Which Is More Likely to Achieve Cardiac Synchronization: Left Bundle Branch Pacing or Left Ventricular Septal Pacing?

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INTRODUCTION

In advanced heart failure patients with low left ventricular ejection fraction and left bundle branch block (LBBB), cardiac resynchronization therapy (CRT) via stimulation of both the right ventricle (RV) and the left ventricular lateral wall is a recommended therapeutic strategy (1–3). However, conventional biventricular pacing causes a dyssynchronous cardiac contraction due to non-physiological fusion of paced propagation, with a non-response rate of up to 30% (4, 5). In 2016, Mafi-Rad et al. (6) established the viability of the left ventricular septal pacing (LVSP) via a trans-interventricular septal approach in 10 patients with sinus node dysfunction, which shortened QRS duration and preserved acute left ventricular contractility compared to RV pacing. Huang et al. refined LVSP and introduced first left bundle branch pacing (LBBP) in 2017 (7), which could restore physiological left ventricular contractility in a patient with LBBB by pacing left bundle branch (LBB) immediately beyond the conduction blockage with satisfactory pacing parameters. Many studies have demonstrated the feasibility and stability of LBBP in patients with pacemaker indications, and it has been proposed that LBBP is a novel physiological pacing method for delivering CRT for achieving electric resynchronization in patients with LBBB (8–10).

BRIEF PACING MECHANISMS OF LBBP AND LVSP

Selective LBBP (SLBBP) and non-selective LBBP (NSLBBP) are two subgroups of LBBP. SLBBP, that is, only the LBB trunk or its proximal fascicles is captured (Figure 1A). NSLBBP, that is, concomitant LBB and adjacent myocardium are captured (Figures 1B,E). It is LVSP if just the left ventricular septal myocardium is captured (Figure 1D). Both LVSP and LBBP usually present a paced pseudo right bundle branch block (RBBB) pattern in lead V1 (11), with the percentage of direct evidence that LBBP captured LBB ranging between 60 and 90% (12–14). Therefore, LBBP described in some previous studies was actually LVSP. A method to measure the time from stimulus to left ventricular activation at high and low outputs in lead V5 or V6 (Stim-LV AT) to distinguish LBBP from LVSP with a specificity of 100% has recently been presented (11). If the Stim-LV AT remains shortest and constant (prolonged ≤ 10 ms) as the pacing output decreases, it must be LBBP, because LBBP directly captures the LBB resulting in physiologically LV excitation; otherwise LVSP can be considered, because LVSP excites left ventricular septum first, rather than LBB. SLBBP and NSLBBP can be distinguished by the discrete component and isoelectric interval between the pacing artifact and V wave on intracardiac electrogram with unchanged Stim-LVAT (11).
In the paper published in Frontiers in Cardiovascular Medicine, Zhu et al. (15) used ultra-high-frequency electrocardiography to compare ventricular depolarization in SLBBP, NSLBBP, and LVSP in 57 bradycardia patients, which were rigorously distinguished by Stim-LVAT. They concluded that LVSP preserved interventricular synchrony and had the same or better local depolarization durations than NSLBBP and SLBBP. Furthermore, they investigated two different types of NSLBBP capture, namely, NSLBBP with LBB and adjacent myocardium captured (Figure 1B), and NSLBBP with LBB and left septal myocardium captured (Figure 1E). NSLBBP with LBB and adjacent myocardium captured, that is, NSLBBP is converted to SLBBP with a shortest and constant Stim-LVAT while decreasing the pacing outputs. NSLBBP with LBB and left septal myocardium captured, that is, NSLBBP is converted to LVSP with prolonged Stim-LVAT while decreasing the pacing outputs. They evaluated the two types of NSLBBP capture and found no statistical difference in Stim-LVAT between the two types, but NSLBBP with LBB and left septal myocardium captured showed greater interventricular synchronization.

Then, which pacing strategy is more physiological, LBBP or LVSP? SLBBP and NSLBBP, unlike LVSP, capture the intrinsic conduction system and rapidly excite LV to maintain left ventricular synchrony at levels comparable to intrinsic left ventricular activation (16). At the same time, activation propagates slowly from left to right in the interventricular septum to excite RV, resulting in interventricular dyssynchrony. LVSP, on the other hand, captures left ventricular septal myocardium, resulting in direct left-to-right septal activation, preserving interventricular dyssynchrony. The terminal R′/r′ wave duration in lead V1, which indicates delayed right ventricular excitation, was significantly longer in LBBP than in LVSP (17), also indicating that LBBP caused more pronounced interventricular dyssynchrony than LVSP. However, this interventricular synchrony of LVSP may not be physiological. Instead of using the same stimulation marker, such as the pacing artifact, Curila et al. calculated interventricular dyssynchrony in SLBBP, NSLBBP, and LVSP as the difference between the first and last activation (15). There is no doubt that Stim-LVAT of LVSP is significantly longer than that of LBBP, implying that the LV excitation in LVSP occurs later than in LBBP. As a result, the improved interventricular synchronization of LVSP is attributable to greater overlap of LV and RV activation produced by delayed activation of both the LV and the RV (18).

