



SAGENT™

Sagent Pharmaceuticals, Inc.

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CUSTOMER NOTIFICATION/ RECALL COMMUNICATION
URGENT: ONDANSETRON in 5% DEXTROSE INJECTION, 32mg/50mL RECALL
MANUFACTURED BY CLARIS LIFESCIENCES

June 1, 2010

Dear Valued Customer:

The Food and Drug Administration issued a Public Health Alert notifying Healthcare Professionals of Claris Lifesciences' Recall of several premix Intravenous products including Ondansetron in 5% Dextrose Injection, 32mg/50mL. As a result of Claris' recall this letter is to inform you that Sagent Pharmaceuticals, Inc. is recalling the following product:

Product	Lot Number(s)	NDC Number	Distribution Dates
Ondansetron in 5% Dextrose Injection, 32mg/50mL	A090309, A090310, A090311, A090312	25021-776-50	August 5, 2009 – May 27, 2010

This recall is being made with the knowledge of the Food and Drug Administration and has been initiated due to Claris' Recall. The Recall is a result of reported foreign material and non-sterility in one lot of product.

To implement this recall, please take the following actions:

1. Immediately examine your inventory and quarantine product subject to recall.
2. Weigh the product. (Required for generation of call tag).
3. Immediately discontinue use and distribution of the attached listed lots. A credit memo will be issued covering the quantity of your product returned.
4. Return product to:
DDN Logistics
ATTN: Returns Department
4580 S. Mendenhall
Memphis, TN 38141

NOTE: A call tag, a pre-printed, pre-paid return label will be provided to you for product return; return shipment is free of charge. For the call tag, contact DDN Customer Service at 1-866-625-1618, option 1. You will be asked for the product weight which is required for generation of the call tag.

5. If you may have further distributed this product, please identify those customers and notify them at once of this product recall. Your notification to your customers should include a copy of this recall notification letter.
6. Please complete and return the enclosed "Customer Recall Return Response Form" as soon as possible and fax the form to us at 1-847-908-1827.

This recall should be carried out to the user level.

Your assistance is appreciated. I apologize for any inconvenience this may cause you.

If you have any questions, please do not hesitate to call our Customer Service at 1-866-625-1618 which was specifically set-up to address any concerns that you may have.

Sincerely,

Sheila Moran
Vice President Quality Assurance & Facility Compliance
Sagent Pharmaceuticals, Inc.
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