



December 22, 2010

URGENT MEDICAL DEVICE RECALL

Aseptic Control Products, Inc.
3831 Industrial Avenue
Unit D
Rolling Meadows, IL 60008

Dear Customer:

This is to inform you of a product recall involving **Sterile Lubricating Jelly** manufactured by Triad Group.

This recall has been initiated due to concerns expressed by the Food and Drug Administration regarding the validation of the gamma radiation sterilization cycles for these products. We are initiating this recall because use of inadequately sterilized product might result in patient infection.

This recall extends to all Lots of Sterile Lubricating Jelly remaining within their labeled expiration dating (three years), including all Lot numbers beginning with the digits 7, 8, 9, or 0. We began shipping the Lots of product subject to this recall in January, 2007

Please immediately examine your inventory and quarantine product subject to recall – any stock of Sterile Lubricating Jelly manufactured by Triad Group. In addition, if you may have further distributed this product, please identify your customers and notify them at once of this product recall. Your notification to your customers may be enhanced by including a copy of this recall notification letter.

This recall should be carried out to the user level. Your assistance is appreciated and necessary to prevent potential patient harm.

Please complete and return the enclosed response form as soon as possible and return the recalled product to Triad Group.

If you have any questions please call Triad Group Customer Service Monday through Friday, between the hours of 8:30 A.M. and 4:00 P.M. Central Time: 262-538-2900 ext 2761.

This recall is being made with the knowledge of the Food and Drug Administration.

Yours truly,

Jack Waterman

Jack Waterman
Regulatory Affairs Manager



Recall Acknowledgement

Please scan and E-mail the following information to:
recall.coordinator@triad-group.net

Or FAX to 262-538-2947

Or mail to:
Recall Coordinator
Triad Group
700 West North Shore Drive
Hartland, WI 53029

Thank you for your cooperation.

I / We have the following stocks of Sterile/Non-Sterile Alcohol Prep Pads and/or Alcohol Swabsticks on hand:

Lot Number	Quantity on Hand

I / We have no stocks of Alcohol Prep Pads and/or Swabsticks on hand

Please have Customer Service contact me with instructions on returning this product

Name / Title	
Firm	
Address	
City, State, ZIP	
Telephone	



Notice of Destruction

PRODUCT DESIGNATED FOR DESTRUCTION

Product Description:

Item Number:		NCR No.:	
Lot Code:		RGA No.:	
Quantity/unit:		PO No.:	

Reason for Destruction:

Product Location:

Company Name:

CERTIFICATE OF DESTRUCTION

THIS PRODUCT HAS BEEN DESTROYED IN A MANNER RENDERING IT INCAPABLE OF USE.

Method of Destruction:

- Deface - Compactor / dumpster - land fill
- Separate Solids From Liquids
- Segregated & Packaged by Disposal Service for Destruction
- In House Wastewater Treatment
- Other _____

Destroyed By:

Title:

Verified By:

Title:



January 3, 2011

URGENT DRUG RECALL

Nurse Assist
3400 Northern Cross Blvd
Ft Worth, TX 76137

Dear Customer:

This is to inform you of a voluntary product recall involving **ALL LOTS** of ALCOHOL PREP PADS, ALCHOLOL SWABS, and ALCOHOL SWABSTICKS manufactured by Triad Group but which has been private labeled for many accounts. This recall involves those products marked as STERILE as well as non-sterile products.


This recall has been initiated due to concerns from a customer about potential contamination of the products with an objectionable organism that may or may not be related to Triad's manufacture of these products. We are, out of an abundance of caution, recalling these lots and revalidating our production lines to ensure that we are not the source of these contamination issues.

Please immediately examine your inventory and quarantine product subject to the recall. In addition, if you have further distributed this product, please identify your customers and notify them at once of this product recall. Your notification may be enhanced by including a copy of this recall notification letter. This recall should be carried out to the user or consumer level. Your assistance is appreciated and necessary to prevent potential patient harm.

