



Medical Devices Bureau, Ministry of Health Malaysia
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Medical Device / Equipment ALERT- PRODUCT CORRECTION

Date Issued : 5th February 2009

Ref:MDB/A/2009/001

IMMEDIATE ACTION	√
ACTION	√
UPDATE	
INFORMATION REQUEST	

PRODUCT	spesific STERRAD 50 Sterilizer
CLASS	Not Mentioned
USE	Sterilizers
SOURCE OF MEDICAL DEVICE RECALL / ALERT	Communication by electronic mail sent to Medical Device Bureau via Johnson & Johnson Sdn. Bhd., Medical Division Malaysia.
ALERTING FIRM	Johnson&Johnson Sdn. Bhd., Medical Division
REASON FOR RECALL/ALERT	<i>Refer attachments for details</i>
SCENARIO IN MALAYSIA	<i>Refer attachments for details</i>
ACTION	Please distribute this product notification to all Sterrad 50 system users in your facility (if available) <i>Refer attachments for details</i>
RECOMMENDATION	Users of the abovementioned device (if available) should contact the distributors/supplier of this device and inform the Medical Devices Bureau, Ministry of Health providing the following information:- a. Name of healthcare centre/hospital/clinic

	<p>b. Contact person and contact number c. Numbers of units available d. Name of supplier(s)</p>
CONTACT/ENQUIRIES	<p>Ong Yean Ting, Regulatory Affairs Manager Johnson&Johnson Sdn. Bhd., Ground Floor, G.01, Block B, 10, Jalan Bersatu 13/4 46200 Petaling Jaya, Selangor.</p> <p>T: 603 79555108 F: 603 79553290</p>
REFERENCES FOR DETAILS	<p><i>Refer attachments for details</i></p>