OTIS CLAPP VALIHIST- acetaminophen,chlorpheniramine maleate, phenylephrine hydrochloride tablet, film coated
MEDIQUE MEDICIDIN D- acetaminophen, chlorpheniramine maleate, phenylephrine hydrochloride tablet, film coated
Unifirst First 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.

Medique Medicidin D/ Otis Clapp Valihist

Drug Facts

Active ingredients (in each tablet)

Acetaminophen 325 mg

Chlorpheniramine Maleate 2 mg

Phenylephrine Hydrochloride 5 mg

Purpose

Pain reliever/fever reducer

Antihistamine

Nasal decongestant

Uses

Temporarily relieves
- minor aches and pains
- headache
- nasal congestion
- sinus congestion and pressure
- runny nose
- sneezing

Temporarily
- reduces fever
- relieves runny nose and reduces sneezing, itching of the nose and throat and itchy, watery eyes due to hay fever
- helps clear nasal and/or sinus passages

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, which is the maximum daily amount
- 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.

Do not use

- with any other drug containing acetaminophen (prescription or nonprescription). If you are not sure whether a drug contains acetaminophen, ask a doctor or pharmacist.
- for more than 10 days for pain unless directed by a doctor
- if you ever had an allergic reaction to this product or any of its ingredients
- 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
thyroid disease
diabetes
high blood pressure
heart disease
glaucoma
chronic bronchitis or emphysema
difficulty in urination due to enlargement of the prostate gland

Ask a doctor or pharmacist before use if you are

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

When using this product

- **do not exceed recommended dosage**
- excitability may occur, especially in children
- may cause drowsiness
- alcohol, sedatives and tranquilizers may increase drowsiness
- avoid alcoholic beverages
- use caution when driving a motor vehicle or operating machinery

Stop use and ask a doctor if

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

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 (1-800-222-12220. Quick 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**

Adults and children (12 years and older): Take 2 tablets every 4 to 6 hours. Do not take more than 10 tablets in 24 hours.

Children under 12 years: Ask a doctor

Other information

- store at room temperature 59º-86ºF (15º-30ºC)
- tamper evident sealed packets
- do not use any opened or torn packets
- avoid excessive heat and humidity

Inactive ingredients

corn starch, crospovidone, hypromellose, microcrystalline cellulose, polyethylene glycol, povidone, pregelatinized starch, stearic acid

Questions or comments? 1-800-634-7680

Principal Display Panel - Otis Clapp Valihist Label

Otis Clapp

Quality and Integrity Since 1840

VALIHIST ™

Decongestant-Antihistamine

Pain Reliever-Fever Reducer

Antihistamine- Decongestant

For Allergies, Colds, Runny Nose

See Warnings and Directions on Side Panel

This Package is for Households without Young Children.

