Ivabradine Might Improve Exercise Capacity in Dilated Cardiomyopathy in Comparison With Bisoprolol: Assessment With the Ergospirometry Test

La ivabradina podría mejorar la capacidad funcional frente al bisoprolol en la miocardiopatía dilatada: valoración con test de ergospirometría

To the Editor,

The 2012 Guidelines for Heart Failure include an indication for ivabradine according to the original design of the SHIFT study, and recommend that this drug be used in patients who, despite optimal treatment and maximum tolerated doses of beta-blockers, angiotensin-converting enzyme inhibitors, and aldosterone antagonists, have a heart rate (HR) in sinus rhythm above 70 bpm (Level of Evidence: Class IIa, Level B). Over the past 2 decades, a vast amount of information has accumulated on the importance of ergospirometry in the functional assessment of patients with dilated cardiomyopathy. The role of this test has begun to be recognized in the clinical context, where it is now considered an essential tool. The primary aim of this study was to assess functional capacity by analyzing several ergospirometry parameters and the Keteyian chronotropic index in patients with dilated cardiomyopathy and severe left ventricular systolic dysfunction treated with ivabradine compared with those treated with bisoprolol.

The study was based on a retrospective design and was conducted at a teaching hospital. All ergospirometry tests performed between July 2012 and September 2013 were reviewed, and we recruited all patients with dilated cardiomyopathy and severe left ventricular systolic dysfunction who had been receiving drug therapy for at least 6 months and were clinically stable. We excluded patients with atrial fibrillation and a cardiac resynchronization therapy device. The study was approved by the Clinical Research Ethics Committee of the hospital. The patient cohort was divided into 2 groups according to the treatment received: group I (beta-blockers) and group II (ivabradine). Patients in group II were not receiving beta-blockers due to documented pulmonary intolerance (asthma or a history of severe chronic obstructive pulmonary disease [Gold 3]). The clinical data analyzed included age, sex, smoking habits, hypertension, diabetes mellitus, hypercholesterolemia, drug therapy, body mass index, maximum oxygen uptake (VO₂max) (expressed as mL/kg/min), linear regression slope of the ventilatory equivalent (VE) for carbon dioxide (VCO₂), and the Keteyian chronotropic index. The formula of this index is defined as follows: (HR at peak exercise – resting HR) / [119 + (resting HR/2) – (age/2) – (5 x 0) – resting HR]. The statistical analysis was performed using SPSS 15.0 (Chicago, Illinois, United States).

Our study population included 26 patients, 13 in each group. All study participants achieved respiratory exchange above 1.1 (which indicates adequate exertion). At the time the ergospirometry was performed, the patients had an HR below 60 bpm. The characteristics of the study population are listed in the Table. There were no statistically significant differences between the 2 groups in age, sex, cardiovascular risk factors, body mass index, or drug therapy. In terms of ergospirometry parameters, patients with dilated cardiomyopathy and severe left ventricular systolic dysfunction treated with ivabradine achieved a higher VO₂max

<table>
<thead>
<tr>
<th>Variables</th>
<th>Group I (BB)</th>
<th>Group II (ivabradine)</th>
<th>p</th>
</tr>
</thead>
<tbody>
<tr>
<td>Age, mean (SD), y</td>
<td>57 (14)</td>
<td>59 (13)</td>
<td>0.72</td>
</tr>
<tr>
<td>Men</td>
<td>8 (6.15)</td>
<td>7 (53.8)</td>
<td>0.69</td>
</tr>
<tr>
<td>Hypertension</td>
<td>3 (23.1)</td>
<td>4 (30.8)</td>
<td>0.65</td>
</tr>
<tr>
<td>Diabetes mellitus</td>
<td>0 (0)</td>
<td>2 (15.4)</td>
<td>0.14</td>
</tr>
<tr>
<td>Smoker</td>
<td>1 (7.7)</td>
<td>0 (0)</td>
<td>0.3</td>
</tr>
<tr>
<td>Hypercholesterolemia</td>
<td>1 (7.7)</td>
<td>2 (15.4)</td>
<td>0.53</td>
</tr>
</tbody>
</table>

