RAKMHSU/RAK Research & Ethics Committee

 Ras Al Khaimah, UAE

Date: DD/MMM/YYYY

**Faculty Research Application**

**(Please note in students research proposals, faculty/staff can be supervisors and/or co-investigators)**

Name of Coordinating/Principal Investigator:

Department:

Phone number:

Email address:

Title of the project:

|  |  |  |  |
| --- | --- | --- | --- |
| Key personnel on the project | Institution | Study Role | Contact details (Phone/email) |
|   |  |  |  |
|  |  |  |  |
|  |  |  |  |

Have you applied for funding for the project? Yes No

If “Yes”, indicate sourse(s) of funding:

Has this research undergone external scholarly review (e.g by granting agency)? Yes No

**If “Yes,” specify results of review:**

**Proposed start date of your research:**

**(Consider 45-60 days for the review process to complete):**

**Faculty: RAKCOMS/RAKCODS/RAKCOPS/RAKCONS/RAKMHSU**

**Principal Investigator Signature**

**Forwarded by Dean (Signature)**

For Research Ethics Committee Use Only:

|  |
| --- |
| RAKMHSU REF NO.Date of submission:Date of Approval: |

RAKMHSU/RAK Research & Ethics Committee

 Ras Al Khaimah, UAE

Research Proposal

Title:

Investigators:

Aims/Objectives

Research question/Research issue:

Plain Language SUMMARY of project

(2-3 sentences using language that is understood by a non-medical person)

Background knowledge (Need for the study):

Methodology:

Duration:

Study setting:

Study design:

Study population:

Sampling method:

Sample size:

Selection method:

Inclusion criteria:

Exclusion criteria:

Study materials: Equipment, questionnaire, data collection forms.

Data collection procedure: Methods used to collect the data

Access issues (relating to patients or their medical records):

Proposed data analysis (This should clarify what the outcomes of interest are, and what comparisons will be made, using which statistical test):

Scope of the study:

Limitations/ expected challenges:

List of REFERENCES: Use Vancouver or APA reference style.

Ethical Considerations

1. How will participants be recruited or cases be identified?
2. How will informed consent be obtained from participants?
3. What procedures will be followed to maintain confidentiality and privacy of participant’s data?
4. Provide details of who will have access to the data.
5. Does the project involve any of the following risks? Indicate yes or no.

|  |  |  |
| --- | --- | --- |
|  | Yes | No |
| * 1. Physical risk to participants
 |  |  |
| * 1. Psychological risks to participants
 |  |  |
| * 1. Does the study require participants to release information of a sensitive or personal nature?
 |  |  |
| * 1. Are there medical conditions, which increase the participants risk while in the study?
 |  |  |
| * 1. Does this study require participants to release information, which may reveal illegal activity?
 |  |  |
| * 1. Are there any risks to the researchers?
 |  |  |
| * 1. Any other risks?
 |  |  |

Detail in this area explanations of an answer of “yes” to any of the above

1. In undertaking this research, do any “conflict of interest” issues arise? Yes/No
	1. Are the participants in a dependent relationship with the investigators?

Yes/ No

* 1. Are the participants staff at the institution where the study is taking place?

Yes/ No

1. Does each member of the research team have ICH-GCP Training?