

## Laboratory-identified MDRO or CDI Event

Instructions for this form are available at: <u>http://www.cdc.gov/nhsn/forms/instr/57_128.pdf</u>		
Page 1 of 1 Facility ID:		*required for savin Event #:
*Patient ID:		Social Security #:
Secondary ID:		Medicare #:
Patient Name, Last:	First:	Middle:
*Gender: M F	1 11 51.	*Date of Birth:
Ethnicity (Specify):		Race (Specify):
Event Details		
*Event Type: LabID		*Date Specimen Collected:
*Specific Organism Type: (Check one)		
		□ C. difficile □ CephR-Klebsiella
CRE-E. coli CRE-Entero	bacter 🗌 CRE-K	Klebsiella 🗌 MDR-Acinetobacter
	*Specimen Body Sit	e/System: *Specimen Source:
*Outpatient: Yes No	. ,	
*Date Admitted to Facility:	_ *Location:	*Date Admitted to Location:
Last physical overnight location of patient immediately prior to arriving into facility (applies to specimen(s) collected in outpatient setting or <4 days after inpatient admission) (Check one):		
□ Nursing Home/Skilled Nursing Facility □ Personal residence/Residential care		
□ Other Inpatient Healthcare Setting (i.e., acute care hospital, IRF, LTAC, etc.) □ Unknown		
*Has patient been discharged from <u>your</u> facility in the past 3 months?		
If Yes, date of last discharge from your facility:		
Has patient been discharged from <u>another</u> facility in the past 4 weeks?  Yes No		
If Yes, from where (Check one):		
□ Nursing Home/Skilled Nursing Facility □ Other Inpatient Healthcare Setting (i.e., acute care hospital,		
Unknown IRF, LTAC, etc.)		
Custom Fields		
Label		Label
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Comments		
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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 15 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 D-74, Atlanta, GA 30333, ATTN: PRA (0920-0666).		
CDC 57.128 Rev6, v8.3		