Analysis of deaths in patients awaiting heart transplantation: impact on patient selection criteria

Guy A Haywood, Peter R Rickenbacher, Pedro T Trindade, Lars Gullesstad, Joseph P Jiang, John S Schroeder, Randall Vagelos, Philip Oyer, Michael B Fowler

Abstract

Objective—To analyse the clinical characteristics of patients who died on the Stanford heart transplant waiting list and to develop a method for risk stratifying status 2 patients (outpatients).

Methods—Data were reviewed from all patients over 18 years, excluding retransplants, who were accepted for heart transplantation over an eight year period from 1986 to 1994.

Results—548 patients were accepted for heart transplantation; 53 died on the waiting list, and 52 survived on the waiting list for over one year. On multivariate analysis only peak oxygen consumption (peak VO₂; 11.7 (SD 2.7) vs 15.1 (5.2) ml/kg/min, P = 0.02) and cardiac output (3.97 (1.03) vs 4.79 (1.06) litres/min, P = 0.04) were found to be independent prognostic risk factors. Peak VO₂ and cardiac index (CI) were then analysed in the last 141 consecutive patients accepted for cardiac transplantation. All deaths and 88% of the deteriorations to status 1 on the waiting list occurred in patients with either a CI < 2.0 or a VO₂ < 12. In those with a CI < 2.0 and a VO₂ < 12, 38% died or deteriorated to status 1 in the first year on the waiting list. Patients with CI ≥ 2.0 and a VO₂ ≥ 12 all survived throughout follow up. Using a Cox’s proportional hazards model with CI and peak VO₂ as covariates, tables were constructed predicting the chance of surviving for (a) 60 days and (b) 1 year on the waiting list.

Conclusions—These data provide a basis for risk stratification of status 2 patients on the heart transplant waiting list.

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Keywords: heart failure; prognosis; heart transplantation

Waiting lists for heart transplants are increasing in centres throughout the USA and in many other countries,1,2 the average length of time patients wait before undergoing transplantation is increasing, and the ratio of patients dying on the waiting list to patients being transplanted has risen from 0.07 in 1983 to 0.21 in 1989.3 Within the waiting list population the group that suffers most from the worsening imbalance between the demand for transplants and the supply of donor organs is the patients in status 2 (patients not requiring inotropic or mechanical circulatory support) who in most centres are transplanted in the chronological order in which they are accepted on to the waiting list. Patients who present in status 1 (requiring inotropic or mechanical circulatory support) or who deteriorate from status 2 to status 1, are transplanted ahead of status 2 patients because of their high mortality if they remain on the waiting list on circulatory support. Selecting status 1 patients for urgent transplantation has been shown to be an appropriate use of donor organs in terms of the impact on overall mortality,4 but has resulted in a high percentage of donor organs that become available going to these severely ill patients, with fewer organs remaining for implantation into status 2 patients.

In this study we examined the mortality in all patients on the waiting list for heart transplant over an eight year period in order to define the length of time that patients who died had been waiting and to determine how many of the deaths occurred in patients who had progressed from status 2 to status 1. We also compared two groups from within the status 2 patients on the waiting list who were at opposite ends of the spectrum for prognostic risk: those who died on the waiting list; and those who survived for more than one year on the waiting list without adverse outcome. By using this approach it was possible to identify clinical variables that were independent prognostic indicators and to propose a strategy to improve our ability to target the available donor organs for transplant not only to status 1 patients, but also to status 2 patients with a high risk of death on the waiting list. The policy suggested has the potential to decrease overall mortality in patients referred for heart transplant and to avoid performing heart transplants on patients who are unlikely to gain survival benefit from the procedure.

Methods

The study was a retrospective review of the clinical records of patients aged 18 years or older who were placed on the Stanford University Hospital heart transplant waiting list between 1 June 1986 and 31 May 1994. Patients who were accepted for repeat heart transplants were excluded from the study. Data drawn from the pretransplant evaluation database were supplemented from the clinical files which are kept for each patient referred for heart transplantation. In instances where
the clinical variables to be assessed were incomplete in these files, the inpatient hospital medical records were reviewed.

