|  |  |  |  |  |
| --- | --- | --- | --- | --- |
| 1. | JKEUPM Ref. No. |  | | |
| 2. | Study Title |  | | |
| 3. | i. Principal investigator  a. Name  b. Address  c. Tel.No  d. Email  ii. List of co-investigators |  | | |
| 4. | Name Of Funding Agency |  | | |
| 5. | Study Site |  | | |
| 6. | Total number of eligible subjects in study site |  | | |
| 7. | Recruitment of subjects in study site (*include CONSORT flowchart for RCT studies only*)   1. Number of participants recruited: 2. Number of participants completing trial/ study: 3. Proposed in original application: 4. Number of withdrawals from trial to date due to: 5. withdrawal of consent 6. no response from participants 7. loss to follow-up 8. death (not the primary outcome)   Total study withdrawals:  v. Number of treatment failures to date (Prior to reaching primary outcome) due to:   1. adverse events 2. lack of efficacy   Total treatment failures: |  | | |
| 8. | Duration of study |  | | |
| 9. | Protocol Violation or Deviation |  | | |
| 10. | Executive summary *(Summary of research background, objectives, methodology, findings (include a table if applicable) and conclusion of the research project) - maximum 500 words)*  \*Committee may request additional information if required. |  | | |
| 11. | Signature of Principal Investigator |  | Date |  |