Author guidelines

Archivos Argentinos de Pediatría is the official publication of the Sociedad Argentina de Pediatría SAP) and has been issued uninterruptedly since 1930. Its publication is bimonthly. It is part of the SciELO network and of the Core Collection of Argentine Scientific Journals. It is indexed in Medline, the Science Citation Index Expanded, Active Embase Journals and LILACS. Openaccess full-text articles can be found at the website of the Sociedad Argentina de Pediatría (www.sap. org.ar/publicaciones/archivos.html) available online since 1970, or at the SciELO (Scientific Electronic Library Online) Argentina website (www.scielo. org.ar).

Archivos Argentinos de Pediatría follows the Recommendations for the Conduct, Reporting, Editing and Publication of Scholarly Work in Medical Journals of the International Committee of Medical Journal Editors (ICMJE).

In relation to ethical standards, Archivos follows those established by the Committee on Publication Ethics (COPE).

Archivos Argentinos de Pediatría publishes articles related to perinatal health, child and adolescent health, and to other relevant fields of medical practice.

Editors reserve the right to publish accepted articles in the printed or just in the electronic version of Archivos Argentinos de Pediatría.

Review process

Archivos Argentinos de Pediatría is a peerreviewed scientific journal. All original articles, special articles, brief reports, clinical case presentation, and any other document deemed relevant by editors are sent to at least two independent reviewers. The identity of authors and reviewers is kept confidential. The editorial process of submitted articles can be followed online through the link http://archivos.sap. org.ar/index.php/AAP, with the username and password created at the time of generating the manuscript submission.

INSTRUCTIONS FOR AUTHORS Terms and conditions for publication

Articles must be unpublished. Prior publication is only accepted as a summary in conference proceedings or other scientific meetings or as a preprint.

Pre-publication of a primary research manuscript is the deposit on a preprint server, institutional or author websites, and open communications between researchers, either on community preprint servers or comment platforms before formal peer review in a journal.

Pre-publication will not be a disqualifier for submission. When submitting a manuscript for review, authors should mention the details of the preprint, including the DOI and license terms. If accepted for publication, the authors are requested to update the pre-publication version with the DOI and a URL link to the published version of the article on the journal website.

Preprints can be cited in the reference list of articles submitted for review.

Authors who pre-publish must adhere to the media communications policy. They can provide an explanation or clarification of the paper or information about its context. In these circumstances, media coverage will not hamper the editorial process. Researchers should make it clear that the article has not yet undergone a peer review, that the findings are provisional, and that the conclusions may change.

Articles by international organizations (WHO, PAHO or other similar entities) will be published if their free reproduction is specifically authorized.

Articles already published in other journals will be considered if editors find it is justifiable. In such case, authors will require the approval of editors of both journals.

Detection during editorial and peer review of duplication of manuscripts, plagiarism, selfplagiarism and fragmentation of the results of an investigation will be considered ethical misconducts. This detection will be carried out through the search for previous publications on the same subject or other similar publications of the authors.

How to submit articles

All articles should be sent through the link to http://archivos.sap.org.ar/index.php/AAP, with the username and password created at the time of generating the manuscript submission.

When a manuscript is accepted for publication, authors must pay an article processing charge.

Files attached should be in MSWord (.doc or .docx) format, and include:

- 1. The manuscript written as per the instructions detailed below, and the file name should be the main author's last name.
- 2. A Letter to the Editor including the name, telephone number, e-mail address and signature of all authors, requesting the evaluation of the article for its potential publication. The letter should clearly state the following:
- The article submitted has not been published in any other medium and shall not be submitted to a different scientific journal or any other publication for as long as *Archivos Argentinos de Pediatría* is evaluating it.
- All authors should state whether they have any disclaimers to make regarding conflicts of interest. If there is any conflict of interest, authors shall declare such interest or financial commitment in the article (see below).
- All sources of external funding should be clearly detailed. If the study did not receive external funding, this should be clearly stated.
- Compliance with the checklist should be indicated before submitting the material.
- It should be stated that if the article gets published, all authors will transfer their copyrights to the Sociedad Argentina de Pediatría.

