

## **Serious Reportable Event (SRE)**

Submit no later than (7) seven calendar days from discovery of event

Please complete all sections of this form and submit to the Patient Safety Surveillance & Improvement System (PSSIS) administered by Vermont Program for Quality in Health Care, Inc. via secure email at <a href="mailto:sre@vpqhc.org">sre@vpqhc.org</a>.

For questions regarding the Patient Safety Surveillance & Improvement System (PSSIS) contact:

Vermont Program for Quality in Health Care, Inc. (VPQHC)

132 Main Street Montpelier, VT 05602

Phone: 802-229-2152 Email: sre@vpqhc.org

1.	Facility Identification
	Facility name:
2.	<b>Contact Information</b>
	Title of person submitting report:
	Telephone number:
3.	Patient Information
	Date of facility encounter/admission:
	Patient age:
4.	When did the event occur?
	Date the event occurred:
	Date the hospital Patient Safety Staff became aware of the event:
	Date of event report to the (PSSIS):

**5. Brief factual narrative about event:** (*If you prefer, you may attach a separate document containing this information.*)

Revised 08/2024



## Patient Safety Surveillance & Improvement System

6.	Where did the event occur?	
	☐ Emergency Department	☐ Labor and Delivery
	☐ Medical/Surgical Floor	☐ Radiology
	☐ Intensive Care Unit	☐ Surgical Services Department
	☐ Pediatrics	☐ Laboratory
	Grounds	Other:
	☐ Inpatient Psychiatry	
7.	What Happened?  National Quality Forum (NQF): List  Serious Reportable Events in Healthc	of Serious Reportable Events (SREs) <u>NOF:</u> are 2011 (qualityforum.org)
	<b>Surgical or Invasive Procedure</b>	Events
	☐ <b>A.</b> Surgery or other invasive pro	ocedure performed on the wrong site.
	☐ <b>B.</b> Surgery or other invasive pro	ocedure performed on the wrong patient.
	☐ C. Wrong surgical or other inva	asive procedure performed on a patient.
	□ <b>D.</b> Unintended retention of a fo	reign object in a patient after surgery or
	other invasive procedure.	
	☐ E. Intra-operative or immediate an ASA Class I patient.	ely postoperative/ post procedure death in
	<b>Product or Device Events</b>	
	☐ A. Patient death or serious injur	y associated with the use of contaminated
	drugs, devices, or biologics p	rovided by the healthcare setting.
	☐ <b>B.</b> Patient death or serious injur	y associated with the use or function of a
	device in patient care, in which as intended.	ch the device is used or functions other than
	☐ C. Patient death or serious injury	y associated with intravascular air embolism for in a healthcare setting.

Revised 08/2024 2



Patient Protection Events
☐ A. Discharge or release of a patient/resident of any age, who is unable to
make decisions, to other than an authorized person.
☐ <b>B.</b> Patient death or serious injury associated with patient
elopement (disappearance).
☐ C. Patient suicide, attempted suicide, or self-harm that results in serious
injury, while being cared for in a healthcare setting.
Care Management Events
☐ <b>A.</b> Patient death or serious injury associated with a medication error (e.g.,
errors involving the wrong drug, wrong dose, wrong patient, wrong time,
wrong rate, wrong preparation, or wrong route of administration).
☐ <b>B.</b> Patient death or serious injury associated with unsafe administration of
blood products.
☐ C. Maternal death or serious injury associated with labor or delivery in a low
risk pregnancy while being cared for in a healthcare setting.
$\Box$ <b>D.</b> Death or serious injury of a neonate associated with labor or delivery in a
low-risk pregnancy.
☐ E. Patient death or serious injury associated with a fall while being cared for
in a healthcare setting.
☐ <b>F.</b> Any Stage 3, Stage 4, and unstageable pressure ulcers acquired after
admission/presentation to a healthcare setting.
$\Box$ <b>G.</b> Artificial insemination with the wrong donor sperm or wrong egg.
$\Box$ <b>H.</b> Patient death or serious injury resulting from the irretrievable loss of an
irreplaceable biological specimen.
$\square$ I. Patient death or serious injury resulting from failure to follow up or
communicate laboratory, pathology, or radiology test results.
<b>Environmental Events</b>
☐ A. Patient or staff death or serious injury associated with an electric

Revised 08/2024 3



## Patient Safety Surveillance & Improvement System

	shock during a patient care process in a healthcare setting.
	B. Any incident in which systems designated for oxygen or other gas
	to be delivered to a patient contains no gas, the wrong gas, or are contaminated by toxic substances.
	C. Patient or staff death or serious injury associated with a burn
	incurred from any source during a patient care process in a healthcare setting.
	<b>D.</b> Patient death or serious injury associated with the use of physical
	restraints or bedrails while being cared for in a healthcare setting.
Ra	diological Events
	A. Death or serious injury of a patient or staff associated with the
	introduction of a metallic object into the MRI area.
Pու	tential Criminal Events
	A. Any instance of care ordered by or provided by someone
Ш	, ,
	impersonating a physician, nurse, pharmacist, or other licensed
	healthcare provider.
	<b>B.</b> Abduction of a patient/resident of any age.
	C. Sexual abuse/assault on a patient or staff member within or on the
	grounds of a healthcare setting.
	<b>D.</b> Death or serious injury of a patient or staff member resulting from a
	physical assault (i.e., battery) that occurs within or on the grounds of a healthcare setting

Revised 08/2024