Absence of diastolic mitral annular oscillations is a marker for relaxation-related diastolic dysfunction

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Riordan MM, Kovács SJ. Absence of diastolic mitral annular oscillations is a marker for relaxation-related diastolic dysfunction. Am J Physiol Heart Circ Physiol 292: H2952–H2958, 2007. First published February 16, 2007; doi:10.1152/ajpheart.01356.2006—Although Doppler tissue imaging frequently indicates the presence of mitral annular oscillations (MAO) following the E’ wave (E’ wave, etc.), only recently was it shown that annular “ringing” follows the rules of damped harmonic oscillatory motion. Oscillatory model-based analysis of E’ and E waves provides longitudinal left ventricular (LV) stiffness (k’), relaxation/viscoelasticity (c’), and stored elastic strain (εc’). We tested the hypothesis that presence (MAO+) vs. absence (MAO−) of diastolic MAO is an index of superior LV relaxation by analyzing simultaneous echocardiographic-hemodynamic data from 35 MAO+ and 20 MAO− normal ejection fraction (EF) subjects undergoing cardiac catheterization. Echocardiographic annular motion and transmitral flow data were analyzed with a previously validated kinematic model of filling. Invasive and noninvasive diastolic function (DF) indexes differentiated between MAO+ and MAO− groups. Specifically, the MAO+ group had a shorter time constant of isovolumic relaxation [τ; 51 (SD 13) vs. 67 (SD 27) ms; P < 0.01] and isovolumic relaxation time [63 (SD 16) vs. 82 (SD 17) ms; P < 0.001] and greater ratio of peak E-wave to peak A-wave velocity [1.19 (SD 0.31) vs. 0.97 (SD 0.31); P < 0.05]. The MAO+ group had greater peak lateral mitral annulus velocity [E’; 17.5 (SD 3.1) vs. 13.5 (SD 3.8) cm/s; P < 0.001] and LVEF [71.2 (SD 7.5) vs. 65.4 (SD 9.1%); P < 0.05] and lower heart rate [65 (SD 9) vs. 74 (SD 9) beats/min, P < 0.001]. Additional conventional and kinematic modeling-derived indexes were highly concordant with these findings. We conclude that absence of early diastolic MAO is an easily discernible marker for relaxation-related diastolic dysfunction. Quantitation of MAO via stiffness and relaxation/viscoelasticity parameters facilitates quantitative assessment of regional (i.e., longitudinal) DF and may improve diagnosis of diastolic dysfunction.

E’ wave; mitral annular motion; echocardiography; mathematical modeling; diastolic function; Doppler tissue imaging

HEART FAILURE with a normal left ventricular (LV) ejection fraction (EF), or “diastolic heart failure,” is a growing epidemic (34). Diastolic dysfunction (DD) itself has substantial adverse prognostic significance, demonstrating the importance of proper diagnosis of DD (35). While invasive diastolic function (DF) characterization typically involves the measurement of LV pressure and volume via cardiac catheterization, noninvasive DF characterization is usually achieved with echocardiography (2). Although new techniques for DF assessment, including strain, strain-rate, and color Doppler M mode, are evolving (30), evaluation of the transmitral flow pattern in concert with Doppler tissue imaging (DTI) remains a common method for DF assessment (2). Certain geometric features of the E and A waves, usually approximated as triangles, have been correlated with dysfunction [i.e., ratio of peak E-wave to peak A-wave velocity (E/A), deceleration time (DT), acceleration time (AT), velocity-time integral (VTI), etc.]. However, the known load dependence of global indexes, the realization that they may not be as sensitive to subtle mechanical dysfunction as regional indexes, and the evolution of imaging technology (strain, strain-rate, color Doppler M-mode, etc.) have led to increased interest in segmental or regional DF characterization. In particular, recent DTI studies have found that the long-axis (longitudinal) motion of the LV during early filling is much less load dependent than the E wave (13, 29, 30). This finding has motivated more routine measurement of the peak E’-wave velocity (E’) as an index of longitudinal DF. However, similar to E and A wave-based indexes, longitudinal DTI-based indexes currently rely only on geometric features (i.e., peaks) of the annular velocity contour, approximated as a triangle, and do not utilize the information content of the curvilinear velocity contour. Furthermore, E/E’, commonly reported as a noninvasive correlate of LV filling pressure, is computed from the nonsimultaneous peak values of the E and E’ waves (26).

MODELING LONGITUDINAL DIASTOLIC FUNCTION

To more completely characterize longitudinal DF in terms of stiffness and relaxation/viscoelasticity, we previously applied and validated damped simple harmonic oscillatory (SHO) motion as the kinematic principle that governs mitral annular oscillations (MAO) (38). Because LV filling is initiated by mechanical suction (defined as a simultaneous decrease in pressure and increase in volume, i.e., dP/dV < 0 immediately after mitral valve opening), a component of which must manifest as longitudinal elastic recoil, the proposed kinematic model for annular motion complements a previously developed and validated global lumped-parameter model that accurately predicts transmitral flow patterns (21, 24). This global model has validated relaxation during early filling as an important indicator of disease (9, 22, 37) and elucidated its relationship to isovolumic relaxation time (IVRT) and the time constant of isovolumic relaxation (τ) (8).

