Urgent: Sanofi is initiating a voluntary recall of all batches within expiry of Regular Strength Zantac® 75, Maximum Strength Zantac® 150, Maximum Strength Zantac® 150 Cool Mint Tablets

Dear ASD Healthcare Customer:

Sanofi is initiating, as a precautionary measure, a voluntary recall of all batches within expiry of Regular Strength Zantac® 75, Maximum Strength Zantac® 150, Maximum Strength Zantac® 150 Cool Mint Tablets.

This recall is being taken due to a possible contamination with a nitrosamine impurity called N-nitrosodimethylamine (NDMA). It is being conducted with the knowledge of the U.S. Food and Drug Administration.

Sanofi provided the following affected product information:

<table>
<thead>
<tr>
<th>Product Description</th>
<th>NDC Numbers</th>
<th>Expiration Date</th>
</tr>
</thead>
<tbody>
<tr>
<td>Regular Strength Zantac 75</td>
<td>41167-0300-0, 1, 2, 3, 5, 6, 7, 8 41167-0310-0, 1, 2, 3, 4, 5, 6, 7, 8, 9 41167-0320-0, 1, 2, 3, 4, 5, 6, 7 0597-0120-06, 08, 09, 24, 38, 50, 76, 78, 80, 82, 87</td>
<td>All product within expiry</td>
</tr>
<tr>
<td>Maximum Strength Zantac 150</td>
<td>0597-0121-01, 06, 08, 09, 11, 24, 38, 50, 64, 66, 68 0597-0121-78, 80, 82, 85, 90, 94 0597-0122-01, 08, 13, 34, 37, 40, 54, 61, 81, 96 66715-9736-2, 3, 8 67751-151-01 67751-152-01, 02 68151-2584-0 502-220-25, 50269-222-25</td>
<td></td>
</tr>
<tr>
<td>Maximum Strength Zantac 150 Cool Mint Tablets</td>
<td></td>
<td></td>
</tr>
</tbody>
</table>

We ask for your cooperation in taking the following action:

- Please examine your stock immediately to determine if you have the above-mentioned products. If you have product, cease use immediately and quarantine the affected items.
- To obtain a return kit, contact INMAR at 877-275-0993 Option1.
- If you prefer to fax this request, the fax number is 336-499-8145; alternatively the email contact is zantacrecall@inmar.com.

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: VOLUNTARY DRUG RECALL
Depth of Recall: Retail and ECommerce Outlets
NDC Number: 41167-0300-0, 1, 2, 3, 5, 6, 7, 8
          41167-0310-0, 1, 2, 3, 4, 5, 6, 7, 8, 9
          41167-0320-0, 1, 2, 3, 4, 5, 6, 7
          0597-0120-06, 08, 09, 24, 38, 50, 76, 78, 80, 82, 87
          0597-0121-01, 06, 08, 09, 11, 24, 38, 50, 64, 66, 68,
          0597-0121-78, 80, 82, 85, 90, 94
          0597-0122-01, 08, 13, 34, 37, 40, 54, 61, 81, 96
          66715-9736-2, 3, 8
          67751-151-01
          67751-152-01, 02
          68151-2584-0
          502-220-25, 50269-222-25

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<thead>
<tr>
<th>Product Description</th>
<th>Expiration Date</th>
</tr>
</thead>
<tbody>
<tr>
<td>Regular Strength Zantac 75°</td>
<td>All product within expiry</td>
</tr>
<tr>
<td>Maximum Strength Zantac 150°</td>
<td>All product within expiry</td>
</tr>
<tr>
<td>Maximum Strength Zantac 150° Cool Mint Tablets</td>
<td>All product within expiry</td>
</tr>
</tbody>
</table>

To Whom It May Concern,

This letter is to inform you that Sanofi is initiating as a precautionary measure a voluntary recall of all batches within expiry of Regular Strength Zantac® 75, Maximum Strength Zantac® 150, Maximum Strength Zantac® 150 Cool Mint Tablets.

This recall is being taken due to possible contamination with a nitrosamine impurity called N-nitrosodimethylamine (NDMA).

Our records indicate that you have received product that is subject to this recall. Distribution of recalled product by Sanofi started on 01/04/2017 and ended on 10/11/2019.

Required Actions:

- Immediately discontinue distribution of Regular Strength Zantac 75 mg, Maximum Strength Zantac 150 mg, Maximum Strength Zantac 150 mg Cool Mint Tablets.

- Please notify any of your customers who have purchased Regular Strength Zantac® 75 mg, Maximum Strength Zantac® 150 mg, Maximum Strength Zantac® 150 mg Cool Mint Tablets and instruct them to discontinue use and return any product within expiry to your retail outlet.
• Sanofi will reimburse retail outlets for product returned to Sanofi. Sanofi will not be able to refund individual consumers.

• To obtain a return kit please contact INMAR at 877-275-0993 Option 1

• If you prefer to fax this request, the fax number is 336-499-8145; alternatively the email contact is zantacrecall@inmar.com.

Sanofi will issue retail outlets credit for product within 10-15 days after the receipt of the returned recalled product by INMAR.

This recall is being conducted with the knowledge of the U.S. Food and Drug Administration. Your assistance with this product recall is greatly appreciated. Direct Accounts may contact Sanofi Consumer Healthcare if you require additional information.

All questions of a medical or clinical nature, or regarding product quality should be sent to Sanofi Medical Information Services in the U.S. at 1-800-633-1610, option #. If you have questions on any other topics, please call 877-275-0993 Option 2

Thank you.

Sincerely,

[Signature]

Allison Steele
Director, Regulatory Compliance
Recall Leader, Sanofi U.S.