Curila et al. only evaluated the LBBP with unipolar pacing configuration, not bipolar pacing configuration (15). Lin et al. developed a bilateral bundle branch area pacing strategy that involves stimulating the cathode and anode in various pacing configurations to capture both LBB and right bundle branch (RBB) area, which can diminish delayed right ventricular activation caused by LBBP and result in more physiological ventricular activation (19). It is essentially LBBP with bipolar pacing configuration (Figure 1C), with the cathode tip capturing LBB and the anode ring capturing RBB area. Shimeno et al. also revealed that the terminal R′/r′ wave duration of LBBP with bipolar pacing configuration is shorter than that of LVSP, presumably due to the contribution of the anodal capture during bipolar pacing (17). In addition, some previous studies and case reports have shown that LBBP can shorten the QRS duration of intrinsic RBBB or even completely correct RBBB (19–23), while LVSP cannot, but the underlying mechanism remains unclear and needs further study.

CONCLUSION

Compared with LVSP, LBBP is a more ideal pacing strategy for CRT, and many studies have confirmed its safety, stability, and efficacy. Future study will focus on how to diminish RBBB associated with LBBP in order to obtain better physiological interventricular synchrony. For example, adjusting the atioventricular delay to combined LV stimulation by LBBP with intrinsic RV excitation in patients with normal RBBB conduction, or modifying the interelectrode distance of pacing lead to better complete bilateral bundle branch area pacing in patients with RBBB. Although LVSP in close proximity to
LBB can be an alternative choice, clinically, this is essentially NSLBBP. The pacing output necessary to convert LVSP to NSLBBP, on the other hand, had not been investigated, and it was unknown if this output would have an adverse effect on pacemaker battery longevity. The long-term clinical effects of LVSP and LBBP remains unclear. Current studies solely examine the differences in electrophysiologic characteristics between LVSP and LBBP, such as Stim-LVAT, QRS duration, terminal R’ wave duration, QRS area, etc. In the future, it will be necessary to evaluate the echocardiographic activation of LVSP and LBBP, encompassing not only intraventricular synchronization, but also interventricular synchronization.

AUTHOR CONTRIBUTIONS

KZ wrote the original manuscript and conceptualized the idea. DC and QL supervised and wrote and edited the manuscript for publication. All authors contributed to the article and approved the submitted version.

