# BC SINUS CONGESTION AND PAIN- acetaminophen, chlorpheniramine maleate, phenylephrine hydrochloride powder Medtech Products Inc.

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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#### **BC Sinus Pain and Congestion**

#### Drug Facts

#### Active ingredients (in each powder)

Acetaminophen 650 mg Chlorpheniramine maleate 4mg Phenylephrine hydrochloride 10mg

#### Purposes

Pain reliever/Fever reducer Antihistamine Nasal decongestant

#### Uses

- temporarily relieves these allergy and cold symptoms:
  - sinus congestion and pressure
  - headache
  - nasal congestion
  - runny nose
  - sneezing
  - minor anches and pains
- temporarily reduces fever

#### Warnings

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

- more than 4,000 mg 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.

#### 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.
- if you are now taking a prescription monoamine oxidase inhibitor (MOAI) (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
- high blood pressure
- diabetes
- thyroid disease
- glaucoma
- heart disease
- a breathing problem such as emphysema or chronic bronchitis
- trouble urinating due to an enlarged prostate gland

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

- the blood thinning drug warfarin
- sedatives or tranquilizers

#### When using this product

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

#### Stop use and ask a doctor if

- pain or nasal congestion gets worse or lasts more than 7 days
- fever gets worse or lasts more than 3 days
- you get nervous, dizzy, or sleepless
- redness or swelling is present
- new symptoms occur

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**: Taking more than the recommended dose can cause serious health problems. In the case of overdose, get medical help or contact a Poison Control Center (1-800-222-1222) right away. 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 take more than directed (see overdose warning)
- **adults and children 12 years of age and over**: place 1 powder on tongue every 4-6 hours, while symptoms persist.
- do not take more than 6 powders in 24 hours unless directed by a doctor.
- children under 12 years of age: ask a doctor.

#### Other information

Store below 25°C (77°F)

#### **Inactive ingredients**

magnesium stearate, maltodextrin, silica

#### **Questions?**

## 1-866-255-5197 bcpowder.com TAMPER EVIDENT: DO NOT USE IF PACKET IS DAMAGED OR OPEN.

#### PRINCIPAL DISPLAY PANEL

BC<sup>™</sup> Sinus Pain & Congestion

AcetaminophenPain Reliever / Fever ReducerChlorpheniramineMaleate AntihistaminePhenylephrine HClNasal Decongestant

#### 12 Powders



# **BC SINUS CONGESTION AND PAIN**

acetaminophen, chlorpheniramine maleate, phenylephrine hydrochloride powder

#### **Product Information**

Product Type		HUMAN OTC DRUG Item Coo		ode (So	urce)	NDC:63029-221		
Route of Administra	ation	ORAL						
Active Ingredien	t/Active Moi	ety						
Ingredient Name					<b>Basis of Strength</b>		Strengt	
ACETAMINOPHEN (UNII: 36209ITL9D) (ACETAMINOPHEN - UNII:36209ITL9D)					ACETAMINOPHEN		650 mg	
CHLORPHENIRAMINE MALEATE (UNII: V1Q0O9OJ9Z) (CHLORPHENIRAMINE - UNII:3U6IO1965U)					CHLO RPHENIRAMINE MALEATE		4 mg	
PHENYLEPHRINE HYDRO CHLO RIDE (UNII: 04JA59TNSJ) (PHENYLEPHRINE - UNII: 1WS297W6 MV)					PHENYLEPHRINE HYDROCHLORIDE		10 mg	
Inactive Ingredie	ents							
Ingredient Name							Strength	
						50101	igtii	
MAGNESIUM STEAR	<b>ATE</b> (UNII: 7009					54141	igtii	
MALTO DEXTRIN (UI	NII: 7CVR7L4A2E	7M6I30) ))				5.111	igtii	
MALTO DEXTRIN (UI	NII: 7CVR7L4A2E	7M6I30) ))					igtii	
MALTO DEXTRIN (UI	NII: 7CVR7L4A2E	7M6I30) ))					igui	
MALTO DEXTRIN (UI	NII: 7CVR7L4A2E	7M6I30) ))					ig til	
MALTO DEXTRIN (UI SILICON DIO XIDE (U	NII: 7CVR7L4A2E	7M6I30) ))					igen	
MALTO DEXTRIN (UI SILICON DIO XIDE (U Packaging	NII: 7CVR7L4A2E JNII: ETJ7Z6XBU	7M6I30) ))		Marke	ting Start Date			
MALTO DEXTRIN (UI SILICON DIO XIDE (U Packaging # Item Code	NII: 7CVR7L4A2E JNII: ETJ7Z6XBU	7M6I30) )) (4)		<b>Marke</b> 0 3/0 1/20	-			
MALTODEXTRIN (UI SILICON DIO XIDE (U Packaging I tem Code NDC:63029-221-04	NII: 7CVR7L4A2E JNII: ETJ7Z6XBU 4 in 1 CARTRID	7M6I30) )) [4) Package Description	Product		0 16			
MALTODEXTRIN (UI SILICON DIO XIDE (U Packaging H Item Code NDC:63029-221-04	NII: 7CVR7L4A2E JNII: ETJ7Z6XBU 4 in 1 CARTRID	7M6I30) )) (4) <b>Package Description</b> GE; Type 0: Not a Combination	Product	03/01/20	0 16			
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MALTODEXTRIN (UI SILICON DIO XIDE (U Packaging H Item Code NDC:63029-221-04	NII: 7CVR7L4A2E JNII: ETJ7Z6XBU 4 in 1 CARTRID 12 in 1 CARTON	7M6I30) )) (4) <b>Package Description</b> GE; Type 0: Not a Combination	Product	03/01/20	0 16			
MALTODEXTRIN (UI   SILICON DIO XIDE (U   Packaging   # Item Code   1 NDC:63029-221-04   2 NDC:63029-221-12	NII: 7CVR7L4A2E JNII: ETJ7Z6XBU 4 in 1 CARTRID 12 in 1 CARTON	7M6I30) )) (4) <b>Package Description</b> GE; Type 0: Not a Combination	n Product roduct	0 3/0 1/20 0 3/0 1/20	0 16		; End Date	

# Labeler - Medtech Products Inc. (122715688)

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Medtech Products Inc.