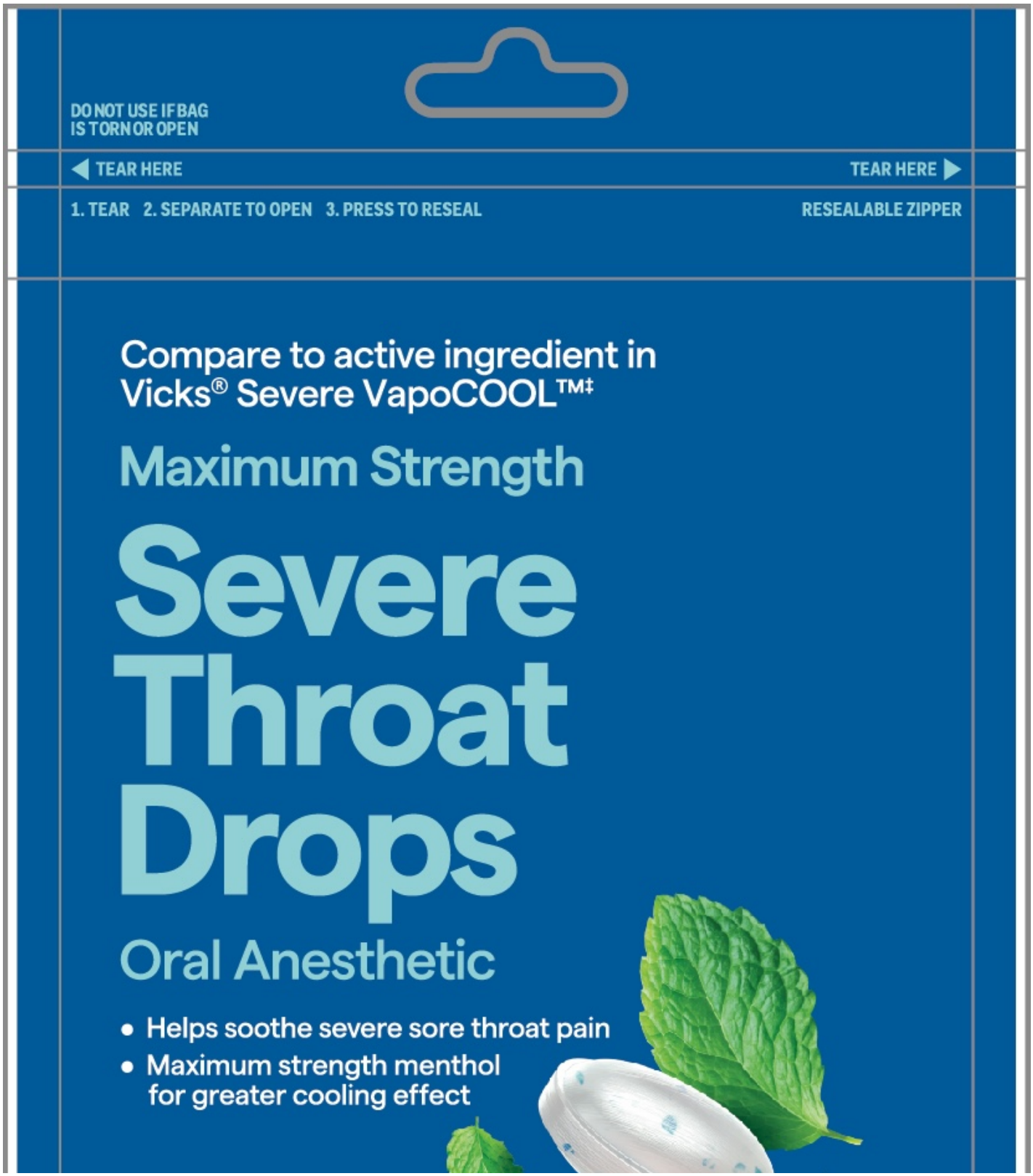
SEVERE THROAT DROPS- menthol ice pastille
Target Corporation Inc.

Severe Throat Drops

Active ingredient 20 mg

Oral anesthetic



Uses temporarily relieves ocassional minor irritation and pain due toL
- Sore throat
- Sore mouth

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Warnings

Sore throat warning: severe or persistence sore throat accompanied by high fever, headache, nausea and vomiting may be serious. Consult a doctor promptly. Do not use more than 2 days or administer to children under 12 years of age.

Ask a doctor before use if you have
- a severe throat accompanied by difficulty in breathing or that last more than 2 days
- a sore throat accompanied by fever, headache, rash, swelling, nausea or vomiting

Stop use and consult a doctor if
- sore mouth symptons do not improve in 7 days or if irritation, pains or redness persists or worsens.

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

Keep this and all drugs out of the reach of children.

