

Impact and Benefits of the MELD Scoring System for Liver Allocation

Richard B. Freeman Jr

Division of Transplantation, Tufts Medical Center, Boston, MA, USA

Abstract

The MELD score was described in 2000 and was employed as part of the US liver allocation policy in 2002. There were many reasons behind the decision to use the MELD score and there have been many benefits to the liver allocation system by doing so. Measuring any system used for allocation of scarce resources can be difficult, but evaluation of the justice and utility of such a system provides some framework for assessing its effectiveness. In this review, the benefits realized from the MELD-based allocation system will be assessed according to justice and utility parameters. In organ allocation, individual justice is served when patient-specific variables are used to assign waiting list priority rather than using physician-based observations or behaviors. Utility should be measured, not just in terms of patient survival after transplant, but also in terms of the overall utility of the system for assigning organs to those most in need and giving little priority to those who will be harmed or have little benefit.

The impact of adoption of this system can be measured by the number of the publications that include MELD or liver allocation in their data. The world has recognized the relative objectivity of the MELD score and the ability to communicate among widely diverse groups using this common language. Perhaps this is the most important impact of the MELD "era" (Trends in Transplant. 2009;3:70-6)

Corresponding author: Richard B. Freeman, rfreeman@tufts-nemc.org

Key words

MELD score. Liver allocation. Survival benefit.

Correspondence to:

Richard B. Freeman
Division of Transplantation, Box 40
Tufts-New England Medical Center
750 Washington Street
Boston, MA 02111, USA
E-mail: rfreeman@tufts-nemc.org

Introuduction

There are many approaches to the allocation of scarce resources. Systems designed for prioritizing the recipients of these resources can be weighted, on the one hand, toward individual justice or, on the other hand, toward maximizing systemic utility. Thus, in the case of organ allocation, donor livers can be directed to the most deserving candidates, based on individual assessments of need, or to those candidates most likely to achieve the best outcome overall. However, implementations of systems at either end of this justice/utility spectrum necessarily carry undesirable consequences. For example, allocating donor livers purely to individuals deemed to “need the organ most” requires that some measurement of need be defined. To maintain fairness, this measure should not be subjective since other patients’ estimations of need will also be affected. Moreover, observer biases in determining need also can influence the interpretation of what is just. At the other end of the spectrum, allocating organs only to patients most likely to achieve the best outcome means that many other patients with good but not necessarily the best characteristics for success will go unserved, and potentially many more will die while waiting in such a pure utilitarian system. Patients with the best outcome also are likely to have reasonably good outcomes without the transplant, so less is gained by allocating the limited number of donor livers to patients who are going to have better outcomes whether or not they get the transplant. Moreover, for patients with stable, relatively mild liver disease, the risks of the transplant surgery and medications often are greater than the liver disease itself, making transplantation a more risky proposition than waiting until the liver disease progresses further.

In the remainder of this review, I will highlight how the model for end-stage liver disease (MELD)-based liver allocation system in the USA has provided benefit in better achieving

a balance between individual justice and utility, and subsequently outline the worldwide impact that the MELD score and allocation systems based on MELD have provided.

MELD score advantages

Partly due to dissatisfaction with existing allocation systems and partly due to improved predictive models, the U.S. liver allocation system was revised in 2002 in an effort to better balance the justice and utility tradeoffs¹. The MELD score, originally developed to predict the risk of death in patients undergoing transjugular intrahepatic portosystemic shunt (TIPS) procedures², was validated as a reproducible, reasonably accurate predictor of mortality in several different populations of patients with chronic liver disease³. From an individual justice point of view, the MELD score offered several advantages for prioritizing waiting liver transplant candidates.

