

**DAYTIME NIGHTTIME SEVERE COLD COUGH AND FLU RELIEF- acetaminophen, dextromethorphan hbr, phenylephrine hcl, diphenhydramine hcl**  
**Rite Aid Corporation**

*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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**Rite Aid Corporation Daytime Nighttime Severe Cold Cough & Flu Relief Drug Facts**

**Active ingredients (in each packet) - Daytime**

Acetaminophen 500 mg

Dextromethorphan HBr 20 mg

Phenylephrine HCl 10 mg

**Purpose**

Pain reliever/fever reducer

Cough suppressant

Nasal decongestant

**Uses**

- temporarily relieves these symptoms due to a cold:
- minor aches and pains
- minor sore throat pain
- headache
- nasal and sinus congestion
- cough due to minor throat and bronchial irritation
- temporarily reduces fever

**Warnings**

**Liver warning:** This product contains acetaminophen. Severe liver damage may occur if you take

- more than 4,000 mg of acetaminophen in 24 hours
- with other drugs containing acetaminophen
- 3 or more alcoholic drinks every day while using this product

**Allergy alert:** Acetaminophen may cause severe skin reactions. Symptoms may include:

- skin reddening
- blisters
- rash

If a skin reaction occurs, stop use and seek medical help right away.

**Sore throat warning:** If sore throat is severe, persists for more than 2 days, is accompanied or followed by fever, headache, rash, nausea, or vomiting, consult a doctor promptly.

### **Do not use**

- in a child under 12 years of age
- if you have ever had an allergic reaction to this product or any of its ingredients
- with any other drug containing acetaminophen (prescription or nonprescription). If you are not sure whether a drug contains acetaminophen, ask a doctor or pharmacist.
- 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**

- liver disease
- heart disease
- high blood pressure
- thyroid disease
- diabetes
- trouble urinating due to an enlarged prostate gland
- cough that occurs with too much phlegm (mucus)
- cough that lasts or is chronic such as occurs with smoking, asthma, or emphysema
- a sodium-restricted diet

### **Ask a doctor or pharmacist before use if you are**

taking the blood thinning drug warfarin

### **When using this product**

**do not exceed recommended dosage**

### **Stop use and ask a doctor if**

- nervousness, dizziness, or sleeplessness occurs
- fever gets worse or lasts more than 3 days
- redness or swelling is present
- new symptoms occur
- pain, cough or nasal congestion gets worse or lasts more than 7 days
- cough comes back or occurs with 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.**

**Overdose warning:** In case of overdose, get medical help or contact a Poison Control Center right away (1-800-222-1222). Prompt medical attention is critical for adults as well as for children even if you do not notice any signs or symptoms.

**Directions**

- **do not use more than directed (see overdose warning)**
- take every 4 hours, while symptoms persist. Do not take more than 6 packets in 24 hours unless directed by a doctor.

| Age  | Dose       |
|--|------------|
| adults and children 12 years of age and over | one packet |
| children under 12 years of age               | do not use |

- dissolve contents of one packet into 8 oz. hot water: sip while hot. Consume entire drink within 10-15 minutes.
- if using a microwave, add contents of one packet to 8 oz. of cool water: stir briskly before and after heating. Do not overheat.

**Other information**

- **each packet contains:** potassium 10 mg and sodium 25 mg
- **phenylketonurics:** contains phenylalanine 22 mg per packet
- store at 20-25°C (68-77°F). Protect product from heat and moisture.

**Inactive ingredients**

acesulfame potassium, anhydrous citric acid, aspartame, colloidal silicon dioxide, D&C yellow #10, FD&C blue #1, FD&C red #40, flavors, maltodextrin, pregelatinized starch, sodium citrate, sucrose, tribasic calcium phosphate

**Questions or comments?**

**1-800-719-9260**

**Active ingredients (in each packet)**

Acetaminophen 650 mg

Diphenhydramine HCl 25 mg

Phenylephrine HCl 10 mg

## **Purposes**

Pain reliever/fever reducer

Antihistamine/cough suppressant

Nasal decongestant

## **Uses**

- temporarily relieves these symptoms due to a cold:
- minor aches and pains
- minor sore throat pain
- headache
- nasal and sinus congestion
- runny nose
- sneezing
- itchy nose or throat
- itchy, watery eyes due to hay fever
- cough due to minor throat and bronchial irritation
- temporarily reduces fever

