

# **RESEARCH ARTICLE**

### ROLE OF 2D ECHO IN DIAGNOSING ACUTE PULMONARY EMBOLISM IN COPD PATIENTS

### Dr. Sarang Patil

#### Manuscript Info Abstract ..... ..... Manuscript History 2d Echo is a noninvasive bedside test which can be easily done in Received: 10 October 2019 COPD patients admitted with breathlessness and often misdiagnosed as Final Accepted: 12 November 2019 acute exacerbation of COPD. It avoids the contrast and radiation Published: December 2019 CTPA, chest CT or conventional angiography hazards of Echocardiography is an attractive imaging modality to diagnose PTE. Also, echocardiography allows visualization of the aorta and the LV to evaluate for other etiologies of chest pain, thus making it a more attractive test.

Copy Right, IJAR, 2019,. All rights reserved.

.....

### Introduction:-

- 1. In patients with pulmonary embolism (newly diagnosed) right ventricular hypertrophy often occurs which an be detected by echocardiography.
- 2. The 60/60 sign on 2d ECHO can help in diagnosis of pulmonary embolism
- 3. Also McConnell's sign can be used in as an important diagnostic tool in diagnosing pulmonary embolism in patients with or without pre-existing cardiorespiratory disease

#### Study design:-

Prospective observational cohort study

#### Sample size:-

100 patients

#### Inclusion criteria:-

- 1. Patients consenting for the study
- 2. Patients with high probability of pulmonary embolism as evidence by clinical examination

#### Exclusion criteria:-

- 1. Patients not consenting for the study
- 2. Patients with active pulmonary TB
- 3. Patients with recent myocardial infarction
- 4. Age less than 35 years

### Methodology:-

2D ECHO was done

- 1. 60/60 sign was looked for:
- 2. Pulmonary valve acceleration time ≤60ms

**Corresponding Author:- Dr. Sarang Patil** 

- 3. Tricuspid regurgitation pressure gradient ≤60mmHg
- 4. McConnell sign was looked for:
- 5. Normokinesia and hyperkinesia of the apical segment of the Right ventricle
- 6. Signs of Right ventricular hypertrophy

### **Results:-**

Pulmonary embolism was confirmed in 67 patients

- 1. 60/60 sign
- 2. Seen in 17/67 patients with pulmonary embolism
- 3. 2 false positives were observed
- 4. McConnell Sign
- 5. Seen in 13/67 patients with pulmonary embolism
- 6. RV pressure hypertrophy
- 7. Seen in 54/67 patients with pulmonary embolism
- 8. 18 were false positives



## **Conclusion:-**

2d Echo is thus helpful and an handy tool in correctly diagnosing pulmonary embolism.

## **Discussion:-**

Patients with COPD develop right ventricular hypertrophy and pulmonary hypertension and are at a risk of developing pulmonary embolism. 2D Echo is an important tool in diagnosing pulmonary hypertension and cardiac function. The most important variables in 2D Echo includes RA area, PASP, pericardial effusion, eccentricity index and TAPSE. PAH patients who have RA area  $\geq$ 27 cm2 are at higher risk of mortality or the need for heart transplant than those with a smaller RA area. The 2015 ESC/ERS guidelines recommend that RA area be used to stratify patients with PAH who are at risk of clinical worsening or death ; RA area <18 cm2 is suggested to confer low risk (<5%), 18–26 cm2 intermediate risk (5–10%), and >26 cm2 high risk (>10%). RA area indexed to BSA was shown to be an independent predictor of mortality in patients with PAH with WHO FC III-IV. Pericardial effusion was also an independent predictor of mortality. Serial 2D Echo monitoring can measure pericardial effusion. RV function is important because it predicts survival . TAPSE provides information about the status of RV systolic function based on the longitudinal function of RV myocardial fibers.TAPSE<1.8 cm was seen in patients who had more severe RV systolic dysfunction, right heart remodeling, and disproportionate RV size relative to the LV; these patients with TAPSE <1.8 cm, 1- and 2-year survival estimates were 94 and 88%, respectively, compared to 60 and 50%, respectively, when TAPSE <1.8 cm.

TAPSE thus can predict risk of severe disease and death in PAH.Additionally, both low pulmonary vascular capacitance and 2D strain of the basal RV free wall have been shown to be predictors of poor prognosis.

