**Project Reference:**

**Project Title:**

**Principal Applicant:**

**Contact Person:**

*Please supply a brief update on progress of your project.  Short comments are fine for each section.*

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| --- |
| 1. **Progress of analysis** |
| 1. **Have new variables been generated for study participants?** *This could be by analysis of samples or derived from existing data (e.g. individual risk scores). Also include datasets arising from computation, such as imputation or sequencing analysis ‘pipelines’:* |
| 1. **Incidental findings in participants of METADAC studies: genetic variants of potential clinical significance** *(see <link> for more information*) 2. Has your analysis revealed any variants involved with serious illness/impairment **YES/NO**   **If YES:** please answer questions (ii) and (iii) below and give brief details of the findings.   1. Variants with a strong scientific case that they are causative **YES/NO** 2. Variants with a possible intervention (i.e. benefit to knowing it is present) **YES/NO**   Brief details: ● |
| 1. **Named researchers**   Are there any changes to your research team **YES/NO**  **If YES**, please note the changes here to apply for METADAC’s approval for the amendment. |
| 1. **Please list any publications and planned publications** *(see* [*www.metadac.ac.uk*](http://www.metadac.ac.uk) *for required data acknowledgements)* |
| 1. **Anticipated timescale for completion of project and publications.** *(Note that any extension to the existing approval will be under current conditions of use.)* |

This proforma was completed by ………………………………………………… on ……………………..(date) and is correct to the best of my knowledge.

**A note on open data access**

*Many funders and journals require that data be made ‘open’ for other researchers to use. Please note that the data for your project cannot be placed in ‘open’ repositories without breaching the conditions of study participants. However, funders and publishers alike accept the validity of ‘managed access’ to ensure bona fide researchers can access the data in accordance with participants’ consent. Please contact the secretariat if you require advice on acceptable repositories for any data arising from your study.*