Treatments used included Ramipril (5 mg/12 h) 13 (100) 13 (100) 1
Furosemide (80 mg/24 h) 13 (100) 13 (100) 1
Eplerenone (25 mg/24 h) 13 (100) 13 (100) 1
Rosuvastatin (20 mg/24 h) 13 (100) 13 (100) 1
Body mass index, mean (SD)    | 25.21 (4.62) | 27.52 (4.68)           | 0.21|
VO₂max, mean (SD), mL/kg/min  | 16.60 (3.88) | 20.77 (3.68)           | 0.01|
Oxygen pulse, mean (SD)       | 11.86 (3.23) | 14.70 (6.51)           | 0.17|
VE/VCO₂ slope, mean (SD)      | 33.35 (3.64) | 27.44 (4.31)           | 0.001|
Keteyian chronotropic index,  | 0.96 (0.08)  | 0.86 (0.09)            | 0.01|
mean (SD)  

BB, beta-blockers; VCO₂, carbon dioxide output; VE, ventilatory equivalent; VO₂max, maximum oxygen uptake.

Unless otherwise indicated, the data are expressed as No. (%).

lower VE/VCO₂ slope, and a lower Keteyian chronotropic index than those treated with bisoprolol.

Our study can be considered a proof-of-concept study, as it shows that patients with dilated cardiomyopathy and severe left ventricular systolic dysfunction after 6 months of ivabradine therapy exhibit better functional and ventilatory capacity. Our results are consistent with those of a study by Votterlani et al., which showed that ivabradine alone or in combination is more effective than carvedilol in improving exercise tolerance among patients with heart failure.

Our study is unique in that all patients had an HR below 60 bpm in the ergospirometry test, unlike that by Votterlani et al., who observed higher HR values. Likewise, we used a new equation, the Keteyian chronotropic index. This is a recently defined equation to predict maximal HR in patients with heart failure who take beta-blockers. Our analysis showed that exercise capacity improved among patients receiving ivabradine therapy, possibly due to a slower increase in HR during exercise, as expressed by the Keteyian chronotropic index. From the pathophysiologic point of view, this may slow the increase in left ventricular filling pressure, delaying the onset of dyspnea and the end of exercise.

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Table: Baseline Characteristics of the Study Population
REFERENCES


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Ventricular Septal Defect as an Unusual Complication of Percutaneous Transcatheter Implantation of an Aortic Valve Prosthesis: Two-year Follow-up

Comunicación interventricular como complicación inusual del implante percutáneo de prótesis biológica aórtica: evolución a 2 años

To the Editor,

We report the case an 87-year-old man with severe aortic stenosis and dyspnea, New York Heart Association functional class III, and a class I indication for valve replacement. His medical history included hypertension, chronic obstructive pulmonary disease, and bilateral knee replacement; his logistic EuroSCORE was 13.92%. The technique chosen by the medical-surgical team was transfemoral transcatheter aortic valve implantation.

Preimplantation transthoracic echocardiography showed a critical aortic stenosis of 0.40 cm² (0.25 cm²/m² when corrected for body surface area), generalized moderate left ventricular hypertrophy, and mild systolic dysfunction. Computed tomography angiography showed a sparsely calcified right common femoral artery of good caliber and a low degree of tortuosity in the aortoiliac segment. The minimum diameters of the right common femoral, right external iliac, and right common iliac arteries were 7 mm, 9 mm, and 11 mm, respectively. The distance from the annulus to the coronary ostia was 11.5 mm from the left coronary artery and 13 mm from the right coronary artery. There was asymmetric moderate valvular calcification and the left coronary and noncoronary leaflets were the most affected; the annulus was 21 × 23 mm and there was no subannular aortic calcification (Figure 1).

Transesophageal echocardiography showed an annulus of 21 to 22 mm and asymmetric severe valvular calcification mainly of the left coronary and noncoronary leaflets (Figure 1).

Figure 1. Image of the left ventricular outflow tract and the aortic valve by transesophageal echocardiography and computed tomography before transcatheter aortic valve implantation. Note the asymmetric calcification of the leaflets and the absence of subannular aortic calcification.