An Assessment of the Psychosocial Evaluation for Early Liver Transplantation in Patients With Acute Alcoholic Hepatitis in the Context of Alcohol Use Disorder, a Case-Control Study

Aryeh Dienstag1, Penina Dienstag2, Kanwal Mohan3, Omar Mirza4, Elizabeth Schubert5, Laura Ford5, Margot Edelman5, Gene Im5 and Akhil Shenoy6

1Department of Psychiatry, Hadassah Hebrew University School of Medicine, Jerusalem, Israel. 2Department of Anesthesia, Hadassah Hebrew University School of Medicine, Jerusalem, Israel. 3Department of Psychiatry, University of Calgary, Calgary, Canada. 4Department of Psychiatry, Harlem Hospital Center, New York, NY, USA. 5Recanati-Miller Transplantation Institute, Mount Sinai Hospital, New York, NY, USA. 6Department of Psychiatry, Columbia University Medical Center, New York, NY, USA.

ABSTRACT

BACKGROUND: Severe acute alcoholic hepatitis (AAH) has an extremely poor prognosis with a high short term mortality rate. As a result, many centers, including our own, have allowed transplant patients to be listed for transplantation prior to achieving 6-months of sobriety. Several scoring systems, designed to target patients with a minimal period of sobriety, have been proposed to identify patients with alcohol use disorder (AUD), who would be predisposed to relapse after liver transplantation. We investigated whether these scoring systems corroborated the results of the non-structured selection criteria used by our center regarding decision to list for transplant.

METHODS: We conducted a retrospective case-control study of 11 patients who underwent early liver transplantation for AAH matched with 11 controls who were declined secondary to low insight into AUD. Blinded raters confirmed the severity of the diagnosis of DSM-5 and scored the patients on a variety of structured psychometric scales used to predict alcohol relapse. These included the High Risk for Alcohol Relapse Scale (HRAR), Stanford Integrated Psychosocial Assessment Tool (SIPAT), Alcohol Relapse Risk Assessment (ARRA), Hopkins Psychosocial Scale (HPSS), Michigan Alcoholism Prognosis Score (MAPS), Alcohol Use Disorders Identification Test - Consumption (AUDIT-C), and Sustained Alcohol Use Post-Liver Transplant (SALT) scales. All patients who underwent transplantation were followed for harmful and non-harmful drinking until the end of the study period.

RESULTS: The transplant recipients had significantly favorable MAPS, HRAR, SIPAT, ARRA, and HPSS scores with cutoffs that matched their previous research. The SALT and AUDIT-C scores were not predictive of our selection of patients for transplantation. Despite an expedited evaluation and no significant period of sobriety, our case cohort had a 30% relapse to harmful drinking after an average of 6.6 years (5-8.5 years) of follow-up.

DISCUSSION: Despite the rapid assessment and the short to no period of sobriety, the patient cohort demonstrated a 30% relapse to harmful drinking, consistent with the 20% to 30% relapse to drinking rate reported after liver transplantation for all forms of alcoholic liver disease. Average scores from maps, HRAR, SIPAT, ARRA, and HPSS corroborated our current stratification procedures, with lower mean risk scores found in the transplanted group.

CONCLUSION: Patients with AUD and severe AAH who obtain new insight into their disease and possess other favorable psychosocial factors have low rates of AUD relapse post-liver-transplantation. The psychosocial selection criteria for patients with alcoholic hepatitis in our institution are consistent with 4 of the 5 scoring systems investigated in their prediction of sobriety post-transplant.

KEYWORDS: Alcoholism, alcohol use disorder, psychosomatic medicine, alcoholic liver disease, acute alcoholic hepatitis, predict alcohol relapse, traditional psychosocial selection criteria

Introduction

Risk factors associated with alcohol relapse after liver transplantation have been extensively studied.1-4 Length of sobriety has been shown to be a significant protective factor in sustained abstinence, therefore, historically, patients who require liver transplantation for alcohol-associated liver disease (ALD) have been asked to complete 6 months of sobriety before being considered for liver transplantation.5 Patients suffering from severe acute alcoholic hepatitis (AAH) unresponsive to medical therapy have a mortality rate of over 70% by 6 months.6
In this subset of patients, the 6 months sobriety rule is there for a virtual death sentence.

In highly selected patients, early liver transplantation for AAH has been shown to achieve excellent clinical outcomes with low impact on the donor pool.7-9 We previously studied mortality in a case-control study of patients selected for transplant after a first episode of AAH8 applying the selection criteria as originally outlined by Mathurin et al.7 The psychosocial team sought candidates with a first liver decompensating event, new information on alcohol use disorder (AUD), motivation for sobriety, and strong social support. Special emphasis was placed on having good insight in the face of new liver disease as a practical way to measure readiness for transplantation. These basic criteria have become standard protocol in further studies of patients presenting with AAH.10,11

In an attempt to predict alcohol relapse after LT for those with alcohol use disorder (AUD) and ALD multiple scoring systems have both been created12 and assessed for validation.13 Commonly used scales to assess AAH patients for appropriateness for liver transplant include the High Risk for Alcohol Relapse Scale (HRAR), Alcohol Relapse Risk Assessment (ARRA), Hopkins Psychosocial Scoring System (HPSS), Sustained Alcohol Use post-LT (SALT), Stanford Integrated Psychosocial Assessment Tool (SIPAT), Michigan Alcoholism Prognosis Scale (MAPS), and the Alcohol Use Disorder Identification Test – Concise (AUDIT-C).5,14,15

These scoring systems were each created to address disparate factors and clinical concerns. The MAPS was conceived through a conceptual review of the alcohol addiction literature; and was intended to help treatment planning in liver transplant candidates.16 The ARRA was retrospectively created using a regression analysis of 25 risk factors. Nine were found to strongly correlate with post-LT alcohol relapse; less than 6 months of sobriety was not associated with relapse rates in the multivariate model.17 The HPSS and SALT were also created at transplant centers utilizing a retrospective review of risk factors, designed for the explicit purpose of assessing risk for relapse in LT recipients with alcohol-associated hepatitis (AH).8,18 The HRAR was originally proposed to predict relapse in a non-transplant population but was later adapted by Yates et al to help listing decisions when the patient was in early remission.19 The AUDIT-C was designed as a screening tool for AUD by the World Health Organization.20 The SIPAT is a general psychosocial assessment tool for transplant recipients.21 Table 1 summarizes the factors and scores assessed by the respective assessment scales.

