

Patient Data Collection Tool for Recommended Quality Measures

1)	Name of person doing data collection:			
2)	Date:			
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	king Information			
3)	Patient Identifier:			
4)	DOB://			
5)	Gender:			
Race	:			
6)	American Indian or Alaskan Native	YES	NO	
7)	Asian, Hawaiian or other Pacific Islander	YES	NO	
8)	Black or African American	YES	NO	
9)	Caucasian or White	YES	NO	
10) Some other race or races	YES	NO	
11) Hispanic Ethnicity:	YES	NO	UNABLE TO DETERMINE
12) Admission date:/	_		
13) Has this patient been discharged?		YES	NO
14) If yes, date of discharge://			
15) If yes, was the discharge due to death?	YES	NO	N/A

Health	History
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16) Primary reason for hospice admissi	on:				
 Heart Failure 	□ Heart Failure				
□ Cancer					
Neurologic Condition (e.g.: stro	oke, dementia)				
 Respiratory Condition 					
□ Other					
Indicate the presence or absence of any co	ommunication impairn	nents at	the time of admission:		
17) Deafness	YES	NO	UNABLE TO DETERMINE		
18) Dementia	YES	NO	UNABLE TO DETERMINE		
19) Non-English speaking	YES	NO	UNABLE TO DETERMINE		
20) Confused, sedated or non verbal	YES	NO	UNABLE TO DETERMINE		
21) Documentation of at least one famile 22) If yes, date of first family meeting: _	ly meeting is in chart	YES	NO Patient has no family		
Problem Screening, Assessment and Ma	anagement				
23) Was patient prognosis documented	? YES NO	UNAE	BLE TO DETERMINE		
24) If yes, what date was it documented	d?//_				
25) Was patient functional status asses	sed on admission?	YES	NO UNABLE TO		
DETERMINE					
26) If yes, on what date is functional sta	atus documented?	/	/		
27) Was patient screened for any phys	i cal symptoms (e.g.: ¡	pain, dys	spnea, nausea, constipation)?		
YES NO UNAE	BLE TO DETERMINE	If no	or UTD, skip to Question 69		
If yes, please answer the questions below		_			

28) Was patient screened for pain? YES NO UTD If no or UTD, skip to Question 45
29) Date of first screening:/
30) Type of pain scale used: NUMERIC VERBAL OBSERVATION NONE
31) For the pain screening were translated materials/interpreter used? YES NO N/A
32) If numeric scale used, what was pain score?
If non-numeric scale used, what was pain severity rating?
NONE MILD MODERATE SEVERE
33) Did the screening indicate the patient was in pain? YES NO If no, skip to
Question 45
34) Was a clinical assessment conducted to determine source of pain?
YES NO UNABLE TO DETERMINE
35) If yes , date clinical assessment for pain completed?/
36) Was treatment for pain initiated? YES NO If no, skip to next Question 45
37) Date first pain treatment initiated://
38) Type of pain treatment initiated (check all that apply):
□ Scheduled medication, non-opioid
□ Scheduled medication, opioid
□ PRN medication
□ Non-medication therapy
39) Was a bowel regimen initiated? YES NO N/A If no or N/A, skip to Question 41
40) Date of bowel regimen initiated:/
41) Was a second assessment of pain done? YES NO If no, skip to Question 45
42) Date of second pain assessment:/
43) Was pain improved on the second assessment? YES NO

44) If a numeric scale was used, what was pain score on second assessment?				
If a non-numeric scale was use, what was the pain severity on second assessment?				
NONE MILD MODERATE SEVERE				
45) Was patient screened for dyspnea? YES NO UTD If no or UTD, skip to Question 59				
46) Date of first screening:/				
47) Type of scale used: NUMERIC VERBAL OBSERVATION NONE				
48) If a numeric scale was used, what was dyspnea rating (using 0 to 10)?				
If a non-numeric scale was use, what was the dyspnea severity?				
NONE MILD MODERATE SEVERE				
49) Did the screening indicate the patient had dyspnea? YES NO If no, skip to Question 59				
50) Was a clinical assessment conducted to determine the source of dyspnea?				
YES NO UNABLE TO DETERMINE				
51) If yes , date clinical assessment completed?/				
52) Was treatment for dyspnea initiated? YES NO If no, skip to Question 59				
53) Date treatment for dyspnea initiated:/				
54) Was a second assessment of dyspnea done? YES NO If no, skip to Question 59				
55) Date of second assessment:/				
56) Was dyspnea improved at the second assessment? YES NO				
57) Was dyspnea treated or patient satisfied within 4 hours? YES NO				
58) If a numeric score was used, what was dyspnea score on second assessment?				
				
