

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: *Patient ID:	Event #:			
Secondary ID #:	Medicare #:	Social Security #:		
Patient Name, Last:	First:			
*Gender: F M Other		*Date of Birth:		
Sex at Birth: F M Unknown		Gender Identity (Specify):		
Ethnicity (Specify):		Race (Specify):		
*Date of Encounter (MM/DD/YYYY):				
Surgical Site Infection (SSI)				
*Event Type: SSI				
*Date of Event:// *Primary CPT Code: *NHSN Procedure Code:				
*SSI Level:				
☐ Superficial Incisional Primary (SIP) ☐ Deep Incisional Primary (DIP) ☐ Organ/Space				
\square Superficial Incisional Secondary (SIS) \square Deep Incisional Secondary (DIS)				
*Specify SSI Criteria Used (check all that appl Signs & Symptoms	y):	Laboratory		
☐ Abscess ☐ Localized sv	velling	☐ Organism(s) identif	ïed	
\square Erythema or redness \square Pain or tende	erness	☐ Culture or non-cultuperformed	ire based testing not	
\square Fever (>38°C) \square Purulent dra	inage	☐ Imaging test evidence of infection		
☐ Heat ☐ Wound spon	taneously			
☐ Incision deliberately opened/drained				
☐ Other evidence of infection found on invasive procedure, gross anatomic exam, or histopathologic exam		Clinical Diagnosis		
		☐ Diagnosis of superficial SSI by surgeon or physician		
*Pathogens Identified: ☐ Yes ☐ No				
If Yes, indicate up to 3 pathogens:				
			Continue>>>	
Assurance of Confidentiality: The information obtained in this surveillance sy be held in strict confidence, will be used only for the purposes stated, and wi accordance with Sections 304, 306 and 308(d) of the Public Health Service /	ill not otherwise be disclosed	or released without the consent of the indiv		
Public reporting burden of this collection of information is estimated to averal gathering and maintaining the data needed, and completing and reviewing the respond to a collection of information unless it displays a currently valid OME information, including suggestions for reducing this burden to CDC, Reports	he collection of information. A B control number. Send comr	An agency may not conduct or sponsor, and ments regarding this burden estimate or an	d a person is not required to ny other aspect of this collection of	

CDC 57.405 (Front), v8.8



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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:		
\square Review of patient's medical record	☐ Patient/caregiver contacts facility to report		
\square Post-discharge surgeon survey	\square Patient returns to outpatient facility for follow-up		
\square Post-discharge patient letter	☐ Surgeon contacts facility to report		
\square 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		
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11			
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