

**WALMART DAYTIME AND NIGHTTIME MS SEVERE RELIEF VALUE PACK-
acetaminophen, dextromethorphan hbr, diphenhydramine hcl
WAL-MART STORES INC**

equate Daytime & Nighttime relief Value Pack

Active ingredients (in each packet) - Daytime Severe Cold Relief

Acetaminophen 500 mg

Dextromethorphan hydrobromide 20 mg

Active ingredients (in each packet) - Nighttime Severe Cold Relief

Acetaminophen 650 mg

Diphenhydramine hydrochloride 25 mg

Purpose - Daytime Severe Cold Relief

Pain reliever/fever reducer

Cough Suppressant

Purpose - Nighttime Severe Cold Relief

Pain reliever/fever reducer

Antihistamine/Cough suppressant

Uses - Daytime Severe Cold Relief

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

Uses - Nighttime Severe Cold Relief

- temporarily relieves these symptoms due to a cold:
- minor aches and pains
- minor sore throat pain
- headache
- 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 - Daytime Severe Cold Relief /Nighttime Severe Cold Relief

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 - Daytime Severe Cold Relief

- in a child under 12 years of age
- if you are allergic to acetaminophen
- with any other drug containing acetaminophen (prescription or nonprescription). If you are not sure whether a drug contains acetaminophen, ask a doctor or a 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.

Do not use - Nighttime Severe Cold Relief

- in a child under 12 years of age
- if you are allergic to acetaminophen
- with any other drug containing acetaminophen (prescription or nonprescription). If you are not sure whether a drug contains acetaminophen, ask a doctor or a 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 - Daytime Severe Cold Relief

- liver disease
- 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 before use if you have - Nighttime Severe Cold Relief

- liver disease
- 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 - Daytime Severe Cold Relief

- taking the blood thinning drug warfarin

Ask a doctor or pharmacist before use if you are - Nighttime Severe Cold Relief

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

When using this product - Daytime Severe Cold Relief

do not exceed recommended dosage

When using this product - Nighttime Severe Cold Relief

- 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 - Daytime Severe Cold Relief /Nighttime Severe Cold and Cough

fever gets worse or lasts more than 3 days

- redness or swelling is present
- new symptoms occur
- pain, or cough 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.

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 - Daytime Severe Cold Relief

- **do not use more than directed**
- 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

Directions Nighttime Severe Cold Relief

- **do not use more than directed**
- 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 - Daytime Severe Cold Relief

- **each packet contains:**potassium 6 mg
- Aspartame free, sodium free
- store at room temperature. Protect from excessive heat and moisture.

Other information - Nighttime Severe Cold Relief

- **each packet contains:** potassium 6 mg
- Aspartame free, sodium free
- store at room temperature. Protect from excessive heat and moisture

Inactive ingredients - Daytime Severe Cold Relief

anhydrous citric acid, caramel, flavors, maltodextrin, potassium chloride, silica, sucralose, sucrose

Inactive ingredients - Nighttime Severe Cold Relief

anhydrous citric acid, F D & C yellow #6 , flavors, maltodextrin, potassium chloride, silica ,sucralose, sucrose,

Questions or comments?

1-866-467-2748

Package/Label Principal Display Panel

NDC# 49035-948-12

DAYTIME & NIGHTTIME RELIEF VALUE PACK

Compare to Theraflu® Multi Symptom Severe Cold Relief active ingredients* Nighttime Severe Cold Relief active ingredients*

