

**EZ FOAM 6793- triclosan soap
OSCEOLA SUPPLY, INC.**

Disclaimer: Most OTC drugs are not reviewed and approved by FDA, however they may be marketed if they comply with applicable regulations and policies. FDA has not evaluated whether this product complies.

EZ Foam 6793 Drug Facts and Label

Drug Facts Box OTC-Active Ingredient Section

Triclosan 0.3%

Drug Facts Box OTC-Purpose Section

Antiseptic

Drug Facts Box OTC-Indications & Usage Section

for hand-washing to decrease bacteria on the skin

Drug Facts Box OTC-Warnings Section

For external use only

Drug Facts Box OTC-When Using Section

do not get into eyes

if contact occurs, rinse eyes thoroughly with water

Drug Facts Box OTC-Stop Use Section

irritation and redness develop

Drug Facts Box OTC-Keep Out of Reach of Children Section

if swallowed, get medical help or contact a Poison Control Center right away

Drug Facts Box OTC-Dosage & Administration Section

wet hands and forearms

apply 5 milliliters (teaspoonful) or palmful to hands and forearms

scrub thoroughly for 1 minute and rinse

Drug Facts Box OTC-Inactive Ingredient Section

water, propylene glycol, alcohol denat., disodium cocoamphodipropionate, lauric acid, ethanolamine, lactic acid, isopropyl alcohol, tetrasodium EDTA, polyquaternium 10, PEG-4, fragrance, methylparaben, propylparaben, FDC red 4, acid red 1

OSCEOLA

EZ Foam

High Foaming Antiseptic Hand Wash

Drug Facts

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Triclosan 0.3%.....	Antiseptic

Use for hand-washing to decrease bacteria on the skin

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Directions ■ wet hands and forearms ■ apply 5 milliliters (teaspoonful) or palmful to hands and forearms ■ scrub thoroughly for 1 minute and rinse

Inactive Ingredients Water, Propylene Glycol, Alcohol denat., Disodium Cocoamphodipropionate, Lauric Acid, Ethanolamine, Lactic Acid, Isopropyl Alcohol, Tetrasodium EDTA, Polyquaternium-10, PEG-4, Fragrance, Methylparaben, Propylparaben, FD&C Red #4, Acid Red 1

Re-Order # 4607F-6-1000

SOLD BY: OSCEOLA SUPPLY

P.O. Box 13503, Tallahassee, FL 32317

Ph.: (850) 580-9800 • www.osceolasupply.com

PACKAGED FOR USE IN A WALL MOUNT DISPENSER

NET CONTENTS: 1000 ML (33.8 FL. OZ.)

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6793P6L513 6.041817

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6793P6L513.6.041817

EZ Foam 6793 1000mL

EZ FOAM 6793

triclosan soap

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:62672-793
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
TRICLOSAN (UNII: 4NM5039Y5X) (TRICLOSAN - UNII:4NM5039Y5X)	TRICLOSAN	3.0 mg in 1 mL

Inactive Ingredients

Ingredient Name	Strength
WATER (UNII: 059QF0KO0R)	
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)	
ALCOHOL (UNII: 3K9958V90M)	
LAURIC ACID (UNII: 1160N9NU9U)	
MONOETHANOLAMINE (UNII: 5KV86114PT)	
LACTIC ACID (UNII: 33X04XA5AT)	
ISOPROPYL ALCOHOL (UNII: ND2M416302)	
EDETATE SODIUM (UNII: MP1J8420LU)	
POLYQUATERNIUM-10 (400 CPS AT 2%) (UNII: HB1401PQFS)	
POLYETHYLENE GLYCOL 200 (UNII: R95B8J264J)	
METHYLPARABEN (UNII: A218C7H9T)	
PROPYLPARABEN (UNII: Z8IX2SC1OH)	
FD&C RED NO. 4 (UNII: X3W0AMIJLX)	
DISODIUM COCOAMPHODIPROPIONATE (UNII: 6K8PRP397M)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:62672-793-06	1 in 1 BOX	04/18/2017	
1		800 mL in 1 BAG; Type 0: Not a Combination Product		
2	NDC:62672-793-08	1 in 1 BOX	04/18/2017	
2		1000 mL in 1 BAG; Type 0: Not a Combination Product		
3	NDC:62672-793-17	532 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	04/18/2017	
4	NDC:62672-793-24	118 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	04/18/2017	
5	NDC:62672-793-01	1200 mL in 1 CARTRIDGE; Type 0: Not a Combination Product	04/18/2017	
6	NDC:62672-793-03	350 mL in 1 CARTRIDGE; Type 0: Not a Combination Product	04/18/2017	
7	NDC:62672-793-05	540 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	04/18/2017	
8	NDC:62672-793-07	700 mL in 1 BAG; Type 0: Not a Combination Product	04/18/2017	
9	NDC:62672-793-09	2000 mL in 1 CARTRIDGE; Type 0: Not a Combination Product	04/18/2017	
10	NDC:62672-793-10	1000 mL in 1 CARTRIDGE; Type 0: Not a Combination Product	04/18/2017	
11	NDC:62672-793-11	1000 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	04/18/2017	
12	NDC:62672-793-12	1000 mL in 1 BAG; Type 0: Not a Combination Product	04/18/2017	
13	NDC:62672-793-13	800 mL in 1 BAG; Type 0: Not a Combination Product	04/18/2017	
14	NDC:62672-793-14	3785 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	04/18/2017	
15	NDC:62672-793-15	946 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	04/18/2017	

16	NDC:62672-793-28	149 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	04/18/2017	
17	NDC:62672-793-27	800 mL in 1 CARTRIDGE; Type 0: Not a Combination Product	04/18/2017	
18	NDC:62672-793-55	208200 mL in 1 DRUM; Type 0: Not a Combination Product	04/18/2017	
19	NDC:62672-793-16	236 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	04/18/2017	
20	NDC:62672-793-18	50 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	04/18/2017	
21	NDC:62672-793-31	750 mL in 1 BAG; Type 0: Not a Combination Product	04/18/2017	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333E	04/18/2017	

Labeler - OSCEOLA SUPPLY, INC. (809050479)

Registrant - ABC Compounding Co., Inc. (003284353)

Establishment

Name	Address	ID/FEI	Business Operations
ABC Compounding Co., Inc.		003284353	manufacture(62672-793)

Revised: 4/2017

OSCEOLA SUPPLY, INC.