ANTI NAUSEA- dextrose, fructose and phosphoric acid liquid
H E B

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.

HEB Anti-Nausea Drug Facts

Active ingredients (per 5 mL)

Dextrose (glucose) 1.87 g

Levulose (fructose) 1.87 g

Phosphoric acid 21.5 mg

Purpose

Upset stomach reliever

Uses

- relieves nausea due to upset stomach from intestinal flu, stomach flu, and food or drink indiscretions

Warnings

- this product contains fructose and should not be taken by persons with hereditary fructose intolerance (HFI)

Do not use if you have

- allergic reactions to any of the ingredients in this product

Ask a doctor before use if you have

- diabetes

Stop use and ask a doctor if

- symptoms persist, return or get worse

If pregnant or breast-feeding,

ask a health professional before use.

Keep out of reach of children.

In case of overdose, get medical help or contact a Poison Control Center right away. (1-800-222-1222)

Directions

- for maximum effectiveness never dilute or drink fluids of any kind immediately before or after

taking this product

- adults and children 12 years of age and over: one or two tablespoons
- children 2 to under 12: one or two teaspoons
- repeat dose every 15 minutes or until distress subsides
- do not take more than 5 doses in 1 hour without consulting a doctor.

Other information

- store at 20°-25°C (68°-77°F)

Inactive ingredients

FD&C red no. 40 aluminum lake, flavor, glycerin, methylparaben, purified water

Questions or comments?

1-800-719-9260

Package/Label Principal Display Panel

Compare to Emetrol® active ingredients

ANTI-NAUSEA

For Children and Adults

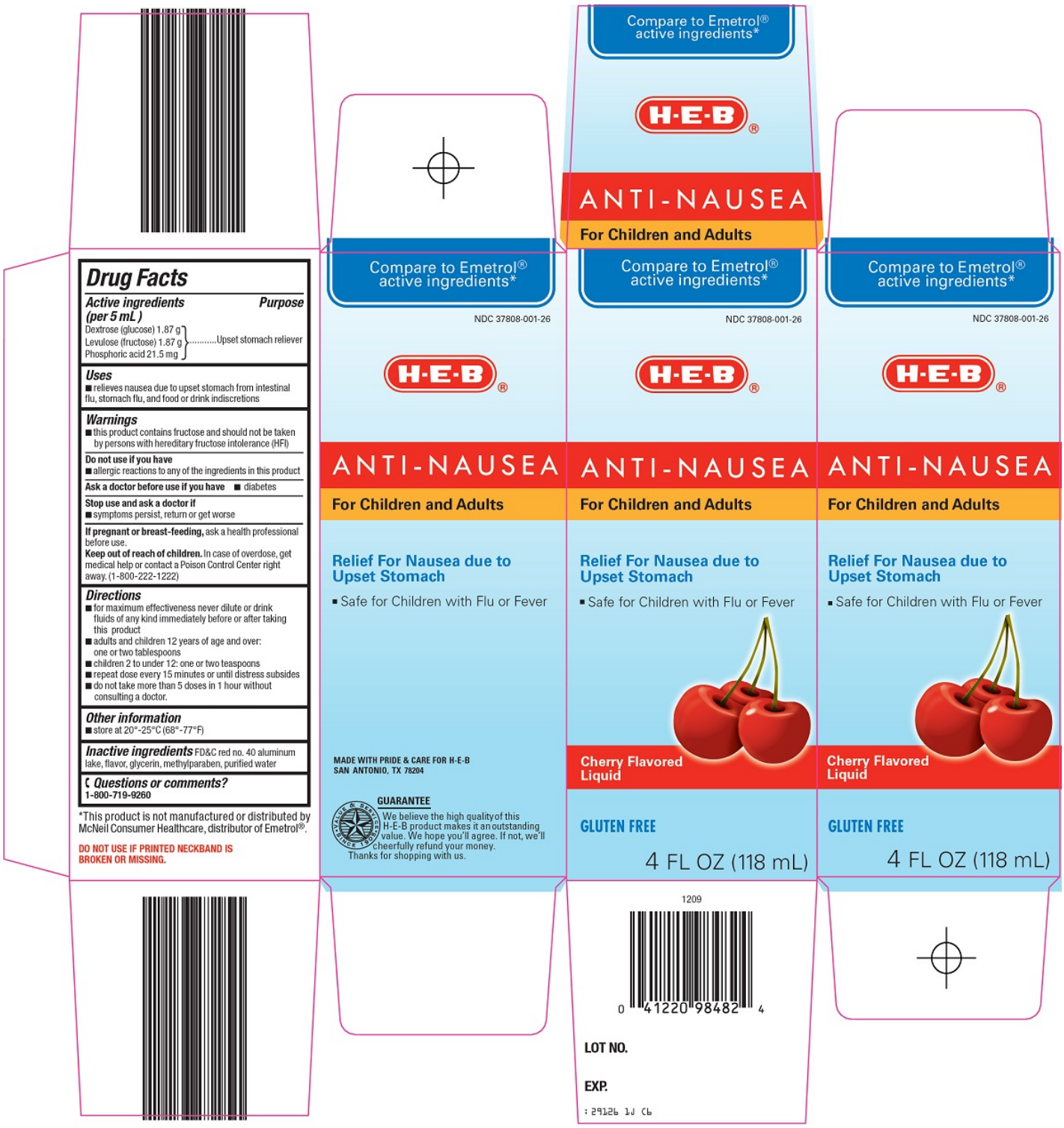
Relief For Nausea due to Upset Stomach

Safe for Children with Flu or Fever

Cherry Flavored

Liquid

GLUTEN FREE



ANTI NAUSEA

dextrose (glucose), levulose (fructose), phosphoric acid liquid

Product Information			
Product Type	HUMAN OTC DRUG LABEL	**Item Code (Source)**	NDC:37808-001
Route of Administration	ORAL	**DEA Schedule**	

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
DEXTROSE (DEXTROSE)	DEXTROSE	1.87 g in 5 mL
FRUCTOSE (FRUCTOSE)	FRUCTOSE	1.87 g in 5 mL
PHOSPHORIC ACID (PHOSPHORIC ACID)	PHOSPHORIC ACID	21.5 mg in 5 mL

Inactive Ingredients

Ingredient Name	Strength
GLYCERIN	
METHYLPARABEN	
WATER	

Product Characteristics

Color	RED (clear)	**Score**	
Shape		**Size**	
Flavor	CHERRY	**Imprint Code**	
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:37808-001-26	1 in 1 CARTON		
1		118 mL in 1 BOTTLE		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
Unapproved drug other		11/22/1995	

Labeler - HEB (007924756)

Revised: 2/2013 HEB