Multiple studies have examined the predictive value of the MAPS with mixed results.16,24,25 In one study, the HRAR was found to predict AUD relapse in liver transplant patients26 but its predictive ability in subsequent studies has been uneven. The SIPAT has been shown to predict morbidity post-transplant27 as well as AUD relapse.13 The ARRA was designed to predict relapse in a retrospective review of AUD patients after liver transplant17 but this has not been replicated. The HPSS helped to identify patients relapsed to harmful drinking after a median follow-up of 1.5 years in a cohort of 17 transplanted patients.8 In a cohort of 138 patients a score greater than 7 on the SALT scale was associated with relapse to any alcohol use post liver transplant.13 In one study the AUDIT-C was found to be predictive of excessive alcohol consumption post-liver transplant.28 In one study the AUDIT-C was found to be predictive of excessive alcohol consumption post-liver transplant.28 Table 2 summarizes prior studies that have been carried out to assess the validity of the respective assessment scales with regards to predicting outcomes post liver transplant for ALD.

In our initial study, we prospectively rated inpatient candidates as having good, developing, or poor insight as they presented to our center with severe AAH. No scales were used in selecting patients with the characteristic of good or developing insight. We hypothesize that the psychometric properties of the scales included here, will validate our process of selecting patients based on either good or developing insight about their alcohol misuse. Furthermore, we present the 6-year follow up outcomes along with the individual psychometric scores of the first transplanted patients at our center, as compared to patients not transplanted.

**Methods**

Between 1 January 2012 and 6 January 2015 the psychosocial team at the Recanati-Miller Transplantation Institute at Mount Sinai Hospital evaluated 81 AAH patients, with less than 3 months of sobriety, for early liver transplantation. Twenty-two (27%) were psychosocially cleared for expedited listing with 11 patients eventually transplanted. Both psychiatry and social work independently evaluated all potential candidates with this presentation in the hospital. In addition to the patient’s interview, the level of addiction was corroborated with family and friends. The control group (n = 11) were age, sex and year-matched patients from the cohort who were also evaluated as inpatients but declined for psychosocial reasons (n = 59). Two psychosomatic fellows (AD and KM) retrospectively reviewed the psychiatric and social work data that confirm the diagnosis of AUD DSM-5 and scored HRAR, ARRA, HPSS, and SIPAT. The assessors were blinded to the evaluation decision and transplant results. The MAPS, AUDIT-C and SALT were scored by one of the authors (AS) who had originally evaluated all patients. Scores of psychosocially accepted cases and declined controls were compared using 2-tailed r-tests with 95% confidence intervals. The sensitivity and specificity for the cutoff points used for these scoring systems was calculated. The mean psychometric scores of cases and controls were compared to patient populations in the reviewed literature.

The follow-up data of all patients who underwent AAH transplantation have been collected for harmful drinking for a minimum of 3 years. All patients were seen every 2 weeks in the first 3 months after transplantation, monthly for the next
| INSTRUMENT NAME | TARGET OF INSTRUMENT | RISK FACTORS ASSESSED | POINTS | PROPOSED INTERPRETATION OF SCORE |
|-----------------|----------------------|-----------------------|--------|----------------------------------|
| **Michigan Alcoholism Prognosis Score (MAPS)** | Prediction of relapse to alcohol use for patients undergoing liver transplant for alcoholic liver disease | Known risk factors for poor outcomes in alcohol use disorder (including): Insight Patient and family Patient only Family only Neither Prognostic indices/psychological health 1. Substitute activities, Yes/No 2. Behavioral consequences, Yes/No 3. Hope/Self-esteem, Yes/No 4. Rehab relationship, Yes/No Social stability/Isolation 1. Steady job 2. Stable residence 3. Does not live alone 4. Stable marriage | 4 3 2 1 3/1 3/1 3/1 1 3/1 3/1 3/1 1 1 1 1 | Total score range: 5-20 Higher score indicates reduced risk for relapse |
| **High-Risk Alcoholism Relapse Scale (HRAR)** | Prediction of relapse and time to relapse for patients suffering from alcohol use disorder Not specific for organ transplant | Known risk factor for relapse in alcohol use disorder (including): Duration of heavy drinking (y) < 11 11-25 > 25 Usual number of daily drinks < 9 9-17 > 17 Number of prior alcoholism inpatient treatment experiences 0-1 | 0 1 2 1 2 1 2 0 1 2 | Total score range: 0-6 Low alcoholism risk > 4 High alcoholism risk |
| **Alcohol Use Disorders Identification Test -Consumption (AUDIT-C)** | Screening test to identify patients who are hazardous drinkers or have active alcohol use disorders based on previously validated tools used to screen for problematic alcohol use. Not specific for organ transplant or patients who had reduced or attempted to reduce alcohol consumption | Questions specifically relevant to present heavy alcohol consumption How often did you have a drink containing alcohol in the past year? Never Monthly or less Two to four times a month Two to three times a week Four or more times a week How many drinks did you have on a typical day when you were drinking in the past year? None, I do not drink 1 or 2 3 or 4 5 or 6 7-9 10 or more How often did you have six or more drinks on one occasion in the past year? Never Less than monthly Monthly Weekly Daily or almost daily | 0 1 2 3 4 0 1 2 3 4 0 1 2 3 4 | Total score range: 0-12 Low Risk: 0-3 points Moderate Risk: 4-5 points High Risk: 6-7 points Severe Risk: 8-12 points |
| **Stanford Integrated Psychosocial Assessment for Transplantation (SIPAT)** | Comprehensive psychosocial assessment used to predict psychosocial outcomes in patients undergoing solid organ transplant Not specific for alcohol use disorder | Patient’s readiness level and illness management 1. Knowledge/understanding of the medical illness 2. Knowledge/understanding of transplantation 3. Willingness/desire for treatment (transplant) 4. History of treatment adherence/compliance 5. Lifestyle factors Social support system level of readiness 6. Availability of social support system 7. Functionality of social support system 8. Appropriateness of living space and environment | 0 0 0-4 0 0 0 0 0 0 0-4 0 0 0 0 0 0 0 0-8 0 0 0-4 0 0 0 0 0 0 0 0-8 | Total score range: 0-110 Excellent candidate > 100 Good candidate 70-99 Minimally Acceptable Candidate 40-68 High Risk candidate 0-69 Poor Candidate |

