A novel echocardiographic estimate of pulmonary vascular resistance employing the hydraulic analogy to Ohm’s law

Ashwin Venkateshvaran a,∗, Erik Tossavainen b, Charlie Borneteg c, Hande Oktay Tureli c, Davide Vanoli c,d, Lars H. Lund a, Frank Flachskampf d, Per Lindqvist c

a Department of Medicine, Cardiology Unit, Karolinska Institute, Stockholm, Sweden
b Department of Cardiology, Public Health & Clinical Medicine, Umeå University, Umed, Sweden
c Department of Clinical Physiology, Surgical & Perioperative Sciences, Umeå University, Umeå, Sweden
d Department of Medical Sciences, Uppsala University, and Clinical Physiology and Cardiology, Uppsala University Clinic, Uppsala, Sweden

ARTICLE INFO

Keywords:
Doppler echocardiography
Right heart catheterization
Pulmonary hypertension
Heart failure

ABSTRACT

Background: Assessment of pulmonary vascular resistance (PVR) is critical for accurate diagnosis and optimal pharmacotherapy in pulmonary hypertension. We aimed to test the diagnostic performance of a novel, Doppler-based method to evaluate PVR based on Ohm’s law (PVR echo) using pragmatic estimates of pulmonary capillary wedge pressure (PCWP).

Methods and results: Simultaneous right heart catheterization (RHC) and echocardiography was performed in a derivation cohort of 111 patients in sinus rhythm referred for PH evaluation and PVR echo independently validated in 238 patients. PVR echo was calculated using pulmonary artery mean pressure estimates (PAMP echo) obtained from peak tricuspid gradient employing a fixed right atrial pressure estimate, PCWP echo was estimated as 10 or 20 mmHg using age-related mitral E/A cut-offs and cardiac output from left ventricular outflow. In the derivation cohort, both PAMP echo and PCWP echo estimates demonstrated excellent agreement with catheterization measurements. PVR echo was highly feasible, demonstrated negligible bias and excellent agreement with PVR RHC (Bias = −0.58, SD 2.2 mmHg) and outperformed the Abbas method to identify PVR > 3WU (AUC = 0.85 vs. 0.70; p = 0.02). In the validation cohort, PVR echo preserved good invasive agreement with negligible bias, displayed strong diagnostic performance (AUC = 0.84) and significant ability to distinguish isolated post-capillary from combined post- and pre-capillary pulmonary hypertension (PH) subgroups (AUC = 0.77).

Conclusion: PVR echo based on Ohm’s law employing pragmatic estimates of PCWP echo demonstrates excellent agreement with invasive reference standard measurements and strong diagnostic ability to identify elevated PVR RHC. This novel approach may be useful during therapy selection to distinguish PH hemodynamic subgroups.

1. Background

Pulmonary hypertension (PH) is a chronic, progressive disease common in multiple clinical disorders and associated with poor long-term outcomes. Hemodynamic classification of patients with PH necessitates estimation of pulmonary vascular resistance (PVR), a static index of impedance that reflects pathological remodeling of the distal arterioles and alterations to the pulmonary vascular bed. Accurate quantification of PVR is important for a number of reasons. As a hemodynamic diagnostic indicator, PVR is integral to classifying PH subjects as having isolated post-capillary or combined post- and pre-capillary PH. [1] Further, PVR is an independent risk factor in the setting of heart failure (HF) and a strong predictor for reduced exercise capacity. [2] In multiple randomized clinical trials, reduction in PVR is associated with improvements of traditional risk stratification indices such as 6-minute walk test, WHO functional class and NT-proBNP. [3,4]

Reference-standard PVR is assessed using invasive right heart catheterization (RHC). Doppler-based approaches have been proposed [5–9], and present distinct advantages of being non-invasive, low-cost and highly accessible. However, their accuracy has been debated [10] and clinical utility may be limited by method complexity. [9] We have previously presented a novel, Doppler-based approach to assess PVR in a
pre-capillary PH cohort based on the hydraulic analogy to Ohm’s relationship. In that study, we employed a fixed, non-elevated PCWP estimate in all patients considering their pre-capillary PH status.[11] In the current study, we hypothesized that incorporation of a reliable, clinically relevant and simplified estimate of PCWP would allow wider application of this approach to the general PH population. We aimed to evaluate the accuracy of a Doppler-derived algorithm based on Ohm’s law to evaluate PVR using routinely-assessed echocardiographic variables in a general population of symptomatic patients referred for PH evaluation.