In the first phase of the study data were used to define the two patient groups: status 2 patients who died on the waiting list and status 2 patients who survived for more than one year on the waiting list without adverse outcome. Accepted outcomes in the latter group were: heart transplantation; removal from the waiting list because of clinical improvement; or continued listing for transplantation at the end of the study period without deterioration to status 1. This analysis was performed in February 1994 and the two subgroups were taken from patients accepted onto the waiting list between 1 June 1986 and 31 December 1993. Data collected from these two subgroups are shown in table 1. These data was subjected to univariate and multivariate analysis.

To devise a formula enabling the prognostic indicators identified by this technique to be used in risk stratification of status 2 patients, we then performed a further analysis on all patients in status 2 at entry who were accepted onto the waiting list between 1 January 1991 and 31 May 1994. (Thirty six of the patients in this cohort of 114 consecutive patients were also used as part of the first phase subgroup analysis.) This time period was used to ensure that the formula would be based on patients who had received modern medical treatment for heart failure. Statistical analyses using Kaplan Meier survival curves and a Cox’s proportional hazards model were performed.

### Table 1 Data collected for univariate and multivariate analysis

<table>
<thead>
<tr>
<th>Variable</th>
<th>Status 2 patients</th>
<th>Status 1 patients</th>
</tr>
</thead>
<tbody>
<tr>
<td>Age</td>
<td></td>
<td></td>
</tr>
<tr>
<td>Diagnosis (ischaemic, dilated, valvular or other)</td>
<td></td>
<td></td>
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<tr>
<td>Resting heart rate</td>
<td></td>
<td></td>
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<tr>
<td>Systolic blood pressure</td>
<td></td>
<td></td>
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<tr>
<td>Ejection fraction</td>
<td></td>
<td></td>
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<tr>
<td>Serum sodium</td>
<td></td>
<td></td>
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<tr>
<td>Serum creatinine</td>
<td></td>
<td></td>
</tr>
<tr>
<td>ACE inhibitor therapy</td>
<td></td>
<td></td>
</tr>
<tr>
<td>Central haemodynamics:</td>
<td></td>
<td></td>
</tr>
<tr>
<td>Cardiac output</td>
<td></td>
<td></td>
</tr>
<tr>
<td>Systolic pulmonary artery pressure</td>
<td></td>
<td></td>
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<tr>
<td>Mean pulmonary artery pressure</td>
<td></td>
<td></td>
</tr>
<tr>
<td>Pulmonary capillary wedge pressure</td>
<td></td>
<td></td>
</tr>
<tr>
<td>Pulmonary vascular resistance</td>
<td></td>
<td></td>
</tr>
<tr>
<td>Exercise variables:</td>
<td></td>
<td></td>
</tr>
<tr>
<td>Percentage of maximum predicted heart rate achieved at peak exercise</td>
<td></td>
<td></td>
</tr>
<tr>
<td>Peak oxygen consumption on exercise</td>
<td></td>
<td></td>
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<tr>
<td>Arrhythmias:</td>
<td></td>
<td></td>
</tr>
<tr>
<td>History of a sustained supraventricular or ventricular arrhythmia (including atrial fibrillation, but excluding frequent ectopy)</td>
<td></td>
<td></td>
</tr>
<tr>
<td>Antiarrhythmic drug administration (other than digoxin)</td>
<td></td>
<td></td>
</tr>
</tbody>
</table>

#### STATISTICAL METHODS

All data were entered into Statview 4.01 (Abacus Concepts). Differences between the two subgroups (6/86–12/93) in which multiple clinical variables were assessed (patients who died on the list and patients who survived for over one year) were analysed using the two tailed Mann-Whitney U test for continuous variables and $\chi^2$ test for nominal variables. Significance was measured at the 5% level. Variables that were found to have significantly different mean values or rates in the two subgroups on univariate analysis were then subjected to multivariate analysis using a least squares analysis (Statview 4.01, Abacus Concepts) and a stepwise logistic regression analysis (SPSS for Windows release 5) to determine which factors were independently associated with outcome.

