Recall detail

Type of Producti	Medical Device
TGA Recall Reference	RC-2014-RN-00949-1
Product Name/Description [®]	ZOLL ProPadz Sterile Multi-Function Electrodes
	Product code 8900-4055-40 and 8900-4052-40
	Lot Number: 2314
	ARTG Number: 218497
Recall Action Leveliv	Hospital
Recall Action Classification ^v	Class II
Recall Action Commencement Date ^{vi}	1/09/2014
Responsible Entity ^{vii}	Zoll Medical Australia Pty Ltd
Reason / Issue ^{vni}	During internal sterility testing Zoll Medical Corporation has determined that the Zoll ProPadz Sterile Multi-Function Electrodes from the lot listed may not be sterile.
Recall Action ^{ix}	Recall
Recall Action Instructions ^x	Zoll Medical Australia is requesting their customer to quarantine all affected stock. Arrangements will be made for the recovery of the devices.
Contact Information ^{xi}	1800 605 555 - Zoll Medical Australia

Footnotes

- ¹ Type of Product: Medicine, Medical Device, or Biological
- ii TGA Recall Reference: Unique number given by the TGA
- iii Product Name/Description: Brand name (including active ingredient for medicines) and may include generic reference for the kind of medical devices. Includes all necessary information such as affected: catalogue / model and / or batch / serial numbers.
- iv Recall Action Level: The level to which the recall action is to be undertaken. This is based on the significance of the risk and the channels through which the goods have been distributed. The recall action levels are / Wholesale / Hospital / Retail / Consumer.
 - Wholesale includes wholesalers and state purchasing authorities.
 - Hospital includes nursing homes and institutions, hospital pharmacists, ambulance services, blood and tissue banks and laboratories as well as wholesale as appropriate.
 - Retail includes retail pharmacists, medical, dental and other health care professionals as well as wholesale
 and hospital as appropriate.
 - Consumer includes patients and consumers, as well as wholesale, hospital and retail levels as appropriate.