

11-26-2020

Effects of left ventricular assist device on pulmonary functions and pulmonary hemodynamics: A meta-analysis

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World Journal of *Cardiology*

World J Cardiol 2020 November 26; 12(11): 513-598



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INDEXING/ABSTRACTING

The *WJC* is now abstracted and indexed in Emerging Sources Citation Index (Web of Science), PubMed, PubMed Central, Scopus, China National Knowledge Infrastructure (CNKI), China Science and Technology Journal Database (CSTJ), and Superstar Journals Database.

RESPONSIBLE EDITORS FOR THIS ISSUE

Production Editor: Jia-Hui Li; Production Department Director: Xiang Li; Editorial Office Director: Jia-Ping Yan.

NAME OF JOURNAL

World Journal of Cardiology

ISSN

ISSN 1949-8462 (online)

LAUNCH DATE

December 31, 2009

FREQUENCY

Monthly

EDITORS-IN-CHIEF

Ramdas G Pai, Dimitrios Tousoulis, Marco Matteo Ciccone

EDITORIAL BOARD MEMBERS

<https://www.wjgnet.com/1949-8462/editorialboard.htm>

PUBLICATION DATE

November 26, 2020

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<https://www.wjgnet.com/bpg/gerinfo/208>

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<https://www.wjgnet.com/bpg/gerinfo/242>

STEPS FOR SUBMITTING MANUSCRIPTS

<https://www.wjgnet.com/bpg/GerInfo/239>

ONLINE SUBMISSION

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Effects of left ventricular assist device on pulmonary functions and pulmonary hemodynamics: A meta-analysis

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Author contributions: Ullah W contributed to conceptualization, analysis, and manuscript writing; Meizinger C contributed to manuscript writing and data collection; Ali Z contributed to revision; Panchal A contributed to data curation; Saeed R contributed to editing; Haas DC and Rame E contributed to supervision.

Conflict-of-interest statement: The authors have no conflict of interest.

PRISMA 2009 Checklist statement: The authors have read the PRISMA 2009 Checklist, and the manuscript was prepared and revised according to the PRISMA 2009 Checklist.

Open-Access: This article is an open-access article that was selected by an in-house editor and fully peer-reviewed by external reviewers. It is distributed in accordance with the Creative Commons Attribution NonCommercial (CC BY-NC 4.0) license, which permits others to distribute, remix, adapt, build

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Abstract

BACKGROUND

Given current evidence, the effect of left ventricular assist device (LVAD) implantation on pulmonary function tests remains controversial.

AIM

To better understand the factors contributing to the changes seen on pulmonary function testing and the correlation with pulmonary hemodynamics after LVAD implantation.

METHODS

Electronic databases were queried to identify relevant articles. The summary effect size was estimated as a difference of overall means and standard deviation on a random-effects model.

RESULTS

A total of four studies comprising 219 patients were included. The overall mean forced expiratory volume in one second (FEV1), forced vital capacity (FVC) and diffusion lung capacity of carbon monoxide (DLCO) after LVAD implantation were significantly lower by 0.23 L (95%CI: 0.11-0.34, $P = 0.0002$), 0.18 L (95%CI: 0.03-0.34, $P = 0.02$), and 3.16 mmol/min (95%CI: 2.17-4.14, $P < 0.00001$), respectively. The net post-LVAD mean value of the cardiac index was significantly higher by 0.49 L/min/m² (95%CI: 0.31-0.66, $P < 0.00001$) compared to

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Manuscript source: Unsolicited manuscript

Specialty type: Cardiac and cardiovascular systems

Country/Territory of origin: United States

Peer-review report's scientific quality classification

Grade A (Excellent): 0
Grade B (Very good): 0
Grade C (Good): C, C, C, C
Grade D (Fair): D
Grade E (Poor): 0