From the consecutive patient cohort (1/91 to 5/94) Kaplan Meier survival curves were constructed using Survival tools for Statview 1.0 (Abacus Concepts). Times of death or deterioration to status 1 were recorded as event times and removal from the waiting list because of transplantation, or any other reason, was treated as a censoring event. Dichotomisation points for the independent prognostic variables identified by multivariate analysis were used to compare patients with values above and below the selected values. Differences in survival were determined using a log rank (Mantel Cox) test. In addition, a Cox proportional hazard model was constructed using the same independent prognostic variables as covariates. Tables giving the predicted chance of surviving for (a) 60 days and (b) one year on the waiting list without death or deterioration to status 1 were calculated for a range of combinations of individual values of the independent prognostic variables using the equation:

$$\text{Chance of survival to time } t = [S_0(t)]^{exp(b' x_1 + b'' x_2)}$$

where $S_0(t)$ = the baseline survival function, $b'$ = the coefficient of hazard calculated for the first prognostic variable, $b''$ = the coefficient of hazard for the second prognostic variable, and $x_1$ and $x_2$ = differences from the baseline values for each value respectively. Values in text and table 2 are given as mean (SD)

### Results

#### DEMOGRAPHIC ANALYSIS

A total of 1114 patients was assessed in the period from June 1986 to December 1993, and from these 548 (49% of the total) were accepted onto the waiting list. During the same study period 53 died on the waiting list, giving a waiting list mortality rate of 9.7%. Mortality has been rising during this period, as shown by comparison with the figures for 1 January 1991 to 31 May 1994, during which time 23 patients (16%) died. Mean time on the waiting list for patients accepted between June 1986 and 31 December 1993 was 127 d, whereas from 1 January 1991 to 31 May 1994 the mean waiting time was 272 d.

#### Criteria for acceptance for heart transplantation

The majority of patients accepted on to the heart transplant waiting list had been referred because of severe ventricular dysfunction and recurrent or refractory symptoms of congestive heart failure. Small numbers of patients were also accepted who had severe recurrent angina pectoris or ventricular tachyarrhythmias which
were considered technically unsuitable or an unacceptable risk for other forms of treatment including surgical intervention. Patients were excluded if they were suffering from significant co-morbid conditions that were likely to limit life expectancy independently to less than five years. Other exclusions were pharmacologically irreversible pulmonary hypertension (pulmonary vascular resistance greater than three Wood units despite intravenous sodium nitroprusside), a history of severe psychiatric illness, continuing drug or substance abuse, or other severe psychosocial dysfunction. During the period January 1991 to May 1994 the mean cardiac output and peak oxygen consumption of status 2 patients on medical treatment was 2·1 (SD 0·6) litres/min/m² and 13·6 (3·8) ml/kg/min respectively. At the time these measurements were made, 86% were receiving angiotensin converting enzyme (ACE) inhibitor therapy.

Death rate
Twenty one of the 53 deaths that occurred between June 1986 and December 1993 were patients who were classified as status 1 at the time of acceptance, and 32 were patients who were status 2 when accepted on to the waiting list. The time distribution of deaths in the status 2 patients is shown in fig 1. The bulk of the deaths occurred early after acceptance onto the list: 69% of all deaths in status 2 patients occurred in the first two months after acceptance. Ten patients who were status 2 at the time of acceptance progressed to status 1 before death on the waiting list (19% of all those who died; fig 2). A high proportion of the status 2 deaths (66%) were sudden cardiac deaths.

Univariate analysis
Table 2 shows a comparison of the clinical variables measured in status 2 patients who died on the waiting list (n = 32) compared with those of status 2 patients who survived for more than one year on the waiting list without an adverse outcome (n = 52). With univariate analysis using a two tailed Mann Whitney U test there were four variables that showed a significant difference between the two groups: systolic blood pressure, cardiac output, peak oxygen consumption, and serum sodium.

Differences in other variables such as ejection fraction, pulmonary capillary wedge pressure (PCWP) and pulmonary vascular resistance (PVR) did not reach significance. The two groups were closely matched for age (51·3 (9·0) in those who died, 51·2 (9·2) in those who survived). In addition to these variables we compared: gender, aetiology of heart failure, arrhythmic history, and ACE inhibitor treatment. Patients were predominantly male in both groups (88% of those who died, 96% of those who survived, NS). Ischaemic heart disease was the commonest diagnosis in each group (63% v 65%, NS) with the next commonest aetiology being idiopathic dilated cardiomyopathy (28% v 25%, NS). There was also no significant difference in the frequency of a history of arrhythmia (59% in patients who died, v 49% in those who survived, NS) or in the proportion of patients receiving ACE inhibitor treatment (86% in patients who died, v 88% in those who survived, NS).

