Clinical Trials in Oncology: clinical, management and operative aspects.

Professional outlets: At the end of the program, the participant will be able to: cooperate to the drafting of clinical trials protocols in oncology; coordinate clinical trials in compliance with regulations, ethics, methodology and Good Clinical Practice(GCP); coordinate trial activities with respect to the participants' safety and rights; manage data accurately and coordinate operational aspects of commercial and investigator initiated clinical trials. The Master course trains professionals as Clinical Research Coordinators (CRC), data managers and Research Nurses for hospitals, University Hospital, University polyclinics, IRCCS, ASL, private health facilities, pharmaceutical companies and CROs.
Director: Pierfranco Conte
Level: 1
Duration: one-year
Period: November 2020 / September 2021
Teaching method: taught class / distance learning
Language: Italian
Short Specialization degree's location: Istituto Oncologico Veneto IOV - c/o Ospedale Busonera - Via Gattamelata, 64 - 35128 Padova
Places available: min: 10 / max: 25
Registration fee: Euro 2.649,00 (first payment: 1.649,00 euro / second payment: 1.000,00 euro)
Benefits / Scholarship: n. 10 possible study awards
Criteria for selection: evalutation of qualifications
Application submission deadline: October 2nd, 2020
Website: www.discog.unipd.it
For information: 049 8211803; master.sperclinonc@unipd.it