Accordingly, longitudinal tissue motion at the annulus (i.e., the E’ wave) can be characterized to excellent approximation.

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in analogy to the motion of a previously displaced (stretched) oscillator recoiling from rest by considering the balance of inertial, elastic, and viscous forces during early filling (38). This paradigm, the parameterized diastolic filling (PDF) formalism, has been successfully applied to E-wave analysis and can be similarly applied to annular motion. It provides longitudinal indexes of DF by using the clinical E′-wave contour as input and generating unique values for the initial spring displacement ($x_0$), damping constant ($c'$), and spring constant ($k'$) of an equivalent SHO. The physiological analogs of $x_0$, $c'$, and $k'$ are the VTI of the E′ wave, longitudinal myocardial relaxation/viscoelasticity, and longitudinal myocardial stiffness, respectively. Importantly, besides its observer-independent aspects, a key advantage of the PDF formalism is that it is “predictive” rather than “accommodative” in characterizing the E′-wave contour (25).

Interestingly, in addition to permitting solution to the “inverse problem” of longitudinal DF (38), the paradigm of a damped SHO for the E′ wave naturally predicts, and easily accounts for, the frequently observed reversal in the direction of motion (i.e., “ringing”) of the annulus. This oscillatory motion can be described by the mass-normalized solution to the equation of motion for a SHO:

$$v(t) = -\frac{k'x_0}{\omega^2} e^{-\omega t} \sin(\omega t)$$  (1)

where $\alpha' = c'/2$, $\omega' = [(4k' - c'^2)^{1/2})/2]$. For underdamped oscillation, where relaxation/viscoelastic effects are dominated by stiffness (i.e., $c'^2 < 4k'$), a MAO (described by the E′ wave) is predicted. Conversely, for overdamped or critically damped motion, in which relaxation/viscoelastic effects dominate stiffness (i.e., $c'^2 \geq 4k'$), the model predicts absence of mitral annular oscillations (MAO−). Therefore, the physics governing annular kinematics naturally dichotomizes subjects into those with and those without annular oscillations based on the relative values of $c'$ and $k'$. Furthermore, this kinematic analysis must be obeyed physiologically in the sense that a given LV either will or will not exhibit MAO.

The presence of MAO (i.e., the E′ wave) has been previously observed in humans (19, 42), but elucidation and characterization beyond noting mere presence have been lacking. Accordingly, we sought to test the hypothesis that if subjects were grouped according to the kinematically mandated presence (MAO+) or absence (MAO−) of annular oscillations, the dichotomization would elucidate DF differences between the groups. Specifically, we hypothesized that the absence of annular oscillations (MAO−) is the consequence of greater longitudinal relaxation/viscoelasticity effects relative to longitudinal stiffness and, therefore, implies worse DF than the presence of annular oscillations (MAO+).

**METHODS**

**Patient selection.** A sample of 55 normal LVEF (i.e., >50%) subjects with high-quality contemporaneous echocardiographic recordings of transmitial Doppler flow and DTI recordings of the lateral mitral annulus and high-fidelity (Millar) LV pressure were obtained from an existing database (8, 27). All subjects gave informed consent in accordance with a protocol approved by the Washington University Medical Center Human Studies Committee (Institutional Review Board) before data acquisition and catheterization. Before data acquisition and catheterization, all subjects gave informed consent according to Washington University Medical Center Human Studies (Institutional Review Board) guidelines. Subjects with abnormalities that could affect MAO, such as mitral stenosis, calcification, or regurgitation, were excluded. Subjects with heart rate (HR) sufficiently high to cause merging of the E′ and A′ waves (precluding identification of MAO) were also excluded. The subjects, ranging in age from 32 to 72 yr [54.5 (SD 9.9) yr], were dichotomized according to the presence (MAO+, n = 35) or absence (MAO−, n = 20) of MAO (i.e., the E′ wave). If a subject exhibited clear annular oscillations on at least three beats, he/she was entered into the MAO+ group. If a subject had no annular oscillations on any recorded beat, he/she was entered into the MAO− group. Subjects with one or two oscillatory beats were not included in the study. In general, annular oscillations were highly reproducible from beat to beat in the MAO+ group. The groups did not differ with respect to age, height, weight, race, or gender, and the number of subjects on diuretics, beta blockers, alpha blockers, angiotensin-converting enzyme inhibitors, and nitrates also did not differ between groups. Both groups included subjects with various nonvalvular comorbidities including, but not limited to, hypertension, diabetes, angiographically diagnosed coronary artery disease (CAD), previous myocardial infarction, wall motion abnormalities on ventriculography, cardiomyopathy, and a history of transient renal dysfunction. Hypertension, diabetes, and CAD were the most common comorbidities and, like the others, were comparably present in each group. Importantly, none of the subjects in either group had active ischemia at the time of data acquisition. A total of 20 subjects had elevated filling pressures defined as LV end-diastolic pressure (LVEDP) > 18 mmHg, and 28 subjects had impaired relaxation defined as $\tau > 50$ ms. Five MAO+ and seven MAO− subjects had elevated $\tau$ in addition to elevated LVEDP based on these criteria. Notably, diagnosis of DD does not require simultaneous presence of both elevated LVEDP and prolonged $\tau$. In contrast, subjects without clinical evidence of advanced DD may exhibit impaired stiffness or relaxation in isolation (12). However, we note that in general, subjects with elevated filling pressures tend to have impaired relaxation, although the inverse is often not true. Table 1 displays the relevant demographic and clinical information for each group.