(Continued)
Table 1. (Continued)

| INSTRUMENT NAME | TARGET OF INSTRUMENT | RISK FACTORS ASSESSED | POINTS | PROPOSED INTERPRETATION OF SCORE |
|-----------------|----------------------|------------------------|--------|----------------------------------|
| Alcohol Relapse Risk Assessment (ARRA)\textsuperscript{17} | Prediction of relapse to alcohol use in patients suffering from alcohol use disorder undergoing orthotopic liver transplant | Known risk factors for alcohol relapse in liver disease | One point for every factor present | Total score range: 0-9 0 points: minimal risk 1-3 points: mild risk 4-6 points: moderate risk 7-9 points: severe risk |
| Hopkins Psychosocial Scale (HPSS)\textsuperscript{8} | Prediction of alcohol relapse for patients undergoing LT for severe alcohol associated hepatitis. Utilizing unique factors specific for patients with severe alcoholic hepatitis in addition to Known risk factors for relapse in AUD. | Protective Characteristics | 0-2 | Total score range: -12 to 10 High-risk HPSS score \leq 0 Low-risk HPSS score >0 |
| | | AT Risk Characteristics | | |
| | | 1. Psychiatric comorbidity | 0-2 |
| | | 2. History of other substance abuse | 0-2 |
| | | 3. History of failed rehab attempt | -2-0 |
| | | 4. Family history of alcoholism | -2-0 |
| | | 5. Employment just prior to presentation | -2-0 |
| | | 6. Legal History related to alcohol | -2-0 |
| Sustained Alcohol Use Post-Liver Transplant (SALT)\textsuperscript{18} | Prediction of brief relapse ("slip") and sustained relapse to alcohol use in patients undergoing early liver transplant for severe alcohol associated hepatitis | Simplified psychosocial assessment of known risk factors specific for patients with severe alcoholic hepatitis | Higher the score the greater risk for relapse (no specific a priori cut off) |
| | | 1. \textgreater 10 drinks/day at presentation | +4 |
| | | 2. \textgreater=2 prior failed rehabilitation attempts | +4 |
| | | 3. Any history of prior alcohol-related legal issues | +2 |
| | | 4. Hx of non-THC illicit substance abuse | +1 |

Modified from Shenoy et al,\textsuperscript{14} Im et al,\textsuperscript{5} and Lim and Sundaram.\textsuperscript{12}

6 months, and every 3 to 6 months depending on stability in the first 3 years. Regular post-transplant interviews, routine, and random urine ethyl glucuronide (uETG) tests, participation of social workers and corroboration from family and outpatient providers helped with the evaluation of relapse. IRB approval was obtained for a review of the chart of all patients evaluated for early LT at Mount Sinai Hospital.

