National Mastectomy & Breast Reconstruction Audit Datasheet - Mastectomy +/- Immediate Reconstruction

Patient Registration data				
Surname	Forename			
NHS/Private Hospital Number	er Date of birth			
Postcode	Ethnicity			
Patient-reported outcomes of	onsent			
Has this patient consented to	o being sent outcome questionnaires?			
☐ Patient has consented to receive questionnaires				
☐ Patient does not want to receive questionnaires				
☐ Patient judged incapable of completing a written questionnaire in English				
☐ Patient was capable	but not asked whether they were happy to receive questionnaire			
Reason patient was judged i	ncapable of completing the questionnaires (if applicable):			
□ Poor eyesight				
☐ Literacy or language comprehension problems				
☐ Cognitive impairment				
DO NOT SUBMIT DATA ELECTRONICALLY UNTIL THIS SECTION IS COMPLETED				
Previous treatment data				
Date of breast cancer diagno	sis:			
Date of decision to treat (mastectomy):				
Treatments for ipsilateral breast cancer prior to this admission (please select all that apply):				
☐ None	☐ Breast-conserving surgery			
☐ Axillary surgery (includ	ing Sentinel Node Bx)			
☐ Chemotherapy	☐ Hormone Therapy			
Co-morbidity data				
Smoking status:	ASA Grading (from pre-operative assessment):			
☐ Current smoker	☐ I – Normal healthy individual			
☐ Ex-smoker	☐ II – Mild systemic disease that does not limit activity			
□ Never smoked	☐ III – Severe systemic disease that limits activity but is not			
Body mass index:	incapacitating			
Weight/kg	☐ IV – Incapacitating systemic disease which is constantly			
Height/m	life-threatening			
BMI (W/H ²)	Pre-operative performance status (ECOG/WHO):			
Diabetes status:	☐ 0 - Fully active			
□ Not diabetic	1 - Light/office work			
☐ Type I diabetes	☐ 2 – Ambulatory / self care, up and about > 50% of the time			
☐ Type II diabetes	☐ 3 - Limited self care, confined to bed / chair > 50% waking hours			
	4 - Completely disabled, no self care and totally confined to bed /			
	chair			











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Operative data			
Date of admission for surgery:			
Date of mastectomy:			
Type of mastectomy (please select one opti	ion only):		
☐ Simple mastectomy			
☐ Subcutaneous or skin sparing mastect	omy via circumareolar approach (nipple excised)		
☐ Subcutaneous or envelope mastectomy via lateral or submammary approach (nipple spared)			
☐ Total mastectomy with excision of any part of pectoralis muscle			
☐ Total mastectomy with excision of both	n pectoral muscles + part of chest wall		
Type of axillary surgery (please select one	option only):		
□ None	☐ Level 1 axillary clearance		
☐ Sentinel node biopsy	Level 2 axillary clearance		
☐ Axillary sampling	Level 3 axillary clearance		
Type of immediate <u>primary</u> reconstruction performed (please select all that apply):			
□ None	☐ SIEA free flap		
☐ Tissue expander	☐ TDAP flap		
☐ Fixed volume implant	☐ TMG/TUG free flap		
☐ Latissimus Dorsi flap	☐ SGAP free flap		
☐ TRAM pedicle flap	☐ IGAP free flap		
☐ TRAM free flap	☐ Nipple reconstruction		
☐ DIEP free flap			
Type of contralateral symmetrisation surgery performed (please select all that apply):			
□ None	☐ Reduction mammoplasty		
☐ Tissue expander	☐ Mastopexy (skin reduction only)		
☐ Augmentation mammoplasty			
Diamad adjuvent tractments.			
Planned adjuvant treatments: Pl	anned <u>secondary</u> reconstructive procedures:		
Radiotherapy	☐ Tissue expansion of breast mound		
☐ Chemotherapy	Exchange of expander for fixed volume implant		
☐ Hormone therapy	☐ Nipple reconstruction		
Specialist palliative care	☐ Areolar tattooing		
	☐ Symmetrisation procedure		
	☐ Exchange of implant/expander for autologous flap		











