• aaps Workshops



60TH ANNUAL PHARMACEUTICAL ANALYSIS CONFERENCE



AUGUST 10-13, 2020 • MEMORIAL UNION, MADISON, WI



Call for Papers

- Email your poster abstract in a Word (.doc) file to abstracts@aaps.org
- Poster presentations will be pre-recorded with audio. Each presenter will be scheduled for a 30-min online video chat session to answer questions from attendees.
- 30 Poster slots available. (First to submit and register to attend will fill the slots.)
- The submission deadline is extended to June 30, 2020*.

Poster Abstract Submission Timeline

Submission Deadline
Registration Deadline for Presenting Authors
Notification (Accept or Reject)
Schedule Notification Sent
Pre-recorded Presentation Due

June 30, 2020*
June 30, 2020
sent as received/registered
July 7, 2020
July 27, 2020

*Attention: Slots are filled first-come/first-serve by qualified submissions, so authors are cautioned that all slots may be filled before June 30, 2020.

Submission Requirements

1. Conditions and Policies

Persons submitting an abstract for presentation at the AAPS workshop are doing so with the understanding that they agree to abide by the conditions and policies provided below, as well as the decisions of AAPS staff. Questions should be directed to abstracts@aaps.org.

2. Registration Requirement for Publication

The presenting author is required to pre-record a short presentation (audio + poster image) for attendee viewing. In addition, the author is required to be available for a 30-min video chat session where attendees can connect with questions. If the presenting author is unable to virtually attend the meeting, a co-author is responsible for the video chat. Registration is required for the author virtually presenting the poster at the meeting.

3. Cancellation policy

If the author that will be presenting the paper cancels his/her registration between the dates of June 30, 2020, and July 27, 2020, AAPS will impose a \$50 administrative fee. After July 28, 2020 no refunds will be issued. Registrant substitutions from the same company are allowed and may be submitted in writing at any time without penalty. If the membership status of the substitute differs from that of the original registrant, a refund or additional charge may apply.

4. Ethics Statement

Final poster presentations must include all methods used and data resulting from the research. Actions such as omission of data from a poster presentation that was included in the submitted abstract will be considered unethical. Authors and organizations violating these requirements will be subject to penalties such as restrictions from presenting future posters.

Poster Abstract Policies and Procedures

Permissions/Clearances

It is the responsibility of the author(s) to obtain the necessary permissions and clearances for all research prior to submission of the abstract. AAPS assumes no liability or responsibility for the publication of any material that is submitted.

Acceptance Criteria

Acceptance of the abstract for presentation will be based on the concise, accurate presentation of new data that is relevant to the field of pharmaceutical research. It is imperative that data is presented in the results section so that AAPS screeners can judge the scientific value of your abstract. Include all research information, data, charts and graphs in your submission so that it can be reviewed in its entirety.

Rejection Criteria

Criteria for rejection include:

- lack of data,
- stating data will be included in the poster presentation,
- commerciality,
- inconsistent or ambiguous data,
- reviews of literature.
- lack of novelty or innovation,
- plagiarism
- previously published, but not acknowledged, or
- failure to follow format guidelines (Purpose, Methods, Results, and Conclusion).

Abstract Revisions

Revisions can be made at any time before the June 30, 2020 submission deadline. *Revisions cannot be made after the submission deadline*. Proofread, spell check and verify that all authors are listed on your abstract before submitting.

Notification of Receipt and Verification of Submission

You will receive a confirmation email once your abstract is received. This notification only confirms receipt of your submission and is not a notification of acceptance.

Screening/Presentation

Final assignment to poster sessions will be made at the discretion of the screening committee.

Notification of Acceptance

Status notifications will be sent via email once registration is complete for the presenting author. Notification and updates will be sent to the *submitting author* only.

