TYLENOL SINUS SEVERE- acetaminophen, guaifenesin, and phenylephrine hydrochloride tablet, coated Johnson & Johnson Consumer Inc.

Tylenol ® Sinus Severe

Drug Facts

| Active ingredients (in each caplet) | Purpose |
|-------------------------------------|----------------|
| | Pain |
| Acetaminophen 325 mg | reliever/fever |
| | reducer |
| Guaifenesin 200 mg | Expectorant |
| Phonylophrino UCLE ma | Nasal |
| Phenylephrine HCl 5 mg | decongestant |

Uses

- temporarily relieves these symptoms associated with hay fever or other respiratory allergies, and the common cold:
 - sinus congestion and pressure
 - headache
 - nasal congestion
 - minor aches and pains
- helps loosen phlegm (mucus) and thin bronchial secretions to make coughs more productive
- temporarily reduces fever

Warnings

Liver warning

This product contains acetaminophen. The maximum daily dose of this product is 10 caplets (3,250 mg acetaminophen) in 24 hours. 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.

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 (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.
- if you have ever had an allergic reaction to this product or any of its ingredients.

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
- persistent or chronic cough such as occurs with smoking, asthma, chronic bronchitis, or emphysema
- cough that occurs with too much phlegm (mucus)

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

When using this product do not exceed recommended dose

Stop use and ask a doctor if

- nervousness, dizziness, or sleeplessness occur
- pain, nasal congestion or cough 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
- 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) 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 and over | |

- swallow whole; do not crush, chew or dissolve
- do not take more than 10 caplets in 24 hours

children under 12 years ask a doctor

Other information

- each caplet contains: sodium 3 mg
- store between 20-25°C (68-77°F)
- do not use if blister unit is torn or broken

Inactive ingredients

carnauba wax, croscarmellose sodium, flavor, hydroxypropyl cellulose, hypromellose, magnesium stearate, microcrystalline cellulose, pregelatinized starch, sucralose, titanium dioxide, triacetin

Questions or comments?

call **1-877-895-3665** (toll-free) or **215-273-8755** (collect)

PRINCIPAL DISPLAY PANEL

NDC 50580-507-01 TYLENOL [®] FOR ADULTS

SINUS SEVERE

Acetaminophen, Phenylephrine HCl, Guaifenesin Pain Reliever-Fever Reducer, Nasal Decongestant, Expectorant

DAY NON-DROWSY

- SINUS HEADACHE
- SINUS PRESSURE
- NASAL CONGESTION
- MUCUS + CHEST CONGESTION

Actual Size

24 CAPLETS



TYLENOL SINUS SEVERE

acetaminophen, guaifenesin, and phenylephrine hydrochloride tablet, coated

| Product Information | | | | |
|-------------------------|----------------|--------------------|---------------|--|
| Product Type | HUMAN OTC DRUG | Item Code (Source) | NDC:50580-507 | |
| Route of Administration | ORAL | | | |

| Active Ingredient/Active Moiety | | | | |
|---|--------------------------------|----------|--|--|
| Ingredient Name | Basis of Strength | Strength | | |
| ACETAMINOPHEN (UNII: 36209ITL9D) (ACETAMINOPHEN - UNII:36209ITL9D) | ACETAMINOPHEN | 325 mg | | |
| GUAIFENESIN (UNII: 495W7451VQ) (GUAIFENESIN - UNII:495W7451VQ) | GUAIFENESIN | 200 mg | | |
| PHENYLEPHRINE HYDROCHLORIDE (UNII: 04JA59TNSJ) (PHENYLEPHRINE - UNII: 1WS297W6MV) | PHENYLEPHRINE HYDROCHLORIDE | 5 mg | | |

| Inactive Ingredients | |
|----------------------|----------|
| Ingredient Name | Strength |

| CARNAUBA WAX (UNII: R12CBM0EIZ) | |
|---|--|
| CROSCARMELLOSE SODIUM (UNII: M280L1HH48) | |
| HYDROXYPROPYL CELLULOSE, UNSPECIFIED (UNII: 9XZ8H6N6OH) | |
| HYPROMELLOSE, UNSPECIFIED (UNII: 3NXW29V3WO) | |
| MAGNESIUM STEARATE (UNII: 70097M6I30) | |
| MICROCRYSTALLINE CELLULOSE (UNII: OP1R32D61U) | |
| SUCRALOSE (UNII: 96K6UQ3ZD4) | |
| TITANIUM DIOXIDE (UNII: 15FIX9V2JP) | |
| TRIACETIN (UNII: XHX3C3X673) | |

| Product Characteristics | | | | |
|-------------------------|-------|--------------|--------------|--|
| Color | white | Score | no score | |
| Shape | OVAL | Size | 20mm | |
| Flavor | MINT | Imprint Code | TYLENOL;1072 | |
| Contains | | | | |

| P | Packaging | | | | | |
|---|----------------------|---|-------------------------|-----------------------|--|--|
| # | Item Code | Package Description | Marketing Start Date | Marketing End Date | | |
| 1 | NDC:50580- 507-01 | 2 in 1 CARTON | 05/15/2015 | | | |
| 1 | | 12 in 1 BLISTER PACK; Type 0: Not a Combination Product | | | | |
| 2 | NDC:50580- 507-03 | 2 in 1 POUCH; Type 0: Not a Combination Product | 07/17/2017 | | | |
| 3 | NDC:50580- 507-04 | 50 in 1 CARTON | 07/17/2017 | | | |
| 3 | | 2 in 1 POUCH; Type 0: Not a Combination Product | | | | |
| 4 | NDC:50580- 507-02 | 3 in 1 PACKAGE | 07/06/2020 | | | |
| 4 | | 2 in 1 CARTON | | | | |
| 4 | | 12 in 1 BLISTER PACK; 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 | 05/15/2015 | |
| | | | |

Labeler - Johnson & Johnson Consumer Inc. (878046358)