CLABSI Process Improvement Discovery Tool



Review a minimum of <u>5</u> and a maximum of <u>10</u> medical records.

Focus:

For this review, focus on the review of randomly selected charts of inpatients (e.g., last 5) who had a diagnosis of a CLABSI made while an inpatient. Do not include patients who were admitted with a diagnosis of a CLABSI.

Instructions:

When reviewing the medical record, if documentation is found for the process, mark "Yes" in the box. If documentation is not found for the process, mark "No". If the process being reviewed is not applicable to the medical record, mark "N/A". After completing the review of all records, note the rows with the highest number of "No" responses. This will identify priority focus areas for improvement.

Note: Do not spend more than 20-30 minutes per medical record.

SUBMIT YOUR DISCOVERY TOOL FINDINGS WHEN COMPLETE: Take a 2 minute survey to report your findings. By submitting your findings, you will have taken the time to identify process gaps in which to focus improvement and to guide educational activities.

CLICK HERE TO SUBMIT YOUR FINDINGS

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Medical Records #							
(INSERTION) The patient had:							
	An order to insert a central line						
	A hospital-defined evidence-based clinical indication for a central line						
	Documentation that the central line was inserted using maximum sterile technique (i.e., mask, cap, sterile gown by all involved in procedure; full-body sterile drape during insertion)						
	Catheter placed in site other than the femoral vein						
	A non-sutured securement device utilized						
(MAII	NTENANCE) The patient had documentation of:						
	Assessment and documentation for the clinical necessity for continued use of central line per hospital policy (e.g., daily or every shift)						
	Assessment of insertion site is assessed (e.g., dressing intact, no signs of inflammation) per hospital policy (e.g. daily or every shift)						
	Patency of all central line lumens						
	If daily chlorhexidine gluconate (CHG) bathing is an established practice in the unit where patient was housed (note N/A if not an established practice in your hospital)						
	There is evidence that patient and/or family was educated about risks associated with a central line that is no longer clinically indicated						
(BLO	OD CULTURE COLLECTION) The patient had:						
	Blood drawn from two different peripheral sites, not the central line (lab result should indicate the source of the blood sample)						
Other (specify):							

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