APAP 500

SAFETY DATA SHEET

SECTION 1, IDENTIFICATION

SDS NAME:	APAP 500
SDS NUMBER:	L211
INFORMATION:	ULTRAtab Laboratories Inc.
	50 Toc Drive
	Highland, NY 12528
	(845)691-8361
EMERGENCY CONTACT:	Dyana Baker
	(845)691-8361

North America: CHEMTREC: 1-800-424-9300

Outside North America: 1-703-527-3887

Consumers: Refer to the package insert or product label for appropriate consumer-specific information about this product when used according to the manufacturer's directions.

Recommended use: Analgesic, Antipyretic

SECTION 2, HAZARD IDENTIFICATION

- **Label Warnings:**
- **Alcohol Warning:** If you consume 3 or more alcoholic drinks every day, ask your doctor whether you should take acetaminophen. Acetaminophen may cause liver damage.

Do not use:

- with any other pain reliever/fever reducer
- if you have ever had an allergic reaction to any other pain reliever/fever reducer
- for pain for more than 10 days or for fever for more than 3 days unless directed by a doctor
- with any other product containing acetaminophen

Ask doctor before using if you have:

- asthma
- stomach problems (such as heartburn, upset stomach or stomach pain)
- gastric ulcers
- bleeding problems

Ask a doctor or pharmacist before use if you are taking a prescription drug for:

- anticoagulation (thinning of blood)
- diabetes
- gout
- arthritis

When using this product do not exceed recommended dose

Stop use and ask a doctor if:

- ringing in the ears or loss of hearing occurs
- pain or fever persists or gets worse
- new symptoms occur redness or swelling is present

SECTION 3, COMPOSITION AND INFORMATION ON INGREDIENTS

PRODUCT USE: Drug product

INGREDIENT	CAS NUMBER	PERCENT
Acetaminophen	103-90-2	80
DI H2O	7732-18-5	proprietary
Starch 1551	9005-25-8	proprietary
Microcrystalline Cellulose	9004-34-6	proprietary
Sodium Starch Glycolate	9063-38-1	proprietary
PVP K30	9003-39-8	proprietary
Stearic Acid	57-11-4	proprietary
Opadry II Clear	25322-68-3	proprietary

The formulations for these products are proprietary information. Inactive ingredients of less than 1% not displayed above.

SECTION 4, FIRST-AID MEASURES

INHALATION: Obtain medical attention

SKIN CONTACT: After contact with skin, wash immediately with plenty of water and get medical attention if irritation develops or persists.

EYE CONTACT: In case of contact with eyes, rinse immediately with plenty of water and seek medical advice.

INGESTION: If swallowed, seek medical attention immediately

SECTION 5, FIREFIGHTING MEASURES

FLAMMABILITY DATA:

Flash Point: N/A

Flammable Limits: No data available.

Upper Explosion Limit (UEL%): No data available.

Lower Explosion Limit (LEL%): No data available.

SPECIAL FIRE FIGHTING PROCEDURES: Wear positive pressure self-contained breathing apparatus.

EXTINGUISHING MEDIA: Water, Dry chemical, foam, CO_2.

HAZARDOUS PRODUCTS: Oxides of carbon, salicylic acid vapors.

See section 9 for physical and chemical properties.

SECTION 6, ACCIDENTAL RELEASE MEASURES

PERSONAL PRECAUTIONS: None required.

SPILLRESPONSE/ CLEANUP: Scoop up material, wash area with warm, soapy water.

See Sections 9 and 10 for additional physical, chemical and hazard information.

SECTION 7, HANDLING AND STORAGE

HANDLING AND STORAGE: No special handling or storage necessary. This product should be handled and stored per label instructions to insure product integrity.

SECTION 8, EXPOSURE CONTROLS AND PERSONAL PROTECTION

The following guidance applies to the handling of the active ingredient(s) in this formulation. The end-user should perform an appropriate risk assessment when handling other forms or formulations of this active ingredient.

EXPOSURE CONTROLS: None required.

RECOMMENDED PERSONAL PROTECTIVE EQUIPMENT (PPE): None required.

Respiratory Protection: None required.

Skin Protection: None required.

Eye Protection: None required.

Ventilation: None required.

Contaminated Equipment: None required.

SECTION 9, PHYSICAL AND CHEMICAL PROPERTIES

FORM: Tablet

COLOR: White

ODOR: Odorless

SOLUBILITY IN H_2O (g/100ml @ 25°C): 90

pH: N/A

See Section 5 for flammability/explosivity information.

SECTION 10, STABILITY AND REACTIVITY

STABILITY/REACTIVITY: Stable under normal conditions.

INCOMPATIBLE MATERIALS/CONDITIONS TO AVOID: Strong oxidizing agents.

HAZARDOUS DECOMPOSITION PRODUCTS/REACTIONS: Oxides of carbon, salicylic acid vapors.

SECTION 11, TOXICOLOGICAL INFORMATION

The information below pertains to the formulated product unless indicated otherwise.

ACUTE TOXICITY DATA: Acetaminophen may cause liver damage.

SECTION 12, ECOLOGICAL CONSIDERATIONS

ECOTOXICITY: This product has not been tested for Eco toxicity.

ENVIROMENTAL DATA: There is no environmental data available for this product.

SECTION 13, DISPOSAL CONSIDERATIONS

MATERIAL WASTE:

Disposal must be in accordance with applicable federal, state/provincial, and/or local regulations. Incarceration is the preferred method of disposal, when appropriate. Operations that involve the crushing or shredding of waste materials or returned goods must be handled to meet the recommended exposure limit(s).

PACKAGING AND CONTAINERS:

Disposal must be in accordance with federal, state/provincial, and/or local regulations.

SECTION 14, TRANSPORT INFORMATION

This material is not subject to the transportation regulations of DOT, IATA, and the ADR.

SECTION 15, REGULATORY INFORMATION

AGENCY	LISTING
TSCA	Listed
OSHA	Not Listed
ACGIH	Not Listed
IARC	Not Listed
NTP	Not Known

STATE REGULATIONS

For details on your regulatory requirements you should contact the appropriate agency in your state.

SECTION 16, OTHER INFORMATION

The information above is believed to be accurate and represents the best information currently available to us. However, we make no warranty of merchantability or any other warranty, express or implied, with respect to such information, and we assume no liability resulting from its use. Users should make their own investigations to determine the suitability of the information for their particular purposes. In no event shall ULTRAtab Laboratories Inc.be liable for any claims, losses, or damages of any third party or for lost profits or any special, indirect, incidental, consequential or exemplary damages, howsoever arising, even if ULTRAtab Laboratories Inc., has been advised of the possibility of such damages. This SDS is for a bulk product. This SDS is for the exclusive use of Prestige Packaging Inc.

DEPARTMENT ISSUING SDS: ULTRATAB LABORATORIES INC.
50 TOC DRIVE
HIGHLAND, NY 12528

COMPANY SDS HELPLINE: (845)691-8361

SDS CREATION DATE: April 9, 2015

SDS VERSION: Original