**NIHR-MRC Trials Methodology Research Partnership Trial Conduct Working Group: Expression of Interest**

**Co-leads:** Catherine Arundel (University of York) , Nicola Harman (University of Liverpool)

The Trial Conduct working group provides a forum to facilitate the development of research and improvements across key aspects of trial conduct and will seek to minimise duplication of effort within relevant areas. We consider Trial Conduct to cover activities that happen from study set-up to reporting of results. Development of methods and/or good working practice across this spectrum of conduct activities will be considered within this working group.

We welcome new members currently active in relevant trials methodology across a wide range of groups including patient research partners, academic researchers (from a range of relevant disciplines and any career stage), trialists, health professionals, postgraduate students and industry representatives.

Please complete this form and return it to the working group co-leads with the subject **TCWG Expression of Interest** you will then be added to the relevant topic group/s on Basecamp.

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| --- | --- | --- | --- | --- |
| **Name** |  | **Job title/role** |  | |
| **Organisation** |  | | | |
| **Email** |  | We will use this email to add you to the basecamp platform. We will also use it to contact you about relevant trial conduct working group meetings. | | |
| **What country are you based in?** |  | | | |
| **Please indicate if applicable:** | I am a PhD student | | | |
| I am an early career researcher | | | |
|  | | | | |
| **Topic group/s that you would like to be a member of**  *(tick all that apply)* | **Inclusivity** (ensuring trials are conducted in an inclusive way) | | |  |
| **Recruitment** to trials (Informed by the Priority I research agenda) | | |  |
| **Retention** to trials (Informed by the Priority II research agenda) | | |  |
| **Data quality and monitoring** (how to maximise good data and best monitor trials) | | |  |
| **Qualitative Research within trial conduct** (e.g. generation of good working practice documents, costing models, connectivity and learning across existing data sets, capacity building) | | |  |
| **Alternative and Complex Consent** (methodological and ethical challenges encountered by underserved groups) | | |  |
| **Greener trials** (research in the area of environmentally sustainable clinical trials) | | |  |
| **What sector do you work in?** | Health Care Organisation | | |  |
| Higher Education Institution | | |  |
| Industry | | |  |
| Third Sector | | |  |
| Other (please specify) | | |  |
| **Are you linked to any of the following** | A UK registered Clinical Trials Unit | | |  |
| UK TMN | | |  |
| HRB-TMRN | | |  |
| Global Health Network | | |  |
| Other TMRP Working group (please specify which groups) | | |  |

**Please return your completed form to** methodologyhubs@liverpool.ac.uk