

SOOTHEEZ ORGANIC WILD CHERRY- pectin lozenge
BFY LLC

Sootheez Organic Wild Cherry

Sootheez Wild Cherry - Organic 25CT 3714-2

Drug Facts

Active Ingredient (in each drop)

Pectin 11.5 mg

Purpose

Oral Demulcent

Uses

for temporary relief of minor discomfort of the throat.

Warnings

Sore throat warning: if sore throat is severe, persists for more than 2 days, is accompanied or followed by fever, headache, rash, nausea, or vomiting, consult a doctor promptly. If sore mouth symptoms do not improve in 7 days, see your dentist or doctor promptly.

Stop use and ask a doctor if

- sore throat does not improve in 7 days.
- irritation, pain, or redness persists or worsens.

Keep out of reach of children.

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Directions

- **Adults and children 10 years of age and older:** allow one drop to dissolve slowly in mouth.
- **May be repeated as needed** or as directed by a doctor.
- **Children under 10 years of age:** ask a doctor.

Other information

store in a dry place between 65°-75°F with pouch sealed.

Inactive ingredients

citric acid, organic cane sugar, organic cherry juice concentrate, organic honey, organic natural flavor, organic rice syrup, organic strawberry juice concentrate, organic

strawberry powder, organic tart cherry powder.

Questions?

Visit our website at www.theEEZco.com, or write us at 450 Heritage Rd., Southbury, CT 06488.

Principal Display Panel





SOOTHEEZ ORGANIC WILD CHERRY

pectin lozenge

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:72427-3714
Route of Administration	ORAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
PECTIN (UNII: 89NA02M4RX) (PECTIN - UNII:89NA02M4RX)	PECTIN	11.5 mg in 3.5 g

Inactive Ingredients

Ingredient Name	Strength
HONEY (UNII: Y9H1V576FH)	

Product Characteristics

Color	red	Score	no score
Shape	OVAL	Size	23mm
Flavor	CHERRY (Wild Cherry)	Imprint Code	P
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:72427-3714-2	88 g in 1 POUCH; Type 0: Not a Combination Product	01/06/2026	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M022	01/06/2026	

Labeler - BFY LLC (080651312)

Revised: 1/2026

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