Received: April 28, 2020

Peer-review started: April 28, 2020

First decision: July 5, 2020

Revised: July 18, 2020

Accepted: October 30, 2020

Article in press: October 30, 2020

Published online: November 26, 2020

P-Reviewer: Caceres-Loriga FM, Soriano-Ursúa MA, Teragawa H

S-Editor: Huang P

L-Editor: A

P-Editor: Wang LYT



pre-LVAD value. The pulmonary capillary wedge pressure and pulmonary vascular resistance were significantly reduced after LVAD implantation by 8.56 mmHg (95%CI: 3.78-13.35, $P = 0.0004$), and 0.83 Woods U (95%CI: 0.11-1.55, $P = 0.02$), respectively. There was no significant difference observed in the right atrial pressure after LVAD implantation (0.61 mmHg, 95%CI: -2.00 to 3.32, $P = 0.65$). Overall findings appear to be driven by studies using HeartMateII devices.

CONCLUSION

LVAD implantation might be associated with a significant reduction of the spirometric measures, including FEV1, FVC, and DLCO, and an overall improvement of pulmonary hemodynamics.

Key Words: Pulmonary function tests; Left ventricular assist device; Spirometry; Ventricular assist device

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Core Tip: Left ventricular assist device (LVAD) implantation can cause worsening of forced expiratory volume in one second, forced vital capacity, and diffusion lung capacity of carbon monoxide in post-LVAD patients. However, the benefits of LVAD implantation outweigh its risks in the context of pulmonary hemodynamics. More large scale studies are required to validate our findings.

Citation: Ullah W, Meizinger C, Ali Z, Panchal A, Saeed R, Haas DC, Rame E. Effects of left ventricular assist device on pulmonary functions and pulmonary hemodynamics: A meta-analysis. *World J Cardiol* 2020; 12(11): 550-558

URL: <https://www.wjgnet.com/1949-8462/full/v12/i11/550.htm>

DOI: <https://dx.doi.org/10.4330/wjc.v12.i11.550>

INTRODUCTION

Since the advent of the left ventricular assist device (LVAD) in 1988, literature has appropriately focused on its survival benefits in heart failure patients^[1,2]. Over the years, with an increase in its use, many LVAD related complications also started surfacing together. Arena *et al*^[3] in 1999 were the first to present data, implicating a decrease in spirometric measures in patients with LVAD implantation. These observation was based on the comparison of pre and post-LVAD findings in a 58-year old gentleman^[3]. Later on, more observational studies attempted to elucidate the complex interplay of pathophysiological dynamics between LVAD and pulmonary system, but had conflicting findings^[4-7]. Until recently, there has been no concrete evidence in the form of a randomized controlled trial (RCT) or meta-analysis to identify the net clinical benefit of LVAD in the context of worsened pulmonary functions and improved pulmonary hemodynamics.

MATERIALS AND METHODS

Search strategy and study collection

A literature search was performed up to December 2019, using PubMed, EMBASE, and Cochrane databases. There was no language or time restriction. The search strategies included various combinations of medical subject headings (MeSH) to generate two subsets of citations: One for LVAD and other for pulmonary function tests (PFTs). The terms from the two subsets were combined in 1:1 combination using boolean operators, and results from all possible combinations were screened for relevant articles. Based on our research question, articles from the reference lists pertinent to the clinical question were also evaluated by an independent author (backward snow bowling). Studies comparing changes in PFTs and pulmonary hemodynamics after LVAD implantation were included in the final analysis.

Pulmonary hemodynamics included cardiac output (CO), right atrial pressure (RAP) and pulmonary capillary wedge pressures (PCWP) assessment; pulmonary vascular resistance (PVR) was calculated by the standard formula $PVR = (mPAP - PCWP)/CO$. The studies with insufficient data, case reports, review articles, and conference papers were excluded after detailed discussion (Figure 1).

Quality of included studies

The overall methodological quality of the included studies was low. Due to the inclusion of single-center retrospective studies, the risk of selection bias could not be assessed. Similarly, randomization and allocation concealment of the subjects at the level of individual study could not be performed. Most studies, however, mentioned the baseline characteristics of the included population except by Imamura *et al*^[5]. The risk of reporting bias across all studies was minimal due to adequate reporting of outcomes. Only patients who had post-LVAD spirometry measurements were included in the study, minimizing the risk of attrition bias. All included studies used a self-control model in which the same population was assessed before and after an intervention (LVAD implantation); hence, the risk for detection bias was negligible. The detailed and summary bias graphs are given in Figure 2.

