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## Gender Differences in Appraisal of Stress and Coping 5 Years after Heart Transplantation

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### Abstract

**OBJECTIVES**—We examined whether gender differences exist regarding stress, symptom distress, coping, adherence, and social support 5 years after heart transplantation.

**BACKGROUND**—Differences exist in health-related quality of life outcomes by gender after heart transplantation; women report poorer outcomes.

**METHODS**—Patients (n=210, female=42), were from a prospective, multi-site, study of health-related quality of life long-term after heart transplantation. Patients completed self-report instruments 5 years after heart transplantation (mean=4.98±0.17 years after transplant). Statistical analyses included two-sample t-tests, Chi-square or Fisher's exact test, and multivariable modeling.

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**RESULTS**—Women did not report more overall stress or symptom distress, but reported more difficulty adhering to the transplant regimen, yet more actual adherence than men. Women reported using more negative coping styles, but reported more satisfaction with social support.

**CONCLUSIONS**—Gender differences exist regarding appraisal of stress, coping styles, and coping resources long-term after heart transplantation. These differences may guide tailoring therapy regarding stress, poor coping, and lack of resources.

### Keywords

stress and coping; heart transplant; gender

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## INTRODUCTION

Long-term survival after heart transplantation is similar for men and women.<sup>1</sup> Other outcomes after heart transplantation differ by gender. We have previously reported that being female was related to worse functional ability both early and later after heart transplantation<sup>2,3</sup> and depression later after heart transplantation.<sup>4</sup> Gender was not related to work status<sup>5</sup> or overall satisfaction with health-related quality of life, although being female was related to less satisfaction with health and functioning long-term after transplant.<sup>6</sup> Given differences in these outcomes by gender long-term after heart transplantation, we questioned whether factors that may influence these outcomes might also differ by gender.

Previous incidental findings of gender differences regarding stress, symptom distress, coping, and adherence<sup>7,8,9</sup> supported our undertaking of a more rigorous examination of the influence of gender on these factors. Importantly, stress, symptom distress, coping, support, and adherence have been related to outcomes (e.g., quality of life, depression, physical function, and survival) after heart transplantation.<sup>1,3,4,6,10</sup> Understanding whether gender affects stress related to heart transplantation, appraisal of transplant-related stress, and coping may provide guidance in tailoring long-term care after heart transplantation which may improve outcomes.

Using the Lazarus and Folkman Stress, Appraisal, and Coping framework,<sup>11</sup> we examined whether gender differences exist regarding stress (e.g., stress related to illness and treatment [i.e., heart transplantation]), appraisal of stress (e.g., symptom distress, and perceived difficulty adhering to the heart transplant treatment regimen), and coping (e.g., use and effectiveness of coping styles, perceived adherence to the heart transplant regimen, and coping resources [e.g., social support]) at 5 years after heart transplantation, which are related to outcomes (Figure 1). This framework has appropriately guided our research, as per Lazarus and Folkman, outcomes, including quality of life, are affected by the stresses of living, evaluation of those stresses, and how individuals cope.<sup>11</sup> We hypothesized that female heart transplant recipients would report more overall stress and symptom distress, use of more negative coping styles, more difficulty adhering to the transplant medical regimen, less adherence to the transplant regimen, and less satisfaction with social support than men.

## METHODS

### Design

The study used a prospective, multi-site, longitudinal, observational design.

### Sample

Patients for this secondary analysis were from a study of HRQOL outcomes at 5 – 10 years after heart transplantation. The study cohort was from a pool of 1,437 adult patients who were transplanted between July 1, 1990 and June 30, 1999 at 4 medical centers in the United States. Inclusion criteria were age  $\geq$  21 years, ability to read and write English, and physically able to participate. Of the 1,437 patients, 884 were potentially eligible to enroll in our study, 597 were enrolled, and 555 patients completed one or more booklets of questionnaires over time. Two hundred eighty-seven patients were not enrolled, and 127 of them chose not to participate. Reasons for non-enrollment are documented elsewhere.<sup>6</sup> When consented patients (n=597) were compared to patients who did not consent (n=127), there were no statistically significant differences between groups for gender.<sup>6</sup> Two hundred ten of the 555 patients were 5 years post heart transplantation and comprise the sample for this report (figure 2). Of the 210 patients, 42 (20%) were women, which is similar to the 22% of women who underwent heart transplantation during the era of data collection for our report, per the International Society of Heart and Lung Transplantation Registry.<sup>12</sup>

