Intensive Care Management of Acute Liver Failure: Considerations While Awaiting Liver Transplantation

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Abstract
Acute liver failure is a unique clinical phenomenon characterized by abrupt deterioration in liver function and altered mentation. The development of high-grade encephalopathy and multisystem organ dysfunction herald poor prognosis. Etiologic-specific treatments and supportive measures are routinely employed; however, liver transplantation remains the only chance for cure in those who do not spontaneously recover. The utility of artificial and bioartificial assist therapies as supportive care—to allow time for hepatic recovery or as a bridge to liver transplantation—has been examined but studies have been small, with mixed results. Given the severity of derangements, intensive critical care is needed to successfully bridge patients to transplant, and evaluation of candidates occurs rapidly in parallel with serial reassessments of operative fitness. Psychosocial assessment is often suboptimal and relative contraindications to transplant, such as ventilator-dependence may be overlooked. While often employed to guide evaluation, no single prognostic model discriminates those who will spontaneously recover and those who will require transplant. The purpose of this review will be to summarize approaches in critical care, prognostic modeling, and medical evaluation of the acute liver failure transplant candidate.

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Introduction
Acute liver failure (ALF) is an infrequent condition in which rapid deterioration of liver function results in altered mentation and coagulopathy in individuals without pre-existing liver disease.1 Approximately 2,000 cases are thought to occur in the USA annually.2 Studies estimating ALF prevalence and incidence in the European Union are lacking; however, a review of the European Liver Transplant Registry identified ALF as the primary indication for transplant in only 8% of transplants performed over a 20-year period.3,4 A key element in identifying ALF is the absence of preexisting liver disease (distinguishing it from acute-on-chronic liver disease) and international societies have guided definitions for severity of liver injury.5

A widely accepted working definition of ALF includes: an international normalized ratio (INR) >1.5 and any degree of mental alteration (encephalopathy) in a subject without preexisting cirrhosis and with illness of <26 weeks’ duration.6 Currently, overall short-term survival (2 years) in those undergoing transplantation approaches 90%.7 Liver transplantation (LT) remains the only definitive treatment for subjects who fail to spontaneously recover or respond to etiology-specific medical therapy. Moreover, patients are often managed in an intensive care unit (ICU) setting while awaiting LT. The approach to the diagnostic evaluation of ALF has been described elsewhere.8,9 The purpose of this review will be to summarize recent advances in critical care management, prognostic modeling, and evaluation of the ALF transplant candidate.

Etiology-specific therapy
Etiology-specific treatment for ALF is often administered in an ICU setting (Table 1).10 Both acetaminophen and non-acetaminophen drug-induced liver injury, acute viral hepatitis, autoimmune hepatitis, Wilson’s Disease, and vascular disorders represent examples for which etiology-specific therapy can be attempted to regain parenchymal function.11

In instances of known or suspected acetaminophen ingestion, N-acetylcysteine (NAC) can be given as an antidote orally or intravenously; its efficacy and safety is well-established.12,13 NAC should be given as early as possible, most commonly via intravenous administration (loading dose of 150 mg/kg in 5% dextrose over 15 m; maintenance dose of 50 mg/kg given over 4 h, followed by 100 mg/kg administered over 16 h or 6 mg/kg/h) but may have value 48 h or more after ingestion.13 At this time, there is no consensus as to whether a standard 72-h period is optimal or the further continuation based on clinical course. Medications other than acetaminophen may be implicated as the causative agent for ALF.15 These are generally thought to represent examples of idiosyncratic drug hepatotoxicity. Investigation has suggested that a course of NAC (dosage outlined above) may abrogate non-acetaminophen-related drug-induced liver injury.16 In a recent randomized case-control study of 80 subjects with non-acetaminophen-induced liver failure, use of NAC was associated with a significantly lower mortality

Keywords: Acute liver failure; Thromboelastography; Intracranial hypertension; Liver assist therapy; Transplantation.

