November 6, 2019

Urgent: Grifols Therapeutics issues voluntary market withdrawal of one lot of Gamunex®-C 10%

Dear ASD Healthcare Customer:

Grifols Therapeutics is initiating a voluntary market withdrawal of one lot of Gamunex-C 10% with the knowledge of the U.S. Food and Drug Administration (FDA), Center of Biologics Evaluation and Research. The affected lot information is listed below.

Grifols’ actions to withdraw Gamunex-C 10% is being conducted due to a higher rate of allergic/ hypersensitivity type reactions, a small number of which were considered medically significant. Hypersensitivity and anaphylactic/ anaphylactoid reactions are a known risk with IVIG products.

Grifols provided the following affected lot information:

<table>
<thead>
<tr>
<th>Product Description</th>
<th>NDC Number</th>
<th>Lot Number</th>
<th>Expiration Date</th>
</tr>
</thead>
<tbody>
<tr>
<td>Gamunex-C 10%</td>
<td>13533-800-24</td>
<td>A1GLD00622</td>
<td>18 May 2022</td>
</tr>
</tbody>
</table>

To comply with the withdrawal, we ask for your cooperation in taking the following action:

- Please examine your stock immediately to determine if you have the above-mentioned lot on hand. If you have product from this lot, cease use immediately and quarantine the affected product.
- Use the Request a Return tool to get started. Return product to ASD Healthcare’s warehouse (address listed below)

Warehouse Supervisor
ASD Healthcare
345 International Blvd, #400A
Brooks, KY 40109

For the full recall notice, please view the attached document on the following pages.

If you have any questions or concerns, please contact ASD Healthcare's customer service at 800.746.6273.
URGENT DRUG WITHDRAWAL

November 5, 2019

This is to inform you that Grifols Therapeutics is initiating a voluntary withdrawal of one lot of Gamunex®-C 10%, as detailed below.

As a precautionary measure, this voluntary withdrawal is being conducted due to a higher rate of allergic/hypersensitivity type reactions, a small number of which were considered medically significant. Hypersensitivity and anaphylactic/anaphylactoid reactions are a known risk with IVIG products.

This withdrawal is being conducted with the knowledge of the U.S. Food and Drug Administration, Center for Biologics Evaluation and Research.

Gamunex®-C 10%, lot affected by this withdrawal is:

<table>
<thead>
<tr>
<th>Lot Number</th>
<th>Material Number</th>
<th>Expiration Date</th>
<th>Market Release</th>
<th>NDC Number</th>
</tr>
</thead>
<tbody>
<tr>
<td>A1GLD00622</td>
<td>729688</td>
<td>18 MAY 2022</td>
<td>20 JUN 2019</td>
<td>13533-800-24</td>
</tr>
</tbody>
</table>

The withdrawal is required to be conducted to the consumer/user level. Please notify your subsidiary customers of this withdrawal, and instruct them to return product as noted below.

Please examine your stock immediately to determine if you have any of the above-mentioned lots on hand. If you have product from this lot, cease use of the product immediately. If you have further distributed the product, you must immediately send notification of this withdrawal to your customers. This lot was distributed within the United States only. Return product to the authorized distributor of record for this lot. Distributors should return product to Grifols Therapeutics, 8368 US Hwy Bus 70 West, Clayton, NC 27520. For any questions concerning return of the product, contact US Customer Service at (800) 243-4153.

After receipt of returned product by Grifols, you will be credited by your distributor for the returned goods. If you have any technical or clinical questions, please contact US Clinical Communications at (800) 520-2807. Your prompt attention to this notice is appreciated.

Sincerely,

Clark Zervos
Vice President, Quality
Grifols Therapeutics