



SWATA Free Communications and Research Committee

2017 Abstract Guidelines

APPLICATION/ABSTRACT

The mission of the Free Communications and Research Committee (FCRC) is to encourage and facilitate the scholarly development of student and professional members of the Southwest Athletic Trainers' Association (SWATA) by providing an avenue for presentation and funding of their work, helping them to become well-rounded clinicians, and enhancing the evidence base within District Six. The guidelines for the 2017 Free Communications and Research Committee Abstract Competition will adhere to the National Athletic Trainers' Association Research and Education Foundation guidelines. Faculty and clinicians are welcome to submit an abstract, but only undergraduate, entry-level masters, and post-professional masters and doctoral students will be eligible for awards. To assure the exchange of valuable information, the FCRC utilizes a blinded peer-review process for abstracts following standardized guidelines, but expects abstracts to be submitted at a quality worthy of publication. Each submitted abstract will be blinded and reviewed by a minimum of two committee members. Due to the potential number of abstracts reviewed it is essential for investigators or clinicians submitting abstract follow these instructions precisely and copy edit their own work. Abstracts that are not submitted in accordance with the instructions below have a strong likelihood of being rejected.

SWATA FREE COMMUNICATIONS AND RESEARCH COMMITTEE

Call for Abstracts

**Southwest Athletic Trainers' Association Annual Meeting & Clinical Symposia
San Marcos, TX July 19-22, 2017**

Instructions for Abstract Preparation and Submission

Please read all instructions before preparing and submitting the abstract. Individuals may submit only one **Original Research Abstract** or **Clinical Case Report Abstract** as the primary (presenting) author, but may submit unlimited abstracts as a secondary author. All abstracts will undergo blind review. All presentations must be of original work (not previously presented).

The **Original Research Abstract** must be written to the accepted scientific standards of a research area and should present findings pertaining to healthcare issues related to the athletic training profession. The **Clinical Case Report Abstract** should present a unique individual athletic injury case of general interest to the SWATA membership. The **Critically Appraised Topic** abstract must include the selection of a clinically focused question answered on the basis of evidence collection with a clinically applicable bottom line as the end result.



SWATA Free Communications and Research Committee

2017 Abstract Guidelines

Formatting Instructions

Cover sheet for the entire submission – please include the following information.

- a. Lead Author's Name
- b. Lead Author's Institution
- c. Year in school (students only)
- d. NATA member number
- e. Lead Author's mailing address
- f. Lead Author's email address
- g. Lead Author's telephone number
- h. Title (brief and to the point – no longer than 16 words)
- i. Contest category that the paper is being submitted to:
 - * Clinical Case Report
 - * Original Research
- e. Supervising athletic trainer's name and their contact information (email address/phone number) (students only)

Abstract page

Prepare your abstract (on your computer) in accordance with the following instructions. You will later be directed to upload your abstract file from your computer to the Abstract Manager system.

1. Top, bottom, right, and left margins of the body of the abstract (in a WORD file) should be set at 1" using the standard 8.5" x 11" format. Use either Arial or Helvetica 12pt. font with single spacing. Provide the title of the paper or project starting at the top left margin.
2. On the next line, indent 3 spaces and provide the names of all authors, with the author who will make the presentation listed first. Enter the last name, then initials (without periods), followed by a comma, and continue the same format for all secondary authors (if any), ending with a colon.
3. On the same line following the colon, indicate the name of the institution (including the city and state) where the research was conducted. If primary author is not at the institution where the work was completed place an * after their name and following the institution where the research was conducted
4. The primary author can indicate their present institution (including the city and state). For collaborative projects where portions of the project were conducted at different institutions, list all authors as described above (#3), then list institutional affiliations using the following consecutive symbols (*, †, ‡, §, †, #, **, etc.)
5. Double space and begin entering the body of the abstract flush left in a single paragraph with no indentions. **The text of the body must be structured** (with the headings as indicated in the various formats below). Do not justify the right margin. Do not include tables or figures. **The body of the abstract for Original Research is limited to 450 words. The body of the abstract for a Clinical Case Report is limited to 600 words.** A word count generated by MS Word must be included at the bottom of the abstract. The word count should include the body of the abstract and structured headings.
6. The required formats for the structured abstracts are listed below. For further clarification, authors should consult the AMA Manual of Style 10th edition and the instructions for authors in the Journal of Athletic Training.



SWATA Free Communications and Research Committee

2017 Abstract Guidelines

7. Abstracts fall into one of the following 7 categories; the author is responsible for determining the most applicable category for structuring their abstract. Each is provided with examples where applicable but the examples are not all encompassing and some may overlap. Authors should choose the format that seems to best fit and present their data or case study.

