Published Online: 2025 March 30

Research Article



Correlation of LVEDP with Severity of Coronary Artery Involvement in Patients Referred to Modarres Hospital with Coronary Artery Disease

Morteza Safi¹, Fariba Bayat 🔟 ¹, Latif Gachkar 🔟 ², Mohammad Reza Shahri^{1,*}

¹ Cardiovascular Research Center, Shahid Beheshti University of Medical Sciences, Tehran, Iran
² Infectious Disease and Tropical Medicine Research Center, Shahid Beheshti University of Medical Sciences, Tehran, Iran

* Corresponding Author: Cardiovascular Research Center, Shahid Beheshti University of Medical Sciences, Tehran, Iran. Email: mrs.xoy@gmail.com

Received: 25 January, 2025; Accepted: 10 March, 2025

Abstract

Background: Diastolic dysfunction due to ischemia leads to heart failure and significantly affects hospitalization rates and mortality. Non-invasive evaluation of left ventricular end-diastolic pressure (LVEDP) provides crucial data about left ventricular diastolic function.

Objectives: This study aimed to investigate the correlation between the severity of coronary artery involvement and LVEDP parameters.

Methods: In this descriptive study, 90 patients referred to Modarres Hospital with a diagnosis of coronary artery disease (CAD) were evaluated for LVEDP parameters using echocardiography and angiography. The presence and severity of CAD were determined by cardiac catheterization and the Gensini score. After collecting patient clinical data, the correlation between the Gensini score and LVEDP was examined.

Results: The findings of the present study revealed a direct, weak, and significant correlation between LVEDP assessed by echocardiography and the Gensini score. There was also a significant, direct, and weak correlation between LVEDP assessed by angiography and the Gensini score. Additionally, a direct and strong correlation was observed between LVEDP assessed by echocardiography and angiography.

Conclusions: The present study demonstrated that LVEDP is significantly associated with the severity and extent of CAD. The LVEDP can be assessed non-invasively by echocardiography before and after percutaneous coronary intervention (PCI) without additional complications, and can therefore be used to rapidly assess improvements in diastolic function.

Keywords: Coronary Artery Disease, Diastolic Dysfunction, Echocardiography, LVEDP

1. Background

Coronary artery disease (CAD) is the leading cause of mortality globally, resulting from several subsequent issues such as hemodynamic problems or ventricular dysfunction (1). Coronary artery disease affects left ventricular (LV) systolic and diastolic function, leading to high LV filling pressures. Validating and comparing the correlation of CAD with different LV filling pressure waveforms may be helpful in the prognosis, diagnosis, and treatment of CAD (2). Diastolic dysfunction leads to ineffective emptying of the left atrium and filling of the left ventricle, reducing the ability to increase cardiac output, increasing pulmonary pressure, and resulting in symptoms of heart failure (3). Diastolic dysfunction significantly affects mortality and hospitalization and contributes to the development of heart failure, hospitalization, and death (4, 5). Compared to systolic function, diastolic function has a longer recovery period and is more susceptible to ischemia (6).

Non-invasive assessment of LV diastolic function by left ventricular end-diastolic pressure (LVEDP) using transmitral doppler echocardiography and tissue doppler imaging provides clinicians with important information about LV diastolic function. However, both methods have relatively low sensitivity and specificity (7).

In individuals with CAD, evaluating LVEDP provides an assessment of hemodynamic status and helps guide appropriate management and therapeutic interventions (8). This parameter is assessed using

Copyright © 2025, International Journal of Cardiovascular Practice. This open-access article is available under the Creative Commons Attribution-NonCommercial 4.0 (CC BY-NC 4.0) International License (https://creativecommons.org/licenses/by-nc/4.0/), which allows for the copying and redistribution of the material only for noncommercial purposes, provided that the original work is properly cited. various techniques, including invasive (cardiac catheterization) and non-invasive (echocardiography) approaches (9, 10). The main advantage of doppler echocardiography assessments is their ability to noninvasively calculate hemodynamic parameters. In this regard, the calculation of LVEDP, which includes early diastolic parameters, has been proposed (11). Abnormal LV diastolic filling, as assessed by transthoracic echocardiography (TTE), is associated with a worse prognosis (12, 13). Elevated LVEDP and abnormal LV relaxation have been observed in patients who underwent invasive evaluation for flow-limiting coronary artery stenosis (14, 15). To date, the correlation of non-invasively measured CAD with LVEDP has not been well evaluated. Increased ischemic load delays diastolic filling independently of clinical variables and ejection fraction (EF) (16). Indeed, subclinical atherosclerosis might induce subclinical ischemia, resulting in diastolic filling abnormalities (17, 18). It has been proposed that hypertrophy and thickening of the LV wall play a compensatory role in LV function preservation. On the other hand, diastolic dysfunction might be a non-anginal manifestation of CAD. Therefore, CAD may be a false target for treatment in patients with elevated LVEDP and consequent dyspnea (18).

