Misinterpretation of the Borg’s Rating of Perceived Exertion Scale by patients with panic disorder during ergospirometry challenge

Ricardo William Muotri, Marcio Antonini Bernik, Francisco Lotufo Neto

ABSTRACT

Introduction The most widely used instrument to measure perceived exertion or exercise intensity is the Borg’s Rating of Perceived Exertion (RPE) Scale. Panic attacks are aversive experiences that may be triggered by bodily sensations such as palpitations, breathlessness or dizziness due to increasing autonomic distress, prior sensitisation to panic and fear conditioning. The consequence is catastrophic interpretation of bodily sensations of arousal in general, which can lead to misinterpretation of exertion or exercise intensity.

Purpose To verify the effectiveness of the Borg’s RPE Scale as a measure of perceived effort in panic disorder subjects.

Methods The study enrolled 72 subjects: the control group (C, healthy sedentary subjects, n=30) and patients with panic disorder (PD, n=42). All subjects were submitted to an ergospirometry challenge. Perceived exertion RPE scores and heart rate at 90% VO2max were compared between groups.

Results Patients with PD showed lower levels of maximal oxygen consumption, when compared with C group [VO2max (mL/kg/min): PD=29.42±6.50xC=34.51±5.35; Student’s t-test=−3.51; p<0.05]. Furthermore, among PD subjects, the maximum heart rate during ergospirometry challenge was lower than expected [predict max HR (bpm)=200.85±6.33xheart rate test (bpm)=178.86±7.28; Z=−5.64; p<0.05]. Perceived exertion, as measured by RPE, was also overestimated in relation to heart rate at 90% of maximum oxygen consumption compared with controls (RPE90% VO2max: PD=18.93±0.55xC=16.67±0.60; U=8.00; Z=7.42; p<0.05).

Conclusions The present study findings suggest that the Borg’s RPE Scale may not be an appropriate measure of subjective exertion among subjects with panic disorder performing a cardiopulmonary exercise test.

INTRODUCTION

The most widely used instrument to measure perceived exertion or exercise intensity is the Borg’s Rating of Perceived Exertion (RPE) Scale (6–20).1 Objective measures of effort, such as heart rate (HR) and oxygen uptake, have been used in conjunction with RPE scores in clinical settings such as cardiac rehabilitation2–5 and with patients receiving β-blocker therapy.6 Using the Borg’s RPE scale improves the safety of the challenge test when used in conjunction with objective measures.

According to its author, RPE scores correlate well with both physiological measures of stress and arousal (eg, HR, ventilatory
The subjective weighting of these physiological changes generates a response, which is measured by the RPE scale to provide a final score of subjective exertion.

The putative similar perception of the exercise intensity (corresponding to anaerobic thresholds) among different individuals makes it possible to adequately prescribe exercise intensity.

Of the many exercise challenge protocols available, Heck's test is the most widely used in sedentary subjects, due to its high reproducibility. This protocol determines the anaerobic threshold of the individual, which is the point at which the rate of lactate production exceeds that of removal.

It is possible that patients with panic disorder do not perceive exertion in the same way as the general population. Spontaneous panic attacks (PAs) are bouts of inappropriately released fear. However, it is more likely that the underlying physiological mechanism is another brain defence reaction such as a mammalian suffocation alarm. Accordingly, a PA is usually associated with marked air hunger, which is not characteristic of external danger-induced fear. Another important difference is the lack of the emergency activation of the hypothalamic–pituitary–adrenal axis with lower levels of adrenocorticotropic hormone and cortisol observed in PAs.

Further situational PAs and interpanic tonic anticipatory anxiety evolve due to several factors, one of which is fear conditioning. Over time, bodily sensations of arousal such as palpitations, breathlessness or dizziness can come to trigger PA due to increasing autonomic distress, sensitisation to panic and fear conditioning leading to catastrophic interpretation of those sensations as symptoms of an impending medical problem. Consequently, avoidance of physical exercise, which share with fear activation the same arousal bodily sensations, has been previously reported (eg, Muotri and Bernik), and was observed in this sample.

Putatively, catastrophic interpretation of exercise-induced bodily sensations can alter the perception of current state of fatigue. Fatigue compromises exercise performance and is determined by central and peripheral mechanisms. Perceived exertion and fatigue are extremely important in the regulation of self-paced physical activity. These mechanisms have been proposed to interfere with the self-evaluation in the RPE Scale through an altered sensory tolerance limit.