Abbreviations: ALF, acute liver failure; CPI, cerebral perfusion pressure; ICH, intracranial hypertension; ICP, intracranial pressure; ICU, intensive care unit; INR, international normalized ratio; KCC, King’s College Criteria; LT, liver transplantation; MAP, mean arterial pressure; MELD, model for end-stage liver disease; NAC, N-acetylcysteine; rFVIIa, recombinant activated factor seven.© 2019 Authors. This article has been published under the terms of Creative Commons Attribution-NonCommercial 4.0 International (CC BY-NC 4.0), which permits noncommercial unrestricted use, distribution, and reproduction in any medium, provided that the following statement is provided: "This article has been published in Journal of Clinical and Translational Hepatology at DOI: 10.14218/JCTH.2019.00032 and can also be viewed on the Journal's website at http://www.jcthnet.com".
Table 1. Etiology-specific therapies for ALF. Directed therapy of ALF when etiology is known may increase transplant-free survival.

| Etiology                                      | Therapy a                  |
|----------------------------------------------|---------------------------|
| Acetaminophen and non-acetaminophen drug-induced liver injury | NAC: loading dose is 150 mg/kg in 5% dextrose over 15 m; maintenance dose is 50 mg/kg given over 4 h followed by 100 mg/kg administered over 16 h or 6 mg/kg/h |
| Hepatitis B                                  | Antiviral therapy with nucleos(t)ide analogue: entecavir or tenofovir |
| Herpes (herpes simplex virus)                 | Acyclovir 5–10 mg/kg intravenous every 8 h for at least 7 days |
| Autoimmune hepatitis                         | Prednisone 40–60 mg PO daily |
| Wilson’s disease                              | Albumin dialysis, continuous hemofiltration, plasmapheresis, or plasma exchange |
| Budd-Chiari syndrome                         | Attempts at hepatic vein recanalization with transjugular or direct portosystemic shunt; systemic anticoagulation |
| Mushroom Poisoning (Amanita phalloides) ingestion | Gastric lavage, activated charcoal NAC; silymarin |
| Acute fatty liver of pregnancy                | Delivery of the fetus |

aEtiology-directed therapy is administered concomitantly with intensive care support and medical evaluation for transplant.

Abbreviations: ALF, acute liver failure; NAC, N-acetylcysteine.

(28% in NAC group vs. 53% control, p = 0.02) and shorter length of stay.11

Acute viral hepatitis acquisition with hepatitis A, B, D or E may progress to ALF.7 In instances of serologically identified hepatitis A and E, management is largely supportive. Nucleos(t)ide analogue therapy can be attempted in acute hepatitis B or D; however, reversal of severe injury with early agents (e.g. lamivudine) has rarely been achieved.18 Improved efficacy has been reported with new-generation therapy (entecavir and tenofovir); a recent investigation found higher short-term efficacy with tenofovir in cases of acute-on-chronic liver failure with suspected hepatitis B virus reactivation.19 Hepatitis C virus infection is an uncommon cause of ALF but has been reported in both immunocompromised and immunocompetent subjects.20,21 Empiric treatment should be considered utilizing acyclovir (5–10 mg/kg every 8 h for at least 7 days). Cytomegalovirus, varicella-zoster, Epstein-Barr, and adenovirus have all been reported as viral causes of ALF, albeit rare, particularly in pediatric or immunocompromised populations.22,24

Autoimmune hepatitis may present with ALF.6 Initiation of steroid therapy may be considered for some patients (prednisone starting at 40–60 mg/day). However, in some patients this may be deleterious, and transplant evaluation should not be delayed while awaiting a response.25

Wilson’s disease, an inherited metabolic disorder in copper transport, is a relatively infrequent cause of ALF.26 This fulminant presentation is considered uniformly fatal without transplantation. Treatment to acutely lower serum copper and to limit further hemolysis can include albumin dialysis, continuous hemofiltration, plasmapheresis, or plasma exchange; although, such copper lowering measures can be employed, recovery is very rare without LT.27

Vascular disorders of the liver may also be associated with ALF. Budd-Chiari syndrome, characterized by acute hepatic outflow obstruction, may precipitate severe acute liver injury requiring consideration of hepatic vein recanalization or shunting; preexisting hypercoagulable state must be investigated and mandates treatment with systemic anticoagulation.28 Acute ischemic injury to the liver may cause ALF; typically in patients with preexisting cardiac dysfunction rendering hepatic tissue sensitive to minor reductions in systolic blood pressure—treatment in these instances generally focuses on restoration of normal hemodynamics.