Please complete and return the attached response form as soon as possible. **DO NOT RETURN THE PRODUCT ON YOUR OWN.** A return authorization number will be issued once the attached response form has been received. No product will be accepted for a return without a return authorization number. We will credit your account for the returned product and pay for the shipping costs to return to Triad. **We ask that you fill out and return the response form even if you have no products subject to this recall, so it will help in our reporting to FDA.**

If you have any questions please call Triad Group Customer Service Monday through Friday between the hours of 8:30 A.M. and 4:00 P.M. Central Time: 262-538-2900 x2761 or e-mail at the following address: recall.coordinator@triad-group.net.

This recall is being conducted with the knowledge of the U.S. Food & Drug Administration.

Yours truly,
 S. Haertle/COO

700 West North Shore Drive
800.288.1288 262.538.2900

Hartland, WI 53029
info@triad-group.net



3400 Northern Cross Boulevard
Fort Worth, TX 76137 U.S.A.
Toll Free (800) 649-6800
Phone (817) 231-1300
Fax (817) 231-1500

Date: 01/20/11

Dear Customer,

This is to inform you that Nurse Assist, Inc. has received a recall notice from Triad Group. This recall has been initiated due to a recall of the **Alcohol Prep Pads** manufactured by Triad Group that are contained in certain procedure trays. See the enclosed recall notice sent by the Triad Group.

This recall involves procedure trays containing Alcohol Prep Pads that were produced by Triad Group in the last 3 years and remaining within their labeled expiration dating. The enclosed list defines the products and lot #s involved. It is important to note that not all lots of product contain Alcohol Prep Pads manufactured by the Triad Group. **Only the product and lots identified on the attached list are affected.**

Please examine your inventory immediately and quarantine the product associated with this recall. We will provide bright labels to affix to your affected inventory. In addition, you need to identify your customers and notify them at once of this product recall. Provide your customers with labels as well.

This recall should be carried out to the user level. Your assistance is appreciated and necessary to prevent potential patient harm.

Please complete and return the enclosed response form as soon as possible.

If you have any questions or to request labels please call Nurse Assist Customer Service Monday through Friday, from 8:00 am to 5:00 pm Central Time at 1-800-649-6800.

Sincerely,

William Kanewske

William Kanewske
Regulatory Affairs Manager

www.nurseassist.com



WELCON

3400 Northern Cross Boulevard
Fort Worth, TX 76137 U.S.A.
Toll Free (800) 649-6800
Phone (817) 231-1300
Fax (817) 231-1500

RECALL ACKNOWLEDGEMENT

Please FAX to 817-231-1500

Or mail to:
Nurse Assist, Inc.
3400 Northern Cross Blvd.
Fort Worth, TX 76137

We have the following stocks of recalled product on hand:

Product Number	Lot Number	Quantity on Hand

We have affixed labels to our current inventory and sent labels to our customers.

We have no stock of recalled product on hand

Name/Title	
Firm	
Address	
City, State, ZIP	
Telephone #	

www.nurseassist.com

Customer #	W5500	TWIN MED, LLC	
Product	Lots		
6112	908311, 909190, 911414		
6122	909103, 1001118, 1002217, 1005019		

Cross to
Twin Med

016C-4044



January 3, 2011

Facts: Alcohol Recall

Thank you for visiting our recall information page. You may have received recent communication about a recall of alcohol wipes produced by Triad. There has been ONE report of a potential contaminant out of hundreds of millions of products sold. This situation is currently being investigated. As a precautionary measure Triad has voluntarily recalled all of our alcohol wipe products. If you have any of these products, please return it to the place of purchase for a full refund.

Please note that Kroger product manufactured in China by Becton Dickinson is NOT a part of this recall. You can find this information on the outside of the box.

Symptoms: Possible rash or skin infection.

If you are experiencing any of the above symptoms, please contact your physician immediately.

Brands affected:

Best Choice	Care One	CVS	Discount Drug Mart
Equaline	Equate	Exchange Select	Exact
Good Sense	Healthcare	Healthy Generations	Kroger
Leader	Life Brand	Longs	Major
MEIJER	Medicine Shoppe	Personelle	Publix
Premier Value	Quality Choice	Rite Aid	Reli-On
Remedy RX	Rexall	Safeway	Select Brand
Shoppers Drug	Sunmark	Up&Up	Top Care
Triad	Triad Sterile	Uniprix	Valu Plus
Western Family	Walgreens		

If you have further questions, please call (800) 288-1288

Yours truly,

The Triad Group, Inc.