OnCore Communication Memo

Memo #: 36 – Addition of Clinical Research Coordinator at Subject Consented

To: All CRC's, CRA's, Research Managers, CTO-AD From: System Administrator Effective Date: February 8, 2018

Purpose: The Workload Calculator report was originally designed to calculate workload based on the lead Clinical Research Coordinator listed on the PC Console. However, after further review it was determined the report would more accurately reflect the true workload if it is calculated using the Clinical Research Coordinator on the Subject Console>On Study tab. Therefore, in order to accurately track subjects assigned to each CRC throughout the subject lifecycle on the protocol, the addition of CRC will now be required at the time of consent, and entered in the On-Study tab.

Notes: After entering the Consent Date on the Consent tab, navigate to the On-Study tab and enter in the Clinical Research Coordinator under the Subject Staff section of the On-Study tab, as well as entering the already required Disease Site and/or Histology. Please note that an error message will pop up: "Warning: Eligibility verification is not done for the subject." The data will still be entered so this message can be ignored. Please remember to change the Clinical Research Coordinator on this tab if/when the subject is transferred to another CRC/CRA. Only one Clinical Research Coordinator should be listed on this tab. Use Clinical Research Coordinator-Secondary for any additional coordinators.

Warning: Eligibility verification is not done for the subject.									
Message: Record has been updated									
★ Subject Console									?
Protocol No.: SWOG-S1400G						tatus: OPEN TO ACCRU			Subject Status: CONSENTED
MRN: 55555555					Su	bject Name: Ginger Len	non		Sequence No.:
Epic									
Switch Subject	Subject On Study Upda	ate							
Type here to search	Seq	quence No.]			On Study Date	117
Summary	Di	isease Site	ung & Bro	nchus				Ready for Registration	
Demographics		Histology	Гуре here t		×				
Consent	Diag	nosis Date			•				
	Oncol	logy Group	Thoracic •	·				ZIP at Registration	
Eligibility		Study Site	Froedtert Hospital					Transferred Date (MM/DD/YYYY)	
On Study		Comments							
Treatment						/_			
Follow-Up				Additional Protocol Subject lo	dentifiers				
				Identifier Type		Identifier			
SAEs				Identifier Type		Identifier			
			=	•		No information er	atorod	Add Cancel	
Payments						NO INOMIAUON EI	ntereu		
Deviations	Subject Staff								See All
Documents/Info »	Role			Staff Name		Start Date			Team
Documents/Into »	Type here to sear	ch 💌		Type here to se	earch				Add
Protocols	Dala				Last Name	First Name	Malata 1-30-1	Home Organization	
MRN	Role Clinical Research Co	oordinator			Last Name Bien	Collette	Middle Initial	Medical College of Wiscon	sin
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