### Instruments

Self-report questionnaires used for this report were completed by paper and pencil at 5 years after heart transplantation. Questionnaires, aligned with our Stress and Coping Framework, measured the following constructs: (1) frequency and intensity of stress (Heart Transplant Stressor Scale<sup>13</sup>); (2) appraisal of stress (symptom frequency and distress (Heart Transplant Symptom Checklist<sup>14</sup>) and perceived difficulty with adherence to the medical regimen (Assessment of Problems with the Heart Transplant Regimen<sup>15</sup>); and (3) coping and coping resources (coping use and perceived effectiveness (Jalowiec Coping Scale<sup>16</sup>), perceived actual adherence with the medical regimen (Assessment of Problems with the Heart Transplant Regimen<sup>15</sup>), and satisfaction with social support (Social Support Index<sup>17</sup>).

**Stress**—We defined stress as “a relationship between the person and the environment that is appraised by the person as taxing or exceeding his or her resources and endangering his or her well-being”.<sup>11</sup> The Heart Transplant Stressor Scale includes 81 items that measure stress related to physical, psychological, self-care, family, work/school/financial, and hospital/clinic dimensions. (0=not stressful at all to 3=very stressful).<sup>13</sup> Psychometric support is adequate for this instrument. Homogeneity reliability is supported (Cronbach’s alpha coefficients = 0.95 for the entire scale and 0.78–0.90 for the dimensions).<sup>13</sup> Validity was also acceptable.<sup>13</sup>

**Appraisal of stress**—Evaluation of stress included measures of symptom distress and difficulty adhering to the medical regimen. Symptom distress occurs in response to the perception of an abnormal physical, emotional, or cognitive state (e.g., cramps in feet, mood swings, and problems with memory).<sup>18</sup> The Heart Transplant Symptom Checklist has 89

items measuring symptom frequency (yes/no) and symptom distress (0=not bothered at all to 3=very bothered for symptom distress).<sup>14</sup> There are six subscales: cardiopulmonary, gastrointestinal, neuromuscular, genitourinary, dermatologic, and psychological). Cronbach's alpha was acceptable (0.95 for the entire scale and range = 0.91 – 0.68 for five of the six subscales). The genitourinary subscale had a Chronbach's alpha of only 0.46 which we potentially attribute to fewer items on the subscale.<sup>14</sup> Content and construct validity were supported.<sup>14</sup>

Adherence is “the extent to which a person's behavior, following the medical regimen, corresponds with the agreed recommendations of a healthcare provider”.<sup>19</sup> Perceived difficulty with adherence to the transplant regimen is measured by the Assessment of Problems with the Heart Transplant Regimen instrument. Part A measures difficulty (1=no difficulty to 4=a lot of difficulty) regarding taking medications, lifestyle changes, and appointment keeping.<sup>15</sup> Test-retest reliability was acceptable for this instrument.<sup>15</sup> This instrument did not meet requirements for internal consistency reliability. Given the multiple distinct items in the instrument, related to the post transplant medical regimen, we did not expect this instrument to have adequate internal consistency reliability. Validity was acceptable, with support for both content and concurrent validity.<sup>15</sup>

**Coping and coping resources**—Coping is defined as “constantly changing cognitive and behavioral efforts to manage specific external and/or internal demands that are appraised as taxing or exceeding the resources of the person”.<sup>11</sup> Coping use and effectiveness are measured by the 60-item Jalowiec Coping Scale (0=never used to 3=often used and 0=not helpful to 3=very helpful) which has 8 subscales (confrontive, evasive, optimistic, fatalistic, emotive, palliative, supportant and self-reliant).<sup>16</sup> Two summary scores were calculated from subscale scores: (1) positive coping styles (confrontive, optimistic, self-reliant and supportant subscales) and (2) negative coping styles (fatalistic, evasive, and emotive subscales). This widely used instrument has alpha reliability coefficients of 0.75–0.86 for the total and subscale scores and acceptable validity.<sup>16</sup>

