Comparison of demographics, cardiovascular risk factors profile and prevalence of coexistent atherosclerotic vascular disease in patients with severe aortic stenosis stratified according to dichotomized stenosis severity

Beata Bobrowska1,2, Wojciech Zasada1, Artur Dziewierz1,2, Olga Kruszelnicka3,*, Andrzej Surdacki1,2,*, Dariusz Dudek1,2,*
1Second Department of Cardiology, Jagiellonian University Medical College, Krakow, Poland
2Second Department of Cardiology and Cardiovascular Interventions, University Hospital, Krakow, Poland
3Department of Coronary Artery Disease and Heart Failure, John Paul II Hospital, Krakow, Poland
*Joint senior authors on this work.

Introduction
Degenerative calcific aortic stenosis (AS) is the most frequent valvular heart disease in Europe and North America. The prevalence of this AS form is especially pronounced in the elderly, representing a major public health issue [1–5]. The choice of optimal management of AS patients requires appropriate identification of subjects with severe AS, and echocardiography is a first-line method to diagnose AS and estimate its severity [5]. According to clinical practice guidelines, echocardiographic criteria of severe AS include a calculated aortic valve area (AVA) < 1.0 cm², mean transaortic valve pressure gradient > 40 mm Hg or maximal flow velocity > 4 m/s [5]. Inconsistencies in AS grading were confirmed by cardiac catheterization and a calculated AVA of 0.8 cm² corresponded to a mean transvalvular pressure gradient of 40 mm Hg by echocardiography in patients with severe AS and a normal left ventricular stroke volume [6].

Aim
Our aim was to compare clinical characteristics of patients with severe degenerative AS according to the degree of AVA narrowing categorized by an AVA cut-off value of 0.8 cm².

Material and methods
We retrospectively analyzed data of 145 previously described [7] consecutive patients (66 women and 79 men) with severe degenerative AS who were admitted to our tertiary center between January 2003 and October 2012. The diagnosis of severe AS was based on recognized echocardiographic criteria [5]. All examinations were performed by one of two well-experienced sonographers with the highest level of competence in our department. All patients underwent elective coronary angiography and carotid ultrasonography as a part of routine diagnostic work-up. Patients’ characteristics obtained during the index hospitalization were recorded. The ethics committee of our university was notified about the registry and no objection was raised. A detailed study design was published previously, as well as results of a data analysis primarily focused on the determinants of coexistent coronary and carotid atherosclerosis [7].

Cardiovascular risk factors (arterial hypertension, hypercholesterolemia, diabetes mellitus and self-reported current smoking habit) were defined according to current recommendations as reported previously [7]. Estimated glomerular filtration rate (eGFR) was calculated according to the simplified Modification of Diet in Renal Disease Study formula. Significant coronary artery disease (CAD) was defined as a history of coronary revascularization or the presence of ≥1 diameter stenosis of ≥50% of at least one major epicardial coronary artery on coronary angiography [7]. Relevant internal carotid artery stenosis (ICAS) was defined as stenosis of ≥50% of at least one internal carotid artery [7].

Statistical analysis
Data are shown as mean and standard deviation (SD) for continuous characteristics and numbers (n) and per-
centages for categorical variables. The patients were divided into two groups according to the AVA value: group A with an AVA in the range 0.8–1.0 cm², and group B with an AVA < 0.8 cm². Intergroup differences were calculated by a two-sided Student’s t test for continuous data and two-tailed Fisher’s exact test for proportions.

In order to identify independent predictors of categorized stenosis severity, multiple logistic regression was performed with a dichotomized AVA (≤ 0.8 cm² vs. 0.8–1.0 cm²) as a dependent variable, and age, gender, eGFR and the number of selected traditional risk factors (hypertension, hypercholesterolemia, diabetes mellitus and current smoking habit) as potential predictors. The Hosmer-Lemeshow test was performed to verify the goodness of fit of the regression equation.

In addition, because we had previously described an effect of the age-gender interaction on the prevalence of significant CAD [7], we evaluated whether a similar interaction might affect categorized AS severity. Accordingly, we created an interaction term which was set to 1 in women older than 76 years, and equalled 0 in the remainder. Then the significance of the possible age-gender interaction was tested by entering the interaction term, gender and categorized age (according to a median of 76 years) into a supplemental logistic regression analysis with a dichotomized AVA as a dependent variable.

### Statistical analysis

A p-value below 0.05 was assumed significant. All analyses were performed using a licensed statistical software: Statistica (data analysis software system), version 12 (StatSoft, Inc., 2014, Tulsa, OK, USA).