In the present study, we hypothesise that patients with panic disorder (PD) overestimate their exertion rendering the RPE Scale scores inappropriate for evaluating cardiovascular (CV) capacity in the patient with PD population.

### MATERIALS AND METHODS

#### Ethical approval and study registration

Patients and controls were aware of the objectives, methods, potential risks and benefits of the study and signed consent forms. The Department of Psychiatry, University of São Paulo Medical School and the Hospital Ethics Committee approved the study. The study received a grant towards its total costs by the São Paulo State Foundation for the Development of Science.

#### PARTICIPANTS

The controls were mostly graduate students and staff of the Institute of Psychiatry, recruited after an internal ad (n=54). All were healthy, sedentary and devoid of any psychiatric disorders (using the Mini International Neuropsychiatric Interview; MINI). Patients with PD were recruited in the Emergency Room of the Institute of Cardiology in our hospital or through self-referrals for treatment to our group (Anxiety Disorders Program) after radio announcements, social media outreach and newspaper ads (n=50).

Fifty patients with PD were selected for this trial. Eight patients were excluded from the study for not having PD as the main diagnosis. Thus, 34 healthy, sedentary subjects, aged 18–55 years, who were devoid of any psychiatric disorders, were selected for this trial as controls. Four of these healthy sedentary subjects were unable to complete the challenge test due to personal reasons (figure 1).

The MINI interview was used for the psychiatric diagnosis (or lack of it) in both groups.

Other exclusion criteria were lifetime cardio-circulatory, respiratory or endocrine diseases including hypertension, pregnancy and epilepsy.

Seventy-two subjects were included in this study—patients: 22 women and 20 men (mean age 33.02 years, SD 8.21 years) and controls: 20 women and 10 men (mean age 36.17 years, SD 7.25 years).

There were no differences in baseline characteristics between the PD group and the control (C) group (n=72). The analysed characteristics were (mean±SD): age=34.33±7.92 years, Z=-1.83, U=469.50; body mass index (BMI)=23.35±3.81kg/m², t=1.40; gender (58.33% women, χ²=1); marital status (61.11% married, χ²=0.107) and sedentary status (72.22% sedentary, χ²=0.127).

There were differences between groups in terms of occupational status (73.66% working outside the home, n=26 PD group, n=27 C group, χ²=7.11, p=0.013) and schooling level (26.38%, below the university level, n=16 PD group, n=03 C group, χ²=7.11, p=0.013). No other differences were observed between groups, including the frequency of lifetime smoking (patients, 12 out of 42 and controls, 3 out of 30, χ²=3.66).

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Subjects were free from psychotropic medications for at least 4 weeks before the evaluation, alcohol free for at least 36 hours and xanthine or caffeine free for at least 8 hours. Smokers were instructed not to smoke for at least 2 hours before the evaluation, and all subjects consumed a light snack 3 hours before the test. The test took place between 11:00 and 00:00.

Eighty-nine per cent of the subjects (patients: 38; controls: 26) had previously undertaken an ergospirometry challenge.

The tests took place in an ambient temperature of 22°C–25°C.

Before the test, the Physical Activity Readiness Questionnaire was used for assessment of CV risk even though the Heck’s test protocol presents a very low risk for cardiac events (1/10,000–1/20,000).

The ergospirometry challenge assessed CV capacity. The stress test was conducted on a medical grade treadmill (IMBRAMED, ATL – 10 200) with variable speed (km/hour) and slope (%). A modified version of the Heck’s stress test protocol was used with fixed speed and increasing slope increments at a rate of 2% per minute.

The test velocity was selected after two pilot tests with different speeds, performed minutes before the test, which also allowed subjects to familiarise themselves with the settings. Once the speed was chosen, the subjects had a 1 min rest period, and then began the protocol at the speed previously chosen. In the recovery phase, which lasted 3 mins, the speed was decreased steadily, in 1 min increments.

The perceived exertion was evaluated by the patient at each stage of the cardiopulmonary exercise test on a linear scale with 15 points (6–20) as described by Borg, Alhoua’i et al and Faulkner and Eston (figure 2).

