

# Medical Device Alert

## Action

Ref: MDA/2009/073 Issued: 11 November 2009 at 15:00

### Device

Oxygen therapy.

Multifit nebulizer and Multifit nebulizer with BOC adaptor.

Manufactured by Teleflex.



### Problem

Failure to provide humidification due to a manufacturing defect. This could result in patients developing dry airways, causing discomfort and irritation.

The manufacturer is recalling specific batches. No other batch numbers are affected.

### Action

- Check batch numbers to identify whether you have any affected devices.
- Quarantine and do not use affected devices.
- Contact Teleflex to arrange for their replacement.

### Action by

All staff responsible for the use of these devices.

### CAS deadlines

Action underway: 25 November 2009  
Action complete: 09 December 2009

### Contact

**Manufacturer**  
Helene Sauvage  
Teleflex Medical  
Tel: 01494 532 761  
Fax: 01494 524 690  
E-mail: [hsauvage@teleflexmedical.com](mailto:hsauvage@teleflexmedical.com)

## Link to full Medical Device Alert

<http://www.mhra.gov.uk/Publications/Safetywarnings/MedicalDeviceAlerts/CON062677>