

Change in Heart Rate from Pre-Implant to Discharge is Associated with Mortality and Admissions in LVAD Patients- A Substudy of the ENDURANCE trial

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BACKGROUND

- The effect of heart rate (HR) on prognosis has been studied in almost every field of cardiology.
- However, no study had evaluated the effects of HR on prognosis and admissions in patients on LVAD support.

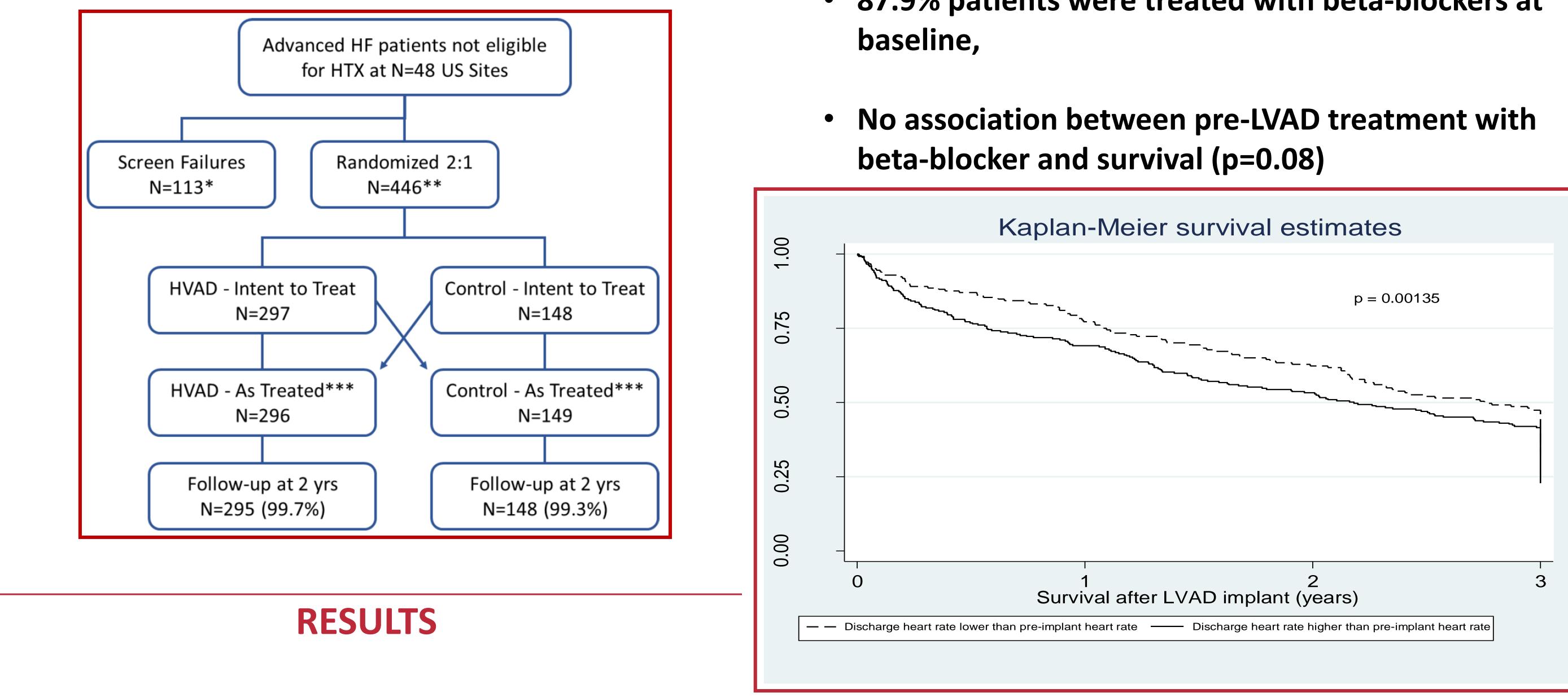
RESULTS (Cont...)

Change in Heart Rate and Survival:

- A decrease in HR at discharge post-implant \bullet compared to HR pre-implant was associated with improved survival (p=0.001, figure).
- This remained significant in a multivariate analysis (OR for mortality 1.01 per 1 bpm increase in HR,

METHODS

- The ENDURANCE trial is a multicenter randomized trial involving 445 patients who were assigned, in a 2:1 ratio, to the study (centrifugal-flow) device or the control (axial-flow) device.
- Using the data collected in the ENDURANCE trial, we analyzed the association between HR, mortality and hospitalization.



95% confidence interval 1.00 – 1.03, p=0.01).

- A decrease in HR at discharge post-implant compared to HR pre-implant was associated with a longer time to first admission (p=0.001).
- HR at discharge or 3-months post-implant was not associated with mortality.

Beta blockers and Survival

- 87.9% patients were treated with beta-blockers at

A total of 445 patients were included in the ENDURANCE trial and in this analysis (78.4% male, mean age 64.7 years +11.2).

CONCLUSION

- A decrease in HR in the early period after LVAD implantation is associated with improved survival
- 296 (66.5%) patients were treated with HVAD and 149 (33.5%) patients with HMII devices.
- The median HR pre-implant was 81 beats per minute (bpm) (interquartile range 73-93) and median HR at discharge post-implant was 88 bpm (interquartile range 79-95).

Right Ventricular Failure and Heart Rate:

No correlation was noted between HR and early right ventricular failure (p=0.30)







and reduced risk of admission.

Future studies will determine if HR reduction strategies could emerge as a therapeutic paradigm to improve outcomes with LVAD therapy.

DISCLOSURES

- **EYB** Consultant: Luitpold Pharmaceuticals, Inc.
- SS None.
- MAA None.
- PA None.
- JT Consultant: Medtronic, Abiomed, CareDX; CEC: Abbott.
- **RK C; Consultant; Medtronic.**
- EB None.
- CM Investigator / Consultant: Medtronic, Abbott, Abiomed.
- NM Consultant: Medtronic, Abbott, Syncardia.
- FP None.
- JR None.
- TV Employee, Medtronic.
- JER PI, Endurance Trial (Medtronic). PI, Momentum 3 Trial (Abbott).