DEPARTMENT OF HEALTH & HUMAN SERVICES Centers for Medicare & Medicaid Services 7500 Security Boulevard, Mail Stop C2-21-16 Baltimore, Maryland 21244-1850



Office of Clinical Standards & Quality/Survey & Certification Group

REF: S&C: 12-32-Hospital

DATE: May 18, 2012

TO: State Survey Agency Directors

FROM: Director

Survey & Certification Group

SUBJECT: Patient Safety Initiative Pilot Phase – Revised Draft Surveyor Worksheets

Memorandum Summary

- Patient Safety Initiative: The Centers for Medicare & Medicaid Services (CMS) is testing
 three revised surveyor worksheets for assessing compliance with three hospital Conditions
 of Participation (CoPs): Quality Assessment and Performance Improvement (QAPI),
 Infection Control, and Discharge Planning. We are focusing on compliance with these
 CoPs as a means to reduce hospital-acquired conditions (HACs), including healthcare
 associated infections (HAIs), and preventable readmissions.
- *Draft Worksheets Made Public:* Via this memorandum we are making these revised draft worksheets publicly available. We emphasize there may be additional revisions based on information gathered during the pilot test phase, which will end sometime in FY 2013.

Patient Safety Initiative Pilot Phase

The Survey & Certification Group (SCG) Patient Safety Initiative has begun pilot testing three revised surveyor worksheets designed to help surveyors assess compliance with the hospital CoPs for QAPI, infection control, and discharge planning. In S&C-12-01, released October 14, 2011, we made available to the public copies of the original surveyor worksheets, which were used during a pre-test phase of the SCG initiative that began in September 2011. The pre-test included testing one or more of the worksheets in eleven volunteer State Survey Agencies (SAs). Based on feedback obtained from SA surveyors and, CMS Central Office (CO) and Regional Office (RO) observers, the worksheets have been revised. The underlying CoPs for QAPI, infection control, and discharge planning have not changed. These regulations are the basis for any deficiencies that may be cited, not the worksheet per se. The worksheets are simply designed to assist surveyors (and hospital staff) to better identify when and where there are issues in compliance with the CoPs.

During the pilot phase of the initiative, surveyors in all State Survey Agencies (SA) initially will be testing each of the three surveyor worksheets in separate surveys. Similar to the pre-test

phase, hospitals will be selected for survey based on risk-adjusted all-cause readmissions data and/or other factors. Hospitals with higher readmission rates, as compared to other hospitals in their State, may be at greater risk for noncompliance with the three CoPs.

As in the pre-test phase, unless an immediate jeopardy is found during a pilot phase survey, we expect that all other deficiency findings will be cited as standard-level (i.e., less serious) deficiencies. Because these surveys are being conducted as part of a test of the worksheets, surveys with only standard-level citations will not require submission of a plan of correction to the SA (although hospitals are free to do so). SAs will, however, be preparing and issuing the Form CMS 2567, Statement of Deficiencies and Plan of Correction. Not only will this facilitate our evaluation of how the observations recorded on the worksheets are translated into survey findings, but we believe the survey findings will also prove educational for the hospitals surveyed.

The pilot phase involving tests by all states of each tool in a separate survey will be completed by the end of September. We may, based on ongoing feedback we receive, make further revisions to the worksheets at the beginning of FY 2013. The pilot phase will continue through the first half of FY 2013, with SAs testing use of the three worksheets in combination on at least one survey. Thereafter it is our expectation that the Patient Safety Initiative will continue, using the same process for selection of hospitals for survey as in the pilot, but employing normal enforcement practices. .

Public Distribution

The three hospital surveyor worksheets are being publicly distributed via this memo. We encourage hospitals to utilize the worksheets for self assessment of their practices related to QAPI, infection control and discharge planning. Feedback from hospitals after utilizing the worksheets is welcome. We also welcome feedback from the hospital industry at large, from patients and consumer groups, and others committed to quality and patient safety.

Please forward all questions, concerns, or comments to PFP.SCG@cms.hhs.gov.

/s/ Thomas E. Hamilton

Attachments: (3)

Patient Safety Initiative Pilot Hospital QAPI Worksheet 2012 Patient Safety Initiative Pilot Hospital Infection Control Worksheet 2012 Patient Safety Initiative Pilot Hospital Discharge Planning Worksheet 2012

cc: Survey & Certification Regional Office Management

PRE-DECISIONAL SURVEYOR WORKSHEET

Assessing Hospital Compliance With the

Condition of Participation for

Quality Assessment & Performance Improvement (QAPI)

Pilot Program Draft Version

Instructions: The following is a list of items, broken down into separate Parts, which must be assessed during the on-site survey in order to determine compliance with the QAPI Condition of Participation. Items are to be assessed primarily by review of the hospital's QAPI program documentation and interviews

State Agency Name

can be assessed in any order. Within each Part there may also	be flexibility to change the o	rder in which the variou	s items are assessed.	•
Citation instructions are provided throughout this inst deficient practices are observed.	rument, indicating the appli	cable regulatory provis	ion to be cited on the Forn	n CMS-2567 when
P/	ART 1 – HOSPITAL CHARACT	ERISTICS		
1.1 Hospital Name (please print)			_	
1.2 Address, State and Zip Code (please print)		Addross		
	ART 1 – HOSPITAL CHARACTERISTICS Address City State Zip			
	City	State	Zip	
1.3 CMS Certification Number (CCN)				

1.4 Date of site visit:
m m d d y y y y m m d d y y y y y
1.5 Number of State Agency surveyors who participated in this survey: 1.6 Approximate time spent on site performing this survey (hours):
O YES 1.7 Does the hospital participate in Medicare via accredited "deemed" status? O NO
O American Osteopathic Association (AOA)/HFAP 1.8a If YES, which AO(s)? (Check all that apply) O DNV Healthcare (DNV) O The Joint Commission (TJC)
1.8b If YES, according to the hospital, what was the end date of the most recent accreditation survey? m m d d y y y y y
1.8c What was the end date of the most recent previous standard (i.e., "full") Federal survey conducted by the State Agency?

NOTE: PART 2 - NEW HOSPITAL WORKSHEET SECTION - PURPOSELY OMITTED FROM PILOT

PART 3: DATA COLLECTION AND ANALYSIS - QUALITY INDICATOR TRACERS

Instructions for Part #3 Questions:

Select 3 quality indicators (not patient safety analyses) and trace them answering the following multipart question. Focus on indicators with related QAPI activities or projects. At least one of the indicators must have been in place long enough for most questions to be applicable.

Elements to be Assessed	Indicator #1	Indicator #2	Indicator #3
3.1 Write in indicator selected.			
Indicator selection identified	0 1	0 1	O 1
through:	0 3	0 3	0 3
	O 5	O 5	O 5
3.1.a Can the hospital provide	O YES	O YES	O YES
evidence that each quality indicator	O NO	O NO	O NO
selected is related to improved			
health outcomes? (e.g. based on	0 1	0 1	0 1
QIO, guidelines from a nationally	0 2	0 2	0 2
recognized organization, hospital	0 3	0 3	0 3
specific evidence, peer-reviewed research, etc.)	O 4 O 5	O 4 O 5	O 4 O 5
research, etc.,			

PART 3: DATA COLLECTION AND ANALYSIS - QUALITY INDICATOR TRACERS (CONTINUED)

Elements to be Assessed	Indicator #1	Indicator #2	Indicator #3
3.1.b Is the scope of data collection appropriate to the indicator, e.g., an indicator related to labor and	O YES	O YES	O YES
	O NO	O NO	O NO
delivery might be appropriate to all areas of that unit and the ED, but indicators related to hand hygiene would require data from multiple parts of the hospital.	O 1 O 2 O 3 O 4 O 5	O 1 O 2 O 3 O 4 O 5	O 1 O 2 O 3 O 4 O 5
3.1.c Is the method (e.g., chart reviews, monthly observations, etc.) and frequency of data collection	O YES	O YES	O YES
	O NO	O NO	O NO
specified?	O 1	O 1	O 1
	O 2	O 2	O 2
	O 3	O 3	O 3
	O 4	O 4	O 4
	O 5	O 5	O 5
3.1.d Is there evidence that the data are actually collected in the manner	O YES	O YES	O YES
	O NO	O NO	O NO
and frequency specified for this indicator? E.g., Is there evidence of late, incomplete, or wrong data collection?	O 1 O 2 O 3 O 4 O 5	O 1 O 2 O 3 O 4 O 5	O 1 O 2 O 3 O 4 O 5

PART 3: DATA COLLECTION AND ANALYSIS - QUALITY INDICATOR TRACERS (CONTINUED)

Elements to be Assessed	Indicator #1	Indicator #2	Indicator #3
3.1.e If unit staff play a role in data	O YES	O YES	O YES
collection, is collection consistent	O NO	O NO	O NO
with the specifications for how the	O N/A	O N/A	O N/A
data are to be collected?	0 1	0 1	0 1
	O 2	0 2	0 2
	O 3	O 3	O 3
	O 4 O 5	O 4 O 5	O 4 O 5
	0 5	0 5	0 5
3.1.f Are data that have been	O YES	O YES	O YES
collected aggregated in accordance	O NO	O NO	O NO
with the hospital methodology			
specified for this indicator?	O 1	0 1	0 1
	O 2	O 2	O 2
	O 3	O 3	0 3
	0 4	0 4	0 4
	O 5	O 5	O 5
3.1.g Are the collected data	O YES	O YES	O YES
analyzed?	O NO	O NO	O NO
	0 1	0 1	0 1
	O 2	O 2	O 2
	O 3 O 4	O 3 O 4	O 3 O 4
	O 5	O 5	O 5
			<u> </u>

PART 3: DATA COLLECTION AND ANALYSIS - QUALITY INDICATOR TRACERS (CONTINUED)

Elements to be Assessed	Indicator #1	Indicator #2	Indicator #3
3.1.h If the indicator is the type that	O YES	O YES	O YES
measures a rate, are rates calculated	O NO	O NO	O NO
for points in time and over time, and	O N/A	O N/A	O N/A
are comparisons made to	O 1	O 1	0 1
performance benchmarks when	O 2	O 2	O 2
available (e.g. established by	O 3	O 3	0 3
nationally recognized	O 4	O 4	O 4
organizations)?	O 5	O 5	O 5
24:34	O MEG	0. 7.6	O 1/50
3.1.i When feasible, are aggregated	O YES	O YES	O YES
data broken down into subsets that	O NO O N/A	O NO O N/A	O NO O N/A
allow comparison of performance among hospital units covered by the	O 1	O 1	O 1
indicator? For example, a hand	0 2	0 2	O 2
hygiene indicator should allow	0 3	O 3	O 3
comparison among different	0 4	0 4	0 4
inpatient units.	0 5	0 5	0 5
If no to any of 3.1.a through 3.1.i, cite	at 42 CFR 482.21(a)(1), (a)(2), (b)(1), &	(b)(3) (Tag A-273)	

PART 3: DATA COLLECTION AND ANALYSIS - QUALITY INDICATOR TRACERS (CONTINUED)

Elements to be Assessed	Indicator #1	Indicator #2	Indicator #3
 3.1.j If the data analysis identified areas needing improvement, is there evidence that the hospital instituted interventions (activities and/or projects) to address them? Check N/A if analysis did not lead to interventions, but the hospital could demonstrate that other areas were of higher priority. Check NO if analysis did not lead to interventions and the hospital could not demonstrate that other improvement activities were of higher priority. 	O YES O NO O N/A O 1 O 2 O 3 O 4 O 5	O YES O NO O N/A O 1 O 2 O 3 O 4 O 5	O YES O NO O N/A O 1 O 2 O 3 O 4 O 5
3.1.k Are interventions evaluated for success?	O YES O NO O N/A O 1 O 2 O 3 O 4 O 5	O YES O NO O N/A O 1 O 2 O 3 O 4 O 5	O YES O NO O N/A O 1 O 2 O 3 O 4 O 5

PART 3: DATA COLLECTION AND ANALYSIS - QUALITY INDICATOR TRACERS (CONTINUED)

Elements to be Assessed	Indicator #1	Indicator #2	Indicator #3
3.1.I If interventions taken were not successful, were new interventions developed?	O YES O NO O N/A O 1 O 2 O 3 O 4 O 5	O YES O NO O N/A O 1 O 2 O 3 O 4 O 5	O YES O NO O N/A O 1 O 2 O 3 O 4 O 5
3.1.m If interventions were successful, did evaluation continue longer to assess if success was sustained?	O YES O NO O N/A O 1 O 2 O 3 O 4 O 5	O YES O NO O N/A O 1 O 2 O 3 O 4 O 5	O YES O NO O N/A O 1 O 2 O 3 O 4 O 5

Manner of Assessment Code: 1-Interview 2-Observation 3- QAPI Documentation 4- Medical Record Review 5- Other

PART 4 - APPLYIN	NG QU	IALITY IND	ICATOR I	NFORMATION - ACTIVITIES AND PROJECTS
Elements to be Assessed				Manner of Assessment Code (Enter all that apply) & Surveyor Notes
4.1 Can the hospital provide evidence that its improvement activities focus on areas that are high risk (severity), high volume (incidence or prevalence), or problem-prone?		YES NO	O 1 O 2 O 3 O 4 O 5	
If no to 4.1, cite at 42 CFR 482.21(c)(1)(i) & (ii) (Tag A				
4.2 Can the hospital provide evidence that it conducts distinct performance improvement projects?		YES NO	O 1 O 2 O 3 O 4 O 5	
4.3 Is the number of projects proportional to the scope and complexity of the hospital's services and operations? No fixed ratio is required, but smaller hospitals with a smaller number of distinct services would be expected to have fewer projects than a large hospital with many different services.		YES NO	O 1 O 2 O 3 O 4 O 5	

PART 4 - APPLYING QUALITY INDICATOR INFORMATION – ACTIVITIES AND PROJECTS (CONTINUED)

Elements to be Assessed				Manner of Assessment Code (Enter all that apply) & Surveyor Notes
4.4 Does the scope of projects reflect the scope and complexity of the hospital's services and operations? For example, if the hospital offers more complex services, such as neonatal intensive care, or open heart surgery, have there been QAPI project(s) related to any of those services?	00	YES NO	00000	
If no to any of 4.2 through 4.4, cite at 42 CFR 482.21(c	d)(1)	(Tag A-297)	
4.5 Can the hospital provide evidence showing why each project was selected? (NOTE: If the project is a QIO cooperative project or an IT project, such as computer ordered physician entry for medications or an electronic medical record, no rationale is required. Check N/A in these cases)	000	YES NO N/A	0000	

PART 5 – PATIENT SAFETY – ADVERSE EVENTS AND MEDICAL ERRORS				
Elements to be Assessed		Manner of Assessment Code (Enter all that apply) & Surveyor Notes		
5.1 In this multipart question evaluate if the hospital's	s leadership se	ets expectations for patient safety? Specifically:		
5.1.a Is there evidence of widespread staff training or communication to convey expectations for patient safety to all staff? (e.g. training related to steps to take in a situation that feels unsafe, how to report medical errors (including near misses/close calls) adverse events, etc.)	O YES O NO	O 1 O 2 O 3 O 4 O 5		
5.1.b Is there evidence that the hospital has adopted policies supporting a non-punitive approach to staff reporting of medical errors (including near misses/close calls), adverse events, and situations they consider unsafe?	O YES O NO	O 1 O 2 O 3 O 4 O 5		
5.1.c On each unit surveyed, can staff explain what the hospital's expectations are for their role in promoting patient safety?	O YES O NO	O 1 O 2 O 3 O 4 O 5		
f no to 5.1.a, 5.1.b, or 5.1.c, cite at 42 CFR 482.21(e)(3) (Tag A-286)				

Elements to be Assessed			Manner of Assessment Code (Enter all that apply) & Surveyor Notes
5.2. In this multipart question evaluate if the hospital events on an ongoing basis? Specifically:	has a	a systematio	process to identify medical errors (including near misses/close calls) and adverse
5.2.a On each unit/program surveyed, can staff describe what is meant by medical errors (including near misses/close calls) and adverse events?	00	YES NO	O 1 O 2 O 3 O 4 O 5
5.2.b On each unit/program surveyed, can staff explain how and/or to whom they should report medical errors (including near misses/close calls) and adverse events?	0 0	YES NO	O 1 O 2 O 3 O 4 O 5
5.2.c Does the hospital employ methods, in addition to staff incident reporting, to identify possible medical errors (including near misses/close calls) and adverse events? (Examples of other methods include, but are not limited to, retrospective medical record reviews, review of claims data, unplanned readmissions, or patient complaints/grievances, interview or survey of patients, etc.)	0 0	YES NO	O 1 O 2 O 3 O 4 O 5

Elements to be Assessed		Manner of Assessment Code (Enter all that apply) & Surveyor Notes
5.2.d Can the hospital provide evidence of medical errors (including near misses/close calls) and adverse events identified through staff reports or other methods?	O YES O NO	O 1 O 2 O 3 O 4 O 5
If no to any 5.2.a through 5.2.d, cite at 42 CFR 482.21	(a)(2) & 482.21	(c)(2) (Tag A-286)
5.3 Is there QAPI program collaboration with infection control officer(s) to identify and track avoidable healthcare-acquired infections?	O YES O NO	O 1 O 2 O 3 O 4 O 5
5.4 Is there evidence that problems identified by infection control officer(s) are addressed through QAPI program activities? If no to 5.3 or 5.4, cite at 42 CFR 482.42(b)(1) (Tag A-	O YES O NO	0 1 0 2 0 3 0 4 0 5

Elements to be Assessed		Manner of Assessment Code (Enter all that apply) & Surveyor Notes
5.5 Does the QAPI program identify and track medication administration errors, adverse drug reactions, and drug related incompatibilities?	O YES O NO	O 1 O 2 O 3 O 4 O 5
If no to 5.5, cite at 42 CFR 482.25(b)(6) (Tag A-508) a	and possibly also	o at 42 CFR 482.21(a)(2) (Tag A-286)
5.6 Is there a QAPI program process for staff to report blood transfusion reactions, and reviews of reported blood transfusion reactions to identify medical errors (including near misses/close calls) and/or adverse events?	O YES O NO	O 1 O 2 O 3 O 4 O 5
If no to 5.6, cite at 42 CFR 482.23(c)(4) (Tag A-410) a		
5.7 Did the survey team have prior knowledge of, or identify while on-site, serious preventable adverse events that the hospital failed to identify?	O YES O NO	O 1 O 2 O 3 O 4 O 5
If yes to 5.7, cite at 42 CFR 482.21(a)(2) (Tag A-286)		

