

Quality Improvement Plan

Developed following the unannounced monitoring assessment for the National Standards for the Prevention and Control of Healthcare Associated Infections by HIQA on August 9th 2016.

Approved by:

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01/12/2016



Tús Álte do Shábháilteacht Othar Patient Safety 2 First

Standard 3 - Environment and Facilities

The physical environment, facilities and resources are developed and managed to minimise the risk of service users, staff and visitors acquiring a Healthcare Associated Infection (HCAI).

Criterion 3.1 - The design and layout of the facility is based on needs assessment which reflects the size, complexity and specialties of the service provided.

Opportunities for Improvement	Agreed Actions	Responsibility	КРІ	Target Date
Improve the infrastructure and ventilation facilities in the Delivery Suite Operating Theatre	Develop design plans for a modular build extension incorporating the redevelopment of this theatre	Master and Property Committee	 Plans developed and submitted. 	• Q4 2016
	to meets required standards.			Target
	 Seek funding and planning permission for the modular build development. 	Secretary General Manager	 Funding and planning permission obtained. 	• Q1 2017
	Complete tender process for the development project	Secretary General Manager	Contractors appointed	• Q 4 2017
	 Construction and commissioning of the new facility 	• Master	 New theatre commissioned and operational 	• Q4 2018

Criterion 3.6 – The cleanliness of the physical environment is effectively managed and maintained according to relevant national guidelines and legislation; to protect service-user dignity and privacy and to reduce the risk of the spread of HCAIs.

Opportunities for	Agreed Actions	Responsibility	KPI	Target Date
Improvement		0.440: 0.1	200/linner an	- Noviac
 Ensure all patient equipment is clean and well maintained 	 Review the cleaning schedules for patient equipment and recording of same 	CMM3 in Delivery, IPC team	 >90% compliance on audits of medical equipment cleaning 	Nov'16, ongoing
	 Ensure equipment requiring replacement is identified and replaced 	Delivery Suite staff	 All equipment is of a satisfactory standard 	Ongoing
 Ensure point of care blood testing equipment is clean 	 Decontamination wipes are located beside the fetal blood analyser 	CMM3 in DS	 Decontamination wipes always available 	• Nov'16
	Point of care equipment checks to include cleanliness assessment	Point of Care Co- ordinator	100% POC checks completed	Dec 16 and Ongoing
 Ensure the Delivery Suite infrastructure and storage facilities are appropriate 	Finalise the plans for the DS refurbishment	 Master and Property Committee 	 Plans finalised and approved by Property Committee 	• Dec'16
	Tender for works and appoint contractor	 Secretary General Manager 	Contractor appointed	March '17
	Commence/complete the phased refurbishment of the Dept	• Master	Rooms renovated	• Dec '17

Standard 6 - Hand Hygiene

Hand hygiene practices that prevent, control and reduce the risk of the spread of Healthcare Associated Infections are in place.

Criterion 6.1 – there are evidence-based best practice policies, procedures and systems for hand hygiene practices to reduce the risk of the spread of HCAIs.

Opportunities for Improvement	Agreed Actions	Responsibility	КРІ	Target Date
Hand hygiene sinks comply with relevant standards	 See the QIPs identified under Standard 3 	Master	 Compliant hand hygiene sinks in D/S 	• Dec '17
			Compliant facilities in Theatre	• Dec '18

Standard 8 - Invasive Medical Device Related Infections

Invasive medical device related infections are prevented or reduced.

Criterion 8.1 – invasive medical devices are managed in line with evidence-based best practice and national and international guidelines.

Opportunities for Improvement	Agreed Actions	Responsibility	КРІ	Target Date
 Anaesthetic medications are prepared in accordance with best practice recommendations 	 Collaborate with Pharmacy to get pre prepared drugs available for emergency c/sections. 	Consultant Anaesthetist & Chairman, Department of Anaesthesia	 Anaesthetic drugs are constituted within appropriate timeframes 	• Mar '17
 Care bundle components are included in all intravenous cannulation / care policies 	Review and update relevant policies and guidelines	 IPC Team, Practice Development Team, Clinical Guidelines Committee 	 Policies are updated and distributed via Q- Pulse 	• Mar'17

Standard 11- Surveillance

Healthcare associated infections and antimicrobial resistance are monitored, audited and reported through a systematic surveillance programme.

Opportunities for Improvement	Agreed Actions	Responsibility	KPI	Target Date
Enhance surveillance of surgical site infection post caesarean section	 Increase surveillance timeframe up to 30 days post surgery 	IPC Team	Surveillance reports record incidents up to 30 days post surgery	Jan'17, ongoing