

**Kogenate FS**

Product Name	NDC No.	Lot No.	Exp. Date	Quantity Returned
Kogenate FS 2000 IU w/ Vial Adapter	0026-3786-65	27118RK	06/2021	
Kogenate FS 2000 IU w/ Vial Adapter	0026-3786-65	27119CG	06/2021	

## BUSINESS REPLY CARD

Bayer  
Drug Recall Program  
7/18/2019 (rev 8/21/2019)

*Your timely response to this drug warning notification is requested. Please fill out and return this form within five (5) business days. Please FAX or EMAIL form to:*

**ATTN: INMAR - RECALL**  
**Email: BayerRecalls@inmar.com**  
**PHONE: 855-707-7518**  
**FAX: 817-868-5362**

Please make one selection in each section below.

Section A: <input type="checkbox"/> We <b>have</b> stock of the above recalled product and will return the stock. <input type="checkbox"/> We <b>do not have</b> stock of the above recalled product and will not be making a return.	Section B: <input type="checkbox"/> We <b>have</b> notified all of our Consignees to the retail level to return the recalled product. <input type="checkbox"/> We <b>do not have</b> any Consignees at the retail level for this lot.
Section C: <input type="checkbox"/> We <b>have</b> received recalled patient-level product and it <b>will be</b> replaced <input type="checkbox"/> We <b>have</b> received recalled patient-level product and it <b>will not be</b> replaced <input type="checkbox"/> We <b>have not</b> received recalled patient-level product.	

**Bayer Account No. to Credit:** \_\_\_\_\_

<b>Date:</b>
<b>Name:</b>
<b>Company:</b>
<b>DEA Number:</b>
<b>Phone No.:</b>
<b>Email:</b>
<b>Signature:</b>

Return Tracking No. For Reference:

*\*\*IF you have recalled product under multiple Bayer Account numbers, please complete this form for each account.*