URGENT DRUG RECALL

April 13, 2017

Dear Valued Moore Medical Customer:

Mylan Specialty L.P. has notified Moore Medical of an Urgent Drug Recall regarding specific lots of their EpiPen and EpiPen Jr Auto-Injectors manufactured by Meridian Medical Technologies, a Pfizer Company. This notice has been issued due to the receipt of two previously disclosed reports outside of the U.S. of failure to activate the device due to a potential defect in a supplier component. The potential defect could make the device difficult to activate in an emergency and have significant health consequences for a patient experiencing a life-threatening allergic reaction. The incidence of the defect is extremely rare and testing and analysis across the potentially impacted lots has not identified any units with a defect. Affected product first shipped December 1, 2015.

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

For clinical inquiries, please contact Mylan at (800) 796-9526.

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 on page 2 for a list of affected item(s) and lot number(s) distributed by Moore Medical

Moore Medical Customer Instructions:

1.) Immediately discontinue use of any product matching the affected item(s) and lot number(s) listed on page 2.

2.) A copy of the Urgent Drug Recall from Mylan has been included for reference.

3.) Regardless of whether 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 affected product(s) from the affected lots listed. Replacement items will not be sent. To ensure timely credit to your account and support the completion of this recall, please respond within 30 days.

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.

(Continued on next page)
## Table of Affected Products

<table>
<thead>
<tr>
<th>Moore Medical #</th>
<th>NDC #</th>
<th>Description</th>
<th>Affected Lot(s)</th>
<th>Exp. Date</th>
</tr>
</thead>
<tbody>
<tr>
<td>66033</td>
<td>49502-500-02</td>
<td>EPIPEN 0.3MG AUTO-INJECTOR 2PKUM PK SZ 2 0.3MG</td>
<td>5GM631, 5GM640, 6GM072, 6GM081, 6GM082, 6GM087, 6GM088, 6GM091, 6GM198, 6GM199</td>
<td>4/30/2017, 5/31/2017, 9/30/2017, 9/30/2017, 10/31/2017, 10/31/2017, 10/31/2017, 10/31/2017, 10/31/2017</td>
</tr>
<tr>
<td>66032</td>
<td>49502-501-02</td>
<td>EPIPEN JR 0.15MG 2 PKUM PK SZ 2 0.15MG</td>
<td>6GN215, 5GN767, 5GN773</td>
<td>9/30/2017, 4/30/2017, 4/30/2017</td>
</tr>
</tbody>
</table>

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