1. The components of the MELD score, three blood tests – bilirubin, international normalized ratio (INR) and creatinine – are much more objective measures directly related to a patient’s condition than many of the previously employed measures for liver disease severity. For example, encephalopathy and ascites are much more subjective measures and depend on the observer’s experience and timing of the observation. They are more specific to the observer and do not directly indicate a patient’s condition. Similarly, time on the waiting list does not define the intrinsic liver disease of the patient and is much more a function of physician practice behavior, referral to the liver transplant program, and access to healthcare in general and much less related to the severity of illness or mortality risk⁴. In addition, location of care, often categorized as home, hospitalized, or in intensive care, is also more related to physician behavior than intrinsic patient condition. These previously employed measures of liver transplant need were much less accurate reflections

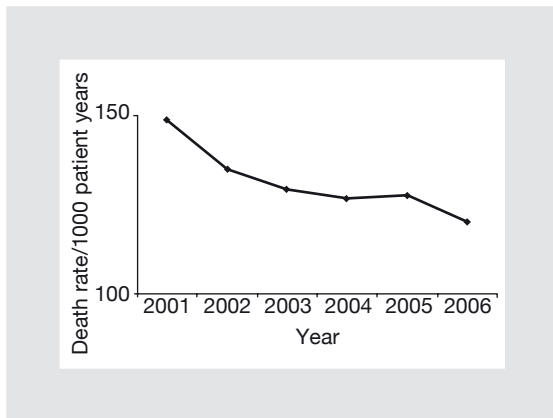


Figure 1. Unadjusted death rates per 1,000 patient-years at risk, 1997-2006. Source: OPTN/SRTR 2007 Annual Report, Table 9.3.

of individual justice because they are not directly derived from the individual patient and much more associated with the treating physicians' observations and/or the system.

2. The MELD score, in defining risk of dying from liver disease, provided a much more precise and transparent measure of liver disease severity than some other measures such as time on the list or location of care delivery.

3. The MELD score is a much more continuous scale than previous categorical systems for prioritizing patients. Previous classification systems using categorical variables tended to group patients together, even though they may have had considerably different underlying clinical characteristics, different need for liver transplantation, and varying risks for death. In addition, disease severity scales such as the Child Turcotte Pugh Score imposed a "ceiling effect" on the more ill candidates since patients with bilirubin of 5 were treated the same as a bilirubin of 15 or 20. Individual justice is better served when individual patients can be discerned and separated more accurately by employing continuous measures of disease severity instead of using categorical classifications.

Because the MELD score provides a relatively simple method for risk adjustment of patient and graft survival after transplantation, the

MELD score also has benefits when measuring liver transplant utility. Many investigators have documented that liver transplant results are associated with MELD score at transplantation⁵⁻⁹. However, although this association is consistent, liver transplant results are also influenced by non-recipient factors such as donor characteristics¹⁰ or surgeon/center experience¹¹. Thus, the MELD score is not nearly as accurate for predicting posttransplant survival as it is for predicting survival without a liver transplant¹². Nonetheless, stratifying patients by MELD score at transplantation has provided important utilitarian observations about the entire liver transplant system when pretransplant and posttransplant survival are combined together to define liver transplant benefit. Merion, et al. measured transplant benefit and found that for patients with lower MELD scores, their projected survival was greater without a liver transplant than could be expected if they were to receive a transplant¹³. For these patients, liver transplantation does not provide a survival benefit. On the other end of the MELD score scale, patients with the highest MELD score always received benefit, even if they receive grafts from donors with higher risks of graft failure¹⁴. These observations again point out the benefits of this relatively objective, transparent disease severity scale.

The most recent liver transplant results have been published in the U.S. Organ Procurement and Transplantation Network/Scientific Registry for Transplant Recipients OPTN/SRTR annual report. Figure 1 indicates the steady decline in death rates since the U.S. liver allocation system was changed in 2002 to a MELD-based system of implementation. Figure 2 displays an equally impressive decline in time to transplantation after the allocation system was changed. These are two extremely important benefits of implementing the MELD system in the USA. Separate analyses have confirmed these results¹⁵. By using a metric defining mortality risk, the MELD-based system more precisely directs organs to those most likely to die in the short term, thereby reducing the waiting

