



CME: Hepatology

Management of acute decompensated cirrhosis[☆]

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ABSTRACT

Acute decompensated cirrhosis (AD) refers to the development of ascites, encephalopathy, gastrointestinal haemorrhage, or any combination of these disorders in a patient with known or previously undiagnosed advanced chronic liver disease. It carries a significant mortality, particularly if associated with organ failure (acute on chronic liver failure (ACLF)). Admissions with AD have increased by 50% over the last decade, and liver-related deaths have increased by 64% in the last 20 years. UK-wide reports and audits, including by the National Confidential Enquiry into Patient Outcome and Death, have revealed unwarranted variation in care and outcomes for patients with AD. This article summarises the management of patients admitted with AD, including the British Society of Gastroenterology (BSG) / British Association for the Study of the Liver (BASL) / Society of Acute Medicine (SAM) decompensated cirrhosis care bundle, developed for completion in the first 6 h of admission to standardise care for these complex patients.

Introduction

Acute decompensated cirrhosis (AD) refers to the development of ascites, encephalopathy and/or gastrointestinal haemorrhage in patients with cirrhosis.¹ It is associated with significant (5–20%) 28-day mortality.²

There has been a 50% increase in liver-related admissions in the last decade and a 64% increase in premature liver-related deaths in the last 20 years.³ Almost half of liver disease deaths occur in people of working age⁴ and liver disease mortality correlates with degree of social deprivation.⁴

Following the 2013 National Confidential Enquiry into Patient Outcome and Death (NCEPOD), which found that only 47% of patients admitted with AD received ‘good’ care,⁵ a decompensated cirrhosis care bundle was developed to guide management in the first 24 h of admission.⁶ This improved care,⁷ but the benefit was limited by low uptake (completed in only 11.4% of AD admissions in a nationwide audit).⁸

Following further recent NCEPOD review,³ the care bundle was revised (see Fig. 1)⁹ and strategies proposed to improve its use.⁹ It provides a structured approach to managing AD across the emergency department and admissions unit, and should be completed in all patients presenting with AD within the first 6 h of admission.

Patients admitted with AD should be cared for on a specialist gastroenterology or hepatology ward and should be seen by a specialist, ideally within 24 h of admission.

History and examination

A careful structured history and examination are important in helping to determine the stage of the patient’s disease, the potential precipitant of the decompensation, and to establish an appropriate ceiling of care. Key points in the history and examination from a patient with AD are summarised in Fig. 2.

Investigations

Initial investigations and treatments are summarised in the care bundle⁹ (Fig. 1). A full non-invasive liver screen (including liver autoantibodies, ferritin and transferrin saturations, immunoglobulins, hepatitis B/C and HIV serology) is only required if not previously performed, but repeat viral hepatitis screen may be appropriate, including hepatitis A and E, particularly if liver enzymes are significantly raised.

A septic screen including an ascitic tap (sent for white cell/polymorph count (PMN) and culture, protein and albumin) should be performed to rule out infection, irrespective of inflammatory markers. Spontaneous bacterial peritonitis (SBP) occurs in approximately 10% of patients admitted with cirrhosis and is frequently asymptomatic, despite carrying a high mortality if treatment is delayed.¹⁰ Diagnosis is based on PMN cell count $>250/\text{mm}^3$ or $0.25 \times 10^9/\text{L}$ ascitic fluid. Coagulopathy and/or thrombocytopenia are not a contraindication to an ascitic tap and checking/correction of