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REFERENCES

1. Cleland JG, Daubert JC, Erdmann E, Freemantle N, Gras D, Kappenberger L, et al. The effect of cardiac resynchronization on morbidity and mortality in heart failure. N Engl J Med. (2005) 352:1539–49. doi: 10.1056/NEJMoa0504968
2. Abraham WT, Fisher WG, Smith AL, Delurgio DB, Leon AR, Loh E, et al. Cardiac resynchronization in chronic heart failure. N Engl J Med. (2002) 346:1845–53. doi: 10.1056/NEJMoa0131688
3. Cazeau S, Leclercq C, Lavergne T, Walker S, Varma C, Linde C, et al. Effects of multisite biventricular pacing in patients with heart failure and intraventricular conduction delay. N Engl J Med. (2001) 344:873–80. doi: 10.1056/NEJM200103223412024
4. Ploux S, Eschalier R, Whinnett ZL, Lumens J, Derval N, Sacher F, et al. Electrical dysynchrony induced by biventricular pacing: implications for patient selection and therapy improvement. Heart Rhythm. (2015) 12:782–91. doi: 10.1016/j.hrthm.2014.12.031
5. Birnie DH, Tang AS. The problem of non-response to cardiac resynchronization therapy. Curr Opin Cardiol. (2006) 21:20–6. doi: 10.1097/01.hco.0000198983.9375599
6. Mafi-Rad M, Luermans JG, Blaauw Y, Janssen M, Crijns HJ, Prinzen FW, et al. Feasibility and acute hemodynamic effect of left ventricular septal pacing by transvenous approach through the interventricular septum. Circ Arrhythm Electrophysiol. (2016) 9:e003344. doi: 10.1161/CIRCEP.115.003344
7. Huang W, Su L, Wu S, Xu L, Xiao F, Zhou X, et al. A novel pacing strategy with low and stable output: pacing the left bundle branch immediately beyond the conduction block. Can J Cardiol. (2017) 33:1736.e1731–e1733. doi: 10.1016/j.cjca.2017.09.013
8. Huang W, Wu S, Vijayaraman P, Su L, Chen X, Cai B, et al. Cardiac resynchronization therapy in patients with nonischemic cardiomyopathy using left bundle branch pacing. JACC Clin Electrophysiol. (2020) 6:849–58. doi: 10.1016/j.jacep.2020.04.011
9. Wu S, Su L, Vijayaraman P, Zheng R, Cai M, Xu L, et al. Left bundle branch pacing for cardiac resynchronization therapy: nonrandomized on-treatment comparison with his bundle pacing and biventricular pacing. Can J Cardiol. (2021) 37:319–28. doi: 10.1016/j.cjca.2020.04.037
10. Guo J, Li L, Xiao G, Ye T, Huang X, Meng F, et al. Remarkable response to cardiac resynchronization therapy via left bundle branch pacing in patients with true left bundle branch block. Clin Cardiol. (2020) 43:1460–8. doi: 10.1002/clc.23462
11. Wu S, Chen X, Wang S, Xu L, Xiao F, Huang Z, et al. Evaluation of the criteria to distinguish left bundle branch pacing from left ventricular septal pacing. JACC Clin Electrophysiol. (2021) 7:1166–77. doi: 10.1016/j.jacep.2021.02.018
12. Heckman LB, Luermans J, Curila K, Van Stipdonk AMW, Westra S, Smisek R, et al. Comparing ventricular synchrony in left bundle branch and left ventricular septal pacing in pacemaker patients. J Clin Med. (2021) 10:882. doi: 10.3390/jcm110404082
13. Jastrzebski M, Kiellbasa G, Curila K, Moskal P, Bednarek A, Rajzer M, et al. Physiology-based electrocardiographic criteria for left bundle branch capture. Heart Rhythm. (2021) 18:935–43. doi: 10.1016/j.hrthm.2021.02.021
14. Jastrzebski M, Kiellbasa G, Moskal P, Bednarek A, Kusiak A, Sondej T, et al. Fixation beats: a novel marker for reaching the left bundle branch area during deep septal lead implantation. Heart Rhythm. (2021) 18:562–9. doi: 10.1016/j.hrthm.2020.12.019
15. Curila K, Jurak P, Vernooij K, Jastrzebski M, Waldauf P, Prinzen F, et al. Left ventricular myocardial septal pacing in close proximity to LBB does not prolong the duration of the left ventricular lateral walldepolarization compared to LBB pacing. Front Cardiovasc Med. (2021) 8:787414. doi: 10.3389/fcvm.2021.787414
16. Vijayaraman P, Patel N, Colburn S, Beer D, Napierkowski A, Subrposha FA. His-Purkinje conduction system pacing in atrioventricular block: new insights into site of conduction block. JACC Clin Electrophysiol. (2022) 8:73–85. doi: 10.1016/j.jaecp.2021.07.007
17. Shimeno K, Tamura S, Nakatsuji K, Hayashi Y, Abe Y, Naruko T. Characteristics and proposed mechanisms of QRS morphology observed during the left bundle branch pacing procedure. Pacing Clin Electrophysiol. (2021) 44:1987–94. doi: 10.1111/pac.14382
18. Huang W, Zhou X, Ellenbogen KA. Pursue physiological pacing therapy: a better understanding of left bundle branch pacing and left ventricular septal myocardial pacing. Heart Rhythm. (2021) 18:1290–1. doi: 10.1016/j.hrthm.2021.05.013
19. Lin J, Chen K, Dai Y, Sun Q, Li Y, Jiang Y, et al. Bilateral bundle branch area pacing to achieve physiological conduction system activation. Circ Arrhythm Electrophysiol. (2020) 13:e008267. doi: 10.1161/CIRCEP.119.008267
20. Li X, Li H, Ma W, Ning X, Liang E, Pang K, et al. Permanent left bundle branch area pacing for atrioventricular block: feasibility, safety, and acute effect. Heart Rhythm. (2019) 16:1766–73. doi: 10.1016/j.hrthm.2019.04.043
21. Li X, Fan X, Li H, Ning X, Liang E, Ma W, et al. ECG patterns of successful permanent left bundle branch area pacing in bradycardia patients with typical bundle branch block. Pacing Clin Electrophysiol. (2020) 43:781–90. doi: 10.1111/pace.13982
22. Zhu K, Sun Y, Cai B, Li L, Li G, Liu J, et al. Left bundle branch pacing in patients with right bundle branch block. Kardiol Pol. (2021) 79:1127–9. doi: 10.33963/KP.a2021.0091
23. Zhu K, Lin M, Li L, Chang D, Li Q. Left bundle branch pacing shortened the QRS duration of a right bundle branch block. J Electrocardiol. (2021) 68:153–6. doi: 10.1016/j.jelectrocard.2021.08.018

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