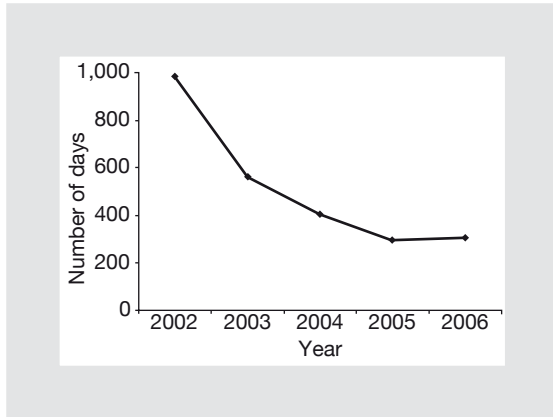


Figure 2. Median time to transplant for new liver waiting list registrations, 2002-2006. Source: OPTN/SRTR 2007 Annual Report, Table 1.5.

list death rate. Secondly, by removal of the time-based component from the old liver allocation system that had directed organs to patients longest on the list, time to transplantation has been dramatically reduced.

Impact of MELD score and MELD-based liver allocation

Patients have not seen any decrement in posttransplant outcome since implementing the MELD system. Figure 3 displays the most recent data for adjusted patient survival by year from the OPTN/SRTR 2007 Annual report¹⁶. The survival curves are virtually identical, indicating that despite using a more direct disease severity scale, liver transplant patient survival has not changed compared with allocation policy in effect prior to the MELD score system. Thus, the MELD-based liver allocation system has impacted the justice and utility of the system as described above, but it has not had an adverse impact on patient (Fig. 3) or graft (data not shown) survival rates.

The MELD model was first published in 2001². In the ensuing nine years, the impact of this score can be measured by a recent MEDLINE search, using MELD as the search term

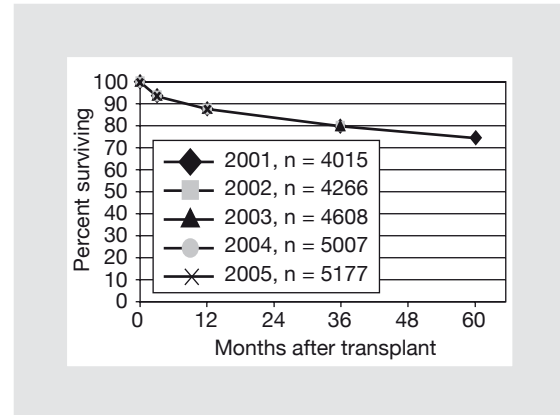


Figure 3. Liver transplant adjusted patient survival stratified by year of transplant. Compared with results from 2001, there is no difference in one-year survival rates compared with 2005. Source: OPTN/SRTR 2007 Annual Report, Table 9.13a.

in which 689 citations were retrieved. Several studies have found a close correlation between preoperative MELD score and early postoperative mortality for patients undergoing routine general surgical procedures^{17,18} and for patients undergoing hepatic resection¹⁹⁻²¹. One recent study stratified outcome data for patients treated with simultaneous ileoanal pouch and liver transplantation using the MELD score²².

In China, investigators have found that the MELD score predicts mortality for patients with decompensated hepatitis B²³ and primary biliary cirrhosis²⁴ related liver disease, and that it can be used to improve outcome prediction for patients with hepatocellular carcinoma (HCC) and cirrhosis undergoing locoregional treatments^{25,26} or waiting on the transplant list²⁷. Chinese ultrasonographers have also shown that the MELD score correlated with an ultrasound-derived measure of liver disease severity²⁸.

The MELD score has been correlated with many other manifestation of liver disease such as hepatic venous pressure gradient in Spain²⁹, findings on MRI scans³⁰, cognitive impairments³¹, brain blood flow in Italy³², TIPS flow velocities³³, response to medical treatment for refractory ascites in Canada³⁴, soluble vascular cell adhesion molecule-1 levels in Spain³⁵, and liver disease-

related quality of life in Brazil³⁶. The MELD score has also been used extensively to stratify study subjects and risk adjust for outcome analyses in studies of renal failure-related mortality in cirrhotics in France³⁷ and Spain³⁸, for primary hepatitis C virus (HCV) treatment³⁹, for treatment of HCC in Taiwan⁴⁰, for predicting sobriety and mortality in alcoholic liver disease in Sweden⁴¹, and for evaluating outcome after treatment for severe liver failure in Singapore⁴². Studies of acute liver failure have found that MELD is closely correlated with mortality risks for cases where the etiology of the acute liver failure is unknown⁴³, and is useful for stratifying patients treated with artificial liver support devices⁴⁴.

