

# Medical Device Alert

## Action

Ref: MDA/2009/047 Issued: 27 July 2009 at 16:00

## Device

Stopcocks (catalogue numbers W20058 and W21372) and specific products containing non-lipid resistant stopcocks, manufactured by Arrow®.



## Problem

Risk of leakage or air embolism due to hub cracking at the female Luer connector during delivery of infusions containing lipids.

## Action

- Identify if you have any of the affected devices.
- Do not administer lipid-containing infusions using these devices.

## Action by

All Medical and nursing staff.

## Contact

### Manufacturer/supplier

Helen Sauvage

UK customer Service

Teleflex Medical

Tel: +353 (0) 90 646 0804

E-mail: [hsauvage@teleflexmedical.com](mailto:hsauvage@teleflexmedical.com)

## CAS deadlines

Action underway: 17 August 2009

Action complete: 07 September 2009

## Link to full Medical Device Alert

<http://www.mhra.gov.uk/Publications/Safetywarnings/MedicalDeviceAlerts/CON052020>