MCKESSON OBSTETRICAL ANTISEPTIC TOWELETTE- benzalkonium chloride swab McKesson Medical-Surgical

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 ingredient

Benzalkonium Chloride, 0.13% w/v

Purpose

First Aid Antiseptic

Use

Towelette to help prevent the risk of skin infection.

Warnings

For external use only.

Do not insert

• into the vagina.

Do not

• use in the eyes or apply over large areas of the body.

Consult a doctor in case of deep or puncture wounds, animal bites, or serious burns. Stop use and consult a doctor if the condition persists or gets worse.

Do not use longer than 1 week unless directed by a doctor.

Keep out of reach of children. If swallowed, get medical help or contact a Poison Control Center right away.

Directions

- Tear open packet
- Unfold and clean the affected area
- Discard after single use

Inactive ingredients

water, sodium bicarbonate, fragrance

Questions?

Call 1-800-777-4908

Principal Display Panel - 0.13 g/100 g Case Label

NDC 68599-5806-3

268

McKESSON

Benzalkonium Chloride Towelettes

ANTISEPTIC | GERMICIDAL | STERILE

100

PER BOX

10

BOXES PER CASE

DO NOT REUSE

Store at 59–86°F (15–30°C). Contents STERILE in unopened, undamaged inner package. Not made with natural rubber latex.

Distributed By McKesson Medical-Surgical Inc.

Richmond, VA 23233

Rev. 00 04/16

Made in China

STERILE R

MFR # 268



Principal Display Panel - 0.13 g/100 g Box Label

NDC 68599-5806-2

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benzalkonium chloride swab

Product Information				
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:68599-5806	
Route of Administration	TOPICAL			

Active Ingredient/Active Moiety				
Ingredient Name	Basis of Strength	Strength		
BENZALKO NIUM CHLO RIDE (UNII: F5UM2KM3W7) (BENZALKONIUM - UNII:7N6 JUD5X6 Y)	BENZALKONIUM CHLORIDE	0.13 g in 100 g		

Inactive Ingredients		
Ingredient Name	Strength	
WATER (UNII: 059QF0KO0R)		
SODIUM BICARBONATE (UNII: 8 MDF5 V39 QO)		

ı	Packaging				
ı	# Item Code	Package Description	Marketing Start Date	Marketing End Date	
ı	1 NDC:68599-5806-3	10 in 1 CASE	07/29/2016		
ı	1 NDC:68599-5806-2	100 in 1 BOX			
ı	1 NDC:68599-5806-1	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	07/29/2016		

Labeler - McKesson Medical-Surgical (023904428)

Establishment					
Name	Address	ID/FEI	Business Operations		
Jiangsu Province Tech (Shanghai)		530968767	manufacture(68599-5806)		

Revised: 8/2020 McKesson Medical-Surgical