

1 **Repeated echocardiographic imaging of aortic stenosis:**

2 **Real-life lessons**

3 Jonathan J. H. Bray MBChB, BSc^{1,2}, Adrian Ionescu MD, MRCP, FRCP (Edin.)¹

4 1. Morriston Cardiac Centre, Morriston Hospital, Heol Maes Eglwys, Cwmrhydyceirw,
5 Swansea, UK.

6 2. Institute of Life Sciences 2, Swansea Bay University Health Board and Swansea University
7 Medical School

8 Address for correspondence

9 Dr A Ionescu

10 Consultant Cardiologist

11 Morriston Cardiac Centre

12 Morriston SA6 6NL

13 adrian.ionescu@wales.nhs.uk

14

15 Abstract word count: 227

16 Word count: 2536

17 Tables: 4

18 Figures: 1

19 Supplementary tables: 5

20 Key words: 2D echocardiography, aortic stenosis, haemodynamic subsets, accuracy

21

22

23

24

25

26

27 **Abstract**

28 **Background**

29 Timing of aortic valve intervention is dependent on the accuracy and reproducibility of
30 echocardiographic (ECHO) parameters. We aimed to assess haemodynamic subsets of aortic
31 stenosis (AS), their change over time, and variability of ECHO parameters.

32 **Method**

33 This retrospective, longitudinal study compared sequential ECHO over 15 months to identify
34 concordant or discordant aortic valve area (AVA) and mean pressure gradient (MPG) in order
35 to determine the real world variability of echocardiographic indices.

36 **Results**

37 We included 143 patients with a mean age of 76.0 years. The median length of time between
38 studies was 112 days (IQR 38-208). Initially participants were classified as 9 (6.4%) mild, 47
39 (33.6%) moderate and 84 (60.0%) severe AS. In 80 (55.9%) AVA and MPG were
40 concordant; stroke volume index (SVi) was $<35\text{ml/m}^2$ in 53 (74.6%). AS severity was
41 downgraded in 29 (20.7%) patients. MPG was most consistent and AVA was the least
42 consistent between successive investigations (intraclass correlation coefficients $R=0.86$ and
43 $R=0.76$, respectively). Even small variations in left ventricular outflow tract (LVOT)
44 measurement of 1 standard deviation reclassified up to 67% of participants from severe to
45 non-severe.

46 **Conclusion**

47 Almost half of patients with AS have valve area/ gradient discordance. Variations in LVOT
48 diameter measurement commensurate with clinical practice reclassified AS severity in up to
49 2/3 of cases. Change in AS severity should only be accepted following careful scrutiny of all
50 available ECHO data.

51

52 **Introduction**

53 Aortic stenosis (AS) is the most common valvular heart disease in high-income
54 countries, and its prevalence is increasing as the population ages (1). Untreated, symptomatic
55 AS has worse survival than many cancers, but timely aortic valve intervention returns the
56 mortality curve to that normal for the population at large (2). Careful follow-up to allow
57 appropriate timing of valve intervention is essential, in order to avoid adverse outcomes
58 associated with advanced disease.

59 With the advent of percutaneous treatments for aortic valve disease, increasing
60 numbers of patients are considered for intervention (3, 4) with a commensurate increase in
61 the number of patients referred for echocardiographic (ECHO) surveillance of their AS.
62 Recommendations for the ECHO follow-up of patients with aortic stenosis (AS) differ in the
63 prescribed frequency of echocardiographic follow-up and are not always applied consistently
64 (5).

65 The reproducibility and accuracy of repeated ECHO measurements is rarely reported
66 or taken into account in ‘routine’ clinical practice; inter- or intra-observer variation may lead
67 to misdiagnoses such as spurious worsening of haemodynamic parameters when different
68 operators perform sequential scans. Accurate and reproducible ECHO measurements are
69 particularly important in the current era, when the proliferation of AS haemodynamic subsets
70 has markedly increased reliance on ECHO for clinical decision-making (6).