Results

Of the 81 patients evaluated by the psychosocial team for alcoholic hepatitis, 11 (14%) were psychosocially cleared and transplanted. All cases (n = 11) and controls (n = 11) met the criteria for AUD and had similar durations of sobriety prior to evaluation (mean 35 vs 22 days, \( P = .08 \)). Transplanted cases versus controls tended to present with their first liver decompensation (73% vs 27%) and with good or developing insight (91% vs 27%). Both groups had similar levels of acceptable social support (100% vs 73%). Three cases presented with their second liver decompensation and were transplanted due to overwhelming support from the recipient review committee. A case with poor insight was cleared and transplanted similarly. The number of drinks per day, years of drinking, failed rehabilitation history, and family history were not different between the groups (Table 3).
| Study | Tool(S) Used | Study Population | Study Type and Follow Up Time | N | Results |
|-------|--------------|------------------|------------------------------|---|---------|
| Lucey et al<sup>25</sup> | MAPS | LT recipients 02/1987-01/1991 at University of Michigan | Retrospective cohort Follow-up between 4- and 8-y | 50 (14 F) | No results provided. MAPS didn’t distinguish between those who abstained from alcohol and those that used alcohol post-transplant. P = not significant |
| Coffman et al<sup>24</sup> | MAPS | LT patients 8/1989-8/1995 in Cedars Sinai Los Angeles, California | Prospective cohort Length of follow up not described | 91 | Mean score for patients who did not relapse was 14.5, and that for the patients who resumed drinking was 12.2 (P = .05) |
| Yates et al<sup>23</sup> | HRAR | Pretransplant patients from the university of Iowa liver clinic or transplant service AH or Cirrhosis | Cross sectional Cohort between 1993 and 1996 | 91 (28 F) | Cutoff allowing a 5% 6-mo relapse risk demonstrated a theoretical 79% agreement (K = 0.56) between the HRAR score and the 6-mo sobriety rule. |
| DiMartini et al<sup>29</sup> | HRAR | Patients transplanted after evaluation for OLT for ALD at the Thomas E starzl institute between March 1993 and December 1994. | Prospective cohort study Regular follow-up for first year, subsequently follow-up as medically necessary | 72 (18 F) | HRAR not predictive of recidivism in transplant sample (P = .174) |
| De Gottardi et al<sup>26</sup> | HRAR | Underwent Liver Transplantation for Alcoholic Liver Disease | Retrospective Cohort study Follow-up time was 61.2 ± 47.5 mo | 387 (92 F) | HRAR score > 4, a duration of abstinence of less than 6 mo before wait-listing for LT and the presence of psychiatric comorbidities were all associated with relapse to Harmful alcohol consumption after LT. In patients with none of these factors, alcohol relapse was 5%, while the presence of 1, 2, or 3 factors was associated with relapse rates of 18%, 64%, and 100% of the patients, respectively. |
| Egawa et al<sup>30</sup> | HRAR | Patients with ALD who underwent LT in Japan from 11/1997 to 12/2011. With information available re alcoholic relapse | Retrospective multi-center cohort | 139 (52 F) | HRAR not predictive of recidivism (P = .48 for relapse; P = .24 for harmful relapse) |
| Zhou et al<sup>31</sup> | HRAR | Outpatient post LT course of 12 wk starting Nov 2011 | Prospective cohort study Follow up to 12 y post LT | 35 (6 F) | HRAR not predictive of recidivism. Sensitivity of the HRAR scale was 17%, the specificity was 90% and the negative predictive value was 84% |
| Lee et al<sup>8</sup> | HPSS (only assessed on AAH group) | LT patients transplanted for ALD exclusively (other liver diseases excluded) 10/2012-06/2015 | Retrospective cohort | AAH – 17 (4 F) | Alcohol cirrhosis – 26 (9 F) | Average Follow-up 1.5 y | HRAR was not predictive of relapse in either group. HPSS identified those with sustained alcohol relapse in post-hoc analysis. AAH Group: No alcohol relapse = 13, HPSS > 3 (+1 to +8), Alcohol Relapse “slip” = 2, HPSS > 1.5 (+1 to +2), P = .09 Sustained alcohol relapse = 2, HPSS > 2 (−4 to −1), P = .03 Study did not validate HPSS because of small sample size |
| Weeks et al<sup>10</sup> | HPSS | All transplants for ALLD 1/10/2012-31/7/2017 | Retrospective cohort study Median follow-up time of 532 d (interquartile range 281-998 d) | 46 (13 F) | Severe alcoholic hepatitis 34 (12 F) | Alcohol cirrhosis | High-risk HPSS found to be predictive of any alcohol relapse in AAH. Hazard ratio = 3.63 (95% CI: 1.16-11.3); p = 0.03 No alcoholic cirrhosis patients had a High-risk HPSS score |
| STUDY                       | TOOL(S) USED | STUDY POPULATION                                                                 | STUDY TYPE AND FOLLOW UP TIME | N          | RESULTS                                      |
|-----------------------------|--------------|----------------------------------------------------------------------------------|------------------------------|------------|----------------------------------------------|
| Lombardo et al32            | HRAR         | All consecutively diagnosed AUD patients for LT 1/2004-4/2016 at hospital clinic of Barcelona, Spain (deaths in first month excluded) | Prospective Cohort Followed until 4/2017 or death Median follow-up of 68 mo (IQR, 35-102 mo) | 309 (31 F) | At an equal duration of abstinence before LT, a moderate-to-high HRAR score (>3) was associated with a 138% increased risk of heavy alcohol relapse Odds ratio = 2.39 (1.02-5.56) P = .04 |
| López-Pelayo et al33        | HRAR         | Patients admitted to the Liver Unit of the Hospital Clinic of Barcelona from 1999 to 2012 with an episode of AAH | Case-control study Follow-up 24 mo | 120 (40 F) | HRAR >3 (OR 2.9) and a history of psychiatric disorders (OR 2.6) predicted long-term treatment retention HRAR >3 (OR 3.0) and previous treatment for AUD (OR 2.9) increased the risk of relapse in the short term. |
| Yano et al28                | AUDIT-C      | LT patients 7/2001-10/2013 in Hiroshima outpatient clinic                           | Cross sectional              | 99 (36 F)  | AUDIT-C - Predictive of post-LT excessive alcohol consumption P = .001 |
| Deutsch-Link et al13        | SALT         | LT patients transplanted between 2011 and 2017 for ALD (chronic)                 | Retrospective Cohort study    | 155 (43 F) | SALT assessed on 138 patients SALT scores ≥7 associated with relapse to any alcohol use post-transplant P = .03 |
| SIPAT                       |              |                                                                                  |                              | 61         | SIPAT assessed on 61 patients SIPAT score ≥21 associated with relapse to any alcohol use post-transplant P = .03 |
| Rodrigue et al17            | ARRA         | Adult primary liver or liver-kidney transplants who suffered from AUD at Beth Israel Deaconess Medical Center from 2002 to 2011 | Retrospective cohort          | 118 (17 F) | ARRA III and ARRA IV were predictive of alcohol relapse Relapse rates were 0% for the ARRA I, 8% for the ARRA II, 57% for the ARRA III, and 75% for the ARRA IV group (P < .001) ARRA III was associated Low and moderate intensity relapse ARRA IV was associated High intensity relapse (χ² = 15.7, P = .003). |
| Rodrigue et al24            |              |                                                                                  |                              |            | A higher ARRA score [β = .88, odds ratios = 2.41 (95% confidence interval = 1.8-3.3), P < .001] and no post-LT SA treatment [β = 21.71, odds ratios = 0.18 (95% confidence interval = 0.04-0.74), P = .02] predicted post-transplant relapse |
| Lee et al18                 | SALT         | LT recipients for AH between January 2012 and March 2017 from 12 U.S. LT centers | Prospective cohort Median post-LT follow-up was 1.6 y (IQR: 0.7-2.8) | 134 (38 F) | The SALT score successfully identified candidates with AH for early LT who were at low risk for sustained alcohol use posttransplant SALT score ≥5 had a 25% positive predictive value (95% CI: 10%-47%) SALT score of <5 had a 95% negative predictive value (95% CI: 89%-98%) for sustained alcohol use post-LT |
Table 3. Characteristics of cases and controls with scoring systems.

