

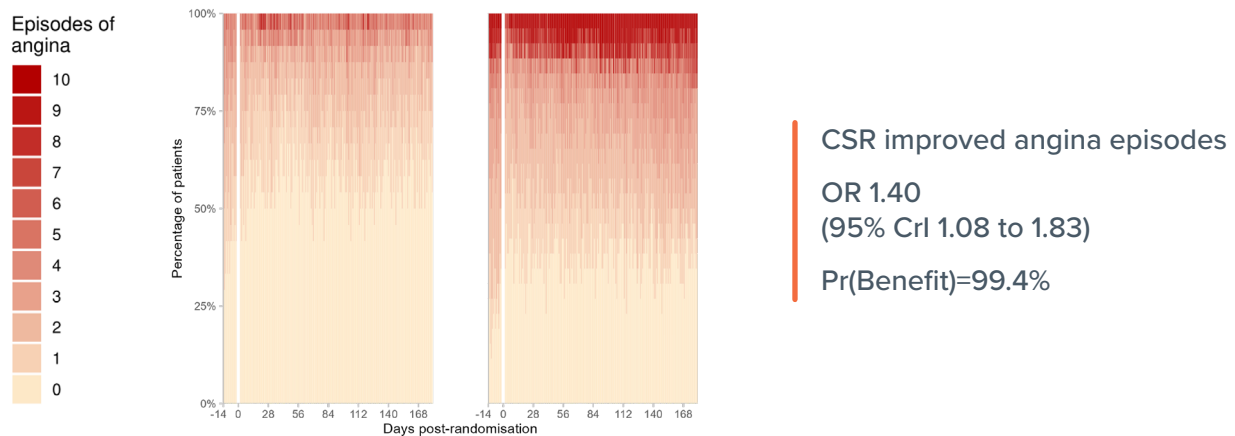
ORBITA-COSMIC RESULTS SHOW THE SHOCKWAVE REDUCER REDUCES ANGINA FREQUENCY AND IMPROVES HEART DISEASE RELATED QUALITY OF LIFE WHILE IMPROVING SUBENDOCARDIAL PERFUSION¹



ORBITA-COSMIC is a randomized, placebo-controlled trial of the coronary sinus Reducer for the treatment of refractory angina.

PRIMARY OUTCOMES

SYMPTOM OUTCOME

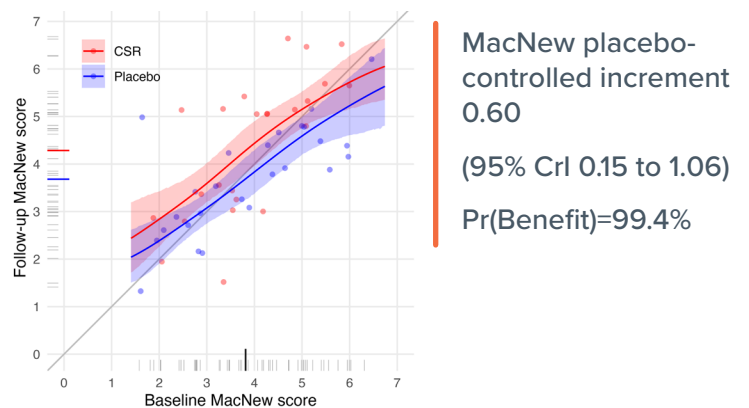
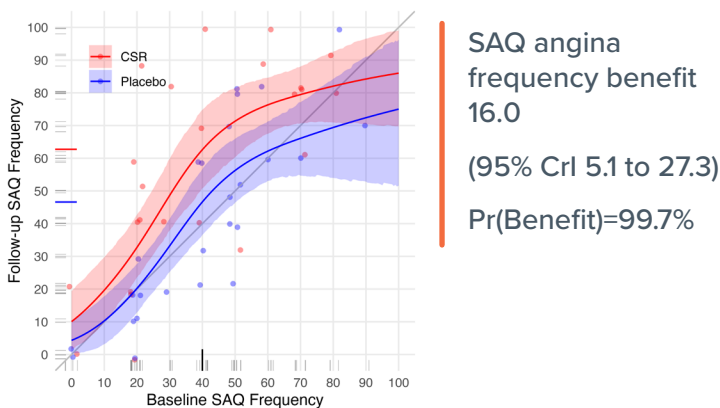


“The symptoms didn’t just improve on the angina symptom app, they also improved on the symptom questionnaires and health-related quality of life questionnaires.”

