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Miller rooth Dongs a root New card) B Saved queue	Description Do QUI enter any identifiable information, such as: - Names of people - Names of	TERTFY/LEVELS/IO PALIFIER BHT UP information from planning was not clear which resulted in the patients being on the bed a long time plum delays in treatment for other patients. The information provided did not clearly state that only the AF tailoo was used as a reference.		
	 Action taken Enter the action taken at the time of the incident. 	Checked the shifts on accordingly	oncentra and annotated the set up sheet	
	Incident classification			
	Category @	Patient Information (records, o	locuments, test results, scans) 📼	
	Detail	Information - other		
	Adverse event	Documentation (including reco	rds, identification) other	
	Is it MHRA reportable?			
	Additional Information			
	Was any equipment involved in the incident?	E.		
	Is this a Medical Devices related incident?		-	
	* Was this a medication incident?	No	*	
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College London Hosp	oitals NHS Foundation Trust - Incident Reportin	g Form - IE7 Virtual Browser fo	r use at UCLH			
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		expect if to happen or recur but it is possible that it might do so	•	•	•	•
		1 Rare - This will probably never happen/recur	•	•	•	•
			Grade: Low risk			
	NPSA NRLS - Please ensure you complete this secti	on -				
	Did any actions minimise the impact of the incident?	Yes	-			
	Additional information - For completion by Risk Management T	eam only -				
	Is this an SI?	No				
	Is a root cause analysis required?	No				
	Is an Action Plan required?	No	*			
	Does this incident need to be reported to the NPSA's NRLS?	Ves				
	Is this incident RIDDOR reportable?	No				
	CQC reportable?	No	-			
	Select any additional external agencies that need to be notified	National Patient Safety Age	ncy (NRLS)			
	CQUIN Reporting For completion by Risk Management T	eam only -				
	CQUIN Exempt?					
			Save Cancel			
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Scenario: On the final fraction of treatment for Mrs G's 3 field oesophagus plan (45Gy in 25#'s) the R&V link failed and disconnected during treatment of a LAO beam. The linac control box indicated 30 monitor units had been given. The R&V system was exited and the Linac control screen froze resulting in the R&V system not recording the delivered 30 monitor units that had been given. Floor superintendent was called to reinstate the beam. The plan had two LAO beams at the same gantry angle and the wrong one had been filled in to indicate the partial beam delivery on the treatment sheet. Radiographers became aware of error whilst selecting remaining beams to be treated. The calculated total dose discrepancy was deemed insignificant and no corrective action required. University College London Hospitals NHS ⁺UCL