2. Objectives

Therefore, our study aimed to examine the correlation between the severity of coronary artery involvement and LVEDP parameters, which may improve the understanding of the pathophysiological relationship between CAD and diastolic dysfunction.

3. Methods

In the present descriptive cross-sectional study, 90 patients were consecutively selected from those diagnosed with CAD and referred to the heart department or clinic of Modarres Hospital, Tehran, Iran, for angiography in 2023. Inclusion criteria included a diagnosis of CAD, indication for angiography, age over 18 years, and willingness to participate in the study. Patients with a history of acute myocardial infarction, congestive heart failure, hypertrophic cardiomyopathy, left ventricular hypertrophy (LVH) more severe than mild. valvular heart disease more severe than moderate. and congenital heart disease were excluded from the investigation. The study was approved by the Ethics Committee of Shahid Beheshti University of Medical Sciences (IR.SBMU.RETECH.REC.1402.441). The LVEDP assessment was performed using echocardiography and angiography. Echocardiographic assessment was

conducted by a cardiologist and professor of echocardiography. The presence and severity of CAD were determined by an interventional specialist using cardiac catheterization. Additionally, demographic information was collected based on medical history after obtaining patient consent. Therefore, the data collection technique was observational.

3.1. Coronary Angiography

The CAD was assessed by coronary angiography (CAG) based on the maximum lumen stenosis and was defined as the presence of at least one stenosis of 50% or more in at least one of the 15 coronary segments of the three main coronary arteries (19).

3.2. Gensini Score

The severity of CAD was evaluated using the Gensini score (20). The Gensini score was calculated by assigning a severity score to each coronary stenosis based on the degree of lumen stenosis and its significance. A reduction in lumen diameter of 25%, 50%, 75%, 90%, and 99%, as well as complete occlusion, were scored as 1, 2, 4, 8, 16, and 32, respectively. Each main vascular segment was assigned a coefficient based on the functional importance of the myocardial region, including a coefficient of 5 for the left main (LM) artery, 2.5 for the proximal segment of the left anterior descending (LAD) artery. 2.5 for the proximal segment of the left circumflex (LCX) artery, 1.5 for the mid-portion of the LAD, 1 for the right coronary artery (RCA) and distal part of the LAD, the posterolateral artery, and the solitary marginal artery, and 0.5 for the other vascular segments.

3.3. Statistical Analysis

Statistical analyses were performed using SPSS software, version 22. In the descriptive section, quantitative parameters were expressed as mean and standard deviation, while qualitative parameters were presented as number and percentage. The Pearson correlation test was conducted to evaluate the correlation between data. In all tests, a P-value of less than 0.05 was considered significant.

4. Results

4.1. Patient Demographic and Clinical Characteristics

In this study, 90 patients with CAD were examined, with an age range of 36 to 83 years. Among them, 52 (57.8%) were male. The distribution of qualitative and quantitative characteristics of patients based on

uriables	Values
ualitative Parameters	
Grouping (Gensini score)	
1	47(52.2)
2	14 (15.6)
4	10 (11.1)
8	6 (6.7)
16	10 (11.1)
32	3 (3.3)
Gender (male)	52 (57.8)
Comorbidities (yes)	84 (93.3)
CAD (yes)	52 (57.8)
Hypertension (yes)	67 (74.4)
Diabetes mellitus (yes)	26 (28.9)
Hyperlipidemia (yes)	44 (48.9)
Smoking (yes)	34 (37.8)
Familial history (yes)	24 (26.7)
uantitative Parameters	
LVEDP via catheterization (mmHg)	17.16 ± 3.18 (10 - 23)
LVEDP via echocardiography (mmHg)	14.75 ± 2.80 (9 - 20)
LVEF (%)	56.33±2.68 (50-65)
DBP (mmHg)	73 ± 8.92 (55 - 95)
SBP (mmHg)	119.83 ± 14.94 (90 - 155)
Age (y)	61.63 ± 10.63 (36 - 83)

Abbreviations: LVEDP, left ventricular end-diastolic pressure; CAD, coronary artery disease.