Statistical analysis

The statistical analysis was performed using the random-effects model (inverse variance) to calculate the mean difference and SD for continuous variables. The probability value of $P < 0.05$ was considered statistically significant. The “test for overall effect” was reported as z value corroborating the inference from the 95% CI. Higgins I^2 -squared (I^2) statistic model was used to assess variations in outcomes of the included studies. I^2 values of 50% or less corresponded to low to moderate, and 75% or higher indicated large amounts of heterogeneity. Publication bias was illustrated graphically using a funnel plot. The quality assessment of the included articles was performed using the Cochrane guidelines for the systematic review and meta-analysis, where each study was screened for five different types of bias (selection, performance, detection, attrition, and reporting bias). All statistical analysis was performed using the Cochrane Review Manager (RevMan) version 5.3.

RESULTS

Search results and study selection

The initial search on multiple databases revealed 1086 articles. After removal of irrelevant and duplicate items, 132 articles deemed relevant for full-text review. We further excluded 128 articles based on our selection criteria. Multiple attempts to authors to retrieve data were unsuccessful. Four articles qualified for final analysis^[4-7]. The Preferred Reporting Items for Systematic Reviews and Meta-Analyses (PRISMA) flow diagram is shown in Figure 1.

A total of 219 patients, with a mean age of 54 ± 11 years, were included. About 73% of the included subjects were males. Two thirds (74%) of the overall population had 2nd generation continuous flow devices (HeartMateII), while 26% of patients had a 3rd generation pulsatile flow device (HeartWare). The overall mean basal metabolic index was 26 ± 12 kg/m². The mean left ventricular ejection fraction was 10%. The indications for LVAD implantation were non-ischemic and ischemic cardiomyopathy in 62% (136/219) and 36% (79/219) of patients, respectively. Sajgalik *et al*^[6] were the only ones reporting hypertrophic cardiomyopathy in four patients. The functional status of the included population was defined either by a 6-min walk test (Raheja *et al*^[7]) or by classification of the New York Heart Association (NYHA) (Sajgalik *et al*^[6]). The mean brain nitric peptide level was 1695 pg/mL. Of the reported comorbidities, diabetes, chronic kidney disease, and hypertension were the most common. Approximately 22% (50/219) of the total population carried a diagnosis of chronic obstructive lung disease (COPD). The mean follow-up duration and timing of post-LVAD spirometry ranged from 6 mo to 12 mo. The detailed study characteristics are given in Supplementary Table 1.

Pooled estimate of spirometric changes with LVAD implantation

The overall mean forced expiratory volume in one second (FEV1) and forced vital capacity (FVC) values were significantly lower after LVAD implantation by 0.23 L (95% CI: 0.11-0.34, $P = 0.0002$) and 0.18 L (95% CI: 0.03-0.34, $P = 0.02$), respectively. The

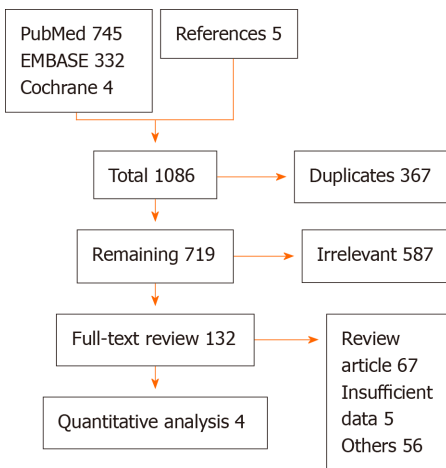


Figure 1 PRISMA flow diagram of the included studies.