71 Our aims were: i) to ascertain the prevalence of haemodynamic subsets of AS in a
72 ‘real-world’ practice, ii) to interrogate their trends of AS parameters on sequential
73 transthoracic echocardiography (TTE), and iii) to model the clinical impact of LVOT
74 measurement variability on grading the severity of AS.

75

76

77 **Setting**

78 Morrison Cardiac Centre is a tertiary academic institution with a catchment
79 population of approximately 1,000,000 and performs approx. 18,000 TTEs/year.
80 Echocardiograms and the corresponding reports are stored in digital format using a
81 commercially available package (Change Healthcare, Nashville, TN, USA). All
82 echocardiographic assessments were made by British Society of Echocardiography (BSE)
83 accredited cardiac physiologists.

84

85 **Methods**

86 **Inclusion and exclusion criteria**

87 We retrospectively searched our digital ECHO database using as inclusion criteria:
88 ‘study performed between 01/01/2019-31/03/2020’ **and** ‘indication for study = assessment of
89 aortic stenosis (AS)’. We identified patients who had >1 study during this period, and
90 retained for further analysis only those who had 2 or 3 studies, after ascertaining that the
91 number of those with >3 exams was small. We excluded patients that had received frequent
92 scans over a concentrated interval because of suspected or confirmed infective endocarditis.
93 All patients had transthoracic echocardiographic (TTE) examinations according to the BSE
94 minimum dataset for TTE (7). We assessed the values of, and sequential changes in: mean
95 aortic valve (AV) gradient (MPG), peak AV velocity (PkV), AV area (AVA - by the
96 continuity equation), stroke volume index (SVi) and left ventricular outflow tract (LVOT)
97 diameter between successive examinations. We considered LVEF (cut-off 50%), SVi
98 (35ml/m²), AVA (1cm²), MPG (40mmHg) and PkV (4m/s) and, in accordance with the
99 literature, identified four haemodynamic subsets of AS, according to AVA (cut-off 1cm²) and
100 MPG (cut-off 40 mm Hg). We classified AS severity using AVA and MPG: Mild - AVA
101 >1.5 cm² and MPG <20 mmHg, moderate - AVA = 1-1.5 cm² or MPG ≥20-<40, and severe

102 when $<1\text{cm}^2$ or $\text{MPG} \geq 40$ (8). We posited that, as the LVOT diameter is relatively fixed in a
103 given patient even if the AS progresses, it can be used to test variability for repeated
104 measurements.

105 We explored the potential clinical significance of variations in the measured LVOT
106 diameter by calculating continuity AVA with the values at the extremes of the range of
107 diameters measured sequentially, and identified the proportion of patients who would have
108 been reclassified from severe to non-severe with the new LVOT values.

109

110 **Statistics**

111 Analyses were performed using IBM SPSS v. 25 (Chicago, IL, USA). Continuous
112 numerical variables are presented as mean (\pm standard deviation ($\pm\text{SD}$)) and range. Non-
113 normally distributed data are presented as median (\pm interquartile range ($\pm\text{IQR}$)). First and
114 second ECHO studies were compared using paired samples 2-tailed t-tests, and first, second
115 and third studies were compared using repeated measures ANOVA. Where parametric
116 assumptions were not met, we used Wilcoxon Signed Rank test (1st vs 2nd) and Friedman's
117 two-way ANOVA (1st vs 2nd vs 3rd). The threshold for significance was set at $p < 0.05$. We
118 calculated the coefficient of variation and of repeatability of the echocardiographic
119 measurements. The coefficient of repeatability (CR) was calculated as within-subject
120 standard deviation (SW) $\times 2.77$ ($\sqrt{2 \times 1.96}$) (9). Dedicated software v. 2019b (Origin Lab,
121 Northampton, MA, USA) was used to produce graphical representations of data. We assessed
122 the variability of repeated measurements using Bland-Altman analysis and linear regression.

