

Medical Device Alert

Ref: MDA/2012/005 Issued: 16 February 2012 at 15:00

Device

Stapler
Duet TRS™ universal straight
and articulating single use
loading units (SULU)



Manufactured by Covidien.

Problem

Covidien has issued [advice](#) following reports of serious injuries that have occurred following the use of these devices in the thoracic cavity.

These stapler loading units have the potential to injure adjacent anatomical structures within the thorax, which may result in life threatening post-operative complications.

Action

- Do not use these Duet TRS™ Loading Units for any adult or paediatric thoracic surgery procedures.
- Return any products purchased solely for thoracic use to the manufacturer, following the instructions in the manufacturer's [Field Safety Notice \(FSN\)](#).

Action by

Thoracic surgeons, all surgical and theatre staff involved in the use of these devices.

CAS deadlines

Action underway: 01 March 2012

Action complete: 08 March 2012

Contact

Manufacturer

Amanda Woolven
Covidien Commercial (UK) Ltd

Tel: 01329 224 435

Fax: 01329 224 418

Email: Amanda.woolven@covidien.com

Device

Product codes and descriptions of affected devices:

Product code	Description
DUET4535	Duet TRS™ 45 3.5mm straight SULU
DUET4535A	Duet TRS™ 45 3.5mm articulating SULU
DUET4548	Duet TRS™ 45 4.8mm straight SULU
DUET4548A	Duet TRS™ 45 4.8mm articulating SULU
DUET6035	Duet TRS™ 60 3.5mm straight SULU
DUET6035A	Duet TRS™ 60 3.5mm articulating SULU
DUET6048	Duet TRS™ 60 4.8mm straight SULU
DUET6048A	Duet TRS™ 60 4.8mm articulating SULU

Distribution

This MDA has been sent to:

- NHS trusts in England (Chief Executives)
- Care Quality Commission (CQC) (Headquarters) for information
- HSC trusts in Northern Ireland (Chief Executives)
- NHS boards in Scotland (Equipment Co-ordinators)
- NHS boards and trusts in Wales (Chief Executives)

Onward distribution

Please bring this notice to the attention of relevant employees in your establishment. Below is a suggested list of recipients.

Trusts

CAS and SABS (NI) liaison officers for onward distribution to all relevant staff including:

- Cardiothoracic departments
- Cardiothoracic surgeons
- Cardiothoracic surgery directors
- Clinical governance leads
- General surgeons
- General surgery
- General surgical units, directors of
- Paediatric surgeons
- Paediatric surgery, directors of
- Risk managers
- Supplies managers
- Theatre managers
- Theatre nurses
- Theatres
- Thoracic surgeons
- Thoracic surgery departments
- Thoracic surgery directors

Independent distribution

Establishments registered with the Care Quality Commission (CQC) (England only)

This alert should be read by:

- Hospitals in the independent sector
- Independent treatment centres

Please note: CQC and OFSTED do not distribute these alerts. Independent healthcare providers and social care providers can sign up to receive MDAs directly from the Department of Health's Central Alerting System (CAS) by sending an email to: safetyalerts@dh.gsi.gov.uk and requesting this facility.

Contacts

Manufacturer

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England

If you are in England, please send enquiries about this notice to the MHRA, quoting reference number **MDA/2012/005** or **2012/001/016/081/009**

Technical aspects

Sally Mounter
Medicines & Healthcare products Regulatory Agency
Floor 4
151 Buckingham Palace Road
London SW1W 9SZ

Tel: 020 3080 7168

Fax: 020 8754 3965

Email: sally.mounter@mhra.gsi.gov.uk

Clinical aspects

Nicola Lennard
Medicines & Healthcare products Regulatory Agency
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Email: nicola.lennard@mhra.gsi.gov.uk

How to report adverse incidents

Please report via our website <http://www.mhra.gov.uk>

Further information about **CAS** can be found at <https://www.cas.dh.gov.uk/Home.aspx>

Northern Ireland

Alerts in Northern Ireland will continue to be distributed via the NI SABS system.

Enquiries and adverse incident reports in Northern Ireland should be addressed to:

Northern Ireland Adverse Incident Centre
Health Estates Investment Group
Room 17
Annex 6
Castle Buildings
Stormont Estate
Dundonald BT4 3SQ

Tel: 02890 523 704

Fax: 02890 523 900

Email: NIAIC@dhsspsni.gov.uk

<http://www.dhsspsni.gov.uk/index/hea/niaic.htm>

How to report adverse incidents in Northern Ireland

Please report directly to NIAIC, further information can be found on our website <http://www.dhsspsni.gov.uk/niaic>

Further information about **SABS** can be found at <http://sabs.dhsspsni.gov.uk/>

Scotland

Enquiries and adverse incident reports in Scotland should be addressed to:

Incident Reporting and Investigation Centre
Health Facilities Scotland
NHS National Services Scotland
Gyle Square
1 South Gyle Crescent
Edinburgh EH12 9EB

Tel: 0131 275 7575

Fax: 0131 314 0722

Email: nss.irc@nhs.net

<http://www.hfs.scot.nhs.uk/online-services/incident-reporting-and-investigation-centre-irc/>

Wales

Enquiries in Wales should be addressed to:

Dr Sara Hayes
Senior Medical Officer
Medical Device Alerts
Welsh Assembly Government
Cathays Park
Cardiff CF10 3NQ

Tel: 029 2082 3922

Email: Haz-Aic@wales.gsi.gov.uk