Perceived adherence to the transplant regimen, a behavioral effort to manage the demands of heart transplantation, is also measured by the Assessment of Problems with the Heart Transplant Regimen instrument. Part B measures actual adherence (1=all of the time to 4=hardly ever) for taking medications, lifestyle changes, and appointment keeping.<sup>15</sup>

Social support (a coping resource) is the degree of perceived satisfaction with support provided by others; thus, the individual appraises whether support is acceptable or satisfactory at times of need.<sup>20, 21</sup> The Social Support Index has 15 items that include both tangible and emotional support. Patients respond to their level of satisfaction with support on a scale from 1=very satisfied to 4=very dissatisfied.<sup>17</sup> Cronbach's alpha reliability was supported for the total scale (0.84) and subscales (tangible support=0.78 and emotional support=0.69).<sup>17</sup> Validity for this instrument was acceptable, as well.<sup>17</sup>

## Medical records

Medical records data at 5 years after transplant were used for this report as well as data available to us from the Cardiac Transplant Research Database (CTRD), a large registry

located at the University of Alabama, Birmingham. Clinical data from medical records and the CTRD included medical conditions diagnosed per cardiologist and associated tests (e.g., acute rejection, infection, renal disease, and endocrine disorders [e.g., diabetes mellitus]) and psychiatric conditions diagnosed per psychologist / psychiatrist (e.g., depression and anxiety). The co-principal investigator, a heart transplant clinician, reviewed small samples of medical records data from all sites to ensure accuracy of chart review.

## Procedures

Sites received Institutional Review Board approval prior to participation in our larger HRQOL study and the CTRD. Patients were consented either face-to-face at a clinic visit or while hospitalized or via mail. Patients who provided written informed consent completed booklets of questionnaires every 6 months from 5 to 10 years after heart transplantation. Research coordinators collected medical records data. Data used for this report were from patient completed self-report questionnaires and medical records at approximately 5 years (mean=4.98±0.17 years, median=4.99 years) after transplant, which for the cohort of patients in this study was from 7/1/95 – 6/30/99. Data were screened and cleaned and then sent to our Data Coordinating Center at the University of Alabama, Birmingham for data entry and analyses.

## Statistical Analyses

Continuously distributed measurements are summarized using means and standard deviations and compared using two-sample t-tests. Categorical variables are shown as counts/percentages and group comparisons are based on the Chi-square or Fisher's exact test.

To assess the hypothesized gender differences, multivariable models were created based on a simplified version of *bagging*,<sup>22</sup> a widely employed machine learning ensemble method. This approach alleviates some of the most common shortcomings of automatic model building methods, such as instability, sensitivity to outliers or lack of reproducibility. For every factor (statistical outcome), a two-stage process was created: first, the original dataset was bootstrap resampled (with replacement) 1,000 times. Within each bootstrap sample, stepwise variable selection with 0.05 entry/exit criteria occurred, and gender was included, by default, in every model. The following predictors were entered into the model: gender, age, race group (white, non-white), marital status (married, non-married), heart disease etiology (idiopathic, ischemic, other), education ( <12 years (high school), >12 years), diabetes, orthopedic illness, gout, renal disease, cardiovascular disease, co-existing oncologic illness, gastrointestinal disease, co-existing psychological illness and coronary artery disease.

In stage two, the short list of predictors significant at a two-sided 5% level in at least 200 of the 1,000 resamples (20%) was compiled. Only these predictors were included in the final regression model created based solely on the *original* dataset. The 20% threshold choice is intended to produce a parsimonious and stable final model. Currently, there are no metric-driven guidelines for the choice of such thresholds, yet common choices in practice may range between 20% and 50%. All analyses were performed using SAS v9.3. Throughout,

statistical significance was established at the two-sided 5% alpha level, and no adjustments for multiplicity were made.

## RESULTS

Differences by gender were detected for demographic and clinical characteristics at 5 years after heart transplantation (Table 1). More female patients (n=42) were significantly younger, non-white, and not married than male patients (n=168). More women than men had dilated cardiomyopathy as the etiology of heart failure. Additionally, women had less concomitant renal disease, as compared to men.

### Appraisal of Heart Transplant-related Stress

Appraisal of heart transplant-related stress is presented in Table 2. Regarding overall stress, as compared to men, women had borderline higher overall intensity of stress by 0.02 points on average. In addition, patients without a psychiatric condition (e.g., depression and anxiety) or diabetes experienced a lower overall intensity of stress.

