



URGENT PRODUCT RECALL

April 18, 2011

To Our Customers,

This letter is to inform you that PSS World Medical, Inc. has received a letter from Smith & Nephew, Inc. Wound Management notifying about the recall of affected lots for several types of Wipes. Refer to the enclosed recall notice from Smiths & Nephew for further reasoning of this recall.

Please reference Attachment A for a list of Affected Item(s) and Lot Number(s)

HEALTHCARE PROVIDER RETURN AND CREDIT INSTRUCTIONS:

- 1.) Immediately examine your inventory and quarantine the affected items and lot numbers referenced in this recall.
- 2.) If your facility has further distributed any of the referenced item(s) and lot number(s) to another healthcare facility, immediately inform them of this notification and request they return the affected product directly to you.
- 3.) Contact your Gulf South Customer Care team via telephone (800-347-2456) or email (customercare@gsms.com) to Request a Return Goods Authorization and to place an order for replacement product.

We sincerely apologize for any inconvenience this notice may have caused you and your staff.

Sincerely,

Quality Assurance Department

Enclosed:

- 1.) Attachment A
- 2.) Smith & Nephew Recall Notification

ATTACHMENT A

GSMS #	MFG #	Product Description
801930	402300	ADHESIVE REMOVER WIPES 50/BX
Affected Lot #		
0F197	0K208	0F195
0F223	0K209	0K233
0F224	0K234	0M150
0F228	0M136	0M151
0F229	0M137	0M238
0F239	0M149	1B129
0H185	0H228	0H229

GSMS #	MFG #	Product Description
805434	403100	ADHESIVE REMOVER WIPES 50/BX
Affected Lot #		
0J232	0E145	
0L147	0E210	
0M177	0E211	
1A117	0G138	
1B112	0H248	
0J190		

GSMS #	MFG #	Product Description
822667	59420600	SKIN PREP WIPE NO STING
Affected Lot #		
0J292	0E230	
9K150	0E231	
9K151	0J290	
9L169	0J291	
9L170		

GSMS #	MFG #	Product Description
805464	420400	SKIN PREP WIPE PROTECT DRESS
Affected Lot #		
0G137	0K271	0L242
0G165	0K272	0M199
0G184	0L164	0M200
0G282	0L165	1A246
0H266	0L206	1A247
0J125	0L225	1A248
0J124	0L226	1A256
0J126	0L227	0J85
0K118	0K119	0K120
0K254	0K255	



TRIAD
GROUP

January 3, 2011

received by Medikmark on January 10, 2011

URGENT DRUG RECALL

Medikmark
3600 Burr Wood Dr
Waukegan, IL 60085

Dear Customer:

This is to inform you of a voluntary product recall involving ALL LOTS of ALCOHOL PREP PADS, ALCOHOL 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,
Eric Haertle/COO

700 West North Shore Drive Hartland, WI 53029
800.288.1288 262.538.2900 info@triad-group.net

Wound Management
Smith & Nephew, Inc.
970 Lake Carillon Drive
Suite 110
St. Petersburg, FL 33716

Recall Hotline: 1-888-613-0271
F 727 392-6914
www.smith-nephew.com

April 6, 2011

URGENT: DEVICE RECALL

ATTENTION: Recall Department
Gulf South Medical Supply Inc
4345 Southpoint Boulevard
JACKSONVILLE, FL 32216

RE: Product Names: REMOVE^o Universal Adhesive Remover Wipes, UNI-SOLVE^o Adhesive Remover Wipes, SKIN-PREP^o Protective Wipes, PERI-PREP Protective Wipes, and NO-STING SKIN-PREP^o Protective Wipes
Product Code Numbers: 420400, 420471, 59420425, 403100, 402300, 59403125, 59420600
Lot Numbers: -See Attached Form

Dear Customer:

Smith & Nephew, Inc., Advanced Wound Management Division is voluntarily recalling multiple lot numbers of the following products: **REMOVE^o Universal Adhesive Remover Wipes, UNI-SOLVE^o Adhesive Remover Wipes, SKIN-PREP^o Protective Wipes, PERI-PREP Protective Wipes, and NO-STING SKIN-PREP^o Protective Wipes.** Our shipping records indicate that you received one or more of the affected lots of these wipes as indicated on the attached inventory Form.

The decision was made to recall these affected lots after a thorough review of available information regarding conditions at the manufacturing site of the contract manufacturer, H&P Industries dba The Triad Group. It has been determined that the recalled lots were made in the same facility as other wipe products, including wipes, swabs and swab sticks, that are the subject of other voluntary recalls due to a potential bacterial contamination issue of these products.

Testing has been performed on some Smith & Nephew products lots already distributed and/or under quarantine and test results received to date show no contamination.

Out of an abundance of caution, however, Smith & Nephew has decided to voluntarily recall the affected wipe products identified in this notice. Smith & Nephew is committed to providing our customers with high quality, reliable products. We are taking this action to remove the product lots identified in this notice from the market as a prudent, cautious and conservative measure to assure patient safety and product performance.

URGENT DEVICE RECALL
Smith & Nephew, Inc.

Please note this recall is limited to specific lots of the products identified in this notice. The lot numbers specified in the attached Form are the affected products that our records indicate were shipped to you from **01/07/2010 through 02/23/2011**.

The purpose of this letter is to alert you to this situation and advise you that we will conduct a voluntary recall process per strict FDA guidelines. This action is proposed as a Class II action and is being conducted to all customer levels (purchaser/distributor, hospitals/health facilities, kit packers, individual end users/consumers) due to the potential health risk. Accordingly, we are asking that you conduct a full sub-recall of the affected products from all of your customers to whom you supply such products.