^a Values are expressed as No. (%) or mean ± SD (min - max).

clinical, echocardiographic, and angiographic findings is presented in Table 1.

4.2. Correlation of LVEDP with Patients' Gensini Score

As shown in Table 2, the correlation between LVEDP and patients' Gensini scores was examined. Analyses using the Spearman test revealed a direct and weak correlation between LVEDP assessed by echocardiography and the Gensini score, which was statistically significant. There was also a significant, direct, and weak correlation between LVEDP assessed by angiography and the Gensini score (Figure 1A). Additionally, as a secondary finding, a direct and strong correlation was observed between LVEDP assessed by echocardiography and angiography (Figure 1B).

5. Discussion

Diastole is an energy-dependent process and requires an adequate energy source (21). In myocardial ischemia, the energy source is decreased or becomes unavailable. Diastolic function is generally more vulnerable to injury than systolic function, meaning that diastolic dysfunction often emerges before systolic dysfunction and tends to persist longer during episodes of ischemia (22). In individuals with CAD, areas experiencing ischemia exhibit heightened myocardial stiffness. This increased stiffness, combined with a reduction in wall thinning and active pressure during ischemic events, leads to alterations in the thickness and pressure of the left ventricular wall as well as changes in the left ventricular pressure-volume relationships, ultimately raising LVEDP. As a result, CAD has a more significant effect on diastolic dysfunction (14).

In this study, we aimed to evaluate the correlation of LVEDP parameters with the severity of coronary artery involvement in CAD patients. According to the main findings, the Gensini score showed a weak, direct correlation between LVEDP assessed by echocardiography and angiography. It was also observed that there was a strong, direct correlation between LVEDP assessed by echocardiography and angiography. The occurrence of preclinical diastolic dysfunction in the overall adult population is estimated to range from 20% to 30%. This prevalence increases with

Parameters	R Coefficient	P-Value
LVEDP via catheterization vs. Gensini score	0.5	0.0001
LVEDP via echocardiography vs. Gensini score	0.45	0.0001
LVEDP via echocardiography vs. LVEDP via catheterization	0.9	0.0001

advancing age and is further heightened by factors such as CAD, cardiovascular disease, and diabetes, all of which act as independent risk factors for the development of diastolic dysfunction (23).

Diastolic dysfunction describes the atypical mechanical characteristics of the myocardium and includes abnormal diastolic dilation of the left ventricle, impaired filling, chamber stiffness, and slow or delayed relaxation (24). Physiologically, any mechanism that interferes with the dissociation of the actin-myosin cross-bridge or the removal of calcium from the cytosol can delay relaxation. Individuals with impaired diastolic function are at increased risk for developing heart failure with preserved ejection fraction (HFpEF). The likelihood of progressing to HFpEF is particularly elevated for those who also suffer from conditions such as anemia, hypertension, diabetes, renal dysfunction, or CAD (25).

Left ventricular ejection fraction (LVEF), which provides a visual assessment of systolic function, has been utilized to categorize individuals recovering from myocardial infarction; however, it is not a perfect measure. Because survival rates are similar between populations with heart HFpEF and heart failure with reduced ejection fraction (HFrEF), it remains unknown whether a more integrated physiologic measure of total LV function, such as LVEDP, would better predict postmyocardial infarction heart failure. A review of three studies, including a total of 5,372 patients, found that elevated LVEDP was linked to more unfavorable outcomes after myocardial infarction, which included higher incidences of heart failure and greater mortality rates. While a lower LVEF is a well-established predictor of negative outcomes in myocardial infarction patients, those with both reduced and preserved LVEF also experience poorer outcomes, raising questions about the full effectiveness of LVEF as a predictor of long-term outcomes compared with LVEDP (26-28).