The ECG was performed with 13 leads (HeartWere, Ergo 13) at rest, during the effort phase and the recovery phase of the exercise test.

Throughout the test, subjects were blind to their current HR.

The airflow and volume analysis was performed by a two-way pre-pressure differential pressure pneumotachometer with high precision and dead space of 39 mL. The pneumotachometer was calibrated prior to each test with a syringe by 10 movements (five expirations and five inspirations) with a capacity of 3 L and dead space of 100 mL, used as a correction factor that determined the reading of the respiratory volume. The expired oxygen pressures (PETO₂) were measured using a zirconia-type cell with a high accuracy and fast response (±0.03% O₂), whereas the expired pressures of carbon dioxide (PET CO₂) were measured with an infrared system with ±0.05% CO₂ accuracy and response <130 ms. The O₂ and CO₂ analysers were calibrated, before and immediately after each test with two known gas mixtures of nitrogen and atmospheric air.

The individuals were positioned on the mat wearing a helmet with a sterilised mouthpiece attached and the nose sealed with a catch.

Throughout the test protocol, expired gases were collected and analysed at each respiratory cycle: pulmonary ventilation, respiratory rate, tidal volume (VC), oxygen consumption (VO₂), carbon dioxide (VCO₂) production, (VCO₂/VO₂), the ventilatory equivalent of oxygen (VE/VO₂), the final expired oxygen pressure (PETO₂), the ventilatory equivalent of carbon
dioxide (VE/VO\textsubscript{2}) and the final expired pressure of dioxide of carbon (PETCO\textsubscript{2}). Ventilatory variables were recorded instantaneously, and then means were calculated over 30-second intervals.\textsuperscript{27}

The amount of VO\textsubscript{2} is calculated from the difference between the PO\textsubscript{2} of the inspired air (atmospheric standard) and the PO\textsubscript{2} of the expired air as well as the expired PCO\textsubscript{2}.\textsuperscript{25–28} The VO\textsubscript{2max} was verified when subjects reached at least three of the following physiological validation criteria: (1) VO\textsubscript{2} plateau, when there was no increase in VO\textsubscript{2} greater than 2.0 mL/kg/min for an increase of inclination between the penultimate and the last stage of the test;\textsuperscript{29} (2) respiratory quotient maximum, equal to or greater than 1.10;\textsuperscript{30} (3) maximum HR, equal to or greater than 95% of the maximum chronotropic response predicted for age to the equation [208 – (0.7 x age)];\textsuperscript{31,32} (4) a value equal to or greater than 18 on the subjective perception scale of Borg\textsuperscript{33} signs of extreme fatigue such as: intense hyperpnoea, excessive sweating, facial flushing or difficulty in maintaining adequate motor coordination with the increment of speed of the treadmill.\textsuperscript{33–34} As a subjective criterion, the Borg’s RPE Scale of fatigue perception was used in all tests as a means of complementing exercise intensity monitoring.\textsuperscript{35}

After the test, patients were referred for treatment in our group. Controls received information about their CV capacity and individualised recommendations for initiating physical exercises.

**STATISTICAL ANALYSIS**

The data were initially tested for normality using the Kolmogorov–Smirnov test.

When normality assumptions were not confirmed or in variables where scores are natural numbers, non-parametric tests were used. The Mann-Whitney U test with Bonferroni correction was used to compare groups when normality assumptions were not confirmed (age, resting HR, predict maximum HR, HR reached during test, HR at 90% of maximal oxygen consumption, Borg’s RPE Scale values at 90% of maximal oxygen consumption and being a smoker). Otherwise, the groups were compared using the Student’s t-test.

For quantitative and qualitative variables, Pearson’s \( \chi^2 \) test was applied to investigate linear correlation. When there were less than six subjects in the comparisons (as in the variables smoking, schooling level and occupational status) the Fisher’s exact test was chosen.

The difference observed between the predicted target HR for the test and the maximal HR in the test for both groups was compared using the Student’s t-test and, when checking a significant difference between the groups, we also used the Wilcoxon W test. The Mann-Whitney U test was used to compare the values of HR at 90% of VO\textsubscript{2max} and the RPE at 90% of VO\textsubscript{2max}.

The level of significance was set at \( \leq 0.05 \) (for two-tailed comparisons) throughout the study. Statistical analyses were performed with Statistical Package for the Social Sciences (SPSS) V.14.0.

