## RESEARCH IDENTIFICATION FORM

**Project title:**

#### \_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_

#### Institution where the project will take place:

#### \_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_

#### Authors: \_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_

#### E-mail and Telephone number for researcher(s):

#### \_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_

#### Address to send material:

#### \_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_

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| --- |
| **Project Links:** |
| ( ) Post Doctorate |
| ( ) Doctorate |
| ( ) Masters |
| ( ) Post graduation (Specialization) |
| ( ) Taught Post-Graduate Courses (linked to the line of research).What? |
| ( ) Study Group/Identifier: |
| ( ) Research Group/Identifier: |
|  |
| **É bolsista?** / **Is it a fellowship?** |
| ( ) yes ( ) no |
| ( ) FAPESP (São Paulo Research Foundation) |
| ( ) CNPQ (National Counsel of Technological and Scientific Development) |
| ( ) CAPES (Coordinator for the Improvement of Higher Education Personnel) |
| ( ) Other (please state which): |

**Request for Material:**

| Quantity | Material Description | Material Code |
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**[INSTITUTION]**

**[/TITLE]**

[Authors]

**[YEAR]**

**Summary**

1. Abstract and Key words
2. Introduction
3. Study objectives (general and specific)
4. Hypothesis
5. Methodology
6. Timeline
7. Budget
8. Final considerations
9. References
10. Annexes

**Abstract and keywords**

It should contain between 150 and 300 words and provide relevant information on the purpose and methodology. Must be typed in a justified manner and in spacing 1,5 Times New Roman 12 or Arial 11 font. It should be written in single paragraph. The use of bibliographic citations should be avoided.

Keywords: Between 3 and 5 words (according to the MeSH vocabulary list (Medical Subject Headings) **-** <https://www.nlm.nih.gov/mesh/MBrowser.html>)

**Introduction**

It should represents the essence of the author's thinking in relation to the subject he intends to develop. Its purpose is to provide the background literature that justifies the study elaboration.

**Objectives**

It should be described in the form of General Objective (clearly defines the study guidelines) and Specific Objectives (represent the hypotheses of the study).

**Methodology**

It should be described in detail aiming that other researchers can accurately evaluate their study proposal and be able to repeat the suggested methodology.

It may be divided into topics according to the preference of the researcher responsible, however it is imperative that you inform, in the case of clinical studies:

• Study design;

• Sample size;

• Inclusion and exclusion criteria;

• Risks and benefits for participating in the sample;

• Variables evaluated;

• Methodology of data analysis;

• Primary and secondary outcome;

• Statistical planning.

**Study Timeline**

It should identify the activities necessary to carry out the study specifying the number of months needed to carry out the study.

Example:

|  |  |  |  |  |  |  |  |  |  |  |  |  |
| --- | --- | --- | --- | --- | --- | --- | --- | --- | --- | --- | --- | --- |
| **Activities Description** | MONTHS | | | | | | | | | | | |
| **1** | **2** | **3** | **4** | **5** | **6** | **7** | **8** | **9** | **10** | **11** | **…** |
| Project Elaboration |  |  |  |  |  |  |  |  |  |  |  |  |
| Litarature Research |  |  |  |  |  |  |  |  |  |  |  |  |
| Data Collection |  |  |  |  |  |  |  |  |  |  |  |  |
| … |  |  |  |  |  |  |  |  |  |  |  |  |
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**Budget**

You should specify the materials, with their values, that will be necessary to carry out the study.

**References**

Identification of the documents consulted. They should follow the Vancouver style. Standards consultation: <http://www.nlm.nih.gov/bsd/uniform_requirements.html>

Example:

Patrias K. Citing medicine: the NLM style guide for authors, editors, and publishers [Internet]. 2nd ed. Wendling DL. technical editor. Bethesda (MD): National Library of Medicine (US). 2007 [citado 2007 Jan 10]. Disponível em: <http://www.nlm.nih.gov/citingmedicine>

Guimarães CA. Normas para manuscritos submetidos às revistas biomédicas: escrita eedição da publicação biomédica (tradução integral do texto). Rev Col Bras Cir. [Internet]. 2006 Out [citado 2008 Jan 11];33(5):318-35. Disponível em: <http://www.scielo.br/pdf/rcbc/v33n5/v33n5a13.pdf>

Rother ET, Braga MER. O novo estilo de Vancouver: o que mudou nas referências. Arq Bras Oftalmol. [Internet]. 2004 Ago [citado 2008 Jan 11];67(4):692-4. Disponível em: <http://www.scielo.br/pdf/abo/v67n4/21423.pdf>.

Universidade Federal do Paraná. Normas para apresentação de documentos científicos. Sistema de Bibliotecas. 2.ed. Curitiba: Ed. UFPR, 2007. 9v: il.

Hulley SB, Cumming SR, Browner WS, Grady DG, Newman TB. Designing clinical research. Philadelphia: Lippincott Willian & Wilking. 2007.

**Annexes**

Assessment sheets, Questionnaires, or other instruments needed to conduct the study. In the case of clinical studies in humans, the "Free and Informed Consent Term" should be included.