Du et al. showed that there was a significant difference in LVEDP between CAD and non-CAD groups and that LVEDP was independently associated with CAD. In different subgroups, LVEDP increased with the increasing number of occluded vessels and showed a positive correlation with the Gensini score. In the non-CAD group, LVEDP was only associated with age, not the Gensini score (29). Aging alters LV diastolic function with increased LV stiffness, increased myocardial fibrosis, decreased filling velocity and amplitude, and ultimately leads to impaired LV diastolic function (30). Therefore, clinical attention should be paid to the underlying diastolic dysfunction in the elderly.

Ren et al. conducted a study involving 693 patients with CAD and discovered that 36% exhibited mild to severe LV diastolic dysfunction. Additionally, they identified that moderate to severe LV diastolic dysfunction was a significant predictor of hospitalization due to heart failure and mortality from heart disease (31). Chronic ischemia can result in diastolic wall motion abnormalities (32). Therefore, it is important to recognize the existence of diastolic dysfunction in patients with CAD.

Lin et al. demonstrated that both obstructive and non-obstructive CAD identified through coronary CT angiography were correlated with elevated LVEDP. Their findings indicated that LVEDP rose with the increasing number of affected vessels and exhibited a positive correlation with the Gensini score, which assesses the extent and severity of CAD. This suggests that a greater burden of coronary disease may contribute to worsening diastolic function and increased pressure in the left ventricle (18). A recent investigation revealed a significant association between the degree of CAD severity and decreased LV compliance (33).

Paul et al. showed that induced LV diastolic dysfunction persists long after the resolution of the ischemic episode (34). The occurrence and degree of diastolic dysfunction are influenced by the level of ischemia. Similarly, impaired diastolic function can serve as an indicator of the severity of ischemic conditions. Perrone-Filardi et al. demonstrated that in patients with CAD, normal resting LV systolic function combined with impaired LV filling is associated with a higher likelihood of ischemia, suggesting a larger area of myocardium at risk (35). Additionally, another investigation found that patients exhibiting impaired LV relaxation tend to have more severe CAD (36).

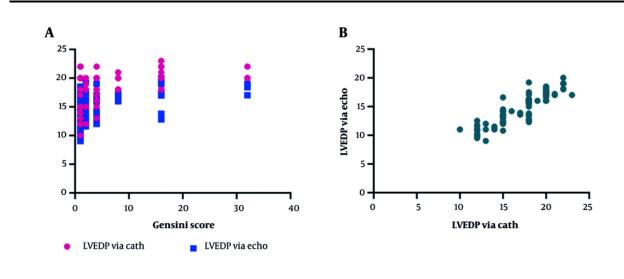


Figure 1. Correlation analysis. A, left ventricular end-diastolic pressure (LVEDP) assessed by echocardiography and angiography with Gensini score; B, LVEDP assessed by echocardiography and angiography.

A recent study revealed a positive correlation between LVEDP and the levels of troponin T, creatine kinase myocardial band (CKMB), and creatine kinase (CK) (37). This finding can be attributed to the fact that diastolic dysfunction of the ventricle resulting from myocardial infarction (MI) leads to an elevation in LVEDP. A study by Kirtane et al. reported an association between high LVEDP, longer hospital stays, and higher rates of heart failure 30 days after ST-elevation myocardial infarction (STEMI). Patients with LVEDP greater than 24 mmHg have been associated with a poorer prognosis and increased mortality (26). These studies indicate that LVEDP is a significant and independent predictor of readmission and prognosis in patients who have experienced a STEMI (38, 39).

Furthermore, research by Şatıroğlu et al. suggests that STEMI can lead to a decrease in left ventricular compliance, resulting in increased LVEDP and impaired left ventricular diastolic function (40). This highlights the importance of LVEDP as a prognostic indicator and its relationship with the pathophysiological changes that occur in the heart following a STEMI. In line with our study, another investigation reported that patients exhibiting high LVEDP were more likely to present with perfusion defects on nuclear scans compared to those with normal LVEDP levels (41). This finding suggests that elevated LVEDP may be associated with impaired myocardial perfusion, reflecting underlying cardiac dysfunction. Such results reinforce the notion that LVEDP can serve as an important marker for assessing coronary perfusion status and overall cardiac health in

patients, particularly following events such as STEMI. This relationship further underscores the prognostic value of LVEDP in the clinical setting.