MULTIVARIATE ANALYSIS
Multivariate analysis of systolic blood pressure, cardiac output, peak oxygen consumption, and serum sodium revealed that only cardiac output (P = 0·023 by logistic regression analysis) and peak VO₂ (P = 0·012 by logistic regression analysis) were independently associated with adverse outcome, whether tested by a least squares or stepwise logistic regression analysis.

Because three of the status 2 patients who died while on the waiting list survived for more than one year on the list before death, data were also analysed excluding these three patients. There was no significant change in the results when these three patients were excluded.

SURVIVAL CURVE ANALYSIS
A total of 141 consecutive patients over 18 years of age who were first time candidates for heart transplantation, accepted onto the waiting list between 1 January 1991 and 31 May 1994, were subjected to survival curve analysis. These 141 patients represented 25% of the 570 assessed during this time. By the end of the period, 74 (52%) of the patients had been transplanted, 23 (16%) died on the waiting list, a further 16 (11%) deteriorated from status 2 to status 1 before transplantation, 12 (9%) were removed from the waiting list because their clinical status had improved, and 32
Table 2  Comparison of patients awaiting first time transplantation who were classified as status 2 at the time of acceptance onto the heart transplant waiting list and who subsequently died on the list (n = 32), and patients from the group who survived for a period of over one year on the waiting list without deterioration to status 1 (n = 52)  (univariate analysis). Values are means (SD).

<table>
<thead>
<tr>
<th>Waiting list deaths</th>
<th>Waiting list survivors</th>
<th>P value</th>
</tr>
</thead>
<tbody>
<tr>
<td>Age (years)</td>
<td>51.3 (9.0)</td>
<td>51.2 (9.2)</td>
</tr>
<tr>
<td>Resting heart rate (beats/min)</td>
<td>88 (15)</td>
<td>84 (16)</td>
</tr>
<tr>
<td>%Maximum heart rate*</td>
<td>73 (14)</td>
<td>80 (15)</td>
</tr>
<tr>
<td>Systolic blood pressure (mm Hg)</td>
<td>98 (16)</td>
<td>106 (14)</td>
</tr>
<tr>
<td>SPAP (mm Hg)</td>
<td>51 (13)</td>
<td>45 (13)</td>
</tr>
<tr>
<td>Mean PAH (mm Hg)</td>
<td>35 (10)</td>
<td>32 (9)</td>
</tr>
<tr>
<td>PCWPS (mm Hg)</td>
<td>23 (7)</td>
<td>22 (9)</td>
</tr>
<tr>
<td>Pulmonary vascular resistance (Wood units)</td>
<td>3.3 (2.2)</td>
<td>2.6 (2.5)</td>
</tr>
<tr>
<td>Cardiac output (litres/min)</td>
<td>3.94 (1.16)</td>
<td>4.81 (1.06)</td>
</tr>
<tr>
<td>Ejection fraction (%)</td>
<td>17 (8)</td>
<td>18 (7)</td>
</tr>
<tr>
<td>Peak Vo2 (mmol/litre)</td>
<td>11.8 (5.1)</td>
<td>15.5 (5.2)</td>
</tr>
<tr>
<td>Serum creatinine (mmol/litre)</td>
<td>122.0 (26.5)</td>
<td>114.0 (26.5)</td>
</tr>
<tr>
<td>Sodium (mmol/litre)</td>
<td>135.6 (4.5)</td>
<td>138.0 (4.9)</td>
</tr>
<tr>
<td>Antiarrhythmics (%)</td>
<td>59</td>
<td>49</td>
</tr>
</tbody>
</table>

*Percent of predicted maximum heart rate achieved.
†SPAP, systolic pulmonary artery pressure.
‡Mean pulmonary artery pressure.
§Pulmonary capillary wedge pressure.
¶Left ventricular ejection fraction.
¶Peak oxygen consumption on treadmill exercise.

(23%) were stable and still waiting for transplantation at the end of the study period.