Abstract Categories

Basic Research

- Basic Sciences (e.g. muscle tissue biopsy, EMG, etc)
- Epidemiology (e.g. cohort, case-control, intervention, clinical trial)
- Biomechanics (e.g. motion analysis, jump landing characteristics)

Survey Research

- Instrument development (e.g. validation and reliability, psychometrics)
- Cross-sectional survey (e.g. paper, web-based, or interview questionnaires)

Meta-Analysis Research & Systematic Reviews

- Meta-analysis (e.g. review and analysis of ACL clinical trials)
- Systematic Review (e.g. review of all clinical trials of the ACL without analysis)

Qualitative Research

- Research using qualitative techniques (e.g. individual interviews, focus groups, field observations, etc.)

Clinical Case Report

- Report of a Single Patient Case (e.g. snake bites football player)

Clinical Case Series

- A series of similar patients numbering between 1 and 10
- Purposely followed to describe their clinical outcomes

Critically Appraised Topic

- Must include a focused clinical question with inclusion and exclusion criteria for articles
- A Clinical Bottom Line must be established with delineation of strength of available evidence



SWATA Free Communications and Research
Committee

2017 Abstract Guidelines

Original

**Review Criteria for All
Research Abstracts:**

- Completeness of requested information in each structured heading.
- Overall clarity of writing
- Originality of research and or contribution to the literature or knowledgebase
- Methods, appropriate statistical analysis and results address the primary objective
- Consistency between data and conclusions
- Adequacy of sample size to support conclusions

Consult the *Journal of Athletic Training* Author's Guide for style information (*AMA Manual of Style* and *Index Medicus* journal abbreviations).

When choosing the appropriate research methodology and statistical analyses please consult your departmental faculty. The additional resources below may assist with choosing the appropriate research methodology and statistical analysis.

Berg, K.E. and Latin, R. W. (2004). *Essentials of research methods in health, physical education, exercise science, and recreation*. 2nd ed. LWW: Baltimore, MD.

Green, S.B., Salkind, N.J. (2010). *Using SPSS for Windows and Macintosh: Analyzing and Understanding Data*. 6th ed. Prentice Hall, Boston, MA.

Hurley, W.L., Denegar, C.R., Hertel, J. (2011). *Research Methods: A Framework for Evidence-Based Clinical Practice*. Lippincott Williams & Wilkins. Philadelphia, PA.

Jewell, D. (2010). *Evidence-Based Physical Therapy Practice*. 2nd ed. Jones & Bartlett, Sudbury, MA.

Norman, G.R. and Streiner, D.L. (2003). *PDQ Statistics*. 3rd ed. BC Decker: Hamilton, ON.

Portney, L.G., Watkins, M.P. (2008). *Foundations of Clinical Research: Applications to Practice*. 3rd ed. Prentice Hall. Boston, MA.

Paszkewicz, J., Webb, T., Waters, B., et. al. (2012) *The Effectiveness of Injury Prevention Programs in Reducing the Incidence of Anterior Cruciate Ligament Sprains in Adolescent Athletes*. *Journal of Sport Rehabilitation*, 2012 (21), 371-377. (Critically Appraised Topic Example)



SWATA Free Communications and Research
Committee

2017 Abstract Guidelines

**Formats For Each
Category**

Abstract

Basic Research Abstracts

The Title of your Abstract Bolded and in Title Case

[3 spaces]Doe JT*, Public JQ†: *First Author's Institution Name, †Second Author's Institution.

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Context: Write a sentence or two summarizing the rationale for the study, providing a reason for the study question and/or uniqueness of study. **Objective:** State the precise objective(s) or question(s) addressed in the report, including a priori hypotheses if applicable. **Design:** Describe the overall study design of the project reported (e.g., randomized controlled trial, crossover trial, cohort or cross-sectional). **Setting:** Describe the environment in which the study was conducted to help readers understand the transferability of the findings, (e.g., patient clinic, research laboratory or field). **Patients or Other Participants:** Describe the underlying target population, selection procedures (e.g., population based sample, volunteer sample or convenience sample) and important aspects of the final subject pool (e.g., number, average age, weight, height and measures of variance, years of experience or gender). Appropriate sample size should be evident. **Interventions:** Interventions are the independent variables in the study. Describe the essential pieces of the experimental methods, types of materials, measurements and instrumentation utilized, data analysis procedures and statistical tests employed. Provide validity and reliability information on novel instrumentation. **Main Outcome Measures:** Clearly identify primary or critical dependent variables that support the primary objective(s) of the study. Indicate the statistical analysis employed to answer the primary research objective(s). **Results:** The main results of the study should be given. Comparative reports must* include descriptive data (e.g., proportions, means, rates, odds ratios or correlations), accompanying measures of dispersion (e.g., ranges, standard deviations or confidence intervals) and inferential statistics data. Results should be accompanied by the exact level of statistical significance. The P value should not exceed 3 digits to the right of decimal. When the exact significance is below $P < .001$, the exact significance should be reported as $P < .001$. **Conclusions:** Summarize or emphasize the new and important findings of the study. The conclusion must be consistent with the study objectives and results as reported and should be no more than three to four sentences. If possible, relate implications of the findings for clinical practice. **Word Count:** Limited to 450 words including headings.