Outside of the liver allocation role MELD plays, many liver transplant researchers have reported transplant results in the context of MELD at the time of transplantation. A descriptive analysis of the waiting list in Iran⁴⁵ stratified by candidates' MELD scores was published in 2006. Selection of liver transplant candidates in Mexico⁴⁶ and Malaysia⁴⁷ using the MELD score has been described. Changes in MELD score over time have been associated with increased waiting list mortality in the USA^{48,49} and in Italy⁵⁰, although the most significant changes tend to occur very late in the course of disease, which limits the prognostic usefulness for this "Delta MELD" measurement⁴⁹. The MELD was shown to be a good predictor for liver transplant waiting list mortality in Romania⁵¹. French investigators recently documented that enterobacteremia following liver transplantation is correlated with MELD score at transplant⁵² and there is a correlation between histopathologic findings in explanted livers after transplant and MELD score at the time of transplant⁵³.

Many aspects of the liver transplant waiting list have been addressed utilizing MELD. Investigators from the University of Pittsburgh reported that MELD underestimates mortality risk for HIV-infected liver transplant candidates⁵⁴. Others have shown that a competing risks approach to estimating mortality risks on the

waiting list should be applied⁵⁵, and that even when these methods are used to calculate waiting list dynamics, the MELD score remains highly predictive of waiting list mortality. A recent report from France indicates that a liver allocation system incorporating some aspects of the MELD-based system from the USA resulted in reduced waiting list deaths and improved access to transplant for the sickest patients⁵⁶.

As mentioned above, lifetime benefit calculations for liver transplant patients combine pre- and post-liver transplant survival rates and stratify these by MELD score at entry to the waiting list⁵⁷. Other investigators from Austria⁵⁸, Belgium⁵⁹, and a European-wide study⁶⁰ have also reported pre- and post-liver transplant survival rates stratified by MELD at waitlist entry and at transplantation⁶¹. There are other liver transplant outcome reports from Israel⁶², Lithuania⁶³, for primary sclerosing cholangitis in Nordic Countries⁶⁴, and for re-transplant recipients⁶⁵. Recent reports from Germany have documented an association between living donor liver transplant recipient hospital mortality and MELD score at transplantation⁶⁶.

MELD has been used to perform liver transplant recipient risk adjustment to enable calculation of an index that characterizes donors in terms of the risk of liver graft failure (Donor Risk Index)¹⁰ and to assess the outcome of using liver grafts from so-called extended criteria donors in Italy⁶⁷ and in the USA⁶⁸. MELD has also been used to stratify results in a paper comparing transfusion requirements among high and low MELD patients⁶⁹ and in living versus deceased donor liver transplant recipients⁷⁰, and to report an association with recurrent HCV after liver transplantation⁷¹.

Recent investigators have found a correlation between MELD score and pretransplant dobutamine stress echo findings and these were predictive of cardiac events during or after liver transplantation surgery⁷². The MELD scores also are directly related to liver

transplant costs and it is clear that the MELD-based liver allocation system has an important impact on overall transplant center costs⁷³.

The MELD score itself has become a useful, widely accepted tool for measuring severity of chronic and acute liver disease. The benefits are many, but in particular, despite some concerns about variations in laboratory values^{74,75}, MELD scores are consistent across all types of patients with chronic liver disease no matter which country or region of the world and no matter what biases a clinical observer might have in otherwise assessing liver disease severity. The recognition of the usefulness of MELD and the impact thereof are represented by the explosion of publications and the implementation of MELD-based or MELD-like liver allocation systems across the world.