No editorial process shall be initiated unless the letter complies with all items indicated above.

GENERAL CONSIDERATIONS FOR MANUSCRIPTS PREPARATION Writing

Manuscripts should be written using a word processor (MSWord), **double-spaced** (including abstracts, references and tables), font: Arial 11. Pages should be **numbered** starting on the cover page in the lower right-hand corner.

First page or cover page. It should include:

Title: in Spanish and English or just in English if the manuscript is written in English. The extension will not exceed 22 words. It should not contain acronyms and abbreviations and be consistent with the manuscript content.

Author(s): first name(s) and last name(s) in a correlative order and with their academic degree indicated in an abbreviated form (M.D., B.S., PhD, DMSc, etc.) with an asterisk to identify the institution to which they belong; name, address and telephone number of the institution

to which the first author belongs. Each of the authors' e-mail address should also be included. The author responsible for receiving the editors' notifications should be indicated separately.

In addition, the participation of each author in the study should be detailed, as required by the International Committee of Medical Journal Editors (ICMJE) in the Recommendations for the Conduct, Reporting, Editing and Publication of Scholarly Work in Medical Journals available in: http://www.icmje.org/icmje-recommendations.pdf

Authors should fully meet the following criteria for authorship:

- Substantial contributions to: the conception or design of the work; or the acquisition, analysis, or interpretation of data for the work.
- Drafting the work or revising it critically for important intellectual content.
- Final approval of the version to be published.
- Agreement to be accountable for all aspects of the work in ensuring that questions related to the accuracy or integrity of any part of the work are appropriately investigated and resolved.

All contributors who do not meet all the criteria for authorship might be listed as Collaborators or cited in the Acknowledgments section.

When a manuscript has been written by a Working Group or members of a multicenter study, the corresponding author should indicate the order in which authors will be mentioned, identify all individual authors and the name of the group. The rest of the group members who do not meet authorship criteria may be listed as Collaborators or cited in the Acknowledgments section.

In the case of multicenter studies, researchers who were not directly involved in drafting the article but who participated in the study development can be included in the Contributors section.

In situations where authorship is not clear enough, editors can directly request the responsible author or a third party to fully comply with the indicated requirements.

If after such request, one or more authors do not meet the authorship criteria, the article will not be accepted for publication.

Subsequent pages

• Ethical-regulatory: As stated in the

Declaration of Helsinki (paragraph 23), all medical research studies in human beings, regardless of experimental or observational nature, including investigation of human material and identifiable information must be submitted for consideration, comment, guidance and approval of the corresponding Ethics Committee before starting the study. The publication of a case/case series report requires, in accordance with current legal provisions, requesting the informed consent for publication, of the patient or their legal representatives, in safeguarding against possible claims or sanctions. Argentine law 26529 on the Rights of the Patient in their Relationship with Health Professionals and Institutions, refers to the right to confidentiality: The patient has the right that any person who participates in the preparation or manipulation of clinical documentation, or has access to the content of the same, keep the due reserve, unless expressly provided otherwise emanating from the competent judicial authority or authorization of the

patient himself. The confidentiality of the patient's data refers to each of the data that allows identification by the place of origin, clinical picture, studies carried out, institution of care, treatment and evolution (see the informed consent form). This is essential to begin the process of reviewing a manuscript submitted to Archivos Argentinos de Pediatría.