The study had some limitations. The patients studied were all clinically suspected of having CAD, which could bias the results to some extent. Additionally, there were unknown confounding factors that may affect the results. Including a control group without CAD would provide more reliable results. A comprehensive assessment of systolic and diastolic function in a large, multicenter cohort followed longitudinally with a focused echocardiographic examination in patients with CAD at all stages and severities is recommended. Furthermore, despite the attempt to examine important demographic aspects related to patients, some determinants such as the type of medication used were not considered. It is recommended that these factors be considered in future studies.

5.1. Conclusions

The present study showed that LVEDP is significantly associated with the severity and extent of CAD. Clinicians should pay attention to diastolic function in patients, especially those with CAD. These findings may have potentially important clinical and therapeutic implications. Given the high correlation between LVEDP values measured by echocardiography and angiography, LVEDP can be measured non-invasively by echocardiography before and after percutaneous coronary intervention (PCI) without any complications and can therefore be used to rapidly assess improvement in diastolic function.

Footnotes

Authors' Contribution: Study concept and design: M. S. and M. R. S.; Acquisition of data: M. R. S.; Analysis and interpretation of data: F. B.; Drafting of the manuscript: M. R. S.; Critical revision of the manuscript for important intellectual content: All authors.; Statistical analysis: L. F.; Administrative, technical, and material support: F. B.; Study supervision: M. S.

Conflict of Interests Statement: The authors declared no conflict of interests.

Data Availability: The dataset presented in the study is available on request from the corresponding author during submission or after its publication.

Ethical Approval: The study was approved by the Ethics Committee of the Shahid Beheshti University of Medical Sciences (IR.SBMU.RETECH.REC.1402.441).

Funding/Support: There was no funding to be declared.

Informed Consent: Informed consent was obtained from all participants.

References

- Rosamond WD. Invited commentary: trends in coronary heart disease mortality-location, location, location. Am J Epidemiol. 2003;157(9):771-3. [PubMed ID: 12727670]. https://doi.org/10.1093/aje/kwg058.
- Barakat A, Amar A, Alsaadi AR. The correlation between coronary artery disease and left ventricular filling pressure: which correlates more LVEDP or LV pre-A wave? *Eur Heart J.* 2022;43(Suppl 1):ehab849.069. https://doi.org/10.1093/eurheartj/ehab849.069.
- El Aidi H, Adams A, Moons KG, Den Ruijter HM, Mali WP, Doevendans PA, et al. Cardiac magnetic resonance imaging findings and the risk of cardiovascular events in patients with recent myocardial infarction or suspected or known coronary artery disease: a systematic review of prognostic studies. J Am Coll Cardiol. 2014;63(11):1031-45. [PubMed ID: 24486280]. https://doi.org/10.1016/j.jacc.2013.11.048.
- Hogg K, Swedberg K, McMurray J. Heart failure with preserved left ventricular systolic function; epidemiology, clinical characteristics, and prognosis. J Am Coll Cardiol. 2004;43(3):317-27. [PubMed ID: 15013109]. https://doi.org/10.1016/j.jacc.2003.07.046.
- Kane GC, Karon BL, Mahoney DW, Redfield MM, Roger VL, Burnett Jr JC, et al. Progression of left ventricular diastolic dysfunction and risk of heart failure. *JAMA*. 2011;**306**(8):856-63. [PubMed ID: 21862747]. [PubMed Central ID: PMC3269764]. https://doi.org/10.1001/jama.2011.1201.
- 6. Mahmarian JJ, Pratt CM. Silent myocardial ischemia in patients with coronary artery disease. Possible links with diastolic left ventricular

dysfunction. *Circulation*. 1990;**81**(2 Suppl):III33-40. [PubMed ID: 2404637].