Mushroom poisoning (most commonly associated with Amanita phalloides) may cause ALF. Typically, gastrointestinal symptoms including profuse vomiting and diarrhea occur after mushroom ingestion. Gastric lavage, supportive measures, as well as various agents with antidotal properties (benzylpenicillin, NAC, silymarin) can be considered.29

Acute fatty liver of pregnancy is characterized by extensive steatosis in the third trimester of pregnancy. Development, in part, is related to inheritance patterns in mother and fetus, particularly of long-chain 3-hydroxyacyl-CoA dehydrogenase (also referred to as LCHAD) deficiency, which is linked to fetal fatty oxidation defects.30 Prompt delivery of the fetus is the preferred management and often reverses liver injury.

Coagulopathy

In ALF, prothrombin time and its derivative INR are elevated and are considered predictors of increased bleeding risk—though magnitude of effect is undefined. Previous investigation has found a concomitant and proportional reduction in plasma levels of both procoagulants and natural anticoagulant proteins, in conjunction with a significant elevation in plasma levels of factors-VIII (referred to as FVIII) and Von Willebrand factor, culminating in an overall efficient, albeit reduced, thrombin generation capacity in comparison with healthy individuals.31 Global hemostasis as assessed with thromboelastography may be normal by several compensatory mechanisms, even in patients with markedly elevated INR.32 In the absence of overt bleeding, measures to correct the INR with plasma are not recommended and may potentially obscure trends marking prognosis.6

Vitamin K deficiency has been reported in patients with ALF33 and vitamin K (10 mg subcutaneously) can be administered (Table 2). Treatment of clotting factor deficiency is generally reserved for clinically significant bleeding or in preparation for a high-risk invasive procedure (e.g., intracranial pressure (ICP) monitor placement). Fresh frozen plasma infusion alone infrequently corrects a severely elevated INR and carries risk of volume overload and transfusion-related lung injury.24 Use of recombinant activated factor seven (referred to as rFVIIa) may be considered, as administration in combination with frozen fresh plasma provides temporary correction of coagulopathy without volume overload.35,36 Important
barriers to the rFVIIa utilization include availability, cost, and reports of thromboembolic phenomenon. There is no consensus regarding the prophylactic administration of platelets in thrombocytopenic patients with ALF and this is generally reserved for cases of overt bleeding or prior to invasive procedures.

Circulatory dysfunction/renal failure

ALF, similar to cirrhosis or sepsis, is characterized by low systemic vascular resistance. Diminished tone may compromise peripheral tissue oxygenation and promote multisystem organ failure. Maintenance of adequate mean arterial pressure (referred to as MAP) is of particular importance in ALF subjects with increased ICP to maintain cerebral perfusion pressure (CPP) [CPP equates to MAP minus ICP]. Intravascular volume depletion is common at ALF presentation and resuscitation is often required. Hypotensive patients with ALF should be resuscitated with normal saline and changed to half-normal saline containing 75 mEq/L sodium bicarbonate if acidic. Volume expanding solutions should contain dextrose to prevent hypoglycemia if present. Subjects remaining hypotensive despite volume resuscitation should be cultured for infection and considered for vasopressor support to maintain a MAP of at least 75 mmHg or a CPP of 60–80 mmHg. While no studies have defined the optimal vasopressor, norepinephrine is known to augment peripheral organ perfusion while minimizing tachycardia and preserving splanchnic/hepatic blood flow. Vasopressin and/or analogues may potentiate effects of norepinephrine and allow a decrease in its infusion rate, mitigating intense vasocostriction in peripheral tissues that may lead to bowel and/or limb ischemia.

Independent of vascular resistance, relative adrenal insufficiency has been commonly observed in over half of patients presenting with ALF. Although a mortality benefit with hydrocortisone usage in ALF subjects has not been demonstrated, utility as an adjunctive measure to reduce systemic vasopressor requirement has been demonstrated. Use of corticosteroids in the context of ALF must be individualized, taking into account etiology of liver injury and risk of infection.

Substantial renal dysfunction (either functional or as a result of acute tubular necrosis) may occur in greater than 50% of patients with ALF. Acute kidney injury more commonly occurs when acetaminophen is the etiology of injury and in the elderly. Renal failure contributes to morbidity and mortality and is associated with a poorer prognosis. When support is required, continuous renal replacement therapy is preferred over conventional hemodialysis, as the former has been shown in randomized trials to result in improved stability in cardiovascular and intracranial parameters compared to the latter. Avoidance of nephrotoxic agents, including intravenous contrast agents, is standard of care.

