FDA Safety Communication: <insert SC title>

PLEASE HELP US (TO HELP YOU)...

We need to hear from you so we can evaluate and improve our Safety Communications as well as the overall effectiveness of the Safety Communication program. Please take a few minutes to answer the questions below. We will publish a summary of the results. All questions relate to this Safety Communication.

Your responses will be kept confidential. Thank you for your assistance.

1.	A. Were you able to identify the problem this Safety Communication addresses? No B. If no, why not?		Yes				
2.	A. Were you able to easily understand the problem addressed in this Safety Communication? Yes B. If no, why not?	No					
3.	A. Did you understand the actions for reducing risk?						
4.	A. Did you find the information contained in this Safety Communication useful? No B. If no, why not?	Yes					
5.	Did you find the information contained in this Safety Communication to be timely?		Yes				
6.	A. Were you aware of the problem addressed in this Safety Communication prior to reading it? \Box	□ No	Yes				
	B. If yes, how did you first become aware of the problem?						
	apersonal experience emanufacturer recall bcoworkers, friends, or family fmanufacturer notification cprofessional bulletin gyour organization's management dprofessional symposium hprint media (e.g., newspaper) i electronic media (e.g., web) j social media (e.g., Twitter, Facebook)						

k	my hea	alth care	provid	er
I.	Other (please s	pecify)	

7. A. Have you taken any actions to eliminate or reduce the risk as a result of the information in this Safety_ Communication? Yes

No

If yos, what actions did you take?

C. If no, why not? a already took action prior to Safety Co b actions planned prior to Safety Comm c actions planned based on Safety Com d risk was never applicable to our operate e felt risk did not warrant action	nunication but not yet taken nmunication but not yet taken
A. Have you signed up to receive future Safe	ety Communications electronically?
B. If no, why not?	
l am a:	
a Hospital Administrator b Risk Manager c Director of Nursing d Biomedical/Clinical Engineer	fQuality Assurance ManagergHome Health Care AdministratorhNursing Home AdministratoriHospice AdministratorjMedical Device Industry RepresentativekStudent (e.g., medical, nursing, public health)l.Health Educatorm.Patientn.Caregiver

10. I have the following suggestions for improving this FDA Safety Communication:

Public reporting burden for this collection of information is estimated to average 0.17 hours per response, including the time for reviewing instructions, searching existing data sources, gathering and maintaining the data needed, and completing reviewing the collection of information. Send comments regarding this burden estimate or any other aspect of this collection of information, including suggestions for reducing this burden to:

> Department of Health and Human Services Food and Drug Administration

An agency may not conduct or sponsor, and a person is not required to respond to, a collection of information unless it displays a currently valid OMB control number.