- Hajahmadi Poorrafsanjani M, Rahimi Darabad B. Evaluate the sensitivity and specificity echocardiography in trans-Doppler and tissue Doppler method in the estimation of left ventricular enddiastolic pressure. *Glob J Health Sci.* 2014;6(7 Spec No):92-7. [PubMed ID: 25363184]. [PubMed Central ID: PMC4796497]. https://doi.org/10.5539/gjhs.v6n7p92.
- Nagueh SF, Sun H, Kopelen HA, Middleton KJ, Khoury DS. Hemodynamic determinants of the mitral annulus diastolic velocities by tissue Doppler. J Am Coll Cardiol. 2001;37(1):278-85. [PubMed ID: 11153752]. https://doi.org/10.1016/s0735-1097(00)01056-1.
- Baan J, van der Velde ET, de Bruin HG, Smeenk GJ, Koops J, van Dijk AD, et al. Continuous measurement of left ventricular volume in animals and humans by conductance catheter. *Circulation*. 1984;**70**(5):812-23. [PubMed ID: 6386218]. https://doi.org/10.1161/01.cir.70.5.812.
- Tanaka N, Dalton N, Mao L, Rockman HA, Peterson KL, Gottshall KR, et al. Transthoracic echocardiography in models of cardiac disease in the mouse. *Circulation*. 1996;**94**(5):1109-17. [PubMed ID: 8790053]. https://doi.org/10.1161/01.cir.94.5.1109.
- Mulvagh S, Quinones MA, Kleiman NS, Cheirif J, Zoghbi WA. Estimation of left ventricular end-diastolic pressure from Doppler transmitral flow velocity in cardiac patients independent of systolic performance. *J Am Coll Cardiol*. 1992;20(1):112-9. [PubMed ID: 1607511]. https://doi.org/10.1016/0735-1097(92)90146-e.
- 12. Owan TE, Hodge DO, Herges RM, Jacobsen SJ, Roger VL, Redfield MM. Trends in prevalence and outcome of heart failure with preserved ejection fraction. *N Engl J Med.* 2006;**355**(3):251-9. [PubMed ID: 16855265]. https://doi.org/10.1056/NEJMoa052256.
- Ahmed A, Perry GJ, Fleg JL, Love TE, Goff Jr DC, Kitzman DW. Outcomes in ambulatory chronic systolic and diastolic heart failure: a propensity score analysis. *Am Heart J.* 2006;**152**(5):956-66. [PubMed ID: 17070167]. [PubMed Central ID: PMC2628474]. https://doi.org/10.1016/j.ahj.2006.06.020.
- Bourdillon PD, Lorell BH, Mirsky I, Paulus WJ, Wynne J, Grossman W. Increased regional myocardial stiffness of the left ventricle during pacing-induced angina in man. *Circulation*. 1983;67(2):316-23. [PubMed ID: 6848219]. https://doi.org/10.1161/01.cir.67.2.316.
- Miyazaki S, Guth BD, Miura T, Indolfi C, Schulz R, Ross Jr J. Changes of left ventricular diastolic function in exercising dogs without and with ischemia. *Circulation*. 1990;81(3):1058-70. [PubMed ID: 2407371]. https://doi.org/10.1161/01.cir.81.3.1058.
- Nakajima Y, Kane GC, McCully RB, Ommen SR, Pellikka PA. Left ventricular diastolic filling pressures during dobutamine stress echocardiography: relationship to symptoms and ischemia. *J Am Soc Echocardiogr.* 2009;22(8):947-53. [PubMed ID: 19524401]. https://doi.org/10.1016/j.echo.2009.04.030.
- Mehta SK, Rame JE, Khera A, Murphy SA, Canham RM, Peshock RM, et al. Left ventricular hypertrophy, subclinical atherosclerosis, and inflammation. *Hypertension*. 2007;**49**(6):1385-91. [PubMed ID: 17404181]. https://doi.org/10.1161/HYPERTENSIONAHA.107.087890.
- Lin FY, Zemedkun M, Dunning A, Gomez M, Labounty TM, Asim M, et al. Extent and severity of coronary artery disease by coronary CT angiography is associated with elevated left ventricular diastolic pressures and worsening diastolic function. *J Cardiovasc Comput Tomogr.* 2013;7(5):289-96 e1. [PubMed ID: 24268115]. https://doi.org/10.1016/j.jcct.2013.08.008.
- Qing X, Furong W, Yunxia L, Jian Z, Xuping W, Ling G. Cystatin C and asymptomatic coronary artery disease in patients with metabolic syndrome and normal glomerular filtration rate. *Cardiovasc Diabetol.* 2012;**11**:108. [PubMed ID: 22978689]. [PubMed Central ID: PMC3473246]. https://doi.org/10.1186/1475-2840-11-108.

