## PRESSURE ULCERS (PU) SIGNIFICANT ADVERSE EVENT (SAE) REVIEW TEMPLATE

Please read the Adverse Event Review Protocol before initiating this review.

Please use this Significant Adverse Event (SAE) template for **review of all NHS Lothian acquired Grade 4 Pressure Ulcers**: For all grade 4 Pressure Ulcers acquired outwith NHS Lothian care, refer to Pathway for reviewing pressure ulcers in Datix record

- 1. **Complete Appendix A** to provide relevant information for inclusion in the report.
- 2. Include details of hospital/ward or community nursing base
- 3. Reviewer(s) names and designations must all be included in full, in designated boxes.
- 4. **Do not include person-identifiable information in the body of the report** identify persons by patient A/ doctor B.
- 5. Save the template securely on your NHS shared drive until complete.
- 6. Once completed, update the document control details (within the footer) and attach document to the relevant Datix adverse event as 'Draft'.

Please complete Appendix A before completing this section

		•				
DATIX ID Number:	Was the pressure ulcer acquired under NHS Care If so, ward/DN/own home	Site, Harm Level & Outcome e.g. Hip, Grade 4 - Major	Date PU noted:			
Reviewer(s):	Date Review Started:	Date Review Completed:	Reported to: Whoever commissioned the review, Names and Designations			
Date referred to Tissue Via (if not, reasons)	ability:	L				
When and how did this (What has changed recently)	Pressure Ulcer occur?					
Background/Timeline to to the event, include date(s).						
Actions Taken as a Con	sequence of the Assess	ment(s)·				
(Was there a current care pla frequency)	-	• •	e, equipment, skin check			
Family/Next of Kin Aware (with patient consent):						
(Date and time, family informed of PU. Were family involved / invited to be involved in the review process?  Please ensure that any questions the family have are considered as part of the review.)						
, 4	,	,	,			

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Review C	Review Outcome (Please indicate the outcome of the review using the codes below)							
1.		opriate care - well plann ery problems identified	ed and de	elivere	ed w	ith no care or ser	vice	
2.	Care or service delivery problems were identified and lessons can be learned though this did not affect the final outcome/ event.							
3.	A diff	erent plan and/or delive	ry of care	may l	have	e resulted in a dif		
4.								
7.	A different plan and or delivery of care, on balance of probability, would have been expected to result in a more favourable outcome, i.e. how case							
		managed had a direct in				•		
After revi	ew d	o you consider the P	U report	ed w	as e	either: (*Definition	ons at end of ter	mplate)
	ng) e.g	ne aspect of NHS asses g. Outcome 3 or 4: Cons				idable* (all care affected develop	_	
Factors v	vhich	could have contribu	ited to th	ne PU	J			
	_	ppendix A contributed				er? This information	on can then be	used to
		op the improvement plar						
Factor Ty				Fa	acto	ors relevant to	the PU	
Patient fac	ctors							
Social Fac	ctors							
Task and	Techn	ology factors						
Individual	(staff	) factors /Team factors	5					
Work Env	ironm	ent Factors						
Organisat	ional	and Management facto	ors					
Please no	ote go	ood practice below:						
Lessons	to Le	arn and Improvemer	nts to be	made	e			
Improvem	ents	Level of Recommendation (Individual, Team, Service Directorate, Organisation)	By Whom	By Whe		Resource Requirements	Evidence of Completion	Completion Sign-off
Author:	Author: Date:							

This SAE <u>cannot</u> be closed on Datix until this report has completed the formal governance approval process. The SAEs will be closed at the end of the process by Quality Improvement Support Team staff.

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This sign-off process can be found on the NHS Lothian intranet under: Healthcare > A-Z > Risk Management > SAE Sign-off process

SIGNED OFF AT OPERATIONAL LEVEL BY: (IF SIGNIFICANT ADVERSE EVENT):				
Acute: Site/Service Director (or nominated member of CMT)  REAS: Director of Operations	Signed:			
H&SCP: Joint Director (or nominated member of JMT)	Date:			
Acute: Site/Services Nurse/Medical Director	Signed:			
REAS: Associate Medical Director/ Chief Nurse	Date:			
H&SCP: Clinical Director/Chief Nurse				
FINAL APPROVAL AT BOARD LEVEL BY:				
NHS Lothian Board Medical Director	Signed: Date:			
NHS Lothian Board Nurse Director	Signed: Date:			

## Appendix A – complete using patient notes/EPR

Factors which could have contributed to the Pressure Ulcer (PU)

Consider factors listed and identify those which contributed to the PU. This information can then be used to inform and develop your improvement plan.