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Reconstructive decision-making data PLEASE COMPLETE IF IMMEDIATE RECONSTRUCTION HAS NOT BEEN PERFORMED ☐ Yes Was immediate reconstruction offered to this patient? If immediate reconstruction was not offered, why was this? (please select all that apply) Patient appropriateness for surgery: ☐ Advanced stage of disease ☐ Concerns about local recurrence ☐ Age of patient ☐ Degree of co-morbidity (e.g. cardio-respiratory disease) ☐ Lifestyle factors (e.g. smoking) ☐ Cognitive impairment ☐ Mental health issues (e.g. psychiatric illness) Treatment pathway issues: ☐ Patient has undergone recent neo-adjuvant chemotherapy ☐ Adjuvant radiotherapy to chest wall anticipated for this patient ☐ Reconstructive surgery would delay other anticipated adjuvant therapies Service access issues: ☐ Immediate reconstruction not available locally ☐ Immediate reconstruction would significantly delay mastectomy surgery ☐ Yes ☐ No Has delayed reconstruction been offered to this patient? ☐ Yes ☐ No If yes, have they accepted the offer? If delayed reconstruction has not been offered, why is this? (please select all that apply) Patient appropriateness for surgery: ☐ Advanced stage of disease ☐ Concerns about local recurrence ☐ Age of patient ☐ Degree of co-morbidity (e.g. cardio-respiratory disease) ☐ Lifestyle factors (e.g. smoking) ☐ Cognitive impairment ☐ Mental health issues (e.g. psychiatric illness) Service access issues: ☐ Delayed reconstruction not available locally











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Peri-operative morbidity data				
Date of discharge:				
Return to theatre during admission	☐ Yes	□ No		
Emergency transfer to HDU or ITU during admission	☐ Yes	□ No		
Death during admission	☐ Yes	□ No		
In-patient complications (please select all that apply):				
Complications requiring therapeutic intervention at:	Mastecto			
	site	site (if applicable)		
None				
Wound infection requiring intravenous antibiotics				
Wound infection requiring surgical debridement				
Skin flap necrosis requiring surgical debridement				
Wound dehiscence requiring re-closure				
Haematoma or seroma requiring aspiration or drainage				
Flap-related complications requiring therapeutic intervention	on:			
Not applicable				
None				
Impaired flap perfusion requiring re-exploration or revision	of anastomosis			
Partial flap necrosis or failure requiring debridement				
Total flap necrosis or failure requiring removal				
Implant/expander-related complications requiring therapeutic intervention:				
Not applicable				
None				
Displaced implant/expander requiring re-positioning				
Infected implant/expander requiring intravenous antibiotic therapy				
Infected implant/expander requiring removal				
Ruptured implant/expander requiring removal				
Distant or systemic complications requiring therapeutic intervention:				
None				
Haemorrhage requiring blood transfusion				
Deep venous thrombosis (DVT) requiring formal anticoagul	ation			
Pulmonary embolism (PE) requiring formal anticoagulation				
Acute myocardial infarction (MI) requiring anticoagulation +	/- thrombolysis			











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Pathology data (from post-operative histology report)				
Tumour laterality:	□ Right	Left		
Invasive status:	\square Invasive	☐ DCIS (ductal carcinoma in situ)		
Grade of DCIS or Invasive Carcinoma: ☐ 1 – low (DCIS) or well differentiated (invasive) ☐ 2 – intermediate (DCIS) or moderately differentiated (invasive) ☐ 3 – high (DCIS) or poorly differentiated (invasive)				
Lymph node involvement:				
(/)				
(number of positive axillary nodes / total number of axillary nodes in pathology specimen)				
Invasive lesion size (mm): Recorded Nottingham Prognostic Index Score (if invasive):				