|                     | TRANSPALNTED (N=11) | CONTROLS (N=11) | P    |
|---------------------|----------------------|-----------------|------|
| Age (y)             | 43.8                 | 45              | .4   |
| Sex (female)        | 55%                  | 55%             | 1.0  |
| Number of drinks per day | 9.9              | 14.7            | .17  |
| Years of alcohol use | 20.9                | 28.5            | .20  |
| HX of failed rehab   | 27%                  | 36%             | .66  |
| Family HX of Alcoholism | 27%              | 45%             | .37  |
| Alcohol Use Disorder Diagnosis | 100%         | 100%            | 1.0  |
| Severe              | 6/11                 | 11/11           |      |
| Moderate            | 3/11                 | 0/11            |      |
| Mild                | 2/11                 | 0/11            |      |
| Sober time prior to evaluation (d) | 35              | 22              | .08  |
| First liver decompensation | 73%            | 27%             | .03  |
| Good or developing insight | 91%            | 27%             | .002 |
| Consistent report with collateral | 91%         | 45%             | .053 |
| Good social support | 100%                 | 73%             | .06  |

Table 4. Scoring system results.

| SCORING SYSTEMS (SCORE RANGE) | MEAN (95% CI) | MEAN (95% CI) | P    |
|-------------------------------|---------------|---------------|------|
| MAPS (5-20)                   | 17.09 (15.41, 18.77) | 10.00 (8.15, 11.85) | <.001 |
| HRAR (0-6)                    | 2.09 (1.68, 2.50) | 3.09 (2.37, 3.81) | .03  |
| AUDIT-C (0-12)                | 9.09 (7.09, 11.09) | 11.27 (10.21, 12.33) | .08  |
| SIPAT (0-110)                 | 23.27 (16.50, 30.04) | 49.45 (45.46, 53.44) | <.001 |
| ARRA (0-9)                    | 2.27 (1.74, 2.80) | 5.70 (4.99, 6.41) | .01  |
| HPSS (−14 to 10)              | 3.27 (0.85, 5.69) | −2.20 (−4.48, −0.08) | .005 |
| SALT (0-12)                   | 3.82 (2.09, 5.55) | 5.00 (3.37, 6.63) | .34  |

The MAPS, HRAR, SIPAT, ARRA, and HPSS discriminated between cases and controls. The mean case score was a higher MAPS ($m = 17.1$), lower HRAR ($m = 2.0$), lower SIPAT ($m = 23.5$), lower ARRA ($m = 2.3$), and higher HPSS ($m = 2.4$). The AUDIT-C and SALT scores were not significantly different between the groups. The AUDIT-C mode was 12 (the maximum score) in both groups (Table 4).

Cut-off scores of: MAPS 14, HRAR 3, SIPAT 40, ARRA 4, and HPSS 0; suggested a classification of the cases and the controls in the expected direction. No single cut-off score on any of the tools would have selected transplanted patients in this cohort or declined patients in the control. A heat map was created to illustrate the wide range of color-coded scores with respect to the risk of relapse (Figure 1).

One transplanted patient died in the first 6 months from postoperative complications. The surviving transplanted cohort ($n = 10$) had positive psychosocial characteristics with low HRAR ($m = 2.0$), low ARRA ($m = 2.3$), high HPSS ($m = 2.4$), low SALT ($m = 3.8$), and low SIPAT ($m = 23.5$). The cohort has been followed from 5 to 8.5 years (mean = 6.6 years). Three patients (#4, #5, and #11) relapsed to regular alcohol use, one dying of liver failure (Figure 1: Heat map).

**Discussion**

In prospective studies of risk factors for relapse in liver transplant patients, a diagnosis of alcohol dependence (severe or moderate AUD), a family history of alcoholism, low social support, and a shorter duration of pretransplant sobriety predicted
relapse. However, in cases of severe AAH in which there is limited time to wait for a longer period of sobriety or to refer to AUD treatment, a new paradigm must be sought. Our transplant cohort, like the control group, had a high burden of alcohol use, addictive behavior, and genetic load, as demonstrated by the high AUDIT-C score in both groups. Outpatient compliance with addiction treatment was not possible because many of these patients were too sick for discharge. Despite this, the primary criteria for selecting patients with new decompensated liver failure, good insight into their addiction, and strong social support helped identify a successful cohort with a 20% relapse rate after a mean follow-up of 6.6 years. This relapse rate is consistent with the reported 20% to 30% relapse to heavy drinking after liver transplantation for all forms of alcoholic liver disease. We cannot comment on the potential relapse outcomes of patients who were not transplanted and did not survive.

Validated cut-off scores of the HRAR < 3, ARRA < 4, HPSS > 0, and SIPAT < 40 would have corroborated the stratification process used in our center. However, given the recent findings from a large multicenter trial that SALT scores below 5 had a 95% NPV for sustained alcohol use post LT, it is possible that our psychosocial clearance was too strict and 5 out of the 11 controls would have been deemed acceptable candidates by this score alone.

The MAPS was highly correlated with our institutional psychosocial assessment, possibly because its emphasis on insight as a protective factor paralleled our use of emerging insight in the face of new liver disease. The use of new information has been a practical way to measure readiness for transplantation. This finding is consistent with previous studies that have identified self-awareness of choice behavior (insight) as a prediction of substance disorder related choices in addiction.