Elements to be Assessed	u			IVIAIIIIEI OI AS	sessifient Code (Enter a	ii tiiat appiy) & Surveyor Notes	
5.8 Has the hospital conducted causal	analyses of all	O YES	0 1				
serious preventable adverse events it		O NO	O 2				
		O N/A	O 3				
Use as your sample all serious prevent	ahla avants	0 11,71	0 4				
, , ,			0 5				
identified by the hospital in the period			0 3				
prior to the survey date? (Note: for ev							
occurred less than 2 months prior to the	•						
date, the hospital may have started, b	ut not yet						
completed a causal analysis.)							
If no to 5.8, cite at 42 CFR 482.21(a)(2	(Tag A-286)						
		PART 5: 0	CAUSAL	ANALYSIS TRAC	ERS		
Instructions for Questions #5.9 and 5	5.10: If the answ	ver to Question	#5.9 is "	yes", select thre	ee causal analyses the h	ospital has completed for adverse	
events or near misses (close calls) du	ring the last 12	- 24 months. A	nalyses r	may be of a sing	le event/near miss or a	group of similar types of events/near	r
misses. Answer the questions in #5.10 for each analysis selected. (For at least one causal analysis selected, there should be sufficient time after					nalysis selected, there	should be sufficient time after	
implementation of preventive measures for the hospital to have evaluated the impact of those measures.) For initial certification surveys of new							
•	ures for the hosi	pital to have eva		the impact of th	ose measures.) For ini	tial certification surveys of new	
implementation of preventive measu		-	aluated t	•		•	
implementation of preventive measure hospitals, this section may not apply	, depending on	whether any se	aluated t rious pre	eventable adver	se events have occurre	•	
implementation of preventive measure hospitals, this section may not apply 5.9 Has the hospital conducted any care	, depending on usal analyses in	whether any se the 12 – 24 mo	aluated t rious pre onths pric	eventable adver		•	
implementation of preventive mease hospitals, this section may not apply 5.9 Has the hospital conducted any casurvey date? If yes continue, if no, skip	, depending on usal analyses in 5.10 and all 5.2	whether any se the 12 – 24 mo 10 sub-question	aluated t rious pre onths pric	eventable adver or to the	se events have occurre O YES O NO	d and been identified.	
implementation of preventive mease hospitals, this section may not apply 5.9 Has the hospital conducted any casurvey date? If yes continue, if no, skip Elements to be Assessed	, depending on usal analyses in 5.10 and all 5.2	whether any se the 12 – 24 mo	aluated t rious pre onths pric	eventable adver or to the	se events have occurre	•	
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implementation of preventive mease hospitals, this section may not apply 5.9 Has the hospital conducted any casurvey date? If yes continue, if no, skip Elements to be Assessed 5.10 Write in selected causal	, depending on usal analyses in 5.10 and all 5.2	whether any se the 12 – 24 mo 10 sub-question	aluated t rious pre onths pric	eventable adver or to the	se events have occurre O YES O NO	d and been identified.	
implementation of preventive mease hospitals, this section may not apply 5.9 Has the hospital conducted any casurvey date? If yes continue, if no, skip Elements to be Assessed 5.10 Write in selected causal analysis, using a code or other means to avoid capturing	, depending on usal analyses in 5.10 and all 5.2	whether any se the 12 – 24 mo 10 sub-question	aluated t rious pre onths pric	eventable adver or to the	se events have occurre O YES O NO	d and been identified.	
implementation of preventive mease hospitals, this section may not apply 5.9 Has the hospital conducted any casurvey date? If yes continue, if no, skip Elements to be Assessed 5.10 Write in selected causal analysis, using a code or other means to avoid capturing identifiable information on this	, depending on usal analyses in 5.10 and all 5.2	whether any se the 12 – 24 mo 10 sub-question	aluated t rious pre onths pric	eventable adver or to the	se events have occurre O YES O NO	d and been identified.	
implementation of preventive mease hospitals, this section may not apply 5.9 Has the hospital conducted any casurvey date? If yes continue, if no, skip Elements to be Assessed 5.10 Write in selected causal analysis, using a code or other means to avoid capturing	, depending on usal analyses in 5.10 and all 5.2	whether any se the 12 – 24 mo 10 sub-question	aluated t rious pre onths pric	eventable adver or to the	se events have occurre O YES O NO	d and been identified.	
implementation of preventive mease hospitals, this section may not apply 5.9 Has the hospital conducted any casurvey date? If yes continue, if no, skip Elements to be Assessed 5.10 Write in selected causal analysis, using a code or other means to avoid capturing identifiable information on this worksheet.	t, depending on a nusual analyses in 5.10 and all 5.2 Causa	whether any se the 12 – 24 mo 10 sub-question	aluated t rious pre onths pric	eventable adver or to the Caus	se events have occurre O YES O NO	Causal Analysis #3	
implementation of preventive mease hospitals, this section may not apply 5.9 Has the hospital conducted any casurvey date? If yes continue, if no, skip Elements to be Assessed 5.10 Write in selected causal analysis, using a code or other means to avoid capturing identifiable information on this worksheet. Causal analysis selection identified	c, depending on pusal analyses in 5.10 and all 5.2 Causa	whether any se the 12 – 24 mo 10 sub-question	aluated t rious pre onths pric	Caus O 1	se events have occurre O YES O NO	Causal Analysis #3	
implementation of preventive mease hospitals, this section may not apply 5.9 Has the hospital conducted any casurvey date? If yes continue, if no, skip Elements to be Assessed 5.10 Write in selected causal analysis, using a code or other means to avoid capturing identifiable information on this worksheet.	Causa Causal analyses in 5.10 and all 5.2 Causa	whether any se the 12 – 24 mo 10 sub-question	aluated t rious pre onths pric	Caus O 1 O 2	se events have occurre O YES O NO	Causal Analysis #3 O 1 O 2	
implementation of preventive mease hospitals, this section may not apply 5.9 Has the hospital conducted any casurvey date? If yes continue, if no, skip Elements to be Assessed 5.10 Write in selected causal analysis, using a code or other means to avoid capturing identifiable information on this worksheet. Causal analysis selection identified	Causa Causa	whether any se the 12 – 24 mo 10 sub-question	aluated t rious pre onths pric	Caus O 1 O 2 O 3	se events have occurre O YES O NO	Causal Analysis #3 O 1 O 2 O 3	
implementation of preventive mease hospitals, this section may not apply 5.9 Has the hospital conducted any casurvey date? If yes continue, if no, skip Elements to be Assessed 5.10 Write in selected causal analysis, using a code or other means to avoid capturing identifiable information on this worksheet. Causal analysis selection identified	Causa O 1 O 2 O 3 O 4	whether any se the 12 – 24 mo 10 sub-question	aluated t rious pre onths pric	Caus O 1 O 2 O 3 O 4	se events have occurre O YES O NO	Causal Analysis #3 O 1 O 2 O 3 O 4	
implementation of preventive mease hospitals, this section may not apply 5.9 Has the hospital conducted any casurvey date? If yes continue, if no, skip Elements to be Assessed 5.10 Write in selected causal analysis, using a code or other means to avoid capturing identifiable information on this worksheet. Causal analysis selection identified	Causa Causa	whether any se the 12 – 24 mo 10 sub-question	aluated t rious pre onths pric	Caus O 1 O 2 O 3	se events have occurre O YES O NO	Causal Analysis #3 O 1 O 2 O 3	
implementation of preventive mease hospitals, this section may not apply 5.9 Has the hospital conducted any casurvey date? If yes continue, if no, skip Elements to be Assessed 5.10 Write in selected causal analysis, using a code or other means to avoid capturing identifiable information on this worksheet. Causal analysis selection identified	Causa O 1 O 2 O 3 O 4	whether any se the 12 – 24 mo 10 sub-question	aluated t rious pre onths pric	Caus O 1 O 2 O 3 O 4	se events have occurre O YES O NO	Causal Analysis #3 O 1 O 2 O 3 O 4	

PART 5: CAUSAL ANALYSIS TRACERS (CONTINUED)

Elements to be Assessed	Causal Analysis #1	Causal Analysis #2	Causal Analysis #3
5.10.a Has the hospital identified potential underlying causes?	O YES	O YES	O YES
	O NO	O NO	O NO
	O 1 O 2 O 3 O 4 O 5	O 1 O 2 O 3 O 4 O 5	O 1 O 2 O 3 O 4 O 5
5.10.b Has the hospital identified all parts of the hospital utilizing similar processes/at similar risk?	O YES	O YES	O YES
	O NO	O NO	O NO
	O 1 O 2 O 3 O 4 O 5	O 1 O 2 O 3 O 4 O 5	O 1 O 2 O 3 O 4 O 5
5.10.c Has the hospital developed and implemented preventive actions based on the analysis in at least one	O YES	O YES	O YES
	O NO	O NO	O NO
area of the hospital?	O 1	O 1	O 1
	O 2	O 2	O 2
	O 3	O 3	O 3
	O 4	O 4	O 4
	O 5	O 5	O 5

PART 5: CAUSAL ANALYSIS TRACERS (CONTINUED)

Elements to be Assessed	Causal Analysis #1	Causal Analysis #2	Causal Analysis #3
5.10.d Has the hospital evaluated the impact of the preventive actions, including tracking	O YES O NO	O YES O NO	O YES O NO
reoccurrences of similar events/near misses?	O 1 O 2 O 3 O 4 O 5	O 1 O 2 O 3 O 4 O 5	O 1 O 2 O 3 O 4 O 5
5.10.e If evaluation showed the intervention(s) did not meet goals, did the hospital implement a revised	O YES O NO O N/A	O YES O NO O N/A	O YES O NO O N/A
intervention and evaluate it?	O 1 O 2 O 3 O 4 O 5	O 1 O 2 O 3 O 4 O 5	O 1 O 2 O 3 O 4 O 5
5.10.f Has the hospital implemented preventive actions found to be effective in all parts of	O YES O NO	O YES O NO	O YES O NO
the hospital utilizing similar processes/at similar risk, unless there are documented reasons for not doing so?	O 1 O 2 O 3 O 4 O 5	O 1 O 2 O 3 O 4 O 5	O 1 O 2 O 3 O 4 O 5

Manner of Assessment Code: 1-Interview 2-Observation 3- QAPI Documentation 4- Medical Record Review 5- Other

PART 6 – BROAD QAPI REQUIREMENTS AND LEADERSHIP RESPONSIBILITIES					
Elements to be Assessed			Manner of Assessment Code (Enter all that apply) & Surveyor Notes		
6.1 Is there evidence that the hospital has a formal QAPI program - including written policies and procedures, budgeted resources, and clearly identified responsible staff - approved by the governing body after input from the CEO and medical staff leadership?	00	YES NO	O 1 O 2 O 3 O 4 O 5		
If no to 6.1, for pilot only, cite at 42 CFR 482.21(e)(1)	<mark>& (2</mark>)	(Tag A-309	9)		
6.1.a Has the hospital maintained and made available for surveyor review evidence of its QAPI program?	0 0	YES NO	O 1 O 2 O 3 O 4 O 5		
If no to 6.1.a, for pilot only cite at 42 CFR 482.21 (Sta					
6.2 In this multipart question evaluate if the hospital's 6.2.a Using information on services offered from the Hospital/CAH Data Base Worksheet, can the QAPI manager provide evidence of QAPI assessment related to each service?	0 0	YES NO	O 1 O 2 O 3 O 4 O 5		
If no to 6.2.a, for pilot only cite at 42 CFR 482.21 (Sta	ndar	d level tag)	(Tag A-308)		

PART 6 – BROAD QAPI REQUIREMENTS AND LEADERSHIP RESPONSIBILITIES (CONTINUED)

Elements to be Assessed		Manner of Assessment Code (Enter all that apply) & Surveyor Notes
6.2.b Using information from the hospital identifying services provided under arrangement (contract), can the QAPI manager provide evidence of QAPI assessment for each service related to clinical care provided under contract or arrangement? (Exclusively administrative contractual services, e.g., payroll preparation, are not required to be included in the QAPI program.)	O YES O NO O N/A	O 1 O 2 O 3 O 4 O 5
If no to 6.2.b, cite at 42 CFR 482.12(e) and 482.21 (fo	<mark>r pilot - Standar</mark>	d level tag) (Tags A-083 and A-308)
6.3 Is there evidence that the governing body, hospital CEO, Medical Staff leadership, and other senior administrative officials, e.g., Director of Nursing, each play a role in QAPI program planning and implementation?	O YES O NO	O 1 O 2 O 3 O 4 O 5
If no to 6.3, cite at 42 CFR 482.21(e)(2) (Tag A-309)	1	
6.4 Is there evidence, e.g. in minutes, that the hospital	al's governing bo	ody:
6.4.a Approves QAPI program indicators selected and frequency of data collection? If no to 6.4.a, cite at 42 CFR 482.21(b)(3) (Tag A-273)	O YES O NO	O 1 O 2 O 3 O 4 O 5

PART 6 – BROAD QAPI REQUIREMENTS AND LEADERSHIP RESPONSIBILITIES (CONTINUED)

Elements to be Assessed			Mai	nner of Assessment Code (Enter all that apply) & Surveyor Notes
6.4.b Ensures the QAPI program annually determines the number of distinct QAPI projects to be conducted in the coming year?	0 0	YES NO	O 1 O 2 O 3 O 4 O 5	
6.4.c Actively reviews the results of QAPI data collection, analyses, activities, projects and makes decisions based on such review?	0 0	YES NO	O 1 O 2 O 3 O 4 O 5	
If no to either, 6.4.b or 6.4.c, cite at 42 CFR 482.21(e)	(2) &	(e)(5) (Tag	\- 309)	
6.4.d Holds the CEO accountable for the effectiveness of the QAPI program?	0 0	YES NO	O 1 O 2 O 3 O 4 O 5	
If no to 6.4.d, cite at 42 CFR 482.21(e)(2) and 482.12(b) (T	ags A-309 8	A-057)	

PART 6 – BROAD QAPI REQUIREMENTS AND LEADERSHIP RESPONSIBILITIES (CONTINUED)

Elements to be Assessed		Manner of Assessment Code (Enter all that apply) & Surveyor Notes
6.5 Regarding resource allocation:		
6.5.a Is there evidence of the amount of resources (funding and personnel) dedicated to the hospital's QAPI program and the functions for which those resources are used?	O YES O NO	O 1 O 2 O 3 O 4 O 5
If no to 6.5.a, cite at 42 CFR 482.21(e)(4) (Tag A-315)		
6.5.b If there are condition-level QAPI program deficiencies, is there evidence that lack of QAPI resources are a significant contributing cause of these deficiencies? If yes to 6.5.b, cite at 42 CFR 482.21(e)(4) (Tag A-315)	O YES O NO O N/A	O 1 O 2 O 3 O 4 O 5

PRE-DECISIONAL SURVEYOR WORKSHEET

Assessing Hospital Compliance with the

Condition of Participation for Infection Control

Pilot Program Draft Version

Name of State Agency:		

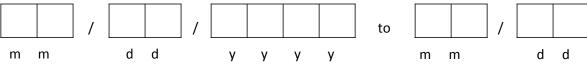
Instructions: The following is a list of items that must be assessed during the on-site survey, in order to determine compliance with the Infection Control Condition of Participation. Items are to be assessed by a combination of observation, interviews with hospital staff, patients and their family/support persons, review of medical records, and a review of any necessary infection control program documentation. During the survey, observations or concerns may prompt the surveyor to request and review specific facility policies and procedures. Surveyors are expected to use their judgment and review only those documents necessary to investigate their concern(s) or to validate their observations.

The interviews should be performed with the most appropriate staff person(s) for the items of interest, as well as with patients, family members, and support persons.

Citation instructions are provided throughout this instrument, indicating the applicable regulatory provision to be cited on Form CMS-2567 when deficient practices are observed.

1. Hospital name: 2. Address, State, Zip Code: City State Zipcode 3. CMS Certification Number (CCN):

Δ	Date	Λf	site	visit



5. Number of State Agency surveyors who participated in this survey:



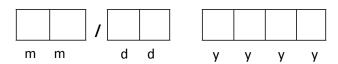
6. Approximate time spent on site performing this survey (hours):



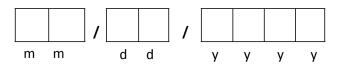
7. Does the hospital participate in Medicare via accredited "deemed" status?



- a. If YES, which Accrediting Organization(s)?
 - i. American Osteopathic Association (AOA)/Healthcare Facilities Accreditation Program (HFAP)
 - ii. Det Norske Veritas Healthcare (DNV)
 - iii. The Joint Commission (TJC)
- b. If YES, according to the hospital, what was the end date of the most recent accreditation survey:



8. What was the end date of the most recent previous State Agency Federal survey:



Module 1: Infection Control/Prevention Program

Section 1. A. Infection control/prevention program and resources Manner of Assessment Code (check all that apply) Elements to be assessed & Surveyor Notes Yes □ 1 1. A.1 The hospital has designated one or more individual(s) as its 2 Infection Control Officer(s). ☐ No \bigcirc 3 4 O N/A \bigcirc 5 If no, cite at 42 CFR 482.42(a) (Tag A-0748) Yes \bigcirc 1 1. A.2 The hospital has evidence that demonstrates the Infection 2 Control Officer(s) is qualified and maintain(s) qualifications □ No \Box 3 through education, training, experience or certification related 4 to infection control consistent with hospital policy. O N/A □ 5 If no, cite at 42 CFR 482.42(a) (Tag A-0748) 1. A.3 The Infection Control Officer(s) can provide evidence that the Yes \bigcirc 1 2 hospital has developed general infection control policies and □ 3 O No procedures that are based on nationally recognized guidelines □ 4 and applicable state and federal law. □ N/A □ 5 If no, cite at 482.42(a) (Tag A-0748) Yes \bigcirc 1 1. A.4 The hospital has infection control policies and procedures relevant to construction, renovation, maintenance, demolition, \square 2 O No 3 and repair. An infection control risk assessment (ICRA) to define 4 the scope of the project and need for barrier measures is □ N/A \bigcirc 5 performed before a project gets underway. If no, cite at 42 CFR 482.42(a) (Tag A-0748)

1. <i>F</i>	A.5 The AIIR meets generally accepted specifications:	Yes	(a)	1	
•	at least 6 (existing facility) or 12 (new construction/renovation)		2	2	
	air changes per hour or per state licensure rules and;	○ No	□ 3	3	
•	direct exhaust of air to outside, if not possible air returned to air		4	4	
	handling system or adjacent spaces if directed through HEPA	□ N/A	□ 5	5	
	filters and;				
•	when AIIR is in use for a patient on Airborne Precautions,				
	documentation that monitoring of air pressure is done daily with				
	visual indicators (smoke tubes, flutter strips), regardless of				
	differential pressure sensing devices (i.e. manometers): and				
•	AIIR door kept closed when not required for entry and exit				
If n	o, cite at 42 CFR 482.42(a)(1) (Tag A-0749)				

Section 1. B. Hospital QAPI systems related to Infection Prevention and Control Manner of Assessment Code (check all that apply) Elements to be assessed & Surveyor Notes The hospital infection prevention program is coordinated into the hospital QAPI program as evidenced by: 1. B.1 The Infection Control Officer(s) can provide evidence that Yes 2 problems identified in the infection control program are 3 addressed in the hospital QAPI program (i.e., development and O No implementation of corrective interventions, and ongoing □ N/A evaluation of interventions implemented for both success and sustainability). If no, cite at 42 CFR 482.42(b)(1) (Tag A-0756) 1. B.2 Is there evidence that the hospital has adopted policies Yes (h) 1 ○ 2 supporting a non-punitive approach to staff reporting of O No hospital acquired infections, adverse events, and situations they consider unsafe? □ N/A If no, cite at 42 CFR 482.21(e)(3) (Tag A-0286)

