

	Event ID:	
Initial Report Date	e (HERF Q1):	

Patient Safety Event Report - Hospital:





PATIENT INFORMATION FORM (PIF)

Use this form only if you are reporting an incident. (When reporting a perinatal incident that affected a mother and a neonate, complete a PIF for the mother and a separate PIF for the neonate.) Highlighted fields are collected for local facility and PSO use. This information will not be forwarded to the Network of Patient Safety Databases (NPSD).

Tac:	uity and PSO use. This information	on will not be forwarded to the Network of Patient Safety Databases (NPSD).					
1.	1. At the time of the event what was the patient's age? CHECK ONE:						
	a. Neonate (0-28 days)	f. Mature adult (65-74 years)					
	b. Infant (>28 days <1 year	g. Older adult (75-84 years)					
	c. Child (1-12 years)	h. Aged adult (85+ years)					
	d. Adolescent (13-17 years)	i. Unknown					
	e. Adult (18-64 years)						
2.	Is the patient's ethnicity Hispan	ic or Latino? CHECK ONE:					
	a. Hispanic or Latino						
	b. Not Hispanic or Latino						
	c. Unknown						
3.	What is the patient's race? CHEC						
	a. American Indian or Alas						
	b. Asian	f. More than one race					
	c. Black or African Americ	6 —					
	d. Native Hawaiian or Othe	er Pacific Islander					
4.	Enter the patient's ICD-9-CM or	ICD-10 CM principal					
7.	diagnosis code at discharge (if						
		ICD-9-CM OR ICD-10-CM CODE					
_	Wee any intervention attempted	lim and an to "manage" the motion to the property to minimize out a manage hours.					
5.	CHECK ONE:	I in order to "rescue" the patient (i.e., to prevent, to minimize, or to reverse harm)?					
	a. Yes 6. Which of the following interventions (rescue) were documented?						
	b. No	CHECK ALL THAT APPLY:					
	c. Unknown	a. Transfer, including transfer to a higher level care area within facility,					
	_	transfer to another facility, or hospital admission (from outpatient)					
b. Monitoring, including observation, physiological examination, laborat							
		testing, phlebotomy, and/or imaging studies					
		c. Medication therapy, including administration of antidote, change in					
		pre-incident dose or route d. Surgical/procedural intervention					
		e. Respiratory support (e.g., ventilation, tracheotomy)					
		f. Blood transfusion					
		g. Counseling or psychotherapy					
		h. Unknown					
		i. Other intervention: PLEASE SPECIFY					

		Initial Report Dat	e (HERF Q1):				
7.	After discovery of the incident, and any subsequent intervention, what was the extent of harm to the patient (i.e., extent to which the patient's functional ability is expected to be impaired subsequent to the incident and any attempts to minimize adverse consequences)? CHECK FIRST APPLICABLE: AHRQ Harm Scale						
	a. Death: Dead at time of assessment.		ANSWER QUESTION 9				
	b. Severe harm: Bodily or psychological injury (including functional ability or quality of life.		ement) that interferes significan	tly with			
	c. Moderate harm: Bodily or psychological injury adversat the level of severe harm.	sely affecting fund	ctional ability or quality of life,	but not			
	d. Mild harm: Minimal symptoms or loss of function, or injury limited to additional treatment, monitoring, and/or increased length of stay.						
	e. No harm: Event reached patient, but no harm was event. Unknown	ident.	ANSWER QUESTION 9				
3.	What is the anticipated duration of the harm to the patient?	CHECK ONE:					
	a. Permanent (one year or greater)						
	b. Temporary (less than one year)						
	c. Unknown						
9.	Approximately when after discovery of the incident was harm	n assessed? CHEC	K ONE:				
	a. Within 24 hours						
	b. After 24 hours but before 3 days						
	c. Three days or later						
	d. Unknown						
LO .	D. Did, or will, the incident result in an increased length of stay	? CHECK ONE:					
	a. Yes						
	b. No (or not anticipated)						
	c. Unknown						
11.	L. After the discovery of the incident, was the patient, patient's	family, or guardi	an notified? CHECK ONE:				
	a. Yes						
	b. No						
	c. Unknown						

Event ID: ___

Thank you for completing these questions.

OMB No. 0935-0143 Exp. Date 10/31/2014

Public reporting burden for the collection of information is estimated to average 15 minutes per response. An agency may not conduct or sponsor, and a person is not required to respond to, a collection of information unless it displays a currently valid OMB control number. Send comments regarding this burden estimate or any other aspect of this collection of information, including suggestions for reducing this burden, to: AHRQ Reports Clearance Officer, Attention: PRA, Paperwork Reduction Project (0935-0143), AHRQ, 540 Gaither Road, Room #5036, Rockville, MD 20850.