MEDI-WIPES- ethyl alcohol, chloroxylenol swab
Afassco 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.

Drug Facts Active Ingredients

Ethyl Alcohol 40.0%

Chloroxylenol (PCMX) 0.5%

Purpose

Antiseptic

Uses:

For hand washing to decrease bacteria on skin when soap and water are not available.

Warnings:

For external use only.

Flammable. Keep away from fire or flame

Directions:

- tear packet open, remove towelette
- wipe hands/wrist areas for 15 seconds and discard

When Using The Product:

- do not get into eyes. If contact occurs rinse eye thoroughly with water

Stop Use And Ask Doctor If:

- irritation or redness develop and persist for more than 72 hours

Inactive Ingredients:

aloe vera, fragrance, purified water

Keep Out Of Reach Of Children

Keep Out Of Reach Of Children:

If swallowed, get medical help or contact a Poison Control center right away. 1-800-222-1222

Product Label Medi-Wipes

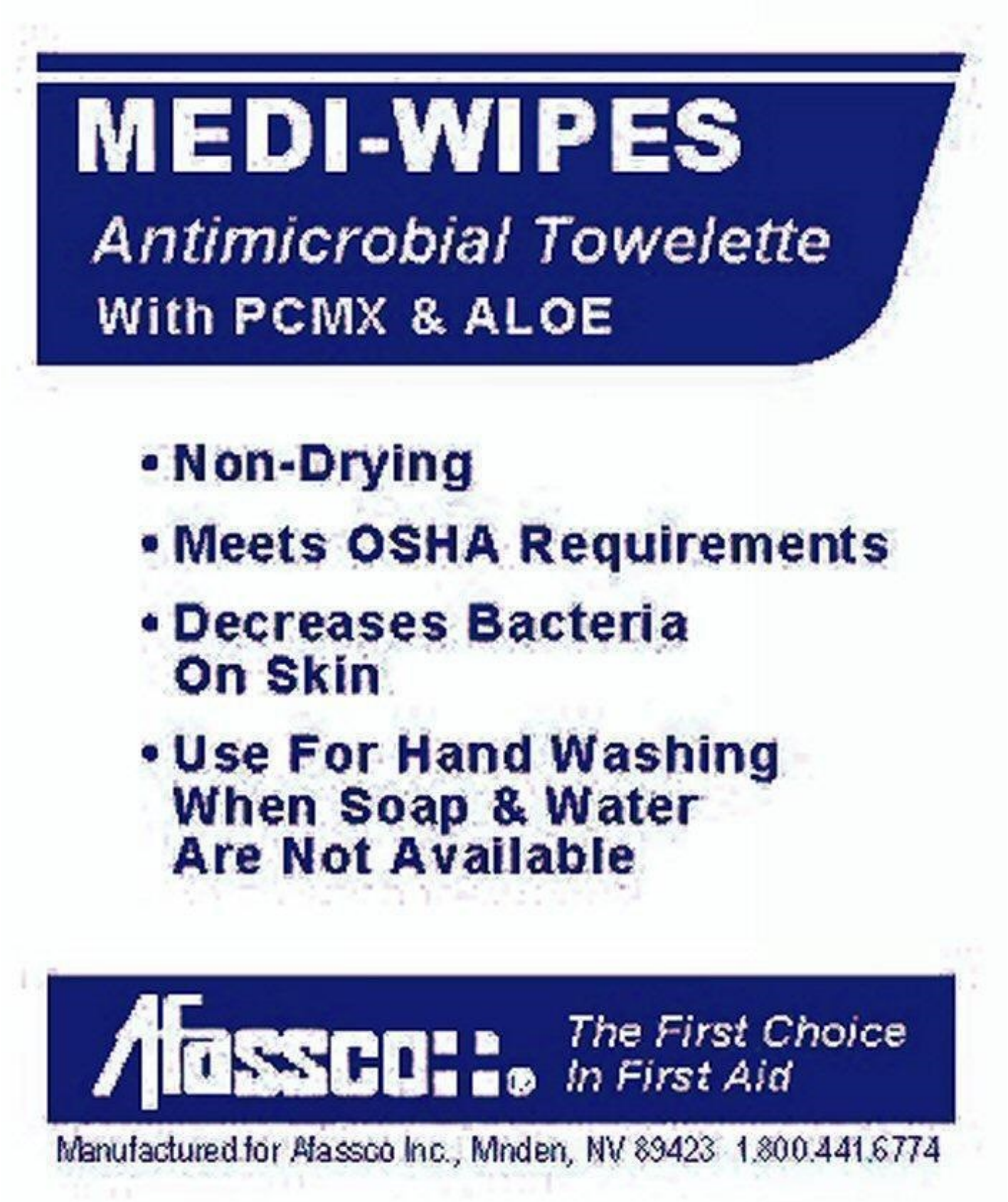
MEDI-WIPES

Antimicrobial Towelette With PCMX and ALOE

Afassco::® The First Choice In First Aid

Manufactured for Afassco Inc, Minden, NV 89423 1.800.441.6774

Non-Drying
Meets OSHA Requirements
Decreases Bacteria On Skin
Use For Hand Washing When Soap And Water Are Not Available



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MEDI-WIPES

ethyl alcohol, chloroxylenol swab

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:51532-5104
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
ALCOHOL (UNII: 3K9958V90M) (ALCOHOL - UNII:3K9958V90M)	ALCOHOL	0.6 g in 1.5 g
CHLOROXYLENOL (UNII: 0F32U78V2Q) (CHLOROXYLENOL - UNII:0F32U78V2Q)	CHLOROXYLENOL	7.5 g in 1.5 g

Inactive Ingredients	
Ingredient Name	**Strength**
ALOE VERA LEAF (UNII: ZY81Z83H0X)	
WATER (UNII: 059QF0KO0R)	

Packaging				
#	**Item Code**	**Package Description**	**Marketing Start Date**	**Marketing End Date**
1	NDC:51532-5104-1	20 in 1 BOX	12/01/2013	
1		1.5 g in 1 PACKET; Type 0: Not a Combination Product		
2	NDC:51532-5104-2	500 in 1 BOX	12/01/2013	
2		1.5 g in 1 PACKET; Type 0: Not a Combination Product		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333E	12/01/2013	

Labeler - Afassco Inc (609982723)

Registrant - Afassco Inc (609982723)

Revised: 12/2019 Afassco Inc