

## **Boot Camp Schedule**

## Interactive Response Technology (IRT) in Clinical Supply Chain

	Greenwich Mean Time	Eastern Time	Pacific Time	Title	Description	Communique Room	Instructors
Monday, Apr 12	14:00 - 16:30 GMT	10:00AM - 12:30PM ET	7:00- 9:30AM PT	- History - Workflow Timing - Understanding Critical Study Parameters - Blinding and Unblinding Controls	This class is geared towards people that have some experience in the clinical trial supplies world. Attendees will be provided with a quick overview and then will take a deeper dive into Interactive Response Technology (IRT). This will include the development path for IRTs; the stages of build, from creating the User Requirement Specification (URS) to quoting, coding, validation, user acceptance testing and go-live; the reasons to use an IRT as well as when NOT to use an IRT; build requirements, including randomization codes and med code IDs; ensuring modules are built to manage individual regulatory country approvals, shipping requests and returns and reconciliation; as well as timelines from concept development to system go-live.		Jan Pieter Kapelle TBC
Tuesday, Apr 13	14:00 - 16:30 GMT	10:00AM - 12:30PM ET	7:00- 9:30AM PT	- Randomization Techniques & Trial Designs - Supply Management Models - IMP Status Management - Managing Site & Depot Supply			Jan Pieter Kapelle TBC
Wednesday, Apr 19	14:00 - 16:30 GMT	10:00AM - 12:30PM ET	7:00- 9:30AM PT	- How to calculate your Drug Management Parameters? - End-of-Study, What should you do? - Benefits of using standards			Jan Pieter Kapelle TBC
Thursday, Apr 20	14:00 - 16:30 GMT	10:00AM - 12:30PM ET	7:00- 9:30AM PT	- Key users, Roles, Responsibilities and Needs - Importance of cross-functional collaboration - When (not) to use RTSM? - IMP management without RTSM - Best practices in Sponsor Service Provider Partnership - Technology platform and IRT Integrations - RTSM usage for New Trends and Products - Summary & Review			Jan Pieter Kapelle TBC