- **Grammar and style.** The rules of the language used for writing should be followed and a style appropriate for scientific writing should be applied. The wording and style should be carefully reviewed before the manuscript submission if possible by a person specially trained for this role.
- **Acronyms and abbreviations.** They should be kept to a minimum and only those usually accepted will be used; it has been demonstrated that excessive acronyms can be confusing to readers. The spelled-out abbreviation followed by the abbreviation in parenthesis should be used on first mention in the manuscript. No acronyms should be used to refer to a single term (e.g., ATB instead of antibiotic). No acronyms should be included in titles and in the abstract written in Spanish (unless strictly justified). Manuscripts in English may

include the acronyms generally used in English publications. If acronyms are used in tables, figures or graphics, they should be written in full at the bottom or in the epigraph, even though they are written in full in the text. In manuscripts that require **5 or more** acronyms

(biochemical determinations, genetic studies, endocrinological studies or similar) it is convenient to place a list of Acronyms and abbreviations after the Abstract, with the meaning of these acronyms, to facilitate the reading.

- Acknowledgments. People or institutions contributing with the article in any way can be included under this heading. Acknowledgement should be written in a formal style and in a separate page from the manuscript.
- **References.** It will be numbered in the order it appears in the text as per the style proposed by the Recommendations for the Conduct, Reporting, Editing and Publication of Scholarly Work in Medical Journals We also recommend to visit the website *Citing Medicine*, by the US National Library of Medicine (http://www.nlm.nih. gov/citingmedicine), as a reference for bibliographic citations.

References should strictly comply with the guidelines of the following examples:

a. Medical Journals

Salhab WA, Wyckoff MH, Laptook AR, Perlman JM. Initial hypoglycaemia and neonatal brain injury in term infants with severe fetal acidemia. Pediatrics. 2004;114(2):361-6. For articles with more than four authors, "et al." should be added after the name of the fourth author.

b. Textbooks

Bradley El. Medical and surgical management. 2nd ed. Philadelphia: W B Saunders; 1982:72-95.

c. Textbook chapter

Stanley F, Blair E, Alberman E. How common are the cerebral palsies? In Bax MC, Hart HM, eds. Cerebral Palsies: epidemiology and causal pathways. London, United Kingdom: Cambridge University Press; 2000:22-39.

d. Journal article published in internet Katheria A, Poeltler D, Durham J, Steen J, et al. Neonatal Resuscitation with an Intact Cord: A Randomized Clinical Trial. *J Pediatr.* 2016 Nov; 178:75-80 e3. doi: 10.1016/j peds 2016.07.053.Epub 2016.Aug26.

e. Internet sites

OPS/OMS. Situación de Salud en las Américas: Indicadores Básicos 2005. Washington DC, 2005. [Accessed on: June .21, 2017]. Available at: http://www.paho.org/spanish/dd/ais/IB-folleto-2005.pdf

- Tables and figures. They should be included in a separate page, one per page and ordered using Arabic numbers. They should have a title on the top, next to their number, and a footnote. Measurement units should be referred to in the text, and in tables and figures.
- **Pictures.** If pictures of patients will be included, efforts should be made to ensure the patient's anonymity. They should be accompanied by the informed consent for publication. No patient identifying information should be present in microscopic observations or imaging studies. If illustrations by other authors are used, whether published or unpublished, the corresponding authorization for reproduction should be attached. Small arrows in a contrasting color should be used for easy identification of what is shown in the picture. Epigraphs or footers should be in a separate page numbered sequentially. Digital images should be legible with a resolution of at least 300 dpi, and in a .jpg format. If additional information is required, please contact the editorial secretary.

SPECIFIC CONSIDERATIONS FOR PREPARING THE MANUSCRIPT

I. Original articles

Original articles refer to a research subject to a specific design, such as: randomized clinical studies, cohort studies, case-control studies, crosssectional studies, epidemiological evaluations, surveys and systematic reviews. Each study component or section has to be submitted in a separate page in the following order:

First page or cover page

It includes the items indicated in "General considerations for manuscripts preparation", and the following aspects should be taken into account:

- Title: it should be brief (no more than 22 words), contain no abbreviations or acronyms and reflect the research objectives and design.
- Authors: see the authorship conditions in "General considerations for manuscripts preparation." All the authors have made major contributions to the design and development of the study, the analysis of results and the manuscript preparation; they approve the final version and are accountable for all aspects of the work.
- Clinical trial registration: Archivos Argentinos de Pediatría adheres to the policies of the World Health Organization (WHO) and the International Committee of Medical Journal Editors (ICMJE), which recommend the registration of clinical trials in a Public Registry of Investigations before the inclusion of the first patient, as a condition to start the review process. It also adheres to the postulates of the Declaration of Helsinki in the sense that "any research study involving humans must be registered in a database accessible to the public before recruiting the first subject".