2. Methods

2.1. Study population

Consecutive patients with unexplained breathlessness referred for clinically-indicated RHC to Norrlands University Hospital between 2010 and 2015 were retrospectively analyzed. Patients with intracardiac or extracardiac shunts and severe valvular disorders were excluded prior to enrollment. Patients with atrial fibrillation or significant arrhythmia and no tricuspid regurgitation (TR) signals on echocardiography were excluded from the final cohort. Ethics committee approval was obtained prior to study enrollment (DNR 07–092M) and all patients provided written informed consent.

2.2. Right heart catheterization

RHC was performed by experienced operators blinded to echocardiographic data. Venous access was obtained by inserting an introducer in a medial cubital vein or in the femoral vein. A retrograde, right-heart catheterization was then performed using a Swan-Ganz pulmonary artery catheter (Edwards Lifesciences). Mean right atrial pressure (RAP), mean pulmonary capillary wedge pressure (PCWP) respectively, and mean pulmonary artery pressure (PAMP) were measured. Blood samples for estimation of oxygen saturation were drawn from the superior and inferior vena cava, as well as right atrium, and samples from the pulmonary and femoral arteries were used for screening for intra-cardiac shunts. Cardiac output (CO) was determined by thermodilution. Pulmonary vascular resistance was calculated using the equation PAMP/RHC – PCWP/RHC (trans-pulmonary gradient) divided by CO/RHC.

2.3. Echocardiography

Doppler Echocardiographic examination was performed by an experienced echocardiographer (PL) with > 15 years’ experience on-table, during RHC using a Vivid 7 system (GE Ultrasound, Horten, Norway) equipped with an adult 1.5–4.3 MHz phased array transducer. Standard views from the parasternal long and short axis and apical views were used in keeping with current recommendations.[12] Gray-scale images were obtained at 50 – 80 frames/sec and Doppler acquisitions at a sweep speed of 100 mm/sec. PASP using echocardiography (PASP/echo) was estimated using Continuous-Wave (CW) Doppler from the tricuspid regurgitation (TR) jet considering the most optimal of signals across multiple acoustic windows. Stroke volume (SV) was measured using Pulse-Wave (PW) Doppler at the level of the LV outflow tract, and CO/echo calculated by multiplying SV with heart rate. Mitral flow interrogation was performed in the 4-chamber view with the PW sample-volume placed between the mitral leaflets and measurements taken at end expiration. Early transmitral (E) and late diastolic (A) velocities were obtained after optimal sample alignment and E/A ratio was subsequently computed. Off-line analysis was performed using a commercially available software system (General Electric, EchoPAC PC version 11.0.0, GE Ultrasound, Waukesha, Wisconsin). Mean of three consecutive tracings were used to estimate a representative measurement.

Assessment of PVR using echocardiography (PVR/echo) was estimated using the hydraulic analogy to the Ohm’s relationship, i.e., PVR = (PAMP – PCWP)/CO employing echocardiographic surrogates for each of the variables employed in conventional equation, i.e transpulmonary gradient and ventricular output. PAMP/echo was calculated using the formula PASP/echo × 0.61 – 2 mmHg according to Chemla et al. [13] PASP/echo was estimated employing the peak trans-tricuspid regurgitate pressure drop adding a fixed right atrial pressure (RAP) of 7 mmHg.[14] Additional analysis was performed to estimate PVR/echo employing current recommendations considering inferior vena cava size and respiratory dynamics.[12] PCWP/echo was estimated based on combination of interpretation of Mitral E/A ratio and age. PCWP/echo was assigned a simplified estimate of 20 mmHg in younger patients (<50 years) if E/A
Table 1

Clinical Characteristics, right heart catheterization and echocardiographic data of patient population in the derivation cohort, grouped by PVR subgroups. Data presented as mean ± SD/ median (Q1; Q3) or number (%).