Acetaminophen 325 mg,

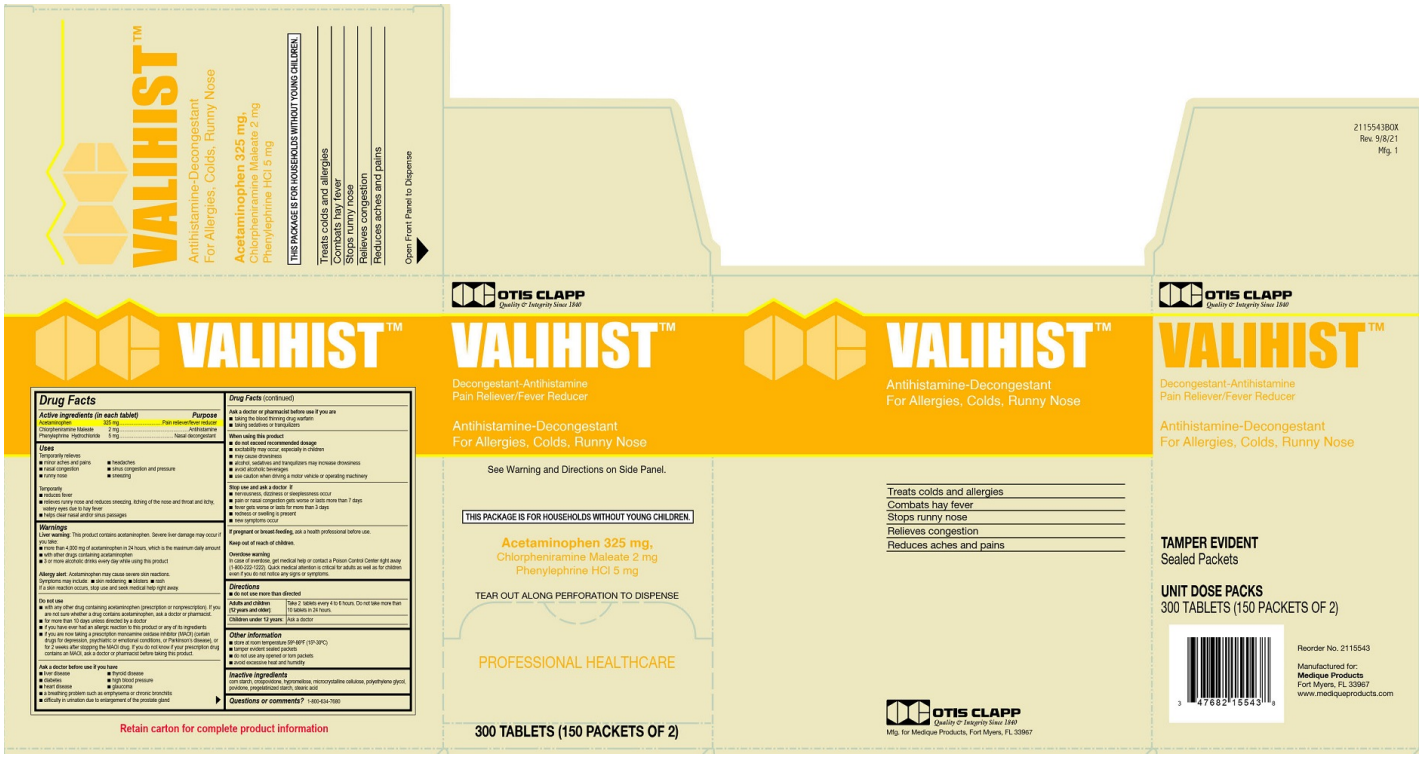
Chlorpheniramine Maleate 2mg

Phenylephrine HCl 5 mg

Tear Out Along Perforation To Dispense

PROFESSIONAL HEALTHCARE

300 TABLETS (150 PACKETS OF 2)



Principal Display Panel - Medique Medicidin D Label

Medique®

Medicidin-D

Cold and Allergy Relief

This Package is for Households without Young Children.

Pull to Open

Pain Reliever/ Fever Reducer ● Acetaminophen 325mg

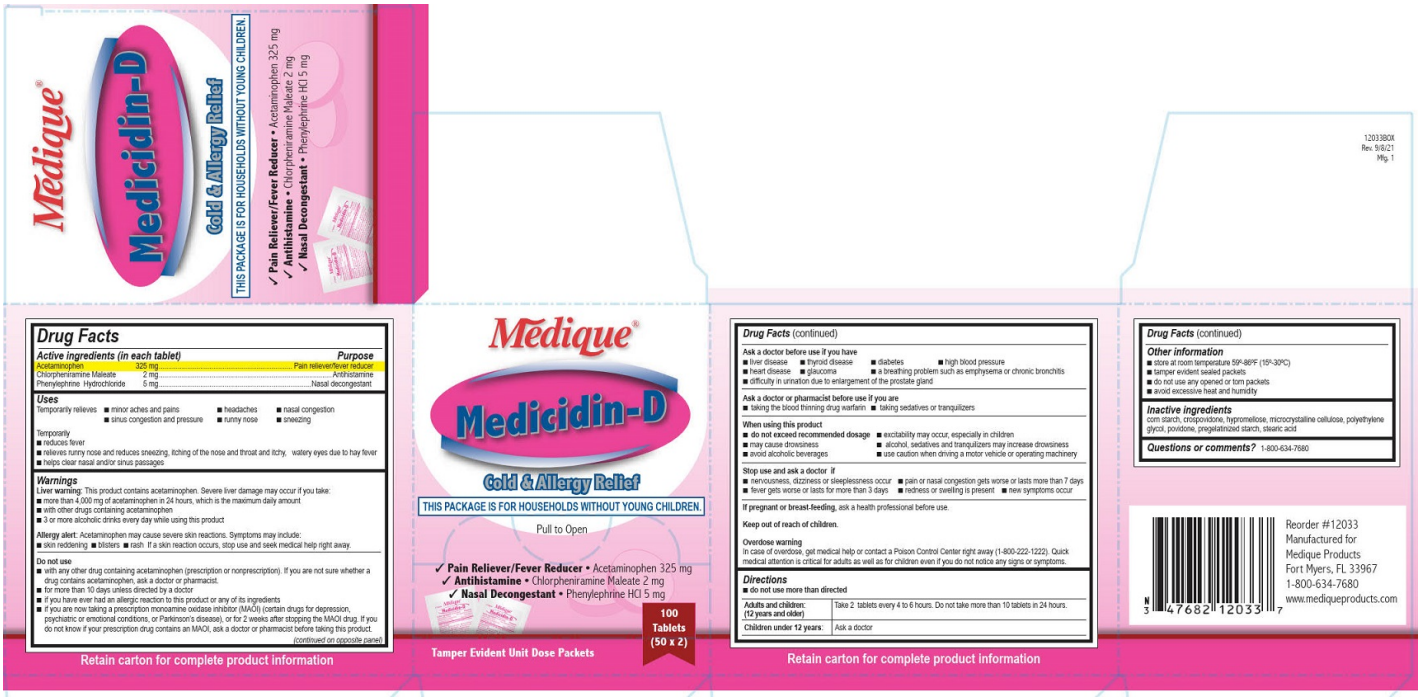
Antihistamine ● Chlorpheniramine Maleate 2mg