123

124 **Results**

125 We included 143 patients: 126 had two, and 17 had three ECHO studies. There were
126 68 females and 75 males, with a mean age of 76.0 years (± 10.0). The median duration

127 between study #1 and study #2 was 112 days (IQR 38-208, range 0-320). Table 1 shows the
128 baseline characteristics and the absence of significant difference in mean ECHO parameters
129 between first and second echo. At inclusion, there were 9 (6.4%) cases of mild, 47 (33.6%) of
130 moderate and 84 (60.0%) severe AS.

131

132 **Haemodynamic subsets**

133 There were two ‘concordant’ subsets (severe AS, with AVA <1 cm² and
134 MPG ≥40mm Hg, and non-severe AS, with AVA ≥1 cm² and MPG ≤40mm Hg) and two
135 ‘discordant’ subsets (one with AVA ≥1 cm² but with MPG >40mm Hg and a ‘low-gradient
136 AS’ subset with AVA <1cm² and MPG ≤40 mm Hg). There were only four patients in the
137 subset with large area/ high gradient. When comparing both subsets, SVi was significantly
138 greater in ‘concordant’ as opposed to ‘discordant’ subsets (40.7 ml/m² [±11.4] versus 33.8
139 [±10.9] p<0.0001, respectively) (Table 2). All other parameters were not significantly
140 different, including LVEF (55.2% [±14.2] versus 55.4% [±12.8], respectively).
141 Supplementary Table 1 shows the ECHO parameters of each subset and supplementary
142 Table 2 displays these, as well as further haemodynamic subsets defined by each of the
143 metrics used. LVEF binary class did not affect haemodynamic parameters, but MPG and PkV
144 were different by both SVi and AVA cut-offs. Supplementary table 3 shows the proportion of
145 patients within each haemodynamic subset.

146

147 **Consistency between repeated measurements**

148 There was no significant difference between the average measurements in those who
149 had either 2 or 3 scans during the study period, and no overall trend was apparent across time.
150 In 68 patients (47.5%) AVA decreased between the first and second ECHO, whereas in 75
151 (52.4%) patients AVA either stayed the same or increased. The group with decreasing AVA

152 compared with the group in which AVA stayed the same or increased, was associated with a
153 significant reduction in SVi between ECHO 1 and 2 (Δ -8.45 ml/m² versus Δ 6.83 ml/m²,
154 respectively, $p < 0.0001$). MPG and PkV were not significantly different. Only 16.8% (24
155 patients) of AVA measurements by continuity changed by $\leq 5\%$.

156 Correlations between repeated measurements were highly significant but only
157 moderate (Table 3). The best correlation between two successive measurements was for the
158 mean transvalvular gradient ($R = 0.86$) and the worst for AVA ($R = 0.76$) with LVOT
159 diameter in an intermediate position ($R = 0.79$).

160 Coefficients of variation and of repeatability for the echocardiographic parameters are
161 given in supplementary Table 4. We assessed agreement between successive
162 echocardiographic measures with Bland-Altman plots, obtaining absolute bias, 95% limits of
163 agreement (LoA) and the proportion of measurement differences falling outside LoA. Bland-
164 Altman plots for MG, AVA, LVOTd, LVOT VTI, PkV were produced (not shown).
165 Absolute bias was small at: -0.54 (± 8.6), 0.049 (± 0.34), -0.004 (± 0.20), -0.38 (± 5.9), -4.7
166 (± 55.6), 0.0003 (± 0.02), respectively. Percentages of measurement differences outside of the
167 95% LoA were consistent with each other at: 7.1%, 6.2%, 5.4%, 5.3%, 7.4% and 8.2%,
168 respectively. There was a tendency for greater disagreement with larger measurements
169 particularly for MG and PkV. Despite this, there was an even spread of observations around
170 the mean difference for all measures indicating no evidence of proportional bias.