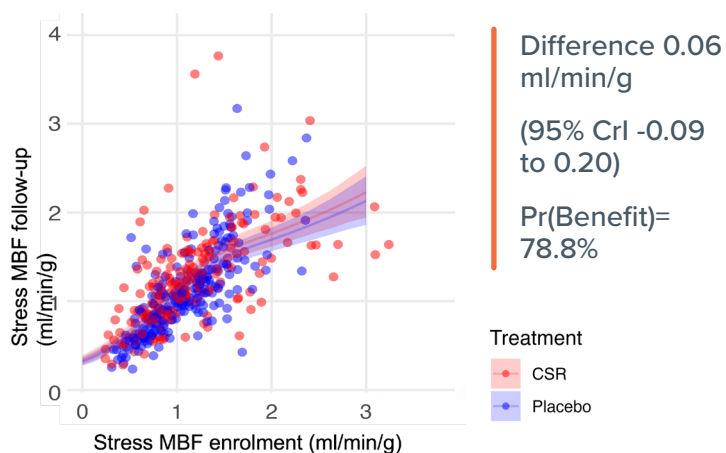
— Dr. Rasha Al-Lamee, Imperial London College, UK²

The Reducer decreased the number of daily angina episodes compared to placebo at 6 months. This benefit emerged at 10 weeks and persisted to 6 month follow up.

Similar to the daily reported angina frequency, there was evidence of benefit on angina frequency reported using the Seattle Angina Questionnaire and on the MacNew heart disease health related quality of life.



IMAGING OUTCOME



Endocardial:Epicardial Ratio		
	Difference at follow up for CSR vs placebo	Probability of benefit with CSR vs placebo*
Endo:epi ratio stress MBF Ischaemic segments	0.09 (95% CrI 0.00 to 0.17)	98.2%
Non-ischaemic segments	-0.02 (95% CrI -0.10 to 0.07)	35.1%
Difference	0.10 (95% CrI 0.02 to 0.19)	99.2%

*Difference is Pr(interaction)

While the primary imaging endpoint showed no improvement in transmural myocardial perfusion, it did show improvement in subendocardial perfusion, supporting the theorized mechanism of action of the Reducer.

“The device seems to work...there is a transfer of blood from the epicardium to the endocardium and this appears to be part of the mechanism of how the reducer works.”

— Dr. Tommaso Gori, University Medical Center, Mainz, Germany³

KEY TAKEAWAYS FROM THE ORBITA-COSMIC RESULTS:

- CSR improves angina and heart disease related quality of life compared to placebo.
- Improvement in myocardial blood flow to subendocardium.
- CSR is a third treatment option for refractory angina.

“With the results of this placebo-controlled trial, we can tell them that their symptoms are more likely to improve with the reducer.”

— Dr. Rasha Al-Lamee, Imperial London College, UK⁴

Late Breaking Clinical Trial presented at ACC 2024 by Michael Foley, MBBS, BSc
Chief investigator: Rasha Al-Lamee MBBS, PhD
Imperial College London

FOOTNOTES

1. Foley et al. The Lancet. 2024 Apr 8; [https://doi.org/10.1016/S0140-6736\(24\)00256-3](https://doi.org/10.1016/S0140-6736(24)00256-3)
2. <https://www.statnews.com/2024/04/08/johnson-and-johnson-shockwave-coronary-sinus-reducer-trial/>
3. <https://www.tctmd.com/news/coronary-sinus-reducer-eases-symptoms-refractory-angina-orbita-cosmic>
4. <https://medicalxpress.com/news/2024-04-coronary-sinus-relieves-angina-unclear.html>

CAUTION In the United States, Shockwave Reducer is an investigational device, limited by United States law to investigational use. The Reducer is subject of Investigational testing and is being studied in the COSIRA II trial in Canada. The Reducer is commercially available in certain countries outside the U.S. and Canada. Please contact your local representative for specific country availability.
Prior to use, please reference the Instructions for Use for more information on warnings, precautions and adverse events:
ifu.neovasc.com

SHOCKWAVE
REDUCER