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Investigator Initiated Studies (IIS) are crucial for new product development. These Investigator Initiated Studies represent new challenges for the Investigators and their team. Indeed, managing a Clinical Trial as sponsor is outside of their day-to-day work. This Investigator Initiated Studies training programme is intended to help them understanding how to implement mandatory Clinical Studies rules for sponsor in practice to ensure patient safety and data quality and overcome all challenges associated to it. If you want to have a clear overview of the IIS set-up and management specificities, then this course is for you!

The programme consists of a series of 14 webinars of 1-2 hours each. The webinars are interactive, to encourage discussions between the subject matter experts and the audience. All sessions will be recorded and available on the ECCRT Virtual Campus in case you have missed one.

Who should attend?

Physicians, acting as Investigators in (academic) hospitals and universities and who want to be involved in setting up Investigator Initiated Studies within their organisation. Principal investigators, sub-investigators, graduated doctors, fellowship, general practitioners will benefit attending this training.

Find out more information on their website and register here:
https://eccrt.com/course_display/investigator-initiated-studies/ [1]

Last update: 23 February 2021

Links
[1] https://eur01.safelinks.protection.outlook.com/?url=https%3A%2F%2Feccrt.com%2Fcourse_display%2Finvestigator-initiated-studies%2F&data=04%7C01%7Cmarta.anguera%40eahp.eu%7C06bc097d7ed54e4f163308d8d80cf7e9%7C2