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**ResMed Issues a Voluntary Recall for Certain S8 Flow Generators**

**FOR IMMEDIATE RELEASE** – San Diego, CA, April 23, 2007 – ResMed today announced a worldwide voluntary recall of approximately 300,000 of its early production S8 flow generators used for the treatment of obstructive sleep apnea. In S8 devices manufactured between July 2004 and May 15, 2006, there is a remote potential for a short circuit in the power supply connector. ResMed plans to work with its distribution partners globally to provide a repaired or if necessary, replacement device to patients who have an affected S8 flow generator.

Patients may continue to use their S8 flow generators until the affected units have been repaired. As with any electrical device, patients should make sure that it is placed on a hard clean surface and that the area around the device is clear during use. Patients should discontinue use of the device if there are any signs of electrical failure such as intermittent power, cracking sounds, sparking or charred smell.

ResMed does not recommend use of supplemental oxygen with an affected device. Patients currently using supplemental oxygen should immediately contact their physician to discuss whether to discontinue use of supplemental oxygen or discontinue using their device until a repaired device or replacement is received. In any event, patients on supplemental oxygen should immediately contact their equipment provider for a repaired or replacement device.

The recall includes the following serial number ranges for all S8 models:

From	To
20040285613	20060283743
20060287568	20060294694

ResMed voluntarily recalled the product after learning that in rare instances – less than two tenths of one percent (0.2%) – a short circuit in the power supply connector, a component supplied by a third party, has caused the devices to fail. In only seven cases worldwide, device failures have led to thermal damage to the device, with a remote potential to ignite material external to the device. No significant property damage or patient injury has been reported.

ResMed has advised the U.S. Food and Drug Administration and other regulatory authorities of this action. ResMed is continuing to discuss this action with those authorities and will finalize its proposed course of action after those discussions are concluded.

ResMed's S8 flow generators are distributed through medical equipment suppliers throughout the world. Affected products can be identified by the serial numbers on the bottom of each device.



ResMed is working in close partnership with its distribution partners and the medical community to ensure that patients are fully aware of the recall program and that patients who have an affected device will receive a repaired or if necessary, replacement S8 flow generator.

Patients will be contacted to arrange for their device to be repaired and are encouraged to visit [www.resmed.com/s8program](http://www.resmed.com/s8program) for more information.