Protocol Title: A Scale-up of treatment for hepatitis C infection among people who inject drugs:
A phase IV, open-label, single arm, multicentre trial of grazoprevir/elbasvir for genotype 1 or 4 in people with chronic hepatitis C virus infection and recent injecting drug

Protocol No: VHCRP1510

Site Signature and Study Responsibilities Log

Site Name: _____ **Site Code:** __ __ __ __ __ **Principal Investigator:** _____

First name	Last name	Signature*	Initials	Date from *dd/mmm/yyyy*	Date to *dd/mmm/yyyy*	Role *(PI, Co-PI, Coordinator, Lab, Pharmacist)*	Responsibility Code(s)*	# CV and ICH-GCP certificate on file	Site PI to initial and date delegation
								Yes ☐ No ☐	
								Yes ☐ No ☐	
								Yes ☐ No ☐	
								Yes ☐ No ☐	
								Yes ☐ No ☐	
								Yes ☐ No ☐	
								Yes ☐ No ☐	
								Yes ☐ No ☐	
								Yes ☐ No ☐	
								Yes ☐ No ☐	

Study responsibilities

1. Obtaining informed consent (investigator's only)
2. Screening assessment; Conduct study visit procedures
3. Eligibility determination (investigator's only)
4. Maintain essential documents
5. eCRF Completion
6. SAE Notification
7. Collection of research samples
8. Processing and storage of research samples
9. Dispensing of study drug
11. Other (specify)_____
12. Other (specify)_____