** The purpose of having both descriptive and inferential data is that it provides the reader with the ability to judge the concluding statements. Descriptive data provides confidence that the data are 'reliable' and provides a gauge to determine whether the inferential statistics and conclusions are meaningful. Studies reporting analysis of larger data bases with multiple variables do not need to report all descriptive data, but should provide descriptive data for those variables which the author(s) believe to be the primary outcome(s) and support the overall conclusions of the study.*



Survey Research

Abstracts

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Context: Write a sentence or two summarizing the rationale for the study, providing a reason for the study question. **Objective:** State the precise objective(s), purpose or question(s) addressed in the report. **Design:** Describe the overall study design of the project reported (e.g., cross sectional, case-control, longitudinal or controlled intervention trial). **Setting:** Describe the environment in which the study was conducted to help readers understand the transferability of the findings, (e.g., population-based, patient clinic, classroom or athletic event). **Patients or Other Participants:** Describe the underlying target population, sample selection procedures (e.g., population based, volunteer or convenience sample, random or systematic sample, or stratified or cluster sampling) and important aspects of the final subject pool (e.g., number, average age, years of experience or gender). Provide the final response rate. **Interventions:** Interventions are the independent variables in the study. Describe the essential pieces of the experimental methods, the mode of survey administration (e.g., in-person interview, telephone, self-administered, online or computer-assisted), details of the survey development (formative research or pre-testing for new instruments), execution and data collection process, and instruments utilized. Provide validity and reliability information for all new instruments. **Main Outcome Measures:** Clearly identify primary or critical dependent variables that support the primary objective(s) of the study. Describe how any data was manipulated (e.g. scoring process for scaled instruments or categorization of variables). Indicate the data and statistical analysis employed to answer the primary research objective(s). **Results:** The main results of the study should be given. Reports must* include descriptive data (e.g., proportions, means, rates, odds ratios or correlations), accompanying measures of dispersion (e.g., ranges, standard deviations or confidence intervals) and inferential statistical data. Results should be accompanied by the exact level of statistical significance. The *P* value should not exceed 3 digits to the right of decimal. When the exact significance is below $P < .001$, the exact significance should be reported as $P < .001$. **Conclusions:** Summarize or emphasize the new and important findings of the study and relate implications of the findings for clinical practice. The statement of your findings must be consistent with the results as reported and should be no more than three to four sentences. **Word Count:** Limited to 450 words including headings.

** The purpose of having both descriptive and inferential data is that it provides the reader with the ability to judge the concluding statements. Descriptive data provides confidence that the data are 'reliable' and provides a gauge to determine whether the inferential statistics and conclusions are meaningful. Studies reporting analysis of larger data bases with multiple variables do not need to report all descriptive data, but should provide descriptive data for those variables which the author(s) believe to be the primary outcome(s) and support the overall conclusions of the study.*



**Meta-Analysis and
Reviews**

Systematic

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Context: Write a sentence or two summarizing the rationale for the study, providing a reason for the study question. **Objective:** State the precise objective(s) or question(s) addressed in the report, including a priori hypotheses if applicable. **Data Sources:** Identify how relevant research papers were identified – include databases and timeframe, key words and search limits. **Study Selection:** Describe the processes through which studies were selected for inclusion for further analysis. **Data Extraction:** Identify the number of investigators, the descriptive and measurement data obtained and if and how the quality of study methods was evaluated. **Data Synthesis:** Describe how the data were organized, the statistical procedures applied (during assessment of heterogeneity) and the results (e.g., effect sizes, odds ratios and 95% confidence intervals) of the analysis. **Conclusions:** Summarize or emphasize the new and important findings of the study and relate implications of the findings for future research and/or for clinical practice and offer an indication as to the strength of the evidence provided. The statement of your findings must be consistent with the results as reported. **Word Count:** Limited to 450 words including headings

Qualitative Research Abstracts

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Context: Briefly explain the rationale for the study—provide a background for the study question. **Objective:** State the precise objective(s) or question(s) addressed in the report. **Design:** Describe the overall study design of the project reported (e.g., case study, phenomenology or grounded theory).

