KS/GCP-201606xx

TRAINING COURSE

THIS IS TO CERTIFY THAT:

HAS PARTICIPATED IN

Good Clinical Practice for Investigators

Identification no. in SLK: CJ495SK

COVERING FOLLOWING TOPICS

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), Declaration of Helsinki, Clinical trial phases

21-JUN-2016

Mgr. Katarína Kováčová
trainer

Credits for attendance: 5

KOVAC SERVICES S.R.O.

The credits are acknowledgments according the regulation of Ministry of Health 366/2005 as amended by the act. This ICH E6 GCP Investigator Site Training meets the Minimum Criteria for ICH GCP Investigator Site Personnel Training identified by TransCelerate BioPharma, Inc. as necessary to enable mutual recognition of GCP training among trial sponsors.