Women reported significantly more difficulty adhering to the heart transplant regimen than men by an average of 0.05 points (Table 2). Additionally, younger patients indicated that they had more difficulty adhering to the heart transplant regimen than older patients. No differences were detected between groups for symptom distress (data not shown).

### Coping Styles and Resources

In terms of coping styles (Table 3), women demonstrated significantly higher scores on the Overall Negative Use subscale than men (mean difference 0.1 points). In addition, women displayed a trend for significantly higher scores on the Overall Positive Use subscale as compared to men, with a mean difference of 0.08 points. Participants with no documented psychiatric condition had lower scores on the Overall Negative use subscale.

Self-report of actual adherence to the heart transplant regimen was significantly higher among women, as compared to men (Table 3), by 0.05 points on average. Similar to difficulty with adherence, younger patients also reported lower adherence than older patients.

Report of overall satisfaction with support (Table 4) was significantly higher among women (mean difference 0.04 points) than men. In addition, women also reported significantly higher levels of satisfaction with support, both on the emotional scale (mean difference 0.09), as well as for tangible support (mean difference 0.06).

## DISCUSSION

We conclude that appraisal of stress related to heart transplantation, coping styles, and coping resources differ between men and women at 5 years after heart transplantation. Our hypotheses were partially supported. Regarding appraisal of heart transplant-related stress, women did not report more overall stress (although there was a strong trend) or symptom distress, but did report more difficulty adhering to the transplant regimen than men. For





with multiple areas of the post transplant regimen, but did find that compliance in most areas worsened over time. A later report by Dew et al.<sup>31</sup> on adherence during the first two years after lung transplantation described better self-care adherence by men than women. Finally, a report from a consensus conference on nonadherence noted that a meta-analysis of risk factors for poor adherence to the medical regimen in other chronic illness populations has typically revealed females being more adherent than males, which is consistent with our findings.<sup>32, 33</sup>

Findings regarding differences by gender to adherence with components of a prescribed medical regimen are somewhat contradictory. Additional research is needed. Yet, our report and other articles in the literature raise important concerns regarding differences in adherence to the medical regimen by demographic characteristics and the inference that time since transplant may influence adherence. Thus, careful monitoring and intervention, as needed, are recommended across time after heart transplantation.

Additionally, per our Stress, Appraisal, and Coping framework, difficulty with adherence to the medical regimen reflects appraisal of stress, which is distinct from actual adherence. While reports in the literature focus primarily on actual adherence, studying difficulty with adherence, as well as actual adherence, can highlight components of the medical regimen that are appraised as taxing and provide clinicians with an opportunity to assist patients to overcome these difficulties. For example, Zikmund-Fisher et al.<sup>34</sup> reported a significant correlation between having problems with taking anti-hypertensive medications and willingness to consider intensification of blood pressure medications in response to an elevated blood pressure. They concluded that paying attention to medication related “issues” can enhance adherence to the medication regimen and downstream outcomes.

More frequent use of negative coping styles as reported by women, as compared to men, long-term after heart transplantation was previously described by our team, using two sample t-tests.<sup>7</sup> Our current examination of coping use by gender, using more rigorous statistical methods, supports our earlier findings. This finding may have important clinical implications after transplant. For example, more use of fatalistic coping and high avoidance coping has been found to be related to poor adherence early after heart transplantation.<sup>15, 30</sup> Our finding that women indicate that they use more negative coping styles than men long term after heart transplantation reinforces the ongoing need for psychosocial monitoring. Whether or not coping styles can be changed is unclear.<sup>35</sup>

Contrary to our hypotheses, women reported being more satisfied with overall social support and experienced higher levels of support (tangible and emotional support) later after transplant than men. Support is an important determinant of outcomes after thoracic organ transplantation. More support is related to better health-related quality of life,<sup>36</sup> while poor family caregiver support is related to an increased risk of depression and psychological distress.<sup>37, 38</sup> Poor support has also been found to be related to immunosuppressant nonadherence after solid organ transplantation.<sup>39</sup> Reports of relationships of social support by gender with outcomes (e.g., survival and health-related quality of life) have also been reported in other populations of chronic illness patients, including those with diabetes,<sup>40</sup> heart failure,<sup>41</sup> and end-stage renal disease.<sup>42</sup> Monitoring satisfaction with support and



actual support for both men and women long-term after transplant may guide development of interventions for those at risk for poor support and potentially poor outcomes.