| Clinical Characteristics | All (n = 111) | PVR ≤ 3WU (n = 60; 54%) | PVR > 3WU (n = 51; 46%) | P-value |
|--------------------------|--------------|-------------------------|-------------------------|---------|
| Age (years)              | 61 ± 14      | 59 ± 15                 | 63 ± 13                 | 0.15    |
| Female                   | 75 (68)      | 42 (70)                 | 33 (65)                 | 0.44    |
| Diabetes                 | 13 (12)      | 10 (17)                 | 3 (6)                   | 0.68    |
| Hypertension             | 40 (36)      | 25 (42)                 | 15 (29)                 | 0.23    |
| Ischemic heart disease   | 14 (13)      | 7 (12)                  | 7 (14)                  | 0.13    |
| Heart rate (bpm)         | 74 ± 14      | 72 ± 15                 | 76 ± 13                 | <0.001  |
| Body surface area (m²)   | 1.86 ± 0.25  | 1.89 ± 0.27             | 1.83 ± 0.21             | 0.30    |
| Systolic blood pressure  | 132 ± 20     | 132 ± 19                | 133 ± 20                | 0.95    |
| (mmHg)                   |              |                         |                         |         |
| Diastolic blood pressure | 77 ± 7       | 75 ± 8                  | 79 ± 10                 | 0.01    |
| (mmHg)                   |              |                         |                         |         |
| NTproBNP (ng/L)          | 477 ± 341    | 341 (152;1375)          | 268 (1933)              | 0.20    |
| Right heart catheterization |          |                         |                         |         |
| RAP (mmHg)               | 7 ± 5        | 7 ± 4                   | 8 ± 6                   | 0.08    |
| PAMP (mmHg)              | 32 ± 15      | 24 ± 9                  | 43 ± 15                 | <0.001  |
| PCWP (mmHg)              | 12 ± 6       | 13 ± 7                  | 11 ± 5                  | 0.05    |
| TPG (mmHg)               | 20 ± 14      | 11 ± 5                  | 30 ± 14                 | <0.001  |
| PVR (WU)                 | 4.2 ± 3.4    | 2.0 ± 0.7               | 6.8 ± 3.3               | <0.001  |
| Cardiac output (L/min)   | 5.3 ± 1.6    | 5.7 ± 1.9               | 4.9 ± 1.9               | 0.01    |
| Echocardiography         |              |                         |                         |         |
| LV end-diastolic volume  | 87 ± 50      | 101 ± 56                | 72 ± 38                 | 0.003   |
| (ml)                     |              |                         |                         |         |
| LV end-systolic volume   | 43 ± 38      | 53 ± 46                 | 32 ± 22                 | 0.005   |
| (ml)                     |              |                         |                         |         |
| LVEF (%)                 | 54 ± 13      | 51 ± 13                 | 57 ± 12                 | 0.03    |
| RV basal diameter (mm)   | 41 ± 8       | 39 ± 9                  | 44 ± 6                  | 0.002   |
| RA area (cm²)            | 19 ± 7       | 18 ± 7                  | 21 ± 6                  | 0.04    |
| TAPSE (mm)               | 20 ± 5       | 21 ± 5                  | 18 ± 4                  | 0.01    |
| RV EL (%)                | 17 ± 7       | 19 ± 7                  | 15 ± 6                  | 0.005   |
| Mitral E wave (m/s)      | 73 ± 27      | 81 ± 22                 | 65 ± 29                 | 0.001   |
| Mitral E/A ratio         | 1.3 ± 0.8    | 1.5 ± 0.9               | 1.1 ± 0.6               | 0.01    |
| Mitral E/e' ratio        | 10 ± 5       | 10 ± 5                  | 10 ± 5                  | 0.41    |
| TR peak velocity (m/s)   | 3.4 ± 0.7    | 2.7 ± 0.3               | 3.7 ± 0.6               | <0.001  |
| RSPV (mmHg)              | 56 ± 22      | 43 ± 16                 | 67 ± 21                 | <0.001  |

