

Emory OnCore

Ask an RCRA Session

April 28, 2022



OnCore Quick Facts



WHAT IS IT: New Emory Clinical Trial Management System/CTMS (*will replace ERMS*)



WHEN DOES IT START: Fall 2022



WHO WILL USE IT: Staff involved with clinical research



HOW WILL I GET ACCESS: After required training



WHY NOW: Standardization, compliance, and alignment with EHC Epic implementation

What studies will go into OnCore?

REQUIRED studies in OnCore:

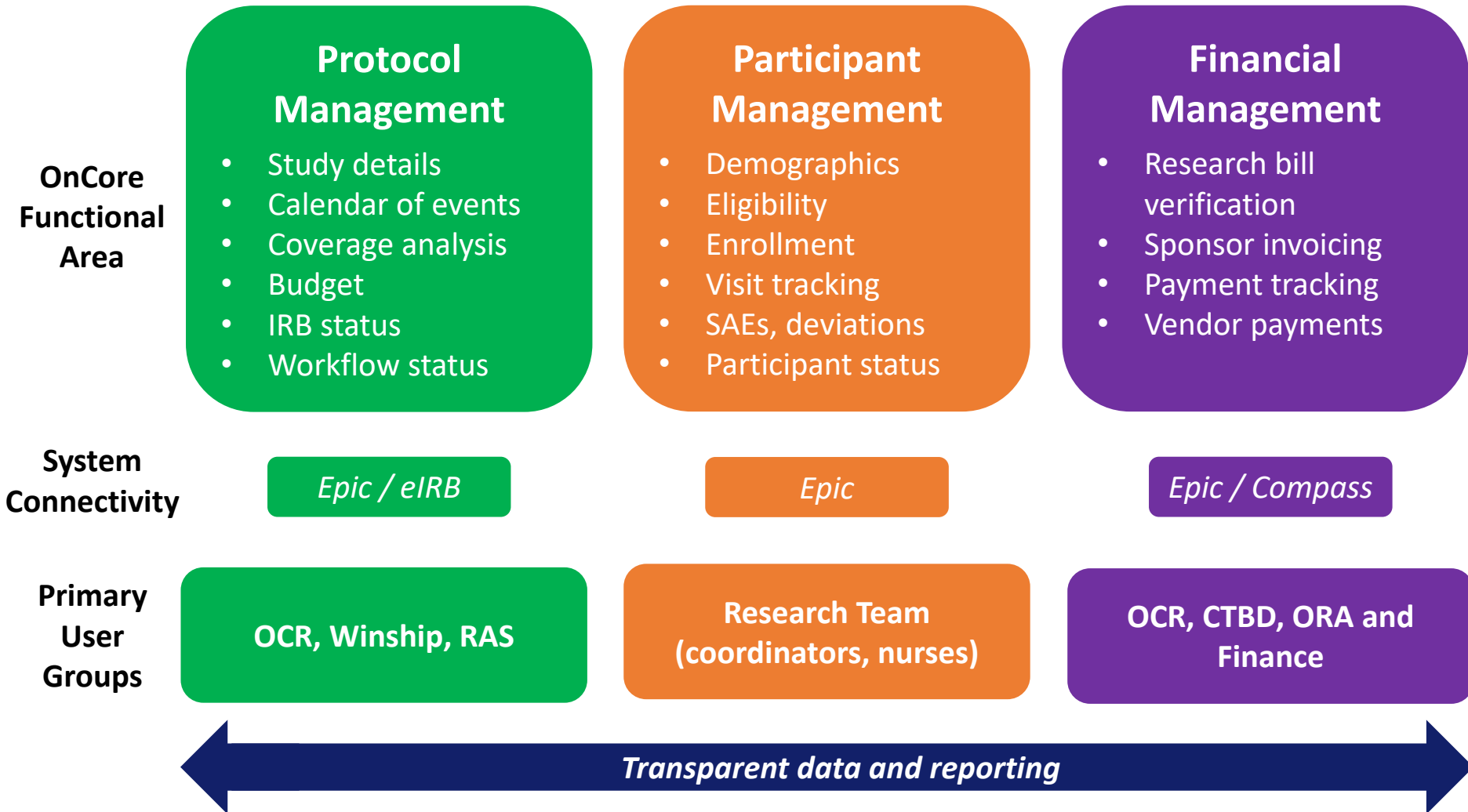
Same as ERMS

- All studies with Emory Healthcare or Grady billable items - regardless of sponsor/funder - requiring a Prospective Reimbursement Analysis (PRA)
- All studies meeting NIH clinical trial definition regardless of billables
- All studies where OCR performs the invoicing
- All studies associated with Winship Cancer Institute

OPTIONAL studies in OnCore: (*“Shell Protocols” will automatically be created with no calendar, PRA, or budget but allows study/subject coordination as needed*):

- Non-interventional clinical research without billable items and no OCR invoicing
- Federal studies without billable items

OnCore Overview



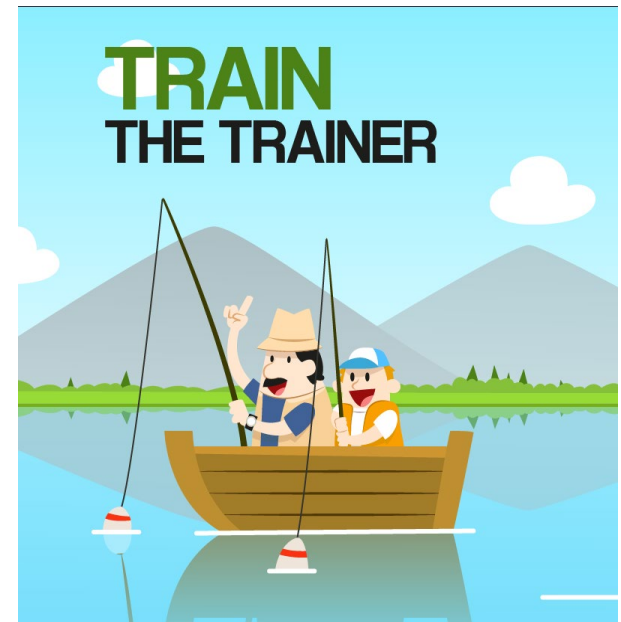
PLEASE NOTE: this represents a **HIGH-LEVEL summary** – additional details, processes, and variations will exist

Training Model

- 40+ departments, 1000+ clinical research professionals
- Train-the-Trainer (TTT/T3) Model
A T3 model is a training framework that turns employees into **subject matter experts/superusers** who can then teach other staff within their department.

Benefits:

- A tailored learning experience using job aids
- Dissemination of information made fast and easy
- Curriculum consistency
- Develops an internal training team



Thank you

For more information, please visit:

<https://emory.sharepoint.com/sites/OnCore>

Save the Date for Emory OnCore Townhall #2

July 13th, 12:00 – 1:00

