System for Australian Recall Actions

Recall detail

Type of Product ⁱ	Medical Device	
TGA Recall Reference ⁱⁱ	RC-2013-RN-01288-1	
Product Name/Description ⁱⁱⁱ	PLUM LifeCare 5000 Series and PLUM XL Familiy of Infusers	
The second	Plum LifeCare 5000 List Number: 02507	
	Plum XL List Number: 11555	
	Plum XLM List Number: 11846	
	Plum XLD List Number: 11859	
	ARTG Number: 138109	
Recall Action Level ^{iv}	Hospital	
Recall Action Classification ^v	Class I	
Recall Action Commencement Datevi	10/12/2013	
Responsible Entity ^{vii}	Hospira Pty Limited	
Reason / Issue ^{viii}	The door roller assembly on the Plum A Lifecare 5000 Series and Plum potential to break which can lead to possible unrestricted flow and/or ov the removal of the IV administration set's cassette from the pump.	
Recall Action ^{ix}	Recall for Product Correction	
Recall Action Instructions ^x	Hospira is requesting hospitals to take the following steps to inspect the prior to loading the cassette: -Open the cassette door by pulling on the lever - Unlatch the cassette door by pushing on the door release tab and pull -Visually inspect the door roller pin for any evidence of the damage or of misalignment.	ing the door down.
	-Ensure that the door roller spins smoothly with a finger touch. If any door rollers or pins appear losse, broken or missing, Hospira is a the device from use.	
Contractile	Hospira is in the process of retiring the Plum LifeCare 5000 and Plum	(L in 2015.
Contact Information ^{xi}	1300 046 774 (Option 1) - Hospira Quality Assurance	

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The TGA publishes Australian recall actions in a searchable database to ensure the public has access to information about therapeutic products that have been recalled from the Australian market. If you are concerned about your health or if you have experienced an adverse event please seek advice from a health professional as soon as possible. Please read all the important information at the beginning of this report.