Elective cardiac catheterization was performed on all subjects at the request of their referring physician on the basis of suspected CAD. Since the primary aim of this study was to determine the stiffness, relaxation/viscoelasticity, and stored elastic strain-based physiologic mechanisms that cause (or prevent) oscillations of the mitral annulus in any heart, we specifically chose not to compare healthy

**Table 1. Subject variables**

<table>
<thead>
<tr>
<th>Attribute</th>
<th>MAO+ Group (n = 35)</th>
<th>MAO− Group (n = 20)</th>
<th>Intergroup Significance</th>
</tr>
</thead>
<tbody>
<tr>
<td>Age, yr</td>
<td>53 (10)</td>
<td>56 (10)</td>
<td>$P = NS$</td>
</tr>
<tr>
<td>Males</td>
<td>25</td>
<td>12</td>
<td>$P = NS$</td>
</tr>
<tr>
<td>Minorities</td>
<td>5</td>
<td>6</td>
<td>$P = NS$</td>
</tr>
<tr>
<td>LVEDP &gt; 18 mmHg and $\tau &lt; 50$ ms</td>
<td>7</td>
<td>0</td>
<td>$P &lt; 0.05$</td>
</tr>
<tr>
<td>LVEDP ≤ 18 mmHg and $\tau &gt; 50$ ms</td>
<td>9</td>
<td>7</td>
<td>$P = NS$</td>
</tr>
<tr>
<td>Hypertensive</td>
<td>10</td>
<td>2</td>
<td>$P = NS$</td>
</tr>
<tr>
<td>Diabetic</td>
<td>6</td>
<td>1</td>
<td>$P = NS$</td>
</tr>
<tr>
<td>CAD</td>
<td>9</td>
<td>5</td>
<td>$P = NS$</td>
</tr>
<tr>
<td>Height, cm</td>
<td>175 (10)</td>
<td>173 (12)</td>
<td>$P = NS$</td>
</tr>
<tr>
<td>Weight, kg</td>
<td>93 (23)</td>
<td>92 (14)</td>
<td>$P = NS$</td>
</tr>
<tr>
<td>HR, beats/min</td>
<td>65 (9)</td>
<td>74 (9)</td>
<td>$P &lt; 0.001$</td>
</tr>
</tbody>
</table>

Clinical parameter values are means (SD), and demographic values are numbers of subjects. MAO+, subjects with mitral annular oscillations; MAO−, subjects without mitral annular oscillations; CAD, coronary artery disease; HR, heart rate; LVEDP, left ventricular end-diastolic pressure; $\tau$, time constant of isovolumic relaxation; NS, not significant.
controls vs. a single pathological group (hypertension, diabetes, CAD, heart failure, etc.). Hence, the MAO\textsuperscript{−} group had multiple clinical comorbidities, as expected.

**Data acquisition.** The methodology has been previously described (8, 27). Briefly, immediately before cardiac catheterization, a full two-dimensional/Doppler examination was performed in the catheterization laboratory with a standard clinical imaging system (Acuson, Mountain View, CA) including DTI of the lateral mitral annulus and transmitial Doppler inflow according to American Society of Echocardiography criteria (15). DTI for 11 of the subjects was performed with somewhat larger sample volumes (5, 9, and 10 mm). Previous DTI studies have used sample volumes ranging from 2 to 10 mm (14, 28–30, 39). All images were captured and stored on magneto-optical disks. A micromonomeric 6-Fr, dual-pressure transducer pigtail catheter (model SPC-560, Millar Instruments, Houston, TX) was advanced into the LV through a 6-F sheath (Arrow, Reading, PA) in the femoral artery. Pressure was recorded via a custom data acquisition system.