**RESULTS**

Patients with PD showed lower levels of maximal oxygen consumption when compared with the control group (VO\textsubscript{2max}(ml/kg/min): PD=29.42 ± 6.50, C=34.51 ± 5.35; \( t=\) −3.51; \( p<0.05 \)) (figure 3 and table 1).

The maximum HR during ergospirometry challenge was lower than expected in patients with PD (predict max HR (bpm)=200.85±6.33xheart rate test (bpm)=178.86±7.28; \( Z=\) −5.64; \( p<0.05 \)).

The perceived exertion characterised by RPE was overestimated in relation to HR at 90% of maximum oxygen consumption in patients with PD when compared with the controls (HR90%VO\textsubscript{2max}(ml/kg/min): PD=160.97±6.55xC=164.73±4.48; \( U=\) 408.00; \( Z=\) −7.28; \( p<0.05 \)) and RPE90%VO\textsubscript{2max}: PD=18.93±0.55xC=16.67±0.60; \( U=\) 8.00; \( Z=\) −7.42; \( p<0.05 \), as shown in figure 4 and table 1.

There were no differences between smokers and non-smokers in relation to neither the subjective perception of effort (RPE scores) nor the VO\textsubscript{2max} (ml/kg/min) (figure 5 and table 1). Accordingly, smokers were not analysed as a separate group as it would reduce the sample size and compromise the data analysis.

**DISCUSSION**

Demographic characteristics of the present study subjects are similar to those reported in the literature for patients with PD who are selected for treatment programmes—predominantly younger patients and women (eg.\textsuperscript{35–38}). In the present study, which focused on physical performance, we observed that patients with PD tend to be sedentary and slightly overweight.
Table 1  Participants’ physical performance results by group

<table>
<thead>
<tr>
<th>Performance characteristics</th>
<th>PD n=42</th>
<th>C n=30</th>
<th>Total n=72</th>
<th>p Value</th>
<th>Value of statistical test</th>
</tr>
</thead>
<tbody>
<tr>
<td>RHR (bpm)</td>
<td>84.00±7.32</td>
<td>85.23±8.09</td>
<td>84.51±7.62</td>
<td>&gt;0.05</td>
<td>Z=-0.69, U=569.00</td>
</tr>
<tr>
<td>PMHR (bpm)</td>
<td>200.85±6.33</td>
<td>207.34±0.95</td>
<td>203.55±5.82</td>
<td>&lt;0.05</td>
<td>Z=-7.00, U=29.00</td>
</tr>
<tr>
<td>HRRDT (bpm)</td>
<td>178.86±7.28</td>
<td>183.03±4.97</td>
<td>180.60±6.71</td>
<td>&lt;0.05</td>
<td>Z=-2.54, U=408.00</td>
</tr>
<tr>
<td>VO₂max (mL/kg/min)</td>
<td>29.42±6.50</td>
<td>34.51±5.35</td>
<td>31.54±6.51</td>
<td>&lt;0.05</td>
<td>t=-3.51</td>
</tr>
<tr>
<td>90%VO₂max (mL/kg/min)</td>
<td>26.48±5.85</td>
<td>31.06±4.81</td>
<td>28.39±5.86</td>
<td>&lt;0.05</td>
<td>t=-3.51</td>
</tr>
<tr>
<td>HR 90%VO₂max (bpm)</td>
<td>160.97±6.55</td>
<td>164.73±4.48</td>
<td>162.54±6.04</td>
<td>&lt;0.05</td>
<td>Z=-2.54, U=408.00</td>
</tr>
<tr>
<td>RPE 90%VO₂max</td>
<td>18.93±0.55</td>
<td>16.67±0.60</td>
<td>17.99±1.26</td>
<td>&lt;0.05</td>
<td>Z=-7.42, U=8.00</td>
</tr>
<tr>
<td>RPE scale to 90% VO₂max and HR measured at 90% VO₂max in the PD group</td>
<td></td>
<td></td>
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</tr>
<tr>
<td>Smoker</td>
<td>2.27±1.28</td>
<td>&gt;0.05</td>
<td>2.27±1.28</td>
<td>&gt;0.05</td>
<td>Z=-0.87, U=148.50</td>
</tr>
<tr>
<td>Non-smoker</td>
<td>1.70±1.34</td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>HRRDT x RPE scale to 90% VO₂max and HR measured at 90% VO₂max</td>
<td>PD</td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>r= -0.511*</td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>C</td>
<td>r=0.005*</td>
<td></td>
<td></td>
<td></td>
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</tr>
</tbody>
</table>

The results are presented as mean and SD.