Our study was limited by survivorship bias. However, our inclusion of four sites across the U.S. enhanced representativeness of our target population. Also, self-report was the only measure of adherence, a measurement method which tends to overestimate actual adherence, and we only examined overall adherence and difficulty with adherence, rather than adherence to components of the medical regimen. Lastly, while these data are old, heart transplant clinical practice has not changed much over time. Immunosuppression today is similar to medications used when our data were collected, except the use of tacrolimus and mycophenolate mofetil has increased, while the use of cyclosporine and azathioprine has decreased. Also, more ventricular assist devices are being implanted as a bridge to heart transplantation. Since our data were collected long-term after transplant, increased use of these devices should not affect our findings. Finally, patients must still adapt to having received a heart transplant and care for themselves long-term after surgery.

## CONCLUSIONS

We conclude that men and women differ on appraisal of stress related to heart transplantation, coping styles, and coping resources long-term after heart transplantation. These differences may guide monitoring and tailoring therapy when stress, symptom burden, poor coping, difficulty with adherence to the medical regimen, and lack of resources are identified later after heart transplantation. We are currently conducting a pilot trial to test an intervention focused on enhancing adherence and maximizing support for an especially vulnerable, high-risk cohort of young adults who receive heart transplantation as children and transition to adult care. Still more research is needed to test interventions (e.g., psychological interventions [related to high stress and poor coping], and behavioral and support-focused interventions [related to adherence with the medical regimen and support] in vulnerable subgroups of patients, after heart transplantation, other solid organ transplantation, and other chronic illnesses (e.g., congenital heart disease).

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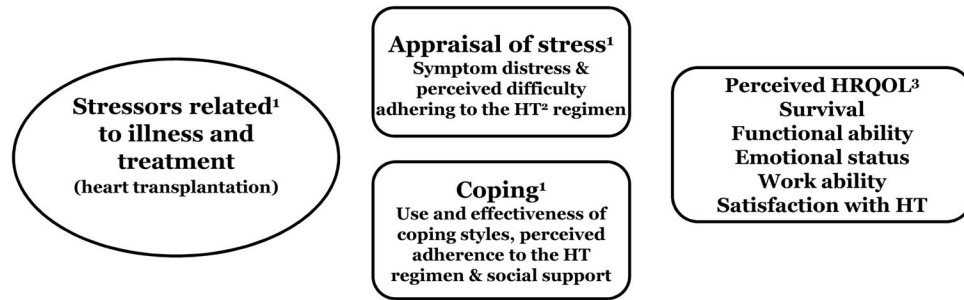
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### Stress and Coping Framework

Antecedent variables → Mediating variables → Outcome variables



<sup>1</sup>Lazarus RS, Folkman S. *Stress, appraisal, and coping*. New York, NY: Springer; 1984.

<sup>1</sup>=variables examined in this report  
<sup>2</sup>HT=heart transplantation  
<sup>3</sup>HRQOL=health-related quality of life