This is particularly relevant to the investigations that contemplate the prospective assignment of persons or groups of people to an intervention, with or without comparison with a control group, to study the relationship between the intervention and the results.

The registration of clinical trials is mandatory in all randomized clinical trials and intervention studies and also, desirable for observational studies.

Interventions are considered any measure used for the purpose of modifying a biomedical result, related to health or behavior (including, but not limited to, medications, surgical procedures, devices, behavioral treatments, educational programs, dietary interventions, interventions to improve quality and changes in care processes).

The name of the Registry, registration number and inscription date should be placed in the manuscript after the Keywords. Registration can be done in an Argentine or international Registry of Clinical Trials (see the addresses available in the list at the end of this Author Guidelines). The review process can not be started if it is not sent the registration number for randomized control trials.

• Funding: indicate all sources of funding and

- the type of contribution made by each of them.
- **Conflicts of interest:** a conflict of interest is when the author, any of his/her relatives, or the institution where he/she works has a financial relationship (employment, counseling, free or fee-for-advice, reports, specialist's report) or a personal relationship with the commercial company that produces the product(s) used or mentioned in the study. A more conceptual definition by the US Institutes of Health reads: "a conflict of interest is a set of circumstances that creates a risk that professional judgment or actions regarding a primary interest will be unduly influenced by a secondary interest" (Institute of Medicine of the National Academies Website. Conflict of interest in medical research, education and practice).
- Sponsor: when a study is sponsored, it should be indicated if the sponsor had any participation in the study design, data collection, discussion, interpretation, manuscript writing or decision to send it for publication.

Abstracts

The second page will include the abstract. If the manuscript is written in Spanish is not required an English abstract.

Characteristics of abstracts:

- Length: the abstract should not exceed 250 words.
- **Key words**: up to **five (5)** key words at the end of each abstract. Use the terms indicated in the Pubmed's Medical Subject Headings (MeSH).
- **Structure:** include the following subheaders:
- Introduction: briefly indicate current knowledge on the research subject and research objectives.
- *Population and methods:* indicate where and when the study was conducted. Mention study design and the studied population. Define the primary outcome measure. Indicate the intervention provided and population assignment (if applicable).
- Results: provide details on the primary outcome measure and describe the rest of the results obtained and their statistical significance.
- Conclusions: provide specific conclusions in relation to proposed objectives and obtained results.

Manuscript writing

The manuscript will have a maximum length of 2500 words (excluding abstracts, acknowledgments references, tables and figures). It will have the following format:

- **Introduction**: provide the study rationale, describe the reasons for conducting the study and any existing unanswered question and point out related studies. In addition, include the research problem (question) together with a theoretical framework supported by bibliography. Do not include data or conclusions of the study being reported.
 - At the end of the introduction, include the research objectives (the purpose of the work and the target population).
- Population and methods: include the following items:
 - Date and time of study conduction and the characteristics of the institution or of the main coordinating site for multicenter studies.
 - **Design**: cursorily indicate the type of design: case series, cross-sectional, casecontrol, cohorts (type), clinical trial (type), systematic review, etc.
 - **Hypothesis**: if applicable, include the research hypothesis.
 - Aspects to be covered: indicate eligibility, inclusion, exclusion and discontinuation criteria applicable to the study population. If randomization processes have been used, indicate the randomization technique. Define the primary and secondary outcome measures in detail, the techniques used for measuring such outcomes and the intervention provided (if applicable).
 - Ethical considerations: all clinical trials should specify if a written informed consent was obtained, the name of the Ethics Committee and the Research Committee, and the date of protocol approval.
 - Statistical analysis: indicate statistical testing and provide enough details so that data may be verified by other researchers, and provide a rationale for using each test. Specify the name of the statistical software used for data processing. Indicate the number of patients calculated for the study (sample size).
- **Results:** describe how many patients were eligible and how many were finally included (patient flow). Specify the main characteristics of the studied population. Provide data of

obtained results. Avoid using the same wording for writing the text regarding data included in a table.