Encephalopathy/ICP management

Cerebral edema and intracranial hypertension (ICH) represent the most serious complications of ALF. Although pathogenesis is only partly understood, there is evidence that both systemic and local inflammation and circulating neurotoxins, including ammonia, play roles. Ammonia infusion causes systemic and local inflammation and circulating neurotoxins, including ammonia, play roles. Ammonia infusion causes cerebral edema in animal models, and an arterial ammonia level >200 µg/dL in humans is strongly associated with cerebral herniation. Encephalopathy can be precipitated by infection or may occur as a result of low systemic blood

| Organ system | Derangement | Supportive measures |
|--------------|-------------|---------------------|
| Hematologic  | Concomitant and proportional reduction in both procoagulants and anticoagulants Significant elevation in plasma levels of factors-VIII (FVIII) and Von Willebrand factor Reduced thrombin generation capacity | Vitamin K 10 mg intravenous x 1 considered in those with nutritional deficiency Fresh frozen plasma, platelets, and rFVIII reserved for active bleeding or invasive procedure |
| Cardiovascular | Low systemic vascular resistance Diminished tone compromising peripheral tissue oxygenation | Hypotensive patients resuscitated with normal saline and changed to half-normal saline containing 75 mEq/L sodium bicarbonate if acidic Volume expanding solutions with dextrose to prevent hypoglycemia Vasopressor (norepinephrine) support to maintain a mean arterial pressure >75 mmHg or a cerebral perfusion pressure of 60–80 mmHg |
| Renal | Prerenal kidney injury from diminished effective circulating volume, acute tubular necrosis, or reduced function | Continuous renal replacement therapy preferred over conventional hemodialysis |
| Neurologic | Systemic and local inflammation and circulating neurotoxins, including ammonia that promote cerebral edema and intracranial hypertension | Consideration of intracranial monitor placement\(^a\) Head of bed elevation Avoid endotracheal suction Mannitol |

\(^a\)Intracranial pressure monitoring placement is variable and based on center expertise. Abbreviation: ALF, acute liver failure.
pressure and vasodilatation. Incidence of cerebral edema and ICH in ALF is directly related to severity of hepatic encephalopathy, with episodes of ICH occurring between 65–75% in those with the highest grade.

As patients progress to higher grades of encephalopathy, intubation and mechanical ventilation are mandatory for airway protection. Propofol is routinely chosen for sedation because it may reduce cerebral blood flow. Frequent neurological evaluation for signs of ICH (e.g., sluggish pupillary reflexes and posturing) should be conducted. To reduce the incidence of ICH, patients should have the head elevated at 30 degrees and stimulation (including endotracheal suctioning) and pain should be minimized.

Invasive and noninvasive monitoring strategies have been utilized to assess progression of cerebral edema. Use of ICP monitoring devices is controversial but employed by many ICUs. The principal concern is for intracranial hemorrhage with ICP placement, particularly in ALF subjects with advanced encephalopathy and active infection. Experience is limited with noninvasive strategies, such as transcranial Doppler, jugular bulb oximetry, and pupillometry. If ICP monitoring is performed, both ICP and CPP are followed with a goal ICP below 20–25 mmHg and CPP maintained above 50–60 mmHg.

Based on the role of arterial ammonia in pathogenesis, investigation has focused on the use of lactulose administration in ALF. In a study of ALF patients who received lactulose compared to a matched group of patients who did not, a small increase in survival time was seen in those receiving lactulose, but there was no difference in the severity of encephalopathy or outcome. One concern regarding the use of lactulose in this setting is the precipitation of bowel gas with excessive usage, thereby complicating subsequent transplant surgery.

Osmotic agents, such as mannitol, can be effective in decreasing cerebral edema. Mannitol was found to transiently correct episodes of elevated ICP in ALF patients and also to improve survival. Administration of intravenous mannitol (1 g/kg over 30 min – 1.0 g/kg) is recommended as first-line therapy of ICH in patients with ALF but repeated doses may trigger serum hyperosmolality. In patients with ALF and severe hepatic encephalopathy, a controlled trial of the prophylactic induction of hypotension with hypertonic saline (to a serum sodium 145–155 mEq/L) suggested a lower incidence of ICH. Utilization of hypertonic saline as treatment for established ICH has not been studied. Barbiturate agents (thiopental or pentobarbital) may also be considered to decrease ICP that has been nonresponsive to other measures. Significant systemic hypotension may limit use and may necessitate vasopressor administration to maintain adequate mean arterial pressure. In addition, barbiturate clearance is markedly reduced in patients with ALF, potentially confounding neurological assessments. Hypothermia may prevent or control ICH in patients with ALF. Pilot studies suggest potential use (core temperature of 33–34°C) as a bridge to LT. Induced hypothermia has not been compared to normothermic conditions in a controlled trial and there is theoretical concern about a negative effect on hepatic regeneration.