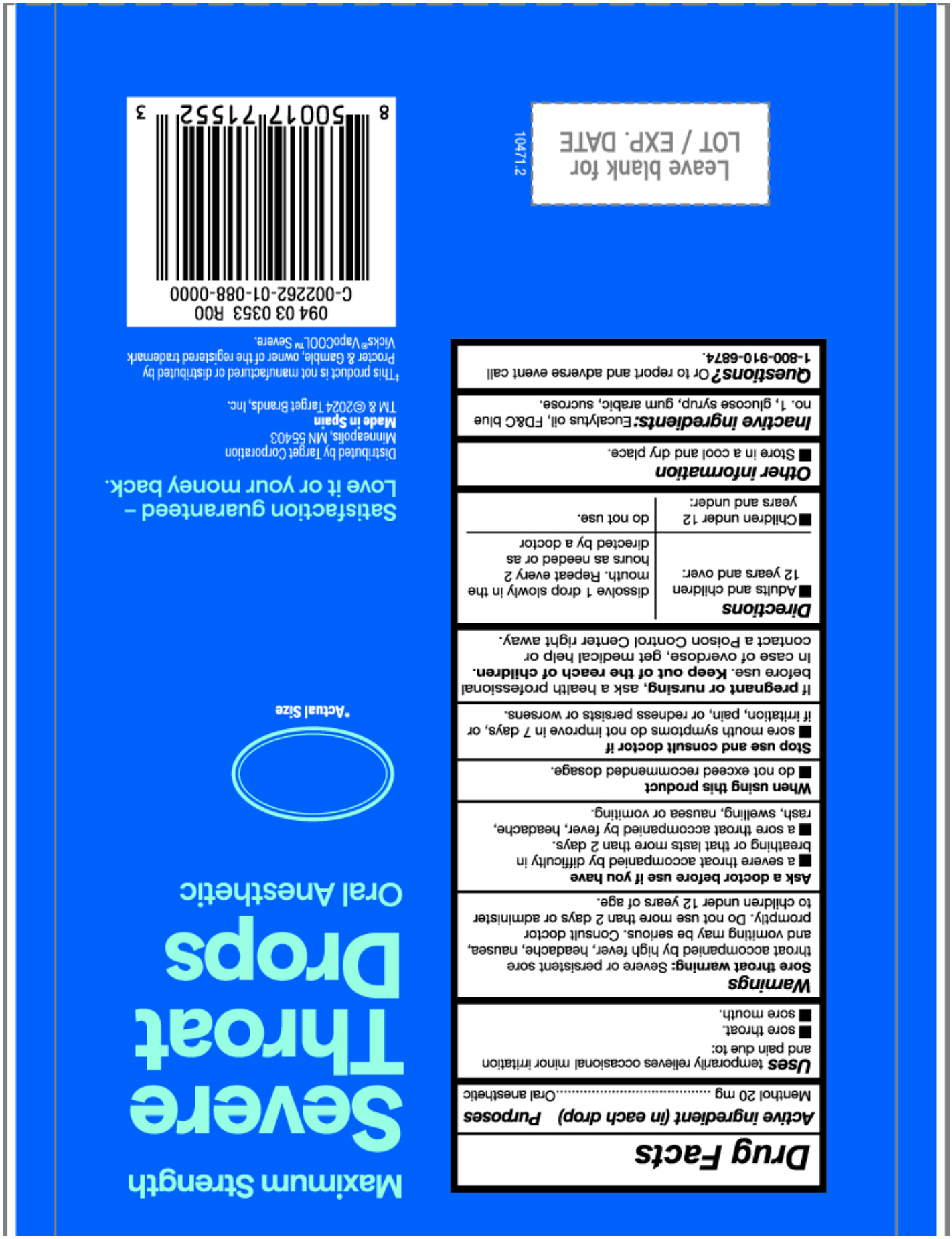
In case of accidental overdose, get medical help or contact a Poison Control Center right away.

Directions
- Adults and children 12 years and over: dissolve 1 drop slowly in mouth. Repeated every 2 hours as needed or as directed by a doctor.
- Children under 12 years and under do not use

- Store in a cool and dry place

eucalytus oil, FD&C blue No.1, glucose syrup, gum arabic, sucrose

Questions? or to report and adverse event call: 1-800-910-68-74



SEVERE THROAT DROPS

menthol ice pastille

Product Information

Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:82442-201
Route of Administration	ORAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
MENTHOL (UNII: L7T10EIP3A) (MENTHOL - UNII:L7T10EIP3A)	MENTHOL	20 mg

Inactive Ingredients

Ingredient Name	Strength
EUCALYPTOL (UNII: RV6J6604TK)	
FD&C BLUE NO. 1 (UNII: H3R47K3TBD)	
ACACIA (UNII: 5C5403N26O)	
SUCROSE (UNII: C151H8M554)	
CORN SYRUP (UNII: 9G5L16BK6N)	

Product Characteristics

Color	white	**Score**	score with uneven pieces
Shape	OVAL	**Size**	25mm
Flavor	MENTHOL	**Imprint Code**	None
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:82442-201-01	45 in 1 BAG; Type 0: Not a Combination Product	05/15/2024	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M012	05/15/2024	

Labeler - Target Corporation Inc. (006961700)

Registrant - Boston Nutraceutical Science SL (466061824)

Establishment

Name	Address	ID/FEI	Business Operations
Boston Nutraceutical Production SL		468121064	manufacture(82442-201) , label(82442-201) , pack(82442-201)

Revised: 7/2024 Target Corporation Inc.