1. B.3 Hospital leadership, including the CEO, Medical Staff, and the Director of Nursing Services ensures the hospital implements successful corrective action plans in affected problem area(s).	C Yes	① 1 ② 2 ② 3 ② 4
	[©] N/A	5
If no, cite at 42 CFR 482.42(b)(2) (Tag A-0756)		
B.4 The hospital utilizes a risk assessment process to prioritize selection of quality indicators for infection prevention and	C Yes	
control.	◯ No	
	□ N/A	O 5
No citation		
		·

Section 1. C. Systems to prevent transmission of MDROs and promote antibiotic stewardship, Surveillance

Elements to be assessed		Manner of Assessment Code (check all that apply) & Surveyor Notes
C.1 The hospital has policies and procedures to minimize the risk of transmission of multidrug-resistant organisms (MDROs)	Yes	
within the hospital (between or amongst patients and health care personnel).	No No	○ 3 ○ 4
	□N/A	
No citation		
C.2 The primary interview participants can provide evidence that the hospital identifies patients with MDROs and has implemented policies and procedures aimed at preventing the development and transmission of MDROs.	☐ Yes ☐ No ☐ N/A	① 1 ○ 2 ○ 3 ○ 4 ○ 5
No citation		
C.3.a Facility has a multidisciplinary process in place to review antimicrobial utilization, local susceptibility patterns, and	C Yes	
antimicrobial agents in the formulary <i>and</i> there is evidence that the process is followed.	O No	□ 3□ 4
	□ N/A	□ 5

 C.3.b Systems are in place to prompt clinicians to use appropriate antimicrobial agents (e.g., computerized physician order entry, comments in microbiology susceptibility reports, notifications from clinical pharmacist, formulary restrictions, evidenced based guidelines and recommendations). 	☐ Yes ☐ No ☐ N/A	1 2 3 4 D 5	
1. C.3.c Antibiotic orders include an indication for use.	☐ Yes ☐ No ☐ N/A	① 1 ② 2 ② 3 ② 4 ② 5	
C.3.d There is a mechanism in place to prompt clinicians to review antibiotic courses of therapy after 72 hours of treatment.	☐ Yes ☐ No ☐ N/A	1 2 3 4 0 5	
C.3.e The facility has a system in place to identify patients currently receiving intravenous antibiotics who might be eligible to receive oral antibiotic treatment.	☐ Yes ☐ No ☐ N/A	1 2 3 3 4 5 5	
No citation for 1.C.3.a through 1.C.3.e	1		
C.4 The hospital has established systems with a clinical microbiology laboratory that ensures prompt notification of IP staff or medical director/designee when a novel resistance pattern is detected.	☐ Yes ☐ No ☐ N/A	1 2 3 C 4 C 5	
No citation			
 C.5 Patients and healthcare personnel identified by laboratory culture as colonized or infected with MDROs are identified and isolated according to facility policies. (Note: The hospital is not required to perform routine surveillance of patients or healthcare personnel). 	☐ Yes ☐ No ☐ N/A	1 2 3 4 0 5	
If no, cite at 42 CFR 482.42(a)(1) (Tag A-0749)		<u>I</u>	

 C.6 The hospital has a system for identifying those present on admission infections in order to control (prevent spread of) those infections and communicable diseases in the hospital. (This does not require the hospital to perform cultures on all patients admitted to the hospital.) 	☐ Yes ☐ No ☐ N/A	① 1 ② 2 ② 3 ② 4 ② 5	
If no, cite at 42 CFR 482.42(a)(1) (Tag A-0749)			
C.7 The Infection Control Officer can provide evidence that an updated list of diseases reportable to the local or state public health authority is available.	☐ Yes ☐ No ☐ N/A	1 2 3 4 5	
No citation			
C.8 The Infection Control Officer can provide evidence that reportable diseases are documented and submitted as required by the local health authority.	☐ Yes ☐ No ☐ N/A	1 2 3 G 4 G 5	
If no, cite at 42 CFR 482.42(a)(1) (Tag A-0749)			
•			

Section 1. D Personnel Education System / Infection Control Training							
Elements to be assessed		Manner of Assessment Code (check all that apply) & Surveyor Notes					
D.1 Healthcare personnel receive job-specific training on hospital infection control practices, policies, and procedures upon hire and at regular intervals	☐ Yes ☐ No ☐ N/A	 □ 1 □ 2 □ 3 □ 4 □ 5 					
D.2 The hospital infection control system trains healthcare personnel that are in contact with bloodborne pathogens on the bloodborne pathogen standards upon hire and when problems are identified.	☐ Yes ☐ No ☐ N/A	 □ 1 □ 2 □ 3 □ 4 □ 5 					

 D.3 The hospital infection control system addresses needle sticks, sharps injuries, and other employee exposure events. 	☐ Yes ☐ No ☐ N/A	1 2 3 4 D 5	
D.4 Following an exposure event, post-exposure evaluation and follow-up, including prophylaxis as appropriate, is available.	☐ Yes ☐ No ☐ N/A	1 2 3 4 C 5	
D.5 The hospital infection control system ensures healthcare personnel with TB test conversions are provided with appropriate follow-up.	☐ Yes ☐ No ☐ N/A	1 2 3 4 0 5	
If no to any of the above (1.D.1-1.D.5), cite at 42 CFR 482.42(a)(1) (Tag	g A-0749)		
D.6 The hospital infection control system ensures the facility has a respiratory protection program that details required worksite-specific procedures and elements for required respirator use.	☐ Yes ☐ No ☐ N/A	1 2 2 3 4 C 5	
D.7 The hospital infection control system ensures that respiratory fit testing is provided at least annually to appropriate healthcare personnel.	☐ Yes ☐ No ☐ N/A	1 2 3 4 C 5	
D.8 Hospital has well-defined policies concerning contact of personnel with patients when personnel have potentially transmissible conditions. These policies should include: * work-exclusion policies that encourage reporting of illnesses and do not penalize with loss of wages, benefits, or job status * education of personnel on prompt reporting of illness to supervisor and occupational health	☐ Yes ☐ No ☐ N/A	1 2 3 4 5	

 D.9 Aggregated rates of TB-test conversion are periodically reviewed by the Infection Control Officer to determine the need for corrective action plans. 	☐ Yes ☐ No ☐ N/A	1 2 3 A 4 D 5	
No citations for 1.D.6 – D.9			
D.10 Healthcare personnel competency and compliance with job- specific infection prevention policies and procedures are ensured through routine training and when problems are identified by the Infection Control Officer.	☐ Yes ☐ No ☐ N/A	1 2 2 3 3 4 C 5	
D.11 If the hospital has had healthcare personnel infection exposure events, the hospital evaluates event data and develops/ implements corrective action plans to reduce the incidence of such events.	☐ Yes ☐ No ☐ N/A	① 1 ② 2 ② 3 ② 4 ② 5	
If no to 1.D.10 or-1.D.11), cite at 42 CFR 482.42(b)(1) (Tag A-0756)	<u> </u>		
D.12 The hospital infection control system provides Hepatitis B vaccine and vaccination series to all employees who have occupational exposure and conducts post-vaccination screening after the third vaccine dose is administered.	☐ Yes ☐ No ☐ N/A	1 2 3 4 D 5	
D.13 The hospital infection control system ensures that all healthcare personnel (paid and unpaid) who have potential for exposure to TB are screened for TB upon hire and, if negative, based upon facility risk classification thereafter.	☐ Yes ☐ No ☐ N/A	1 2 3 4 C 5	
D.14 The hospital infection control system ensures that all healthcare personnel are offered annual influenza vaccination.	☐ Yes ☐ No ☐ N/A	1 2 2 3 4 C 5	
No citations for 1.D.12 - 14			

Module 2: General Infection Control Elements - to be applied to all locations (e.g., general wards, critical care units, labor and delivery, emergency department, endoscopy suites, radiology)

Section 2. A Hand Hygiene Manner of Assessment Code Manner of Assessment Code Elements to be assessed (check all that apply) & Surveyor Notes (check all that apply) & Surveyor Notes Hand hygiene is performed in a manner consistent with hospital infection control practices, policies, and procedures to maximize the prevention of infection and communicable disease including the following: Yes \square 1 Yes \bigcap 1 2. A.1 Soap, water, and a sink are readily 2 □ 2 accessible in patient care areas including but □ 3 □ 3 No No. not limited to direct care areas (such as food 4 4 and medication preparation areas). □ 5 □N/A □N/A <u> 1</u> Yes 2. A.2 Alcohol-based hand rub is readily accessible Yes 2 2 and placed in appropriate locations. □ 3 No. No. 4 4 □ 5 □N/A □ 5 ©N/A 0 1 2.A.3 Healthcare personnel perform hand Yes Yes \bigcap 1 2 2 hygiene: 3 No. **3** No. Before contact with the patient or their 4 4 immediate care environment (even if gloves □ 5 □ N/A ♠N/A □ 5 are worn) Before exiting the patient's care area after touching the patient or the patient's immediate environment (even if gloves are worn) Before performing an aseptic task (e.g., insertion of IV or urinary catheter, even if gloves are worn) After contact with blood, body fluids or contaminated surfaces, (even if gloves are worn)

2. A.4 Healthcare personnel perform hand hygiene using soap and water when hands are visibly soiled (e.g., blood, body fluids) or after caring for a patient with known or suspected <i>C. difficile</i> or norovirus during an outbreak)	☐ Yes ☐ No ☐ N/A	① 1 ② 2 ② 3 ② 4 ② 5	☐ Yes ☐ No ☐ N/A	① 1 ② 2 ② 3 ② 4 ② 5				
*Note: In all other situations, alcohol-based hand rub is preferred.								
2. A.5 Healthcare personnel who have direct	Yes	(i) 1	Yes	O 1				
contact with high-risk patients (e.g., those in intensive care units or ORs) do not wear artificial fingernails or extenders	◯ No ◯ N/A	□ 2□ 3□ 4□ 5	⊖ No	□ 2□ 3□ 4□ 5				
If yo to any of the charge (2, A 1 through 2, A 5) oil	o at 42 CED 492 42	(Co.) (Tog. 8, 0748)						
If no to any of the above (2. A.1 through 2. A.5), cite at 42 CFR 482.42(a) (Tag A-0748)								

Section 2. B Injection Practices and Sharps Safety (Medications, Saline, Other Infusates)

		Manner of Assessment Code		Manner of Assessment Code					
Elements to be assessed		(check all that apply) & Surveyor Notes		(check all that apply) & Surveyor Notes					
Injections are given and sharps safety is managed in a manner consistent with hospital infection control policies and procedures to maximize the prevention of infection and									
communicable disease including the following:			A						
2. B.1 Injections are prepared using aseptic	Yes		C Yes	<u>O</u> 1					
technique in an area that has been cleaned		□ 2		□ 2					
and free of visible blood, body fluids, or	No No	□ 3	☐ No	3					
contaminated equipment.		□ 4		□ 4					
' '	Ĉ N/A	5	Ĉ N/A	5					
2. B.2 Needles are used for only one patient.	Yes	O 1	Yes	O 1					
		2		2					
	No No	3	○ No	3					
		(C) 4		O 4					
	□N/A	5	Ĉ N/A	5					
2. B.3 Syringes are used for only one patient (this	Yes	O 1	Yes	① 1					
includes manufactured prefilled syringes and		(a) 2		2					
insulin pens).	No No	Ö 3	○ No	3					
		(a) 4		Õ 4					
	◯ N/A	5	○ N/A	5					

Interview = 1 Obs

Observation = 2

Infection Control Document Review = 3

Medical Record Review = 4

Other Document Review = 5

2. B.4 The rubber septum on a medication vial is disinfected with alcohol prior to piercing.	☐ Yes ☐ No ☐ N/A	① 1 ② 2 ② 3 ② 4 ① 5	☐ Yes ☐ No ☐ N/A	① 1 ① 2 ② 3 ① 4 ① 5	
2. B.5 Medication vials are entered with a new needle. Note - Reuse of syringes and/or needles to enter a medication vial contaminates the contents of the vial making the vial unsafe for use on additional patients. If a surveyor sees needles or syringes being reused to enter a vial to obtain additional medication for the same patient, no citation should be made if the vial is discarded immediately.	☐ Yes ☐ No ☐ N/A	① 1 ② 2 ② 3 ② 4 ② 5	☐ Yes ☐ No ☐ N/A	① 1 ② 2 ② 3 ② 4 ② 5	
 2. B.6 Medication vials are entered with a new syringe. Note - Reuse of syringes and/or needles to enter a medication vial contaminates the contents of the vial making the vial unsafe for use on additional patients. If a surveyor sees needles or syringes being reused to enter a vial to obtain additional medication for the same patient, no citation should be made if the vial is discarded immediately. 	☐ Yes ☐ No ☐ N/A	① 1 ② 2 ② 3 ② 4 ② 5	☐ Yes ☐ No ☐ N/A	① 1 ② 2 ② 3 ② 4 ② 5	
2. B.7 Single dose (single-use) medication vials are used for only one patient.	☐ Yes ☐ No ☐ N/A	1 2 2 3 4 C 5	☐ Yes ☐ No ☐ N/A	① 1 ② 2 ② 3 ② 4 ② 5	
2. B.8 Bags of IV solution are used for only one patient (and not as a source of flush solution for multiple patients).	☐ Yes ☐ No ☐ N/A	① 1 ② 2 ② 3 ② 4 ② 5	☐ Yes ☐ No ☐ N/A	1 2 3 4 5 5	

2. B.9 Medication administration tubing and	Yes	1		Yes	□ 1	
connectors are used for only one patient.		2			2	
	🗀 No			No No	3	
		4			4	
	Ĉ N/A	□ 5		Ĉ N/A		
	,					
2. B.10 Multi-dose vials are dated when they are	Yes	O 1		Yes	O 1	
first opened and discarded within 28 days		<u>2</u>			2	
unless the manufacturer specifies a different	♠ No	Ö 3		♠ No	<u> </u>	
(shorter or longer) date for that opened vial.		(C) 4		,	O 4	
Note: This is different from the expiration date	Ĉ N/A	□ 5		□ N/A	<u> </u>	
·	₩/A	₩ 3		₩ N/A	₩ 3	
for the vial. The multi-dose vial can be						
dated with either the date opened or the						
discard date as per hospital policies and						
procedures, so long as it is clear what the						
date represents and the same policy is used						
consistently throughout the hospital.						
Consistently throughout the hospital.						
	30	~		3	~	
2. B.11 If multi-dose vials are used for more than	Yes	O 1		Yes	1	
one patient, they do not enter the immediate		<u>2</u>			2	
patient treatment area (e.g., operating room,	🗘 No			○ No	3	
patient room, anesthesia carts).	_	4		_	4	
Note: If multi-dose vials are found in the patient	□ N/A	◯ 5		Ĉ N/A	□ 5	
care area they must be dedicated for single						
patient use and discarded after use.						
'						
2. B.12 All sharps are disposed of in a puncture-	Yes	O 1		Yes	O 1	
resistant sharps container.	₩ TES			₩ 163	<u> </u>	
resistant sharps container.	♠ No	□ 2□ 3		Ĉ No	 2 3	
	⊕ INO	□ 3 □ 4		₩ NO	□ 3 □ 4	
	□ N/A	□ 4 □ 5		Ĉ N/A	□ 4 □ 5	
	₩N/A	₩ 5		₩N/A	₩ 5	
2. B.13 Sharps containers are replaced when the	Yes	1		Yes	O 1	
	₩ Tes			₩ res	<u> </u>	
fill line is reached and disposed of in	◯ No	□ 2 □ 3		○ No	□ 2□ 3	
accordance with State medical waste rules.	₩ NO			₩ NO		
	C NI/A	(C) 4		Pau/a	<u>4</u>	
	□N/A	□ 5		◯ N/A	<u> </u>	
		V V(2) (=		0.5.6.15.//		
If no to any of the above (2.B.1 through 2.B.13), cit	e at 42 CFR 482.42	(a)(1)(Ta	ag A-U749) *See notes on 2.B.5 and	2.B.6 if "no" is ch	ecked.	

Section 2. C Personal Protective Equipment/Standard Precautions

Elements to be assessed		Manner of Assessment Code		Manner of Assessment Code
Personal protective equipment is utilized in a mann	or consistent with	(check all that apply) & Surveyor Notes	duros to maximiza	(check all that apply) & Surveyor Notes
communicable disease including the following:	er consistent with	nospital infection control policies and proce	dures to maximize	the prevention of infection and
C.1 Supplies for adherence to Standard and	Tes Yes	O 1	Yes	O 1
Transmission-based Precautions (e.g., gloves,	₩ 163		₩ 163	
gowns, mouth, eye, nose, and face	□ No		No	<u> </u>
protection) are available and located near	- NO		- NO	Ö 4
point of use.	□N/A	© 5	◯ N/A	<u> </u>
2. C.2 HCP wear gloves for procedures/activities	Yes	O 1	Yes	Q 1
where contact with blood, body fluids,	_	□ 2	-	<u> </u>
mucous membranes, or non-intact skin is	No No		Ĉ No	<u></u>
anticipated.	_	O 4		<u> </u>
	Ĉ N/A	5	□ N/A	₾ 5
2. C.3 HCP change gloves and perform hand	Tes Yes	□ 1	☐ Yes	O 1
hygiene before moving from a contaminated		□ 2		□ 2
body site to a clean body site.	🗖 No	□ 3	O No	□ 3
		□ 4		□ 4
	□ N/A	□ 5	Ĉ N/A	□ 5
2. C.4 Gowns are worn to prevent contamination	Yes	O 1	Yes	O 1
of skin and clothing during		□ 2		□ 2
procedures/activities where contact with	🗀 No	(3	🖾 No	□ 3
blood, body fluids, secretions, or excretions		□ 4		□ 4
are anticipated.	□N/A	5	©N/A	□ 5
2. C.5 Gowns and gloves are removed and hand	Yes	O 1	Yes	O 1
hygiene is performed immediately before		□ 2		□ 2
leaving the patient's environment.	🗀 No	□ 3	No No	□ 3
		4		□ 4
	□ N/A	□ 5	©N/A	□ 5

2. C.6 Appropriate mouth, nose, eye protection is	Yes	O 1	C Yes	O 1
worn for aerosol-generating procedures		□ 2		□ 2
and/or procedures/activities that are likely to	🗀 No	3	🗖 No	
generate splashes or sprays of blood or body		O 4		□ 4
fluids.	[©] N/A	□ 5	□ N/A	፟ 5
2. C.7 Surgical masks are worn by HCP when	Yes	① 1	Yes	O 1
placing a catheter or injecting materials into		□ 2		□ 2
the epidural or subdural space.	O No	3	O No	□ 3
		(a) 4		O 4
	□ N/A	5	□N/A	O 5
If no to any of the above (2.C.1 through 2.C.7), cite	at 42 CFR 482.42(a)(1) (Tag A-0749)		