171

172 **Potential clinical impact of variation in LVOT diameter measurement**

173 Repeated scanning led to a reclassification of the severity of the AS compared to the
174 first scan in 43 patients (30.7%), downgraded in 29 (20.7%) and upgraded in 19 (13.6%)
175 (Figure 1). Change in classification did not appear to be associated with age, sex or specific
176 ECHO parameters (Supplementary Table 4). To assess the contribution of variation in AVA

177 to reclassification, when AVA is kept constant as the value found in the first ECHO and the
178 MPG from ECHO 2 is used, only 3.6% (5/140) of cases are reclassified (2 mild to moderate,
179 1 moderate to severe, and 2 severe to moderate).

180 To model the impact of LVOT diameter, we added or subtracted the SD of the
181 diameter, or the highest difference between sequential diameter measurements, to the
182 diameter reported for each study. We calculated the proportion of patients whose AS severity
183 would have been reclassified as $AVA \geq 1\text{cm}^2$ or indexed $AVA \geq 0.6\text{cm}^2$ if LVOT diameter
184 would have been measured as either bigger or smaller than the actual value (Table 4). The
185 proportion of patients reclassified to non-severe, unindexed $AVA \geq 1\text{cm}^2$, and indexed AVA
186 $\geq 0.6\text{cm}^2$ ranged between 65% to 24%, 67% to 20%, and 54% to 10%, respectively. ±

187

188 **Discussion**

189 We found that in almost 50% of patients with aortic stenosis who had 2 TTE studies
190 in a tertiary centre over the course of 15 months there was a decrease in the AVA of 0.22cm^2
191 (± 0.16) ($p < 0.0001$) between studies. Repeated scanning reclassified the AS severity (defined
192 by AVA and MPG) of 43 patients (30.7%). In 80 patients (55.9%) initial AVA and MPG
193 were concordant, while the discordant scans were in patients with low-gradient AS, and there
194 was a moderate-to-good correlation of repeated measurement of parameters used for
195 calculating aortic valve area, including diameter of the LVOT. We demonstrated that chance
196 variations in the measurement of the LVOT diameter, well within the range encountered in
197 clinical practice, have a major impact on the classification of AS severity, with its corollary
198 of potentially inappropriate or delayed surgical referral.

199 Further haemodynamic subset stratification demonstrated that LVEF, with a cut-off of
200 50%, did not result in significant differences between the metrics monitored, while SVi
201 (35ml/m^2) and AVA (1cm^2) dichotomised observed MPG and PkV.

202

203 **Clinical impact of variation in measurement**

204 The assessment of the reproducibility, reliability and accuracy of echocardiographic
205 measurements is an important component of the quality improvement of ECHO services (10).
206 Repeated measurement of the LVOT diameter provided us with an opportunity to assess the
207 reproducibility (11) of this linear measurement in our clinical practice. We found little
208 variation of LVOT measurement in our lab, and demonstrated that variation within the limits
209 of the standard deviation of the LVOT diameter measurement had a dramatic impact on the
210 classification of AS severity. We focused on the LVOT diameter because it is a major
211 contributor to discrepancies in the assessment of AS severity (12) and because (unlike
212 gradients and areas, which may change during follow-up as the disease progresses) LVOT
213 dimensions are generally static over time. To our best knowledge there is nothing published
214 previously on the reproducibility of ECHO measurements in NHS, clinical non-research
215 settings, and our data represent a step in this direction.

216 The downgrading of AS severity by continuity AVA in over 1/5 of patients at a
217 second ECHO study was unexpected. The vast majority of this change was due to variation
218 between first and second echo in AVA (96.4%). With the natural progression of AS we
219 would expect either no change or worsening of AVA. We did not have global longitudinal
220 strain data from enough patients to explore the possibility that this phenomenon reflected a
221 subclinical change in LV systolic function.

222

223 **Haemodynamic subsets of AS**

224 Since Hachicha et al. introduced the new entity of paradoxical low-flow, low gradient
225 AS (13), the whole field has gained complexity, with the continued proliferation of multiple

226 new indices and haemodynamic patient subsets (14) deemed to have prognostic relevance,
227 although this approach has been questioned (15).