3. Results

Of 145 patients referred for RHC in the derivation cohort, 32 patients with AF or significant arrhythmia and 2 with no TR signals were excluded. In effect, 111 (mean age 61 ± 14 years; 36 males) with sinus rhythm were included in the analysis. In the analyzed patient cohort, trivial or mild TR was seen in 88 patients (79%), moderate in 20 (18%) and severe to 3 (3%). No patients demonstrated free-flowing severe/torrential TR. On catherization, 35 (32%) did not demonstrate PH and 76 (68%) had PH in keeping with the revised hemodynamic definition of PAMP_{RHC} ≥ 20 mmHg at rest. [15] Fifty-one patients (46%) demonstrated elevated invasive PVR_{RHC} (≥ 3 WU). When PH patients were classified by etiology, 36 (32%) demonstrated pulmonary arterial hypertension, 40 (36%) had PH secondary to left heart disease, 9 (8%) had PH due to lung disease, 11 (10%) demonstrated chronic thromboembolic PH, and 15 (14%) demonstrated PH due to multifactorial mechanisms. When classified by hemodynamic status, 46 PH patients (61%) demonstrated pre-capillary PH (PCWP ≤ 15 mmHg) and 30 (39%) demonstrated post-capillary PH (PCWP > 15 mmHg). Among post-capillary PH patients, 18 (60%) demonstrated isolated post-capillary PH (PCWP > 15 mmHg and PVR < 3 WU) and 12 (40%) demonstrated combined post- and pre-capillary PH (PCWP > 15 mmHg and PVR > 3WU).

Baseline characteristics of the derivation cohort are presented in Table 1, stratified by PVR_{RHC} subgroups. Patients with elevated PVR_{RHC} demonstrated significantly smaller LV volumes and higher EF, larger right atrial (RA) and RV size, and lower RV longitudinal function seen both in lower TAPSE and RV free wall strain (p < 0.05 for all group comparisons).

3.1. Feasibility and diagnostic accuracy of PAMP\textsubscript{echo} to represent PAMP\textsubscript{RHC}

TR velocity could be adequately assessed in 96 (86%), and echocardiographic estimates of RAP from inferior vena cava size and collapse in 92 (83%) of patients in the derivation cohort. Applying ASE/EACVI recommended estimates of RAP (12), PAMP_{echo} using the Chemla’s equation demonstrated strong correlation (r = 0.82, r² = 0.67; p < 0.001 for both) and minimal bias (Bias = 0.66; SD 9.22 mmHg) with PAMP_{RHC}. Employing a simplified approach using a fixed, mean RAP_{echo} (7 mmHg), strong correlation (r = 0.80, r² = 0.64; p < 0.001 for both) (Fig. 2a) and excellent agreement with PAMP_{RHC} was preserved with a relatively higher spread of data points (Bias = 0.83; SD 9.56 mmHg) (Fig. 2b).

3.2. Diagnostic accuracy of age-dependent mitral E ratio to represent PCWP_{RHC}

Mitral E/A ratio was highly feasible (95%), demonstrated a strong positive correlation with PCWP_{RHC} (r = 0.65, p < 0.001) and outperformed other echocardiographic surrogates i.e., Mitral E (r = 0.45; p < 0.001), E/e’ (0.46; p < 0.001), TR velocity (r = 0.01; p = 0.90) and LA volume index (r = 0.38; p < 0.001). Further, mitral E/A demonstrated excellent ability to identify elevated PCWP_{RHC} (AUC = 0.84; CI 0.73 to 0.94; p < 0.001) and E/A cut-off > 2 demonstrated 50% sensitivity and 100% specificity to identify elevated PCWP_{RHC} in the total cohort. Eighty-four patients (76%) were ≥ 50 years and 27 (24%) were < 50 years old in the derivation cohort. In the older (≥50 years) sub-group,
mitral E/A \( > 1.4 \) demonstrated 69% sensitivity, 96% specificity, 90% PPV, 87% NPV and 88% accuracy to identify elevated PCWP \( \text{RHC} \) \((\text{AUC} = 0.84, \text{CI} 0.72 \text{ to } 0.96; p < 0.001)\) \((\text{Fig. 3a})\). Lower sensitivity (46%) and accuracy (82%) but excellent specificity (100%) was observed when E/A \( > 2 \) was considered as cut-off in this subgroup. In the younger group \((< 50 \text{ years})\) mitral E/A cut-off \( > 2 \) demonstrated 67% sensitivity, 100% specificity, 100% PPV, 91% NPV and 92% accuracy \((\text{AUC} = 0.87, \text{CI} 0.70 \text{ to } 1.0)\) \((\text{Fig. 3b})\).