Section 2. D Environmental Services							
Elements to be assessed			Manner of Assessment Code (check all that apply) & Surveyor Notes				
Environmental services are provided in a manner consistent with hos communicable disease including the following:		ntrol poli	cies and procedures to maximize the prevention of infection and				
2. D.1 HCP wear appropriate PPE to preclude exposure to	🗖 Yes	0 1					
infectious agents or chemicals (PPE can include gloves, gowns,	-	O 2					
masks, and eye protection).	O No	9 3					
	<i>₽</i>	Q 4					
	□ N/A	<u>O</u> 5					
2. D.2 Objects and environmental surfaces in patient care areas	C Yes	<u>O</u> 1					
that are touched frequently (e.g., bed rails, side table, call	~	Q 2					
button) are cleaned and then disinfected when visibly	O No	<u>3</u>					
contaminated or at least daily with an EPA-registered	7	Q 4					
disinfectant.	□ N/A	O 5					
2. D.3 For terminal cleaning (i.e., after patient discharge), all	Yes	(i) 1					
surfaces are thoroughly cleaned and disinfected and towels		2					
and bed linens are replaced with clean towels and bed linens.	□ No	3					
		4					
	□ N/A	O 5					
2. D.4 Cleaners and disinfectants, including disposable wipes, are	Yes	O 1					
used in accordance with manufacturer's instructions (e.g.,		O 2					
dilution, storage, shelf-life, contact time).	No No	3					
		O 4					
	□ N/A	O 5					

2. D.5 Clean, (laundered if not disposable), cloths are used for each	🗖 Yes	0 1	
room or corridor.		O 2	
	□ No	○ 3	
		O 4	
	□ N/A	O 5	
2. D.6 Mop heads and cleaning cloths are laundered at least daily	Yes	O 1	
using appropriate laundry techniques (e.g., following		O 2	
manufacturer instructions when laundering microfiber items).	□ No		
		O 4	
	□ N/A	O 5	
	,		
2. D.7 The facility decontaminates spills of blood or other body	C Yes	O 1	
fluids according to its policies and procedures.		O 2	
	◯ No	3	
		O 4	
	□ N/A	O 5	
2. D.8 Facility has established and follows a cleaning schedule for	Yes	O 1	
areas/equipment to be cleaned/serviced regularly (e.g., HVAC		O 2	
equipment, refrigerators, ice machines, eye wash stations,	🗖 No		
scrub sinks, aerators on faucets).		O 4	
,	□ N/A	□ 5	
	,		
If no to any of the above (2.D.1 through 2.D.8), cite at 42 CFR 482.42	2(a)(1) (Tag A-074	19)	
Laundry is processed in a manner consistent with hospital infection of	ontrol policies an	d proced	ures to maximize the prevention of infection and communicable disease
including the following:	•	•	·
2. D.9 HCP handle soiled textiles/linens in a manner that ensures	Yes	0 1	
segregation of dirty from clean textiles/linens and ensure that		2	
there is not cross contamination of clean textiles/linens prior	□ No	○ 3	
to use.		□ 4	
	□ N/A	O 5	
2. D.10 Soiled textiles/linens are bagged at the point of collection	Yes	O 1	
and kept in a covered leak-proof container or bag at all times		O 2	
until they reach the laundry facility.	_		
arren ency reach the launary racinty.	No No	(
Note: Covers are not needed on contaminated textile	□ No	34	
Note: Covers are not needed on contaminated textile	□ No		
		4	
Note: Covers are not needed on contaminated textile hampers in patient care areas.	□ N/A	O 4 O 5	
Note: Covers are not needed on contaminated textile hampers in patient care areas. 2. D.11 There is clear separation of soiled laundry space from clean		0 4 0 5	
Note: Covers are not needed on contaminated textile hampers in patient care areas. 2. D.11 There is clear separation of soiled laundry space from clean laundry areas and soiled laundry is maintained under	□ N/A □ Yes	0 4 0 5	
Note: Covers are not needed on contaminated textile hampers in patient care areas. 2. D.11 There is clear separation of soiled laundry space from clean	□ N/A	0 4 0 5	
Note: Covers are not needed on contaminated textile hampers in patient care areas. 2. D.11 There is clear separation of soiled laundry space from clean laundry areas and soiled laundry is maintained under	□ N/A □ Yes □ No	0 4 5 5 C 1 C 2 C 3 C 4	
Note: Covers are not needed on contaminated textile hampers in patient care areas. 2. D.11 There is clear separation of soiled laundry space from clean laundry areas and soiled laundry is maintained under	□ N/A □ Yes □ No □ N/A	0 4 0 5 5 C 4 0 5 5 C 4 0 5 C 5 C 6 C 6 C 6 C 6 C 6 C 6 C 6 C 6 C	

Interview = 1

Observation = 2

Infection Control Document Review = 3

Medical Record Review = 4

Other Document Review = 5

	stent with hospita	I infectio	on control policies and procedures to maximize the prevention of infection
and communicable disease including the following:			
2. D.12 Reusable noncritical patient-care devices (e.g., blood	Yes	□ 1	
pressure cuffs, oximeter probes) are disinfected when visibly soiled		□ 2	
and on a regular basis (such as after use on each patient or once	🟳 No	□ 3	
daily or once weekly), and there is clear delineation of		4	
responsibility for this among healthcare personnel. Note: For	🗖 N/A	□ 5	
patients on Contact Precautions, if dedicated, disposable			
devices are not available, noncritical patient-care devices are			
disinfected after use on each patient.			
·			
2. D.13 Manufacturers' instructions for cleaning noncritical medical	Yes	O 1	
equipment are followed.		□ 2	
• •	🗖 No	□ 3	
		4	
	© N/A	<u> </u>	
If no to any of the above (2.D.12 through 2.D.13), cite at 42 CFR 482.		749)	
2. D.14 Hydrotherapy equipment (e.g., Hubbard tanks, tubs,	C Yes	□ 1	
whirlpools, spas, birthing tanks) are drained, cleaned, and		2	
disinfected using an EPA-registered disinfectant according to	○ No	□ 3	
manufacturer's instructions after each patient use.		4	
·	□ N/A	□ 5	
If no cite at 42 CFR 482.42(a)(1) (Tag A-0749)	<u> </u>		

Module 3: Equipment Reprocessing

Section 3.A. Reprocessing of Semi-Critical Equipment **Manner of Assessment Code Manner of Assessment Code** Elements to be assessed (check all that apply) & Surveyor Notes (check all that apply) & Surveyor Notes High-Level Disinfection of Reusable Instruments and Devices is accomplished in a manner consistent with hospital infection control policies and procedures to maximize the prevention of infection and communicable disease including: Note: Hospital policies should address what to do when there are discrepancies between manufacturer's instructions for a device and manufacturer's instructions for a device reprocessor. 3. A.1 All reusable semi-critical items receive at Yes 0 1 Yes 2 2 least high-level disinfection. □ No 3 O No 3 4 4 □ N/A O N/A □ 5 5

3. A.2 High-level disinfection is performed on-site. Continue if "yes." If "no," skip to 3.A.14.If the response is No, no citation is made in response to this question.	C Yes	① 1 ② 2 ② 3 ② 4 ② 5	Ĉ Yes Ĉ No	1 2 3 6 4 6 5	
 A.3 Flexible endoscopes are inspected for damage and leak tested as part of each reprocessing cycle. 	☐ Yes ☐ No ☐ N/A	1 2 3 4 5	○ Yes○ No○ N/A	① 1 ② 2 ② 3 ② 4 ② 5	
3. A.4 Items are thoroughly pre-cleaned according to manufacturer instructions and visually inspected for residual soil prior to high-level disinfection. Note: for lumened instruments (e.g., endoscopes), pre-cleaning must include all device channels and lumens with cleaning brushes appropriate for size of instrument channel or port.	☐ Yes ☐ No ☐ N/A	1 2 3 4 5	☐ Yes ☐ No ☐ N/A	1 2 3 4 6 5	
3. A.5 Enzymatic cleaner or detergent is used and discarded according to manufacturer's instructions (typically after each use).	Yes No N/A	① 1 ② 2 ② 3 ② 4 ② 5	☐ Yes ☐ No ☐ N/A	1 2 1 3 1 4 5 5	
3. A.6 Cleaning brushes are disposable or cleaned and high-level disinfected or sterilized (per manufacturer's instructions) after each use.	☐ Yes ☐ No ☐ N/A	1 2 3 Q 4 Q 5	☐ Yes ☐ No ☐ N/A	1 2 3 4 C 5	
 3. A.7 For chemicals used in high-level disinfection, manufacturer's instructions are followed for: preparation testing for appropriate concentration replacement (e.g., prior to expiration or loss of efficacy). 	☐ Yes ☐ No ☐ N/A	1 2 3 4 C 5	☐ Yes ☐ No ☐ N/A	① 1 ② 2 ② 3 ② 4 ② 5	

3. A.8 If automated reprocessing equipment is	Yes	O 1	C Yes	O 1
used, proper connectors are used to assure		O 2		① 2
that channels and lumens are appropriately	🗖 No	□ 3	○ No	□ 3
disinfected.		<u> </u>		O 4
	🗖 N/A	<u> </u>	□N/A	
3. A.9 Devices are disinfected for the appropriate	Yes	O 1	Yes	O 1
length of time as specified by manufacturer's		□ 2		① 2
instructions.	🗖 No	<u></u> 3	🖰 No	□ 3
		O 4	_	O 4
	[□] N/A	5	□N/A	<u></u> 5
3. A.10 Devices are disinfected at the appropriate	Yes	① 1	C Yes	① 1
temperature as specified by manufacturer's	0	□ 2		2
instructions.	🖰 No		No No	3
	~	O 4	~	4
	□ N/A	<u> </u>	□N/A	○ 5
3. A.11 After high-level disinfection, devices are	Yes	O 1	C Yes	① 1
rinsed with sterile water, filtered water, or	_	□ 2	-	2
tap water followed by a rinse with 70% - 90%	No No		🗖 No	3
ethyl or isopropyl alcohol.	_	O 4	-	O 4
	□ N/A	O 5	©n/A	
3. A.12 Devices are dried thoroughly prior to	Yes		C Yes	O 1
reuse.	2	<u> </u>	2	
Note: for lumened instruments (e.g., endoscopes)	O No	□ 3	O No	<u></u> 3
this includes flushing channels with alcohol	~	○ 4	€	
and forcing air through the channels.	□ N/A	<u>5</u>	© N/A	<u>0</u> 5
3. A.13 Routine maintenance procedures for high-	Yes	O 1	C Yes	O 1
level disinfection equipment conform to	⊘	□ 2	<u>م</u>	□ 2 □
manufacturer's instruction; confirm	No No	○ 3	O No	
maintenance records are available.		O 4	₽	<u>0</u> 4
	© N/A	5	□N/A	O 5
3. A.14 After high-level disinfection, devices are	Yes	O 1	Yes	
stored in a manner to protect from damage	♠	O 2	₹	0 2
or contamination (Note: endoscopes must be	🖰 No	○ 3	No No	□ 3
hung in a vertical position).	₹ N/A	O 4	₽	O 4
	Ĉ N/A	5	©N/A	O 5
3. A.15 The facility has a system in place to	Yes	0 1	Yes	0 1
identify which instrument (e.g., endoscope)	Ĉ N-	O 2	Ĉ N-	0 2
was used on a patient via a log for each	🗖 No	O 3	No No	O 3
procedure.	□ N/A	□ 4□ 5	PN/A	0 4
If you have great the otherwise (2.4.4 and / a. 2.4.2 d.	,	'E'	©n/A	○ 5
If no to any of the above (3.A.1 and/or 3.A.3 through	gn 3.A.15), cite at 4	42 CFK 482.42(a)(1) (1ag A-0/49)		

Section 3. B Reprocessing of Critical Equipment Sterilization of Reusable Instruments and Devices

Elements to be assessed			Manner of Assessment Code (check all that apply) & Surveyor Notes
•	Hospital policies		spital infection control policies and procedures to maximize the prevention dress what to do when there are discrepancies between manufacturer's
Note: for lumened instruments, pre-cleaning must include all device channels and lumens with cleaning brushes appropriate for size of instrument channel or port.	□ N/A	5	
3. B.2 All reusable critical instruments and devices are sterilized on site.If no, no citation is issued and skip to 3.B.12.	☐ Yes ☐ No ☐ N/A	☐ 1 ☐ 2 ☐ 3 ☐ 4 ☐ 5	
3. B.3 Enzymatic cleaner or detergent is used and discarded according to manufacturer's instructions (typically after each use).	☐ Yes ☐ No ☐ N/A	1 2 3 4 C 5	
3. B.4 Cleaning brushes are disposable or cleaned and high-level disinfected or sterilized (per manufacturer's instructions) after each use.	☐ Yes ☐ No ☐ N/A	① 1 ② 2 ② 3 ② 4 ② 5	
3. B.5 After pre-cleaning, instruments are appropriately wrapped/packaged for sterilization (e.g., package system selected is compatible with the sterilization process being performed, hinged instruments are open, and instruments are disassembled if indicated by the manufacturer).	☐ Yes ☐ No ☐ N/A	1 2 3 4 5	

3. B.6 A chemical indicator (process indicator) is placed correctly in the instrument packs in every load.	☐ Yes ☐ No ☐ N/A	☐ 1 ☐ 2 ☐ 3 ☐ 4 ☐ 5	
3. B.7 A biological indicator is used at least weekly for each sterilizer and with every load containing implantable items.	☐ Yes ☐ No ☐ N/A	1 2 3 4 C 5	
3. B.8 For dynamic air removal-type sterilizers, a Bowie-Dick test is performed each day the sterilizer is used to verify efficacy of air removal.	☐ Yes ☐ No ☐ N/A	1 2 3 4 D 5	
3. B.9 Sterile packs are labeled with the sterilizer used, the cycle or load number, and the date of sterilization.	☐ Yes ☐ No ☐ N/A	1 2 3 4 C 5	
3. B.10 Logs for each sterilizer cycle are current and include results from each load.	☐ Yes ☐ No ☐ N/A	1 2 3 G 4 D 5	
3. B.11 Routine maintenance for sterilization equipment is performed according to manufacturer's instructions (confirm maintenance records are available).	☐ Yes ☐ No ☐ N/A	1 2 3 4 1 5 5	
3. B.12 After sterilization, medical devices and instruments are stored so that sterility is not compromised.	☐ Yes ☐ No ☐ N/A	1 2 3 4 0 5	

B.13 Sterile packages are inspected for integrity and compromised packages are reprocessed prior to use.	☐ Yes ☐ No ☐ N/A	1 1 2 1 3 1 4 1 5	
 3.B.14 If immediate-use steam sterilization is performed, the following criteria are met: The item being sterilized is thoroughly cleaned prior to placing it in the sterilizer container (that is FDA cleared for use with the cycle) or tray The sterilizer cycle being used is one that is approved by both the instrument and sterilizer manufacturer The sterilizer function is monitored with monitors (e.g., mechanical, chemical and biologic) that are approved for the cycle being used The facility maintains a sufficient volume of instruments to meet the surgical volume and permit time to complete all steps of reprocessing 	☐ Yes ☐ No ☐ N/A	1 2 3 4 0 5	
3. B.15 Instruments that are subject to immediate use sterilization procedures are used immediately and handled in a manner to prevent contamination during transport from the sterilizer to the patient. 3. B.16 HCP respond (i.e., recall of device and risk assessment) according to facility policies and procedures in the event of a reprocessing error/failure that could result in the transmission of infectious disease.	☐ Yes ☐ No ☐ N/A ☐ Yes ☐ No	1 6 2 6 3 6 4 6 5	
If no to any of the above (3.B.1 and/or 3.B.3 through 3.B.16), cite at 4	□ N/A 12 CFR 482.42(a)(2	51) (Tag A-	0749)

Section 3. C Single-Use Devices (SUDs)								
Elements to be assessed		Manner of Assessment Code (check all that apply) & Surveyor Notes		Manner of Assessment Code (check all that apply) & Surveyor Notes				
Single use devices are used in a manner consistent wincluding the following:	vith hospital infec	tion control policies and procedures to maxi	mize the prevention	on of infection and communicable disease				
C.1 Single use devices are discarded after use and not used for more than one patient.	☐ Yes ☐ No ☐ N/A	1 2 2 3 4 C 5	☐ Yes ☐ No ☐ N/A	① 1 ① 2 ② 3 ② 4 ① 5				
If no, do not cite and go to 3.C.2								
3. C.2 If the hospital elects to reuse single-use devices, these devices are reprocessed by an entity or a third party reprocessor that is registered with the FDA as a third-party reprocessor and cleared by the FDA to reprocess the specific device in question. The hospital must have documentation from the third party reprocessor confirming this is the case.	☐ Yes ☐ No ☐ N/A	① 1 ② 2 ② 3 ② 4 ① 5	☐ Yes ☐ No ☐ N/A	① 1 ○ 2 ○ 3 ○ 4 ○ 5				
If no, cite at 42 CFR 482.42(a) (Tag A-0748) Module 4: Patient Tracers 4. The hospital develops and implements infection	☐ Yes	□ 1						
control policies and procedures related to the following sections to ensure an environment minimizing risk for spread of infection and maximizing prevention of infection and communicable disease.	☐ No ☐ N/A	© 2 © 3 © 4 © 5						
Do not cite unless the lack of an individual protocol communicable disease. If so, cite at 42 CFR 482.42(per of protocol failures that indicate the abs	ence of an active p	program to control infections and				

Section 4. A Urinary Catheter Tracer Manner of Assessment Code Manner of Assessment Code Elements to be assessed (check all that apply) & Surveyor Notes (check all that apply) & Surveyor Notes Urinary catheters are inserted, accessed, and maintained in a manner consistent with hospital infection control policies and procedures to maximize the prevention of infection and communicable disease including the following: Insertion: 4. A.1 The hospital has guidelines for appropriate 1 **1** Yes Yes 2 **2** indications for urinary catheters. □ No No No **(**) 3 4 **(**) 4 O N/A □N/A □ 5 **7** 5 No citation **1 1** 4. A.2 Hand hygiene performed before and after Yes Yes Yes 2 **(**) 2 insertion. 3 O No ☐ No **(**) 3 4 **(**) 4 \bigcirc 5 □ N/A **(**) 5 **(**) 1 4. A.3 Catheter placed using aseptic technique Yes \bigcirc 1 T Yes 2 (C) 2 and sterile equipment. □ 3 □ No Mo. **(**) 3 4 **4 (**) 5 O N/A \bigcirc 5 □N/A 4. A.4 Catheter secured properly after insertion. Yes \bigcirc 1 T Yes 1 **1** 2 □ 3 **(**) 3 □ No Mo. **1** 4 4 □ 5 **1** 5 □N/A □ N/A If no to 4.A.2 through 4.A.4, cite at 42 CFR 482.42(a)(1) (Tag A-0749) 4. A.5 Catheter insertion and indication Yes (i) Tage Yes 1 1 □ 2 **2** documented. □ 3 **(**) 3 O No No 🏳 4 **4** □ N/A □N/A □ 5 **7** 5 If no, cite at 42 CFR 482.24(c)(2)(vi) (Tag A-0467)

