

RESPOND-CRT Trial

- Contractility Sensor-Guided Optimization of Cardiac Resynchronization Therapy –

Brugada J, Delnoy PP, Brachmann J, Reynolds D, Padeletti L, Noelker G, Kantipudi C, Rubin Lopez JM, Dichtl W, Borri-Brunetto A, Verhees L, Ritter P, Singh JP; RESPOND CRT Investigators. Contractility sensor-guided optimization of cardiac resynchronization therapy: results from the RESPOND-CRT trial. Eur Heart J. 2017 Mar 7;38(10):730-738. doi: 10.1093/eurheartJ/ehw526. PMID: 27941020;

Background & objectives

- → Although cardiac resynchronization therapy (CRT) is effective in patients with systolic heart failure (HF) and a wide QRS interval, a substantial proportion of patients remain non-responsive.
- → The SonR® hemodynamic contractility sensor embedded in the right atrial lead enables individualized weekly automatic optimization of the atrioventricular (AV) and interventricular (VV) timings.

RESPOND-CRT is a prospective, **Objective**: The multicentre, randomized, double-blind, non-inferiority trial study investigated the safety and efficacy of the contractility sensor system in HF patients undergoing de novo CRT implants.

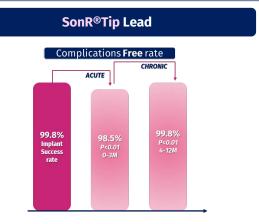
Methodology



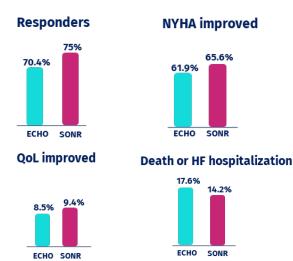
Trial endpoints

- → Clinical responders rate (without HF-related events, with NYHA class improvement of 1 level or QoL improvement of at least 5 points,
- Freedom from acute and chronic atrial lead-related complications.
- Rate of clinically worsened patients
- Composite of all-cause mortality or HF hospitalisation.

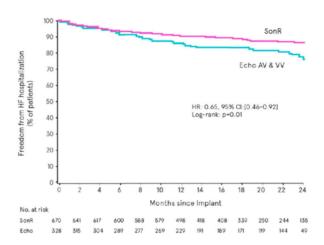
Results



75% CRT responders rate



35% risk reduction in HF hospitalization



Conclusion

- → Automatic AV and VV optimization using SonR® technology was safe and as effective as Echo-guided AV and VV optimization in increasing response to
- → Clinical response for most subgroups was in favour of using the SonR® sensor, especially in patients with a prior history of atrial fibrillation or renal dysfunction.