## **Warnings**

**Liver warning:** This product contains acetaminophen. Severe liver damage may occur if you take

- more than 4,000 mg of acetaminophen in 24 hours
- with other drugs containing acetaminophen
- 3 or more alcoholic drinks every day while using this product

**Allergy alert:** Acetaminophen may cause severe skin reactions. Symptoms may include:

- skin reddening
- blisters
- rash

If a skin reaction occurs, stop use and seek medical help right away.

**Sore throat warning:** If sore throat is severe, persists for more than 2 days, is accompanied or followed by fever, headache, rash, nausea, or vomiting, consult a doctor promptly.

## **Do not use**

- in a child under 12 years of age

- if you have ever had an allergic reaction to this product or any of its ingredients
- with any other drug containing acetaminophen (prescription or nonprescription). If you are not sure whether a drug contains acetaminophen, ask a doctor or pharmacist.
- with any other product containing diphenhydramine, even one used on skin
- 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**

- liver disease
- heart disease
- high blood pressure
- thyroid disease
- diabetes
- glaucoma
- trouble urinating due to an enlarged prostate gland
- a breathing problem such as emphysema or chronic bronchitis
- cough that occurs with too much phlegm (mucus)
- cough that lasts or is chronic such as occurs with smoking, asthma, or emphysema

### **Ask a doctor or pharmacist before use if you are**

- taking sedatives or tranquilizers
- taking the blood thinning drug warfarin

### **When using this product**

- **do not exceed recommended dosage**
- avoid alcoholic drinks
- marked drowsiness may occur
- alcohol, sedatives, and tranquilizers may increase drowsiness
- be careful when driving a motor vehicle or operating machinery
- excitability may occur, especially in children

### **Stop use and ask a doctor if**

- nervousness, dizziness, or sleeplessness occurs
- fever gets worse or lasts more than 3 days
- redness or swelling is present
- new symptoms occur
- pain, cough or nasal congestion gets worse or lasts more than 7 days
- cough comes back or occurs with 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.

**Overdose warning:** In case of overdose, get medical help or contact a Poison Control Center right away (1-800-222-1222). Prompt medical attention is critical for adults as well as for children even if you do not notice any signs or symptoms.

### Directions

- **do not use more than directed (see overdose warning)**
- take every 4 hours, while symptoms persist. Do not take more than 5 packets in 24 hours unless directed by a doctor.

| Age  | Dose       |
|--|------------|
| adults and children 12 years of age and over | one packet |
| children under 12 years of age               | do not use |

- dissolve contents of one packet into 8 oz. hot water: sip while hot. Consume entire drink within 10-15 minutes.
- if using a microwave, add contents of one packet to 8 oz. of cool water: stir briskly before and after heating. Do not overheat.

### Other information

- **each packet contains:** potassium 10 mg and sodium 25 mg
- **phenylketonurics:** contains phenylalanine 13 mg per packet
- store at 20-25°C (68-77°F). Protect product from heat and moisture.

### Inactive ingredients

acesulfame potassium, anhydrous citric acid, aspartame, colloidal silicon dioxide, D&C yellow #10, FD&C blue #1, FD&C red #40, flavors, maltodextrin, pregelatinized starch, sodium citrate, sucrose, tribasic calcium phosphate

