**Yorkshire and Humber Secure Data Environment – Data Availability Form**

**Part 1: Study overview (Mandatory)**

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| --- | --- |
| **Requestor name:** |  |
| **Requester organisation:** |  |
| **Requester’s role:** |  |
| **Requester’s contact email:** |  |
| **Study name:** |  |
| **Disease areas(s):** |  |

|  |
| --- |
| **Study overview and objective(s) (max 500 words):** |
|  |

**Part 2: Data priority uses**

*Please indicate which of the following priority uses cases/ stage of the patient access journey will be addressed by the proposed work (tick all that apply):*

|  |  |  |  |  |  |
| --- | --- | --- | --- | --- | --- |
| **AI/algorithm development** | ☐ |  | **Discovery and development** | ☐ |  |
| **Clinical trial activities** | ☐ | **HTA/ Regulatory** | ☐ |  |
| **Real world studies** | ☐ | **Post launch / in market** | ☐ |  |
| **Translational research** | ☐ |  |  | |
| **Epidemiological studies** | ☐ |  |  | |
| **Health systems research** | ☐ |  |  | |

**Part 3: Data Requirements**

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| **Specific fields / data items required** |
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| **Cohort inclusion and exclusion criteria** |
|  |
| **Time period of interest** |
| *Please indicate if this study relates to patients that underwent treatment and follow-up during a specific period of time, e.g. 2018 to 2022.* |
|  |
| **Indicative patient counts / cohort sizes** |
| *Please list any sub-cohorts and minimum size required if known* |
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