**Doppler analysis.** For each subject, five diastolic intervals containing clear E and A waves were selected and converted to 8-bit grayscale images with Paint Shop Pro 7 (Jasc Software, Minnetonka, MN). Conventional echocardiographic parameters AT, DT, peak E- and A-wave velocity, and E- and A-wave VTI (VTIE and VTIA, respectively) were measured manually with the triangle approximation for wave shape and averaged. IVRT was measured as the time between closure of the aortic valve and opening of the mitral valve (15). E-wave analysis using the PDF formalism was performed via model-based image processing according to previously validated methods to output the (mathematically) unique best-fit kinematic parameters (16, 17, 21). The parameters \( c/(g/s) \) and \( k/(g/s^2) \) denote the SHO (global) damping constant and spring constant (i.e., stiffness, \( dP/dV \); Ref. 27), respectively. The parameter \( x_0 \) (cm) corresponds to the stored elastic strain available at mitral valve opening that facilitates mechanical recoil (21, 23). All parameters are computed per unit mass (\( m \)). Additional SHO-derived indexes include \( kx_0 \), the peak force (equivalent to the peak atrioventricular pressure gradient) (4), \( 1/2kx_0^2 \), the stored elastic strain energy associated with LV recoil during filling, and \( \beta \), a viscoelastic stiffness parameter that quantifies the relative contributions of stiffness \( (k) \) and relaxation/isoviscosity \( (c) \) in determining the E-wave contour \( (\beta = c^2 - 4mk) \).

Three to five DTI cardiac cycles from each subject containing E’ and A’ waves were also selected, clipped, and converted to grayscale images. Peak E’- and A’-wave velocities were determined, as well as E’- and A’-wave VTI (VTIE\textsubscript{E’} and VTIA\textsubscript{A’}, respectively), by triangle approximation. For subjects with annular oscillations (i.e., an E’ wave), peak E’-wave velocity (E’) and E’-wave VTI (VTIE\textsubscript{E’}, respectively) were also determined. Lateral mitral annular excursion during early filling was determined by subtracting VTIE\textsubscript{E’} from VTIE\textsubscript{E} for subjects in the MAO\textsuperscript{−} group. E’-wave analysis was performed analogously to E-wave analysis via model-based image processing and nonlinear least-squares fitting of the model-predicted annular velocity \((Eq. 1)\) to the actual mitral annulus velocity contour as previously detailed (38). The longitudinal parameters \( c’/(g/s) \) and \( k’/(g/s^2) \) denote the longitudinal damping constant and spring constant of the system, respectively, determined from the E’ wave. Additional longitudinal indexes include \( k’x_0’ \), the peak longitudinal force driving recoil, \( 1/2k’x_0’^2 \), the peak stored elastic strain energy, and \( \beta’ \), where \( \beta’ = c’^2 - 4mk’ \) (38).

**Fig. 1** shows an example of the SHO model fit to an oscillatory and a nonoscillatory beat from representative subjects in the MAO\textsuperscript{−} and MAO\textsuperscript{−} groups, respectively.

For the MAO\textsuperscript{−} group, longitudinal \( c’ \) was determined by fitting a decaying exponential from the peak of the E’ wave to the peak of the E’ wave, as previously described (38). For the MAO\textsuperscript{−} group, longitudinal \( c’ \) was determined by assuming critical damping, which has slight limitations, as previously discussed (38).

**Statistical analysis.** All data are displayed as means (SD). Statistical differences between the subjects with and without MAO were determined by two-tailed analysis of variance. Statistical analysis of the nominal data for gender and minority representation, representative comorbidities, DD based on cutoff values for LVEDP and \( \tau \) (see Table 1), and pertinent medications in each group was performed with the two-tailed \( z \)-test for proportions. All statistical calculations were performed in Microsoft Excel 97 (Microsoft, Redmond, WA). Statistical significance was at the \( P < 0.05 \) level.

**RESULTS**

The values of all DTI measurements, parameters, and indexes analyzed for both groups are displayed in Table 3. Conventional Doppler and hemodynamic measurements, parameters, and indexes are shown in Table 2.

**Major findings.** Group comparison showed that the MAO\textsuperscript{−} group had a shorter \( \tau (P < 0.01) \) and IVRT \((P < 0.001)\) and greater E/A \((P < 0.05)\), indicating prolonged relaxation in subjects with MAO\textsuperscript{−}. The MAO\textsuperscript{+} group also had greater E’ \((P < 0.0001)\), slightly greater EF \((P < 0.05)\), and lower HR \((P < 0.001)\). The quantity \( \text{LVP}_{\text{diast}} - \text{LVP}_{\text{min}} \) was greater in the MAO\textsuperscript{−} group.

**Additional concordant findings.** Additional conventional and kinematic parameters/indexes of global and longitudinal LV function corroborate these findings. Specifically, among the...
conventional indexes, the MAO⁺ group had greater E′/A′ (P < 0.05) and VTIE (P < 0.0001), greater annular excursion (VTIE⁺ – VTIE⁻ < P < 0.05), and lower E/E′ (P < 0.05) and A (P < 0.05). In longitudinal PDF parameter/index terms (see Table 3), the MAO⁺ group had lower c (P < 0.0001) and k' (P < 0.05), greater x₀ (P < 0.01), k'x₀ (P < 0.05), and 3k'x₀² (P < 0.01), and more negative β' (P < 0.0001). In global PDF parameter/index terms (see Table 2), the MAO⁺ group had lower c (P < 0.001), lower kx₀ (P < 0.05), and more negative β (P < 0.001).