### Questions or comments?

**1-800-719-9260**

### Package/Label Principal Display Panel

COMBO PACK

Compare to the active ingredients of Theraflu® Severe Cold and Theraflu® Nighttime Severe Cold & Cough

daytime

severe cold cough & flu relief  
acetaminophen 500 mg  
dextromethorphan HBr 20 mg  
phenylephrine HCl 10 mg  
pain reliever/fever reducer  
cough suppressant &  
nasal decongestant

relieves:

nasal & sinus congestion

cough

body ache

sore throat pain

headache

fever

Green Tea & Honey Lemon flavors

6 PACKETS

nighttime

severe cold cough & flu relief

acetaminophen 650 mg

diphenhydramine HCl 25 mg

phenylephrine HCl 10 mg

pain reliever/fever reducer

antihistamine/

cough suppressant

& nasal decongestant

relieves:

nasal congestion

cough

runny nose

sneezing

body ache

sore throat pain

headache

fever

honey lemon infused with white tea flavors

6 PACKETS





# COMBO PACK

\*Compare to the active ingredients of Theraflu® Severe Cold and Theraflu® Nighttime Severe Cold & Cough

## daytime severe cold cough & flu relief

**acetaminophen 500 mg**  
**dextromethorphan HBr 20 mg**  
**phenylephrine HCl 10 mg**

pain reliever/fever reducer  
cough suppressant &  
nasal decongestant

relieves:  
nasal & sinus congestion  
cough  
body ache  
sore throat pain  
headache  
fever



Green Tea &  
Honey Lemon flavors

6 PACKETS

## nighttime severe cold cough & flu relief

**acetaminophen 650 mg**  
**diphenhydramine HCl 25 mg**  
**phenylephrine HCl 10 mg**

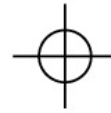
pain reliever/fever reducer  
antihistamine/  
cough suppressant  
& nasal decongestant

relieves:  
nasal congestion  
cough  
runny nose  
sneezing  
body ache  
sore throat pain  
headache  
fever



honey lemon infused  
with white tea flavors

6 PACKETS



DO NOT TAKE BOTH PRODUCTS AT THE SAME TIME OR TAKE MORE THAN 5 DOSES IN TOTAL IN ANY 24 HOUR PERIOD.

DO NOT TAKE A DOSE OF THE NIGHTTIME PRODUCT SOONER THAN 4 HOURS AFTER THE LAST DOSE OF MULTI-SYMPTOM PRODUCT UNLESS DIRECTED BY YOUR DOCTOR.

Multi-Symptom Severe Cold

### Drug Facts

| Active ingredients (in each packet) | Purposes                    |
|-------------------------------------|-----------------------------|
| Acetaminophen 500 mg.....           | Pain reliever/fever reducer |
| Dextromethorphan HBr 20 mg.....     | Cough suppressant           |
| Phenylephrine HCl 10 mg.....        | Nasal decongestant          |

**Uses** ■ temporarily relieves these symptoms due to a cold:  
■ minor aches and pains ■ minor sore throat pain ■ headache  
■ nasal and sinus congestion ■ cough due to minor throat and bronchial irritation  
■ temporarily reduces fever

**Warnings**  
**Liver warning:** This product contains acetaminophen. Severe liver damage may occur if you take  
■ more than 4,000 mg of acetaminophen in 24 hours  
■ with other drugs containing acetaminophen  
■ 3 or more alcoholic drinks every day while using this product  
**Allergy alert:** Acetaminophen may cause severe skin reactions. Symptoms may include:  
■ skin reddening ■ blisters ■ rash  
If a skin reaction occurs, stop use and seek medical help right away.  
**Sore throat warning:** If sore throat is severe, persists for more than 2 days, is accompanied or followed by fever, headache, rash, nausea, or vomiting, consult a doctor promptly.

**Do not use** ■ in a child under 12 years of age  
■ if you have ever had an allergic reaction to this product or any of its ingredients  
■ with any other drug containing acetaminophen (prescription or nonprescription), if you are not sure whether a drug contains acetaminophen, ask a doctor or pharmacist.  
■ 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** ■ liver disease  
■ heart disease ■ high blood pressure ■ thyroid disease ■ diabetes  
■ trouble urinating due to an enlarged prostate gland  
■ cough that occurs with too much phlegm (mucus)  
■ cough that lasts or is chronic such as occurs with smoking, asthma, or emphysema  
■ a sodium-restricted diet