- Gensini GG. A more meaningful scoring system for determining the severity of coronary heart disease. *Am J Cardiol.* 1983;51(3):606. [PubMed ID: 6823874]. https://doi.org/10.1016/s0002-9149(83)80105-2.
- 21. Wijns W, Serruys PW, Slager CJ, Grimm J, Krayenbuehl HP, Hugenholtz PG, et al. Effect of coronary occlusion during percutaneous transluminal angioplasty in humans on left ventricular chamber stiffness and regional diastolic pressure-radius relations. *J Am Coll Cardiol*. 1986;**7**(3):455-63. [PubMed ID: 2936788]. https://doi.org/10.1016/s0735-1097(86)80453-3.
- Amano J, Thomas Jr JX, Lavallee M, Mirsky I, Glover D, Manders WT, et al. Effects of myocardial ischemia on regional function and stiffness in conscious dogs. *Am J Physiol*. 1987;**252**(1 Pt 2):H110-7. [PubMed ID: 3812705]. https://doi.org/10.1152/ajpheart.1987.252.1.H110.
- 23. Wan SH, Vogel MW, Chen HH. Pre-clinical diastolic dysfunction. *J Am Coll Cardiol.* 2014;**63**(5):407-16. [PubMed ID: 24291270]. [PubMed Central ID: PMC3934927]. https://doi.org/10.1016/j.jacc.2013.10.063.
- Deswal A. Diastolic dysfunction and diastolic heart failure: mechanisms and epidemiology. *Curr Cardiol Rep.* 2005;7(3):178-83. [PubMed ID: 15865857]. https://doi.org/10.1007/s11886-005-0074-7.
- Kass DA, Bronzwaer JG, Paulus WJ. What mechanisms underlie diastolic dysfunction in heart failure? *Circ Res.* 2004;**94**(12):1533-42. [PubMed ID: 15217918]. https://doi.org/10.1161/01.RES.0000129254.25507.d6.
- Kirtane AJ, Bui A, Murphy SA, Karmpaliotis D, Kosmidou I, Boundy K, et al. Association of epicardial and tissue-level reperfusion with left ventricular end-diastolic pressures in ST-elevation myocardial infarction. J Thromb Thrombolysis. 2004;17(3):177-84. [PubMed ID: 15353915]. https://doi.org/10.1023/B:THRO.0000040486.10549.f6.
- Bagai A, Armstrong PW, Stebbins A, Mahaffey KW, Hochman JS, Weaver WD, et al. Prognostic implications of left ventricular enddiastolic pressure during primary percutaneous coronary intervention for ST-segment elevation myocardial infarction: Findings from the Assessment of Pexelizumab in Acute Myocardial Infarction study. *Am Heart J.* 2013;**166**(5):913-9. [PubMed ID: 24176448]. https://doi.org/10.1016/j.ahj.2013.08.006.
- Planer D, Mehran R, Witzenbichler B, Guagliumi G, Peruga JZ, Brodie BR, et al. Prognostic utility of left ventricular end-diastolic pressure in patients with ST-segment elevation myocardial infarction undergoing primary percutaneous coronary intervention. *Am J Cardiol.* 2011;**108**(8):1068-74. [PubMed ID: 21798494]. https://doi.org/10.1016/j.amjcard.2011.06.007.
- Du LJ, Dong PS, Jia JJ, Fan XM, Yang XM, Wang SX, et al. Association between left ventricular end-diastolic pressure and coronary artery disease as well as its extent and severity. *Int J Clin Exp Med.* 2015;8(10):18673-80. [PubMed ID: 26770481]. [PubMed Central ID: PMC4694381].
- Bonow RO, Vitale DF, Bacharach SL, Maron BJ, Green MV. Effects of aging on asynchronous left ventricular regional function and global ventricular filling in normal human subjects. *J Am Coll Cardiol.* 1988;11(1):50-8. [PubMed ID: 3335706]. https://doi.org/10.1016/0735-1097(88)90166-0.
- 31. Ren X, Ristow B, Na B, Ali S, Schiller NB, Whooley MA. Prevalence and prognosis of asymptomatic left ventricular diastolic dysfunction in

ambulatory patients with coronary heart disease. *Am J Cardiol.* 2007;**99**(12):1643-7. [PubMed ID: 17560867]. [PubMed Central ID: PMC2778467]. https://doi.org/10.1016/j.amjcard.2007.01.041.