228 A common clinical problem is AS with low area but also with low gradient (16). If
229 LVEF is depressed, the distinction between truly severe and pseudo-severe AS can often be
230 made by low-dose Dobutamine stress echo (17). In the presence of a normal LVEF there may
231 be paradoxical low-flow, low-gradient AS (15). Classifications depend on accurate
232 echocardiographic measurements of multiple haemodynamic parameters, require elimination
233 of alternative causes for symptoms, do not have universally accepted treatment implications,
234 and even in the best laboratories, discrepancies in AS grading may occur, with puzzling
235 clinical implications (18). The prevalence of each haemodynamic subset outside core echo
236 labs is poorly characterised.

237 We describe the 'real world' prevalence of haemodynamic subsets of AS, defined by
238 area, gradient, peak velocity and stroke volume index and show that LVEF (cut-off 50%) is
239 not associated with different values of the haemodynamic metrics, whereas SV_i and AVA
240 dichotomise MPG and PkV.

241

242 **Limitations**

243 This work is retrospective and observational, but as such represents real-life practice.
244 Information on further tests to confirm the severity of AS of participants was not available.
245 Therefore, it was not possible to determine which echocardiographic assessment and thus
246 classification of AS was most accurate. It would have been useful to assess the variation of
247 the dimensionless Otto index in our database, due to its superior reproducibility,(19, 20) but
248 our software did not facilitate collection of this data. Although we did not have access to the
249 clinical files to understand exactly the indication for the echo studies, the fact that in the
250 majority of patients the severity of the AS appeared to have progressed suggests that the

251 indication for echocardiographic surveillance of AS was correct and in keeping with the
252 guidelines (21).

253

254 **Conclusions**

255 In a 'real world' setting, almost half of patients with AS have valve area/gradient
256 discordance. AVA decreased in 45%, together with SVi. Small variations in LVOT diameter
257 measurement reclassified AS severity in up to 2/3 of cases. In over a fifth of cases AS
258 severity was downgraded by a follow-up scan. Clinical decisions should never be based
259 solely on reported echocardiographic AS progression.

260

261 **Declaration of Interest**

262 None.

263 **Funding**

264 None.

265 **CRedit Author Statement Jonathan J. H. Bray:** Methodology, Investigation, Formal
266 analysis, Visualization, Writing – Review & Editing, Project administration. **Adrian Ionescu:**
267 Conceptualisation, Methodology, Investigation, Formal analysis, Writing – Original Draft,
268 Visualization, Writing – Review & Editing.

269 **Acknowledgements**

270 None.

271 **Data availability statement**

272 The data that support the findings of this study are available from the corresponding author upon
273 reasonable request.