Survival curve analysis in relation to peak Vo2 and cardiac index in patients in status 2 at first placement on the waiting list.

In constructing Kaplan Meier survival curves for patients above and below specified values of peak Vo2 or cardiac index, we chose to analyse both death and deterioration in status from status 2 to 1 as a joint end point. This approach allowed us to develop a strategy designed not only to decrease mortality on the waiting list, but also to reduce the number of patients deteriorating from status 2 to status 1 before transplantation. In addition it ensured that the main censoring event, transplantation, was not affected by non-uniform alterations in clinical status during the period of follow-up. By treating deterioration to status 1 as an end point, the only factors affecting the likelihood of undergoing transplantation were time from acceptance on the waiting list and the availability of a donor organ of the correct blood group.

Values for cardiac index and peak Vo2 at the time of acceptance were collected for all patients initially in status 2 (n = 114 patients).

Cardiac index measurements were present for 100% of the patients accepted, and peak Vo2 had been assessed at the time of acceptance in 94 (82%). Cardiac index measurements ranged from 1.15 to 3.86 (mean 2.1 (0.56)) litres/min/m2, and peak Vo2 from 5.7 to 23.9 (mean 13.6 (3.8)) ml/kg/min. There were nine patients with cardiac index greater than or equal to 3 litres/min/m2 and eight with peak Vo2 max 20 ml/kg/min. We found that status 2 patients with a CI ≥ 2 litres/min/m2 at the time of acceptance had a one year survival free from death or deterioration to status 1 of 83%. Status 2 patients with Vo2 ≥ 12 ml/kg/min had a one year survival free from death or deterioration to status 1 of 82%. In both cases survival above and below the dichotomisation points were highly significantly different (fig 3). Neither variable on its own identified all of those who died while on the waiting list. We therefore tried two strategies based on these dichotomisation points: (1) selecting a group in which cardiac index was less than 2.0 litres/ min/m2 and peak Vo2 was less than 12 ml/kg/min, and (2) selecting patients in whom cardiac index was ≥ 2.0 litres/min/m2 and peak Vo2 was ≥ 12 ml/kg/min. The intention of the first strategy was to attempt to define high risk patients to be given priority. The intention of the second strategy was to identify patients with a good prognosis in whom transplantation could be deferred, allowing resources to be concentrated on higher risk patients. The percentage of patients identified by these two different strategies is shown in table 3.

The highest risk group with cardiac index less than 2.0 litres/min/m2 and peak Vo2 less than 12 ml/kg/min had a very poor prognosis, with 38% either dying or deteriorating to status 1 during their first year on the waiting list. Although this group was small, containing only 19% of all status 2 patients on the waiting list, it contained 38% of all patients who died. Conversely, a strategy of selecting patients in

Figure 3  Cumulative survival (Kaplan-Meier curves) without death or deterioration to status 1 in patients who were classified as status 2 at the time of acceptance. (Peak Vo2 = peak oxygen consumption—results shown for the 94 patients with peak oxygen consumption measured at the time of acceptance). Values are shown in days. Percentage survival at one year is shown.

Cumulative survival without death or deterioration to status 1

Impact of cardiac index on prognosis

Cardiac index ≥ 2.0 at time of acceptance

- 83% (n = 66, 56% of all accepted)
- Number of deaths = 4

Impact of peak Vo2 on prognosis

Peak Vo2 ≥ 12.0 at time of acceptance

- 82% (n = 59, 63% of all accepted)
- Number of deaths = 4

Peak Vo2 < 12.0 at time of acceptance

- 82% (n = 35, 37% of all accepted)
- Number of deaths = 9

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### Analysis of deaths in patients awaiting heart transplantation: impact on patient selection criteria

<table>
<thead>
<tr>
<th>CI &lt; 2.0 and V̇O₂max &lt; 12</th>
<th>CI &gt; 2.0 and V̇O₂max &gt; 12</th>
</tr>
</thead>
<tbody>
<tr>
<td>(n = 18)</td>
<td>(n = 27)</td>
</tr>
</tbody>
</table>