### Results

As shown in Table I, patients with more advanced AS (group B: AVA < 0.8 cm²) were older, more frequently women, and with a lower prevalence of smoking habit compared to subjects with an AVA in the range

| Characteristics | Group A AVA: 0.8–1.0 cm² N = 67 | Group B AVA < 0.8 cm² N = 78 | P-value |
|-----------------|---------------------------------|-----------------------------|---------|
| Age and gender: |                                 |                             |         |
| Age [years]     | 72 (9)                          | 77 (9)                      | 0.006   |
| Gender, men/women, n (%) | 44/23 (66/34) | 35/43 (45/55) | 0.013   |
| Clinical characteristics, n (%): |                         |                             |         |
| Arterial hypertension | 58 (87)                     | 64 (82)                     | 0.5     |
| Hypercholesterolemia | 49 (73)                     | 48 (62)                     | 0.16    |
| Diabetes mellitus | 24 (36)                         | 28 (36)                     | 1       |
| Smoking habit | 11 (16)                         | 4 (5)                       | 0.03    |
| Echocardiographic parameters: |                             |                             |         |
| Left ventricular ejection fraction, n (%) | 56 (12) | 56 (14) | 0.9 |
| PG-max [mm Hg] | 80 (26)                         | 98 (26)                     | < 0.001 |
| PG-mean [mm Hg] | 50 (20)                        | 59 (17)                     | 0.003   |
| Coexistent coronary and carotid atherosclerosis, n (%): |               |                             |         |
| Significant CAD | 42 (63)                         | 44 (56)                     | 0.5     |
| Relevant ICAS | 11 (16)                         | 11 (14)                     | 0.8     |
| Biochemical characteristics: |                             |                             |         |
| Total cholesterol [mmol/l] | 4.4 (1.1) | 4.6 (1.2) | 0.3 |
| LDL-C [mmol/l] | 2.6 (1.0)                        | 2.5 (0.9)                   | 0.5     |
| HDL-C [mmol/l] | 1.4 (0.5)                        | 1.3 (0.5)                   | 0.3     |
| Triglycerides [mmol/l] | 1.3 (0.4) | 1.4 (0.8) | 0.3 |
| eGFR [ml/min/1.73 m²] | 78 (29) | 78 (29) | 0.98 |

Data are presented as mean (SD) or n (%). CAD – coronary artery disease, eGFR – estimated glomerular filtration rate, HDL-C – high-density lipoprotein cholesterol, ICAS – internal carotid artery stenosis, LDL-C – low-density lipoprotein cholesterol, PG-max – maximal transvalvular aortic pressure gradient, PG-mean – mean transvalvular aortic pressure gradient.
0.8–1.0 cm² (group A). The two groups did not differ in the proportion of other traditional risk factors or biochemical characteristics. Maximal and mean transaortic pressure gradients were higher in group B, while left ventricle ejection fraction and biochemical characteristics were comparable in both groups. The proportion of subjects with coexistent significant CAD or relevant ICAS was similar across the groups (Table I).

Multiple logistic regression confirmed an independent association of stenosis severity with female gender, while a weak positive effect of age did not reach statistical significance (Table II). Curiously, stenosis severity was not more pronounced with increasing number of classical atherosclerotic risk factors; on the contrary, an opposite, yet not significant, relationship was observed (Table II).

There was no interaction between age and gender in terms of the categorized degree of stenosis severity (p = 0.3). In other words, the proportion of AS patients with an AVA < 0.8 cm² was higher in those with an over-median age (> 76 years) compared to their younger counterparts both in women (31/42 vs. 12/24, i.e. 74% vs. 50%, p = 0.06) and men (14/28 vs. 21/51, i.e. 50% vs. 41%, p = 0.5).

Discussion

Our principal finding was a lack of differences in the prevalence of traditional atherosclerotic risk factors according to the degree of AS. Such a difference might be expected assuming a putative contribution of atherosclerotic risk factors to the development of degenerative AS [8, 9], keeping in mind the progressive pattern of AS. Our multivariate regression analysis showed a lower, albeit insignificantly, number of classical risk factors in patients with more severe AS.