Table 2. Conventional and PDF parameters/indexes

<table>
<thead>
<tr>
<th>Parameter/Index</th>
<th>MAO⁺ Group (n = 35)</th>
<th>MAO⁻ Group (n = 20)</th>
<th>Intergroup Significance</th>
</tr>
</thead>
<tbody>
<tr>
<td>EF, %</td>
<td>71.2 (7.5)</td>
<td>65.4 (9.1)</td>
<td>P &lt; 0.05</td>
</tr>
<tr>
<td>E, cm/s</td>
<td>70.2 (15.0)</td>
<td>66.0 (15.5)</td>
<td>P = 0.34</td>
</tr>
<tr>
<td>AT, ms</td>
<td>87 (13)</td>
<td>92 (13)</td>
<td>P = 0.13</td>
</tr>
<tr>
<td>DT, ms</td>
<td>184 (40)</td>
<td>205 (53)</td>
<td>P = 0.11</td>
</tr>
<tr>
<td>VTIE₆, mm</td>
<td>95 (20)</td>
<td>100 (26)</td>
<td>P = 0.47</td>
</tr>
<tr>
<td>A, cm/s</td>
<td>61.5 (13.6)</td>
<td>74.1 (21.2)</td>
<td>P &lt; 0.05</td>
</tr>
<tr>
<td>VTIE₆, mm⁺</td>
<td>47 (10)</td>
<td>51 (17)</td>
<td>P = 0.37</td>
</tr>
<tr>
<td>E/A⁺</td>
<td>1.19 (0.31)</td>
<td>0.97 (0.31)</td>
<td>P &lt; 0.05</td>
</tr>
<tr>
<td>IVRT, ms</td>
<td>63 (16)</td>
<td>82 (17)</td>
<td>P &lt; 0.001</td>
</tr>
<tr>
<td>τ, ms⁺</td>
<td>51 (13)</td>
<td>67 (27)</td>
<td>P &lt; 0.01</td>
</tr>
<tr>
<td>LVESV, ml†</td>
<td>48 (24)</td>
<td>61 (29)</td>
<td>P = 0.08</td>
</tr>
<tr>
<td>LVEDV, mmHg</td>
<td>16.8 (4.5)</td>
<td>17.9 (5.4)</td>
<td>P = 0.44</td>
</tr>
<tr>
<td>LVP₀, mmHg†</td>
<td>6.3 (3.5)</td>
<td>7.0 (4.3)</td>
<td>P = 0.54</td>
</tr>
<tr>
<td>LVP₂, mmHg⁺</td>
<td>11.0 (4.5)</td>
<td>10.5 (4.1)</td>
<td>P = 0.69</td>
</tr>
<tr>
<td>LVP₀ – LVP₂, mmHg†</td>
<td>4.7 (1.6)</td>
<td>3.6 (1.2)</td>
<td>P &lt; 0.05</td>
</tr>
<tr>
<td>dP/dt₀-min, mmHg/µs</td>
<td>-1.731 (263)</td>
<td>-1.636 (414)</td>
<td>P = 0.30</td>
</tr>
<tr>
<td>c, cm/s²</td>
<td>18.4 (5.9)</td>
<td>27.3 (10.4)</td>
<td>P &lt; 0.001</td>
</tr>
<tr>
<td>k, cm/s²</td>
<td>204 (38)</td>
<td>225 (69)</td>
<td>P = 0.15</td>
</tr>
<tr>
<td>x₀, cm</td>
<td>10.5 (2.9)</td>
<td>11.7 (3.4)</td>
<td>P = 0.17</td>
</tr>
<tr>
<td>kx₀, dyn</td>
<td>2,100 (660)</td>
<td>2,590 (1,000)</td>
<td>P = 0.07</td>
</tr>
<tr>
<td>1/kx₀², erg</td>
<td>11,700 (7,100)</td>
<td>16,100 (10,300)</td>
<td>P = 0.001</td>
</tr>
<tr>
<td>β, x₀² –²</td>
<td>-445 (254)</td>
<td>-54 (542)</td>
<td>P &lt; 0.01</td>
</tr>
</tbody>
</table>

Values are means (SD). EF, ejection fraction; E and A, peak amplitude of E and A wave, respectively; AT and DT, acceleration and deceleration time of E wave, respectively; VTIE and VTIE₆, velocity-time integral of E and A wave, respectively; IVRT, isovolumic relaxation time; τ, time constant of isovolumic relaxation; LVESV, left ventricular (LV) end-systolic volume; LVEDV, LV end-diastolic pressure; LVP₀, minimum LV pressure; LVP₂, diastolic LV pressure; dP/dt₀-min, peak negative rate of pressure decline during isovolumic relaxation; c, damping constant; k, spring constant; x₀, initial displacement of spring before release; β, effect of stiffness vs. relaxation. *E and A waves were merged in 3 of the subjects without annular oscillations, precluding determination of A, E/A, and VTIE for these subjects. †Technical difficulties during data acquisition precluded determination of LVESV. LVP₀, LVP₂, and LVP₀ – LVP₂ in 1 and τ and dP/dt₀-min in 2 of the subjects without annular oscillations, respectively. Bold type indicates significance.