Setting: Describe the environment in which the study was conducted to help readers understand the transferability of the findings, (e.g., clinical setting or educational institution). **Patients or Other Participants:** Describe the underlying target population, selection procedures and important aspects of the final subject pool (e.g., number, average age and measures of variance, years of experience or gender). Describe the essential pieces of the sampling methods (e.g., theoretical sampling and criterion sampling). Comment on why this number of participants was used (e.g., data saturation guided the total number of participants selected for the study). **Data Collection and Analysis:** Describe how the data were collected (e.g., interviews, observations or document analysis), managed (e.g., interviews were recorded and transcribed verbatim; identify if software was utilized) and analyzed (e.g., the interviews were analyzed using an inductive content analysis). Include intercoder agreement information if relevant to the study. Identify any verification strategies used to ensure trustworthiness (e.g., indicate form of triangulation, or use of peer debriefer). **Results:** A short descriptive account of the case or the interpretation of the findings should be provided. This should include identifying and briefly explaining the emergent categories of themes. **Conclusions:** Summarize or emphasize the new and important findings of the study and relate implications of the findings for future research and/or for clinical



SWATA Free Communications and Research Committee

2017 Abstract Guidelines

practice. The statement of your findings must be consistent with the results as reported and should be no more than five sentences. **Word Count:** Limited to 450 words including headings.

Clinical Case Report Abstracts

NOTE: All clinical case report abstracts submitted to Free Communications must have permission of the patient prior to submission.  [\[Click here\]](#) for example.

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Background: Include the individual's age, sex, sport or activity, pertinent aspects of their medical history, a brief history of their complaint and physical findings from the athletic trainer's examination. **Differential Diagnosis:** Include all possible diagnoses suspected based on the history, mechanism of injury, and the initial clinical examination prior to physician evaluation and subsequent diagnostic imaging and laboratory tests. **Treatment:** Include the physician's evaluation and state the results of diagnostic imaging and laboratory results if performed. The final diagnosis of the injury or condition and subsequent treatment and clinical course followed should be clearly detailed. Relevant and unique details should be included, as well as the final outcome of the case. **Uniqueness:** Briefly describe the uniqueness of this case such as its mechanism, incidence rate, evaluate findings, rehabilitation, or predisposing factors. **Conclusions:** Include a concise summary of the case as reported and highlight the case's importance to the athletic training profession and provide the reader with a clinical learning opportunity. **Word Count:** Limited to 600 words including headings.

Clinical Case Series Abstracts

These are series of similar patients typically greater than 1 but less than 10 who have been purposefully followed to describe their clinical case outcomes. The intentions of these projects are to describe occurrences in a like group of patients and share insights on these occurrences. The case series does not have to include a hypothesis nor should a cause and effect conclusion be made due to the observational nature of this information.

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Background: Describe the underlying target population and important aspects of the subject pool (e.g., number, average age, weight, height (with measures of variance, sex, sport or activity, and years of experience). Pertinent aspects of their medical histories should be included. Describe their complaints, MOI, initial clinical examination, diagnostic imaging, lab tests, and their commonality (examples: characteristic, injury, postural/gait abnormality, pathology, MOI). It is encouraged to present information



SWATA Free Communications and Research Committee

2017 Abstract Guidelines

as a group or average (proportions). **Treatment**: The clinical course followed should be clearly detailed. Time frame should be provided and averaged if possible. Relevant and unique details should be included. Specific outcome variables in which all patients within the series were evaluated for should be listed. The final outcome of these variables should be provided in respect to their common characteristic. **Results**: The unique subsequent treatment, prevention program, specific rehab program, special/diagnostic test, outcomes, or predisposing factors, that all subjects experienced is explained. Use of percentages is encouraged. **Uniqueness**: Briefly describe the uniqueness of these cases as a whole. **Conclusions**: Conclusions should recap the most important background, treatment, and uniqueness points for the reader. Your conclusions must be consistent with the final outcome. Statements should concisely describe the most pertinent points of your clinical cases while providing the reader a clinical learning opportunity. Avoid statements of cause and effect since these are observational reports. **Word count**: 600

Critically Appraised Topics Abstracts

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Clinical Scenario: A brief description of the clinical scenario leading to the clinical question. **Clinical Question**: A focused clinical question of importance in athletic training. Search Strategy: Identify how relevant research papers were identified – include databases and timeframe, key words, search limits and the processes through which studies were selected for inclusion for further analysis. **Search Results**: Identify the number of relevant studies found, the number of investigators and how the quality of study methods were evaluated. **Best Evidence**: Indicate how many studies were chosen for inclusion and appraisal in this CAT and provide the reasons that these studies were selected (use Centre for Evidence-Based Medicine's definitions in determining level of evidence). **Clinical Bottom Line**: The most important take-home message from the available evidence. Some statement regarding the level of available evidence and subsequent strength of recommendations is required. **Strength of Recommendation**: A brief description of the strength of evidence summarized following the critical appraisal. **Word Count**: Limited to 450 words including headings.

If you have any questions about an abstract or presentation designation please contact:

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