

WHITUBEN- acetaminophen, dextromethorphan hydrobromide hydrate, guaifenesin, pseudoephedrine hydrochloride, triprolidine hydrochloride hydrate capsule
OASIS TRADING

Disclaimer: This drug has not been found by FDA to be safe and effective, and this labeling has not been approved by FDA. For further information about unapproved drugs, click here.

Acetaminophen 180mg
Dextromethorphan Hydrobromide Hydrate 8mg
DL-Methylephedrine Hydrochloride 12.5mg
Guaifenesin 20.84mg
Pseudoephedrine Hydrochloride 15mg
Triprolidine Hydrochloride Hydrate 0.66mg

relieve your cold symptoms

Keep out of reach of children

- adult and children 15 years of age and older: 2 capsules three times a day
- children under 15 years of age: ask a doctor

Warnings

When using this product

Do not use for a period longer than 1 weeks

Stop use and ask a doctor if nervousness, dizziness, or sleepness occur. Pain, nasal congestion or cough gets worse or lasts more than 7 days.

fever gets worse or lasts more than 3 days

New symptoms occur

These may 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

cellulose, corn starch, magnesium stearate, sodium starch glycolate

For oral use only

WHITUBEN

Drug Facts	
Active Ingredients (in one capsule)	Purpose
Acetaminophen 180g	Aches and pain
Dextromethorphan Hydrobromide Hydrate 8mg	Aches and pain
DL-Methylephedrine Hydrochloride 12.5mg	Aches and pain
Guaifenesin 20.84mg 20.84mg	Aches and pain
Pseudoephedrine Hydrochloride 15mg	Aches and pain
Triprolidine Hydrochloride Hydrate 0.66mg	Aches and pain
Uses	
<ul style="list-style-type: none"> relieve your cold symptoms 	
Warnings	
<p>When using this product Do not use for a period longer than 1 week</p> <p>Stop use and ask a doctor if nervousness, dizziness, or sleepiness occur. Pain, nasal congestion or cough gets worse or lasts more than 7 days.</p> <p>fever gets worse or lasts more than 3 days. New symptoms occur. These may be signs of a serious condition.</p> <p>If pregnant or breast-feeding, ask a health professional before use.</p> <p>Keep out of reach of children, In case of overdose, get medical help or contact a poison control center right away</p>	
Directions	
<ul style="list-style-type: none"> adult and children 15 years of age and older: 2 capsules three times a day children under 15 years of age: ask a doctor 	
Other Information	
<ul style="list-style-type: none"> Store at room temperature, 15–30°C (59–86°F) 	
Inactive Ingredient	
cellulose, corn starch, magnesium stearate, sodium starch glycolate	
Questions or comments ?	
Call weekdays from 9 a.m to 5 p.m EST at (201) 669-8405	
Distributed By: P&K FRONTIER MARKETING CORP.	
329 BROAD AVENUE # 2F, PALISADES PARK, NJ 07650, USA	
Made in South Korea	

WHITUBEN

acetaminophen, dextromethorphan hydrobromide hydrate, guaifenesin, pseudoephedrine hydrochloride, triprolidine hydrochloride hydrate capsule

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:72689-0037
Route of Administration	ORAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
ACETAMINOPHEN (UNII: 362O9ITL9D) (ACETAMINOPHEN - UNII:362O9ITL9D)	ACETAMINOPHEN	180 mg
GUAIFENESIN (UNII: 495W7451VQ) (GUAIFENESIN - UNII:495W7451VQ)	GUAIFENESIN	20.84 mg
PSEUDOEPHEDRINE HYDROCHLORIDE (UNII: 6V9V2RYJ8N) (PSEUDOEPHEDRINE - UNII:7CUC9DDI9F)	PSEUDOEPHEDRINE HYDROCHLORIDE	15 mg
METHYLEPHEDRINE HYDROCHLORIDE, (+/-)- (UNII: 99214P83XM) (METHYLEPHEDRINE, (+/-)- - UNII:SHS9PGQ2LS)	METHYLEPHEDRINE HYDROCHLORIDE, (+/-)-	12.5 mg

TRIPROLIDINE HYDROCHLORIDE (UNII: YAN7R5L890) (TRIPROLIDINE - UNII:2L8T9S52QM)	TRIPROLIDINE HYDROCHLORIDE	0.66 mg
DEXTROMETHORPHAN HYDROBROMIDE (UNII: 9D2RTI9KYH) (DEXTROMETHORPHAN - UNII:7355X3ROTS)	DEXTROMETHORPHAN HYDROBROMIDE	8 mg

Inactive Ingredients

Ingredient Name	Strength
STARCH, CORN (UNII: O8232NY3SJ)	

Product Characteristics

Color	red	Score	no score
Shape	OVAL	Size	14mm
Flavor		Imprint Code	TWLQQ
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:72689-0037-1	10 in 1 BLISTER PACK; Type 0: Not a Combination Product	11/22/2018	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
unapproved drug other		11/22/2018	

Labeler - OASIS TRADING (689991468)

Registrant - OASIS TRADING (689991468)

Establishment

Name	Address	ID/FEI	Business Operations
OASIS TRADING		689991468	manufacture(72689-0037) , label(72689-0037)

Revised: 3/2019

OASIS TRADING