If a non-numeric score was use, what was the dyspnea severity rating on the second assessment				
NONE MILD MODERATE SEVERE				
59) Was patient screened for nausea? YES NO UTD If no or UTD, skip to Question 63a				

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60) Date of first screening:/				
61) Was patient nauseous? YES NO If no, skip to Question 64				
62) Was treatment initiated? YES NO If no, skip to Question 64				
63) Date treatment initiated:/				
a. Was patient screened for constipation? YES NO UTD				
If no or UTD, skip to Question 69				
64) Date of first screening for constipation:/				
65) Was patient constipated? YES NO If no, skip to Question 69				
66) Was treatment initiated? YES NO If no, skip to Question 69				
67) Date treatment initiated:/				
68) Was bowel function assessed at least weekly since initial screening? YES NO				
69) Was patient screened for depression? YES NO UTD If no or UTD, skip to Question 74				
70) Date of first screening:/				
71) Did the patient screen positive for depression? YES NO If no, skip to Question 74				
72) If yes, did the patient receive further assessment, counseling or medication treatment?				
YES NO UTD				
73) Was patient diagnosed with depression?				
74) If yes, date of depression diagnosis:/				
75) Date treatment for depression initiated:/				
76) Was patient screened for anxiety? YES NO UTD If no or UTD, skip to Question 81				
77) Date of first screening:/				
78) Did the patient screen positive for anxiety? YES NO If no, skip to Question 81				
79) Did the patient receive counseling, medication or other treatment for anxiety?				
YES NO If no, skip to Question 81				
80) Date treatment initiated:/				

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81) Is there documentation of a discussion of the patient's <i>spiritual</i> concerns?						
	YES	NO		If no, skip to Q	uestion 83	3
82) If yes:	Date of first of	liscussion of spirit	ual concerns:		_/	· —— ——
Patient Pr	eferences					
83) Does tl	ne patient cha	rt contain docume	ention of the p	atient's preferen	ces for life	sustaining
treatme	ents?				YES NO	UTD
84) Does tl	ne chart indica	ite a preference r	egarding the υ	ise of CPR?	YES NO	DTU C
85) Does tl	ne chart indica	ate a preference re	egarding hosp	italizations?	YES N	O UTD
86) Does tl	ne chart indica	ate a preference re	egarding the υ	se of mechanic	al ventilation	on?
YES I	NO UTD					
87) Does tl	he chart indica	ate a preference re	egarding artific	cial nutrition?	YES N	O UTD
88) Does tl	he chart indica	ate a preference re	egarding the υ	se of antibiotics	? YES N	IO UTD
89) Does tl	ne chart indica	ate preference for	location of de	ath?	YES N	NO UTD
90) Does th	ne patient cha	rt contain an adva	anced directive	e (living will or he	ealth care	power of
attorne	y)? YES N	O UTD				
91) If yes ,	date advance	d directive placed	in chart:	/		
92) If no, is	s there docum	entation of a disc	ussion about a	an advanced dire	ective or th	nat the patient has
no adv	anced directiv	e? YES NO				
93) Does tl	he chart conta	in the name and o	contact inform	ation of the surr	ogate deci	sion maker?
YES	NO	UTD				
94) If yes ,	date placed in	chart:/_	/		UTD	
95) If no, is	s there docum	entation of a disc	ussion that no	surrogate decis	sion maker	is available?
YES	NO					
96) If yes,	date of discus	sion:/	/		UTD	

Care around the time of death

Fo	r the	e next set of questions, please refer to the last week of the patient's life.
97)) Ind	icate the patient's average level of alertness during the final week of life:
		Mostly alert
		Drowsy but aware
		Somnolent but able to be awakened
		Mostly asleep or unconscious
		Unable to determine
98)) Wh	nat was the highest pain severity noted during the patient's final week of life:
		None
		Mild
		Moderate
		Severe
		Unable to determine
99)) In v	what setting did this patient die?
		At home
		Acute care hospital
		Nursing home
		Hospice inpatient facility
		Assisted Living
		Other (please specify)

END OF DATA COLLECTION

This material was prepared by The Carolinas Center for Medical Excellence (CCME), the Medicare Quality Improvement Organization for North Carolina, under contract with the Centers for Medicare & Medicaid Services (CMS), an agency of the U.S. Department of Health and Human Services. The contents presented do not necessarily reflect CMS policy. 8SOW-NC-PEACE-08-06