

Appendix 1: Agenda

Health Informatics Workshop: Use of the electronic health record in the design and conduct of clinical trials

10:00-16:00, Monday 23rd June 2014

Staff House Conference Centre, The University of Manchester

09:30	Registration, tea & coffee
10:00	Welcome and introduction Professor Paula Williamson, NWHTMR, University of Liverpool
10:10	Overview of use of routinely collected health data in trials and an introduction to the 3 topic areas covered in the workshop Professor Ian Ford, Glasgow CTU
10:35	A survey of UK registered CTUs Dr Oluseun Adeogun, University of Liverpool
Topic 1: Feasibility assessments using routine data sources: 2 examples	
10:45	Feasibility assessments and recruitment using routine data sources: experience from the THRIVE trial Dr Chris Bray, Oxford CTSU
11:00	The PLEASANT Trial: a case clinical trial case study using routinely collected data Professor Steven Julious, Sheffield CTU
11:15	Tea/Coffee Break
Topic 2: Identifying and contacting eligible patients: 2 examples	
11:30	Recruitment models for pragmatic point-of-care randomised trials using routinely collected electronic records Professor Tjeerd Van Staa, University of Manchester
11:45	Identification of eligible patients for clinical research within primary care Dr Martyn Lewis, Keele University
Topic 3: Outcome data collection using the EHR:	
12:00	The use of routine data to enhance collection of trial primary and secondary endpoints in the SHIFT trial Prof Amanda Farrin, Leeds CTRU
12:15	Q & A from morning session
12:30	Lunch & Networking
13:30	Including routine linked data outcomes in trials: opportunities, benefits and risks Professor Helen Snooks, Swansea University
13:45	Small group work: Methodological research priorities
14:45	Feedback from small group work
15:15	Tea/Coffee Break
15:30	Future plans
16:00	Close of Meeting