Simplified estimation of PCWP \( \text{echo} \) as being non-elevated \( (10 \text{ mmHg}) \) or elevated \( (20 \text{ mmHg}) \) considering age in addition to mitral E/A as described in our methods demonstrated excellent diagnostic ability to identify \( \text{PCWP}_{\text{RHC}} \) \((\text{AUC} = 0.84; \text{CI} 0.70 \text{ to } 0.94; p < 0.001)\) in addition to good agreement with \( \text{PCWP}_{\text{RHC}} \) \((\text{Kappa coefficient} = 0.69)\). When compared with the current 2016 ASE/EACVI algorithm to determine elevated LV filling pressure, age-dependent mitral E/A demonstrated higher feasibility \((95 \text{ vs } 87\%)\), specificity \((97 \text{ vs } 93\%)\) PPV \((91 \text{ vs } 32\%)\) and modestly higher accuracy \((89 \text{ vs } 87\%)\) \((\text{Table 3})\). An illustration displaying age-dependent mitral E/A ratio and corresponding \( \text{PCWP}_{\text{RHC}} \) in addition to \( \text{PVR}_{\text{echo}} \) and corresponding \( \text{PVR}_{\text{RHC}} \) is provided in \text{Fig. 4}.

### 3.3. Diagnostic accuracy of \( \text{PVR}_{\text{echo}} \)

\( \text{PVR}_{\text{echo}} \) could be estimated in 88 of 111 patients \((79\%)\) employing \( \text{PAMP}_{\text{echo}} \) and \( \text{PCWP}_{\text{echo}} \) in the Ohm’s relationship. When compared with those in whom \( \text{PVR}_{\text{echo}} \) could not be assessed \((n = 23; 21\%)\), patients with quantifiable \( \text{PVR}_{\text{echo}} \) demonstrated higher PA pressures and \( \text{PVR} \) on \( \text{RHC} \), and lower \( \text{TAPSE} \) on echocardiography \((p < 0.05 \text{ for all comparisons})\).

\( \text{PVR}_{\text{echo}} \) demonstrated strong association \((r = 0.78, r^2 = 0.61; p < 0.001)\), negligible bias and excellent agreement with \( \text{PVR}_{\text{RHC}} \) on Bland-Altman analysis \((\text{Bias} = -0.58, \text{SD} 2.2 \text{ mmHg})\) \((\text{Fig. 5a} \& 5b)\). Further, this novel assessment of \( \text{PVR} \) outperformed conventional echocardiographic assessment using Abbas method \((5)\) to identify elevated invasive \( \text{PVR} > 3 \text{WU} \) \((\text{AUC} = 0.85, \text{CI} 0.76 \text{ to } 0.93 \text{ vs. AUC} = 0.70, \text{CI} 0.58 \text{ to } 0.81; p = 0.02 \text{ for comparison of AUC curves})\) \((\text{Fig. 5c})\).

### 3.4. External validation of \( \text{PVR}_{\text{echo}} \)

We then validated the novel \( \text{PVR}_{\text{echo}} \) in an independent database of 238 symptomatic patients with normal sinus rhythm referred for clinically-indicated \( \text{RHC} \) to the PH referral center at the Karolinska...
Table 2
Clinical Characteristics, right heart catheterization and echocardiographic data of patient population in the validation cohort. Data presented as mean ± SD/median (Q1; Q3) or number (%).

| Clinical Characteristics                                      | All Patients (n = 238) |
|---------------------------------------------------------------|------------------------|
| Age (years)                                                  | 58 ± 16                |
| Female                                                       | 120 (50)               |
| Diabetes                                                     | 27 (11)                |
| Hypertension                                                 | 103 (43)               |
| Ischaemic heart disease                                      | 22 (9)                 |
| Heart rate (bpm)                                             | 72 ± 13                |
| Body surface area (m²)                                       | 1.87 ± 0.24            |
| Systolic blood pressure (mmHg)                               | 121 ± 23               |
| Diastolic blood pressure (mmHg)                              | 68 ± 12                |
| NTproBNP (ng/L)                                              | 1395 (349:2765)        |

Right heart catheterization

| Parameter                        | Value |
|----------------------------------|-------|
| RAPmean (mmHg)                   | 7 ± 5 |
| PAPmean (mmHg)                   | 32 ± 13|
| PCWP (mmHg)                      | 14 ± 7 |
| TPG (mmHg)                       | 19 ± 13|
| PVR (WU)                         | 4.3 ± 3.5|
| Cardiac output (L/min)           | 5.3 ± 1.6|