The results included should be related to the objectives and based on the statistical analysis indicated in the Population and Methods section. All central tendency parameters should be accompanied with the corresponding dispersion measure (mean and standard deviation, median, range and mode, percentage, and confidence intervals). It is preferable that the exact value of the significance test is reported so that readers are able to compare it with other p values (instead of p < 0.05, p = 0.03).

- **Discussion:** this section is used for stating comments on the study and comparing results with those obtained by other authors. Here speculations can be made and a new hypothesis can be developed based on the research. Results should not be repeated here. It is important that the study's strengths and weaknesses are pointed out.
- Conclusion(s): these will be brief, accurate statements with no speculations, related to the research objective and based on the results obtained.
- References: the maximum number of citations will be 40 and they should be written according to the specifications indicated in "General considerations for manuscripts preparation."
- Tables and figures: the maximum number will be six (6), and they will be prepared according to the above mentioned specifications. No more than two pictures can be included. If editors consider that it is reasonable to have a higher number of tables, figures, pictures or annexes, these will only be published in the electronic format version.

II. Brief reports

These are descriptions of observations, presentation of clinical situations, preliminary results, medical technology, procedures or other interesting aspects.

Preliminary results are relevant to the purpose of the study but cannot be generalized, either by sample size or other conditions that affect the representativeness of the sample. Also should be considered as preliminary, the results of studies in which the outcome addresses partly to the objective.

Brief reports are also considered studies that even with proper design, are relevant only to the place where the research was conducted.

The registration of clinical trials is mandatory, as specified in the original articles.

The manuscript for brief reports will be written and submitted similarly to what is indicated in "General considerations for manuscripts preparation."

The **structure** will be as follows:

- **Abstract**: if the manuscript is written in Spanish is not required an English abstract. If the manuscript is written in English is not required a Spanish abstract. The maximum length is **150 words**. The abstract does not need to follow a structure. It should include up to **five (5)** key words.
- **Description:** it will have a maximum length of **1400 words** (excluding abstracts, references and tables or figures), and no more than **four (4)** illustrations (tables, graphics, or pictures). Write a brief **introduction** pointing out the relevance of the subject and similar experiences already published. Then describe the methodology of the study, the results found and finally write a discussion or comment.
- References: the maximum number of citations will be 15, in accordance to the instructions provided above.

III. Case reports

Case reports should be accompanied by the Consent for Publication of Personal Information. They refer to the presentation of patients or series of patients with a rare disease or an uncommon clinical condition, whose description is relevant for pediatric practice and complies with at least one of the following criteria:

- Drug side effects or adverse interactions that are rare or have not yet been reported.
- The occurrence of a rare disease.
- New associations or variations in the process of a disease.
- Presentation, diagnosis and/or treatment for a new or emerging disease.
- Unexpected association between diseases or symptoms.
- An unexpected event observed in the clinical course of a disease or its treatment.
- Results or findings that shed new light on the possible pathogenesis of a disease or adverse effect.

The structure of Case reports will be similar to that of Brief reports. The abstract will be written in Spanish and English if the manuscript is written in Spanish. When the manuscript is written in English is not required a Spanish abstract.

In order to be considered the author of a Case reports, the author should have contributed to the idea, intellectual development, writing and review of the report. Having provided care to the patient presented in the report is not a criterion of authorship. This can be mentioned in the Acknowledgments section.

IV. Review articles

These are complete and comprehensive reviews or updates regarding an important issue.

They should point out the advances made in recent years and, if they refer to a disease, they should highlight the clinical, diagnostic, therapeutic and prognostic aspects.