However, our results are consistent with a study by Ortlepp et al. [10], who reported that the presence of coexistent CAD, but not severe calcific AS by itself, was associated with increased prevalence of traditional risk factors. In their study, none of the traditional risk factors, including hypertension, hypercholesterolemia, diabetes mellitus and smoking, showed a significantly increased proportion in patients with severe AS compared to those with a normal aortic valve, irrespective of the presence of significant CAD on angiography. In particular, Ortlepp et al. [10] described a lower prevalence of smoking habit in AS patients compared to non-matched controls without AS, in some analogy to a decreased percentage of smokers in subjects with more severe AS in the present study. Additionally, in that study, CAD prevalence increased gradually with the increasing number of risk factors [10], in agreement with our previous analysis focused on determinants of the presence of coexistent atherosclerosis in the same study group [7]. Accordingly, since in the present study the prevalence of coexistent significant CAD or relevant ICAS was similar in those with an AVA < 0.8 cm² and 0.8–1.0 cm², this is likely to explain comparable percentages of the traditional risk factors in groups A and B.

Admittedly, we did not study control subjects without severe AS. Nevertheless, similar mechanisms appear to account for the development of severe AS and its further progression. Therefore, our results support the notion that the link between AS and atherosclerotic risk factors may be mainly due to their association with accompanying CAD [10]. In line with this concept, statins failed to retard the progression of degenerative AS in a recent meta-analysis [11]. So, despite some similarities, atherosclerosis and degenerative AS may be regulated by different pathways, which was also suggested by a lack of correlation between aortic valve calcification and coronary artery calcium score or ascending aorta calcifications in patients aged ≥60 years [12]. Notably, only age and male gender, but not other risk factors, were significant independent predictors of progression to AS over a mean 5-year follow-up in elderly subjects with normal aortic valves or aortic valve sclerosis without AS at baseline [13]. Furthermore, after exclusion of patients with CAD, traditional risk factors were not associated with subsequent surgery for AS over a mean follow-up of 10.5 years in a recent nested case-referent prospective study [14], which also supports our findings.

That our study patients with more advanced valve stenosis were older by about 5 years is consistent with progression of degenerative AS with age. Additionally, a higher proportion of females in this subgroup is not unexpected, keeping in mind the 8-year longer average life expectancy in women compared to men in Poland [15].

In a previous analysis of the same study group [7], we proposed a “survival bias” to explain an observation that the prevalence of significant coronary atherosclerosis in AS patients with an over-median age (>76 years) was significantly higher compared to their younger counterparts only in men (89% vs. 55%, respectively, p = 0.002), while the corresponding proportions were similar in women, exhibiting, counterintuitively, even a slight opposite tendency (48% vs. 54%) [7]. The hypothetical survival bias

**Table II. Logistic regression analysis of predictors of an aortic valve area below 0.8 cm²**

| Predictor variable | Odds ratio (OR) | Mean OR (95% CI) | P-value |
|--------------------|-----------------|------------------|---------|
| Age (per 10-year increment) | 1.47 (0.97–2.23) | 0.07 |
| Gender (women vs. men) | 2.23 (1.03–4.81) | 0.04 |
| Number of risk factors (per increment of 1) | 0.71 (0.47–1.06) | 0.09 |
| eGFR (per rise of 10 ml/min/1.73 m²) | 1.01 (0.89–1.15) | 0.84 |

CI – confidence interval, eGFR – estimated glomerular filtration rate. p = 0.3 by the goodness-of-fit Hosmer-Lemeshow test for the regression equation.
could result from a high mortality in women with severe AS and coexistent CAD, which might have abolished the expected age-dependent rise in the prevalence of significant CAD in women with severe AS [7].

In contrast, in the present analysis, a higher prevalence of more advanced AS in older AS patients (aged > 76 years) compared to younger AS subjects was observed irrespective of sex, i.e. both in men (50% vs. 41%) and women (74% vs. 50%). This finding strengthens the previously proposed concept of survival bias, being concordant with the notion that a hypothetically altered mortality pattern in women with severe AS could be linked to coexistent CAD. Accordingly, we had previously found an interaction between age and gender with regard to concomitant prevalence of significant coronary atherosclerosis [7]. However, such an interaction was absent in the present study regarding the degree of AS severity. The latter would be expected if that interaction were related to a higher premature mortality of women owing to more severe AS, but not to concomitant CAD.

Conclusions

Within patients with severe AS, the degree of stenosis severity is associated with a similar prevalence of coexistent coronary and carotid atherosclerosis and atherosclerotic risk factors. Thus, a link between AS and risk factors may result from their association with concomitant atherosclerosis rather than AS by itself.

Conflict of interest

The authors declare no conflict of interest.

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