

## Recall detail

Type of Product <sup>i</sup>	Medical Device
TGA Recall Reference <sup>ii</sup>	RC-2014-RN-00567-1
Product Name/Description <sup>iii</sup>	Disposable Ventstar 180cm Circuit (used in connection with the Dräger Oxylog 2000 Transport Ventilator)  Product Description: Packaging Box Set Part Number: 2M86841 Batch Numbers: 330384.001 - 331661.002  Product Description: Hose Packaging Part Number: 2M86791 Batch Numbers: 330384.001 - 331661.002  ARTG Number: 131796
Recall Action Level <sup>iv</sup>	Hospital
Recall Action Classification <sup>v</sup>	Class I
Recall Action Commencement Date <sup>vi</sup>	27/05/2014
Responsible Entity <sup>vii</sup>	Draeger Medical Australia Pty Ltd
Reason / Issue <sup>viii</sup>	Dräger has become aware of cases where prior to the use of the Disposable Ventstar Oxylog 2000 Ventilation Hoses, adhesive residue the size of a finger nail was found loose in the hose or the packaging around the hose; had the hose been used with the residue in place it may have entered the patient's lungs.
Recall Action <sup>ix</sup>	Recall
Recall Action Instructions <sup>x</sup>	Draeger is advising their customers to not use the hose system from the affected batches and to contact customer service to arrange for recovery and replacement.
Contact Information <sup>xi</sup>	03 9244 7248 - Draeger Medical Australia

## Footnotes

<sup>i</sup> Type of Product: Medicine, Medical Device, or Biological

<sup>ii</sup> TGA Recall Reference: Unique number given by the TGA

<sup>iii</sup> 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.

<sup>iv</sup> 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 /