Review articles are usually requested by editors, but unrequested articles of real interest can also be submitted if they follow the indicated guidelines. In these cases, their acceptance will be at the editors' discretion.

Review articles will be written and submitted similarly to what is indicated in "General considerations for manuscripts preparation."

The **specific characteristics** for reviews are as follows:

- Abstract: it will have a maximum length of 150 words, pointing out only the most important aspects of the subject. The abstract does not need to follow a structure. It should include up to five (5) key words.
- **Text**: it will include an introduction, the development of the different aspects and, if applicable, a discussion or relevant comments can be included. The text will have a maximum length of 3000 words (excluding abstracts, references and tables or figures).
- **References:** it will be as thorough as possible, and there is not a maximum number of citations provided that citations included are actually relevant for gaining knowledge on the subject or becoming aware of other authors' experience and, also, for an easy search.

V. Special articles

Special articles have certain differences from the other articles mentioned above. These usually cover especially relevant subjects, which may or may not refer to a specific disease or a particular situation or aspect. The subject may refer to other

disciplines, whether or not related to medicine, but that do concern the human being (philosophy, ethics, anthropology, sociology, etc.). In addition, special articles may cover topics related to research methodologies and technologies, among other issues. The manuscript for special articles will be written and submitted similarly to what is indicated in "General considerations for manuscripts preparation". The specific characteristics of the abstract, text and references will be similar to those of the review articles. but they may have a different structure. The maximum length will be 3000 words.

VI. Comments

There are two types of comments: editorial comments and comments on different topics of interest. Editorial comments usually refer to an article presented in the same journal issue, and are made by request of the editors. Comments on different topics of interest are also usually requested by editors, but they may be freely submitted. In these cases, their acceptance will be at the editors' discretion.

Both types of comments will be published in the same section and will have a maximum length of 1300 words (excluding references). References: the maximum number of citations will be 10.

VII. Practical pediatrics

These will be articles regarding experiences or relevant topics that earn knowledge on the practical aspects of pediatric practice. The manuscript for these articles will be written and submitted similarly to what is stated in "General considerations for manuscripts preparation."

It will have an abstract with a maximum length of 150 words and up to five (5) key words. The manuscript will have a **maximum length** of 2200 words (excluding abstracts, references and tables or figures), but editors may consider a longer article depending on the case. References: the maximum number of citations will be 25.

VIII. Other sections

The manuscripts focusing on medical education, health and social pediatrics and epidemiology, will have structural characteristics similar to those of Practical pediatrics.

IX. What is your diagnosis?

This text will have two parts: the first part will be a brief description of not more than 200 words regarding a clinical condition with five diagnostic options. One (1) figure or picture can be included. The second part will have a brief description of the clinical condition with a maximum length of 100 words, and a description of the disease or disorder regarding the correct diagnosis and differential diagnosis. This will have a maximum length of 800 words, with not more than two (2) illustrations (figures or pictures) and a maximum of five (5) references.

The articles in this section are usually requested by editors, but they can also accept those not requested and their acceptance will be at the editors' discretion.

Informed consent is required for the publication of the clinical case.

X. Letters to the editor

Letters to the editor will refer to a published article or to any other topic of interest, and can include recommendations and criticisms, always in a respectful style. They should be addressed to the editor and have a title. They will have a maximum length of 1000 words, with up to 5 references.

XI. Books

Books reviews are requested by editors and will have a **maximum length of 800 words**. They usually provide a brief description of the characteristics of a book that the editors consider being especially interesting for medicine and other disciplines.

Archivos Argentinos de Pediatría may publish other types of articles and add sections at the editors' discretion.

The Editorial Direction reserves the right to reject any article that does not strictly meet the instructions herein indicated or whose topic does not match the magazine's profile.

The authors will be exclusively responsible for the content and comments of the article.

The Sociedad Argentina de Pediatría will be the copyright holder for any of the articles published, which cannot be reproduced in any other journal, whether partially or in full, without their corresponding authorization.