- Husic M, Norager B, Egstrup K, Lang RM, Moller JE. Diastolic wall motion abnormality after myocardial infarction: relation to neurohormonal activation and prognostic implications. *Am Heart J.* 2005;**150**(4):767-74. [PubMed ID: 16209980]. https://doi.org/10.1016/j.ahj.2004.11.024.
- Strauer BE, Bolte HD, Heimburg P, Riecker G. [Coronary disease. II. Analysis of diastolic pressure-volume correlations and left ventricular elasticity in 110 patients]. *Z Kardiol.* 1975;64(4):311-22. DE. [PubMed ID: 1210520].
- 34. Paul AK, Kusuoka H, Hasegawa S, Yonezawa T, Makikawa M, Nishimura T. Prolonged diastolic dysfunction following exercise induced ischaemia: a gated myocardial perfusion SPECT study. Nucl Med Commun. 2002;23(11):1129-36. [PubMed ID: 12411843]. https://doi.org/10.1097/00006231-200211000-00014.
- Perrone-Filardi P, Bacharach SL, Dilsizian V, Bonow RO. Impaired left ventricular filling and regional diastolic asynchrony at rest in coronary artery disease and relation to exercise-induced myocardial ischemia. *Am J Cardiol.* 1991;67(5):356-60. [PubMed ID: 1994658]. https://doi.org/10.1016/0002-9149(91)90041-i.
- Fukuta H, Ohte N, Wakami K, Goto T, Tani T, Kimura G. Prognostic value of left ventricular diastolic dysfunction in patients undergoing cardiac catheterization for coronary artery disease. *Cardiol Res Pract.* 2012;2012:243735. [PubMed ID: 22567531]. [PubMed Central ID: PMC3332169]. https://doi.org/10.1155/2012/243735.
- Zhou X, Lei M, Zhou D, Li G, Duan Z, Zhou S, et al. Clinical factors affecting left ventricular end-diastolic pressure in patients with acute ST-segment elevation myocardial infarction. *Ann Palliat Med.* 2020;9(4):1834-40. [PubMed ID: 32279513]. https://doi.org/10.21037/apm.2020.03.22.
- Saito D, Nakanishi R, Watanabe I, Yabe T, Okubo R, Amano H, et al. Combined assessment of left ventricular end-diastolic pressure and ejection fraction by left ventriculography predicts long-term outcomes of patients with ST-segment elevation myocardial infarction. *Heart Vessels*. 2018;33(5):453-61. [PubMed ID: 29143103]. https://doi.org/10.1007/s00380-017-1080-6.
- Brienesse SC, Davies AJ, Khan A, Boyle AJ. Prognostic Value of LVEDP in Acute Myocardial Infarction: a Systematic Review and Meta-Analysis. J Cardiovasc Transl Res. 2018;11(1):33-5. [PubMed ID: 29243014]. https://doi.org/10.1007/s12265-017-9776-7.
- 40. Şatıroğlu Ö, Çiçek Y, Bostan M, Çetin M, Bozkurt E. Acute Change in Left Ventricle End-diastolic Pressure after Primary Percutaneous Coronary Intervention in Patients with ST Segment Elevation Myocardial Infarction. Am Heart Hosp J. 2010;8(2):86. https://doi.org/10.15420/ahhj.2010.8.2.86.
- Elhabyan AK, Reyes BJ, Hallak O, Broce M, Rosencrance JG, Lucas BD, et al. Subendocardial ischemia without coronary artery disease: is elevated left ventricular end diastolic pressure the culprit? *Curr Med Res Opin*. 2004;20(5):773-7. [PubMed ID: 15140345]. https://doi.org/10.1185/030079904125003359.