Advanced systems

Over the years, several artificial and bioartificial liver assist therapies have been tested for utility as supportive care to allow time for hepatic recovery or as a bridge to LT. Owing to the relative rarity of the condition, clinical heterogeneity and complexity in administration, studies have been small with mixed results. In the fulmar trial examining use of an extracorporeal device in patients with ALF, there was no improvement in overall survival in the cohort but there was a trend of potential benefit in patients with acetaminophen as the etiology of ALF. Therapeutic plasma exchange has hypothesized applicability in ALF through removal of inhibitors of hepatic regeneration. In a multicenter randomized controlled trial, three 5-l sessions of therapeutic plasma exchange significantly improved survival, although the survival benefit was inferior to that seen with transplantation. Other forms of cellular therapy (including hepatocyte transplantation) have shown potential as future treatments for ALF but are not ready for routine clinical practice.

Recent society guidelines suggest consideration of available support systems in patients with ALF with an expected poor prognosis without transplantation but who have clear medical or psychiatric contraindications to surgery.

Prognostic models

A number of different systems are in current worldwide use to assess prognosis in patients with ALF (Table 3). All systems utilize admission laboratory measures (coagulopathy of highest weight) and recognize the development of encephalopathy and advanced age as markers of poor prognosis. One of the oldest and most utilized tools are the King’s College Criteria (KCC), the original iteration now used for 3 decades, being the first to recognize the importance of etiology of ALF/acetaminophen toxicity carrying higher chance of recovery than others. With over 3 decades of utilization, the KCC is noted to have high specificity (relatively few cases fulfilling criteria would be ‘unnecessarily transplanted’); however, it suffers from relatively low sensitivity (significant number of cases not fulfilling criteria would progress and die without earlier identification and consideration of LT). The latest model from the King’s College is a dynamic outcome prediction model and has been validated for use with ALF from acetaminophen toxicity. The model uses a number of clinical variables sequentially assessed from admission to 72 h and has demonstrated good discrimination between survivors and non-survivors.

Recently, the United States Acute Liver Failure Group prospectively enrolled over 1900 subjects with ALF, managed with and without transplantation. Investigators aimed to develop a model for ALF (all etiologies included) to predict transplant-free survival at 21 days. Clinical demographics and laboratory values were collected at enrollment and recorded serially up to 1 week. Variables of prognostic value adopted in the predictive model included admission comat grade, ALF etiology and vasopressor requirement, and admission INR and bilirubin values. The model correctly predicted outcome of illness in 66.3% of subjects, slightly outperforming KCC and the model for end-stage liver disease (referred to as MELD) score. Performance appeared best in patients with non-acetaminophen etiology and high-grade encephalopathy.

A number of other scoring systems have been proposed to identify candidates most at risk for death and need for LT. Non-liver specific indices, such as the sequential organ failure assessment which is widely used to quantify severity of multiorgan failure in other forms of critical illness, have...
been utilized with comparable performance to KCC in prediction of non-survival; although, their organ “non-specificity” compromises applicability in determining benefit with LT and use is limited.\textsuperscript{77,78} The MELD score, widely used for liver prioritization/allocation in chronic liver disease, has been investigated in ALF with similar performance to that of the KCC.\textsuperscript{79} An emerging theme in ALF prognostication is the need for individualized, dynamic assessments as opposed to historically static ones at presentation. An investigation utilizing the ALF early dynamic model followed clinical variables over 3 days and outperformed KCC and MELD in predicting outcome in an observational study of 380 subjects.\textsuperscript{80}