Accessing/Maintenance:						
4. A.6 Hand hygiene performed before and after	Yes	O 1		C Yes	O 1	
manipulating catheter.		O 2			0 2	
mamparating catheter.	○ No	3		◯ No	3	
	0 110	O 4		0 110	0 4	
	☐ N/A	O 5		◯ N/A	5	
4. A.7 Catheter and collecting tubing are not	Yes	O 1		Yes	O 1	
disconnected (irrigation avoided).		2			O 2	
	□ No	□ 3		O No	○ 3	
		O 4			O 4	
	□ N/A	O 5		□ N/A	O 5	
4. A.8 Urine bag emptied using aseptic technique.	Yes	O 1		Yes	O 1	
	~	O 2		~	O 2	
	○ No	3		O No	○ 3	
		O 4			O 4	
	◯ N/A	O 5		□ N/A	5	
4. A.9 Urine samples obtained aseptically (via	Yes	O 1		Yes	Q 1	
needleless port for small volume).	~	O 2			<u>0</u> 2	
	○ No	3		○ No	<u>0</u> 3	
	_	Q 4			6 4	
	□ N/A	O 5		O _{N/A}	6 5	
4. A.10 Urine bag kept below level of bladder at all	Yes	O 1		Yes	O 1	
times.	-	O 2		-	O 2	
	○ No	O 3		◯ No	○ 3	
		O 4		~	O 4	
	□ N/A	O 5		□ N/A	O 5	
4. A.11 Catheter tubing unobstructed and free of	Yes	0 1		Yes	O 1	
kinking.	-	O 2		-	O 2	
	◯ No	Q 3		□ No		
	~ .	Q 4		~	O 4	
	□ N/A	O 5		□ N/A	O 5	
If no to any of 4.A.6 through 4.A.11, cite at 42 CFR 4	82.42(a)(1) (Tag	A-0749).				
4. A.12 Need for urinary catheters reviewed daily	Yes	① 1		Yes	O 1	
with prompt removal of unnecessary urinary	<u> </u>	<u> </u>		<u> </u>	□ 2	
catheters.	○ No	3		🗖 No	□ 3	
	-	4		-	O 4	
	□ N/A	<u> </u>		□ N/A	□ 5	
	•	_		•		
No citation for 4.A.12			•			

Section 4. B Central Venous Catheter Tracer Manner of Assessment Code Manner of Assessment Code Elements to be assessed (check all that apply) & Surveyor Notes (check all that apply) & Surveyor Notes Central venous catheters are inserted, accessed and maintained in a manner consistent with hospital infection control policies and procedures to maximize the prevention of infection and communicable disease including the following: Insertion: Yes Yes 4. B.1 Hand Hygiene performed before and after 2 <u>2</u> insertion. □ 3 3 □ No No. □ 4 □ 5 \bigcirc 5 ⁽¹⁾ 1 □ 1 4. B.2 Maximal barrier precautions used for Yes Yes <u>2</u> insertion (includes use of cap, mask, sterile □ 3 3 O No □ No gown, sterile gloves, and a sterile full body 4 4 drape). [□] 5 O N/A O N/A \Box 1 1 4. B.3 > 0.5% chlorhexidine with alcohol used for Yes Yes 2 □ 2 skin antisepsis prior to insertion (If □ 3 □ 3 ☐ No No. contraindicated, tincture of iodine, an 4 4 iodophor, or 70% alcohol can be used as □N/A O N/A \Box 5 \bigcirc 5 alternatives). O 1 0 1 Yes 4. B.4 Sterile gauze or sterile, transparent, semi Yes 2 2 permeable dressing used to cover catheter \Box 3 \Box 3 □ No □ No site (may not apply for well-healed tunneled 4 4 catheters). 5 5 □ N/A O N/A If no to any of the above (4.B.1 through 4.B.4), cite at 42 CFR 482.42(a)(1) (Tag A-0749) \bigcirc 1 4. B.5 Central line insertion and indication \Box 1 Yes Yes 2 2 documented. □ 3 □ 3 □ No □ No 4 4 □ N/A \Box 5 □ N/A \Box 5 If no to 4.B.5, cite at 42 CFR 482.24(c)(2)(vi) (Tag A-0467)

Accessing/Maintenance:					
4. B.6 Hand hygiene performed before and after	Yes	1	Yes	O 1	
manipulating catheter (even if gloves worn).		2		2	
	○ No	□ 3	🗀 No	□ 3	
		4		4	
	◯ N/A		🗖 N/A	□ 5	
4. B.7 Dressings that are wet, soiled, or dislodged	Yes	<u> </u>	Yes	<u> </u>	
are changed promptly.	-	<u> </u>	-	<u> </u>	
	○ No	<u>3</u>	🗖 No	<u>3</u>	
		<u> </u>	<u>~</u>	O 4	
	□ N/A	□ 5	[©] N/A	□ 5	
	A		A	A	
4. B.8 Dressing changed with aseptic technique	🗀 Yes	0 1	🗖 Yes		
using clean or sterile gloves.	□ No	<u>2</u>	☐ No	23	
	∪ No	34	□ NO	_	
	[©] N/A	□ 4 □ 5	[⊕] N/A	45	
	□ N/A	⊖ 5	□ N/A	<u></u> 5	
4. B.9 Access port is scrubbed with an appropriate	Yes	□ 1	Yes	O 1	
antiseptic (chlorhexidine, povidone iodine, an		<u> </u>		2	
iodophor, or 70% alcohol) prior to accessing.	O No	☐ 3	☐ No		
		□ 4		4	
	[♠] N/A	<u>5</u>	[©] N/A	□ 5	
4. B 10 Catheter accessed only with sterile	☐ Yes	O 1	☐ Yes	O 1	
devices.	□ 163	\bigcirc 2	□ 163	\bigcirc 2	
devices.	No No	<u>2</u> 3	☐ No	<u>2</u> 3	
	140	O 4	0 110	<u> </u>	
	[□] N/A	<u> </u>	[©] N/A	<u> </u>	
	,		,		
If no to any of 4.B.6 through 4.B.10, cite at 42 CFR 4					
4. B.11 Need for central venous catheters	Yes	<u> </u>	Yes	O 1	
reviewed daily with prompt removal of	~	2	_	<u>2</u> 2	
unnecessary lines.	○ No	<u>3</u>	□ No	<u></u>	
	6	Q 4	£	<u></u> 4	
	◯ N/A	□ 5	□ N/A	□ 5	
No situation for 4 P 11					
No citation for 4.B.11					

Elements to be assessed		N	Manner of Assessment Code		N	Manner of Assessment Code		
			all that apply) & Surveyor Notes	•	all that apply) & Surveyor Notes			
Respiratory procedures are performed in a manner consistent with hospital infection control policies and procedures to maximize the prevention of infection and communicable disease including the following:								
General respiratory therapy practices (apply to pat	ients with and wit	hout ver	itilators):					
4. C.1 Hand hygiene is performed before and after contact with patient or any respiratory equipment used on patient.	☐ Yes ☐ No	① 1 ② 2 ② 3		☐ Yes ☐ No	123			
	□ N/A	45		□ N/A	□ 4□ 5			
C.2 Gloves are worn when in contact with respiratory secretions and changed before	☐ Yes	12		☐ Yes	① 1 ② 2			
contact with another patient, object, or environmental surface.	□ No	☐ 3 ☐ 4		Ĉ No	① 3 ② 4			
environmental surface.	© N/A	5		[©] N/A	O 5			
4. C.3 Only sterile water is used for nebulization.	☐ Yes	1 0 2		C Yes	① 1 ② 2			
	◯ No	① 3 ② 4		□ No	34			
	[©] N/A	<u>5</u>		□ N/A	<u>5</u>			
4. C.4 Single-dose vials for aerosolized	C Yes	☐ 1 ☐ 2		C Yes	O 1			
medications are not used for more than one patient.	€ No	□ 3		◯ No	(i) 2 (i) 3			
	□ N/A	□ 4□ 5		Ĉ N/A	□ 4□ 5			
4. C.5 If multi-dose vials for aerosolized	☐ Yes	O 1		C Yes	<u> </u>			
medications are used, manufacturer's instructions for handling, storing, and	□ No	① 2 ② 3		□ No	23			
dispensing the medications are followed.	□ N/A	① 4 ① 5		[©] N/A	45			

		_				
4. C.6 If multi-dose vials for aerosolized	Yes	¹ 1		Yes	1	
medications are used for more than one	A	<u>2</u>			<u>2</u> 2	
patient, they are restricted to a centralized	☐ No	<u>3</u>		○ No	<u>3</u>	
medication area and do not enter the	□ N/A	☐ 4 ☐ 5		€ N/A	(L) 4	
immediate patient treatment area.	□ N/A	□ 5		™ N/A	□ 5	
'						
4. C.7 Nebulizers (e.g., mask/mouthpiece, cup) are	Yes	O 1		☐ Yes	O 1	
rinsed with sterile water (or with tap water		<u> </u>			2	
followed by isopropyl alcohol) and dried	☐ No	□ 3		○ No	∃ 3	
thoroughly between uses on the same		□ 4			4	
patient.	[□] N/A	<u> </u>		□ N/A	□ 5	
If no to any of the above (4.C.1 through 4.C.7), cite			g A-0749)			
4. C.8 Hospital has a comprehensive oral-hygiene	Yes	(C) 1		Yes		
program (that might include the use of an	2	<u>2</u>		2	2	
antiseptic agent) for patients who are at high	Ĉ No	<u>3</u>		Ĉ No	<u>3</u>	
risk for health-care associated pneumonia.	△ N/A	(C) 4		(n 1/2	O 4	
	□ N/A	() 5		◯ N/A	□ 5	
4. C.9 In the absence of medical	Yes	<u> </u>		Yes	<u> </u>	-
contraindication(s), head of bed is elevated at	103	<u> </u>		₩ 103	_ 1	
an angle of 30-45 degrees for patients at	□ No	<u> </u>		No No	<u>2</u>	
high risk for aspiration (e.g., a person	- 110	<u> </u>		₩ 110	<u> </u>	
receiving mechanically assisted ventilation	□N/A	<u> </u>		□ N/A	<u> </u>	
and/or who has an enteral tube in place).	,)			_ 3	
and, or who has an effect at case in place,.						
No citation for 4.C.8-9						
Ventilators:						
Ventilators are used in a manner consistent with ho	snital infection co	ntrol poli	cies and procedures to maximize th	e prevention of int	ection an	ud communicable disease
including the following:	opital illicotion co.	ici oi poii	ores and procedures to maximize th	e prevention or in	cotion an	a communicable alsease
4. C.10 Ventilator circuit is changed if visibly soiled	🖺 Yes	<u> </u>		Yes	O 1	
or mechanically malfunctioning.		<u>2</u>			<u>2</u>	
	□ No	<u> 3</u>		□ No	Ö 3	
		<u> </u>		-	Ö 4	
	□N/A	<u> </u>		□N/A	5	
	,	· ·		' '		
Interview = 1 Observation = 2	Infection Contro	l Docum	nent Review = 3 Medical Rec	ord Review = 4	Othe	r Document Review = 5

Infection Control Document Review = 3 Medical Record Review = 4 Other Document Review = 5

4. C.11 Sterile water is used to fill bubbling	Yes	O 1		Yes	O 1	
humidifiers (if applicable).		2			2	
, ,	☐ No	□ 3		○ No	3	
		4			4	
	[□] N/A	□ 5		🗖 N/A	□ 5	
4. C.12 Condensate that collects in the tubing of a	C Yes	🗘 1		Yes	O 1	
mechanical ventilator is periodically drained	_	2		_	<u>2</u>	
and discarded, taking precautions not to	□ No	3		○ No	□ 3	
allow condensate to drain toward the patient.	6	4		_	<u> </u>	
	Ĉ N/A	5		□ N/A	<u> </u>	
	A	Α.		20	A	
4. C 13 If single-use open-system suction catheter	C Yes	0 1		🗀 Yes		
is employed, a sterile, single-use catheter is	<u> </u>	<u>Q</u> 2		~ · ·	<u>2</u>	
used.	□ No	<u>3</u>		○ No	<u>3</u>	
	A	Q 4		♠ 1.74	Q 4	
	□ N/A	□ 5		Ĉ N∕A	□ 5	
4. C.14 If multi-use closed-system suction catheter	☐ Yes	O 1	+	C Yes	O 1	
is employed, only sterile fluid is used to	U 163	<u> </u>		₩ 163	<u> </u>	
remove secretions upon reentry into the	□ No	<u>2</u> 3		○ No	<u>2</u> 3	
respiratory tract.	- NO	<u> </u>		₩ 110	□ 4	
respiratory tract.	□ N/A	<u> </u>		◯ N/A	<u> </u>	
	- N/A)		₩ N/A	₩ 3	
If no to any of the above (4.C.10 through 4.C.14), ci	te at 42 CFR 482.4	2(a)(1) ((Tag A-0749)			
4. C.15 Sedation is lightened daily in eligible	Yes	1		Yes	O 1	
patients.		2			□ 2	
	♠ No	☐ 3		🗖 No	<u> </u>	
		4			<u> </u>	
	◯ N/A	<u> </u>		◯ N/A	□ 5	
4. C.16 Spontaneous breathing trials are	C Yes	O 1		Yes	O 1	
performed daily in eligible patients.		2		-	<u> </u>	
	◯ No	<u> </u>		○ No	<u> </u>	
	A .	<u> </u>			<u> </u>	
	□ N/A	□ 5		◯ N/A	□ 5	
N						
No citation for 4.C.15-16						

Interview = 1 Observation = 2

Infection Control Document Review = 3

Medical Record Review = 4

Other Document Review = 5

Section 4. D Spinal Injection Procedures Manner of Assessment Code Manner of Assessment Code Elements to be assessed (check all that apply) & Surveyor Notes (check all that apply) & Surveyor Notes Spinal injection procedures are performed in a manner consistent with hospital infection control policies and procedures to maximize the prevention of infection and communicable disease including the following: Yes Yes 1 4. D.1 Hand hygiene performed before and after 2 2 the procedure. O No 3 O No 4 4 □N/A \square 5 O N/A 5 Yes 4. D.2 The spinal injection procedure is performed 0 1 1 Yes 2 2 using aseptic technique and sterile 3 3 O No O No equipment, including use of sterile gloves. 4 4 □ N/A **6** 5 □ N/A **6** 5 4. D.3 Surgical masks are worn by HCP when Yes 1 Yes \square 1 □ 2 2 placing a catheter or injecting materials into 3 □ No No. the epidural or subdural space. 4 4 ©N/A □N/A □ 5

If no to any of the above (4.D.1 through 4.D.3), cite at 42 CFR 482.42(a)(1) (Tag A-0749)

Section 4. E Point of Care Devices (e.g. Blood Glucose Meter, INR Monitor)