	Random sequence generation (selection bias)	Allocation concealment (selection bias)	Blinding of participants and personnel (performance bias)	Blinding of outcome assessment (detection bias)	Incomplete outcome data (attrition bias)	Selective reporting (reporting bias)	Other bias
Imamura	+	+	+	+	+	+	+
Mohamedali				+	+	+	+
Raheja				+	+	+	+
Sajgalik				+	+	+	+

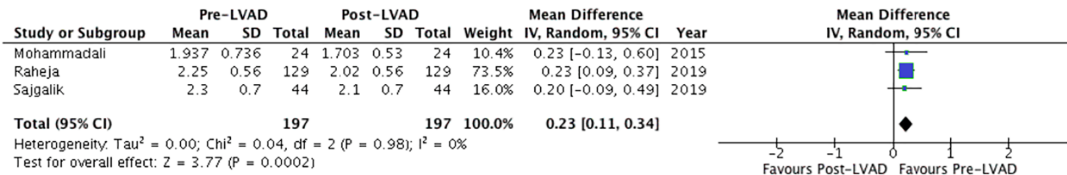
Figure 2 Detailed methodological quality and bias assessment of included studies.

heterogeneity among the outcomes of the included studies was minimal for both FEV1 and FVC with an $I^2 = 0\%$ ($P = 0.98$ and $P = 0.75$, respectively) (Figure 3A and B). The overall mean value of Δ DLCO (diffusion lung capacity of carbon monoxide) was robustly reduced by 3.16 mmol/min (95%CI: 2.17-4.14, $P < 0.00001$) in the post-LVAD state. There was no heterogeneity among the outcomes of individual studies $I^2 = 0\%$ ($P = 0.59$) (Figure 3C).

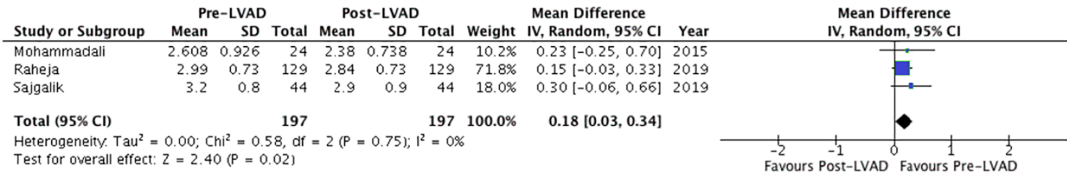
Pooled estimate of pulmonary hemodynamics with LVAD implantation

All indices of pulmonary hemodynamic observed an overall improvement with LVAD implantation. The net post-LVAD implantation mean cardiac index was significantly higher by 0.49 L/min/m² (95%CI: 0.31-0.66, $P < 0.00001$) compared to pre-LVAD values with no heterogeneity ($I^2 = 0$). Imamura *et al*^[5] study contributed more than half to the overall effect size (52% weight) (Figure 4A). Compared to pre-LVAD values, the pooled estimate of PCWP and PVR was significantly reduced after post-LVAD implantation by 8.56 mmHg (95%CI: 3.78-13.35, $P = 0.0004$), and 0.83 Woods U (95%CI: 0.11-1.55, $P = 0.02$), respectively (Figure 4B and C). There was no significant difference observed in RAP in the post-LVAD cohort both at the study level and in the pooled analysis (0.61 mmHg, 95%CI: -2.00 to 3.32, $P = 0.65$). The heterogeneity among the included studies was moderate ($I^2 = 76\%$, $I^2 = 42\%$, $I^2 = 57\%$, respectively) (Figure 4D).

A Δ FEV1



B Δ FVC



C Δ DLCO

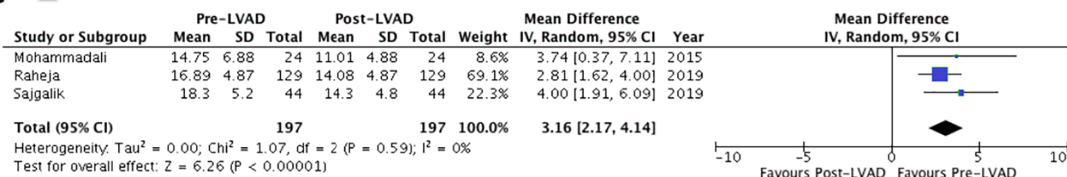


Figure 3 Forest plot depicting the pooled mean difference of forced expiratory volume in one second (A), forced vital capacity (B) and diffusion lung capacity of carbon monoxide (C) between pre- and post-left ventricular assist device patients. FEV1: Forced expiratory volume in one second; FVC: Forced vital capacity; DLCO: Diffusion lung capacity of carbon monoxide; LVAD: Left ventricular assist device.

