

**MRC North West Hub for Trials Methodology Research and The North West Clinical Trials Collaborative**

**On-campus course 'Improving health by improving trials'**

**Programme 16<sup>th</sup> September – 20<sup>th</sup> September 2019**

**Location: The Foresight Centre, University of Liverpool**

(please note the course will be held at the Clinical Trials Research Centre, Alder Hey Children's Hospital on Wednesday 18<sup>th</sup> September)

Monday 16 <sup>th</sup>		Registration and coffee	
		Welcome, introductions and overview of course	
	<b>AM SESSION</b>	<b>What do you need to do to design a trial?</b> <i>Professor Paula Williamson, University of Liverpool</i>	Topics: Identifying, defining and justifying the question; the importance of the protocol
	<b>PM SESSION</b>	<b>General design issues</b> <i>Dr Susanna Dodd, University of Liverpool</i>	Topics: Feasibility, external and internal pilot studies; pragmatic and explanatory designs, internal and external validity; sample size considerations
Tuesday 17 <sup>th</sup>	<b>AM SESSION</b>	<b>Introduction to different designs</b> <i>Dr Chris Sutton, University of Manchester</i> <i>Professor James Wason, Newcastle University/</i> <i>Cambridge MRC Biostatistics Unit</i>	Topics: Adaptive designs; trials of complex interventions; cluster randomised trials
	<b>PM SESSION</b>	<b>Recruitment of trial participants</b> <i>Professor Bridget Young, University of Liverpool</i> <i>Dr Nicola Harman, Clinical Trials Research Centre</i>	Topics: Barriers and facilitators; effective recruitment and retention strategies; recruitment monitoring
Wednesday 18 <sup>th</sup>	<b>AM SESSION</b>	<b>Trial conduct (part 1)</b> <i>Dr Emma Bedson, Clinical Trials Research Centre</i> <i>Ms Katie Neville, Clinical Trials Research Centre</i>	Topics: Ethical, legal and regulatory requirements; pharmacovigilance; barriers and facilitators to setting up sites; data sources; information systems and data management
	<b>PM SESSION</b>	<b>Visit to the Clinical Trials Research Centre, Liverpool</b>	
Thursday 19 <sup>th</sup>	<b>AM SESSION</b>	<b>Trial conduct (part 2)</b> <i>Professor Catrin Tudur Smith, University of Liverpool</i> <b>Keynote talk 'How to be a good Chief Investigator'</b> <i>Professor Tony Marson</i>	Topics: Risk assessment; risk-based monitoring and safety monitoring; trial oversight committees
	<b>PM SESSION</b>	<b>Public and Patient Involvement</b> <i>Professor Peter Bower, University of Manchester</i> <i>Dr Claire Planner, University of Manchester</i> <i>Ailsa Donnelly</i>	Topics: Basic principles of patient centred trials; evidence of benefit
Friday 20 <sup>th</sup>	<b>AM SESSION</b>	<b>Analysis and reporting (part 1)</b> <i>Professor Richard Emsley, King's College London</i>	Topics: Key principles of trial analysis; intention to treat analysis; adjustment for compliance; mechanisms evaluation; stratified medicine trials
	<b>PM SESSION</b>	<b>Analysis and reporting (part 2)</b> <i>Professor Dyfrig Hughes, Bangor University</i> <i>Dr Jamie Kirkham, University of Liverpool</i>	Topics: Basic principles of health economics; methods of economic evaluation; economic outcomes; good practice in trial reporting