CHECKLIST FOR SUBMITTING A MANUSCRIPT

Carefully review if the following steps have been met:

- 1. Carefully read "Terms and conditions for publication" and "General considerations for manuscripts preparation."
- 2. For **writing the letter** requesting the publication of the article, indicate the following:
 - The article submitted has not been published in any other medium and shall not be submitted to a different scientific journal or any other publication for as long as Archivos Argentinos de *Pediatría* is evaluating it.
 - All authors should state whether they have any disclaimers to make regarding conflicts of interest.
 - All sources of external funding are detailed.
 - Indicate compliance with the checklist before submitting the material.
 - If the article gets published, all authors will transfer their copyrights to the Sociedad Argentina de Pediatría.
- 3. In the **first page or cover page**, include the following items:
 - Title.
 - Authors: first name(s), last name(s) and degree indicated in an abbreviated form.
 - The author who will receive editors' notifications.
 - Detail the participation of each author in the study.
 - Name, address and telephone number of the institution where the study was conducted.
 - Registration number of randomized studies with intervention in an Argentine or international Registry of Controlled Clinical Trials (see list).
- 4. The abstracts should comply strictly the indications for each type of article about extension and number of key words.
- 5. An **adequate manuscript presentation**, will be in strict compliance with what is specified in the "General aspects for preparing the manuscript" for each type of article in terms of length, structure, references, tables or figures.
- 6. A copy of the Informed Consent for Publication of personal Information including medical data, pictures, images, etc. or the approval of the Ethics Committee when appropriate.

If additional information is required, please contact the publication office at 54-11 4821-8612, ext. 123/143, Monday through Friday, from 11 AM to 8 PM, or send an email to publicaciones@sap.org.ar.

This Guideline will be in force: as of May 2021

LIST OF CLINICAL TRIAL REGISTRIES

Argentina (among others):

• National Register:

Registro Nacional de Investigaciones en Salud (RENIS) - Sisa https://sisa.msal.gov.ar/sisa/sisadoc/docs/050104/renis intro.jsp

• City of Buenos Aires:

Registro de proyectos de investigación. Comité Central de Ética en Investigación, Ministerio de Salud http://www.buenosaires.gob.ar/salud/docenciaeinvestigacion/investigacion/comite-central-deetica-en-investigacion

• Province of Córdoba:

REPIS- Investigaciones patrocinadas http://www.cba.gov.ar/repis/

• Province of Río Negro:

Ministerio de Salud de la Provincia de Río Negro. Comisión Provincial de Investigación Salud http://www.salud.rionegro.gov.ar/

• Province of Mendoza:

Registro Provincial de Investigaciones en Salud. (RBPRIS) http://salud.mendoza.gov.ar/dependencias/dicyt/

• Province of Neuquén:

Comisión Asesora en Investigación Biomédica en Seres Humanos (CAIBSH) https://bioetica.saludneuquen.gob.ar/caibsh/

• Province of Buenos Aires:

CCIS- Comisión Conjunta de Investigación en Salud

http://www.ms.gba.gov.ar/sitios/investigacion/comision-conjunta-de-investigacion-ccis/

• Autonomous City of Buenos Aires

Plataforma de Registro Informatizado de Investigaciones en Salud de Buenos Aires (PRIISA.BA). https://www.buenosaires.gob.ar/salud/docencia-investigacion-y-desarrollo-profesional/priisaba

ANMAT

• Presentation and authorization of clinical pharmacology studies. http://www.anmat.gov.ar/Medicamentos/investigacion_clinica.asp

Internationals (among others):

• ClinicalTrials.gov

https://clinicaltrials.gov

European Union:

• EU Clinical Trials Register

https://www.clinicaltrialsregister.eu/

• ISRCTN Registry (BioMed Central)

https://www.isrctn.com/

• OPS OMS

International Clinical Trials Registry Platform (ICTRP)

http://www.who.int/ictrp/es/