DISCUSSION

Our meta-analysis shows a substantial decrease in the overall mean spirometric measures (FEV1, FVC, FEV1/FVC, and DLCO) in post-LVAD patients. The pulmonary hemodynamics were, however, significantly improved with an overall mean decrease in the PCWP and PVR. As expected, post-LVAD patients observed a substantial improvement in the cardiac index. Together these changes represented a net decrease in left ventricular filling pressure and an improvement in the left ventricular function. There was no effect seen on the net difference of RAP in patients following LVAD implantation. The detailed pre- and post-LVAD spirometric metrics are given in Figure 5.

Many factors could be accounted for the restrictive pulmonary pattern seen in post-LVAD patients. The initial transient decline in PFTs (FVC and FEV1) could be attributed to anesthesia effects or respiratory muscle weakness due to the major cardiothoracic surgery^[4]. Long term decrease in post-LVAD FVC can partly be explained by an LVAD patient now having an object that occupies approximately 50 mL (HeartWare) to 63 mL (HeartMateII) of intrathoracic space^[4]. In addition to the anatomical limitations, a constellation of mechanical and physiological mechanisms compromises the diaphragmatic functions, such as surgical implantation of HeartMateII below the diaphragm, sternotomy, thoracic scarring and respiratory muscle weakness due to cardiac cachexia^[7]. Moreover, heart failure induced reactive fibrosis, pleural effusion, and cardiomegaly can worsen the post-LVAD spirometric measures by mechanically impeding the lungs from expanding^[6]. Heart failure can also incite bronchial hypersensitivity or can cause interstitial edema, and compressive atelectasis, further compromising the spirometric metrics of post-LVAD patients. Some post-LVAD patients are at high risk of decline in spirometric measures if they have a compelling indication for pneumotoxic medications such as amiodarone^[4-7].

The literature on the post-LVAD pulmonary dynamics is scarce. Our extensive literature search identified four studies for quantitative analysis^[4-7]. All these studies unanimously agreed on the decline of pulmonary functions and the improvement of pulmonary hemodynamics in post-LVAD patients. These studies, however, should be interpreted in the context of its methodological limitations. Most studies used older generation devices (HeartMateII), while more contemporary studies used the newer

