OnCore to PowerTrials Process Steps (Cancer Center Studies)

Responsibilities:
- Regulatory Team
- CRS (Clinical Research Specialists)
- Research Nurses
- System

	Preparation	Open to Accrual	Enrolling Subjects				IRB Study Closure
OnCore	1. Verify that protocol should be in PowerTrials: • Full IRB Review • IU Health MRN • IU Health Study Site • Requires Access for Monitor Visits 2. Add Primary Study Site Contact to PowerTrials Study Contact role	Protocol OPENED to accrual	Monitor & address discrepant subjects				Protocol IRB Study CLOSURE
			ON Study	ON Treatment*	OFF Treatment	OFF Study	
RPE		Manual send	Auto send	Auto send		Auto send	Manual send
PowerTrials	While protocol amendments/revisions DO NOT have to be sent to PowerTrials, you will have to push over any changes to the PowerTrials Study Contact during the course of the trial. See the PowerTrials job aid for assistance if needed.	Protocol created in PowerTrials and is viewable in Research Viewpoint	Subject **ON** Study in PowerTrials	Subject **ON** Treatment in PowerTrials	Subject **OFF** Treatment in PowerTrials	Subject **OFF** Study in PowerTrials	Protocol closed in PowerTrials
PowerChart			Subject ON Study Message center admission notifications to PowerTrials Study Contact	Add Clinical Note for each subject enrolled in the trial.		Subject OFF Study ** Please note that ARM information will be present on the Clinical Research page in PowerChart if sent from OnCore.*	Messages center notifications stop

10.24.2021