

# TRAINING COURSE

THIS IS TO CERTIFY THAT:

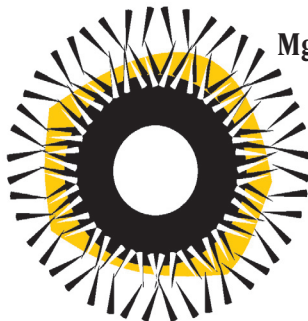
**HAS PARTICIPATED IN** Good Clinical Practice - R2 for Investigators

## 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, Investi-  
gational 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.