**Ask a doctor or pharmacist before use if you are taking the blood thinning drug warfarin**

When using this product do not exceed recommended dosage



acetaminophen 650 mg  
diphenhydramine HCl 25 mg  
phenylephrine HCl 10 mg  
pain reliever/fever reducer  
antihistamine/  
cough suppressant  
& nasal decongestant

# nighttime severe cold & flu relief

Do not use if printed packets are torn or unsealed

acetaminophen 500 mg  
dextromethorphan HBr 20 mg  
phenylephrine HCl 10 mg  
pain reliever/fever reducer  
cough suppressant  
& nasal decongestant

# daytime severe cold & flu relief



### Drug Facts (continued)

#### Stop use and ask a doctor if

- nervousness, dizziness, or sleeplessness occurs
- fever gets worse or lasts more than 3 days
- redness or swelling is present
- new symptoms occur
- pain, cough or nasal congestion gets worse or lasts more than 7 days
- cough comes back or occurs with 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. Overdose warning:** In case of overdose, get medical help or contact a Poison Control Center right away (1-800-222-1222). Prompt medical attention is critical for adults as well as for children even if you do not notice any signs or symptoms.

#### Directions

- do not use more than directed (see overdose warning)
- take every 4 hours, while symptoms persist. Do not take more than 6 packets in 24 hours unless directed by a doctor.

### Drug Facts (continued)

| Age  | Dose       |
|--|------------|
| adults and children 12 years of age and over | one packet |
| children under 12 years of age               | do not use |

■ dissolve contents of one packet into 8 oz. hot water: sip while hot. Consume entire drink within 10-15 minutes.  
■ if using a microwave, add contents of one packet to 8 oz. of cool water: stir briskly before and after heating. Do not overheat.

#### Other information

- each packet contains: potassium 10 mg and sodium 25 mg
- phenylketonurics: contains phenylalanine 22 mg per packet
- store at 20-25°C (68-77°F). Protect product from heat and moisture.

**Inactive ingredients** acesulfame potassium, anhydrous citric acid, aspartame, colloidal silicon dioxide, D&C yellow #10, FD&C blue #1, FD&C red #40, flavors, maltodextrin, pregelatinized starch, sodium citrate, sucrose, tribasic calcium phosphate

**Questions or comments?** 1-800-719-9260

### Drug Facts (continued)

#### Ask a doctor or pharmacist before use if you are

- taking sedatives or tranquilizers
- taking the blood thinning drug warfarin

#### When using this product

- do not exceed recommended dosage
- avoid alcoholic drinks
- alcohol, sedatives, and tranquilizers may increase drowsiness
- be careful when driving a motor vehicle or operating machinery
- marked drowsiness may occur
- excitability may occur, especially in children

#### Stop use and ask a doctor if

- nervousness, dizziness, or sleeplessness occurs
- fever gets worse or lasts more than 3 days
- redness or swelling is present
- new symptoms occur
- pain, cough or nasal congestion gets worse or lasts more than 7 days
- cough comes back or occurs with 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. Overdose warning:** In case of overdose, get medical help or contact a Poison Control Center right away (1-800-222-1222). Prompt medical attention is critical for adults as well as for children even if you do not notice any signs or symptoms.

#### Directions

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| adults and children 12 years of age and over | one packet |
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- dissolve contents of one packet into 8 oz. hot water: sip while hot. Consume entire drink within 10-15 minutes.
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#### Other information

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- store at 20-25°C (68-77°F). Protect product from heat and moisture.

**Inactive ingredients** acesulfame potassium, anhydrous citric acid, aspartame, colloidal silicon dioxide, D&C yellow #10, FD&C blue #1, FD&C red #40, flavors, maltodextrin, pregelatinized starch, sodium citrate, sucrose, tribasic calcium phosphate

**Questions or comments?** 1-800-719-9260

\*These products are not manufactured or distributed by GSK Consumer Healthcare, distributor of Theraflu® Severe Cold and Theraflu® Nighttime Severe Cold & Cough.