Check all that apply & Surveyor Notes Check all that apply & Surveyor Notes	Elements to be assessed		Manner of Assessment Code		Manner of Assessment Code					
4. E.1 Hand hygiene is performed before and after the procedure.	Elements to be assessed		(check all that apply) & Surveyor Notes		(check all that apply) & Surveyor Notes					
4. E.1 Hand hyglene is performed before and after the procedure. O 1		Point of care devices are used in a manner consistent with hospital infection control policies and procedures to maximize the prevention of infection and communicable								
the procedure: No										
4. E.2 Gloves are worn by healthcare personnel when performing the finger stick procedure to obtain the sample of blood and are removed after the procedure (followed by hand hygiene). 4. E.3 Finger stick devices are not used for more than one patient. 4. E.4 If used for more than one patient, the point-of-care device is cleaned and disinfected after every use according to manufacturer's instructions for cleaning and disinfection, then the device should not be used for >1 patient. 4. E.5 Insulin pens are used for only one patient C No C 3 C 2 C C 2 C C C C C C C C C C C C C		Yes		Yes	O 1					
4. E.2 Gloves are worn by healthcare personnel when performing the finger stick procedure to obtain the sample of blood and are removed after the procedure (followed by hand hygiene). 4. E.3 Finger stick devices are not used for more than one patient. Note: This includes both the lancet and the lancet holding device. 4. E.4 If used for more than one patient, the point-of-care device is cleaned and disinfected after every use according to manufacturer's instructions. Note: if manufacturer does not provide instructions for cleaning and disinfection, then the device should not be used for >1 patient. 4. E.5 Insulin pens are used for only one patient C N/A C 5 C 1 C 2 C NO C 3	the procedure.			_	□ 2					
4. E.2 Gloves are worn by healthcare personnel when performing the finger stick procedure to obtain the sample of blood and are removed after the procedure (followed by hand hygiene). 4. E.3 Finger stick devices are not used for more than one patient. Note: This includes both the lancet and the lancet holding device. 4. E.4 If used for more than one patient, the point-of-care device is cleaned and disinfected after every use according to manufacturer's instructions. Note: if manufacturer does not provide instructions for cleaning and disinfection, then the device should not be used for >1 patient. 4. E.5 Insulin pens are used for only one patient 1 Yes		□ No		□ No	□ 3					
4. E.2 Gloves are worn by healthcare personnel when performing the finger stick procedure to obtain the sample of blood and are removed after the procedure (followed by hand hygiene). 4. E.3 Finger stick devices are not used for more than one patient. 4. E.3 Finger stick devices are not used for more than one patient. 4. E.4 If used for more than one patient, the point-of-care device is cleaned and disinfected after every use according to manufacturer's instructions. Note: if manufacturer does not provide instructions for cleaning and disinfection, then the device should not be used for >1 patient. 4. E.5 Insulin pens are used for only one patient 4. E.5 Insulin pens are used for only one patient 4. E.5 Insulin pens are used for only one patient 4. E.5 Insulin pens are used for only one patient 4. E.5 Insulin pens are used for only one patient 4. E.6 Insulin pens are used for only one patient 4. E.7 Insulin pens are used for only one patient 5. Insulin pens are used for only one patient 6. Yes 6. 1 7. Yes 7. 1 7. Yes 9. 1 9. Yes		_			□ 4					
when performing the finger stick procedure to obtain the sample of blood and are removed after the procedure (followed by hand hygiene). 4. E.3 Finger stick devices are not used for more than one patient. Note: This includes both the lancet and the lancet holding device. 4. E.4 If used for more than one patient, the point-of-care device is cleaned and disinfected after every use according to manufacturer's instructions. Note: if manufacturer does not provide instructions for cleaning and disinfection, then the device should not be used for >1 patient. 4. E.5 Insulin pens are used for only one patient 1		□ N/A	□ 5	□ N/A	O 5					
when performing the finger stick procedure to obtain the sample of blood and are removed after the procedure (followed by hand hygiene). 4. E.3 Finger stick devices are not used for more than one patient. Note: This includes both the lancet and the lancet holding device. 4. E.4 If used for more than one patient, the point-of-care device is cleaned and disinfected after every use according to manufacturer's instructions. Note: if manufacturer does not provide instructions for cleaning and disinfection, then the device should not be used for >1 patient. 4. E.5 Insulin pens are used for only one patient 4. E.5 Insulin pens are used for only one patient 4. E.5 Insulin pens are used for only one patient 5. No. 3	4. E.2 Gloves are worn by healthcare personnel	🗇 Yes		🗖 Yes	6 1					
removed after the procedure (followed by hand hygiene). A. E.3 Finger stick devices are not used for more than one patient. Note: This includes both the lancet and the lancet holding device. A. E.4 If used for more than one patient, the point-of-care device is cleaned and disinfected after every use according to manufacturer's instructions. Note: if manufacturer does not provide instructions for cleaning and disinfection, then the device should not be used for >1 patient. A. E.5 Insulin pens are used for only one patient A. E.5 Insulin pens are used for only one patient A. E.5 Insulin pens are used for only one patient A. E.5 Insulin pens are used for only one patient A. E.5 Insulin pens are used for only one patient A. E.6 Insulin pens are used for only one patient A. E.7 Insulin pens are used for only one patient A. E.8 Insulin pens are used for only one patient A. E.8 Insulin pens are used for only one patient A. E.8 Insulin pens are used for only one patient A. E.9 Insulin pens are used for o										
removed after the procedure (followed by hand hygiene). A. E.3 Finger stick devices are not used for more than one patient. Note: This includes both the lancet and the lancet holding device. A. E.4 If used for more than one patient, the point-of-care device is cleaned and disinfected after every use according to manufacturer's instructions. Note: if manufacturer does not provide instructions for cleaning and disinfection, then the device should not be used for >1 patient. A. E.5 Insulin pens are used for only one patient A. E.5 Insulin pens are used for only one patient A. E.5 Insulin pens are used for only one patient A. E.6 Insulin pens are used for only one patient A. E.7 Insulin pens are used for only one patient A. E.8 Insulin pens are used for only one patient A. E.8 Insulin pens are used for only one patient A. E.9 Insulin pens are used for only one patient A. E.8 Insulin pens are used for only one patient A. E.9 Insulin pens are used for only one patient A. E.9 Insulin pens are used for only one patient A. E.9 Insulin pens are used for only one patient A. E.9 Insulin pens are used for only one patient A. E.9 Insulin pens are used for only one patient A. E.9 Insulin pens are used for only one patient A. E.9 Insulin pens are used for only one patient A. E.9 Insulin pens are used for only one patient A. E.9 Insulin pens are used for only one patient A. E.9 Insulin pens are used for only one patient A. E.9 Insulin pens are used for only one patient A. E.9 Insulin pens are used for only one patient A. E.9 Insulin pens are used for only one patient A. E.9 Insulin pens are used for only one patient	to obtain the sample of blood and are	☐ No		🗖 No	□ 3					
4. E.3 Finger stick devices are not used for more than one patient. Note: This includes both the lancet and the lancet holding device. No	removed after the procedure (followed by		<u></u> 4							
than one patient. Note: This includes both the lancet and the lancet holding device. ONO ONA ONA OS ONA ONA ONA OS OS ONA ONA OS OS ONA ONA OS OS ONA OS OS ONA OS OS ONA OS OS OS ONA OS	hand hygiene).	◯ N/A	<u> </u>	© N/A	□ 5					
than one patient. Note: This includes both the lancet and the lancet holding device. ONO ONA ONA OS ONA ONA ONA OS OS ONA ONA OS OS ONA ONA OS OS ONA OS OS ONA OS OS ONA OS OS OS ONA OS										
Note: This includes both the lancet and the lancet holding device. No	4. E.3 Finger stick devices are not used for more	C Yes	O 1	Yes	<u> </u>					
holding device. A. E.4 If used for more than one patient, the point-of-care device is cleaned and disinfected after every use according to manufacturer's instructions. Note: If manufacturer does not provide instructions for cleaning and disinfection, then the device should not be used for >1 A. E.5 Insulin pens are used for only one patient A. E.5 Insulin pens are used for only one patient A. E.5 Insulin pens are used for only one patient A. E.5 Insulin pens are used for only one patient A. E.5 Insulin pens are used for only one patient A. E.5 Insulin pens are used for only one patient A. E.5 Insulin pens are used for only one patient A. E.5 Insulin pens are used for only one patient A. E.5 Insulin pens are used for only one patient A. E.5 Insulin pens are used for only one patient A. E.5 Insulin pens are used for only one patient A. E.5 Insulin pens are used for only one patient A. E.5 Insulin pens are used for only one patient A. E.5 Insulin pens are used for only one patient A. E.5 Insulin pens are used for only one patient A. E.5 Insulin pens are used for only one patient A. E.5 Insulin pens are used for only one patient A. E.6 Insulin pens are used for only one patient A. E.7 Insulin pens are used for only one patient A. E.6 Insulin pens are used for only one patient A. E.7 Insulin pens are used for only one patient A. E.7 Insulin pens are used for only one patient A. E.7 Insulin pens are used for only one patient A. E.7 Insulin pens are used for only one patient	than one patient.		=		□ 2					
A. E.4 If used for more than one patient, the point- of-care device is cleaned and disinfected after every use according to manufacturer's instructions. Note: if manufacturer does not provide instructions for cleaning and disinfection, then the device should not be used for >1 patient. A. E.5 Insulin pens are used for only one patient Yes No Yes No No No No No No No No No N	Note: This includes both the lancet and the lancet	O No	-	🗀 No	□ 3					
4. E.4 If used for more than one patient, the point- of-care device is cleaned and disinfected after every use according to manufacturer's instructions. Note: if manufacturer does not provide instructions for cleaning and disinfection, then the device should not be used for >1 patient. 4. E.5 Insulin pens are used for only one patient Yes No Yes No No No Yes No No No No No No No No No N	holding device.									
of-care device is cleaned and disinfected after every use according to manufacturer's instructions. Note: if manufacturer does not provide instructions for cleaning and disinfection, then the device should not be used for >1 patient. 4. E.5 Insulin pens are used for only one patient 9 Yes 1 0 2 No N/A 1 0 5 N/A 2 0 3 N/A 3 0 4 N/A 5 0 5		[□] N/A	<u> </u>	□ N/A	<u></u> 5					
of-care device is cleaned and disinfected after every use according to manufacturer's instructions. Note: if manufacturer does not provide instructions for cleaning and disinfection, then the device should not be used for >1 patient. 4. E.5 Insulin pens are used for only one patient 9 Yes 1 0 2 No N/A 1 1 0 3 N/A 1 5 5 N/A 2 1 0 2 NO N/A 3 0 4 N/A 4 0 5										
every use according to manufacturer's instructions. Note: if manufacturer does not provide instructions for cleaning and disinfection, then the device should not be used for >1 patient. 4. E.5 Insulin pens are used for only one patient 9 Yes 1	• • • •	Yes		Yes						
instructions. Note: if manufacturer does not provide instructions for cleaning and disinfection, then the device should not be used for >1 patient. 4. E.5 Insulin pens are used for only one patient O Yes O NO O N	of-care device is cleaned and disinfected after									
Note: if manufacturer does not provide instructions for cleaning and disinfection, then the device should not be used for >1 patient. 4. E.5 Insulin pens are used for only one patient O Yes O NO NO NO NO NO NO NO NO NO	every use according to manufacturer's	□ No	=	□ No						
instructions for cleaning and disinfection, then the device should not be used for >1 patient. 4. E.5 Insulin pens are used for only one patient O Yes O NO O N		6		_						
then the device should not be used for >1 patient. 4. E.5 Insulin pens are used for only one patient O Yes O No	·	□ N/A	□ 5	□ N/A	○ 5					
patient. □ Yes □ 1 □ Yes □ 1 □ 2 □ No □ 3 □ No □ 3 □ 4 □ 4 □ N/A □ N/A □ N/A □ N/A □ N/A □ 5 □ N/A □ N/A □ 5 □ N/A □										
4. E.5 Insulin pens are used for only one patient O Yes O NO O										
□ No □ 2 □ 3 □ 4 □ 5 □ No □ 4 □ 5 □ No □ 5 □ 5 □ No □ No □ No □ No □	patient.									
□ No □ 2 □ 3 □ 4 □ 5 □ No □ 4 □ 5 □ No □ 5 □ 10 □ No □ 10 □ 10 □ No □ No □ 10 □ No □ 10 □ No □ No □ 10 □ No □ 10 □ No □ No □ No □ 10 □ No □ 10 □ No □ N	4.551 13	<i>₽</i>		2 v						
□ No □ 3 □ 4 □ 4 □ 5 □ N/A □ 5 □ No □ 3 □ 4 □ 5 □ N/A □ 5	4. E.5 Insulin pens are used for only one patient	Yes		₩ Yes						
$\bigcirc N/A$ $\bigcirc 4$ $\bigcirc 5$ $\bigcirc N/A$ $\bigcirc 5$		€ No.		n Na						
□ N/A □ 5 □ N/A □ 5		I ₩ INO		₩ NO						
		€N/A		P N/A						
If no to any of the above (4 F 1 through 4 F 5) cite at 42 CFR 482 42(a)(1) (Tag Δ-0749)		IN/A	<u> </u>	₩ IN/A	<u> </u>					
	If no to any of the above (4.E.1 through 4.E.5), cite	at 42 CFR 482.42(l							

Section 4. F Isolation: Contact Precautions

Elements to be assessed		N	Nanner of Assessment Code			Manner of Assessment Code		
			all that apply) & Surveyor Notes			k all that apply) & Surveyor Notes		
Patients requiring contact isolation are identified and managed in a manner consistent with hospital infection control policies and procedures to maximize the prevention of								
infection and communicable disease including the following:								
4. F.1 Gloves and gowns are available and located	Yes	◯ 1		Yes	□ 1			
near point of use.		2			2			
	○ No	◯ 3		○ No	<u> </u>			
		4			4			
	□ N/A	□ 5		◯ N/A	□ 5			
4. F.2 Signs indicating patient is on Contact	☐ Yes	O 1		☐ Yes	O 1			
Precautions are clear and visible.		□ 2			2			
	○ No	<u> </u>		No No	□ 3			
		4			4			
	◯ N/A	□ 5		[©] N/A	□ 5			
4. F.3 Patients on contact precautions are housed	☐ Yes	<u> </u>		☐ Yes	O 1			
in single-patient rooms when available or	_	<u> </u>			2			
cohorted based on a clinical risk assessment.	□ No	<u> 3</u>		□ No	<u> </u>			
	A .	6 4		A .	Q 4			
	□ N/A	6 5		[□] N/A	□ 5			
4. F.4 Hand hygiene is performed before entering	C Yes			C Yes				
patient care environment.	O No			No.				
Note: Coop and water must be used when here	₩ INO			U NO				
	∩ NI/A			€ NI/A				
	□ N/A	₩ 5		□ N/A	<u> </u>			
known or suspected C. difficile or norovirus								
during an outbreak. In all other situations,								
ABHR is preferred.								
4. F.5 Gloves and gowns are donned before	Yes	O 1		Yes	O 1			
entering patient care environment.		2						
	□ No			□ No				
	□ N/A	□ 5		□ N/A	□ 5			
patient care environment. Note: Soap and water must be used when bare hands are visibly soiled (e.g., blood, body fluids) or after caring for a patient with known or suspected <i>C. difficile</i> or norovirus during an outbreak. In all other situations, ABHR is preferred.	○ No ○ N/A	2 3 4 5		□ No □ N/A	① 2 ② 3 ② 4 ② 5			

4. F.6 Gloves and gowns are removed and discarded, and hand hygiene is performed before leaving the patient care environment. 4. F.7 Dedicated or dispensable pensitival patient.	☐ Yes ☐ No ☐ N/A ☐ Yes	1 2 3 4 0 5	☐ Yes ☐ No ☐ N/A ☐ Yes	1 2 3 4 5 5	
4. F.7 Dedicated or disposable noncritical patient-care equipment (e.g., blood pressure cuffs) is used or if not available, then equipment is cleaned and disinfected prior to use on another patient according to manufacturer's instructions.	☐ No ☐ N/A	2	□ No □ N/A	2 2 3 0 4 0 5	
F.8 Facility limits movement of patients on Contact Precautions outside of their room to medically necessary purposes.	☐ Yes ☐ No ☐ N/A	1 2 3 4 5	☐ Yes ☐ No ☐ N/A	1	
4. F.9 If a patient on Contact Precautions must leave their room for medically necessary purposes, there are methods followed to communicate that patient's status and to prevent transmission of infectious disease.	☐ Yes ☐ No ☐ N/A	1 2 2 3 4 C 5	☐ Yes ☐ No ☐ N/A	☐ 1 ☐ 2 ☐ 3 ☐ 4 ☐ 5	
4. F.10 Objects and environmental surfaces in patient care areas that are touched frequently (e.g., bed rails, side table, call button) are cleaned and then disinfected when visibly soiled and at least daily with an EPA-registered disinfectant.	☐ Yes ☐ No ☐ N/A	① 1 ② 2 ③ 3 ③ 4 ③ 5	☐ Yes ☐ No ☐ N/A	① 1 ② 2 ② 3 ② 4 ② 5	
4. F.11 For terminal cleaning (i.e., after patient discharge), all surfaces are thoroughly cleaned and disinfected and all textiles are replaced with clean textiles.	☐ Yes ☐ No ☐ N/A	1 2 3 4 5	☐ Yes ☐ No ☐ N/A	1 2 2 3 4 D 5	

4. F.12 Cleaners and disinfectants are labeled and used in accordance with hospital policies and procedures and manufacturer's instructions (e.g., dilution, storage, shelf-life, contact time).	☐ Yes ☐ No ☐ N/A	① 1 ② 2 ② 3 ② 4 ② 5	☐ Yes ☐ No ☐ N/A	① 1 ① 2 ① 3 ① 4 ① 5		
If no to any of the above (4.F.1 through 4.F.12), cite at 42 CFR 482.42(a)(1) (Tag A-0749)						

Section 4. G Isolation: Droplet Precautions Manner of Assessment Code Manner of Assessment Code Elements to be assessed (check all that apply) & Surveyor Notes (check all that apply) & Surveyor Notes Patients requiring Droplet Precautions are identified and managed in a manner consistent with hospital infection control policies and procedures to maximize the prevention of infection and communicable disease including the following: (i) 1 Tes 4. G.1 Surgical masks are available and located \square 1 <u>2</u> 2 near point of use. ☐ No \Box 3 No. 3 □ 4 4 O N/A <u> 5</u> □N/A 5 4. G.2 Signs indicating patient is on Droplet Yes (C) 1 Yes **1** 2 2 Precautions are clear and visible. **(**3 □ No □ 3 □ No. **6** 4 4 5 O N/A □ N/A \Box 5 O 1 4. G.3 Patients on Droplet Precautions are housed Yes Yes 1 6 <u>2</u> in single-patient rooms when available or 2 3 O No No. cohorted based on a clinical risk assessment. 4 4 □ N/A [□] 5 □ N/A 1 1 4. G.4 Hand hygiene is performed before entering Yes Yes 6 2 6 3 2 patient care environment. 3 O No O No 6 4 4 [□] 5 □ N/A □ N/A 5

4. G.5 HCP don surgical masks before entering the	C Yes	O 1	C Yes	O 1	
patient care environment or private room.		□ 2		□ 2	
	○ No	○ 3		□ 3	
		O 4		4	
	Ĉ N/A	□ 5	□ N/A	O 5	
4. G.6 Mask is removed and discarded, and hand	Yes	O 1	☐ Yes	0 1	
hygiene is performed upon leaving the		□ 2		O 2	
patient care environment.	○ No	□ 3	○ No	3	
	_	O 4		O 4	
	□ N/A	○ 5	□ N/A	O 5	
4. G.7 Facility limits movement of patients on	Yes	O 1	Yes	O 1	
Droplet Precautions outside of their room to	~	2		O 2	
medically necessary purposes (note: policy	No No	Q 3	◯ No	Q 3	
should address that patient wear surgical		Q 4		Q 4	
mask when transported).		<u>0</u> 5		<u>O</u> 5	
4. G.8 If a patient on Droplet Precautions must	Yes	O 1	C Yes	O 1	
leave their room for medically necessary	<u> </u>	<u>0</u> 2		O 2	
purposes, there are methods followed to	☐ No	<u>0</u> 3	Ĉ No	Q 3	
communicate that patient's status and to	A .	O 4		<u> </u>	
prevent transmission of infectious disease	Ĉ N/A	○ 5	□ N/A	O 5	
(note that patient should wear surgical mask					
when transported).	A	A			
4. G.9 Objects and environmental surfaces in	C Yes	O 1	C Yes	<u>0</u> 1	
patient care areas that are touched	<u> </u>	□ 2 □ 2 □ 3 □ 4 □ 4 □ 4 □ 7	<u> </u>	O 2	
frequently (e.g., bed rails, side table, call	O No		□ No	<u>3</u>	
button) are cleaned and then disinfected	□ N/A	□ 4□ 5	© N/A	□ 4 □ 5	
when visibly soiled and at least once a day	□ N/A	□ 3	□ N/A	∪ 3	
with an EPA-registered disinfectant.					
4. G.10 During terminal cleaning (i.e., after patient	☐ Yes	O 1	Yes	O 1	
discharge), all surfaces are thoroughly				O 2	
cleaned and disinfected and all textiles are	○ No		□ No	3	
		1 4		O 4	
replaced with clean textiles.	□ N/A	6 5	□ N/A	O 5	
4. G.11 Cleaners and disinfectants are labeled and	C Yes	O 1	Yes	Q 1	
used in accordance with hospital policies and		2		O 2	
procedures and manufacturer's instructions	○ No	2 3	□ No	3	
(e.g., dilution, storage, shelf-life, contact		O 4		A	
time).	◯ N/A	5 5	□ N/A	6 5	
If no to any of the above (4.G.1 through 4.G.11), cit	e at 42 CFR 482.42	(a)(1) (Tag A-0749)			

Section 4. H Isolation: Airborne Precautions

Elements to be assessed		Manner of Assessment Code (check all that apply) & Surveyor Notes		Manner of Assessment Code (check all that apply) & Surveyor Notes
Patients requiring Airborne Precautions are identifie	•	a manner consistent with hospital infection	control policies ar	nd procedures to maximize the prevention
of infection and communicable disease including th	e following:			
4. H.1 NIOSH-approved particulate respirators (N-95 or higher) are available and located near point of use.	☐ Yes ☐ No ☐ N/A	① 1 ② 2 ② 3 ② 4 ② 5	☐ Yes ☐ No ☐ N/A	 □ 1 □ 2 □ 3 □ 4 □ 5
H.2 Signs indicating patient is on Airborne Precautions are clear and visible.	☐ Yes ☐ No ☐ N/A	☐ 1 ☐ 2 ☐ 3 ☐ 4 ☐ 5	☐ Yes ☐ No ☐ N/A	☐ 1 ☐ 2 ☐ 3 ☐ 4 ☐ 5
4. H.3 Patients on Airborne Precautions are housed in airborne infection isolation rooms (AIIR).	☐ Yes ☐ No ☐ N/A	☐ 1 ☐ 2 ☐ 3 ☐ 4 ☐ 5	☐ Yes ☐ No ☐ N/A	1
H.4 Hand hygiene is performed before entering patient care environment.	☐ Yes ☐ No ☐ N/A	□ 1□ 2□ 3□ 4□ 5	☐ Yes ☐ No ☐ N/A	□ 1□ 2□ 3□ 4□ 5
4. H.5 HCP wear a NIOSH-approved particulate respirator (N95 or higher) upon entry into the AIIR for patients with confirmed or suspected TB. Facility policies are followed for other pathogens requiring AIIR.	☐ Yes ☐ No ☐ N/A	☐ 1 ☐ 2 ☐ 3 ☐ 4 ☐ 5	☐ Yes ☐ No ☐ N/A	☐ 1 ☐ 2 ☐ 3 ☐ 4 ☐ 5