Age dependence of MAO. Since the MAO⁺ group had better DF than the MAO⁻ group based on a variety of invasive and noninvasive indexes (including τ, IVRT, E/A, and E′/A′) and DF indexes are known to depend on age (32, 33, 36), we also investigated the age dependence of MAO⁺. Figure 3 displays the percentage of subjects as a function of age expressed in decade intervals. The bar chart shows that the percentage of MAO⁺ subjects decreases monotonically with age, implying that presence or absence of MAO, in concordance with the aforementioned Doppler indexes, depends on age.

DISCUSSION

Cardiac anatomy and physiology. Zaky et al. (42) were the first to note that motion of the mitral annulus may have clinical importance. They noted that annular motion often reverses direction after the initial E’ wave. Much later, Isaz et al. (19) described the continuous, alternating atrially and apically directed motion of the mitral annulus during the cardiac cycle and hypothesized that the balance between stored elastic potential energy and kinetic energy plays a role. Recently, we complemented and extended these descriptive approaches, as well as others (19, 20, 40–42), by introducing and validating a quantitative, kinematics-based method of longitudinal DF assessment (38). This method views longitudinal annular motion during early filling as driven by strain energy (titin, extracellular matrix, etc.) stored during the previous systole; because of inertial effects and depending on the relative magnitudes of stiffness and viscoelasticity, the annulus “over-shoots” its equilibrium position and reverses direction (toward the apex) in certain subjects. The physics governing oscillatory motion predicts an oscillatory (stiffness > viscoelasticity) regime for annular motion as well as a nonoscillatory (stiffness < viscoelasticity) regime. Both regimes are observed

Table 3. Conventional and PDF parameters/indexes of longitudinal diastolic function

<table>
<thead>
<tr>
<th>Parameter/Index</th>
<th>MAO⁺ Group (n = 35)</th>
<th>MAO⁻ Group (n = 20)</th>
<th>Intergroup Significance</th>
</tr>
</thead>
<tbody>
<tr>
<td>E′, cm/s</td>
<td>17.5 (3.1)</td>
<td>13.5 (3.9)</td>
<td>P &lt; 0.0001</td>
</tr>
<tr>
<td>AT, ms</td>
<td>65 (12)</td>
<td>54 (11)</td>
<td>P &lt; 0.01</td>
</tr>
<tr>
<td>DT, ms</td>
<td>79 (17)</td>
<td>67 (20)</td>
<td>P &lt; 0.05</td>
</tr>
<tr>
<td>VTIE₆, mm</td>
<td>12.7 (3.5)</td>
<td>8.3 (3.7)</td>
<td>P &lt; 0.0001</td>
</tr>
<tr>
<td>VTIE₆ – VTIE⁰, mm⁺</td>
<td>10.4 (3.6)</td>
<td>8.3 (3.7)</td>
<td>P &lt; 0.05</td>
</tr>
<tr>
<td>A′, cm/s</td>
<td>16.3 (3.6)</td>
<td>15.5 (3.5)</td>
<td>P = 0.42</td>
</tr>
<tr>
<td>VTIE₀, mm</td>
<td>7.3 (2.3)</td>
<td>8.6 (3.0)</td>
<td>P = 0.06</td>
</tr>
<tr>
<td>E/A’</td>
<td>1.11 (0.28)</td>
<td>0.91 (0.35)</td>
<td>P &lt; 0.05</td>
</tr>
<tr>
<td>E/E’</td>
<td>4.16 (1.15)</td>
<td>5.18 (1.80)</td>
<td>P &lt; 0.05</td>
</tr>
<tr>
<td>c’, cm/s²</td>
<td>18.3 (5.9)</td>
<td>48.6 (7.7)</td>
<td>P &lt; 0.0001</td>
</tr>
<tr>
<td>k’, cm/s²</td>
<td>490 (120)</td>
<td>594 (233)</td>
<td>P &lt; 0.05</td>
</tr>
<tr>
<td>x₀, cm</td>
<td>1.16 (0.32)</td>
<td>0.85 (0.45)</td>
<td>P &lt; 0.01</td>
</tr>
<tr>
<td>k’x₀, dyn</td>
<td>547 (153)</td>
<td>460 (149)</td>
<td>P &lt; 0.05</td>
</tr>
<tr>
<td>1/k’x₀², erg</td>
<td>337 (164)</td>
<td>217 (147)</td>
<td>P &lt; 0.01</td>
</tr>
<tr>
<td>β, x₀² –²</td>
<td>-1,787 (489)</td>
<td>0 (0)</td>
<td>P &lt; 0.0001</td>
</tr>
</tbody>
</table>

