

**FEVERALL CHILDRENS- acetaminophen suppository**  
**Sun Pharmaceutical Industries, Inc.**

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**Feverall**®  
**Children's**

***Drug Facts***

**Active ingredient (in each rectal suppository)**

Acetaminophen, USP 120 mg

**Purposes**

Pain reliever/fever reducer

**Uses**

temporarily

- reduces fever
- relieves minor aches, pains, and headache

**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. Symptoms may include:

- skin reddening
- blisters
- rash

If a skin reaction occurs, stop use and seek medical help right away.

**For rectal use only.**

**Do not use**

- if you are allergic to acetaminophen.
- 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**

- the child has liver disease.
- the child is taking the blood thinning drug warfarin.

**Stop use and ask a doctor if**

- fever lasts more than 3 days (72 hours), or recurs.
- pain lasts more than 5 days or gets worse.
- new symptoms occur.
- redness or swelling is present in the painful area.

These may be signs of a serious condition.

**Keep out of reach of children.** If swallowed or in case of overdose, get medical help or contact a Poison Control Center right away. Quick medical attention is critical in case of overdose for adults and for children even if you do not notice any signs or symptoms.

**Directions**

- **this product does not contain directions or warnings for adult use**
- **do not use more than directed**
- remove wrapper
- carefully insert suppository well up into the rectum

**Dosing Chart**

<b>Age</b>	<b>Dose</b>
under 3 years	Do not use unless directed by a doctor
3 to 6 years	Use 1 suppository every 4 to 6 hours (maximum of 5 doses in 24 hours)

**Other information**

- store at 2°-27°C (35°-80°F)
- do not use if imprinted suppository wrapper is opened or damaged

**Inactive ingredients**

glycerol monostearate, hydrogenated vegetable oil, polyoxyethylene stearate, polysorbate 80

**Questions?**

**call 1-866-923-4914**

Dist. by: **Taro Pharmaceuticals U.S.A., Inc.**  
Hawthorne, NY 10532

**PRINCIPAL DISPLAY PANEL - 120 mg Suppository Blister Pack Carton**

NDC 51672-2115-2

RECTAL  
SUPPOSITORY\*

\*actual size

Doctor Recommended

Pain Reliever/Fever Reducer

- No Parabens or Any Other Preservatives
- No Artificial Colors

CHILDREN'S

ages 3-6 years

FeverAll®

ACETAMINOPHEN SUPPOSITORIES

Pain Reliever/Fever Reducer

6

Rectal

Suppositories

120

mg

each

NDC 51672-2115-2



\*actual size

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CHILDREN'S  
ages 3-6 years

# FeverAll®

ACETAMINOPHEN SUPPOSITORIES

Pain Reliever/Fever Reducer



6 Rectal  
Suppositories

120 mg  
each



Dist. by: Taro Pharmaceuticals U.S.A., Inc.  
Hawthorne, NY 10532 Made in USA  
GW 710011-TBD

Tamper-evident: Suppositories are individually wrapped.  
Do not use if imprinted wrapper is opened or damaged.

**Other information**

- store at 20-27°C (35°-80°F)
- do not use if imprinted suppository wrapper is opened or damaged

**Inactive ingredients**

glycerol monostearate, hydrogenated vegetable oil, polyoxyethylene stearate, polyorbate 80

