

EVENTS FOR SERIOUS SAFETY EVENT (SSE) REVIEW

TGH Risk Department contact name: * . Email: * . Phone #: .

SERIOUS SAFETY EVENT WORKSHEET

Event Date:		MRN:
Event Report Date:	Event Number:	Name:

SERIOUS SAFETY EVENT CATEGORY

<p>SURGICAL/INVASIVE PROCEDURE EVENTS</p> <ul style="list-style-type: none"> <input type="checkbox"/> Surgery or other invasive procedure performed on the wrong site **† <input type="checkbox"/> Surgery or other invasive procedure performed on wrong patient**† <input type="checkbox"/> Wrong surgical or other invasive procedure performed on a patient <input type="checkbox"/> Unintended retention of a foreign object in a patient after surgery or other invasive procedure *† <input type="checkbox"/> The performance of a surgical procedure that is medically unnecessary or otherwise unrelated to the patient's diagnosis or medical condition <input type="checkbox"/> The surgical repair of damage resulting to a patient from a planned surgical procedure, where the damage is not a recognized specific risk, as disclosed to the patient and documented through the informed-consent process <input type="checkbox"/> Intraoperative or immediately postoperative/post-procedure death in an ASA Class 1 patient 	<p>PRODUCT OR DEVICE EVENTS</p> <ul style="list-style-type: none"> <input type="checkbox"/> Patient death or serious injury associated with the use of contaminated drugs, devices, or biologics provided by the healthcare setting <input type="checkbox"/> Patient death or serious injury associated with the use or function of a device in patient care, in which the device is used or functions other than as intended <input type="checkbox"/> Patient death or serious injury associated with intravascular air embolism that occurs while being cared for in a healthcare setting <p>RADIOLOGIC EVENTS</p> <ul style="list-style-type: none"> <input type="checkbox"/> Death or serious injury of a patient or staff associated with the introduction of a metallic object into the MRI area <input type="checkbox"/> Prolonged fluoroscopy with cumulative dose >1500 rads to a single field, or any delivery of radiotherapy to the wrong body region or >25% above the planned radiotherapy dose† 	<p>PATIENT PROTECTION EVENTS</p> <ul style="list-style-type: none"> <input type="checkbox"/> Discharge or release of a patient/resident of any age, who is unable to make decisions, to other than an authorized person*† <input type="checkbox"/> Patient death or serious injury associated with patient elopement (disappearance) <input type="checkbox"/> Patient suicide, attempted suicide, or self-harm that results in serious injury, while being cared for in a healthcare setting, or within 72 hours of discharge*† <p>CARE MANAGEMENT EVENTS</p> <ul style="list-style-type: none"> <input type="checkbox"/> 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) <input type="checkbox"/> Patient death or serious injury associated with unsafe administration of blood products <input type="checkbox"/> Maternal death or serious injury associated with labor or delivery in a low-risk pregnancy while being cared for in a healthcare setting <input type="checkbox"/> Death or serious injury of a neonate associated with labor or delivery in a low-risk pregnancy <input type="checkbox"/> Patient death or serious injury associated with a fall while being cared for in a healthcare setting <input type="checkbox"/> Any Stage 3, Stage 4, and unstageable pressure ulcers acquired after admission/presentation to a healthcare setting <input type="checkbox"/> Artificial insemination with the wrong donor sperm or wrong egg <input type="checkbox"/> Patient death or serious injury resulting from the irretrievable loss of an irreplaceable biological specimen <input type="checkbox"/> Patient death or serious injury resulting from failure to follow up or communicate laboratory, pathology, or radiology test results <input type="checkbox"/> Severe neonatal hyperbilirubinemia (bilirubin >30 mg/dL) † <input type="checkbox"/> Death of a patient as a result of an adverse event <input type="checkbox"/> Brain or spinal damage to a patient as a result of an adverse event
<p>ROOT CAUSE DETERMINATION:</p>		