<table>
<thead>
<tr>
<th>Percent of all accepted</th>
<th>Percent of deaths</th>
<th>Percent of determinations to category 1 or in deaths</th>
</tr>
</thead>
<tbody>
<tr>
<td>19%</td>
<td>38%</td>
<td>42%</td>
</tr>
</tbody>
</table>

### Cox’s proportional hazards model relation to peak V̇O₂ and cardiac index in status 2 patients

The coefficient of hazard for a unit increase from a baseline cardiac index of 2-0 litres/min/m² was −1-493 and the coefficient of hazard

### Table 3 Percentage of status 2 patients defined by combinations of values for cardiac index (CI) (litres/min/m²) and peak oxygen consumption (V̇O₂max: ml/kg/min) as percentages of:

1. all status 2 patients in whom measurements of both cardiac index and peak oxygen consumption were made and at the time of acceptance for heart transplantation (n = 94),
2. all status 2 patients in whom measurements of both cardiac index and peak oxygen consumption were made and at the time of acceptance for heart transplantation who died in the waiting list (n = 13),
3. all status 2 patients in whom measurements of both cardiac index and peak oxygen consumption were made and at the time of acceptance for heart transplantation who deteriorated to status 1 while on the waiting list (n = 26)

<table>
<thead>
<tr>
<th>Cardiac index</th>
<th>Change of survival for 60 days from death or deterioration to status 1</th>
</tr>
</thead>
</table>

### Table 4 Survival functions at 60 days and at one year derived from a Cox proportional hazards model for different values of cardiac index (CI) and peak oxygen consumption (peak V̇O₂)

<table>
<thead>
<tr>
<th>Peak V̇O₂</th>
<th>Cardiac index</th>
<th>Change of survival for 60 days from death or deterioration to status 1</th>
</tr>
</thead>
</table>

### Table 5 Survival relationships for patients on the waiting list for heart transplantation

<table>
<thead>
<tr>
<th>Peak V̇O₂</th>
<th>Cardiac index</th>
<th>Change of survival for 1 year from death or deterioration to status 1</th>
</tr>
</thead>
</table>

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Cumulative survival without death or deterioration to status 1 in patients who were classified as status 2 at the time of acceptance. Patients groups are stratified in relationship to combinations of measurements of cardiac index and peak oxygen consumption (peak \(V_0_2\)). Percentage survival at one and two years is shown.

Discussion

Estimates from the 24th Bethesda conference on heart transplantation were that in the USA alone 40,000 people under 65 who might benefit from a heart transplant die each year, but the number of donor hearts available per year is currently around 2000. The number of patients awaiting heart transplants in 1993 was approximately 2500, with 300 new patients being added and 150 patients being transplanted per month. Stevenson et al. have projected that within less than four years the candidate pool will reach 4000 patients, 93% of whom will be outpatients, but only the 270 critically ill candidates in hospital at any one time will have a chance of receiving a donor organ.

ANALYSIS OF DEATHS ON THE WAITING LIST

The growing imbalance between the number of donor organs available and the number of candidates has resulted in increases in waiting time, and the number of deaths on the waiting list reported to the United Network for Organ Sharing (UNOS) has been rising steadily, from 98 in the last quarter of 1987 to 207 in the last quarter of 1991. The parallel increase in waiting time and rate of death on the waiting list is also seen in the data from our centre reported in this study. The mean waiting time for all patients between 1 January 1991 and 31 May 1994 was 272 days, with a death rate over this period of 16%. Mean waiting time has not been reported by UNOS since 1989 but figures for the overall percentage ratio between patients accepted onto the waiting list in 1991 and deaths on the waiting list in the same year (the comparable figure to the rate we report) can be estimated from reported UNOS data to be 22%.

