



3rd Annual Women's Cardiovascular Symposium

Friday, October 11, 2024 | Cincinnati, Ohio

Abstract Submission Form

The Women's Heart Center Program Committee is accepting abstract submission forms through **August 16, 2024**. Completed forms should be emailed to WHC@TheChristHospital.com.

Abstract submissions should be gender- and sex-specific research pertaining to one of the program topics outlined below.

The Program Committee wishes to encourage young scientific investigators and will reward up to 4 abstracts/posters submitted by presenters considered early career (definition provided below). First place will receive \$1000, second place will receive \$500, and two honorable mentions will each receive \$250.

The presenting author will be sent an email with the status of the submission by **August 30, 2024**. If your abstract is accepted, your notification will contain complete presentation information. However, please note the following:

- All human subject research must conform to the principles of the Declaration of Helsinki of the World Medical Association.
- The presenting author should be able to provide documentation of IRB approval if requested.
- The Program Committee is unable to reimburse presenters for travel, hotel, or per diem expenses.
- Submission of an abstract constitutes a commitment by the presenting author (or designee) to present in-person at the symposium on October 11, 2024, during the following times:
 - Registration & Networking: 7:00 – 8:00 am
 - Networking Lunch: 12:00 – 1:00 pm
 - Poster Session Award Announcement: 3:40 – 4:00 pm
- All accepted abstract presenters must register for the symposium via Eventbrite and pay the applicable registration fees (trainees and invited speakers will have the registration fee waived).
- If an author wishes to withdraw an abstract, please email WHC@TheChristHospital.com.

Presenting Author Information

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Early Career (Defined as physicians, scientists, medical students, and other healthcare providers currently in residency or fellowship programs or within three years of training)? Yes No

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Abstract Topic (must be gender- or sex-specific)

- | | | |
|-------------------------------------------------------------|-------------------------------------------------|--------------------------------------------------------|
| <input checked="" type="checkbox"/> Preventative cardiology | <input type="checkbox"/> General cardiology | <input type="checkbox"/> Interventional cardiology |
| <input type="checkbox"/> Heart failure | <input type="checkbox"/> Cardio-oncology | <input type="checkbox"/> Cardio-obstetrics |
| <input type="checkbox"/> Electrophysiology | <input type="checkbox"/> Cardiovascular Imaging | <input type="checkbox"/> Coronary Microvasculature |
| <input type="checkbox"/> Social Determinants of Health | <input type="checkbox"/> Mental Health | <input checked="" type="checkbox"/> Precision Medicine |

Title: Include the full title as it will appear on the poster.

Riloncept Utilization in a Steroid-Sparing Paradigm for Recurrent Pericarditis: Real-World Evidence Demonstrating Increased Adoption

Background: In an initial paragraph, provide relevant information regarding the background and purpose of the study, preferably in no more than two to three sentences.

Recurrent pericarditis (RP) is a chronic autoinflammatory disease mediated by interleukin-1 (IL-1) requiring prolonged treatment, historically anchored on corticosteroids (CS) in refractory patients (pts). Early CS use may prolong disease, and long-term use accumulates side effects. RESONANCE, an ongoing 5-year, 500-pt US registry, has been collecting real-world RP data since 2021. We hypothesized that riloncept availability in RP (2021) would enable a CS-sparing paradigm in pts failing NSAIDs/colchicine.

Methods: Briefly state the methods used.

Pt demographics, disease characteristics, and RP medication sequencing are reported (13 July 2023 cut) in 210 pts with complete data records from 272 active RP pts at 23 sites; median [IQR] 2.1 [1.5:2.5] years (424.5 PY) of observation.

Results: Summarize the results in sufficient detail to support the conclusions.

Mean age was 49.9 ± 15.9 years; 59.5% female. Time from index pericarditis episode was 3.4 [2.1:6.1] years. Of 202 pts with complete medication data, half (101) were managed with IL-1 pathway inhibition, of whom 73% (74) took riloncept (Figure). Over 50% of pts starting riloncept did so after NSAIDs/colchicine (Figure inset).

Conclusions: Concisely state the conclusions reached.

Whereas 2015 ESC guidelines position IL-1 pathway inhibitors after CS, real-world registry data since riloncept availability in RP (2021) identify new adoption of a CS-sparing paradigm in RP, with riloncept use directly after NSAIDs/colchicine. Long-term treatment for full RP duration may obviate iatrogenic complications of chronic steroid use and could inform future treatment guidelines.

Tables/Figures/Graphics: Include images that are part of your submission here. Images should be high resolution and have a file type of “gif”, “jpg”, or “jpeg”.

Figure: Distribution of medication class use in total population (n=202 pts with non-missing data) and (inset) within the 6-month period prior to riloncept initiation.