274 **References**

- 275 1. D'Arcy JL, Coffey S, Loudon MA, Kennedy A, Pearson-Stuttard J, Wilson J, et al. Large-
276 scale community echocardiographic screening reveals a major burden of undiagnosed valvular heart
277 disease in older people: The OxVALVE Population Cohort Study. *European Heart Journal*.
278 2016;37(47).
- 279 2. Hannan EL, Samadashvili Z, Lahey SJ, Smith CR, Culliford AT, Higgins RSD, et al. Aortic
280 Valve Replacement for Patients With Severe Aortic Stenosis: Risk Factors and Their Impact on 30-
281 Month Mortality. *The Annals of Thoracic Surgery*. 2009;87(6):1741-9.
- 282 3. Mack MJ, Leon MB, Thourani VH, Makkar R, Kodali SK, Russo M, et al. Transcatheter
283 Aortic-Valve Replacement with a Balloon-Expandable Valve in Low-Risk Patients. *New England*
284 *Journal of Medicine*. 2019;380(18):1695-705.
- 285 4. Popma JJ, Deeb GM, Yakubov SJ, Mumtaz M, Gada H, O'Hair D, et al. Transcatheter
286 Aortic-Valve Replacement with a Self-Expanding Valve in Low-Risk Patients. *New England Journal*
287 *of Medicine*. 2019;380(18):1706-15.
- 288 5. Zaidi A, Ionescu A, Sharma R, Heatley M. Echocardiographic surveillance of aortic valve
289 stenosis: towards a standardized approach. *J Heart Valve Dis*. 2012;21(6):707-13.
- 290 6. Clavel M-A, Magne J, Pibarot P. Low-gradient aortic stenosis. *European Heart Journal*.
291 2016;37(34):2645-57.
- 292 7. Wharton G, Steeds R, Allen J, Phillips H, Jones R, Kanagala P, et al. A minimum dataset for
293 a standard adult transthoracic echocardiogram: a guideline protocol from the British Society of
294 Echocardiography. *Echo Research and Practice*. 2015;2(1):G9-G24.
- 295 8. Messika-Zeitoun D & Lloyd G 2018. Aortic valve stenosis: evaluation and management of
296 patients with discordant grading. *European Society of Cardiology e-Journal of Cardiology Practice*.
297 **26**; vol. 15. (Found at: <https://www.escardio.org/Journals/E-Journal-of-Cardiology-Practice/Volume-15/Aortic-valve-stenosis-evaluation-and-management-of-patients-with-discordant-grading>.)
- 298
- 299 9. Vaz S, Falkmer T, Passmore AE, Parsons R, Andreou P. The Case for Using the Repeatability
300 Coefficient When Calculating Test–Retest Reliability. *PLOS ONE*. 2013;8(9):e73990.
- 301 10. Masani N. The Echocardiography Quality Framework: a comprehensive, patient-centered
302 approach to quality assurance and continuous service improvement. *Echo Research and Practice*.
303 2018;5(4):G35-G41.
- 304 11. Bunting KV, Steeds RP, Slater LT, Rogers JK, Gkoutos GV, Kotecha D. A Practical Guide to
305 Assess the Reproducibility of Echocardiographic Measurements. *Journal of the American Society of*
306 *Echocardiography*. 2019;32(12):1505-15.
- 307 12. Michelena HI, Margaryan E, Miller FA, Eleid M, Maalouf J, Suri R, et al. Inconsistent
308 echocardiographic grading of aortic stenosis: is the left ventricular outflow tract important? *Heart*.
309 2013;99(13):921.

- 310 13. Hachicha Z, Dumesnil JG, Bogaty P, Pibarot P. Paradoxical Low-Flow, Low-Gradient Severe
311 Aortic Stenosis Despite Preserved Ejection Fraction Is Associated With Higher Afterload and
312 Reduced Survival. *Circulation*. 2007;115(22):2856-64.
- 313 14. Eleid MF, Sorajja P, Michelena HI, Malouf JF, Scott CG, Pellikka PA. Flow-Gradient
314 Patterns in Severe Aortic Stenosis With Preserved Ejection Fraction. *Circulation*. 2013;128(16):1781-
315 9.
- 316 15. Bleakley C, Monaghan MJ. Assessment of Normal-Flow Aortic Stenosis. *Circulation:
317 Cardiovascular Imaging*. 2017;10(12):e007293.
- 318 16. Pibarot P, Dumesnil JG. Aortic Stenosis Suspected to Be Severe Despite Low Gradients.
319 *Circulation: Cardiovascular Imaging*. 2014;7(3):545-51.
- 320 17. deFilippi CR, Willett DL, Brickner ME, Appleton CP, Yancy CW, Eichhorn EJ, et al.
321 Usefulness of dobutamine echocardiography in distinguishing severe from nonsevere valvular aortic
322 stenosis in patients with depressed left ventricular function and low transvalvular gradients. *The
323 American Journal of Cardiology*. 1995;75(2):191-4.
- 324 18. Minners J, Allgeier M, Gohlke-Baerwolf C, Kienzle R-P, Neumann F-J, Jander N.
325 Inconsistencies of echocardiographic criteria for the grading of aortic valve stenosis. *European Heart
326 Journal*. 2008;29(8):1043-8.
- 327 19. Sacchi S, Dhutia NM, Shun-Shin MJ, Zolgharni M, Sutaria N, Francis DP, et al. Doppler
328 assessment of aortic stenosis: a 25-operator study demonstrating why reading the peak velocity is
329 superior to velocity time integral. *European Heart Journal - Cardiovascular Imaging*.
330 2018;19(12):1380-9.
- 331 20. Geibel A, Görnandt L, Kasper W, Bubenheimer P. Reproducibility of Doppler
332 echocardiographic quantification of aortic and mitral valve stenoses: Comparison between two
333 echocardiography centers. *American Journal of Cardiology*. 1991;67(11):1013-21.
- 334 21. Nishimura RA, Otto CM, Bonow RO, Carabello BA, Erwin JP, Guyton RA, et al. 2014
335 AHA/ACC Guideline for the Management of Patients With Valvular Heart Disease: Executive
336 Summary. *Circulation*. 2014;129(23):2440-92.