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Patient details		  	
Decompensated Cirrhosis Care Bundle - First 6 hours			
ALL patients presenting with symptoms/signs of decompensated cirrhosis¹	→	<i>Bloods:</i> FBC, U/Es, clotting, calcium, phosphate, CRP, lactate, glucose ☐ <i>Septic screen:</i> CXR ☐ Urine dipstick ☐ Blood cultures ☐ <i>Clinical ascites:</i> Ascitic tap ESSENTIAL² ; send for MCS and protein ☐ Request USS abdomen including Doppler of hepatic and portal vein ☐ VTE prophylaxis, unless platelets <50 or active bleeding ☐ Referral to dietetics made +/- oral nutritional supplements prescribed ☐ Specialist review: refer to GI/liver team at earliest opportunity ☐	
Ongoing alcohol intake? <i>(if NO- move to next section)</i>	→	IV Pabrinex/Thiamine as per hospital guidelines ☐ Commence CIWA or GMAWS according to hospital guidelines ☐	N/A ☐
Spontaneous Bacterial Peritonitis? <i>(if NO- move to next section)</i>	→	<i>(Diagnosis: Ascitic neutrophils >250/mm³ or >0.25 x 10⁹/L)</i> Prescribe 1.5g/kg of 20% Human Albumin Solution (HAS) ☐ Antibiotics as per hospital guidelines ☐	N/A ☐
Acute Kidney Injury as per KDIGO criteria³ <i>(if NO- move to next section)</i>	→	Suspend all diuretics and nephrotoxic drugs ☐ Fluid resuscitate with crystalloid in 250ml boluses ☐ Strict urine output monitoring ☐ At 6 hours, if deteriorating despite this, obtain senior review and consider escalation to ITU/HDU.	N/A ☐
GI bleeding AND varices suspected? <i>(if NO- move to next section)</i>	→	Target Hb 7-8 g/L, but if massive bleeding aim for Hb >8g/L ☐ <i>It is not recommended to routinely correct INR/APTT with blood products (unless on anticoagulants).</i> Terlipressin: if no clear contraindications ⁴ prescribe 2mg stat IV followed by 2mg QDS ☐ <i>(If contraindication to Terlipressin, contact on-call GI bleed team and consider dose reduction/alternate agent⁴)</i> Prescribe prophylactic antibiotics as per hospital guidelines ☐	N/A ☐
Symptoms/signs of Hepatic Encephalopathy?	→	Lactulose 20-30mls QDS or phosphate enema ☐ If clinical concern, for CT head to rule out a subdural haematoma ☐	N/A ☐
Name Grade Date Time ^(1,2,3,4) Please turnover for additional information			

Fig. 1. British Society of Gastroenterology, British Association for the Study of the Liver and Society of Acute Medicine Decompensated Cirrhosis Care Bundle – first 6 h.⁹

History	
Background	- Known advanced liver disease? aetiology & mode of diagnosis (clinical/biopsy/TE/imaging) - Previous decompensation episodes and treatment - Last endoscopy – endoscopic treatment/beta blocker - Last surveillance liver USS: date,result - Alcohol history: current use, dependence (AUDIT score), prior history, abstinence/relapse - Functional status/frailty score pre-admission - Advanced care planning in place? Resuscitation status? - Referred or listed for liver transplant?
History Presenting Complaint	- Confusion/poor concentration/sleep-wake disturbance - Bowel frequency - Melaena/haematemesis - Swelling (legs, abdomen)/breathlessness - Appetite and oral intake
Past Medical History	- Other significant comorbidities
Drug History	- Recent antibiotics or new medications - Over-the-counter or non-prescribed drugs/supplements - Medication concordance
Systemic Enquiry	- Infective symptoms: cough, breathlessness, back pain, joint pain/swelling, diarrhoea, urinary symptoms, toothache, abdominal pain - Weight loss
Examination	
Signs Of Decompensation	- Jaundice - Ascites (e.g., shifting dullness)/peripheral oedema - Encephalopathy signs: GCS, asterixis - Rectal exam: melaena
Liver disease stigmata	- Spider naevi - Gynaecomastia - Palmar erythema –Splenomegaly
Signs Of Infection	- Respiratory: chest signs - Cardiac: murmurs - Skin: cellulitis, ulcers, pressure sores - Joints: swollen/red joints - Abdomen: tenderness - Oral: poor dentition
Wernicke's / Alcohol Brain Injury	- Nystagmus - Ophthalmoplegia - Cerebellar signs - Gait disturbance
Nutritional Status	- Evidence of sarcopenia - Weight/BMI (adjusted for ascites/oedema)

Fig. 2. History and examination in patients presenting with acute decompensated.

coagulation is not required prior to the procedure.¹¹

A Doppler ultrasound scan of the abdomen should be performed to look for hepatic/portal vein thrombosis, hepatocellular carcinoma and ascites.

A CT head scan should be considered in patients with reduced consciousness or focal neurology, even if encephalopathy is suspected, particularly if there is a history of head injury or falls.