### **References:-**

- Badesch DB, Champion HC, Sanchez MA, Hoeper MM, Loyd JE, Manes A, et al. Diagnosis and assessment of pulmonary arterial hypertension. J Am Coll Cardiol. (2009) 54(1 Suppl.):S55–66. doi: 10.1016/j.jacc.2009.04.011
- 2. Runo JR, Loyd JE. Primary pulmonary hypertension. Lancet. (2003) 361:1533-44. doi: 10.1016/S0140-6736(03)13167-4
- Simonneau G, Gatzoulis MA, Adatia I, Celermajer D, Denton C, Ghofrani A, et al. Updated clinical classification of pulmonary hypertension. J Am Coll Cardiol. (2013) 62(25 Suppl):D34–41. doi: 10.1016/j.jacc.2013.10.029
- 4. Dellegrottaglie S, Ostenfeld E, Sanz J, Scatteia A, Perrone-Filardi P, Bossone E. Imaging the right heartpulmonary circulation unit: the role of MRI and computed tomography. Heart Failure Clin. (2014) 14:377–91. doi: 10.1016/j.hfc.2018.03.004
- 5. Raymond TE, Khabbaza JE, Yadav R, Tonelli AR. Significance of main pulmonary artery dilation on imaging studies. Ann Am Thorac Soc. (2014) 11:1623–32. doi: 10.1513/AnnalsATS.201406-253PP
- 6. Farber HW, Miller DP, Poms AD, Badesch DB, Frost AE, Muros-Le Rouzic E, et al. Five-Year outcomes of patients enrolled in the REVEAL Registry. Chest. (2015) 148:1043–54. doi: 10.1378/chest.15-0300
- Humbert M, Sitbon O, Chaouat A, Bertocchi M, Habib G, Gressin V, et al. Survival in patients with idiopathic, familial, and anorexigen-associated pulmonary arterial hypertension in the modern management era. Circulation. (2010) 122:156–63. doi: 10.1161/CIRCULATIONAHA.109.911818
- Benza RL, Miller DP, Gomberg-Maitland M, Frantz RP, Foreman AJ, Coffey CS, et al. Predicting survival in pulmonary arterial hypertension: insights from the Registry to Evaluate Early and Long-Term Pulmonary Arterial Hypertension Disease Management (REVEAL). Circulation. (2010) 122:164–72. doi: 10.1161/CIRCULATIONAHA.109.898122
- 9. Humbert M, Gerry Coghlan J, Khanna D. Early detection and management of pulmonary arterial hypertension. Eur Respir Rev. (2012) 21:306–12. doi: 10.1183/09059180.00005112
- Tonelli AR, Arelli V, Minai OA, Newman J, Bair N, Heresi GA, et al. Causes and circumstances of death in pulmonary arterial hypertension. Am J Respir Crit Care Med. (2013) 188:365–9. doi: 10.1164/rccm.201209-1640OC
- Schulze-Neick I, Lange PE, Haas NA. Intravenous epoprostenol for primary pulmonary hypertension. N Engl J Med. (1996) 334:1477; author reply -8
- 12. Savarese G, Paolillo S, Costanzo P, D'Amore C, Cecere M, Losco T, et al. Do changes of 6-minute walk distance predict clinical events in patients with pulmonary arterial hypertension? A meta-analysis of 22 randomized trials. J Am Coll Cardiol. (2012) 60:1192–201. doi: 10.1016/j.jacc.2012.01.083
- Voelkel NF, Quaife RA, Leinwand LA, Barst RJ, McGoon MD, Meldrum DR, et al. Right ventricular function and failure: report of a National Heart, Lung, and Blood Institute working group on cellular and molecular mechanisms of right heart failure. Circulation. (2006) 114:1883–91. doi: 10.1161/CIRCULATIONAHA.106.632208
- Matthews JC, McLaughlin V. Acute right ventricular failure in the setting of acute pulmonary embolism or chronic pulmonary hypertension: a detailed review of the pathophysiology, diagnosis, and management. CurrCardiol Rev. (2008) 4:49–59. doi: 10.2174/157340308783565384
- 15. Haddad F, Hunt SA, Rosenthal DN, Murphy DJ. Right ventricular function in cardiovascular disease, part I: Anatomy, physiology, aging, and functional assessment of the right ventricle. Circulation. (2008) 117:1436–48. doi: 10.1161/CIRCULATIONAHA.107.653576.
- 16. OSA in obese S.Patil et.al CHEST journal volume 155, issue 6, page A383.