NDC 79903-350-12

DAYTIME SEVERE MULTI- SYMPTOM

COLD RELIEF

Acetaminophen - Pain Reliever/ Fever Reducer

Dextromethorphan HBr - Cough Suppressant

Hot Liquid Therapy That Relieves

- Cough
- Head & Body
- Sore throat pain
- ache
- Fever

Honey Lemon Flavor

Infused with chamomile and white tea flavors

Naturally and Artificially Flavored

6 PACKETS

NIGHTTIME SEVERE MULTI-SYMPATOM

COLD Relief

Acetaminophen - Pain Reliever/ Fever Reducer

Dextromethorphan HBr - Cough Suppressant

Hot Liquid Therapy That Relieves

- Cough
- Head & Body
- Sore throat pain
- Ache
- Runny nose
- Sneezing
- Fever

Honey lemon flavor

Infused with chamomile & white tea flavors

6 PACKETS

12 TOTAL PACKETS

**READ ALL WARNINGS AND DIRECTIONS ON CARTON BEFORE USE .KEEP
CARTON FOR REFERENCE, DO NOT DISCARD.**

**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 DAYTIME PRODUCT UNLESS DIRECTED BY
DOCTOR.**

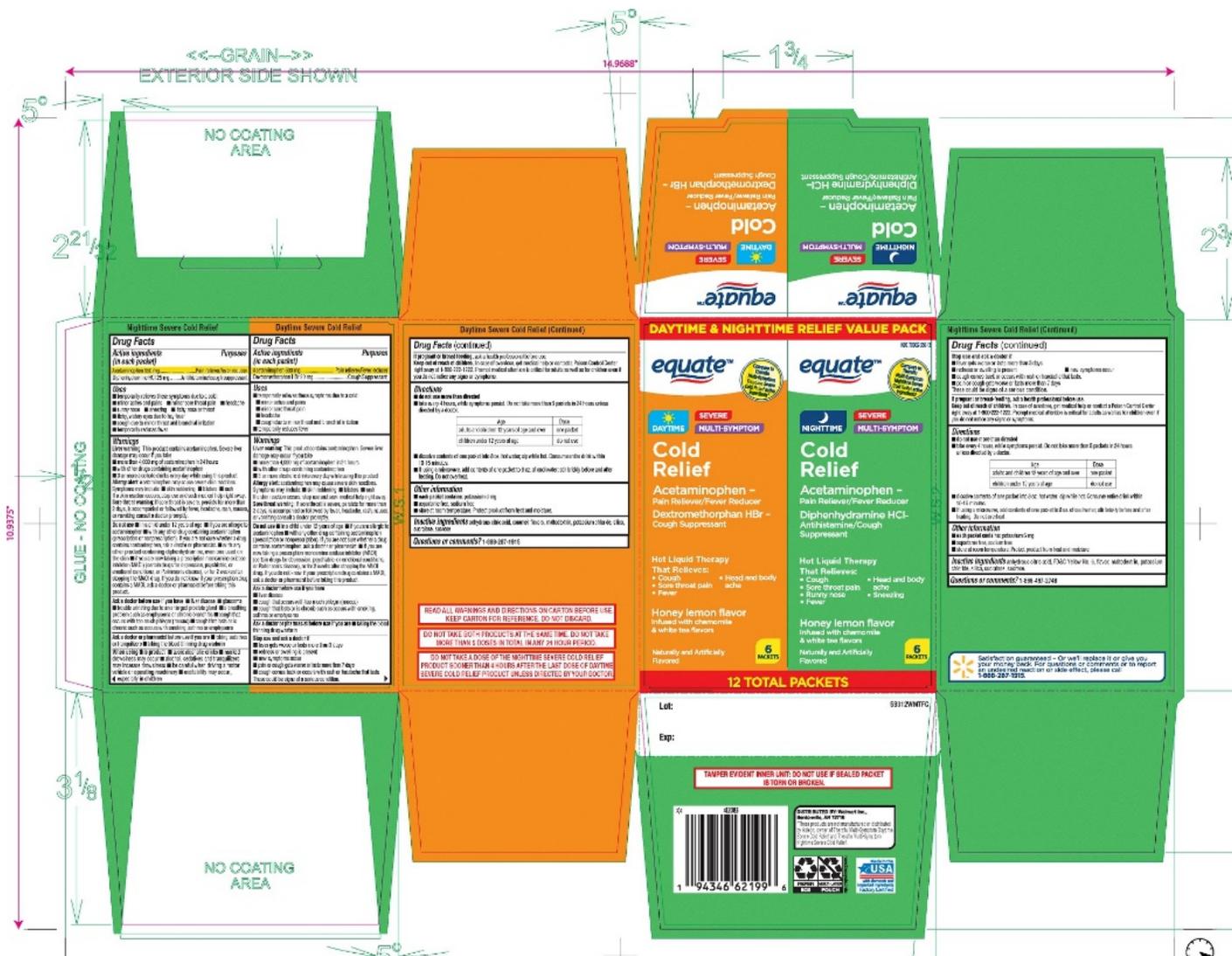
*These Products are not manufactured or distributed by Haleon, owner of Theraflu® Multi-Symptom Daytime Severe Cold & Theraflu® Multi-Symptom Nighttime Severe Cold Relief.

