



Professional Education Program



Introduction to Good Clinical Practice V2.0

(Includes the 2016 updates to ICH E6 R2)



This is to certify that

Llewellyn Mills

Completed the following training programs:

- GCP The Standards and why we have them
- Study set up responsibilities, approvals and essential documents
- The process of informed consent
- Case Report Form, source data and data entry completion
- Safety reporting

The equivalent of 6 hours of study was completed.

Pass mark 100%

GCP TRAINER: SOPHIE MEPHAM

Date of training 28th May 2019



"This ICH E6 GCP Investigator Site Training meets the Minimum Criteria for ICH GCP Investigator Site Personnel Training identified by TransCelerate BioPharma as necessary to enable mutual recognition of GCP training among trial sponsors."