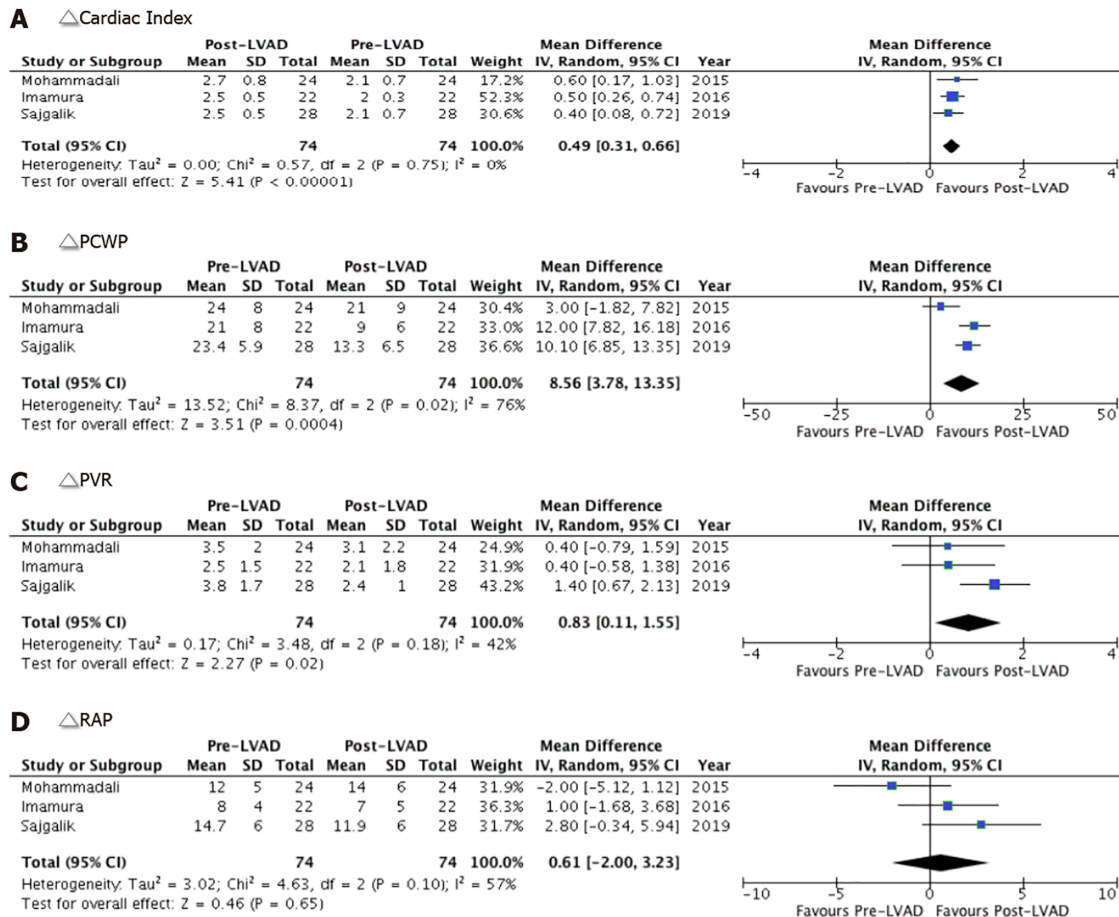


Figure 4 Forest plot showing pooled comparison of mean difference in cardiac index (A), pulmonary capillary wedge pressures (B), pulmonary vascular resistance (C), and right atrial pressure (D) between pre- and post-left ventricular assist device groups. PCWP: Pulmonary capillary wedge pressures; PVR: Pulmonary vascular resistance; RAP: Right atrial pressure; LVAD: Left ventricular assist device.

generation pulsatile LVAD devices (HeartWare). This potentially skewed the overall results of the studies. Mohamedali *et al*^[4] in 2015, were the first to recognize that a decrease in the spirometric measures (FEV1, FVC, and DLCO) was actually driven by the HeartMateII devices, rather than HeartWare. They attributed these findings to the surgical resection related paralysis of the diaphragm, post-procedure lung fibrosis, and restriction of the left hemidiaphragm by a relatively larger device size in the former^[4]. Raheja *et al*^[7] subsequently validated the overall results of the prior study. They also quantified the effect of LVAD on the functional status of heart failure patients using a six-minute walk distance (MWD). A significant correlation between the MWD and LVAD induced change in FEV1 ($P = 0.0466$), and DLCO ($P = 0.0032$) was demonstrated; however, these findings were never adjusted for the duration of follow up^[7]. Sajgalik *et al*^[6] hurdled these limitations by performing a stratified analysis based on different follow-up durations. Although a substantial amount of decline in FEV1 ($P = 0.04$) and FVC ($P = 0.01$) was observed at 12-mo, there was no significant decrease in PFTs on a shorter follow-up duration of 6-mo (FVC $P = 0.07$, FEV1 $P = 0.27$). Similarly, the observed predictive effect of survival with the change of DLCO at 6-mo ($P = 0.03$) was attenuated at an extended follow-up duration of 12-mo ($P = 0.22$)^[6].

Mohamedali *et al*^[4] exclusively used the 2nd generation (HeartMateII) devices, while about 77% LVADs in the Sajgalik *et al*^[6] and 88% of devices used by Raheja *et al*^[7] study were HeartMateII devices. Despite a higher percentage of HeartMateII devices in the later studies, the overall outcomes were never stratified based on the device type^[5-7]. Therefore, a clear distinction between the relative safety of HeartMateII and HeartWare could not be ascertained. This, along with the fact that all studies had substantial limitations due to non-randomized data, call into question the generalizability of individual results.