Action required by you:

Please **check your inventory immediately** to identify and return the lot numbers of the following products shipped to you and identified on the attached Form: REMOVE[®] Universal Adhesive Remover Wipes, UNI-SOLVE[®] Adhesive Remover Wipes, SKIN-PREP[®] Protective Wipes, PERI-PREP Protective Wipes, and NO-STING SKIN-PREP[®] Protective Wipes. Product returns to Smith & Nephew must be conducted according to the instructions detailed below.

Please report the amount of affected lot numbers using the inventory Form and **(1) mail the Form back to us using the postage-paid, self-addressed envelope provided; (2) also fax your response to: 727-392-6914.**

It is very important that you fill in the requested information on the enclosed form, fax the form to the preceding number and return the original to us within ten (10) business days after receipt of this Notice. Even if you do not have any of the identified product lots on hand, please complete the Form and fax it to us along with sending the original copy to us in the enclosed envelope.

Distributors or Kit Packers:

Smith & Nephew is advising all distributors and kit packers who utilize REMOVE[®] Universal Adhesive Remover Wipes, UNI-SOLVE[®] Adhesive Remover Wipes, SKIN-PREP[®] Protective Wipes, PERI-PREP Protective Wipes, and NO-STING SKIN-PREP[®] Protective Wipes to return unused inventories of affected product lots to the address below. This recall should be carried out to the end user/consumer level.

You are advised to take appropriate action to notify your customers to discontinue use of all affected lots. Your customers should be directed to either **destroy or return the product to you. Note: Smith & Nephew will not accept product returns from customers who do not have an account with us.**

You should notify your customers/patients to **discontinue the use of the affected lots of these products immediately and seek an alternative product.**

URGENT DEVICE RECALL
Smith & Nephew, Inc.

All Smith & Nephew customers:

Replacement Product: Smith & Nephew expects to promptly restore its inventory to normal levels with product supplied from a new source. Upon receipt of your inventory Form Smith & Nephew will contact you to review options for inventory adjustments. Please indicate on the attached inventory Form the contact information for the person in your organization responsible for purchasing and/or inventory management.

Product Return: Please return the affected product by shipping it back to Smith & Nephew via UPS (United Parcel Service). Under payment terms, please ship 'Bill 3rd Party', and enter account number 93A25X **Please mark the box:** *SKIN-PREP^o Protective Wipes, REMOVE^o Universal Adhesive Remover Wipes, and UNI-SOLVE^o Adhesive Remover Wipes, - CORRECTION AND REMOVAL.* Please return the product to:

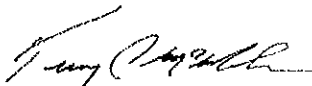
Smith and Nephew
c/o APL Logistics
2925 Shawnee Industrial Way
Suite # 100
Suwanee, GA 30024-4609

If you have any questions concerning this notification, please contact our Recall Hotline Center at 1-888-613-0271.

RECALL NOTIFICATION
Enclosure/As Stated

We apologize for any inconvenience this may have caused you, your customers or patients.

Sincerely,



Terry C. McMahon
Director, Regulatory Affairs and Quality – North America
Smith & Nephew, Inc.
Advanced Wound Management



URGENT PRODUCT RECALL

March 10, 2011

To Our Customers,

This letter is to inform you that PSS World Medical, Inc. has received a recall letter from Medikmark, Inc. regarding the Triad Recall of Sterile Lubrication Jelly and Alcohol Prep Pads contained in certain Medikmark procedure trays.

This recall involves Medikmark procedure trays containing Sterile Lubricating Jelly or Alcohol Prep Pads that were produced by Triad Group in the last 3 years and remain within the labeled expiration date. "Attachment A" defines the products and lot numbers involved.

It is important to note that not all lots of Medikmark procedure trays containing Sterile Lubricating Jelly packs or Alcohol Prep Pads are manufactured by the Triad Group.

The affected products and lot numbers are identified on "Attachment A".

Required Action:

- 1.) **Examine your inventory immediately and quarantine the product associated with this recall.**
- 2.) **If your facility/agency has further distributed any of the referenced item(s) and lot number(s) to another healthcare facility, immediately inform them of this notification and request they return the affected product directly to you.**
- 3.) **Contact your Gulf South Customer Care team via telephone (800-347-2456) or email (customercare@gsms.com) to Request a Return Goods Authorization and to place an order for replacement product.**

INSTRUCTIONS TO PATIENT/CONSUMER

- 4.) **If your facility/agency has provided any of the reference item(s) and lot number(s) to a patient/consumer, instruct them to dispose of the affected product.**
- 5.) **Please fill out the enclosed Notice of Destruction (ATTACHMENT B) form with the reorder number along with the number of kits the patient/end user destroyed and FAX to (800-827-2002).**

ATTACHMENT A

GSMS Item #	Mfg #	Description	Lot Numbers
801532	OR3207	TRAY FOLEY INSERTION 30CC	908519
801532	OR3207	TRAY FOLEY INSERTION 30CC	908526
801532	OR3207	TRAY FOLEY INSERTION 30CC	909502
801532	OR3207	TRAY FOLEY INSERTION 30CC	909516
825927	3T8000	IRRIGATION TRAY	9E022
825927	3T8000	IRRIGATION TRAY	9A024
825927	3T8000	IRRIGATION TRAY	9B012
825927	3T8000	IRRIGATION TRAY	9C013
825927	3T8000	IRRIGATION TRAY	9C024
825927	3T8000	IRRIGATION TRAY	9G004
825927	3T8000	IRRIGATION TRAY	9H027