Values are means (SD). E′ and A′, peak amplitude of E′ and A′ wave, respectively; AT and DT, acceleration and deceleration time of E′ wave, respectively; VTIE and VTIE⁰, velocity-time integral of E′ and E wave, respectively; VTIE₀, velocity-time integral of A′ wave; E′, peak amplitude of E wave; c′, longitudinal damping constant; k’, longitudinal spring constant; x₀, initial longitudinal displacement of spring before release; β, effect of longitudinal stiffness vs. relaxation. E’ waves were absent in subjects without annular oscillations. †β’ was computed for the MAO⁻ subjects by assuming critical damping as discussed in METHODS. Bold type indicates significance.
ROC Curves for Discriminating \( \tau > 50 \) ms

<table>
<thead>
<tr>
<th>Method</th>
<th>AUC</th>
</tr>
</thead>
<tbody>
<tr>
<td>MAO&lt;</td>
<td>0.74</td>
</tr>
<tr>
<td>IVRT &gt; 80 ms</td>
<td>0.64</td>
</tr>
<tr>
<td>DT &gt; 220 ms</td>
<td>0.71</td>
</tr>
<tr>
<td>MAO&lt; and DT &gt; 220 ms</td>
<td>0.79</td>
</tr>
</tbody>
</table>

Fig. 2. Receiver-operator characteristic curves for absence of MAO (MAO<), isovolumic relaxation time (IVRT) > 80 ms, deceleration time (DT) > 220 ms, and MAO< and DT > 220 ms (in combination) for predicting time constant of isovolumic relaxation (\( \tau \)) > 50 ms (relaxation-related diastolic dysfunction). Note that the area under the MAO< curve (AUC) is the greatest for all curves determined from a single marker, implying that absence of MAO is more predictive of \( \tau > 50 \) ms than either IVRT > 80 ms or DT > 220 ms. The associated sensitivity and specificity of MAO< for \( \tau > 50 \) ms are 62.2% and 74.2%, respectively. However, when the MAO< and DT > 220 ms markers are used in tandem, the AUC increases by 5%.

The elevated \( \tau \), IVRT, \( c \), and \( c' \) indicate that LV relaxation during both isovolumic relaxation and early filling is impaired in MAO< subjects relative to MAO+ subjects. Although most of the indexes determined in this study are load dependent, the fact that load, as assessed by LVEDP, did not differ between groups indicates that the observed concordance of conventional and kinematic parameters/indexes that differentiated between the groups did so based on differences in the underlying physiology rather than load. Additionally, we emphasize that absence of MAO is indicative of prolonged \( \tau \), but not of elevated LV filling pressures (i.e., LVEDP) per se. Therefore, absence of MAO is best viewed as a marker for DD related to impaired LV relaxation.

Because the presence or absence of MAO is determined by whether \( c'^2 \) is less than or greater than \( 4k' \) (i.e., the sign of the parameter \( b' \)), there is no absolute cutoff for \( c' \) or \( k' \) for determination of MAO+ or MAO<. However, we note that lack of an absolute cutoff value does not limit clinical utility of annular “ringing.” Given the reasonable sensitivity and specificity of MAO< for identifying abnormal relaxation-related DF, noting the mere presence or absence of MAO may prove useful as an additive indicator in characterizing relaxation-related DD, particularly since the presence of MAO depends highly on LV (global and longitudinal) relaxation. Indeed, as pointed out by Aurigemma et al. (3), diagnosis of DD and diastolic heart failure requires the examination of multiple Doppler-echo parameters/indexes in concert since standard noninvasive DF indexes do not correlate well with invasive pressure-derived indexes. We stress that there is no need for off-line analysis of annular motion contours to utilize the information conveyed by MAO; all that is required is determining the presence or absence of MAO by visual inspection of the DTI image.

**Limitations.** Sample volume positioning during DTI may affect the shape of the DTI contour and thus longitudinal PDF parameters and indexes. However, the generous sample size...
should minimize positioning-imposed effects. Neither the po-

sition nor size of the imaging sample length should affect the exis-
tence of MAO.

Only DTI of the lateral annulus was obtained, rather than
other aspects, because, in general, it is echocardiographically
easier to localize the sample volume at the lateral aspect and
thereby minimize the likelihood of including tissue (myoco-
dial) velocities. Several other studies have found that the
motion of the septal and other aspects of the mitral annulus
differs from the motion of the lateral annulus. Investigators
have reported that the peak velocity of septal E’ waves is
generally lower than that of lateral E’ waves and that the
total annular excursion during early filling is lower for the septal
aspect (1, 19). Based on prior observations, septal E’ and E”
are generally of lower amplitude than their lateral counterparts,
and E” is sometimes not observed even though oscillations of
the lateral annulus may be seen in the same subject. We note
that our modeling approach (the model, equations, methodology
of computing parameters, etc.) does not depend on which
aspect of the annulus is imaged.

