**Florence - Case Study Template**

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| **Case Study Theme/Protocol** |
| Clinical service that the case study relates to:Aim:Target cohort: |
| **Submission Organisation**  |
| Name: |  |
| Address: |   |
| **Case Study Essential Criteria:** |
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| *1.* | Patient consent details: the intended use of anonymised information i.e. reports/publication/conference use *(please highlight response)* | *Yes* | *No* |
| 3. | Lead IndividualDetails: | *Name :*  |  |
| *Role/Position:*  |  |
| *Contact Tel Nr:* |  |
| *Email:* |  |
| 4. | *Please provide a brief overview of your case study highlighting the aspects you feel are most relevant:* |

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| **Case Study Desirable Criteria:** |
| 1. | Patient approval of case study: patient proofed the case study before signing consent *(please highlight response)* | *Yes* | No | N/A |
| 2. | *Please provide a brief description of the clinical and non-clinical outcomes as well as structure & process related to the case study:* |
| 3. | *Please provide a brief description of any proxy measures and immediate clinical objectives (eg lowering BP) related to the case study:* |
| 4. | *Please provide a brief description of how supporting literature (if you know it exists) shows the potential longer term impact of faster/better outcomes:* |
| ***Approach to Show Casing the Case Study*** |
| *Please briefly describe how you propose to show case this case study:* |