Recently, investigation has focused on the use of biomarkers in prediction of outcome with ALF. Circulating blood levels of caspase-cleaved and uncleaved cytokeratin K18 (referred to as CK18), an apoptosis cell death marker, has shown promise.\textsuperscript{81} Subjects with spontaneous recovery of ALF have been demonstrated to have higher levels of caspase activation than in subjects who required transplant or died.\textsuperscript{82} Circulating levels in concert with standard blood measures of coagulation and renal function demonstrated superior sensitivity and specificity to KCC in predicting ALF outcome; although, further studies are needed.\textsuperscript{83} HLA-DR monocyte expression has been identified as a potential biomarker of ALF severity and outcome in acetaminophen-related ALF.\textsuperscript{84}

A number of shortcomings may be present despite thorough evaluation. Living donor donation for ALF (particularly in areas of the world where access to cadaveric grafts may be low) raises ethical concern of coercion for potentially compatible family members. In addition, given the acuity of illness and associated multisystemic organ dysfunction, determining futility or when the patient is too sick to transplant may be a clinical challenge, as widely accepted relative contraindications to LT (bacteremia or ventilatory-dependent respiratory failure) may be overlooked if responding to critical care measures.

Adult ALF subjects who meet program criteria for listing in the USA are granted status 1A, placing the patient at the highest priority on the waiting list. In order to be listed as status 1A, the ALF candidate must be at least 18 years-old, with life expectancy of less than 7 days. Additionally, he or she must be in the ICU and meet at least 1 of the following criteria: 1) ventilator dependence; 2) requiring renal replacement therapy; 3) INR greater than 2.0.\textsuperscript{88} The mean time between listing and LT is 2 days in the USA.\textsuperscript{89}

Currently, ALF accounts for approximately 8% of all liver transplants, as per data from the Scientific Registry of Transplant Recipients and the European Liver Transplant Registry, with 1-year survival rates of 84% in the USA and 79% in Europe, respectively.\textsuperscript{4,90} Outcomes are inferior when compared with patients receiving a transplant due to chronic liver disease; though survival rates even out beyond 1 year.\textsuperscript{91}

In a series of over 1400 patients from the United Network for Organ Sharing database of LT for ALF, body mass index greater than 30, serum creatinine greater than 2 mg/dL, recipient age greater than 50 years, and a history of life support were independent factors of poor post-transplant outcomes, with a survival of less than 50% in recipients with all factors pretransplant.\textsuperscript{92} An analysis of the European Liver Transplant Registry identified male gender, older age of the donor (>60 years-old), older age of the recipient (>50 years-old), incompatible graft, and reduced graft size as predictors of poor prognosis.\textsuperscript{4}

**Conclusions**

ALF represents abrupt deterioration in hepatic function from diverse etiologies. Associated multisystem organ dysfunction and dense encephalopathy with chance for progression to brainstem herniation mandates prompt recognition and transfer to a liver transplant center. While etiologic-specific

### Table 3. Prognostic models for ALF

| Study          | Prognostic model | ALF etiology | Subjects studied, n | Subjects died or transplanted | Sensitivity | Specificity | AUROC |
|----------------|------------------|--------------|---------------------|-------------------------------|-------------|-------------|-------|
| McPhail et al.\textsuperscript{79} | KCC          | All          | 2153                | Unknown                       | 0.55        | 0.79        | 0.76  |
| McPhail et al.\textsuperscript{79} | MELD         | All          | 2153                | Unknown                       | 0.74        | 0.67        | 0.78  |
| Koch et al.\textsuperscript{76}   | ALFSG        | All          | 1974                | 987 (50%)                     | Not reported| Not reported| 0.84  |
| Cholongitas et al.\textsuperscript{78} | SOFA       | APAP only    | 125                 | 58 (46%)                      | 0.67        | 0.80        | 0.79  |

**Abbreviations:** ALF, acute liver failure; ALFSG, Acute Liver Failure Study Group; APAP, acetaminophen; AUROC, area under the receiver operating curve; KCC, King's College Criteria; MELD, model for end-stage liver disease score; SOFA, sequential organ failure assessment score.
Seetharam A.: Liver transplantation for acute liver failure treatments and a number of systemic supportive measures can be employed. LT remains the only chance for cure in those who do not spontaneously recover. While often employed to guide evaluation, no single prognostic model discriminates those who will spontaneously recover and those who will require transplant. Given the severity of derangements, intensive critical care is needed to bridge ALF patients to transplant; furthermore, evaluation of potential candidates must occur rapidly and in the context of serial reassessment of fitness for surgery.

Conflict of interest
The author has no conflict of interest related to this publication.

Author contributions
Contributed to data review and writing of manuscript (AS).

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