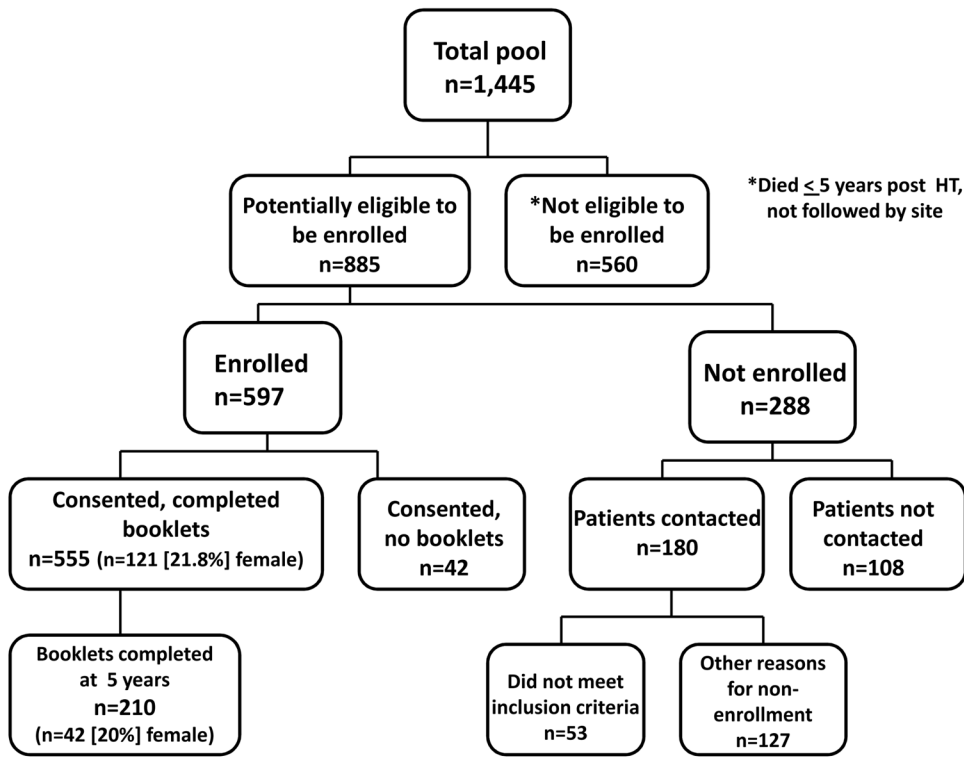
**Figure 1.** Stress and coping framework for patients who undergo heart transplantation

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**Figure 2.**  
Flow chart of study sample

**Table 1**

Comparison of Demographic and Pre- and Post-operative Characteristics by Gender

Variable	Women (N=42)	Men (N=169)	P-value
<b><u>Demographic characteristics</u></b>			
Age of recipient at transplant (years)	48.8 ± 12.0	57.1 ± 8.5	<.001
Race			0.008
White	34 (81%)	160 (95%)	
Black	8 (19%)	7 (4%)	
Asian	0 (0%)	2 (1%)	
Marital Status			<.001
Divorced/separated	9 (21%)	17 (10%)	
Married	24 (57%)	139 (82%)	
Single	7 (17%)	5 (3%)	
Widowed	2 (5%)	8 (5%)	
Education			0.85
High school degree or less	16 (38%)	67 (40%)	
College degree or more	26 (62%)	102 (60%)	
<b><u>Pre-operative characteristics</u></b>			
Heart Disease Etiology			<.001
Idiopathic	24 (57%)	38 (22%)	
Ischemic	9 (21%)	116 (69%)	
Other	9 (21%)	15 (9%)	
<b><u>Post-operative characteristics</u></b>			
Co-morbid conditions			
Diabetes	8 (20%)	57 (34%)	0.07
Orthopedic Illness	7 (17%)	36 (22%)	0.53
Gout	5 (12%)	32 (19%)	0.30
Cardiovascular Disease	8 (20%)	31 (19%)	0.89
Oncologic Illness	7 (17%)	40 (24%)	0.35
GI Illness	8 (20%)	39 (23%)	0.60
Renal Illness	5 (12%)	74 (44%)	<.001
Psychiatric Condition	14 (34%)	38 (23%)	0.13
Coronary Artery Disease	10 (24%)	65 (38%)	0.08
Coronary Heart Failure	0 (.%)	2 (100%)	.
Hypertension	30 (73%)	143 (86%)	0.06
Hyperlipidemia	29 (71%)	125 (75%)	0.59
Pulmonary Disease	6 (15%)	16 (10%)	0.35
Infection Co-Existing Illness	3 (7%)	14 (8%)	0.82
Total number of rejection episodes	2.1 ± 1.9	1.6 ± 1.6	0.06
Total number of infection episodes	0.6 ± 1.0	0.8 ± 1.3	0.24
At least on rejection episode	35 (83%)	119 (70%)	0.09
At least one infection episode	14 (33%)	78 (46%)	0.13

<b>Variable</b>	<b>Women (N=42)</b>	<b>Men (N=169)</b>	<b>P-value</b>
New York Heart Functional Class			0.45
I	23 (55%)	111 (65%)	
II	17 (40%)	54 (32%)	
III	0 (0%)	1 (1%)	
missing	2 (5%)	3 (2%)	

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**Table 2**

Multivariable Models for Appraisal of Heart Transplant-related Stress. Dependent Variables (Outcomes) are “Difficulty with Adherence to the HT Regimen” and “Intensity of Overall Stress”. Independent Variables are Listed for Each Outcome and No Additional Variables Were Controlled For.

