**MRC-NIHR TMRP Health Informatics Working Group: Membership Expression of Interest**

The Health Informatics Working Group will focus on the methodology needed to realise the potential of digital technology to improve trial design, conduct and analysis.

We are seeking members currently active in relevant trials methodology to contribute to:

1. Oversight activities, via a small steering group;
2. Research and dissemination activities, via a number of sub-groups focusing on topic areas;

We will also maintain a mailing list of ‘affiliate’ members, to provide regular updates about working group activity.

Membership will be inclusive and reviewed annually. We wish to include key collaborators, academic researchers, trialists, clinicians, early career researchers, TMRP PhD students, PPI, industry representatives and ensure a mix of gender, career stage, geographical location and disciplines.

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| **If you wish to connect with Working Group, please complete the information below:** |
| **Name** |  |
| **Institution** |  |
| **Job Title**  |  |
| **Email**  |  |
| **If you wish to be an affiliate member only, please tick box 🡪** |  |
| **If you wish to additionally apply for full membership, please complete the following details:** |
|  |
| **Gender**  |  |  **Are you an Early Career Researcher?** | *Yes / No* |
|  |
| **Are you linked to any of the following?*****Tick all that apply*** | **HDR-UK** |  |
| **NHS Digital** |  |
| **Registered CTU** |  |
| **UKCRC Registered CTU Network** e.g. Task & Finish Groups*(please specify)* |  |
| **Other TMRP Working group** *(please specify)* |  |
| **MRC TMRP MRC PhD student** |  |
|  |
| **I can contribute to the Working Group in the following topic areas:*****Tick all that apply*** | **Use of electronic healthcare records / routine administrative datasets** in support / conduct / design of clinical trials  |  |
| **Use & assessment of information collected directly from trial participants** (e.g. apps wearables, online patient-reported outcome measurement etc) |  |
| **Use & assessment of technology to support trial conduct** (e.g. interactive eCRF, electronic consent tools) |  |
| **Clinical trials data sharing** (e.g. methods for anonymising, sharing, optimised storage of trial data  |  |
| **Novel trial designs** for intervention evaluation alongside routine data collection |  |
| Methods for trial design / conduct when **evaluating digital health interventions** |  |
| Provide a brief statement, outlining your research activity aligned to the Working Group’s topic areas  |  |
| **Please return this Expression of Interest to a.j.farrin@leeds.ac.uk**  |