

## **KIDS ORAL CARE VANILLA FLAVOR- xylitol, calcium lactate spray TOOTHFILM INC.**

*Disclaimer: This drug has not been found by FDA to be safe and effective, and this labeling has not been approved by FDA. For further information about unapproved drugs, click here.*

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**Kids Oral Care Vanilla Flavor**

Active ingredients: Xylitol 2.0mL/20mL; Calcium Lactate 0.02mL/20mL

Inactive ingredients: Water, Glycerin, Sodium Levulinate, Sodium Anisate, Trehalose, Sodium Chloride, Dicalcium Phosphate, Silica, Calcium Oxide

Purpose: Anticavity

keep out of reach of children: For external use only. Keep out of reach of children. Avoid contact with eyes. Discontinue use if signs of irritation or rash occur.

Spray after meals and feedings. Don't forget to brush kid's teeth for the best results! It's fluoride-free, preservative-free, and safe to swallow when used as directed!

Apply the spray on the kiddo's teeth surface to form a protective coating. • Do not spit the spray out. • Do not rinse with water after spray. This spray is not intend to replace brushing or flossing.

Warnings: Avoid spraying in eyes. In case of contact, rinse eyes thoroughly with water. Stop use and ask a doctor or pharmacist if you have an allergic reaction to this product or any of its ingredients.

Helps protect against cavities. Spray 2 to 3 times towards children's teeth to prevent plaque build up and cavities.