Management

Alcohol

Patients with alcohol-related liver disease (ARLD) who continue to consume alcohol should be assessed for signs and symptoms of

Wernicke's encephalopathy (WE) and be prescribed parenteral thiamine (for at least 72 h in those with features of WE, or 24 h for those without features of WE)⁹ before stepping down to oral thiamine.¹²

Patients at risk of alcohol withdrawal should be assessed with a validated tool such as the Clinical Institute Withdrawal Assessment-Alcohol scale (CIWA)¹³ or the Glasgow Modified Assessment and Management of Alcohol tool (GMAWS)¹⁴ and treated with a symptom-triggered regimen of benzodiazepines. Chlordiazepoxide can accumulate in people with liver impairment/failure; consider using reduced doses or lorazepam (shorter acting and less likely to accumulate in patients with advanced liver disease) as an alternative or in combination.

All patients with alcohol dependence should be assessed by the alcohol care team and receive ongoing support with alcohol reduction/abstinence.¹⁵

Infections

Liver disease and alcohol both increase susceptibility to infections – patients may not display typical signs of infection or have significantly raised inflammatory markers, so a high index of suspicion is required.¹⁶ Multidrug-resistant organisms are a growing problem, therefore empirical antibiotics are not recommended.¹ If suspicion of infection or signs of sepsis are present, a thorough history, examination and septic screen should be performed, followed by broad-spectrum antibiotics based on likely source in line with local guidelines, early de-escalation to narrow-spectrum antibiotics based on cultures, and cessation of antibiotics once infection is excluded.

Patients with suspected SBP should have prompt broad-spectrum antibiotics according to local guidelines, modified according to ascitic fluid culture results. SBP increases the risk of hepatorenal syndrome, therefore intravenous 20% albumin should be administered (1.5 g/kg albumin at diagnosis and 1 g/kg at 72 h).¹ Routine albumin is not recommended in patients with other sources of infection.¹⁷

Patients treated for SBP should start secondary prophylaxis on completing their course of antibiotics, according to local protocol (eg cotrimoxazole 960 mg daily).¹⁸

Acute kidney injury (AKI)

AKI affects 20–30% of patients with AD, and it is associated with a poor outcome.¹⁹ The majority of AKI is pre-renal (47%) with 32% acute tubular necrosis (ATN) and 23% hepatorenal syndrome (HRS) (<1% post-renal).²⁰

Diuretics, beta blockers and other nephrotoxic medication should be suspended and hypovolaemia corrected with crystalloids in 250 mL boluses, aiming for urine output (UO) >0.5 mL/kg/hr.²⁰ Early treatment of infection is key. In patients with hypotension, UO <0.5 mL/kg/hr, worsening lactate or development of respiratory failure following 6 h of fluid resuscitation, consider escalation to a high dependency unit. Renal replacement therapy should be considered on a case-by-case basis – only approximately 25% recover following dialysis.²¹

HRS-AKI describes AKI associated with haemodynamic changes in patients with advanced liver disease and ascites. Without treatment, it carries a mortality approaching 100%.¹⁰ It is diagnosed in patients with AD and ascites, following volume expansion, treatment for any infection, and suspension of nephrotoxic medication, in the absence of structural renal disease (ie no proteinuria or haematuria, and normal kidneys on ultrasound imaging).¹⁰ It can co-exist with other causes of AKI. Treatment is with terlipressin (2–8 mg daily in divided doses or as continuous infusion, titrated to response) and 20% albumin 100 mL twice daily.¹ Terlipressin should be used with caution in older patients (>70 years old), and patients with peripheral vascular disease or ischaemic heart disease (as it can cause ischaemia),²² as well as those with severe renal or liver impairment, due to increased risk of respiratory complications and sepsis.²³

Acute variceal haemorrhage (AVH)

Acute upper gastrointestinal bleeding in people with known cirrhosis, or first presentation AD, should be treated as AVH until proven otherwise. AVH carries a 15% 30-day mortality.²⁴ Resuscitation with blood according to the major haemorrhage protocol should be instituted in patients with haemodynamic instability and life-threatening bleed, aiming for a mean arterial pressure >65 mmHg (permissive hypotension may reduce rate of bleeding). A restrictive transfusion strategy is recommended in those who are haemodynamically stable (transfusion below 70 g/dL aiming for 70–80 g/dL).²⁵ Platelet transfusion and fresh frozen plasma are not recommended outwith the major haemorrhage protocol, as this can increase portal hypertension and exacerbate bleeding.²⁵ Tranexamic acid and proton pump inhibitors are of no benefit.²⁶

