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**Agreement of Principal Investigator to abide by Guidelines**

**for Access to Data from the CRPS UK Registry**

**Project title submitted to CRPS UK Registry Steering Committee:**

**…………………………………………………………………………………………**

**Project number (to be added by CRPS administrator): ……….**

**Undertaking by Principal Investigator**

I have read and understood the guidelines Access to Data from the CRPS UK Registry. I undertake to abide by these Guidelines.

**Signature ………………………………………………………....**

**Name ………………………………………………………………………..**

**Position ……………………………………………………………………..**

**Institute Address**

**…………………………………………………………………………………………**

**…………………………………………………………………………………………**

**…………………………………………………………………………………………**

**E-mail ……………………………………………..**

**Phone and Fax Numbers ………………………………………………..**

**Date …………………………..**

**Approval by CRPS UK Registry Steering Committee**

The Steering Committee has agreed to approve this project access to the CRPS-UK Registry data requested:

**Steering Committee Chairman: …………………………………….**

**Date: …………………………..**

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**Data access form**

|  |  |  |
| --- | --- | --- |
|  |  | Please expand boxes, if necessary |
| 1 | Study title: |  |
| 2 | Primary hypotheses: |  |
| 3 | Please provide brief background and protocol for proposed analysis: | *May be provided on separate sheet.* |
| 4 | Please list all personnel involved in the study. |  |
| 5 | What data do you require? *Please include timescales for recruitment and study end date.* |  |
| 6 | What is the timescale for analysis? |  |
| 7 | Has funding and / or ethics been obtained? |  |
| 8 | Has this protocol been peer-reviewed? | *If so, please indicate where.* *Please list the reviewer(s), if known.* |

**Upon completion of analysis, a report will be required to be submitted, regardless of whether publication is anticipated.**