References

- Freeman RB, Wiesner RH, Harper A, et al. The New Liver Allocation System: moving towards evidence-based transplantation policy. *Liver Transpl.* 2002;8:851-8. **Original description of the MELD based allocation system in the USA.*
- Malinchoc M, Kamath PS, Gordon FD, Peine CJ, Rank J, ter Borg PC. A model to predict poor survival in patients undergoing TIPS. *Hepatology.* 2000;31:864-71. ***Original description of the MELD score.*
- Wiesner RH, McDiarmid SV, Kamath PS, et al. MELD and PELD: application of survival models to liver allocation. *Liver Transpl.* 2001;7:567-80. ***Important paper documenting that MELD is an accurate predictor of mortality among several different patient cohorts and without inclusion of subjective ascites and encephalopathy variables.*
- Freeman RB, Edwards EB. Liver transplant waiting time does not correlate with waiting list mortality: implications for liver allocation policy. *Liver Transpl.* 2000;6:543-52.
- Freeman RB, Harper A, Edwards EB. Excellent liver transplant survival rates under the MELD/PELD system. *Transplant Proc.* 2005;37:585-8.
- Waki K. UNOS Liver Registry: ten year survivals. *Clin Transpl.* 2006;29:39. **Comprehensive report of long term survival for liver transplantation in the USA.*
- Freeman RB, Steffick DE, Guidinger MK, Farmer DG, Berg CL, Merion RM. Liver and intestine transplantation in the United States, 1997-2006. *Am J Transplant.* 2008;8:958-76. ***This paper provides a good summary of recent liver transplant results from the US OPTN/SRTR database.*
- Nagler E, Van Vlierberghe H, Colle I, Troisi R, de Hemptinne B. Impact of MELD on short-term and long-term outcome following liver transplantation: a European perspective. *Eur J Gastroenterol Hepatol.* 2005;17:849-56.
- Silberhumer GR, Hetz H, Rasoul-Rockenschaub S, et al. Is MELD score sufficient to predict not only death on waiting list, but also posttransplant survival? *Transpl Int.* 2006;19:275-81.
- Feng S, Goodrich NP, Bragg-Gresham JL, et al. Characteristics associated with liver graft failure: the concept of a donor risk index. *Am J Transplant.* 2006;6:783-90. ***Provides a continuous scale that defined risk of liver graft failure according to donor characteristics.*
- Edwards EB, Roberts JP, McBride MA, Schulak JA, Hunsicker LG. The effect of the volume of procedures at transplantation centers on mortality after liver transplantation. *N Engl J Med.* 1999;341:2049-53.
- Habib S, Berk B, Chang CC, et al. MELD and prediction of post-liver transplantation survival. *Liver Transpl.* 2006; 12:440-7.
- Merion RM, Schaubel DE, Dykstra DM, Freeman RB, Port FK, Wolfe RA. The survival benefit of liver transplantation. *Am J Transplant.* 2005;2:307-13. ***This is a seminal paper that defines the concept of survival benefit in liver transplantation.*
- Schaubel DE, Sima CS, Goodrich NP, Feng S, Merion RM. The survival benefit of deceased donor liver transplantation as a function of candidate disease severity and donor quality. *Am J Transplant.* 2008;8:419-25. ***This paper points out that, for patients with high MELD scores, the risk of death without a transplant is much higher than the risk of any graft these patients might receive.*