After analysis, was this event considered preventable? Yes No

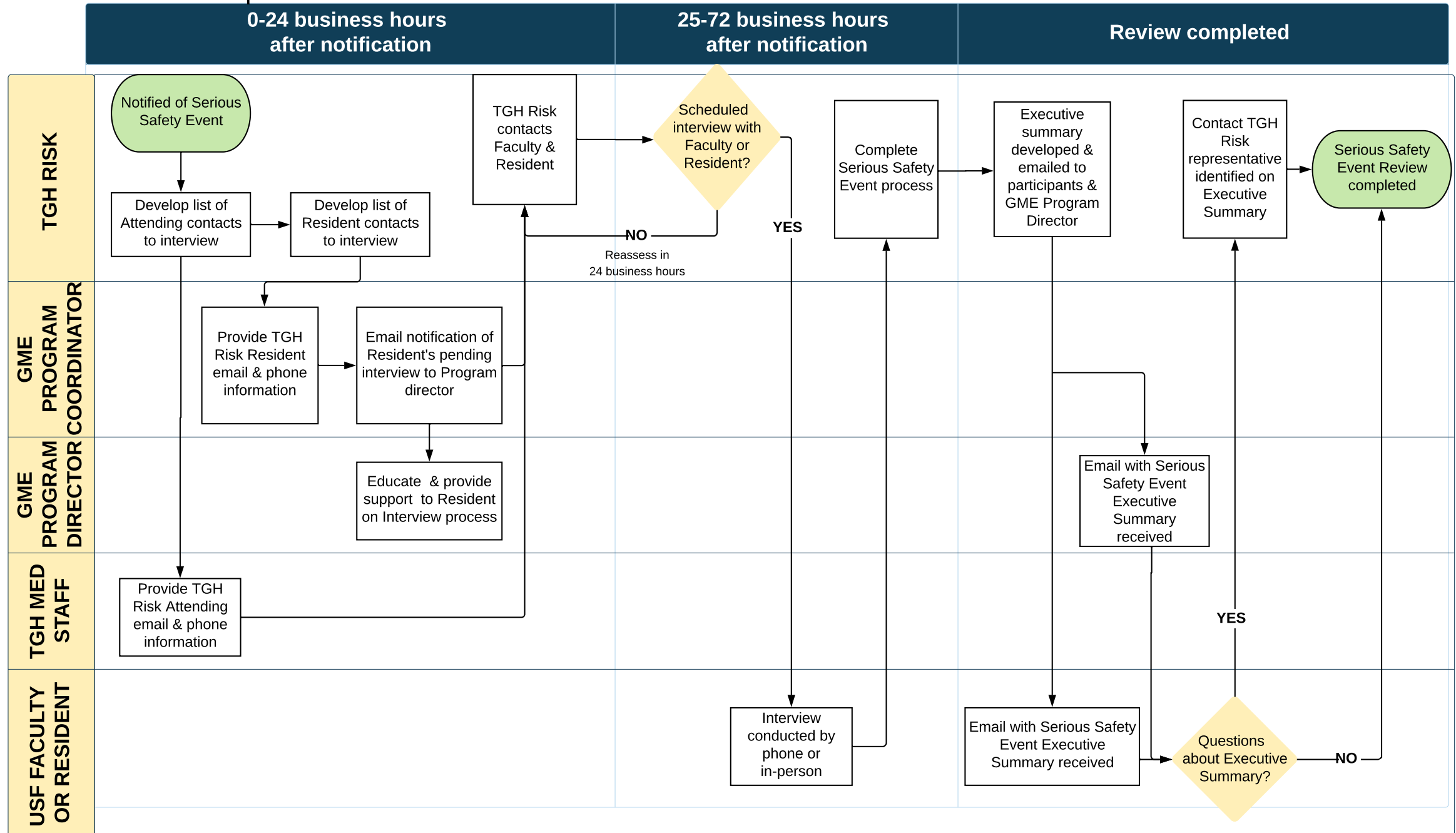
Disclosure and follow-up to the family? Yes No

†Joint Commission Sentinel Event; **Sentinel Event and NQF Never Event; AHCA Code 15 in red

PROCESS FOR SERIOUS SAFETY EVENTS AT TGH

v2. 1/2019

TGH Risk Department contact name: * . Email: * . Phone # .



INFORMATION FOR SERIOUS SAFETY EVENT (SSE) INTERVIEW

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EVENT TIMELINE

Date & Time	Source (Med Record, interview, etc.)	Event (Description & Response) What happened?	Deviation from Expected Practice (if applicable) What normally happens? What do policies/procedures require?	What usually happens? How often does this type of deviation occur (rare, common, very frequently)?		
		<p style="color: red; border: 2px solid red; padding: 10px;"> - TGH's Risk Department will contact any USF Faculty and Residents who were present during the Serious Safety Event. - TGH's Risk Department will email and/or phone to schedule Serious Safety Event Review interviews (phone or in-person) that will last approximately 10-15 minutes. USF GME and TGH expect Faculty and Residents to respond to TGH's Risk Department within 48 business hours unless on official leave. - Open-ended questions will be asked to complete the Event Timeline. - Any USF Faculty and Residents interviewed will be emailed an Executive Summary of the Serious Safety Event once an action plan has been developed. Any questions regarding this Executive Summary can be directed to TGH's Risk Department (Contact: *, Phone #*). The Resident's Program Director should be cc'd on any email sent to Residents. </p>				