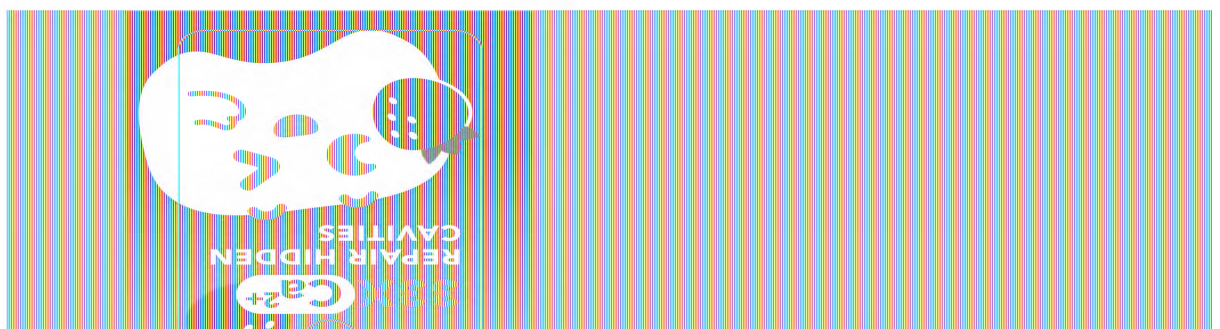
Question: support@lab52.com

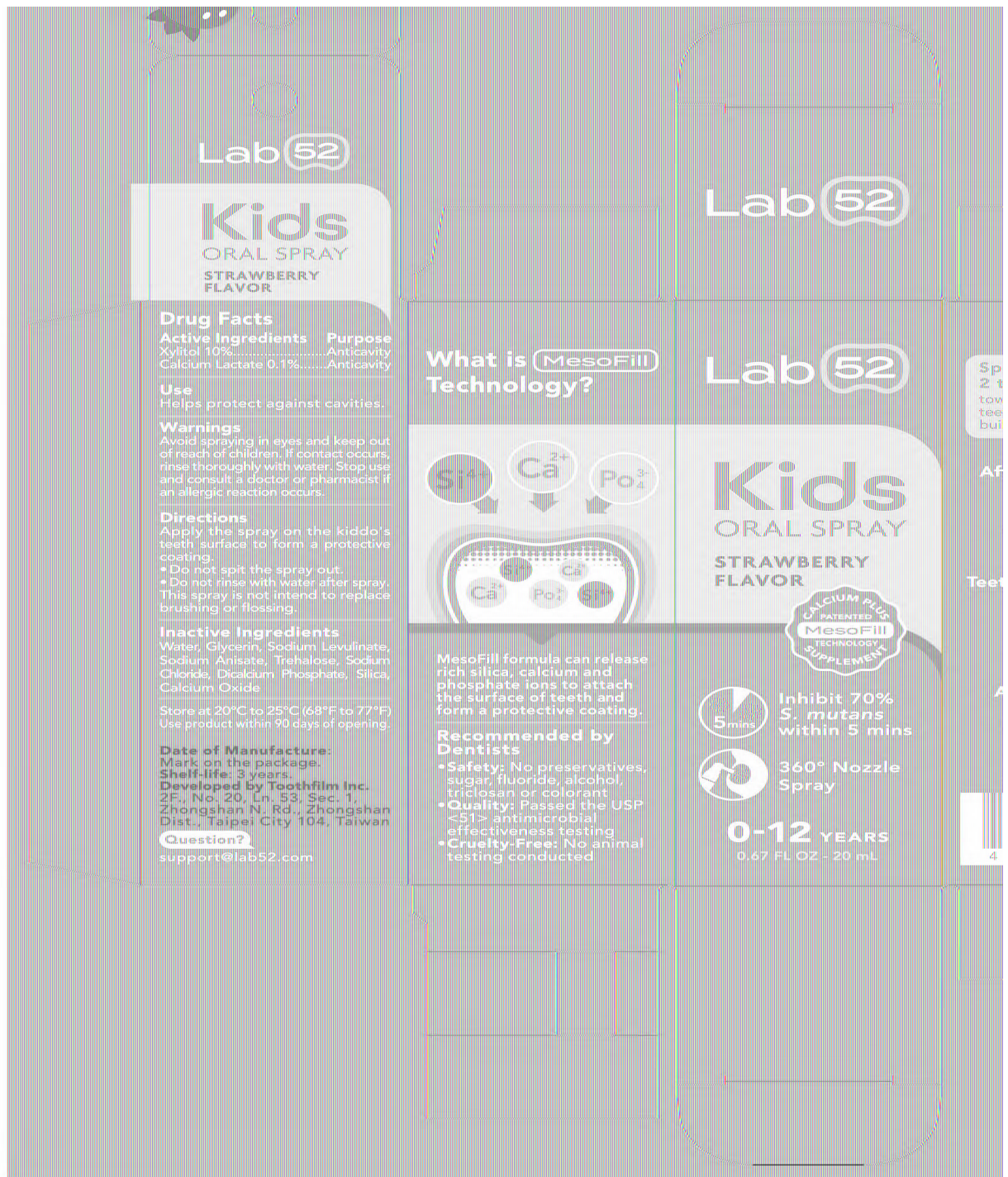
Other: Store at 20°C to 25°C (68°F to 77°F) Use product within 90 days of opening.

齒妍堂\_英文版兒童噴霧外盒\_草莓



877C 1915C 局部光





See Label

## KIDS ORAL CARE VANILLA FLAVOR

xylitol, calcium lactate spray

### Product Information

<b>Product Type</b>	HUMAN OTC DRUG	<b>Item Code (Source)</b>	NDC:82711-103
<b>Route of Administration</b>	DENTAL		

**Active Ingredient/Active Moiety**

Ingredient Name	Basis of Strength	Strength
<b>CALCIUM LACTATE</b> (UNII: 2URQ2N32W3) (CALCIUM CATION - UNII:2M83C4R6ZB)	CALCIUM LACTATE	0.02 mg in 20 mg
<b>XYLITOL</b> (UNII: VCQ006KQ1E) (XYLITOL - UNII:VCQ006KQ1E)	XYLITOL	2 mg in 20 mg

**Inactive Ingredients**

Ingredient Name	Strength
<b>SODIUM ANISATE</b> (UNII: F9WFJ28MV9)	
<b>SODIUM LEVULINATE</b> (UNII: VK44E1MQU8)	
<b>SODIUM CHLORIDE</b> (UNII: 451W47IQ8X)	
<b>CALCIUM OXIDE</b> (UNII: C7X2M0VVNH)	
<b>GLYCERIN</b> (UNII: PDC6A3C0OX)	
<b>WATER</b> (UNII: 059QF0KO0R)	
<b>SILICA</b> (UNII: ETJ7Z6XBU4)	
<b>DICALCIUM PHOSPHATE</b> (UNII: L11K75P92J)	
<b>TREHALOSE</b> (UNII: B8WCK70T7I)	

**Packaging**

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:82711-103-02	1 in 1 BOX	03/13/2025	
1	NDC:82711-103-01	20 mg in 1 BOTTLE; Type 0: Not a Combination Product		

**Marketing Information**

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
unapproved drug other		03/13/2025	

**Labeler** - TOOTHFILM INC. (656036549)**Establishment**

Name	Address	ID/FEI	Business Operations
JUN DA BEAUTY-TEC CO., LTD.		658442561	manufacture(82711-103) , label(82711-103)

Revised: 3/2025

TOOTHFILM INC.