

**MAX RELIEF JUNIOR- acetaminophen liquid**  
**ATLANTIC BIOLOGICALS CORP.**

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**MAX relief junior**  
**Dye-free Children's Pain Reliever and fever reducer**  
**Acetaminophen 160 mg per 5 mL**  
**Alcohol Free, Aspirin Free**  
**For Ages 2 to 11 years**

**Active ingredient (in each 5 mL)**

Acetaminophen 160 mg

**Purpose**

Pain Reliever/Fever Reducer

**Uses**

- temporarily
- reduces fever
- relieves minor aches and pains due to:
  - the common cold
  - flu
  - headache
  - sore throat
  - toothache

**Warnings**

**Liver Warning:** This product contains acetaminophen. Severe liver damage may occur if your child takes:

- more than 5 doses in 24 hours, which is the maximum daily amount
- with other drugs containing acetaminophen

**Allergy alert:** Acetaminophen may cause severe skin reactions.

**Soar throat warning:** if sore throat is severe, persists or more than 2 days, is accompanied or followed by fever headache, rash, nausea, or vomiting, consult a doctor promptly.

**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.

**Ask a doctor before use if your child has** liver disease.

**Ask a doctor or pharmacist before use if your child is** taking the blood thinning drug warfarin.

**When using this product: Do not exceed recommended dose (see overdose warning)**

**Stop use and ask a doctor if**

- pain gets worse or lasts more than 5 days
- fever gets worse or lasts more than 3 days
- new symptoms occur
- redness or swelling is present. These could be signs of a serious condition.

**Keep out of reach of children.**

**If pregnant or breast-feeding,** ask a health professional before use

**Overdose warning.** In case of overdose, get medical help or contact a Poison Control Center (1-800-222-1222) right away. Quick medical attention is critical even if you do not notice any signs or symptoms.

**Directions**

- **this product does not contain direction or complete warnings for adult use**
- **Shake well before using**
- ml = milliliter
- find right dose on chart below
- if possible, use weight to dose; otherwise use age
- use only the enclosed dosing cup designed for use with this product. Do not use any other dosing device.
- if needed, repeat dose every 4 hours while symptoms last
- do not give more than 5 times in 24 hours
- do not give more than 5 days unless directed by doctor.

<b>Weight (lbs.)</b>	<b>Age (yrs.)</b>	<b>Dose (tsp or mL)</b>
under 24	under 2	ask a doctor
24 to 35	2 to 3	1 tsp or 5 mL
36 to 47	4 to 5	1 1/2 tsp or 7.5 mL
48 to 59	6 to 8	2 tsp or 10 mL
60 to 71	9 to 10	2 1/2 tsp or 12.5 mL
72 to 95	11	3 tsp or 15 mL

Other Information store at room temperature 15°-30°C (59°-86°F). Protect from Freezing. Protect from light.

**Inactive ingredients:** amydrrous citric acid, bubble gum flavor, glycerin, polyethylene glycol400, punfied water, saccharin sodium, sodium benzoate, sodium citrate, sorbtol solution, sucralose.

**DISTRIBUTED BY:**

**ATLANTIC BIOLOGICALS CORP.**

**MIAMI, FL 33179**

This product is not manufactured by or distributed by McNeil Consumer Healthcare, owner of the registered trademark Tylenol Eixir.







## Inactive Ingredients

Ingredient Name	Strength
<b>ANHYDROUS CITRIC ACID</b> (UNII: XF417D3PSL)	
<b>GLYCERIN</b> (UNII: PDC6A3C0OX)	
<b>POLYETHYLENE GLYCOL 400</b> (UNII: B697894SGQ)	
<b>PROPYLENE GLYCOL</b> (UNII: 6DC9Q167V3)	
<b>WATER</b> (UNII: 059QF0KO0R)	
<b>SACCHARIN SODIUM</b> (UNII: SB8ZUX40TY)	
<b>SODIUM BENZOATE</b> (UNII: OJ245FE5EU)	
<b>SODIUM CITRATE</b> (UNII: 1Q73Q2JULR)	
<b>SORBITOL</b> (UNII: 506T60A25R)	
<b>SUCRALOSE</b> (UNII: 96K6UQ3ZD4)	

## Product Characteristics

<b>Color</b>		<b>Score</b>	
<b>Shape</b>		<b>Size</b>	
<b>Flavor</b>	BUBBLE GUM	<b>Imprint Code</b>	
<b>Contains</b>			

## Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:17856-0025-1	50 in 1 BOX, UNIT-DOSE	02/28/2024	
1	NDC:17856-0025-5	20.31 mL in 1 CUP, UNIT-DOSE; Type 0: Not a Combination Product		
2	NDC:17856-0025-2	72 in 1 BOX, UNIT-DOSE	02/28/2024	
2	NDC:17856-0025-6	5 mL in 1 CUP, UNIT-DOSE; Type 0: Not a Combination Product		
3	NDC:17856-0025-3	72 in 1 BOX, UNIT-DOSE	02/28/2024	
3	NDC:17856-0025-7	15 mL in 1 CUP, UNIT-DOSE; Type 0: Not a Combination Product		
4	NDC:17856-0025-4	48 in 1 BOX, UNIT-DOSE	02/28/2024	
4	NDC:17856-0025-8	5 mL in 1 SYRINGE; Type 2: Prefilled Drug Delivery Device/System (syringe, patch, etc.)		

## Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M	05/14/2023	

**Labeler** - ATLANTIC BIOLOGICALS CORP. (047437707)

**Registrant** - ATLANTIC BIOLOGICALS CORP. (047437707)

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## Establishment

Name	Address	ID/FEI	Business Operations
UNIT DOSE SOLUTIONS		360804194	repack(17856-0025)

Revised: 2/2024

ATLANTIC BIOLOGICALS CORP.