*A brief final study report should be sent to the Ethics Committee and the MHRA (if applicable) within 12 months of the end of the study. Note: The end of the study is defined in the study protocol.*

*All instructions are highlighted in yellow and should be deleted when the report is finalised. This template is for both clinical trials and clinical research. Delete sections that are not applicable.*

**FINAL STUDY REPORT**

**Study Title:** *Insert*

**REC Ref/ CTA No:** *Insert*

**Chief Investigator:** *Insert*

**Sponsor:** *Insert*

|  |  |
| --- | --- |
| **List of Principal Investigators and Sites** |  |
| **List of Publications (or plans for publications) including those for patients (if applicable)** |  |
| **Study Start and End Dates** |  |
| **Study Design** |  |
| **No. of Patients (planned and analysed)** |  |
| **Main inclusion/exclusion criteria** |  |
| **Investigational Medicinal Product(s) (including comparator, if applicable), mode of administration and batch number(s)** |  |
| **Duration of Treatment** |  |
| **Primary and Secondary Objective(s)** |  |
| **Endpoints/ Outcome Measure(s)** |  |
| **Statistical Methods** |  |
| **Conclusions**  |  |

**Authorised by: \_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_**

**Signature: \_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_**

**Date: \_\_\_\_\_\_\_\_\_\_\_\_\_**