MAX RELIEF JUNIOR- acetaminophen liquid
ATLANTIC BIOLOGICALS CORP.

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 Paoison 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 hourswhile symptoms last
- do not give more than 5 times in 24 hours
- do not give more than 5 days unless directed by doctor.

Weight (lbs.)	Age (yrs.)	Dose (tsp or mL)
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 15º-30ºC (59º-86ºF). Protect from Freezing. Protect from light.

Inactive ingredients: amydrous citric acid, bubble gum flavor, glycerin, polyethylene glycol400, punfied water, saccharin sodium, sodium benzoate, sodium citrate, sorbttol 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.

17856-0046-01
ACETAMINOPHEN
ORAL ELIXIR
160 mg PER 5 mL
DELIVERS 64mg/2 mL

UDOSE

See package insert for indications and dosage schedule

Bubble Gum Flavor. Alcohol/Aspirin/Dye/Sugar Free. Shake Well Before Using. Store at Room Temperature 15°-30°C(59°-86°F). Protect from freezing. Protect from Light.
Keep this and all Medication out of the reach of children

217856004601

17856-0046-01 **Dosage : 64 mg / 2 mL**

ACETAMINOPHEN **Qty: 120 SYRINGES**

GTIN: 00117856004615
S/N: XXXXXXXXXXX
Exp: 02/27/24
Lot: XXXXXXXXXX

OTC

Packaged by:

Distributed by: Atlantic Biologicals Corp.
Miami, FL 33179

Rev.08/21

Call to Reorder:

MAX RELIEF JUNIOR

acetaminophen liquid

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:17856-0046(NDC:71399-0021)
Route of Administration	ORAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
ACETAMINOPHEN (UNII: 362O9ITL9D) (ACETAMINOPHEN - UNII:362O9ITL9D)	ACETAMINOPHEN	160 mg in 5 mL

Inactive Ingredients

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

Product Characteristics

Color		**Score**	
Shape		**Size**	
Flavor	BUBBLE GUM	**Imprint Code**	
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:17856-0046-1	120 in 1 BOX, UNIT-DOSE	02/28/2024	
1	NDC:17856-0046-2	2 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)

Establishment

Name	Address	ID/FEI	Business Operations
UNIT DOSE SOLUTION		360804194	repack(17856-0046)

Revised: 2/2024 ATLANTIC BIOLOGICALS CORP.