Echocardiography

| Parameter                        | Value |
|----------------------------------|-------|
| LV end-diastolic volume (ml)     | 114 ± 58 |
| LV end-systolic volume (ml)      | 55 ± 53 |
| LVEF (%)                         | 55 ± 15 |
| RV basal diameter (mm)           | 41 ± 8 |
| RA area (cm²)                    | 20 ± 7 |
| TAPSE (mm)                       | 17 ± 6 |
| RV SL (%)                        | 17 ± 8 |
| Mitral E wave (m/s)              | 86 ± 32|
| Mitral E/A ratio                 | 1.6 ± 1.3|
| Mitral E/e prime                 | 10 ± 5 |
| TR peak velocity (m/s)           | 3.5 ± 0.8|
| RVSP (mmHg)                      | 56 ± 22|

NOTproBNP, N-terminal pro-B-type natriuretic peptide; RAP, right atrial pressure; PAP, pulmonary artery pressure; PCWP, pulmonary capillary wedge pressure; TPG, transpulmonary gradient; PVR, pulmonary vascular resistance; LV, left ventricle; EF, ejection fraction; RV, right ventricular; TAPSE, tricuspid annular plane systolic excursion; TR, tricuspid regurgitation.

University Hospital. Baseline characteristics of this cohort are presented in Table 2. This population demonstrated a higher proportion of patients with PH (n = 192; 81% vs. 68% in the Umeå cohort) and elevated PVR (61% vs. 48% respectively). Among those with PH, 121 (63%) demonstrated pre-capillary PH and 71 (37%), post-capillary PH. Among post-capillary PH patients, 121 (63%) demonstrated minimal bias and excellent agreement with RHC (Fig. 6a) and strong diagnostic ability to identify PVR (AUC = 0.84, CI 0.78 to 0.90; p < 0.001) (Fig. 6b). PVR echo > 3WU demonstrated 88% sensitivity, 54% specificity, 69% PPV and 79% NPV to identify PH (Kappa coefficient 0.43). A relatively higher agreement with RHC was obtained when PVR echo > 4.6WU was employed as cut-off (Kappa coefficient 0.55; 72% sensitivity, 83% specificity, 84% PPV and 71% NPV). Further, PVR echo was significantly higher among combined post- and pre-capillary PH patients when compared with isolated post-capillary PH (5.9 ± 3.7 vs 2.8 ± 2.7WU, p = 0.001) and demonstrated good diagnostic performance to discriminate these two groups (AUC = 0.77, CI 0.64 to 0.89; p = 0.001).

4. Discussion

We propose a novel echocardiographic approach to assess PVR employing variables routinely obtained in daily clinical practice using the hydraulic analogy to Ohm’s law. Simplified Doppler-based estimates of PCWP and PAMP employed in this equation demonstrated negligible bias and excellent agreement with corresponding invasive measurements. PVR echo obtained using this approach was highly feasible, demonstrated strong diagnostic performance and outperformed traditional echocardiographic algorithms to assess PVR RHC. When validated in an independent hemodynamic database of patients referred for PH evaluation, PVR echo preserved strong agreement with RHC measurements, showed excellent ability to identify elevated PVR RHC and strong diagnostic capability to differentiate isolated post-capillary from combined post- and pre-capillary PH. Our findings showcase PVR echo as a promising, non-invasive surrogate of reference-standard PVR that may be useful in diagnosis and regulating PH therapy.

4.1. Age-dependent mitral E/A ratio to represent PCWP

While the mitral E/A ratio is highly feasible and integral to the assessment of diastolic dysfunction, it demonstrates well-recognized limitations that prevent its use as an independent surrogate of elevated LV filling pressures as per current recommendations.[16] First, the E/A ratio showcases a U-shaped relation with LV diastolic function. In the specific setting of normal LV function, both subjects with normal and elevated PCWP RHC can demonstrate E/A ratio between 1 and 2. However, for values over 2, a sensitivity of 43% and specificity of 99% for identifying elevated PCWP RHC has been reported.[17] Further, both age and gender are known to significantly affect mitral doppler indices of diastolic dysfunction and age has been earlier shown to be the strongest independent predictor of mitral E/A.[18] An observed shift from a normal transmitral filling pattern to an ‘abnormal’ relaxation pattern is not unusual with aging, suggesting that absolute cut-offs may not be suited to the diagnosis of diastolic dysfunction. More complex algorithms have been recently proposed to evaluate PCWP RHC. Recently, a model combining TR velocity, E/e’, LV EF, RV fractional area change, IVC diameter and LA volume demonstrated a sensitivity of 92%, specificity of 93% and area under the curve of 0.97 to estimate elevated PCWP RHC.[9] However, such an algorithm necessitates acquisition of several measures incorporating considerable inter- and intra-observer variability in the approach. Our data suggests that considering age in addition to mitral E/A (which demonstrated strongest correlation with PCWP RHC) offers a simple, pragmatic measure with strong diagnostic performance.

