Near elimination of ventricular pacing in SafeR mode compared to DDD modes: a randomized study of 422 patients.

RESULTS FROM THE SAVE-R STUDY

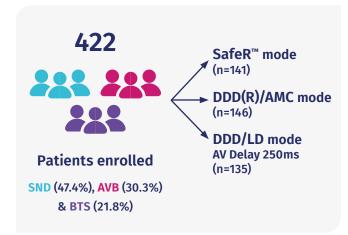
Davy JM, Hoffmann E, Frey A, Jocham K, Rossi S, Dupuis JM, Frabetti L, Ducloux P, Prades E, Jauvert G. Pacing Clin Electrophysiol. 2012 Apr;35(4):392-402. doi: 10.1111/j.1540-8159.2011.03314.x. Epub 2012 Feb 6. Erratum in: Pacing Clin Electrophysiol. 2012 Jul;35(7):909. PMID: 22309303. April 2015.

Background & objectives

→ SafeR[™] performance versus DDD / automatic mode conversion (DDD/AMC), based on AV delay hysteresis, and DDD with a 250 ms atrioventricular (AV) delay (DDD/LD) modes were assessed toward ventricular pacing (VP) reduction.

Objective: The main endpoint was the percentage of VP (%VP) at 2 months and 1 year after randomization, ascertained from device memories.

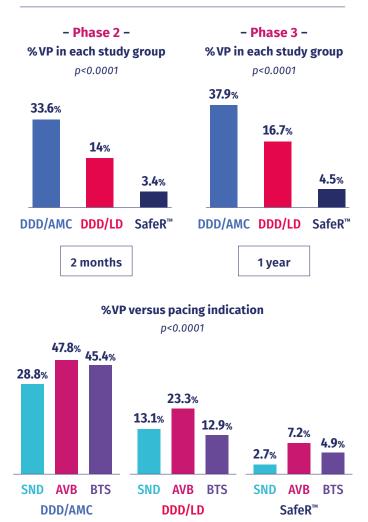
Methodology



Study was divided into 3 phases:

- > Phase 1: Before assignment to one of the three study groups, all patients were paced in SafeR™ mode (with bipolar atrial sensing mandatory) to confirm the proper functioning of the pacing system and the presence of predominant spontaneous AV conduction.
- > Phase 2: interim evaluation and interrogation of the pacemaker 2 months after random assignment
- > Phase 3: final evaluation at 1 year

Results



Conclusion

- → This randomized trial confirmed the superiority of SafeR[™] in the prevention of ventricular pacing in patients without fixed high-degree AV block compared with DDD pacing.
- → In patients with preserved AV conduction, SafeR[™] significantly decreased the percentage of ventricular-paced events compared with DDD pacing modes and eliminated ventricular pacing in high proportions of patients paced for paroxysmal AV block, SND, or BTS.