337

338

339

340

341

342

343

344

345 **Figure and table legends**

346 **Table 1.** Mean (standard deviation) and [range] (*median (interquartile range) and [range])
347 for echocardiographic parameters of aortic stenosis at successive scans. Difference between
348 values of echocardiographic parameters at 2 successive time points expressed as a mean. All
349 comparisons compared with echo 1 are statistically non-significant. Coefficient of
350 Repeatability (within-subjects SEM). Other rhythm includes ventricular paced rhythm,
351 junctional rhythm and non-reported rhythms.

352

353 **Table 2.** Comparison and associations of discordant (echo 1: AVA ≥ 1 cm², MPG >40 mm
354 Hg and AVA <1 cm², MPG \leq 40 mm Hg) versus concordant and reclassified (changed AS
355 classification of mild, moderate or severe from echo 1 to echo 2) versus non-reclassified
356 severity subgroups. Abbreviations: LVOTd – Left ventricular outflow tract diameter, LVEF –
357 Left ventricular ejection fraction, SVi – Stroke volume index, MPG – Mean pressure gradient
358 and PkV – peak velocity.

359

360 **Table 3.** Correlation coefficients of repeated measurements in patients who had 2 ECHO
361 studies. Abbreviations: LVOT – left ventricular outflow tract; AV – aortic valve.

362

363 **Figure 1.** Reclassification of Aortic Stenosis (AS) severity between ECHO 1 and ECHO 2.

364

365

366

367 **Table 4.** Impact of variation in the measurement of the LVOT diameter on classification of
368 AS severity. R is the radius of the LVOT; 0.23 cm is the SD of the measurement of the
369 diameter of the LVOT, and 0.04 cm is the largest difference between successive
370 measurements of the LVOT diameter. AVA – Aortic valve area.

371 **Table 1.**

	1st Echo n = 143	2nd Echo n = 143	2nd Echo (>90 day interval) n = 89	Coefficient of Repeatability 1st echo vs. 2nd echo
Age		76.0 (±10.0)		
Sex (Male)		n = 75 (52.4%)		
Rhythm		Normal sinus rhythm: n = 76 Atrial fibrillation: n = 21 Sinus rhythm with bundle branch block: n = 16 Sinus bradycardia: n = 14 Other: n = 16		
Technical quality (Good or adequate visualisation)	n = 79 (55.2%)		n = 69 (48.3%)	
AV area (cm²)*	1.00 (0.76-1.20) [2.46]	1.05 (0.75-1.23) [2.65]	1.01 (0.78-1.23) [2.03]	0.542 (0.0161)
Mean Gradient (mmHg)*	25.9 (18.0-33.0) [54.4]	24.0 (18.0-33.0) [59.0]	26.1 (18.0-33.2) [59.0]	23.95 (0.694)
Peak velocity (mmHg/s)*	336 (288-390) [337]	324 (287-381) [337]	327 (288-393) [305]	154.09 (4.56)
Systolic Volume index (ml/m²)*	35.1 (30.6-44.1) [70.6]	36.2 (28.7-44.5) [66.5]	39.3 (30.3-45.1) [50.5]	16.43 (0.483)
LVOT Diameter (cm²)	2.05 (0.16) [0.52]	2.09 (0.21) [0.86]	2.08 (0.23) [1.45]	1.081 (0.0322)