### PARENTS:

Learn about teen medicine abuse

[www.StopMedicineAbuse.org](http://www.StopMedicineAbuse.org)

DISTRIBUTED BY: RITE AID  
30 HUNTER LANE  
CAMP HILL, PA 17011

MADE IN MEXICO



IF YOU'RE NOT  
SATISFIED, WE'LL  
HAPPILY REFUND  
YOUR MONEY.





# CODE AREA

1E655 83 C1

## DAYTIME NIGHTTIME SEVERE COLD COUGH AND FLU RELIEF

acetaminophen, dextromethorphan hbr, phenylephrine hcl, diphenhydramine hcl kit

### Product Information

|                     |                |                           |                |
|---------------------|----------------|---------------------------|----------------|
| <b>Product Type</b> | HUMAN OTC DRUG | <b>Item Code (Source)</b> | NDC:11822-2050 |
|---------------------|----------------|---------------------------|----------------|

### Packaging

| # | Item Code        | Package Description                              | Marketing Start Date | Marketing End Date |
|---|------------------|--|----------------------|--------------------|
| 1 | NDC:11822-2050-1 | 1 in 1 CARTON; Type 0: Not a Combination Product | 06/24/2019           |                    |

### Quantity of Parts

| Part # | Package Quantity | Total Product Quantity |
|--------|------------------|------------------------|
| Part 1 | 6 PACKET         | 6                      |
| Part 2 | 6 PACKET         | 6                      |

## Part 1 of 2

### DAYTIME SEVERE COLD COUGH AND FLU RELIEF

acetaminophen, dextromethorphan hbr, phenylephrine hcl powder, for solution

### Product Information

|                                |      |
|--------------------------------|------|
| <b>Route of Administration</b> | ORAL |
|--------------------------------|------|

### Active Ingredient/Active Moiety

| Ingredient Name  | Basis of Strength             | Strength |
|--|-------------------------------|----------|
| <b>ACETAMINOPHEN</b> (UNII: 362O9ITL9D) (ACETAMINOPHEN - UNII:362O9ITL9D)                    | ACETAMINOPHEN                 | 500 mg   |
| <b>DEXTROMETHORPHAN HYDROBROMIDE</b> (UNII: 9D2RTI9KYH) (DEXTROMETHORPHAN - UNII:7355X3ROTS) | DEXTROMETHORPHAN HYDROBROMIDE | 20 mg    |
| <b>PHENYLEPHRINE HYDROCHLORIDE</b> (UNII: 04JA59TNSJ) (PHENYLEPHRINE - UNII:1WS297W6MV)      | PHENYLEPHRINE HYDROCHLORIDE   | 10 mg    |

### Inactive Ingredients

| Ingredient Name                                     | Strength |
|---|----------|
| ACESULFAME POTASSIUM (UNII: 23OV73Q5G9)             |          |
| ANHYDROUS CITRIC ACID (UNII: XF417D3PSL)            |          |
| ASPARTAME (UNII: Z0H242BBR1)                        |          |
| SILICON DIOXIDE (UNII: ETJ7Z6XBU4)                  |          |
| D&C YELLOW NO. 10 (UNII: 35SW5USQ3G)                |          |
| FD&C BLUE NO. 1 (UNII: H3R47K3TBD)                  |          |
| FD&C RED NO. 40 (UNII: WZB9127XOA)                  |          |
| MALTODEXTRIN (UNII: 7CVR7L4A2D)                     |          |
| SODIUM CITRATE, UNSPECIFIED FORM (UNII: 1Q73Q2JULR) |          |
| SUCROSE (UNII: C151H8M554)                          |          |
| TRIBASIC CALCIUM PHOSPHATE (UNII: 91D9GV0Z28)       |          |

### Product Characteristics

|          |                                       |              |  |
|----------|---------------------------------------|--------------|--|
| Color    |                                       | Score        |  |
| Shape    |                                       | Size         |  |
| Flavor   | HONEY (green tea) , LEMON (green tea) | Imprint Code |  |
| Contains |                                       |              |  |

### Packaging

| # | Item Code | Package Description                              | Marketing Start Date | Marketing End Date |
|---|-----------|--|----------------------|--------------------|
| 1 |           | 1 in 1 PACKET; Type 0: Not a Combination Product |                      |                    |

