# 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 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 15°-30°C (59°-86°F). 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

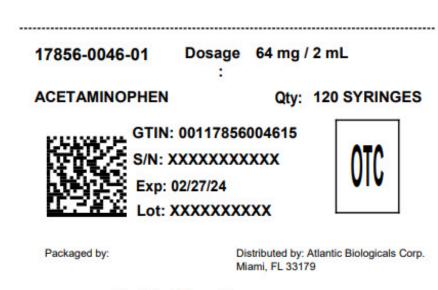


See package insert for indications and dosage schedule

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\*\*



Rev.08/21 Call to Reorder:

MAX RELIEF JUNIOR acetaminophen liquid				
<b>Product Information</b>				
Product Type	HUMAN OTC DRUG	Item Code (Sourc	e) NDC:17856-0046(N	DC:71399-0021)
Route of Administration	ORAL			
Active Ingredient/Active	Moiety			
Ingree	lient Name		<b>Basis of Strength</b>	Strength
ACETAMINOPHEN (UNII: 36209ITL	9D) (ACETAMINOPHE	N - UNII:36209ITL9D)	ACETAMINOPHEN	160 mg in 5 mL

Shape Size				Ingredient Name				Strength
PROLYETHYLENE GLYCOL 400 (UNII: 6bG78945GQ)       Image: Section of the	AN	IHYDROUS C	ITRIC ACI	D (UNII: XF417D3PSL)				
PROPYLENE GLYCOL (UNII: 6DC9Q167V3) WATER (UNII: 059QF0KO0R) SACCHARIN SODIUM (UNII: SB8ZUX40TY) SODIUM BENZOATE (UNII: 0245FESEU) SODIUM CITRATE (UNII: 1Q73Q2JULR) SORBITOL (UNII: 506T60A25R) SUCRALOSE (UNII: 96K6UQ3Z D4) Product Characteristics Color Shape Size I Flavor BUBBLE GUM Imprint Code Contains I Package Description Marketing Start Date Packaging Marketing Information Marketing Information Number or Monograph Marketing Start Marketing End	GL	YCERIN (UNI	: PDC6A3C	COOX)				
WATER (UNII: 059QF0K00R)       SACCHARIN SODIUM (UNII: SB8Z UX40TY)       SACCHARIN SODIUM (UNII: 0245FE5EU)         SODIUM BENZOATE (UNII: 1073Q2JULR)       SOBITOL (UNII: 506T60A25R)       SSOBITOL (UNII: 506T60A25R)         SUCRALOSE (UNII: 96K6UQ3Z D4)       STOR       SSOB         Product Characteristics         Color       Score         Size         Flavor       BUBBLE GUM         Imprint Code         Packaging         Marketing Start Date         Marketing Information         Marketing Information         Marketing       Application Number or Monograph       Marketing Marketing Marketing End	PC	LYETHYLEN	E GLYCOL	<b>400</b> (UNII: B697894SGQ)				
SACCHARIN SODIUM (UNII: SB8ZUX40TY) SODIUM BENZOATE (UNII: 0)245FE5EU) SODIUM CITRATE (UNII: 1073Q2JULR) SORBITOL (UNII: 506T60A25R) SUCRALOSE (UNII: 96K6UQ3ZD4) Product Characteristics Color Score Shape BUBBLE GUM Imprint Code Flavor BUBBLE GUM Imprint Code Contains Marketing Start Date Package Description Marketing Start Marketing End Date Marketing Information Marketing Information Marketing Start Marketing End	PR	OPYLENE GI	YCOL (UN	III: 6DC9Q167V3)				
SODIUM BENZOATE (UNII: 0)245FE5EU) SODIUM CITRATE (UNII: 1Q73Q2JULR) SORBITOL (UNII: 506T60A25R) SUCRALOSE (UNII: 96K6UQ3Z D4) Product Characteristics Color Shape BUBBLE GUM BUBBLE G	N	ATER (UNII: 0	59QF0KO0	R)				
SODIUM CITRATE (UNII: 1Q73Q2JULR) SORBITOL (UNII: 506T60A25R) SUCRALOSE (UNII: 96K6UQ3ZD4) Product Characteristics Color Shape Size I In SUBBLE GUM Imprint Code Size I In Subble GUM Imprint Code I In Subble GUM Imprint Code Packaging I In Subble GUM Imprint Code I In Subble GUM Imprint GU	SA	CCHARIN SC	DIUM (UN	III: SB8ZUX40TY)				
