

Once completed, please email this form to: nss.irc@nhs.scot

SUBJECT:

* Indicates a mandatory field. For more information about using this form [click here](#).

1. Reporter details	
Your Ref	
Your role*	
Profession*	
First name*	
Last name*	
Email address*	
Telephone number	

2. Workplace details	
The organisation which delivers the service where the incident happened	
Organisation name	
Incident location	
Incident location (if not on list)	

3. Local contact	
Please give details of someone who can be contacted by us or the manufacturer	
Designated contact*	
First name	
Last name	
Email	
Other contact information	

4. Product information	
Please don't send contaminated devices by mail - contact IRIC to make alternative arrangements	
Workstream*	
Operator of the device	
Device manufacturer name	
Equipment description*	
Type of device	
Any additional device or accessory that may also have been involved	
Unique device identifier (UDI)	

IRIC Adverse Incident Report Form

adverse incidents | near misses | safety concerns

4. Product information (continued)	
Model	
Catalogue number	
Serial number	
Lot or batch number	
Firmware version	
Software version	
Where is the device now? *	
Location if quarantined	
Containment / corrective actions	
Manufacturer/supplier contacted?	Yes No Unknown
Manufacturer's case Ref (if contacted)	

5. The incident		Please give as much information as possible about what happened			
Date incident occurred*					
Please describe what happened*					
Suspected problem					
Was any injury or illness involved?					
Health impact					
Additional comments					
Persons who were put at risk (tick all that apply)	Patient Visitor	Service user Carer	Staff Other	Contractor None	

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When to use of this form

This form is only to be used when it is not possible to use the standard webform, e.g. when local IT security measures prevent access. Refer to the [IRIC website](#) for more information on reporting adverse incidents.

This form can be used to report adverse incidents, near misses or safety concerns involving medical devices, in vitro diagnostic medical devices, estates, facilities, social care equipment and personal protective equipment used in Scotland's health and social care services. We will normally send you an email acknowledgement within three working days of receiving your report. However, please get in touch at nss.irc@nhs.scot if you haven't heard from us after five working days.

What we do with information you send us on this form

All incident reports become live data points on our trending system as soon as they are received. This helps us identify and prioritise the most pressing issues. However, we also individually assess and triage each incident report so, depending on the circumstances, we may open an investigation without waiting for a trend to develop. You can find out more by clicking here: [Incident Trending](#).

Regardless of how we triage your incident report we always recommend you do two things:

- report the incident to your local adverse event management system (often referred to as Datix)
- raise a complaint with the equipment supplier and send us a copy of their reply.

Who we share incident information with

We may share information on this form, e.g., with manufacturer or regulatory bodies, for the purposes of investigation and safety surveillance. We work in partnership with the Medicines and Healthcare products Regulatory Agency (MHRA) and we are responsible for informing them about medical device incidents occurring in Scotland.