CLINICAL VARIABLES AS PROGNOSTIC INDICATORS

Reporting on the current situation regarding the use of clinical variables to risk stratify the heart transplant waiting list, Task Force 3 at the 1993 Bethesda conference observed that “a few studies have used multivariate techniques to predict survival, but no consistent objective clinical criteria have emerged”. The variable which has drawn most attention is the peak \(V_0_2\), although values of 10, 11, 12, 14 and 20 have all been proposed as useful dichotomisation points to identify high or low risk subsets, the value has varied widely. The accepted derives from a joint observation by ourselves and by Mancini et al. that patients with peak \(V_0_2\) greater than 14 ml/kg/min were a relatively good prognosis group in whom heart transplantation could be deferred. In the Task Force 3 report, a \(V_0_2\) less than 14 ml/kg/min is given as a probable indication for heart transplantation, although the report did not go so far as to include this figure in its recommendations. The use of the policy suggested by Mancini et al. of deferring transplantation in patients with \(V_0_2\) greater than 14 ml/kg/min would have excluded 33% of the status 2 patients from the waiting list (slightly less than the 39% excluded by the application of cardiac index \(> 2.0 \text{l/min/m}^2\) and peak \(V_0_2 \geq 12 \text{ml/kg/min in our study}"); however 15% of all deaths in the status 2 patients occurred in patients with \(V_0_2\) greater than 14. Furthermore these deaths all occurred within two months from the time of acceptance; the three monthly reassessment of peak \(V_0_2\) advised in conjunction with this policy would therefore not have helped these patients. Using the strategy of cardiac index \(> 2.0 \text{l/min/m}^2\) and peak \(V_0_2 \geq 12 \text{ml/kg/min}, there were no deaths during follow up in patients satisfying both criteria.

While cardiac output has been identified as an independent prognostic indicator in at least one other study, several other variables such as low serum sodium, raised pulmonary artery diastolic pressure, left ventricular end diastolic dilatation, permanent pacing, etiology, New York Heart Association class, third heart sound, pulmonary wedge pressure, and mean systemic blood pressure have also been found to be independent risk factors for death in patients either accepted or referred for heart transplantation. Some of these variables were significant univariate predictors of death in our series, but only peak \(V_0_2\) and cardiac index were indepen-
dently significant on multivariate analysis. Another study of prognosis in patients evaluated, but not necessarily accepted, for heart transplantation did not find cardiac index to be a useful prognostic indicator. However, the measurements of cardiac index analysed in that series were made after a period of intensive medical treatment guided by pulmonary flotation catheter monitoring. Thus the mean values for pulmonary capillary wedge pressure in that group, were approximately 15 mm Hg, considerably lower than the mean values of 23–24 mm Hg seen in our patients at the time of initial evaluation. Interestingly the same investigators observed that in their series, cardiac index values above or below 2.5 litres/min/m² had a significant effect on survival (P = 0.0005) in patients who were receiving hydralazine and nitrates rather than angiotensin converting enzyme inhibitors and nitrates. While the ability to respond to treatment by achieving pulmonary capillary wedge pressures below 15 mm Hg may be a further useful prognostic indicator, this type of intensive inpatient “tailoring” of treatment is expensive and a prognostic strategy that can be applied without the need to perform therapeutic manipulation guided by central haemodynamic monitoring may be of practical value in many centres.

One of the main problems in applying fixed indices as practical tools for making decisions over which patients should go on to the heart transplant waiting list has been the tendency for most studies to report the results as survival curves for groups of patients termed low or high risk. The problem facing a transplant cardiologist is to apply mean survival values for a group, which includes a wide range of different values, to a patient who may have a measured value just above or below the cut off value. Although extrapolation from the results of this study must be undertaken with caution as the patient population may not be representative of all centres and the data have not been tested prospectively, the tables giving the predicted chance of survival without death or deterioration to status 1 for patients with specific values of cardiac index and peak VO₂ may be useful in determining which patients may have a worse chance of surviving one year if transplanted than if main- tained on medical treatment. It should be emphasised that the single centre nature of the study and the limited sample size may result in the power of risk stratification achieved by the use of cardiac index and peak VO₂ varying when applied to different populations.

It has been suggested that patients who have survived on the waiting list for over nine months do not to derive survival benefit from transplantation, and it has been proposed that patients who have survived on the list for this time should be removed from the waiting list. It is clearly preferable, however, to exclude patients who will not gain survival benefit from transplantation at the time of evaluation, rather than after nine months on the list. While reduction of the current candidate pool may be facilitated by such a removal policy, it may be difficult for patients to feel confidence in a decision removing them after they have been on the waiting list for nine months. There is also a risk in centres with small waiting lists and few patients with less common blood groups that chance fluctuations in the availability of hearts of a particular blood group may result in patients with a good medium term outlook being transplanted inappropriately within the nine month period. Survival probability tables of the type presented here provide an alternative approach to the management of such patients who are borderline for placement on the heart transplant waiting list.