- Austin MT, Poulouse BK, Ray WA, Arbogast PG, Feurer ID, Pinson CW. Model for end-stage liver disease: did the new liver allocation policy affect waiting list mortality? *Arch Surg.* 2007;142:1079-85. **Confirms reductions in waiting list mortality and time to transplant with the MELD system.*
- See www.ustransplant.org
- Befeler AS, Palmer DE, Hoffman M, Longo W, Solomon H, DiBisceglie AM. The safety of intra-abdominal surgery in patients with cirrhosis: model for end-stage liver disease score is superior to Child-Turcotte-Pugh classification in predicting outcome. *Arch Surg.* 2005;140:650-4.
- Northup PG, Wanamaker RC, Lee VD, Adams RB, Berg CL. MELD predicts non-transplant surgical mortality in patients with cirrhosis. *Ann Surg.* 2005;242:244-51. **Validates MELD as a preoperative predictor of surgical mortality in non-liver transplant surgery.*
- Teh SH, Christein J, Donohue J, et al. Hepatic resection of hepatocellular carcinoma in patients with cirrhosis: MELD score predicts perioperative mortality. *J Gastrointest Surg.* 2005;9:1207-15.
- Schroeder RA, Marroquin CE, Bute BP, Khuri S, Henderson WG, Kuo PC. Predictive indices of morbidity and mortality after liver resection. *Ann Surg.* 2006;243:373-9.
- Cucchetti A, Ercolani G, Vivarelli M, et al. Impact of MELD score on prognosis after hepatectomy for HCC on cirrhosis. *Liver Transpl.* 2006;12:966-71.
- Mathis KL, Dozois EJ, Larson DW, et al. Ileal pouch-anal anastomosis and liver transplantation for ulcerative colitis complicated by primary sclerosing cholangitis. *Br J Surg.* 2008;95:882-6.
- Chan HL, Chim AM, Lau JT, Hui AY, Wong VW, Sung JJ. Evaluation of MELD for prediction of mortality in decompensated chronic hepatitis B. *Am J Gastroenterol.* 2006; 101:1516-23.
- Wong GL, Hui AY, Wong VW, Chan FK, Sung JJ, Chan HL. A retrospective study on clinical features and prognostic factors of biopsy-proven primary biliary cirrhosis in Chinese patients. *Am J Gastroenterol.* 2005;100:2205-11.
- Huo TI, Huang YH, Lin HC, et al. Proposal of a modified Cancer of the Liver Italian Program staging system based on the MELD for patients with HCC undergoing loco-regional therapy. *Am J Gastroenterol.* 2006;101:975-82.
- Huo TI, Huang YH, Su CW, et al. Validation of the HCC-MELD for dropout probability in patients with small hepatocellular carcinoma undergoing locoregional therapy. *Clin Transplant.* 2008 [Epub ahead of print].
- Freeman RB, Edwards EB, Harper AM. Comparison of liver transplant waiting list removal rates among patients with chronic and malignant liver diseases. *Am J Transpl.* 2006; 6:1416-21. **Develops the concept of a continuous score for prioritizing HCC candidates using MELD and tumor variables.*
- Yan G, Duan Y, Ruan L, Chao T, Yang Y. A study on the relationship between ultrasonographic score and clinical score (MELD, CPT) in cirrhosis. *Hepatogastroenterology.* 2005;52:1329-33.
- Ripoll C, Banares R, Rincon D, et al. Influence of hepatic venous pressure gradient on the prediction of survival of patients with cirrhosis in the MELD era. *Hepatology.* 2005; 42:793-801.
- Bilaj F, Hyslop WB, Rivero H, et al. MR imaging findings in autoimmune hepatitis: correlation with clinical staging. *Radiology.* 2005;236:896-902.
- Sorrell JH, Zolnikov BJ, Sharma A, Jinnai I. Cognitive impairment in people diagnosed with end-stage liver disease evaluated for liver transplantation. *Psychiatry Clin Neurosci.* 2006;60:174-81.