Similarly, SIPAT and HPSS, with their focus on readiness for transplantation candidates with lower risk of relapse. As the study population was focused on patients undergoing expedited transplant listing, tools that utilized extended abstinence as a variable, such as a recently developed tool that required follow up time to observe if the patient followed up with an intensive outpatient program (IOP) were not utilized in this study.

As the medical community has moved to view alcohol use disorder as a disease and not a vice, it has become universally accepted that patients with ALD should not be automatically excluded from receiving a liver transplant. While the assessment of patients with ALD has been criticized as

---

**Figure 1.** Heat map of cases and controls with 5+ year follow up.

| CASES: Transplanted | CONTROLS: Declined for Psychosocial Reasons |
|---------------------|--------------------------------------------|
| MAPS                | 20 18 16 20 13 20 14 13 18 16             | 13 9 15 11 10 8 6 15 8 7 8 |
| HRAR                | 2 2 2 2 1 2 3 1 3 2 3                     | 1 1 3 3 4 5 4 3 3 3 4 |
| AUDIT-C             | 6 6 6 12 4 6 12 12 12 12 12              | 11 6 12 12 12 12 12 12 12 11 12 |
| SIPAT               | 8 37 26 14 21 17 21 34 18 14 46           | 46 49 50 59 38 55 42 46 52 60 47 |
| ARRA                | 1 1 2 2 4 2 2 3 3 2 3                     | 5 7 5 6 5 6 7 3 7 7 5 |
| HPSS                | 9 1 -3 6 5 3 2 4 2 5 -8                    | -5 -5 2 -6 -3 -8 -8 -1 1 -2 -8 |
| SALT                | 0 0 7 4 0 4 4 4 6 4 9                     | 5 1 4 5 5 9 4 4 1 8 |

| Patient # | 1 | 2 | 3 | 4 | 5 | 6 | 7 | 8 | 9 | 10 | 11 |
|------------|---|---|---|---|---|---|---|---|---|----|----|
| Years alive| 8.5| 8.5| 7.5| 3| 6.5| 0.5| 6| 5.5| 5.5| 5 |
| D = Died   | D | D | D | D | D | D | D | D | D | D |
| R = Relapse| NR | NR | S | R | R | NR | NR | NR | R |
| S = Slip   | NR | NR | S | R | R | NR | NR | NR | R |
| NR = No Relapse | | | | | | | | | | |

---

Substance Abuse: Research and Treatment
somewhat subjective\textsuperscript{42} and inconsistent,\textsuperscript{43,44} the use of numeric scoring systems partially alleviates these ethical challenges by introducing a numeric score that can be used consistently to portray a patient’s alcohol related behaviors and relapse risk. In our study, scores from the objective scales largely reflected the risk stratification that our institution employed in our initial AAH transplant population.

Limitations of the case-control include the retrospective nature of our data collection and our small sample size of patients cleared and transplanted. The small sample size was largely a function of the high mortality associated with severe AAH and that only 27\% of all potential transplant candidates evaluated psychosocially were deemed acceptable by the methods used at our center. This clearance rate was similar to the Franco-Belgian study done by Mathurin et al\textsuperscript{7} using similar criteria for clearance. Larger cohorts that include a wide range of risk scores will be necessary to validate the use of any of these tools, as well as to analyze which individual factors can prognosticate a favorable candidate in this unique population.

Conclusions

Patients with AUD and new information on their addiction, social support, and readiness for transplantation at the time of evaluation for transplantation have low rates of alcohol relapse after transplantation. Scoring systems may approximate and assist in directing this traditional selection process. Pre-existing scoring systems may have varying utility in their ability to assist in making this determination. Patients with ALD should instead be evaluated to stratify risk for selection for transplantation and should be referred for AUD treatment and post-LT follow-up. Centers that perform liver transplants for patients with alcoholic hepatitis should include a psychosocial team with addiction experience and consider known risk factors for AUD relapse in their initial assessments.

Author contributions

AD was involved in preparing the manuscript, as well as retrospectively reviewing the psychiatric and social work data confirming the DSM-5 diagnosis of AUD and scored the HRAR, ARRA, HPSS, and SIPAT. PD was involved in preparing the manuscript. KM was involved in retrospectively reviewing the psychiatric and social work data confirming the DSM-5 diagnosis of AUD and scored the HRAR, ARRA, HPSS, and SIPAT. OM was involved in preparing the manuscript. ES performed the initial pre-transplant psychosocial evaluation. LF performed the initial pre-transplant psychosocial evaluation. ME performed the initial pre-transplant psychosocial evaluation. GI performed the pre- and post-transplant medical evaluations. AS was involved in editing the manuscript, performed the initial pre-transplant psychiatric evaluations and retrospectively scored the MAPS, AUDIT-C and SALT psychometric scales.

ORCID iD

Aryeh Dienstag https://orcid.org/0000-0001-8120-4846

REFERENCES

1. Dew MA, DiMartini AF, Steel J, et al. Meta-analysis of risk for relapse to substance use after transplantation of the liver or other solid organs. Liver Transpl. 2008;14:159-172.
2. Rice JP, Lucey MR. Should length of sobriety be a major determinant in liver transplant selection? Curr Opin Organ Transplant. 2013;18:259-264.
3. Zhao J, Hetrick SE, Prabhakar M, Musselman D. Risk factors for alcohol relapse following orthotopic liver transplantation: a systematic review. Psychosomatics. 2015;56:21-35.
4. DiMartini A, Dew MA, Day N, et al. Trajectories of alcohol consumption following liver transplantation. Am J Transplant. 2010;10:2305-2312.
5. Im GY, Cameron AM, Lucey MR. Liver transplantation for alcoholic hepatitis. J Hepatol. 2019;70:328-334.
6. Hosseini N, Shor J, Szabo G. Alcoholic hepatitis: a review. Alcohol Alcohol. 2019;54:408-416.
7. Mathurin P, Moreno C, Samuel D, et al. Early liver transplantation for severe alcoholic hepatitis. N Engl J Med. 2011;364:1900.
8. Lee BP, Hsu C, Haugen C, et al. Three-year results of a pilot program in early liver transplantation for severe alcoholic hepatitis. Ann Surg. 2017;265:20-29.
9. Im GY, Kim-Schluger L, Shenoy A, et al. Early liver transplantation for severe alcoholic hepatitis in the United States—a single-center experience. Am J Transplant. 2016;16:841-849.