Nasal Decongestant ● Phenylephrine HCl 5mg

100Tablets

(50 x 2)

Tamper Evident Unit Dose Packets



OTIS CLAPP VALIHIST

acetaminophen,chlorpheniramine maleate, phenylephrine hydrochloride tablet, film coated

Product Information

Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:47682-543
Route of Administration	ORAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
ACETAMINOPHEN (UNII: 362O9ITL9D) (ACETAMINOPHEN - UNII:362O9ITL9D)	ACETAMINOPHEN	325 mg
CHLORPHENIRAMINE MALEATE (UNII: V1Q0O9OJ9Z) (CHLORPHENIRAMINE - UNII:3U6IO1965U)	CHLORPHENIRAMINE MALEATE	2 mg
PHENYLEPHRINE HYDROCHLORIDE (UNII: 04JA59TNSJ) (PHENYLEPHRINE - UNII:1WS297W6MV)	PHENYLEPHRINE HYDROCHLORIDE	5 mg

Inactive Ingredients

Ingredient Name	Strength
CELLULOSE, MICROCRYSTALLINE (UNII: OP1R32D61U)	
HYPROMELLOSES (UNII: 3NXW29V3WO)	
POLYETHYLENE GLYCOL, UNSPECIFIED (UNII: 3WQ0SDW1A)	
STARCH, CORN (UNII: O8232NY3SJ)	
STEARIC ACID (UNII: 4ELV7Z65AP)	
CROSPOVIDONE (UNII: 68401960MK)	
POVIDONE (UNII: FZ989GH94E)	

Product Characteristics

Color	white (white)	**Score**	no score
Shape	ROUND (ROUND)	**Size**	11mm
Flavor		**Imprint Code**	AZ;275
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:47682-543-03	150 in 1 BOX	12/30/2008	
1	NDC:47682-543-99	2 in 1 PACKET; Type 0: Not a Combination Product		
2	NDC:47682-543-99	2 in 1 PACKET; Type 0: Not a Combination Product	12/30/2008	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph final	part341	12/30/2008	

MEDIQUE MEDICIDIN D

acetaminophen, chlorpheniramine maleate, phenylephrine hydrochloride tablet, film coated

Product Information

Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:47682-120
Route of Administration	ORAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
ACETAMINOPHEN (UNII: 362O9ITL9D) (ACETAMINOPHEN - UNII:362O9ITL9D)	ACETAMINOPHEN	325 mg
CHLORPHENIRAMINE MALEATE (UNII: V1Q0O9OJ9Z) (CHLORPHENIRAMINE - UNII:3U6IO1965U)	CHLORPHENIRAMINE MALEATE	2 mg
PHENYLEPHRINE HYDROCHLORIDE (UNII: 04JA59TNSJ) (PHENYLEPHRINE - UNII:1WS297W6MV)	PHENYLEPHRINE HYDROCHLORIDE	5 mg

Inactive Ingredients

Ingredient Name	Strength
CELLULOSE, MICROCRYSTALLINE (UNII: OP1R32D61U)	
HYPROMELLOSES (UNII: 3NXW29V3WO)	
POLYETHYLENE GLYCOL, UNSPECIFIED (UNII: 3WQ0SDW1A)	
STARCH, CORN (UNII: O8232NY3SJ)	
STEARIC ACID (UNII: 4ELV7Z65AP)	
CROSPOVIDONE (UNII: 68401960MK)	
POVIDONE (UNII: FZ989GH94E)	

Product Characteristics			
Color	white (white)	**Score**	no score
Shape	ROUND (ROUND)	**Size**	11mm
Flavor		**Imprint Code**	AZ;275
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:47682-120-64	12 in 1 BOX	12/30/2008	
1		2 in 1 PACKET; Type 0: Not a Combination Product		
2	NDC:47682-120-33	50 in 1 BOX	12/30/2008	
2		2 in 1 PACKET; Type 0: Not a Combination Product		
3	NDC:47682-120-47	100 in 1 BOX	12/30/2008	
3		2 in 1 PACKET; Type 0: Not a Combination Product		
4	NDC:47682-120-13	250 in 1 BOX	12/30/2008	
4	NDC:47682-120-99	2 in 1 PACKET; Type 0: Not a Combination Product		
5	NDC:47682-120-99	2 in 1 PACKET; Type 0: Not a Combination Product	12/30/2008	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph final	part341	12/30/2008	

Labeler - Unifirst First Aid Corporation (832947092)

Revised: 3/2022 Unifirst First Aid Corporation