32. Panzera P, Cicco G, Memeo R, et al. MELD predictive value of alterations of brain perfusion during liver transplantation. *Transplant Proc.* 2005;37:2622-5.
33. Harrod-Kim P, Waldman DL. Abnormal portal venous flow at sonography predicts reduced survival after TIPS creation. *J Vasc Interv Radiol.* 2005;16:1459-64.
34. Tandon P, Tsuyuki RT, Mitchell L, et al. The effect of 1 month of therapy with midodrine, octreotide-LAR and albumin in refractory ascites: a pilot study. *Liver Int.* 2008 [Epub ahead of print].
35. Lo Iacono O, Rincón D, Hernando A, et al. Serum levels of soluble vascular cell adhesion molecule are related to hyperdynamic circulation in patients with liver cirrhosis. *Liver Int.* 2008 [Epub ahead of print].
36. Dias Teixeira MC, de Fatima Gomes de Sa Ribeiro M, Strauss E. A new insight into the differences among non-cirrhotic and cirrhotic patients using the liver disease quality of life instrument (LDQOL). *Ann Hepatol.* 2005;4:264-71.
37. du Cheyron D, Bouchet B, Parienti JJ, Ramakers M, Charbonneau P. The attributable mortality of acute renal failure in critically ill patients with liver cirrhosis. *Intensive Care Med.* 2005;31:1693-9.
38. Terra C, Guevara M, Torre A, et al. Renal failure in patients with cirrhosis and sepsis unrelated to spontaneous bacterial peritonitis: value of MELD score. *Gastroenterology.* 2005; 129:1944-53.
39. Everson GT, Trotter J, Forman L, et al. Treatment of advanced hepatitis C with a low accelerating dosage regimen of antiviral therapy. *Hepatology.* 2005;42:255-62. *Uses MELD score to stratify patients receiving interferon therapy.
40. Huo TI, Lin HC, Huang YH, et al. The MELD-based Japan Integrated Scoring system may have a better predictive ability for patients with hepatocellular carcinoma undergoing locoregional therapy. *Cancer.* 2006;107:141-8.
41. Kalaitzakis E, Wallskog J, Björnsson E. Abstinence in patients with alcoholic liver cirrhosis: A follow-up study. *Hepatol Res.* 2008 [Epub ahead of print].
42. Wagholikar GD, Lee KH, Pandey D, Leong SO, Singh R, Tan KC. Pre-transplant optimization by Molecular Adsorbent Recirculating System in patients with severely decompensated chronic liver disease. *Indian J Gastroenterol.* 2007;26:110-2.
43. Kremers WK, van Ijperen M, Kim RW, et al. MELD score as a predictor of pre and post transplant survival in OPTN/ UNOS status 1 patients. *Hepatology.* 2004;39:764-9. *Indicates the MELD is a good predictor of mortality for patients with non acetaminophen acute liver failure.
44. Hassanein TI, Tofteng F, Brown RS, et al. Randomized controlled study of extracorporeal albumin dialysis for hepatic encephalopathy in advanced cirrhosis. *Hepatology.* 2007;46:1853-62.
45. Saberifiroozi M, Serati AR, Malekhosseini SA, et al. Analysis of patients listed for liver transplantation in Shiraz, Iran. *Indian J Gastroenterol.* 2006;25:11-3.
46. Vilatoba M, Eckhoff DE, Contreras JL. [Recipient selection for liver transplantation]. *Revista de Investigacion Clinica.* 2005;57:244-51.
47. McCormick A, Sultan J. Liver transplantation--patient selection and timing. *Med J Malaysia.* 2005;60:83-7.
48. Merion RM, Wolfe RA, Dystra DM, Leichtman AB, Gillespie B, Held PJ. Longitudinal assessment of mortality risk among candidates for liver transplantation. *Liver Transpl.* 2003; 9:19-21.
49. Bambha K, Kim WR, Kremers WK, et al. Predicting survival among patients listed for liver transplantation: an assessment of serial MELD measurements. *Am J Transplant.* 2004; 4:1798-804.
50. Giannini EG, Rizzo D, Caglieri S, Testa R. Longitudinal modifications of the MELD score have prognostic meaning in patients with liver cirrhosis. *J Clin Gastroenterol.* 2005; 39:912-4.
51. Gheorghe L, Iacob S, Iacob R, Gheorghe C, Popescu I. Variation of the MELD score as a predictor of death on the waiting list for liver transplantation. *J Gastrointest Liver Dis.* 2007;16:267-72.
52. Bellier C, Bert F, Durand F, et al. Risk factors for Enterobacteriaceae bacteremia after liver transplantation. *Transpl Int.* 2008 [Epub ahead of print].
53. Khettry U, Azabdaftari G, Simpson MA, et al. Impact of MELD scoring system on pathological findings at and after liver transplantation. *Liver Transpl.* 2006;12:958-65.
