

RESEARCH CERTIFICATION EXAM REVIEW GROUP

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SCORE Mtg – 12NOV2019

GENERAL CONSIDERATIONS

Thoroughly review ACRP/SoCRA exam info

Assess your own experience/skill

Strengths? Weaknesses?

Assess your other time commitments & comfort level

Start early & plan ahead

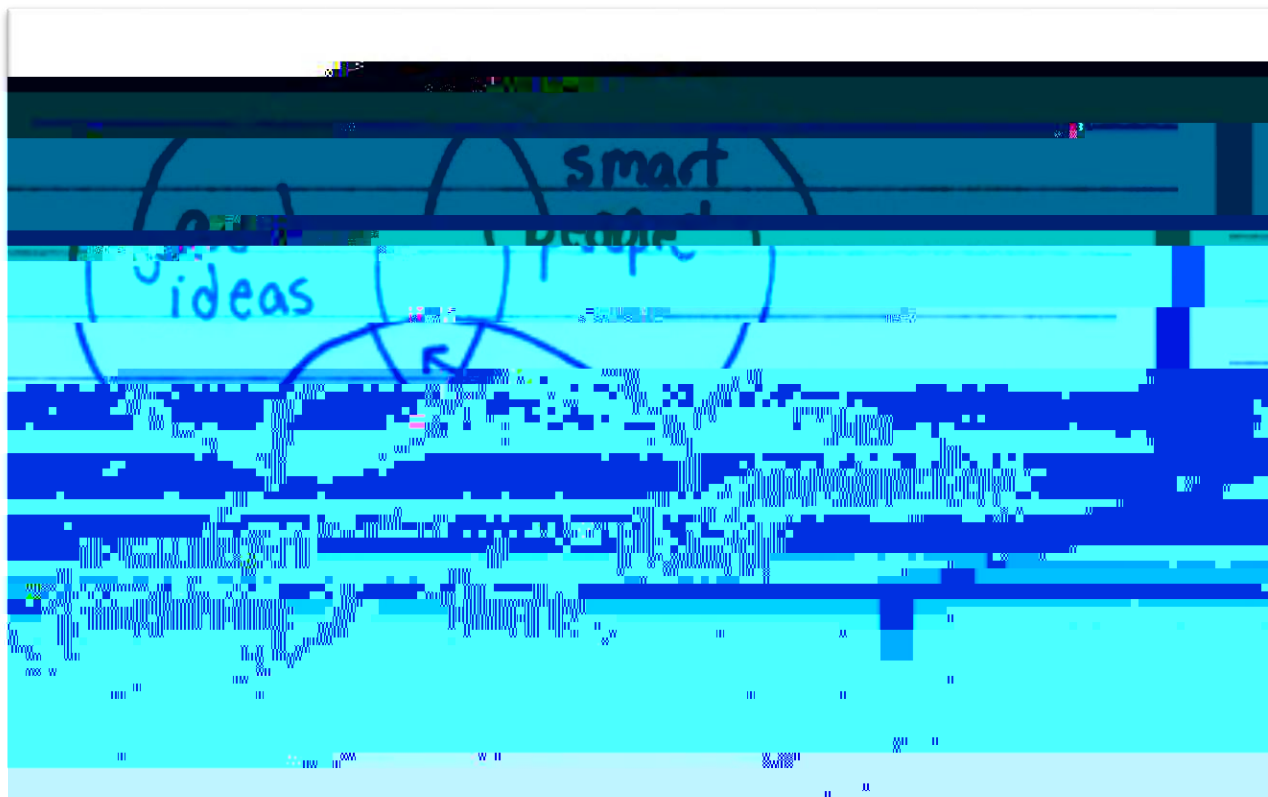
Schedule your study time

Choose study methods that are right for you

GOALS & EXPECTATIONS

Study Group Goal =

DISCLAIMER



FORMAT & SCHEDULE

Weekly Sessions, January 6 – February 24

Mondays, 4-6pm, SRB 1404 (Or Zoom)

Format (Most Sessions):

Assigned materials to review *prior* to each session

- Content area reading assignments and slide sets

Facilitated session review:

- Review assigned slide sets to answer questions, clarify content, emphasize important content
- Review practice questions

Independent group study for remainder of session

Additional resources provided: glossaries/abbreviation lists, guidelines & regulations, textbooks, online training options/resources, sources for practice questions

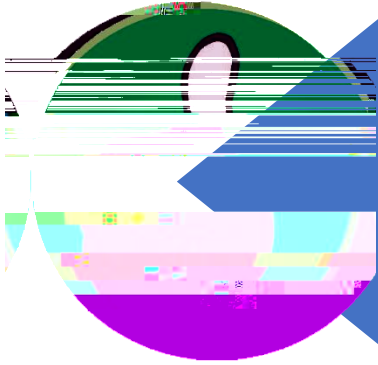
SCHEDULE

Weekly Sessions, January 6 – February 24
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SESSION	CONTENT
1	Introduction & RCERP Overview
2	Historical Perspectives; Study Design Basics & Drug/Device Development; Federal Regulations & ICH GCP Overview; FDA-Regulated Research
3	The Study Protocol; Study Feasibility; Institutional Review Boards; Roles & Responsibilities; Recruitment & Retention; Informed Consent
4	Study Activities; Essential Documentation & GDPs; Adverse Events & Unanticipated Problems; Data & Safety Monitoring Plans
5	Guest Attendees
6	Monitoring, Audits & Inspections; Investigational Product Management; Data Management; Genetic Research; International Trials
7	Practice Questions & Wrap-Up

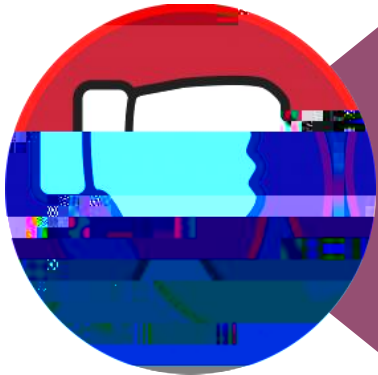
REVIEW MATERIALS





Interested?

Sign-up sheet on side table (or email Kelly)



No Thanks!

Independent review materials are available