Title: A phase IIIb, open-label, multicentre, international randomised controlled trial of simplified treatment monitoring for 8 weeks glecaprevir (300mg)/pibrentasvir (120mg) in chronic HCV treatment naïve patients without cirrhosis

Protocol No: VHCRP1701

Site Staff Training Log

Site Name: ______________________________ **Site Code:** ______________ **Principal Investigator:** ______________________________

Staff Name (First and Last Name)	Role	Date Started on Study	Date of Training	Topics Covered (1 – 8)	Conducted by (Name of Trainer/s)	Staff Signature
		__ __/ __ __/ __ __	__ __/ __ __/ __ __			
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All site staff must be trained in the study. The following topics are required:

1	Background & Overview	2	Study Protocol	3	ICH/GCP and consent
4	Study Visits & Procedures	5	Study questionnaires / Data Management	6	Study Documentation
7	Research Samples & Labkey	8	Other:	9	Other:

To be completed at Site Initiation Visit and updated whenever new staff joins the study.
Original to be filed in Investigator Site File and a copy sent to the SMART-C Project Coordinator.