Limitations of the case-control include the retrospective nature of our data collection and our small sample size of patients cleared and transplanted. The small sample size was largely a function of the high mortality associated with severe AAH and that only 27% of all potential transplant candidates evaluated psychosocially were deemed acceptable by the methods used at our center. This clearance rate was similar to the Franco-Belgian study done by Mathurin et al\textsuperscript{7} using similar criteria for clearance. Larger cohorts that include a wide range of risk scores will be necessary to validate the use of any of these tools, as well as to analyze which individual factors can prognosticate a favorable candidate in this unique population.

Conclusions

Patients with AUD and new information on their addiction, social support, and readiness for transplantation at the time of evaluation for transplantation have low rates of alcohol relapse after transplantation. Scoring systems may approximate and assist in directing this traditional selection process. Pre-existing scoring systems may have varying utility in their ability to assist in making this determination. Patients with ALD should instead be evaluated to stratify risk for selection for transplantation and should be referred for AUD treatment and post-LT follow-up. Centers that perform liver transplants for patients with alcoholic hepatitis should include a psychosocial team with addiction experience and consider known risk factors for AUD relapse in their initial assessments.

Author contributions

AD was involved in preparing the manuscript, as well as retrospectively reviewing the psychiatric and social work data confirming the DSM-5 diagnosis of AUD and scored the HRAR, ARRA, HPSS, and SIPAT. PD was involved in preparing the manuscript. KM was involved in retrospectively reviewing the psychiatric and social work data confirming the DSM-5 diagnosis of AUD and scored the HRAR, ARRA, HPSS, and SIPAT. OM was involved in preparing the manuscript. ES performed the initial pre-transplant psychosocial evaluation. LF performed the initial pre-transplant psychosocial evaluation. ME performed the initial pre-transplant psychosocial evaluation. GI performed the pre- and post-transplant medical evaluations. AS was involved in editing the manuscript, performed the initial pre-transplant psychiatric evaluations and retrospectively scored the MAPS, AUDIT-C and SALT psychometric scales.

ORCID iD

Aryeh Dienstag https://orcid.org/0000-0001-8120-4846

REFERENCES

1. Dew MA, DiMartini AF, Steel J, et al. Meta-analysis of risk for relapse to substance use after transplantation of the liver or other solid organs. Liver Transpl. 2008;14:159-172.
2. Rice JP, Lucey MR. Should length of sobriety be a major determinant in liver transplant selection? Curr Opin Organ Transplant. 2013;18:259-264.
3. Zhao J, Hetrick SE, Prabhakar M, Musselman D. Risk factors for alcohol relapse following orthotopic liver transplantation: a systematic review. Psychosomatics. 2015;56:21-35.
4. DiMartini A, Dew MA, Day N, et al. Trajectories of alcohol consumption following liver transplantation. Am J Transplant. 2010;10:2305-2312.
5. Im GY, Cameron AM, Lucey MR. Liver transplantation for alcoholic hepatitis. J Hepatol. 2019;70:328-334.
6. Hosseini N, Shor J, Szabo G. Alcoholic hepatitis: a review. Alcohol Alcohol. 2019;54:408-416.
7. Mathurin P, Moreno C, Samuel D, et al. Early liver transplantation for severe alcoholic hepatitis. N Engl J Med. 2011;364:1900.
8. Lee BP, Hsu C, Haugen C, et al. Three-year results of a pilot program in early liver transplantation for severe alcoholic hepatitis. Ann Surg. 2017;265:20-29.
9. Im GY, Kim-Schluger L, Shenoy A, et al. Early liver transplantation for severe alcoholic hepatitis in the United States—a single-center experience. Am J Transplant. 2016;16:841-849.
10. Weeks SR, Sun Z, McCaul ME, et al. Liver transplantation for severe alcoholic hepatitis, updated lessons from the world’s largest series. J Am Coll Surg. 2018;226:549-557.
11. Sundaram V, Wu T, Klein AS, et al. Liver transplantation for severe alcoholic hepatitis: report of a single center pilot program. Transplant Proc. 2018;50:3527-3532.
12. Lim J, Sundaram V. Risk factors, scoring systems, and interventions for alcohol relapse after liver transplantation for alcoholic liver disease. Clin Liver Dis. 2011;15:105-110.
13. Deutsch-Link S, Weinrieb RM, Jones LS, Solga SF, Weinberg EM, Serper M. Prior relapse, ongoing alcohol consumption, and failure to engage in treatment predict alcohol relapse after liver transplantation. Dig Dis Sci. 2020;65:2089-2103.
14. Shenoy A, Dienstag A, Dienstag P, et al. Scoring systems to assess relapse risk in alcohol use disorder presenting for early liver transplantation: a systematic review. Gen Hosp Psychiatry. 2021;72:23-30.
15. Lim J, Curry MP, Sundaram V. Risk factors and outcomes associated with alcohol relapse after liver transplantation. World J Hepatol. 2017;9:771-780.
16. Lucey MR, Merion RM, Henley KS, et al. Selection for and outcome of liver transplantation in alcoholic liver disease. Gastroenterology. 1992;102:1736-1741.
17. Rodrigue JR, Hanto DW, Curry MP. The alcohol relapse risk assessment: a scoring system to predict the risk of relapse to any alcohol use after liver transplant. Prog Transplant. 2013;23:310-318.
18. Lee BP, Vittinghoff E, Hsu C, et al. Predicting low risk for sustained alcohol use after early liver transplant for acute alcoholic hepatitis: the sustained alcohol use post-liver transplant score. Hepatology. 2019;69:1477-1487.
19. Yates WR, Martin M, LaBeccique D, Hillebrand D, Voigt M, Pfähl D. A model to examine the validity of the 6-month abstinence criterion for liver transplantation. Alcohol Clin Exp Res. 1992;16:513-517.
20. Bradley KA, McDonell MB, Bush K, Kivlahan DR, Diehr P, Fihn SD. The AUDIT alcohol consumption questions. Alcohol Clin Exp Res. 1999;22:1842.
21. Maldonado JR, Dubois HC, David EE, et al. The Stanford Integrated Psychosocial Assessment for Transplantation (SIPAT): a new tool for the psychosocial evaluation of pre-transplant candidates. Psychosomatics. 2012;53:123-132.
22. Beresford TP, Schwartz J, Wilson D, Merion R, Lucey MR. The short-term psychological health of alcoholic and non-alcoholic liver transplant recipients. Alcohol Clin Exp Res. 1992;16:996-1000.
23. Yates WR, Booth BM, Reed DA, Brown K, Masterson BJ. Descriptive and predictive validity of a high-risk alcoholism relapse model. J Stud Alcohol. 1993;54:645-651.
24. Coffman KL, Hoffman A, Sher L, Rojter S, Vierling J, Makowka L. Treatment of the postoperative alcoholic liver transplant recipient with other addictions. *Liver Transplant Surg*. 1997;3:322-327.