54. Ragni MV, Egthesad B, Schlesinger KW, Dvorchik I, Fung JJ. Pretransplant survival is shorter in HIV-positive than HIV-negative subjects with end-stage liver disease. *Liver Transpl.* 2005;11:1425-30.
55. Kim WR, Therneau TM, Benson JT, et al. Deaths on the liver transplant waiting list: an analysis of competing risks. *Hepatology.* 2006;43:345-51. *Competing risks analysis is an important concept when analyzing waiting list outcomes.
56. Jacquelinet C, Audry B, Pessione F, Antoine C, Loty B, Calmus Y. [Rules for allocation of livers for transplantation]. *Presse Med.* 2008 [Epub ahead of print].
57. Adler M, De Gendt E, Vereerstraeten P, et al. Value of the MELD score for the assessment of pre- and post-liver transplantation survival. *Transplant Proc.* 2005;37:2863-4.
58. Silberhumer GR, Hetz H, Rasoul-Rockenschaub S, et al. Is MELD score sufficient to predict not only death on waiting list, but also posttransplant survival? *Transplant Int.* 2006; 19:275-81.
59. Nagler E, Van Vlierberghe H, Colle I, Troisi R, de Hemptinne B. Impact of MELD on short-term and long-term outcome following liver transplantation: a European perspective. *Eur J Gastroenterol Hepatol.* 2005;17:849-56.
60. Ekka-Zohar A, Zitser-Gurevich Y, Mandel M, et al. Graft survival and its determinants: a 3 year national experience with liver transplantation in Israel. *Isr Med Assoc J.* 2006; 8:400-5.
61. Habib S, Berk B, Chang CC, et al. MELD and prediction of post-liver transplantation survival. *Liver Transpl.* 2006;12: 440-7.
62. Ekka-Zohar A, Zitser-Gurevich Y, Mandel M, et al. Graft survival and its determinants: a 3 year national experience with liver transplantation in Israel. *Isr Med Assoc J.* 2006; 8:400-5.
63. Sumskiene J, Kupcinskis L, Pundzius J, Sumskas L. Prognostic factors for short and long-term survival in patients selected for liver transplantation. *Medicina (Kaunas).* 2005;41:39-46.
64. Brandsaeter B, Friman S, Broome U, et al. Outcome following liver transplantation for primary sclerosing cholangitis in the Nordic countries. *Scand J Gastroenterol.* 2003; 38: 1176-83.
65. Onaca N, Levy MF, Ueno T, et al. An outcome comparison between primary liver transplantation and retransplantation based on the pretransplant MELD score. *Transpl Int.* 2006;19:282-7.
66. Tsui TY, Scherer MN, Schnitzbauer AA, Schliitt HJ, Obed A. Adult living donor liver transplantation: body mass index and MELD score of recipients are independent risk factors for hospital mortality. *Langenbecks Arch Surg.* 2008 [Epub ahead of print].
67. Avolio AW, Agnes S, Gasbarrini A, Nure E, Siciliano M, Castagneto M. Prognostic value of MELD score and donor quality in liver transplantation: implications for the donor recipient match. *Transplant Proc.* 2006;38:1059-62.
68. Renz JF, Kin C, Kinkhabwala M, et al. Utilization of extended donor criteria liver allografts maximizes donor use and patient access to liver transplantation. *Ann Surg.* 2005; 242:556-63.
69. Xia VW, Du B, Braunfeld M, et al. Preoperative characteristics and intraoperative transfusion and vasopressor requirements in patients with low vs. high MELD scores. *Liver Transpl.* 2006;12:614-20.
70. Frasco PE, Poterack KA, Hentz JG, Mulligan DC. A comparison of transfusion requirements between living donation and cadaveric donation liver transplantation: relationship to MELD score and baseline coagulation status. *Anesth Analg.* 2005;101:30-7.
71. Carmiel-Haggai M, Fiel MI, Gaddipati HC, et al. Recurrent hepatitis C after retransplantation: factors affecting graft and patient outcome. *Liver Transpl.* 2005;11:1567-73.
72. Umphrey LG, Hurst RT, Eleid MF, et al. Preoperative dobutamine stress echocardiographic findings and subsequent short-term adverse cardiac events after orthotopic liver transplantation. *Liver Transpl.* 2008;14:886-92.
73. Axelrod DA, Koffron AJ, Baker T, et al. The economic impact of MELD on liver transplant centers. *Am J Transplant.* 2005; 5:2297-301.
74. Lisman T, van Leeuwen Y, Adelmeijer J, et al. Interlaboratory variability in assessment of the model of end-stage liver disease score. *Liver Int.* 2008 [Epub ahead of print].
75. Trotter JF, Olson J, Lefkowitz J, Smith AD, Arjal R, Kenison J. Changes in international normalized ratio (INR) and MELD based on selection of clinical laboratory. *Am J Transplant.* 2007;7:1624-8.