4. H.6 Facility limits movement of patients on Airborne Precautions outside of their room to medically-necessary purposes (note: policy should address that patient wear surgical mask).	☐ Yes ☐ No ☐ N/A	1 2 3 1 4 1 5		☐ Yes ☐ No ☐ N/A	1 2 3 4 6 5	
4. H.7 If a patient on Airborne Precautions must leave their room for medically necessary purposes, there are methods followed to communicate that patient's status and to prevent transmission of infectious disease (note: policy should address that patient wear surgical mask when transported).	☐ Yes ☐ No ☐ N/A	① 2 ② 3 ② 4 ⑥ 5		☐ Yes ☐ No ☐ N/A	1 2 2 3 4 6 5	
If no to any of the above (4.H.1 through 4.H.7), cite	at 42 CFR 482.42	(a)(1) (Tag /	A-0749)			

Section 4. I Surgical Procedure Tracer Manner of Assessment Code Manner of Assessment Code Elements to be assessed (check all that apply) & Surveyor Notes (check all that apply) & Surveyor Notes Surgical procedures are performed in a manner consistent with hospital infection control policies and procedures to maximize the prevention of infection and communicable disease including the following: Yes 4. I.1 Healthcare personnel perform a surgical Yes ⁰ 1 0 1 2 □ 2 scrub before donning sterile gloves for □ 3 3 ♠ No. O No surgical procedures (in OR) using either an 4 4 antimicrobial surgical scrub or an FDA-□ N/A □ N/A \bigcirc 5 5 approved alcohol-based antiseptic surgical hand rub. Note: If hands are visibly soiled, they should be prewashed with soap and water before using an alcohol-based surgical scrub. O 1 4. I.2 After surgical scrub, hands and arms are Yes Yes 1 2 2 dried with a sterile towel (if applicable), and □ 3 O No 3 O No sterile surgical gown and gloves are donned 4 4 in the OR. [□]N/A O N/A 5 □ 5

4. I.3 Surgical attire (e.g., scrubs) and surgical caps/hoods covering all head and facial hair are worn by all personnel in semi restricted and restricted areas.	☐ Yes ☐ No	① 1 ② 2 ② 3 ③ 4 ⑥ 5	☐ Yes ☐ No	1 2 2 3 4 4 6 9 9 9 9 9 9 9 9 9 9 9 9 9 9 9 9 9				
Note: Restricted area includes ORs, procedure rooms, and the clean core area. The semi restricted area includes the peripheral support areas of the surgical suite.	[©] N/A	5	□ N/A	5				
4. I.4 Surgical masks are worn (and properly tied, fully covering mouth and nose) by all personnel in restricted areas where open sterile supplies or scrubbed persons are located.	☐ Yes ☐ No ☐ N/A	1	☐ Yes ☐ No ☐ N/A	1				
4. I.5 Sterile drapes are used to establish sterile field.	☐ Yes ☐ No ☐ N/A	○ 4	☐ Yes ☐ No ☐ N/A	1 2 2 3 4 Q 5				
 4. I.6 Sterile field is maintained and monitored constantly. Ensure that: Items used within sterile field are sterile. Items introduced into sterile field are opened, dispensed, and transferred in a manner to maintain sterility. Sterile field is prepared in the location where it will be used and as close as possible to time of use. Movement in or around sterile field is done in a manner to maintain sterility. 	☐ Yes ☐ No ☐ N/A	1 2 3 4 1 5	☐ Yes ☐ No ☐ N/A	1 2 3 0 4 0 5 5				
4. I.7 Traffic in and out of OR is kept to minimum and limited to essential staff.	☐ Yes ☐ No ☐ N/A	□ 1□ 2□ 3□ 4□ 5	☐ Yes ☐ No ☐ N/A	① 1 ② 2 ② 3 ⑤ 4 ⑤ 5				
4. I.8 Surgical masks are removed when leaving the sterile areas and are not reused when returning.	☐ Yes ☐ No ☐ N/A	① 1 ② 2 ② 3 ② 4 ② 5	☐ Yes ☐ No ☐ N/A	1 2 2 3 4 C 5				
If no to any of the above (4.I.1 through 4.I.8), cite at 42 CFR 482.42(a)(1) (Tag A-0749)								

Processes ensuring infection control in the OR are accomplished in a	manner consister	it with ho	spital infection control policies and procedures to maximize the prevention of
infection and communicable disease including the following:			
4. I.10 Cleaners and EPA-registered hospital disinfectants are used	Yes	O 1	
in accordance with hospital policies and procedures and		2	
manufacturer's instructions (e.g., dilution, storage, shelf-life,	O No	□ 3	
contact time).		4	
	□ N/A	□ 5	
4. I.11 Cleaners and EPA-registered disinfectants, when in use, are	C Yes	O 1	
labeled, diluted according to manufacturer's instructions, and		2	
are dated.	O No	3	
		4	
	☐ N/A	O 5	
4. I.12 All horizontal surfaces (e.g., furniture, surgical lights, booms,	T Yes	O 1	
equipment) are damp dusted before the first procedure of the		2	
day using a clean, lint-free cloth and EPA-registered hospital	🗖 No	3	
detergent/disinfectant.		4	
	© N/A	O 5	
4. I.13 High touch environmental surfaces are cleaned and	🗀 Yes	O 1	
disinfected between patients.		O 2	
	O No	O 3	
	6	O 4	
	[©] N/A	O 5	
4. I.14 Anesthesia equipment is cleaned and disinfected between	Yes	O 1	
patients.		O 2	
	□ No	Q 3	
	<u>~</u> .	O 4	
	Ĉ N/A	O 5	
4. I.15 Reusable noncritical items (e.g., blood pressure cuffs, ECG	C Yes	0 1	
leads, tourniquets, oximeter probes) are cleaned and	O .:	□ 2 □ 2 □ 2 □ 3 □ 3 □ 4 □ 4 □ 5 □ 6 □ 7	
disinfected between patients.	☐ No	<u>3</u>	
	€ N/A	Q 4	
	□ N/A	□ 5	
4 L1C ODs are townsinally closured after last was adding of the day	Yes	0 1	
4. I.16 ORs are terminally cleaned after last procedure of the day	U res	\bigcirc 2	
(including weekends) and each 24-hour period during regular	□ No	□ 2 □ 3	
work week. Terminal cleaning includes wet-vacuuming or	₩ INU	① 4	
mopping floor with an EPA-registered disinfectant.	□ N/A	<u> </u>	
	- IV/A	_ 5	
	1	1	

ceilings have cleanable surfaces, are visibly clean, and there is evidence that all surfaces are cleaned regularly in accordance	□ No	23	
with hospital policies and procedures.		O 4	
	©n/A	□ 5	
4. I.18 Internal components of anesthesia machine breathing circuit	☐ Yes	O 1	
are cleaned regularly according to manufacturer's instructions.	163	<u> </u>	
are steamed regularly according to manaracturer of motifications.	◯ No	□ 3	
		□ 4	
	[©] N/A	□ 5	
	O	A .	
4. I.19 Ventilation requirements meet the following:	C Yes	12	
Positive pressure, 15 air exchanges per hour (at least 3 of	□ No	□ 2□ 3	
which are fresh air)			
 90% filtration (HEPA is optional), air filters checked regularly and replaced according to hospital policies and procedures 	© N/A	<u></u>	
Temperature and relative humidity levels are	,		
maintained at required levels			
Doors are self-closing			
Air vents and grill work are clean and dry.			
If no to any of the above (4.I.10 through 4.I.19), cite at 42 CFR 482.42	2(a)(1) (Tag A-074	9)	

Module 5: Special Care Environments

Section 5. A Protective Environment (e.g. Bone Marrow patients) Manner of Assessment Code Manner of Assessment Code Elements to be assessed (check all that apply) & Surveyor Notes (check all that apply) & Surveyor Notes For patients requiring a Protective Environment - the hospital ensures: 5. A.1 Positive pressure [air flows out to the Yes Yes ① 1 2 2 corridor]. 3 □ 3 O No O No 4 4 5 □ N/A □ 5 N/A

5. A.2 Twelve (12) air changes per hour.	Yes	O 1		Yes	O 1	
		□ 2		_	<u> </u>	
	□ No	□ 3		☐ No	☐ 3	
	6	<u> </u>			<u> </u>	
	□ N/A	□ 5		🗖 N/A	<u> </u>	
	O			₽		
5. A.3 Supply air is HEPA filtered.	☐ Yes	12		C Yes	0 1	
	□ No	□ 2 □ 3		☐ No	□ 2 □ 2 □ 3	
	U NO	\bigcirc 3		□ NO	34	
	Ĉ N/A	<u>0</u> 5		Ĉ N/A	□ 4□ 5	
	□ IV/A	□ 3		□ N/A	□ 3	
5. A.4 Well sealed rooms so that there are no	☐ Yes	O 1		Yes	O 1	
penetration spaces in walls, ceilings, or		□ 2			2	
windows.	☐ No	☐ 3		Ĉ No	□ 3	
		4			4	
	Ĉ N/A	<u> </u>		☐ N/A	<u> </u>	
	-	_			_	
5. A.5 Self closing door that fully closes on all	C Yes	0 1		C Yes		
room exits.	2 v	□ 2□ 3		A	2	
	◯ No	☐ 3 ☐ 4		□ No	34	
	🗖 N/A	O 5		Ĉ N/A	<u>4</u> 5	
	₩/A			□ N/A	<u></u>)	
5. A.6 Documents and demonstrates that failures	☐ Yes	(i) 1		☐ Yes	O 1	
are addressed.		<u> </u>			2	
	☐ No	<u> 3</u>		No No		
		□ 4			4	
	Ĉ N/A	□ 5		🗖 N/A	C 5	
If no to any of the above (5.A.1 through 5.A.6) cite			g A-0749)	-	-	
5. A.7 For patients requiring a Protective	Yes	<u> </u>		Yes	<u> </u>	
Environment, the hospital ensures that	20	<u>2</u> 2			<u>2</u> 2	
ventilation specifications are monitored using	O No	<u>3</u>		O No	3	
visual methods (e.g. flutter strips, smoke	□ N/A	Q 4		♠ N/A	① 4	
tubes) and observations documented daily.	🗖 N/A	□ 5		O N/A	፟ 5	
If no, cite at 42 CFR 482.42(b)(2) (Tag A-0756)	<u> </u>					
,						

Interview = 1 Observation = 2

PRE-DECISIONAL SURVEYOR WORKSHEET

Assessing Hospital Compliance with the

Condition of Participation for Discharge Planning

Pilot Program Draft Version

Instructions: The following is a list of items that must be assessed during the on-site survey, in order to determine compliance with the Discharge

Name of State Agency: _

Planning Condition of Participation. Items are to be assessed by a combination of observation, review of the hospital's discharge planning program documentation, interviews with hospital staff, patients and their family/support persons, and review of medical records.						
suppoi	The interviews should be performed with the most appropriate staff person(s) for the items of interest, as well as with patients, family members, and rt persons.					
deficie	Citation instructions are provided throughout this instrument, indicating the applicable regulatory provision to be cited on Form CMS-2567 when ent practices are observed.					
Sectio	n 1 Hospital Characteristics					
1.	Hospital name:					
2.	Address, State, Zip Code:					
3.	CMS Certification Number (CCN):					

4.	Date of site visit:
	/
5.	Number of State Agency surveyors who participated in this survey:
6.	Approximate time spent on site performing this survey (hours):
7.	Does the hospital participate in Medicare via accredited "deemed" status?
	a. If YES, which Accrediting Organization(s)?
	i. O American Osteopathic Association (AOA)/Healthcare Facilities Accreditation Program (HFAP)
	ii. O Det Norske Veritas Healthcare (DNV)
	iii. O The Joint Commission (TJC)
	b. If YES, according to the hospital, what was the end date of the most recent accreditation survey:
8.	What was the end date of the most recent previous State Agency Federal survey:

Section 2 Discharge Planning – Policies and Procedures						
Elements to be assessed		Manner of Assessment Code (list all that apply) & Surveyor Notes				
2.1 Are discharge planning policies and procedures in effect for	all inpatient	. ,,,,,				
Specifically:						
2.1a For every inpatient unit surveyed is there evidence of	O Yes	0 1				
applicable discharge planning activities?	O No	O 2				
		O 3				
		0 4				
		O 5				
2.1b Are staff members responsible for discharge planning	O Yes	0 1				
activities correctly following the hospital's discharge	O No	O 2				
planning policies and procedures?		O 3				
		O 4				
		O 5				
NOTE: If no for either 2.1a or 2.1b cite the applicable standard	for identific	cation of patients needing discharge planning, 42 CFR 482.43(a) (Tag A-0800);				
		ng and implementing the discharge plan, 42 CFR 482.43(c) (Tag A-0817)				
2.2 Does the discharge planning process apply to certain	O Yes	0 1				
categories of outpatients?	O No	O 2				
		O 3				
		O 4				
		O 5				
If yes, check all that apply:						
O Observation patients who are not subsequently admi	O Same day surgery patients					
O be patients who are not subsequently admitted O ED patients who are not subsequently admitted						
O Other						
_	skip to	0 1				
	estion 2.8	0 2				
O No,		O 3				
·	estion 2.4	O 4				
		O 5				
NOTE: No citation is made related to questions 2.2 and 2.3						

Elements to be assessed			Manner of Assessment Code (list all that apply) & Surveyor Notes			
2.4 For patients	s not initially identified as in need of a discharge p	olan, is ther	re a process for updating this determination based on changes in the patient's			
condition or	circumstances? Specifically,					
2.4a Does t	he discharge planning policy address changes in	O Yes	O 1			
patien	t condition that would call for the development	O No	O 2			
of a di	scharge plan in patients not previously		O 3			
identif	fied as in need of one?		O 4			
			O 5			
2.4b Are in	patient unit staff aware of how, when, and	O Yes	O 1			
whom	to notify of such changes in patient condition?	O No	O 2			
			O 3			
			O 4			
			O 5			
NOTE: If no to e	either 2.4a or 2.4b, cite 42 CFR 482.43(a) (Tag A-	0800)				
2.5 Is there a pr	rocess for patients, or their representatives, and I	physicians t	to request a discharge planning evaluation? Specifically,			
2.5a Does t	he hospital have a standard process for	O Yes	O 1			
notifyi	ing patients (or their representative if	O No	O 2			
applica	able) and physicians that they may request a		O 3			
discha	rge planning evaluation and that the hospital		O 4			
will co	nduct an evaluation upon request?		O 5			
2.5b Can bo	oth discharge planning and unit nursing staff	O Yes	O 1			
persor	nnel describe the process for a patient or the	O No	O 2			
patien	t's representative to request a discharge		O 3			
planni	ng evaluation?		O 4			
			O 5			
	ew patients (or their representatives if	O Yes	0 1			
	able). If they say they were not aware they	O No	O 2			
	request a discharge planning evaluation, can		O 3			
	spital provide evidence the patient or		O 4			
•	sentative received notice they could request an		O 5			
evalua	ition?					

Elements to be assessed			Manner of Assessment Code (list all that apply) & Surveyor Notes
2.5d Interview attending physicians. If they are not aware	O Yes	0 1	
they can request a discharge planning evaluation, can	O No	O 2	
the hospital provide evidence of how it informs the		O 3	
medical staff about this?		O 4	
		O 5	
NOTE: If no to any part of question 2.5, cite 42 CFR 482.43(b)(1)		•	
2.6 Interview attending physicians. If they are not aware they	O Yes	0 1	
can request a discharge plan regardless of the outcome of	O No	O 2	
the completed evaluation, can the hospital provide		O 3	
evidence of how it informs the medical staff about this?		0 4	
		O 5	
NOTE: If yo to 2.6 site 42 CER 492 42(a)(2) /Tog 4 0910)			
NOTE: If no to 2.6, cite 42 CFR 482.43(c)(2) (Tag A-0819) 2.7 Can discharge planning personnel describe a process for	O Yes	0 1	
physicians to order a discharge plan to be completed on a	O No	0 2	
patient, regardless of the outcome of the patient's	O NO	0 3	
evaluation?		0 4	
Evaluation:		0 5	
NOTE: If no to 2.7, cite 42 CFR 482.43(c)(2) (Tag A-0819)		<u> </u>	
2.8 Does the hospital discharge planning policy include a	O Yes	0 1	
process for ongoing reassessment of the discharge plan	O No	O 2	
based on changes in patient condition, changes in available		O 3	
support, and/or changes in post-hospital care		O 4	
requirements?		O 5	
NOTE: If no to 2.8, cite 42 CFR 482.43(c)(4) (Tag A-0821)			

Section 3 Discharge Planning – Reassessment and QAPI		
Section 5 Discharge Flamming Reassessment and QAFF		
Elements to be assessed		Manner of Assessment Code (list all that apply) & Surveyor Notes
3.1 Does the hospital review the discharge planning process in	O Yes	0 1
an ongoing manner?	O No	O 2
		O 3
		O 4
		O 5
3.2 Does the hospital track its readmissions as part of its review	O Yes	0 1
of the discharge planning process?	O No	O 2
		O 3
		O 4
		O 5
3.2a Does the assessment of readmissions include an	O Yes	0 1
evaluation of whether the readmissions were	O No	O 2
potentially preventable?		O 3
		O 4
		O 5
3.3 If the hospital identified preventable readmissions where	O Yes	O 1
problems in the discharge planning process were identified	O No	0 2
as a possible cause, did it make changes to its discharge	O N/A	0 3
planning process to address the problems?	.,,,	0 4
process to data see the process.		O 5
NOTE: If no to any one of 3.1 through 3.3, cite 42 CFR 482.43(e)	(Tag A-084	43). Consider citing QAPI 42 CFR 482.21(c) (Tag A-0283)
3.4 Does the hospital have a process for collecting and	O Yes	O 1
considering feedback from post-acute providers in the	O No	O 2
community about the effectiveness of the hospital's		O 3
discharge planning process?		O 4
		O 5
No citation is made related to this question		

Section 4 Discharge Planning Tracers

In this section, survey 1-2 current inpatients and review the closed medical records of 2-3 discharged patients. When possible, include one inpatient who was readmitted within 30 days of a previous admission. For closed records, be sure to select a record that includes a discharge planning evaluation and a discharge plan, and do not choose N/A instead of a Yes or No response. Note key at bottom of page for Manner of Assessment code.