SORBITOL (UNII: 506T60A25R)   SUCRALOSE (UNII: 96K6UQ3ZD4)   Product Characteristics   Color   Shape   Size   Shape   Flavor   BUBBLE GUM   Imprint Code   Pockaging   #   Item   Code   1   NDC:17856-   2mL in 1 SYRINGE; Type 2: Prefilled Drug Delivery   0046-2   2mL in 1 SYRINGE; Type 2: Prefilled Drug Delivery   0046-2   Marketing Information   Marketing Application Number or Monograph Marketing Start Marketing End	5C	DIUM BENZ	OATE (UNI	I: OJ245FE5EU)				
SUCRALOSE (UNII: 96K6UQ3ZD4)  Product Characteristics  Color Shape Size Size Size Contains BUBBLE GUM Imprint Code  Marketing Start Date Marketing Code 1 NDC:17856 120 in 1 BOX, UNIT-DOSE 1 NDC:17856 120 in 1 BOX, UNIT-DOSE 1 NDC:17856 2 mL in 1 SYRINGE; Type 2: Prefilled Drug Delivery Device/System (syringe, patch, etc.)  Marketing Marketing Application Number or Monograph Marketing Start Marketing Start Marketing Start Marketing Start Marketing Marke	5C	DIUM CITRA	TE (UNII: 1	LQ73Q2JULR)				
Product Characteristics         Score         Size         Size         BUBBLE GUM         Imprint Code         Contains         Marketing         Marketing Marketing End Date         Package Description       Marketing Marketing End Date         Marketing Information         Marketing Information Number or Monograph       Marketing Start       Marketing End	5C	RBITOL (UNI	I: 506T60A	25R)				
Score         Size         Shape       Size         Flavor       BUBBLE GUM       Imprint Code         Contains         Packaging         #       Marketing Code       Marketing Marketing End Date         #       Item Code       Package Description       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.)       02/28/2024	SU	ICRALOSE (U	NII: 96K6U	Q3ZD4)				
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Implication       Marketing         Backaging       Package Description       Marketing       Marketing         Implication       Marketing       Marketing       Marketing         Implication       Marketing       Marketing       Marketing         Implication       Marketing       Marketing       Marketing         Implication       Marketing       Marketing <tht< td=""><td colspan="3">Shape</td><td></td><td>Siz</td><td>е</td><td></td><td></td></tht<>	Shape				Siz	е		
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Item Code       Package Description       Marketing Start Date       Marketing End Date         1       NDC:17856- 0046-1       120 in 1 BOX, UNIT-DOSE       02/28/2024       02/28/2024         1       NDC:17856- 0046-2       2 mL in 1 SYRINGE; Type 2: Prefilled Drug Delivery Device/System (syringe, patch, etc.)       02/28/2024       Image: Comparison of the system of the syst	Co	ontains						
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Marketing Application Number or Monograph Marketing Start Marketing End	T	0046-1		DX, UNIT-DOSE			02/28/2024	
Marketing Application Number or Monograph Marketing Start Marketing End	1	NDC:17856- 0046-2	2 mL in 1 Device/Sy					
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Labeler - ATLANTIC BIOLOGICALS CORP. (047437707)

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

## Establishment

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

Revised: 2/2024

ATLANTIC BIOLOGICALS CORP.