25. Lucey MR, Carr K, Beresford TP, et al. Alcohol use after liver transplantation in alcoholics: a clinical cohort follow-up study. *Hepatology*. 1997;25:1223-1227.

26. De Gottardi A, Spahr L, Gelez P, et al. A simple score for predicting alcohol relapse after liver transplantation: results from 387 patients over 15 years. *Arch Intern Med*. 2007;167:1183-1188.

27. Maldonado JR, Sher Y, Lolak S, et al. The Stanford integrated psychosocial assessment for transplantation: a prospective study of medical and psychosocial outcomes. *Psychosom Med*. 2015;77:1018-1030.

28. Yano T, Ohira M, Sakamoto R, et al. Alcohol Use Disorders Identification Test consumption predicts the risk of excessive alcohol consumption after liver transplantation. *Transplant Proc*. 2019;51:1934-1938.

29. DiMartini A, Magill J, Fitzgerald MG, et al. Use of a high-risk alcohol relapse scale in evaluating liver transplant candidates. *Alcohol Clin Exp Res*. 2000;24:1189-1201.

30. Egawa H, Nishimura K, Teramukai S, et al. Risk factors for alcohol relapse after liver transplantation for alcoholic cirrhosis in Japan. *Liver Transplant*. 2014;20:298-310.

31. Zhou M, Wagner LM, Diffo T, Naegle M. Implementation of the high-risk alcoholism relapse scale in a liver transplant clinic. *Gastroenterol Nurs*. 2015;38:447-454.

32. Lombardo-Quezada J, Colmenero J, López-Pelayo H, et al. The High-Risk alcoholism relapse score predicts alcohol relapse among liver transplant candidates with less than six months of abstinence. *Liver Transplant*. 2019;25:25460.

33. López-Pelayo H, Miquel L, Ahmariro J, et al. Treatment retention in a specialized alcohol programme after an episode of alcoholic hepatitis: impact on alcohol relapse. *J Psychosom Res*. 2019;116:75-82.

34. Rodrigo JR, Hanto DW, Curry MP. Substance abuse treatment and its association with relapse to alcohol use after liver transplantation. *Liver Transplant*. 2013;19:1387-1395.

35. Kim JS, Kim GJ, Lee JM, Lee CS, Oh JK. HAIS (Hanil alcohol Insight Scale): validation of an insight-evaluation instrument for practical use in alcoholism. *J Stud Alcohol*. 1998;59:52-55.

36. Beresford TP, Turcotte JG, Merion R, et al. A rational approach to liver transplantation for the alcoholic patient. *Psychosomatics*. 1990;31:241-254.

37. Moeller SJ, Hajcak G, Parvaz MA, Dunning JP, Volkow ND, Goldstein RZ. Psychophysiological prediction of choice: relevance to insight and drug addiction. *Brain*. 2012;135:3481-3494.

38. Havassy BE, Hall SM, Wasserman DA. Social support and relapse: commonalities among alcoholics, opiate users, and cigarette smokers. *Addict Behav*. 1991;16:235-246.

39. Moon TJ, Mathias CW, Mullen J, et al. The role of social support in motivating reductions in alcohol use: a test of three models of social support in alcohol-impaired drivers. *Alcohol Clin Exp Res*. 2019;43:123-134.

40. Marroni CA, Fleck AM Jr, Fernandes SA, et al. Liver transplantation and alcoholic liver disease: history, controversies, and considerations. *World J Gastroenterol*. 2018;24:2785-2805.

41. Mellingler JL, Volk ML. Transplantation for alcohol-related liver disease: is it fair? *Alcohol Alcohol*. 2018;53:173-177.

42. Solga SF, Serper M, Young RA, Forde KA. Transplantation for alcoholic hepatitis: are we achieving justice and utility? *Hepatology*. 2019;69:1798-1802.

43. Singhvi A, Welch AN, Levitsky J, Singhvi D, Gordon EJ. Ethical considerations of transplantation and living donation for patients with alcoholic liver diseases. *AMA J Ethics*. 2016;18:163-173.

44. Webb K, Shepherd L, Day E, Masterton G, Neuberger J. Transplantation for alcoholic liver disease: Report of a consensus meeting. *Liver Transplant*. 2006;12:301-305.