**Questions?** call 1-866-923-4914

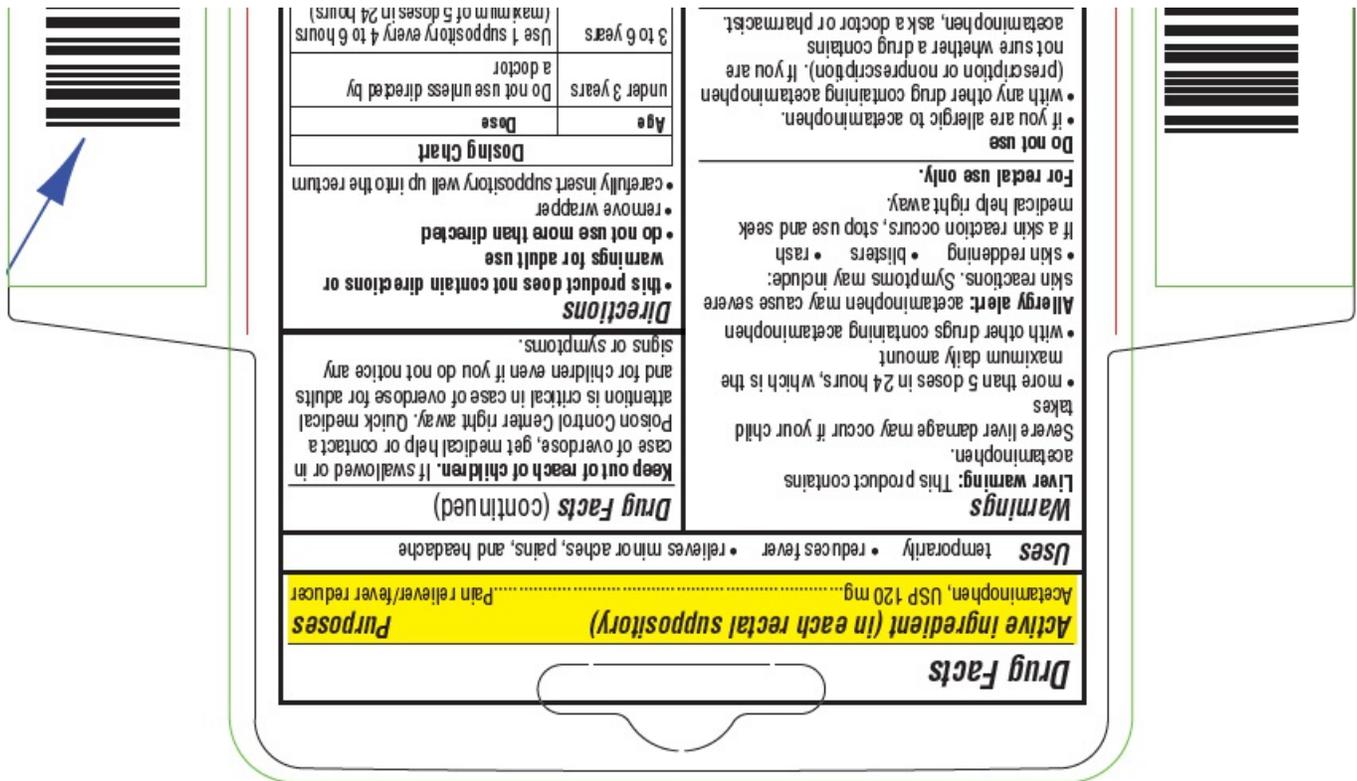
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- new symptoms occur.
- redness or swelling is present in the painful area.

These may be signs of a serious condition.



## FEVERALL CHILDRENS

acetaminophen suppository

### Product Information

<b>Product Type</b>	HUMAN OTC DRUG	<b>Item Code (Source)</b>	NDC:51672-2115
<b>Route of Administration</b>	RECTAL		

### Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
<b>ACETAMINOPHEN</b> (UNII: 362O9ITL9D) (ACETAMINOPHEN - UNII:362O9ITL9D)	ACETAMINOPHEN	120 mg

### Inactive Ingredients

Ingredient Name	Strength
<b>GLYCERYL MONOSTEARATE</b> (UNII: 230OU9XXE4)	
<b>HYDROGENATED PALM KERNEL OIL</b> (UNII: FM8D1RE2VP)	
<b>PEG-100 STEARATE</b> (UNII: YD01N1999R)	
<b>POLYSORBATE 80</b> (UNII: 6OZP39ZG8H)	

### Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:51672-2115-2	6 in 1 CARTON	12/12/2013	

<b>1</b>	NDC:51672-2115-0	1 in 1 BLISTER PACK; Type 0: Not a Combination Product		
<b>2</b>	NDC:51672-2115-4	50 in 1 CARTON	12/12/2013	
<b>2</b>	NDC:51672-2115-0	1 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
NDA	NDA018337	12/12/2013	

**Labeler** - Sun Pharmaceutical Industries, Inc. (146974886)

## Establishment

Name	Address	ID/FEI	Business Operations
G&W NC Laboratories, Inc.		079419931	manufacture(51672-2115)

Revised: 2/2026

Sun Pharmaceutical Industries, Inc.