication Number or Monograph DateMarketing Start Date  |   |                         |                            |         |              |               |     |  |
| ANHYDROUS CITRIC ACID (UNII: XF417D3PSL)<br>EDETATE DISODIUM (UNII: 7FL091C86K)<br>PROPYL GALLATE (UNII: 8D45NN7V92)<br>SODIUM BENZOATE (UNII: 0J245FE5EU)<br>SODIUM CITRATE (UNII: 0J245FE5EU)<br>SODIUM CITRATE (UNII: 0J245FE5EU)<br>SODIUM CITRATE (UNII: 0J245FE5EU)<br>SORBITOL (UNII: 50°T60A25R)<br>FD&C RED NO. 40 (UNII: WZ B9127X0A)<br>GLYCERIN (UNII: 90°C6A3C00X)<br>PROPYLENE GLYCOL (UNII: 6DC90167V3)<br>WATER (UNII: 05°QF0KO0R)<br>SUCRALOSE (UNII: 96K6UQ3ZD4)<br>XANTHAN GUM (UNII: TTV12P4NEE)<br>PACKaging<br># tem Code Package Description Marketing Start Date Date<br>Marketing Combination Product<br>Marketing Combination Number or Monograph Marketing Start Date Marketing End<br>Category Application Number or Monograph Marketing Start Date Marketing End<br>Date Marketing Category Citation Number or Monograph Citation Marketing Start Date Marketing End<br>Date Marketing Category Citation Number or Monograph Marketing Start Date Marketing End<br>Date Marketing Category Citation Number or Monograph Citation Marketing Start Marketing End<br>Date Marketing Category Citation Marketing Start Marketing Category Citation Marketing Start Marketing Category Citation Marketing Category Citation Category Citation Marketing Category Citation Category Category Citation Category Cita | Inactive Ingre  | edients                 |                            |         |              |               |     |  |
| EDETATE DISODUM (UNII: 7FLD91C86K)       Image: Content of the state  |   | Strength                |                            |         |              |               |     |  |
| PROPYL GALLATE (UNII: 8D4SNN7V92)       Solum BENZOATE (UNII: 0J245FE5EU)         SODIUM CITRATE (UNII: 1Q73Q2JULR)       SORBITOL (UNII: 506T60A25R)         FD&C RED NO. 40 (UNII: WZB9127X0A)       GLYCERIN (UNII: PDC6A3C00X)         FD&C RED NO. 40 (UNII: WZB9127X0A)       GLYCERIN (UNII: PDC6A3C00X)         PROPYLENE GLYCOL (UNII: 6DC9Q167V3)       WATER (UNII: 059QF0K00R)         SUCRALOSE (UNII: 96K6UQ3Z D4)       XANTHAN GUM (UNII: TTV12P4NEE)         Packaging       Marketing Start Date         #       Item Code       Package Description Date         1       NDC:36800- 0000000000000000000000000000000000  | ANHYDROUS CITP  | RIC ACID (UNII:         | XF417D3PSL)                |         |              |               |     |  |
| SOLUM BENZOATE (UNII: 0/245FESEU)       Image: Contract (UNII: 10/30/2)ULR)         SOLUM CITRATE (UNII: 10/30/2)ULR)       Image: Contract (UNII: 506T60A25R)         FD&C RED NO. 40 (UNII: WZ B9127X0A)       Image: Contract (UNII: PDC6A3C00X)         GLYCERIN (UNII: PDC6A3C00X)       Image: Contract (UNII: 60590167V3)         WATER (UNII: 0590F0K00R)       Image: Contract (UNII: 96K60032D4)         SUCRALOSE (UNII: 96K60032D4)       Image: Contract (UNII: 707204)         XANTHAN GUM (UNII: TTV12P4NEE)       Image: Contract (UNII: 707204)         Marketing Code (UNII: 11 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product         NDC:36800- 605-06       1/7 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product       04/30/2016         Marketing Category         Marketing Start (Category)       Application Number or Monograph (Citation)       Marketing Start (Date (UNII: 707204))  | EDETATE DISODI  | UM (UNII: 7FLD9         | 1C86K)                     |         |              |               |     |  |
| SOLUM CITRATE (UNII: 107302JULR)       Image: Control of the state of   | PROPYL GALLATE  | (UNII: 8D4SNN7          | V92)                       |         |              |               |     |  |
| SORBITOL (UNII: 506T60A25R)       Image: Solution of the solution of t   | SODIUM BENZOA   | <b>TE</b> (UNII: OJ2451 | E5EU)                      |         |              |               |     |  |
| FD&C RED NO. 40 (UNII: WZ B9127X0A)       Image: Constant of the state of the sta  | SODIUM CITRATE  | (UNII: 1Q73Q2J          | JLR)                       |         |              |               |     |  |
| Bit Starketing Application Number or Monograph Marketing Start Marketing Start   | SORBITOL (UNII: 5   | 06T60A25R)              |                            |         |              |               |     |  |
| Propriation       Marketing Start Combination Product       Marketing Start Combinatic Product   | FD&C RED NO. 40   | ) (UNII: WZ B912        | 7XOA)                      |         |              |               |     |  |
