TRAINING COURSE

THIS IS TO CERTIFY THAT:

HAS PARTICIPATED IN Good Clinical Practice - R2 for Investigators

COVERING FOLLOWING TOPICS

ICH, ICH E6 article 4 (Investigator qualification and agreement, Adequate Resource, Medical care of Trial Subjects, Communication with IEC, Compliance with Protocol, Investigational Product, Randomization Procedures and Unblinding, Inform Consent of Trial Subjects, Records and Reports, ALCOA, Safety Reporting, Premature Termination or Suspension of Trial, Final Report)

26-OCT-2017

Mgr. Katarína Kováčová

KOVAC SERVICES S.R.O.