Factor	Variable	Estimate	95% CI	P-value
<b>Difficulty with Adherence to the HT Regimen</b>				
	Women	0.05	(0.01, 0.08)	0.01
	Age (years)	-0.001	(-0.003, 0.001)	0.035
	No Renal Disease	0.03	(0.001, 0.05)	0.048
	Idiopathic Etiology	-0.005	(-0.05, 0.04)	0.80
	Ischemic Etiology	0.02	(-0.02, 0.06)	0.27
	Non-White	-0.03	(-0.08, 0.01)	0.14
	High School Degree or less	-0.02	(-0.04, 0.01)	0.23
<b>Intensity of Overall Stress</b>				
	Women	0.02	(0.001, 0.05)	0.052
	Age	-0.001	(-0.002, 0.001)	0.16
	No Psychiatric Condition	-0.05	(-0.07, -0.03)	<.0001
	No Diabetes	-0.02	(-0.04, -0.001)	0.036
	No CAD	-0.01	(-0.03, 0.01)	0.14

**Table 3**

Multivariable Models for Use of Coping Styles. Dependent Variables (Outcomes) are “Overall Positive Use of Coping Styles”, “Overall Negative Use of Coping Styles” and “Actual Adherence to the Heart Transplant (HT) Regimen”. Independent Variables are Listed for Each Outcome and No Additional Variables Were Controlled For.

Factor	Variable	Estimate	95% CI	P-value
<b>Overall Positive Use of Coping Styles</b>				
	Women	0.08	(-0.003, 0.17)	0.06
	No Cardiovascular Illness	-0.07	(-0.15, 0.02)	0.13
	No Oncologic Illness	-0.06	(-0.14, 0.03)	0.17
<b>Overall Negative Use of Coping Styles</b>				
	Women	0.10	(0.03, 0.16)	0.005
	Age (years)	-0.002	(-0.005, 0.001)	0.13
	No Psychiatric Condition	-0.10	(-0.15, -0.04)	0.001
	Idiopathic Etiology	-0.002	(-0.08, 0.08)	0.96
	Ischemic Etiology	0.08	(-0.005, 0.16)	0.07
	No Cardiovascular Illness	-0.04	(-0.10, 0.02)	0.19
<b>Actual Adherence to the HT Regimen</b>				
	Women	0.05	(0.02, 0.08)	0.005
	Age (years)	-0.003	(-0.004, -0.001)	<.0001
	Idiopathic Etiology	0.05	(0.01, 0.09)	0.028
	Ischemic Etiology	0.06	(0.02, 0.11)	0.003
	No Psychiatric Condition	-0.02	(-0.05, 0.004)	0.09
	No Orthopedic Illness	-0.02	(-0.05, 0.01)	0.22
	No Diabetes	-0.02	(-0.04, 0.01)	0.26

**Table 4**

Multivariable Models for Coping Resources. Dependent Variables (Outcomes) are “Overall Satisfaction with Social Support”, “Satisfaction with Emotional Support” and “Satisfaction with Tangible Support”. Independent Variables are Listed for Each Outcome and No Additional Variables Were Controlled For.

Factor	Variable	Estimate	95% CI	P-value
<b>Overall Satisfaction with Social Support</b>				
	Women	0.04	(0.004, 0.08)	0.032
	Married	-0.02	(-0.06, 0.02)	0.26
<b>Satisfaction with Emotional Support</b>				
	Women	0.09	(0.03, 0.14)	0.004
	Idiopathic Etiology	-0.07	(-0.15, 0.004)	0.06
	Ischemic Etiology	-0.04	(-0.12, 0.03)	0.24
<b>Satisfaction with Tangible Support</b>				
	Women	0.06	(0.01, 0.10)	0.008
	No Oncologic Illness	0.03	(-0.01, 0.06)	0.17
	Idiopathic Etiology	0.003	(-0.05, 0.06)	0.93
	Ischemic Etiology	0.03	(-0.02, 0.08)	0.26
	No Cardiovascular Illness	0.02	(-0.02, 0.06)	0.26
	High School Degree or less	-0.02	(-0.05, 0.02)	0.35