One can also argue that spirometric changes observed in post-LVAD patients could

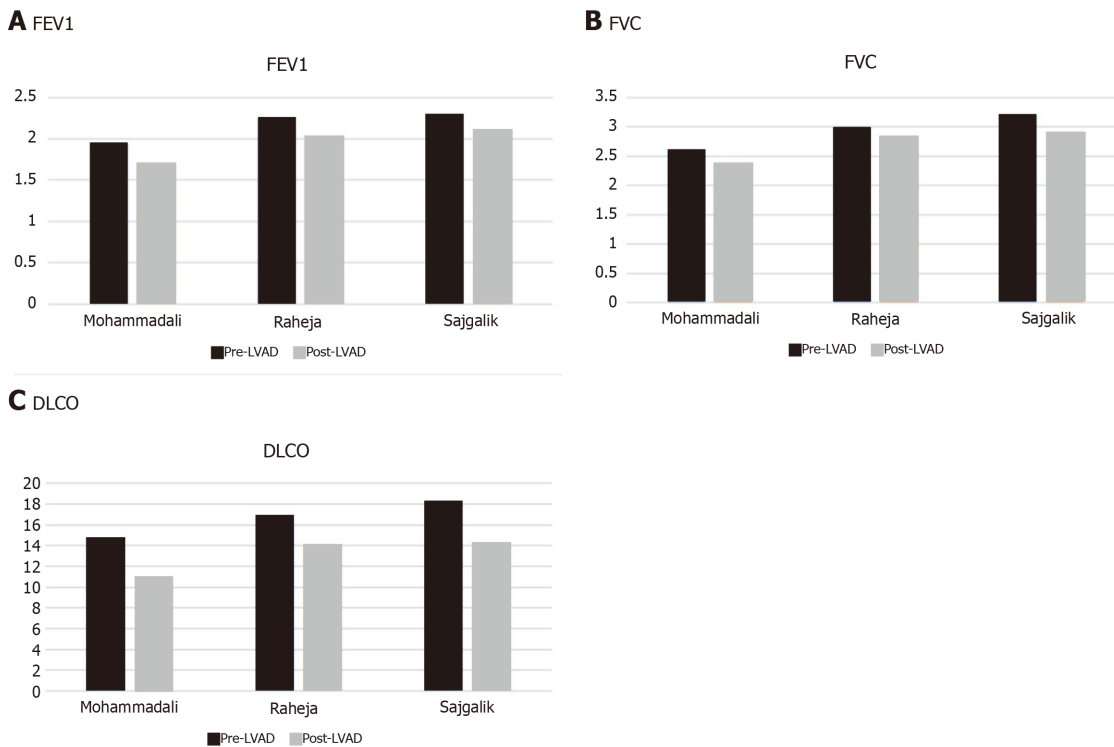


Figure 5 Trend of decrease in forced expiratory volume in one second (A), forced vital capacity (B), and diffusion lung capacity of carbon monoxide (C) in the included studies. FEV1: Forced expiratory volume in one second; FVC: Forced vital capacity; DLCO: Diffusion lung capacity of carbon monoxide; LVAD: Left ventricular assist device.

be a transient decline of the pre-existing lung disease, and might not have long-term consequences. As claimed by Imamura *et al*^[5] in his analysis, that the post-LVAD PFTs worsening is a reversible phenomenon with heart transplantation. However, more studies are needed to shed light on the long-term impacts of LVAD implantation on pulmonary dynamics. It should also be noted that, approximately 36% of patients from Sajgalik *et al*^[6] study had obstructive sleep apnea, while 29% of patients in Mohamedali *et al*^[4], and 22% of Raheja *et al*^[7] patients had COPD. It is unclear if the decrease of post-LVAD spirometric measures in the mentioned cohorts was an actual device effect or was merely worsening of their underlying lung condition.

Given apparent differences in the outcomes and significant effect of covariates on the results, one should be cautious about the clinical interpretations of individual results. This, along with the fact that previous small studies were either underpowered to detect clinical outcomes, or had conflicting findings, prompted us to integrate the findings of these studies.