While τ was significantly increased in the MAO− group
compared with the MAO+ group, LVEDP was similar be-
tween groups. This finding may be at least partially explained by the
fact that MAO occur during early filling (near E-wave termina-
tion), which is governed in part by the rate of LV relaxation,
whereas LVEDP occurs at the end of late atrial filling (end of
Doppler A-wave) and thus is influenced by atrioventricular
interactions. Moreover, MAO occur during the pressure rise
from LV min to LV dist, and the pressure difference from
minimum to diastatic pressure (LVp dist − LVp min) differen-
tiated between the MAO+ and MAO− groups better than
LVEDP. The greater LVp dist − LVp min in the MAO+ group
is evidence of enhanced suction-initiated filling in this group
compared with the MAO− group. This is further corroborated
by the nearly significantly lower LVESV in the MAO+ group
and the additional concordant findings enumerated above.

The slightly increased HR in the MAO− group may be
thought to contribute to the longitudinal and global parameters/
indexes that differentiated between groups. However, in-
creased HR is primarily accommodated by a decrease in the
duration of diastasis (7). E- and A-wave durations are nearly
HR independent, as each decreases by <20% for a 100% increase
in HR (6). DT has also been shown to decrease <20%
for a 100% increase in HR and E to increase <12% (7). In light
of the fact that DT was increased (P = 0.11) and peak E
was decreased (P = 0.34) in the MAO− group having the increased
HR, other factors must be contributing to the decreased E/A in
the MAO− group. It is also notable that IVRT and τ, which
would be expected to decrease with increasing HR, were
actually increased significantly in the MAO− group. The re-
results indicate that increased (longitudinal and global) relax-
ation/viscoelasticity rather than increased HR in the MAO−
group is primarily responsible for the trends observed in these
indexes.

Both groups included subjects with various nonvalvular
comorbidities, notably hypertension, diabetes, and CAD. Be-
cause hypertension and diabetes are detrimental to DF and
would be expected to correlate with the absence of MAO, their
predominance in the subjects with oscillations suggests that
exclusion of subjects with these comorbidities from the anal-
ysis would not alter the main findings. To verify this, we
investigated the extent to which the inclusion of subjects with
documented CAD and wall motion abnormalities on ventricu-
lography influenced our results. Additionally, since LVEDP >
18 mmHg is often characteristic of DD, we investigated the
influence of elevated LVEDP. Therefore, we divided both the
MAO+ and MAO− groups into three subgroups and conducted
subanalyses comparing the various global and longitudinal
parameters and indexes between a given MAO+ subgroup
and its corresponding MAO− subgroup. The subgroups
were 1) absence of wall motion abnormalities on ventriculog-
raphy (n = 45), 2) no history of CAD (i.e., <50% luminal
narrowing on angiography) (n = 41), and 3) LVEDP ≤ 18
mmHg (n = 35). While several supportive parameters/indexes
lost significance, it is noteworthy that IVRT, c’, k’, and c
remained significant in each subgroup, demonstrating that
longitudinal stiffness and longitudinal and global relaxation
during both early filling and isovolumic relaxation are impor-
tant correlates of the absence of annular oscillations. In partic-
ular, it is noteworthy that these kinematic stiffness and relax-
ation/viscoelasticity indexes remained significant in the sub-
group with LVEDP ≤ 18 mmHg. This finding implies that
these indexes are capable of detecting relaxation-related DD
before overt clinical manifestation based on invasively deter-
mined filling pressures. We note that the conventional Doppler
indexes E’ and A remained significant in each subgroup as
well. To determine whether a modified control group with
normal LVEDP and normal τ would have a higher prevalence
of MAO, we selected all subjects included in the study with
LVEDP ≤ 18 mmHg and τ < 50 ms without history of CAD
or wall motion abnormalities. Of the 13 subjects that met these
inclusion criteria, 10 (77%) had MAO.

Although MAO+ and MAO− likely depend on loading
conditions in the LV, the load dependence of MAO could not
be specifically determined in this study because of the similar
LVEDP across groups. Future work is planned to address this
issue.

Conclusions. The existence of oscillations of the mitral
annulus during early filling is predicted by a unifying physical
law that governs annular motion during early filling and whose
parameters convey quantitative information regarding longitudi-
(inal and global) diastolic function. Absence of oscillations
implies elevated viscoelasticity (i.e., impaired relaxation) effects
relative to stiffness effects and reduced longitudinal stored
strain energy to power early filling. The findings collectively
indicate that absence of mitral annular oscillations is a marker
of relaxation-related diastolic dysfunction. Importantly, ab-

cence of annular oscillations is not the cause, but rather is the
consequence, of impaired diastolic function. These findings
underscore the utility of considering the presence or absence
of annular oscillation in concert with more commonly used non-
invasive DF indexes, thereby providing insight into physiolog-
ical mechanisms of dysfunction and facilitating identification
of dysfunction in its preclinical state.

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REFERENCES