The anticipated reduction in status 2 patients of 39% using the cardiac index ≥ 2.0 l/min/m² and peak VO₂ ≥ 12 ml/kg/min criteria (or a similar reduction using a one year death and deterioration survival function of 0.75) also compares favourably with the age restriction to an upper limit of 55 years which it has been suggested would achieve a 30% reduction in the waiting list and would avoid excluding patients on the grounds of age who have been found to have results from transplantation comparable to those of younger patients.

**BENEFITS IN PERIOPERATIVE MORBIDITY AND MORTALITY**

In our study 14% of patients who were status 2 at the time of acceptance were transplanted after progression to status 1. If this number could be reduced, there is evidence that this would decrease perioperative morbidity and mortality. Reports of the influence of requirement for inotropic or mechanical support before cardiac transplantation on mortality following surgery show some variation between centres but overall show worse survival after transplantation in patients transplanted from status 1 than with those transplanted from status 2. One year survival in status 1 patients who have undergone transplantation has been found to be between 69% and 83%, with certain subgroups showing a particularly high postoperative mortality. It appears that the combination of the requirement for status 1 support and the presence of pulmonary hypertension is very unfavourable, with a three month actuarial survival of only 30-3%. Although UNOS data do not allow a direct comparison of status 1 and status 2 patients survival following transplantation, Bethesda task force 1 reported a one year survival of 74% for patients requiring ventilation, intra-aortic counterpulsion, or left ventricular assistance before transplantation, with all other groups having a one year survival greater than 80%.

**MANAGEMENT STRATEGIES FOR HIGH RISK STATUS 2 PATIENTS**

Reducing the size of the status 2 waiting list by avoiding accepting patients who are unlikely to gain survival benefit from transplantation will tend to reduce the time to transplantation for the higher risk status 2 patients who remain on the waiting list. However, it is unlikely that this measure alone will be sufficient to reduce the waiting time to within two months from the time of acceptance. In our patient population, the majority of status 2 patients who died on the waiting list did so within two months from the
time of acceptance for transplantation. To reduce the numbers of status 2 patients dying on the waiting list it would therefore be necessary to identify a high risk subset in which early intervention was warranted. Such a high risk subgroup could be transplanted ahead of lower risk patients on the waiting list rather than waiting in chronological order of acceptance, but this would disadvantage the remaining patients on the waiting list. An alternative approach might be to consider entering particularly high risk patients into prospective trials for new treatments, such as elective implantable left ventricular assistance devices or implantable defibrillator insertion. The value of such devices when applied to this particular population is currently unknown, but data on survival when implantable left ventricular assistance devices are used in status 1 patients are very favourable, particularly after a delay to allow organ recovery, and although such devices have had a less impressive impact when applied to patients with poor left ventricular function and recurrent arrhythmias a prospective trial is warranted. By retrospective interrogation of the devices, this would provide valuable information on the type of arrhythmias experienced in these patients.

SUMMARY

In this study, status 2 patients who were accepted onto the heart transplant waiting list with cardiac index $\geq 2.0$ litres/min/m$^2$ and peak $V_O_2 \geq 12$ ml/kg/min had a one year mortality of zero and a rate of deterioration to status 1 of 12%. Such patients may therefore have a worse prognosis following transplantation than if they continue with medical treatment alone. Conversely, status 2 patients with cardiac index $\leq 2.0$ litres/min/m$^2$ and peak $V_O_2 \leq 12$ ml/kg/min had a very poor prognosis on the waiting list. Using tables of the type constructed here from a Cox proportional hazards model, it may be possible to predict risk of death or deterioration to status 1 from measurements of cardiac index and peak $V_O_2$ in individual patients. This approach could be used to specify patients for whom it would be appropriate either to avoid transplantation and continue with medical treatment, or to consider early transplantation or entry into trials of new treatments. These criteria should be tested retrospectively on waiting list data bases in other populations and could form the basis for a prospective multicentre randomised trial designed to assess the value of the strategy described in the management of status 2 patients being listed for heart transplantation.

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