Patient Factors: Were there any underlying reasons why this patient was more likely to develop a PU?	Y	N	NA	Comments/additional information
Reduced mobility				
Sensory / Cognitive impairment				
Pain				
Extremes of age e.g. young babies, older people				
Extremes of size e.g. obese, very thin				
Underlying circulatory problems e.g. PAD, CVA,				
Spinal injury				
Malnutrition etc.				
Previous healed pressure ulcer in same location on body				
Previous surgery to area affecting tissues e.g. flap surgery				
Increased moisture on skin e.g. incontinence, sweat, other				
body fluids				
Were there any issues identified with patient/carer/family which contradicted the proposed care plan? (e.g. refusing equipment, not agreeing to skin checks or position changes etc.) please specify  If yes, was it documented that advice and support was given				
regarding the need to follow proposed care plan?				
Social factors : External factors related to social situation which may have impacted on PU development	Υ	N	NA	Comments/additional info.
Chaotic lifestyle				
People who inject drugs				
Lack of support				
Interaction with healthcare services etc.				
Was the patient/carer/family involved in the care plan				
development? please specify				
Was the patient/carer/family involved in the delivery of the				
patient's goal(s)? please specify				
Task/Technology factors: Were all aspects of assessment, care planning and review in place?	Υ	N	NA	Comments/additional information
Was Waterlow/PPURA (adults) or Glamorgan (children) risk assessment carried out within 6 hours of admission to an inpatient setting OR first clinician visit in community  If Waterlow/Glamorgan score was ≥10 was a SSKIN Bundle				
or Care Rounding Tool in place for PU prevention?				
Was the Waterlow/Glamorgan/PPURA reassessed according				
to patient's condition or on transfer to another area? e.g.				
changes in patient condition, pre and post- op	1			
Was a PU prevention care plan in place when risk identified?	1			
Was PU care plan followed and updated as changes to patient condition were noted?				
Was the pressure ulcer graded according to the Scottish Adaptation of the EPUAP grading tool?				
Were Care Rounding /SSKIN bundle element review times appropriate to level of risk/skin damage identified?				

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<b>Skin -</b> Was skin assessed as per prescribed time intervals/care plan?				
Was a wound chart completed, accurate and updated?				
Was the wound treatment plan appropriate?				
Surface- Was the patient's mattress suitable for their condition?				
If patient was sitting up, was the chair and/or cushion suitable for the patient?				
Keep Moving - Was repositioning/off-loading carried out as				
per prescribed time intervals/care plan?				
Incontinence - If patient was incontinent has a review of bladder/bowel function/management taken place? Were appropriate skin protectors used as per formulary?				
<b>Nutrition</b> - If patient required additional nutritional support, did they receive this? e.g. assistance with meal, supplements				
Individual/Team factors:	Y	N	NA	Comments/additional information
Was Patient/carer 'Prevent PU' leaflet and/or advice given by staff?				
Staff trained in PU prevention including skin assessment, use of equipment etc.				
Were there any issues around written communication /documentation/ notes?				
Were there any other team factors that may have contributed to the PU?				
Work Environment factors: Equipment working and in place in a timely manner: If no, please explain	Y	N	NA	Comments/additional information
Therapeutic mattresses available e.g. foam (pentaflex), static air (repose), alternating pressure (nimbus etc), low air loss (breeze, ambience)				
Cushions e.g. foam (propad), foam + gel/polymer (flotech, dynatech gel), altering pressure (Aura, Talley BASE)				
Hoist plus appropriate slings				
Glide sheets or other transfer aids				
If patient required additional pressure redistribution equipment was this available? (e.g. heel protectors / heel elevators, dermal-type replacement pads/strips)				
If pressure damage is related to a medical device, was there a care plan in place to reduce associated risk?				
Organisation and Management factors: Were the following available if required:	Υ	N	NA	Comments/additional information
Extra equipment purchased e.g. heel protectors(Repose), heel lift boots, cushions				
Nutritional support e.g. dietetic review, dietary supplements,				
special diet, etc. Specialist input e.g. TVN, Physio, OT, continence team etc.				
Senior management involvement				
Local management structures				
Consider if patient has been boarded and impact on				
communication etc.				
Were there any other organisational and management factors	1	+	1	+
that may have contributed to the PU?				

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e.g. co-morbidities, over activity, non-compliance issues etc.			
Completed by (Name & Designation):	Date:		

## **Definition of Avoidable/Unavoidable Pressure Ulcer**

Avoidable Pressure Ulcer
The person receiving the care developed a pressure ulcer and the provider of care did not do one of the following:

- Evaluate the person clinical condition and pressure ulcer risk factors;
- Plan and implement interventions that are consistent with the persons needs and goals and recognised standards of practice;
- Monitor and evaluate the impact of the interventions; or revise the interventions as appropriate.

Unavoidable Pressure Ulcer
The person receiving the care developed a pressure ulcer even although the provider of the care did the following;

- Evaluated the person's clinical condition and pressure ulcer risk factors;
- Planned and implemented interventions that were consistent with the person's needs and goals;
- Recognised standards of practice; monitored and evaluated the impact of the interventions; and revised the approaches as appropriate;
- There is documented evidence the individual person refused to adhere to prevention strategies in spite of education of the consequences of nonadherence.

Reference: Tissue Viability Society. Achieving Consensus in Pressure Ulcer Reporting. JTV 2012