

Outpatient Procedure Component Surgical Site Infection (SSI) Event

This form is used for reporting data on each patient having a SSI event related to one of the NHSN operative procedures selected for monitoring.

Instructions for this form are available at: https://www.cdc.gov/nhsn/forms/instr/57.405-toi.pdf.

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Facility ID:		Event #:			
*Patient ID:		Social Security #:			
Secondary ID #:		Medicare #:			
Patient Name, Last:		First:		Middle:	
*Gender: F M Other		*Date of Birth:			
Birth Sex: F M Unknown		Gender Identity (Specify):			
Ethnicity (Specify):		Race (Specify):			
*Date of Encounter (MM/DD/YYYY):					
Surgical Site Infection (SSI)					
*Event Type: <u>SSI</u>					
*Date of Event:// *Primary CPT Code: *NHSN Procedure Code:					
*SSI Level:					
□ Superficial Incisional Primary (SIP) □ Deep Incisional Primary (DIP) □ Organ/Space					
Superficial Incisional Se	condary (SIS)	Deep Incision	onal Secondary (DIS)		
*Specify SSI Criteria Used (check all that apply):					
Signs & Symptoms			Laboratory		
	□ Localized swelling		Organism(s) identified		
Erythema or redness	□ Pain or tenderness		Culture or non-culture based testing not performed		
□ Fever (>38°C)	Purulent draina	age	Imaging test evidence of infection		
Heat	Sinus tract				
			□ Organism(s) identified from ≥		
Incision deliberately	Wound sponta	neously	periprosthetic specimens		
opened/drained dehisce			Other positive laboratory test		
			<u>Clinical Diagnosis</u>		
Other evidence of infection found on invasive gross anatomic exam, or histopathologic exar					
		11	Diagnosis of superficial SSI by surgeon or physician		
*Pathogens Identified:	Yes 🗆 No				
If Yes, indicate up to 3 path	logens:				
				Continue>>>	
Assurance of Confidentiality: The voluntarily provided information obtained in this surveillance system that would permit identification of any individual or institution is collected with a guarantee that it will be held in strict confidence, will be used only for the purposes stated, and will not otherwise be disclosed or released without the consent of the individual, or the institution in accordance with Sections 304, 306 and 308(d) of the Public Health Service Act (42 USC 242b, 242k, and 242m(d)).					
Public reporting burden of this collection of information is estimated to average 21 minutes per response, including the time for reviewing instructions, searching existing data sources, gathering, and maintaining the data needed, and completing and reviewing the collection of information. 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 CDC, Reports Clearance Officer, 1600 Clifton Rd., MS H21-8, Atlanta, GA 30333, ATTN: PRA (0920-0666). CDC 57.405 (Front), v8.					



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SSI Event Detected:					
*How did the ASC facility (where the procedure was originally performed) detect/identify the SSI event? (select the method that <i>most closely resembles</i> the method of detection/identification)					
The SSI was detected through the facility's ACTIVE surveillance process:	The SSI was detected through a PASSIVE surveillance process that was not initiated by the facility:				
□ Review of patient's medical record	□ Patient/caregiver contacts facility to report				
Post-discharge surgeon survey	□ Patient returns to outpatient facility for follow-up				
Post-discharge patient letter	□ Surgeon contacts facility to report				
Post-discharge patient phone call	Report from another facility (inpatient, health department, emergency department, etc.)				
Cooperative infection prevention process between facilities					
Custom Fields					
Label	Label				
/ / / / / / / /	/ / / / / / / /				