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Satisfaction guaranteed – or we'll replace it or give you your money back. For questions or comments or to report an undesired reaction or side effect, please call 1-888-287-1915

TAMPER EVIDENT INNER UNIT: DO NOT USE IF SEALED PACKET IS TORN OR BROKEN



WALMART DAYTIME AND NIGHTTIME MS SEVERE RELIEF VALUE PACK

acetaminophen, dextromethorphan hbr, diphenhydramine hcl kit

Product Information

Product Type

HUMAN OTC DRUG

Item Code (Source)

NDC:79903-350

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:79903-350-12	1 in 1 CARTON; Type 0: Not a Combination Product	02/02/2026	

Quantity of Parts

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

Part 1 of 2**WALMART DAYTIME MS SEVERE COLD**

acetaminophen, dextromethorphan hbr powder, for solution

Product Information

Item Code (Source)	NDC:79903-436
Route of Administration	ORAL

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
ACETAMINOPHEN (UNII: 362O9ITL9D) (ACETAMINOPHEN - UNII:362O9ITL9D)	ACETAMINOPHEN	500 mg
DEXTROMETHORPHAN HYDROBROMIDE (UNII: 9D2RT9KYH) (DEXTROMETHORPHAN - UNII:7355X3ROTS)	DEXTROMETHORPHAN HYDROBROMIDE	20 mg

Inactive Ingredients

Ingredient Name	Strength
ANHYDROUS CITRIC ACID (UNII: XF417D3PSL)	
CARMEL (UNII: T9D99G2B1R)	
MALTODEXTRIN (UNII: 7CVR7L4A2D)	
POTASSIUM CHLORIDE (UNII: 660YQ98I10)	
SILICON DIOXIDE (UNII: ETJ7Z6XBU4)	
SUCRALOSE (UNII: 96K6UQ3ZD4)	
SUCROSE (UNII: C151H8M554)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:79903-436-06	6 in 1 CARTON		

1	1 in 1 PACKET; Type 0: Not a Combination Product		
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Marketing Information

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

Part 2 of 2

WALMART NIGHTTIME MS SEVERE COLD

acetaminophen, diphenhydramine hcl powder, for solution

Product Information

Item Code (Source)	NDC:79903-439
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: TC2D6JAD40) (DIPHENHYDRAMINE - UNII:8GTS82S83M)	DIPHENHYDRAMINE HYDROCHLORIDE	25 mg

Inactive Ingredients

Ingredient Name	Strength
ANHYDROUS CITRIC ACID (UNII: XF417D3PSL)	
FD&C YELLOW NO. 6 (UNII: H77VEI93A8)	
MALTODEXTRIN (UNII: 7CVR7L4A2D)	
POTASSIUM CHLORIDE (UNII: 660YQ98I10)	
SILICON DIOXIDE (UNII: ETJ7Z6XBU4)	
SUCRALOSE (UNII: 96K6UQ3ZD4)	
SUCROSE (UNII: C151H8M554)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:79903-439-06	6 in 1 CARTON		
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 Drug	M012	02/02/2026	

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

Labeler - WAL-MART STORES INC (051957769)

Revised: 2/2026

WAL-MART STORES INC