Life-Assist
Helping Heroes Save Lives

URGENT DRUG RECALL

Tuesday, June 7, 2016

Dear Valued Life-Assist Customer,

According to our records you may have purchased an item that has been recalled by the manufacturer. Please examine your stock to determine if you have the following product, with the affected lot number in your possession.

Aspirin, Children's Chewable, 81 mg (blister packs)

Life-Assist, Inc. Product Code	NDC/UPC	Lot #(s)
MD85281	63739-434-01	0110701

Reason for Recall: The manufacturer has issued a voluntary recall of the above lot of Aspirin Chewable Tablets 81mg due to a mix up of secondary packaging labeling.

Affected product first shipped from the manufacturer February 2016.

Please immediately remove and quarantine any of the affected lot from your inventory.

Contact Life-Assist Customer Service at 800-824-6016 or salesservice@life-assist.com for replacement or credit and to obtain a Return Authorization.

We apologize for any inconvenience.

mooremedical
Supporting Health & Care

URGENT DRUG RECALL

June 3, 2016

Dear Valued Moore Medical Customer:

McKesson Packaging Services has notified Moore Medical of an Urgent Drug Recall regarding one lot of their Aspirin Chewable Tablets 81mg. This recall has been issued due to a mix up of secondary packaging labeling. Affected product first shipped February 10, 2016.

This Urgent Drug Recall is being done with the knowledge of the Food and Drug Administration.

For questions regarding this notification, please contact McKesson Packaging Services at (704) 784-4301.

A review of our records indicates that you or your company may have purchased items included in this notification. Carefully review the information in this letter and follow the instructions provided below.

Refer to the table below for a list of affected item(s) and lot number(s) distributed by Moore Medical

Moore Medical #	NDC #	Description	Affected Lot(s)	Exp. Date
85281	63739043401	Aspirin 81 mg UD Chew Tabs	110701	07/31/2017

Moore Medical Customer Instructions:

1.) Review the enclosed Urgent Drug Recall from McKesson Packaging Services for details and a complete listing of the affected product(s).

2.) Quarantine and immediately discontinue use of any product matching the affected item(s) and lot number(s) listed above.

3.) If you have product affected by this recall, fill out the Moore Medical Reply Form and fax it back (do not mail) to our Regulatory Affairs Department at **866.550.1169.** Detailed product return instructions are provided on the reply form. Please note that credit will only be issued for product(s) from the affected lots listed. Replacement items will not be sent.

4.) If you have further distributed any of the item(s) referenced in this notification, provide your accounts with a copy of this notification and request that they return the affected product directly to you.

We sincerely apologize for any inconvenience this product recall may have caused you and your staff. If you have any questions about information provided in this communication, please contact our Regulatory Affairs Department at 800.234.1464 ext. 5407.

Thank you for your prompt attention,

Regulatory Affairs Department
Telephone: 800-234-1464 X5407
Email: MMCregulatoryaffairs@mooremedical.com