DCP = Discharge Planning

	Patient/Record #1	Patient/Record #2	Patient/Record #3	Patient/Record #4
	O Open	O Open	O Open	O Open
	O Closed	O Closed	O Closed	O Closed
4.1 When was the screening done to identify	a.O	a. O	a.O	a. O
whether the inpatient needed a discharge	b.O	b.O	b.O	b.O
planning evaluation?	c. O	c. O	c. O	c. O
	d.O	d.O	d.O	d.O
a. Before or at time of admission				
b. After admission but at least 48 hours	0 1	0 1	0 1	0 1
prior to discharge	O 2	O 2	O 2	O 2
c. N/A – all admitted patients receive a	O 3	O 3	O 3	O 3
discharge plan	O 4	O 4	O 4	O 4
d. None of the above	O 5	O 5	O 5	O 5
NOTE: If response 4.1d is selected, cite 42 CFR	482.43(a) (Tag A-0800)			
4.2 Can hospital staff demonstrate that the	O Yes	O Yes	O Yes	O Yes
hospital's criteria and screening process	O No	O No	O No	O No
for a discharge planning evaluation were				
correctly applied?	O 1	0 1	0 1	0 1
	O 2	O 2	O 2	O 2
	O 3	O 3	O 3	O 3
	O 4	O 4	O 4	O 4
	O 5	O 5	O 5	O 5
NOTE: If no to 4.2, cite 42 CFR 482.43(a) (Tag	4-0800)			

	Patient/Record #1	Patient/Record #2	Patient/Record #3	Patient/Record #4
4.3 If the patient did not meet the hospital's	O Yes	O Yes	O Yes	O Yes
criteria for an evaluation, were the patient	O No	O No	O No	O No
(or patient's representative if applicable)	O N/A	O N/A	O N/A	O N/A
and the patient's physician made aware	O 1	0 1	0 1	0 1
they could still request a discharge	O 2	O 2	O 2	O 2
planning evaluation?	O 3	O 3	O 3	O 3
	O 4	0 4	0 4	O 4
	O 5	0 5	O 5	O 5
NOTE: If no to 4.3, cite 42 CFR 482.43(b)(1) (Ta	g A-0806)			
4.4 Was the discharge planning evaluation	O Yes	O Yes	O Yes	O Yes
and, as applicable, the discharge plan	O No	O No	O No	O No
developed by an RN, Social Worker, or	O N/A	O N/A	O N/A	O N/A
other qualified personnel, as defined in	0 1	O 1	O 1	O 1
the hospital discharge planning policies	O 2	O 2	O 2	O 2
and procedures, or someone they	O 3	O 3	O 3	O 3
supervise?	O 4	0 4	0 4	O 4
·	O 5	0 5	O 5	O 5
NOTE: If no to 4.4, cite 42 CFR 482.43(b)(2) (Ta	g A-0807 - evaluation) and/	or 42 CFR 482.43 (c)(1) (Tag	A-0818 - plan), as applicab	le

	Patient/Record #1	Patient/Record #2	Patient/Record #3	Patient/Record #4
4.5 Are the results of the discharge planning	O Yes	O Yes	O Yes	O Yes
evaluation documented in the medical	O No	O No	O No	O No
record?	O N/A	O N/A	O N/A	O N/A
	0 1	0 1	0 1	0 1
	O 2	O 2	O 2	O 2
	O 3	O 3	O 3	O 3
	O 4	O 4	O 4	O 4
	O 5	O 5	O 5	O 5
NOTE: If no to 4.5, cite 42 CFR 482.43(b)(6) (Ta				
4.6 Did the evaluation include an assessment	O Yes	O Yes	O Yes	O Yes
of the patient's post-discharge care needs	O No	O No	O No	O No
being met in the environment from which				
he/she entered the hospital?	0 1	0 1	0 1	0 1
	O 2	O 2	O 2	O 2
	O 3	O 3	O 3	O 3
	O 4	O 4	O 4	O 4
	O 5	O 5	O 5	O 5

	Patient/Record #1	Patient/Record #2	Patient/Record #3	Patient/Record #4
For patients admitted from home				
4.7 Did the evaluation include an assessment of the patient's ability to perform activities of daily living (e.g. personal hygiene and grooming, dressing and undressing, feeding, voluntary control over bowel and bladder, ambulation, etc.)?	O Yes O No O N/A O 1 O 2 O 3 O 4 O 5	O Yes O No O N/A O 1 O 2 O 3 O 4 O 5	O Yes O No O N/A O 1 O 2 O 3 O 4 O 5	O Yes O No O N/A O 1 O 2 O 3 O 4 O 5
4.8 Did the evaluation include an assessment of the patient's or family/support person's ability to provide self-care/care?	O Yes O No O N/A O 1 O 2 O 3 O 4 O 5	O Yes O No O N/A O 1 O 2 O 3 O 4 O 5	O Yes O No O N/A O 1 O 2 O 3 O 4 O 5	O Yes O No O N/A O 1 O 2 O 3 O 4 O 5

	Patient/Record #1	Patient/Record #2	Patient/Record #3	Patient/Record #4
4.9 Did the evaluation include an assessment	O Yes	O Yes	O Yes	O Yes
of whether the patient will require	O No	O No	O No	O No
specialized medical equipment or home	O N/A	O N/A	O N/A	O N/A
and physical environment modifications?	0 1	0 1	O 1	0 1
	O 2	O 2	O 2	O 2
	O 3	O 3	O 3	O 3
	O 4	O 4	O 4	O 4
	O 5	O 5	O 5	O 5
4.9a If yes, did the evaluation include an assessment of whether the equipment is available or if the modifications can	O Yes O No O N/A	O Yes O No O N/A	O Yes O No O N/A	O Yes O No O N/A
be made to safely discharge the	0 1	0 1	0 1	0 1
patient to that setting?	O 2	O 2	O 2	O 2
	O 3	O 3	O 3	O 3
	O 4	O 4	O 4	O 4
	O 5	O 5	O 5	O 5

	Patient/Record #1	Patient/Record #2	Patient/Record #3	Patient/Record #4
4.10 If the patient or family/support person is	O Yes	O Yes	O Yes	O Yes
unable to meet care needs or there are	O No	O No	O No	O No
additional care needs above their	O N/A	O N/A	O N/A	O N/A
capabilities, did the evaluation include an	0 1	0 1	O 1	0 1
assessment of available community-based	O 2	O 2	O 2	O 2
services to meet post-hospital needs?	O 3	O 3	O 3	O 3
	O 4	O 4	O 4	O 4
	O 5	O 5	O 5	O 5
NOTE: If any no answer to questions 4.6 – 4.10	, cite 42 CFR 482.43(b)(4) (1			
4.11 If applicable, did the hospital provide the	O Yes	O Yes	O Yes	O Yes
patient with lists of Medicare-participating	O No	O No	O No	O No
HHAs or SNFs that provide post-hospital	O N/A	O N/A	O N/A	O N/A
services that could meet the patient's	0 1	0 1	0 1	0 1
medical needs?	O 2	O 2	O 2	O 2
	O 3	O 3	O 3	O 3
	O 4	O 4	O 4	O 4
	O 5	O 5	O 5	O 5

	Patient/Record #1	Patient/Record #2	Patient/Record #3	Patient/Record #4
4.11a Were the lists geographically appropriate for the patient?	O Yes O No O N/A O 1 O 2 O 3 O 4 O 5	O Yes O No O N/A O 1 O 2 O 3 O 4 O 5	O Yes O No O N/A O 1 O 2 O 3 O 4 O 5	O Yes O No O N/A O 1 O 2 O 3 O 4 O 5
NOTE: If no to 4.11 or 4.11a, cite 42 CFR 482.43				
For patients admitted from a nursing home/ski			0.4	0.4
4.12 Did the evaluation assess whether the prior facility has the capability to provide necessary post-hospital services to the	O Yes O No O N/A			
patient (i.e. is the same, higher, or lower	0 1	0 1	0 1	0 1
level of care required and can those needs be met in that facility?)	O 2 O 3 O 4 O 5			

	Patient/Record #1	Patient/Record #2	Patient/Record #3	Patient/Record #4
For all patients				
4.13 Did the evaluation include an assessment of the patient's insurance coverage (if applicable) and how that coverage might or might not provide for necessary services post-hospitalization?	O Yes O No O N/A O 1 O 2 O 3 O 4 O 5	O Yes O No O N/A O 1 O 2 O 3 O 4 O 5	O Yes O No O N/A O 1 O 2 O 3 O 4 O 5	O Yes O No O N/A O 1 O 2 O 3 O 4 O 5
If no to 4.12 or 4.13 cite 42 CFR 482.43(b)(4) (T	ag A-0806)			
4.14 Was the discharge planning evaluation completed in a timely basis to allow for appropriate arrangements to be made for post-hospital care and to avoid delays in discharge? NOTE: If no to 4.14, cite 42 CFR 482.43(b)(5) (T	O Yes O No O N/A O 1 O 2 O 3 O 4 O 5	O Yes O No O N/A O 1 O 2 O 3 O 4 O 5	O Yes O No O N/A O 1 O 2 O 3 O 4 O 5	O Yes O No O N/A O 1 O 2 O 3 O 4 O 5
NOTE: 11 110 to 4.14, cite 42 CFK 482.43(D)(5) (1	ag H-ngini			

	Patient/Record #1	Patient/Record #2	Patient/Record #3	Patient/Record #4
4.15 Was the patient (or the patient's	O Yes	O Yes	O Yes	O Yes
representative, if applicable) involved in a	O No	O No	O No	O No
discussion of the evaluation results?	O N/A	O N/A	O N/A	O N/A
	0 1	0 1	0 1	0 1
	0 2	O 2	O 2	O 2
	0 3	O 3	0 3	O 3
	0 4	O 4	0 4	O 4
	0 5	O 5	O 5	O 5
NOTE: If no to 4.15, cite 42 CFR 482.43(b)(6) (T	ag A-0811). Consider citing		Rights (Tag A-0130)	
4.16 Did the discharge plan match the	O Yes	O Yes	O Yes	O Yes
identified needs as determined by the	O No	O No	O No	O No
evaluation?	O N/A	O N/A	O N/A	O N/A
	0 1	0 1	0 1	0 1
	O 2	O 2	O 2	O 2
	O 3	O 3	O 3	O 3
	O 4	O 4	O 4	O 4
	O 5	O 5	O 5	O 5
NOTE: If no to 4.16, cite 42 CFR 482.43(c)(1) (T	ag A-0817)			

	Patient/Record #1	Patient/Record #2	Patient/Record #3	Patient/Record #4
4.17 If any significant changes in the patient's	O Yes	O Yes	O Yes	O Yes
condition were noted in the medical	O No	O No	O No	O No
record that changed post-discharge needs,	O N/A	O N/A	O N/A	O N/A
was the discharge plan updated	O 1	O 1	O 1	O 1
accordingly?	O 2	O 2	O 2	O 2
	O 3	O 3	O 3	O 3
	O 4	O 4	O 4	O 4
	O 5	O 5	O 5	O 5
NOTE: If no to 4.17, cite 42 CFR 482.43(c)(4) (T	ag Δ-0821)			
4.18 For patients discharged to home, did the h		implementation of the discl	narge plan? Specifically, look	for evidence of the
following, if applicable, based on the dischar	•	implementation of the disci	iarge plant. Specifically, look	Tor evidence or the
4.18a Providing in-hospital training to	O Yes	O Yes	O Yes	O Yes
patient and family/support persons,	O No	O No	O No	O No
using recognized methods. (Examples	O N/A	O N/A	O N/A	O N/A
include teach-back or repeat-back,	0 1	0 1	0 1	0 1
simulation laboratories, etc. but these	0 2	0 2	0 2	0 2
specific methods are not required.)	0 3	0 3	0 3	0 3
specific methods are not required.	0 4	0 4	0 4	0 4
	0 5	0 5	0 5	0 5
	0 3	0 3	0 3	0 3

	Patient/Record #1	Patient/Record #2	Patient/Record #3	Patient/Record #4
4.18b Written discharge instructions that are legible and use non-technical language.	O Yes O No O N/A O 1 O 2 O 3 O 4 O 5	O Yes O No O N/A O 1 O 2 O 3 O 4 O 5	O Yes O No O N/A O 1 O 2 O 3 O 4 O 5	O Yes O No O N/A O 1 O 2 O 3 O 4 O 5
4.18c A list of all medications the patient should be taking after discharge, with clear indication of changes from the patient's pre-admission medications	O Yes O No O N/A O 1 O 2 O 3 O 4 O 5	O Yes O No O N/A O 1 O 2 O 3 O 4 O 5	O Yes O No O N/A O 1 O 2 O 3 O 4 O 5	O Yes O No O N/A O 1 O 2 O 3 O 4 O 5

	Patient/Record #1	Patient/Record #2	Patient/Record #3	Patient/Record #4
4.18d Evidence of education of patients and support persons on admission vs. discharge medications, highlighting changes.	O Yes O No O N/A O 1 O 2 O 3 O 4 O 5	O Yes O No O N/A O 1 O 2 O 3 O 4 O 5	O Yes O No O N/A O 1 O 2 O 3 O 4 O 5	O Yes O No O N/A O 1 O 2 O 3 O 4 O 5
4.18e Referrals to established/new primary care physician or health center.	O Yes O No O N/A O 1 O 2 O 3 O 4 O 5	O Yes O No O N/A O 1 O 2 O 3 O 4 O 5	O Yes O No O N/A O 1 O 2 O 3 O 4 O 5	O Yes O No O N/A O 1 O 2 O 3 O 4 O 5

	Patient/Record #1	Patient/Record #2	Patient/Record #3	Patient/Record #4
4.18f Referrals, if applicable, to	O Yes	O Yes	O Yes	O Yes
specialized ambulatory services, e.g.	O No	O No	O No	O No
PT, OT, HHA, hospice, mental health,	O N/A	O N/A	O N/A	O N/A
etc.	0 1	0 1	O 1	0 1
	O 2	O 2	O 2	O 2
	O 3	O 3	O 3	O 3
	0 4	O 4	O 4	O 4
	O 5	O 5	O 5	O 5
4.18g Referrals, if applicable, to	O Yes	O Yes	O Yes	O Yes
community-based resources other	O No	O No	O No	O No
than health services, e.g. Depts. of	O N/A	O N/A	O N/A	O N/A
Aging, elder services, transportation	0 1	0 1	0 1	0 1
services, etc.	0 2	0 2	0 2	0 2
	0 3	0 3	0 3	0 3
	0 4	0 4	0 4	0 4
	0 5	0 5	0 5	0 5

	Patient/Record #1	Patient/Record #2	Patient/Record #3	Patient/Record #4
4.18h Arranging essential durable medical	O Yes	O Yes	O Yes	O Yes
equipment, e.g. oxygen, wheel chair,	O No	O No	O No	O No
hospital bed, commode, etc., if	O N/A	O N/A	O N/A	O N/A
applicable.	0 1	0 1	0 1	0 1
	O 2	O 2	O 2	O 2
	O 3	O 3	O 3	O 3
	O 4	O 4	O 4	O 4
	O 5	O 5	O 5	O 5
4.40: Canding a second was disal	O V	O Var	O V	O Yes
4.18i Sending necessary medical	O Yes	O Yes	O Yes	
information to providers the patient	O No	O No	O No	O No
was referred to prior to the first post-	O N/A	O N/A	O N/A	O N/A
discharge appointment or within 7	0 1	0 1	0 1	0 1
days of discharge, whichever comes	O 2	O 2	O 2	O 2
first.	O 3	O 3	O 3	O 3
	O 4	O 4	O 4	O 4
	O 5	O 5	O 5	O 5
NOTE: If implementation of the discharge plan	was not initiated, cite 42 CI	R 482.43(c)(3) (Tag A-0820)		

	Patient/Record #1	Patient/Record #2	Patient/Record #3	Patient/Record #4	
4.19 For patients transferred to another	O Yes	O Yes	O Yes	O Yes	
inpatient facility, was necessary medical	O No	O No	O No	O No	
information ready at time of transfer and	O N/A	O N/A	O N/A	O N/A	
sent to the receiving facility with the	0 1	0 1	O 1	0 1	
patient?	O 2	O 2	O 2	O 2	
	O 3	O 3	O 3	O 3	
	O 4	O 4	O 4	O 4	
	O 5	O 5	O 5	O 5	
NOTE: If no to 4.19, cite 42 CFR 482.43(d) (Tag	A-0837)		<u> </u>	1	
4.20 Were there portions of the plan the	O Yes	O Yes	O Yes	O Yes	
hospital failed to begin implementing,	O No	O No	O No	O No	
resulting in delays in discharge?	O N/A	O N/A	O N/A	O N/A	
	0 1	0 1	0 1	0 1	
	0 2	0 2	0 2	0 2	
	0 3	0 3	0 3	0 3	
	0 4	0 4	0 4	0 4	
	0 5	0 5	0 5	0 5	
NOTE: If we to 4.20 site 42 CED 402 42()(2)	T 4 0020\				
NOTE: IT yes to 4.20, cite 42 CFR 482.43(c)(3)	NOTE: If yes to 4.20, cite 42 CFR 482.43(c)(3) (Tag A-0820)				

	Patient/Record #1	Patient/Record #2	Patient/Record #3	Patient/Record #4	
4.21 For information only, were any of the follo	4.21 For information only, were any of the following services initiated while the patient was hospitalized:				
 a. Scheduling follow-up appointments b. Filling prescriptions c. Pharmacist meeting with patient and/or family/support persons to review medication regimen d. Pharmacist reviewing discharge medication orders prior to hospital departure 	a. O b. O c. O d. O e. O f. O g. O	a. O b. O c. O d. O e. O f. O g. O	a. O b. O c. O d. O e. O f. O g. O	a. O b. O c. O d. O e. O f. O g. O	
 e. Home setting visitation by hospital staff f. Transportation arranged for follow-up appointments g. Discharge planning checklists, e.g. CMS, AHRQ, CAPS checklists 	O 1 O 2 O 3 O 4 O 5	O 1 O 2 O 3 O 4 O 5	O 1 O 2 O 3 O 4 O 5	O 1 O 2 O 3 O 4 O 5	
NOTE: Do not cite; these are not required und		_		_	
4.22 Is there documentation in the medical record of providing the results of tests, pending at time of discharge, to the patient and/or post-hospital provider of care, if applicable?	O Yes O No O N/A O 1 O 2 O 3 O 4 O 5	O Yes O No O N/A O 1 O 2 O 3 O 4 O 5	O Yes O No O N/A O 1 O 2 O 3 O 4 O 5	O Yes O No O N/A O 1 O 2 O 3 O 4 O 5	
NOTE: If no to 4.22, cite 42 CFR 482.43(d) (Tag	NOTE: If no to 4.22, cite 42 CFR 482.43(d) (Tag A-0837)				

	Patient/Record #1	Patient/Record #2	Patient/Record #3	Patient/Record #4
4.23 Is there any evidence the patient has	O Yes	O Yes	O Yes	O Yes
been readmitted to this hospital within 30	O No	O No	O No	O No
days of a prior related admission?	O N/A	O N/A	O N/A	O N/A
	0 1	0 1	0 1	0 1
	O 2	O 2	O 2	O 2
	O 3	O 3	O 3	O 3
	O 4	O 4	O 4	O 4
	O 5	O 5	O 5	O 5
NOTE: Do not cite				