4.2. Echocardiographic evaluation of PAMP

In this study, PAMP echo was assessed using the validated relationship proposed by Aduen et al.[19] and Chemla et al.[13] Assessment of pulmonary artery systolic pressure has traditionally been performed by adding an RAP estimate derived from IVC size and respiratory dynamics to the trans-tricuspid gradient.[12] Recent studies, however, suggest that these RAP estimates are frequently inaccurate and do not improve agreement with invasive reference.[14] Application of a fixed,
Fig. 4. Illustration of PCWP_{echo} assessment based on age and mitral E/A ratio and corresponding invasive PCWP and PVR_{echo}.

Fig. 5. (a) Scatter plot displaying association between PVR_{echo} and PVR_{RHC} (b) Bland-Altman analysis demonstrating excellent agreement between PVR_{echo} and PVR_{RHC} in the derivation cohort and (c) comparison of diagnostic performance employing PVR_{echo} by Ohm's relationship (AUC = 0.85) and Abbas algorithm (AUC = 0.70) in the derivation cohort (p = 0.02 for comparison).
represents a limited spread of invasive RAP (Median 6 mmHg, IQR 4 to 10 mmHg) measurements and no significant differences when patients with elevated and normal PVR were compared. In this context, a fixed RAP estimate simplifies assessment of PAMP using the Chemla approach [13], retains strong agreement with invasive measurements, and overcomes inherent technical limitations associated with IVC assessment [20].

One can argue that the assessment of PVR as employed in this study requires assessment of PAMP-PCWP and CO and each variable introduces a margin of error. However, we have chosen a pragmatic, simplified approach to assess highly reproducible variables routinely assessed in echocardiography labs worldwide. The variables chosen demonstrate higher feasibility and our approach demonstrates lower complexity when compared with more recently proposed models [9]. Advanced speckle-tracking has shown promise in estimation of RAP [21] and potentially improve estimation of PA pressures but this approach demonstrates relatively lower reproducibility and is rarely utilized in clinical practice.

4.3. Comparison with other Doppler-based PVR assessment

Our novel approach to assess PVR outperformed the conventional Doppler-based algorithm postulated by Abbas and colleagues. [5] The Abbas algorithm was originally tested in a pre-capillary PH population with preserved EF, and one can speculate that this approach may generate false-positives and showcase lower accuracy in a population that includes HF patients with post-capillary PH. However, comparison with other echocardiographic methods to estimate PVR [6–8] needs to be explored in further studies. Another strength of the current approach is its reasonable ability to distinguish isolated post-capillary PA from combined post- and pre-capillary PH from invasive pressures and identifies elevated PVR with high accuracy. This novel, pragmatic approach to non-invasive PVR assessment may be of value in patient screening, diagnosis and PH therapy regulation.

4.4. Clinical implications

Accurate, non-invasive estimation of PVR employing commonly available echocardiographic variables taking age into consideration may improve patient screening and triaging for invasive catheterization in addition to regulating therapy during follow-up. In addition, this approach may be useful to distinguish PH hemodynamic subgroups where PVR evaluation determines therapeutic management.

4.5. Limitations

Although micromanometer-tipped catheters offer high-fidelity pressure recordings and are considered the invasive standard, we employed standard fluid-filled catheters that are routinely utilized in clinical practice. Analysis of echocardiographic images in the validation and derivation sites were performed by two experienced operators employing standard international recommendations, thereby minimizing inter-evaluator variability.

4.6. Conclusions

PVR estimated employing the hydraulic analogy to Ohm’s Law is highly feasible, demonstrates excellent agreement with invasive measurements and identifies elevated PVR with high accuracy. This novel, pragmatic approach to non-invasive PVR assessment may be of value in patient screening, diagnosis and PH therapy regulation.

Declaration of Competing Interest

The authors declare that they have no known competing financial interests or personal relationships that could have appeared to influence the work reported in this paper.

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