In contrast to individual data, our meta-analysis provides strong evidence that LVAD related beneficial effects outweigh its potential spirometric decline in heart failure patients. A minimal mean decrease in post-LVAD measures of FEV1 and FCV by only 0.23 L and 0.18 L respectively is massively offset by a significantly higher increase in the cardiac index by 0.49 L/min/m², and improvement of PCWP by 8.56 mmHg and PVR by 0.83 Woods U, respectively. Our meta-analysis also illustrates the paucity of randomized data in post-LVAD patients and highlights the limits on our ability to draw definitive conclusions due to the retrospective nature of data.

Our study is constrained by the limitations of the available data. A significant barrier was our inability to perform a stratified subgroup analysis based on different LVAD types and variable follow up durations. The inherent heterogeneity in the inclusion criteria of studies could not be accounted for in the pooled analysis. This, in addition to the biases and confounding factors inherent in observational non-randomized studies, call for caution when interpreting the results of this meta-analysis.

To determine the actual effect of LVAD implantation on spirometric metrics, it is imperative to account for all the confounding variables in a controlled population. One way would be to determine the causation of lung findings on a prospective scale or conduction of a RCT. This will enable the researchers not only to assess the risk of post-LVAD spirometric changes but can also help in the identification of instantaneous

event rates such as mortality and survival benefits.

CONCLUSION

LVAD implantation might be associated with worsening of FEV1, FVC, and DLCO in post-LVAD patients. However, the overall benefits of LVAD in the context of pulmonary hemodynamics outweigh the presumed harm to spirometric metrics. More large scale studies are needed to validate our findings.

ARTICLE HIGHLIGHTS

Research background

Recent studies suggest that left ventricular assist device (LVAD) implantation has not been associated with an improvement in pulmonary function tests. However, the improvement seen in post-LVAD pulmonary hemodynamics outweighs the observed decrease in spirometry.

Research motivation

The studies investigating these parameters are not expansive and the overall methodological quality of the studies available is low. Further inquiry into the effects of the LVAD implantation on pulmonary hemodynamics, objective pulmonary function testing and on the observed clinical outcomes is needed.

Research objectives

This meta-analysis aims to stratify the observed outcomes in studies assessing these parameters, in order to better understand the factors contributing to the changes seen on pulmonary function testing and the correlation with pulmonary hemodynamics.

Research methods

Our study literature search was performed on published data until December 2019, using PubMed, EMBASE, and Cochrane databases. After screening the studies 132 articles deemed relevant were reviewed. 128 articles were excluded based on our selection criteria. Four studies were analysed and included in this meta-analysis.

Research results

A total of four studies comprising 219 patients were included. The overall mean forced expiratory volume in one second (FEV1), forced vital capacity (FVC) and diffusion lung capacity of carbon monoxide (DLCO) after LVAD implantation were significantly lower by 0.23 L (95% CI: 0.11-0.34, $P = 0.0002$), 0.18 L (95% CI: 0.03-0.34, $P = 0.02$), and 3.16 mmol/min (95% CI: 2.17-4.14, $P < 0.00001$), respectively. The pulmonary capillary wedge pressure and pulmonary vascular resistance were significantly reduced after LVAD implantation by 8.56 mmHg (95% CI: 3.78-13.35, $P = 0.0004$), and 0.83 Woods U (95% CI: 0.11-1.55, $P = 0.02$), respectively. There was no significant difference observed in the right atrial pressure after LVAD implantation (0.61 mmHg, 95% CI: -2.00 to 3.32, $P = 0.65$).

Research conclusions

LVAD implantation might be associated with a significant reduction of the spirometric measures, including FEV1, FVC, and DLCO, and an overall improvement of pulmonary hemodynamics.

Research perspectives

The short term and long-term effects of LVAD on the pulmonary hemodynamics on and pulmonary function tests need to be expanded and are essential in order to better assess outcomes. The need for randomized control trials exists to identify confounding factors that may affect the outcomes seen in the studies analyzed. Also, further studies with extended follow-up are needed to assess the clinical outcomes of the changes seen on PFTs and hemodynamics.

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