**PrimDISC application**

**Please submit this application form three weeks ahead of the next PrimDISC meeting via** [**primdisc@phc.ox.ac.uk**](mailto:primdisc@phc.ox.ac.uk)**.**

**Dates can be found on the website:** [**https://www.phc.ox.ac.uk/intranet/better-workplace-groups-committees-open-meetings/primdisc-committee**](https://www.phc.ox.ac.uk/intranet/better-workplace-groups-committees-open-meetings/primdisc-committee)

**Title and Chief Investigator**

**Title of Research:**

**Chief Investigator:**

**Organisation:**

**Co-Applicants:**

**Provide lay summary:**

**Please tick the relevant box which data resource is requested:**

ORCHID-Epidemiology (ORCHID-E)  ORCHID Trials/Prospective studies (T/P)

Clinical Trials Unit (CTU)  Departmental cohort data  Secondary use of data

DPhil/student project  Other (please specify)

**Does this project have any existing ethical approval?**

No  Yes Please attach approved protocol, REC notification of approval and list of approved documents

Please note that ORCHID-E projects do not need their own ethical approval. Studies are covered under the overarching ethical approval of ORCHID-E.

**If it is not ORCHID, please list the data set PI:**

**Experience – please provide a description of your team’s expertise in the different areas**

**Methodological Expertise (Statistics, Machine Learning):**

**Data Management:**

**Funding**

**Has the funding been secured?**

**Funding body:**

**Datasets – please provide a description of the variables required for each of the datasets**

**Protocol**

**What is the scientific justification for the research? What is the background? Why is this an area of importance?**

**Give a brief synopsis / summary of methods and overview of the planned research:**

**What are the principal research questions/objectives?**

**What are the secondary research questions/objectives?**

**What are the principal inclusion criteria? (please justify):**

**What are the principal exclusion criteria? (please justify):**

**What are the health or economic outcomes to be measured?**

**What is the primary outcome measure for the study?**

**What are the secondary outcome measures?**

**Study Design**

**Study Design (e.g. case control, cohort, cross-sectional study etc)**

**Selection of comparison group(s) or controls**

**What are the exposures?**

**Describe the statistical methods and / or other relevant methodological approaches (e.g. for qualitative research) to be used in the analysis of the results.**

**Plan for addressing confounding**

**Plans for addressing missing data**

**Limitations of study design, data sources, and analytic methods**

**Please present your feasibility calculation. Has the size of the study been informed by a formal statistical power calculation?**

**Patient or user group involvement: *Describe how you have involved patient or user groups in the development of the research question? How you plan to involve them in the research and dissemination of the findings of the projects?***

**Data retention, storage and destruction**

**How long do you wish to retain the data?**

**If longer than 12 months, please justify your reasons below (the maximum data retention period is 3 years, with a review annually)**