### Marketing Information

| Marketing Category  | Application Number or Monograph Citation | Marketing Start Date | Marketing End Date |
|---------------------|--|----------------------|--------------------|
| OTC monograph final | part341                                  |                      |                    |

### Part 2 of 2

## NIGHTTIME SEVERE COLD COUGH AND FLU RELIEF

acetaminophen, diphenhydramine hcl, phenylephrine hcl powder, for solution

### Product Information

|                         |      |
|-------------------------|------|
| Route of Administration | ORAL |
|-------------------------|------|

### Active Ingredient/Active Moiety

| Ingredient Name  | Basis of Strength | Strength |
|--|-------------------|----------|
| ACETAMINOPHEN (UNII: 362O9ITL9D) (ACETAMINOPHEN - UNII:362O9ITL9D) | ACETAMINOPHEN     | 650 mg   |
| DIPHENHYDRAMINE HYDROCHLORIDE (UNII: TC2261AD40)                   | DIPHENHYDRAMINE   |          |

|   |                                  |       |
|---|----------------------------------|-------|
| <b>DIPHENHYDRAMINE HYDROCHLORIDE</b> (UNII: 1CZD06JAD4U)<br>(DIPHENHYDRAMINE - UNII:8GTS82S83M) | DIPHENHYDRAMINE<br>HYDROCHLORIDE | 25 mg |
| <b>PHENYLEPHRINE HYDROCHLORIDE</b> (UNII: 04JA59TNSJ) (PHENYLEPHRINE - UNII:1WS297W6MV)         | PHENYLEPHRINE<br>HYDROCHLORIDE   | 10 mg |

### Inactive Ingredients

| Ingredient Name  | Strength |
|--|----------|
| <b>ACESULFAME POTASSIUM</b> (UNII: 23OV73Q5G9)             |          |
| <b>ANHYDROUS CITRIC ACID</b> (UNII: XF417D3PSL)            |          |
| <b>ASPARTAME</b> (UNII: Z0H242BBR1)                        |          |
| <b>SILICON DIOXIDE</b> (UNII: ETJ7Z6XBU4)                  |          |
| <b>D&amp;C YELLOW NO. 10</b> (UNII: 35SW5USQ3G)            |          |
| <b>FD&amp;C BLUE NO. 1</b> (UNII: H3R47K3TBD)              |          |
| <b>FD&amp;C RED NO. 40</b> (UNII: WZB9127XOA)              |          |
| <b>MALTODEXTRIN</b> (UNII: 7CVR7L4A2D)                     |          |
| <b>SODIUM CITRATE, UNSPECIFIED FORM</b> (UNII: 1Q73Q2JULR) |          |
| <b>SUCROSE</b> (UNII: C151H8M554)                          |          |
| <b>TRIBASIC CALCIUM PHOSPHATE</b> (UNII: 91D9GV0Z28)       |          |

### Product Characteristics

|                 |  |                     |  |
|-----------------|--|---------------------|--|
| <b>Color</b>    | WHITE (mixture of white, light yellow-orange particles) , ORANGE | <b>Score</b>        |  |
| <b>Shape</b>    |  | <b>Size</b>         |  |
| <b>Flavor</b>   |  | <b>Imprint Code</b> |  |
| <b>Contains</b> |  |                     |  |

### Packaging

| # | Item Code | Package Description                              | Marketing Start Date | Marketing End Date |
|---|-----------|--|----------------------|--------------------|
| 1 |           | 1 in 1 PACKET; Type 0: Not a Combination Product |                      |                    |

### Marketing Information

| Marketing Category  | Application Number or Monograph Citation | Marketing Start Date | Marketing End Date |
|---------------------|--|----------------------|--------------------|
| OTC monograph final | part341                                  |                      |                    |

### Marketing Information

| Marketing Category  | Application Number or Monograph Citation | Marketing Start Date | Marketing End Date |
|---------------------|--|----------------------|--------------------|
| OTC monograph final | part341                                  | 06/24/2019           |                    |

**Labeler** - Rite Aid Corporation (014578892)