Notification of Presentation Schedule and Poster Number

Notifications will be sent via email by **July 7, 2020**, with the presentation date, time and poster number for the abstract. If you do not receive notification at that time, please contact AAPS at abstracts@aaps.org for an update on the schedule of your abstract. Scheduling notifications will be sent to the submitting author.

Abstract Withdrawal

To withdraw your abstract, a written request must be sent to abstracts@aaps.org. This request should be sent by the submitting author and must include the abstract title and abstract submission ID number. AAPS will acknowledge all withdrawal notifications by email.

Copyright/Permissions

- By submitting this poster abstract to AAPS, I state that all authors listed thereon are aware that they are listed.
- By submitting this poster abstract to AAPS, I give AAPS permission to display and publish the abstract in various ways related to the conference.

Poster Abstract Submission Format

Sample Abstract – Last page of this document

- 1. One page Word (doc) document with 1" margins
- 2. Title of Abstract (this will be your poster title), Times Roman 12, Bold
- 3. Author(s), Affiliation(s), Times Roman 10
- 4. Section Headings: Purpose, Methods, Results, Conclusion, Arial 10
- 5. Reference(s) (if needed) Arial 10
- 6. Acknowledgement(s) (if needed) Arial 10
- Images are allowed. However, all text and images must fit on one page with 1" margins

Poster Abstract Submissions—Topics

The abstract for poster presentation should be related to the objectives of the Conference and appropriate for conference attendees. This three-day conference is to provide an educational forum to discuss issues and applications related to pharmaceutical analysis. The conference connects leaders together to share new scientific and regulatory developments in the areas related to or with direct application for pharmaceutical analysis, product development, product quality and the analytical method lifecycle applications.

Submitting Your Poster Abstract

- 1. Email to abstracts@aaps.org
- 2. Subject of Email: LOL August (or name of event)
- 3. Attach Word (doc) file to email
- 4. Body of Email Indicate the Topic the abstract relates to.
- 5. Include contact information for the submitter of the abstract
 - a. Last Name, First Name, Middle Initial
 - b. Company/Affiliation Name
 - c. Email Address
 - d. Phone Number
- 6. Indicate the Presenting Author of the Poster
 - a. Last Name, First Name, Middle Initial
 - b. Company/Affiliation Name
 - c. Email Address
 - d. Phone Number

Sample Abstract

Quantitative Determination of Compound X in Rat and Dog Plasma using LC/MS/MS Doe, J, AAPS; Smith, J, Blank Corp.

Purpose: To develop and validate an LC/MS/MS method for the determination of compound X, an anti-cancer agent in rat and dog plasma.

Methods: This method utilized a solid phase extraction from 0.1 mL plasma with an Isolute C2 cartridge and methanol as eluent. HPLC separation was carried out on a Phenomenex Luna C18 column (150 x 4.6 mm) column at a flow rate of 0.3 mL/min with an analysis time of 4.5 minutes. Compounds were eluted using a mobile phase of H20/CH3CN/HCOOH: 70/30/0.1 (v/v/v), pH 4.0. Compound Y, a structural analogue of compound X was used as the internal standard to account for differences due to adsorption, extraction, and instrumental performance. Mass spectrometry detection was carried out with a PE Sciex API III+ triple quadrupole mass spectrometer equipped with a Turbo IonSpray as LC/MS interface. ESI mass spectra were acquired in positive ion mode with multiple reaction monitoring.

Results: No matrix interference was observed across the elution windows of X and Y, indicating the specificity of the method. Acceptable intraday and interday assay precision (<5% CV) and accuracy (<10% diff.) were observed over a linear range of 1-1500 nM for both matrices. The mean (n=3) correlation coefficients in rat and dog plasma were 0.9998_0.0001 and 0.9997_0.0002, respectively. The mean extraction recovery was 91.2 2.5% and 90.1 4.6% for rat and dog plasma, respectively.

Conclusion: A validated method which was robust, sensitive, specific, accurate, and reliable was developed. The method has been used to quantify rat and dog plasma for pharmacokinetic and drug safety studies.