| WATER (UNII: 059QF0KO0R)       SUCRALOSE (UNII: 96K6UQ3ZD4)         SUCRALOSE (UNII: 96K6UQ3ZD4)       SUCRALOSE (UNII: 96K6UQ3ZD4)         XANTHAN GUM (UNII: TTV12P4NEE)       SUCRALOSE (UNII: 96K6UQ3ZD4)         Package Description       Marketing Start Date         Marketing Start Orbination Product       Marketing Start Date         Marketing Category       Application Number or Monograph Citation   | GLYCERIN (UNII: P   | DC6A3C0OX)              |                            |         |              |               |     |  |
| SUCRALOSE (UNII: 96K6UQ3ZD4)         A Marketing Code         Package Description       Marketing Start Date       Marketing End Date         Marketing Code       Package Description       Marketing Start Date       Marketing End Date         I MDC:36800-<br>605-06       177 mL in 1 BOTTLE, PLASTIC; Type 0: Not a<br>Combination Product       04/30/2016         Marketing Category       Application Number or Monograph Citation       Marketing Start Date       Marketing End Date   | PROPYLENE GLYC  | OL (UNII: 6DC9          | Q167V3)                    |         |              |               |     |  |
| Warketing Category       Application Number or Monograph Marketing Start Date       Marketing Start Date   | WATER (UNII: 0590   | QF0KO0R)                |                            |         |              |               |     |  |
| Warketing Code       Package Description       Marketing Start Date       Marketing End Date         1       NDC:36800-<br>605-06       177 mL in 1 BOTTLE, PLASTIC; Type 0: Not a<br>Combination Product       04/30/2016       04/30/2016         Marketing Information         Marketing Category       Application Number or Monograph<br>Citation       Marketing Start<br>Date       Marketing End<br>Date   | SUCRALOSE (UNII:  | 96K6UQ3ZD4)             |                            |         |              |               |     |  |
| #       Item Code       Package Description       Marketing Start Date       Marketing End Date         1       NDC:36800-<br>605-06       177 mL in 1 BOTTLE, PLASTIC; Type 0: Not a<br>Combination Product       04/30/2016       04/30/2016         Marketing Treation Product         Marketing Category         Application Number or Monograph Citation       Marketing Start Date       Marketing End Date  | XANTHAN GUM (U  | NII: TTV12P4NE          | Ξ)                         |         |              |               |     |  |
| #       Item Code       Package Description       Marketing Start Date       Marketing End Date         1       NDC:36800-<br>605-06       177 mL in 1 BOTTLE, PLASTIC; Type 0: Not a<br>Combination Product       04/30/2016       04/30/2016         Marketing Treation Product         Marketing Category         Application Number or Monograph Citation       Marketing Start Date       Marketing End Date  |   |                         |                            |         |              |               |     |  |
| **       Item Code       Package Description       Date       Date         1       NDC:36800-<br>605-06       177 mL in 1 BOTTLE, PLASTIC; Type 0: Not a<br>Combination Product       04/30/2016         Marketing Information       Marketing Start       Marketing End<br>Date         Marketing Category       Application Number or Monograph<br>Citation       Marketing Start       Marketing End<br>Date  | Packaging   |                         |                            |         |              |               |     |  |
| Image: Combination Product       04/30/2010         Marketing Information         Marketing Category       Application Number or Monograph Citation  | # Item Code   | Pa                      | ackage Description         |         |              |               | Enc |  |
| Marketing<br>CategoryApplication Number or Monograph<br>CitationMarketing Start<br>DateMarketing End<br>Date   |   |                         |                            | а       | 04/30/2016   |               |     |  |
| Marketing<br>CategoryApplication Number or Monograph<br>CitationMarketing Start<br>DateMarketing End<br>Date   |   |                         |                            |         |              |               |     |  |
| Category Citation Date Date  | Marketing   | Informat                | ion                        |         |              |               |     |  |
| OTC Monograph Drug M012 04/30/2016   |   | Applica                 |                            | Iraph   |              |               | End |  |
|  | OTC Monograph Dr  | ug M012                 |                            | (       | 04/30/2016   |               |     |  |

Labeler - TOP CARE (Topco Associates LLC) (006935977)

Revised: 10/2023