*Pearson’s correlation. Alpha was set to 0.05 for all statistical analyses.

HR, heart rate; HRRDT, heart rate reached during test; HR 90%VO₂max, heart rate at 90% of maximal oxygen consumption; PMHR, predict maximum heart rate; RHR, resting heart rate; VO₂max, maximal oxygen consumption; 90%VO₂max, 90% of the maximal oxygen consumption; RPE 90%Vo2max, Borg’s RPE Scale values at 90% of maximal oxygen consumption.

(BMI=25.35 kg/m²). This has also been reported previously (eg, Lykouras and Michopoulos). There is a wealth of studies reporting on cardiopulmonary exercise testing (CPX) in patients with PD. Evidence suggests that aerobic fitness may be reduced in this clinical condition. As already noted by other authors, VO₂max tends to be lower in patients with PD than in age-matched and sex-matched healthy subjects. This finding has been replicated in the present study (table 1). In addition, patients with PD have been shown to have a significantly higher resting HR and resting tachycardia than healthy subjects.

There are two possible explanations: the first is that this lower aerobic fitness observed in many patients with PD may result from avoidance of strenuous activity that might trigger PAs. This leads to a reduction of exposure to exercise and outside activities that could improve subjects’ aerobic fitness. In fact, regular exercise improves this condition, leading to levels of aerobic fitness similar to that of healthy subjects.

Alternatively, although the mechanism is not clear, performance of maximal CPX may be hampered by panic anxiety during the challenge. Autonomic manifestations naturally triggered by exercise are similar to those present in a PA. It is our view that patients with PD are unwilling to achieve the exhaustion that characterises a maximal CPX due to anticipatory anxiety related to a PA. In accordance with this idea is the fact that patients with PD overestimate their degree of physical exhaustion as observed in our study (figures 3 and 4).

In the control group, there was a clear correlation between the percentage of the target maximum HR reached and the RPE score reported at all score levels (figure 4). This is a finding widely reported in the literature (eg, 48–50). On the other hand, in patients with PD, there was no correlation between the percentages of target HR max with the reported RPE (figure 6 and table 1) in scores lower than 17. This is consistent with the model of ‘catastrophic’ interpretation of autonomic symptoms in panic patients as proposed by Clark and corroborated by Carton and Rhodes and Noble. This finding supports our initial hypothesis.
Finally, in the present study there were no differences between smokers and non-smokers in relation to the subjective perception of effort RPE and VO2max. This is in contrast with previous reports in the literature. One possible explanation is the small sample size for this comparison, which yielded a power (1-β) of <0.8%, which is too small to rule out type II errors.

CONCLUSIONS

Perceived effort comprises several distinct inputs that are perceived differently across the duration of an executed task and among different individuals. Thus, exertion may be only one of many perceptual features experienced during exercise engagement. For some populations, it may be not appropriate to measure effort solely through the conceptual lens of ‘exertion’. This seems to be particularly true for patients with a diagnosis of panic disorder.

Psychological influences on perceived effort are hypothesised to occur when signals arising in the sensory cortex are matched with the so-called ‘perceptual-cognitive reference filter’. Therefore, to evaluate perceived effort with only one physiological index is an oversimplification of the psychophysiological construct.

More research is needed to establish a comprehensive concept of perceived effort and to develop an associated multidimensional instrument, including psychophysiological determinants, to measure perceived effort. One alternative would be to understand the role of the central corollary discharge, a copy of the neural drive from the brain to the working muscles that provides a signal from the motor system to sensory systems and is considered a feedback mechanism that might influence fatigue and consequently exercise performance.

In summary, these findings suggest that in patients with PD, a simple rating of perceived exertion as indexed by the Borg’s RPE Scale may not be sufficient to accurately capture the intensity of physical effort.

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Competing interests None declared.

Ethics approval Ethics Committee for Analysis of Research Projects FMUSP (CAPPESQ).

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Author note This study contributes to a better cardiorespiratory factor analysis in a specific population.

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