372 **Table 1.** Mean (standard deviation) and [range] (*median (interquartile range) and [range]) for echocardiographic parameters of aortic stenosis
373 at successive scans. Difference between values of echocardiographic parameters at 2 successive time points expressed as a mean. All
374 comparisons compared with echo 1 are statistically non-significant. Coefficient of Repeatability (within-subjects SEM). Other rhythm includes
375 ventricular paced rhythm, junctional rhythm and non-reported rhythms.

376

377

378

379

380

381

382

383

384

385 **Table 2.**

	Discordant	Concordant	P value
Age	76.8 (10.3)	76.1 (10.5)	>0.05
Sex	Male = 33 Female = 39	Male = 42 Female = 29	>0.05
LVOTd (cm)	1.98 (0.20)	2.12 (0.23)	>0.05
LVEF (%)	54.4 (12.8)	55.2 (14.2)	>0.05
SVi (ml/m²)	33.8 (10.9)	40.7 (11.4)	<0.0001
MPG (mmHg)	27.1 (8.31)	27.8 (13.9)	>0.05
PkV (mmHg/s)	347 (54.8)	344 (77.7)	>0.05

386

387 **Table 2.** Comparison and associations of discordant (echo 1: AVA \geq 1 cm², MPG >40 mm Hg and AVA <1 cm², MPG \leq 40 mm Hg) versus concordant and
388 reclassified (changed AS classification of mild, moderate or severe from echo 1 to echo 2) versus non-reclassified severity subgroups. Abbreviations: LVOTd

389 – Left ventricular outflow tract diameter, LVEF – Left ventricular ejection fraction, SVi – Stroke volume index, MPG – Mean pressure gradient and PkV –
390 peak velocity.

391

392 **Table 3.**

Parameter	Intra-class correlation coefficient (95% CI)	p-value
LVOT Diameter (cm)	0.787 (0.707 – 0.845)	<0.0001
Mean Gradient (mmHg)	0.859 (0.808 – 0.896)	<0.0001
AV area (cm²)	0.757 (0.665 – 0.823)	<0.0001

393 **Table 3.** Correlation coefficients of repeated measurements in patients who had 2 echo studies. Abbreviations: LVOT – left ventricular outflow
394 tract; AV – aortic valve.

395

396

397

398 **Table 4.**

Proportion of	Original	AVA	AVA	AVA	AVA
non-severe	AVA	R + 0.23	R + 0.04	R - 0.04	R - 0.23
AS	(continuity)				
Non-severe	59/140 (42%)	91/140 (65%)	65/140 (46%)	53/140 (38%)	33/140 (24%)
Unindexed n	62/146 (42%)	98/146 (67%)	67/146 (46%)	57/146 (39%)	29/146 (20%)
$\geq 1\text{cm}^2$ (%)					
Indexed n	28/114 (25%)	61/114 (54%)	36/114 (32%)	23/114 (20%)	11/114 (10%)
$\geq 0.6\text{cm}^2$ (%)					

399 **Table 4.** Impact of variation in the measurement of the LVOT diameter on classification of AS severity. R is the radius of the LVOT; 0.23 cm is
400 the SD of the measurement of the diameter of the LVOT, and 0.04 cm is the largest difference between successive measurements of the LVOT
401 diameter. Non-severe is defined as unindexed AVA $\geq 1\text{cm}^2$ or MPG $< 40\text{mmHg}$. AVA – Aortic valve area.

402

403