Pre-endoscopy, terlipressin (2 mg then 1–2 mg four times a day, unless contraindications) and empirical intravenous antibiotics are recommended in AVH.¹ In those with a contraindication to terlipressin, octreotide (50 µg followed by 50 µg/hr infusion) can be used.²⁷ Recent evidence suggests that stopping terlipressin once haemostasis is confirmed (24 h post-endoscopic treatment with no evidence of bleeding) leads to better outcomes than continuing for 72 h.²⁸ Intravenous erythromycin pre-endoscopy (250 mg) helps to clear blood from the stomach and improve views and endoscopic treatment.²⁹

Endoscopy should be undertaken within 24 h of presentation.²⁹ Prognostic scores such as the Blatchford score are not suitable for use in suspected portal hypertensive bleeding. Patients with active haematemesis, encephalopathy or haemodynamic instability should be intubated prior to endoscopy and discussed with critical care.¹

Hepatic encephalopathy (HE)

Hepatic encephalopathy can be precipitated by dehydration, electrolyte disturbance, infection, sedative medication and constipation. Ammonia levels are not required to diagnose patients with clinical HE, but a normal ammonia can be helpful in differentiating delirium from HE. All potential precipitants should be addressed by correcting electrolyte imbalance, suspending sedating medication, treating infections and by treating constipation with lactulose (aiming for 2–3 soft stools/day) and phosphate enemas if needed. Nasogastric tube placement should be considered in patients unable to take lactulose orally, and patients with reduced levels of consciousness (grade 3/4 encephalopathy) should be considered for escalation of care to critical care. Rifaximin can be added to those with recurrent/chronic encephalopathy to reduce risk of recurrence but does not acutely improve HE.³⁰

Ascites and oedema

Patients with ascites and/or oedema should have a moderately salt-restricted diet, under the supervision of a dietician.

Those with clinically tense ascites require paracentesis. In the absence of tense ascites, and/or where there is peripheral oedema, diuretics should be titrated every 72 h to aim for 0.5–1.0 kg weight loss per day. A combination of spironolactone (starting at 100 mg) and furosemide (starting at 40 mg) can be titrated according to potassium levels, renal function and response.^{1,18}

Nutrition

Malnutrition in AD is associated with increased complications including infections, HE and ascites, and mortality.³¹ Patients should eat every 2–3 h, including a bedtime snack, and have a high-protein diet (1.5 g/kg/day), rarely achievable in hospital without nutritional supplements.³¹ All patients should be reviewed by a dietician, and high protein supplements offered to those with/at risk of malnutrition or insufficient oral intake. Electrolytes (potassium, magnesium and phosphate) should be monitored and replaced if depleted. Enteral feeding should be considered in those who are unable to tolerate sufficient calories orally, but can be risky in very confused/agitated patients.

VTE prophylaxis

Clotting parameters in patients with cirrhosis do not accurately reflect bleeding risk, and patients with AD are at increased risk of thrombosis, so all patients with platelet count >50 should have tinzaparin thromboprophylaxis in the absence of active bleeding.⁹

Alcohol-associated hepatitis (AH)

Alcohol-associated hepatitis is a clinical syndrome characterised by acute-onset jaundice in the setting of long-term heavy alcohol use.

Decompensated Cirrhosis Discharge Bundle

This checklist should be completed by a member of the ward team. It should be started a minimum of 48 hours prior to discharge but can be done earlier and should be completed alongside the discharge letter. The information on the checklist should be reviewed on the consultant ward round prior to discharge.

Named consultant			
Date of liver follow up appointment			
Aetiology of liver disease			
Cause of decompensation (if known)			

Ascites			
Ascites present	Y	N	
Previous SBP	Y	N	
If yes: Date			
Organism (if known)			
Prophylactic antibiotics	Y	N	
If yes: name			
If no: reason why			

Patients with ascites who have had an episode of SBP should be considered for antibiotics (secondary prophylaxis). Co trimoxazole 480mg od first line unless contraindicated

Current management of ascites			
Diuretics	Y	N	
Paracentesis	Y	N	
Weight at discharge and documented in discharge letter		Kg	

If requiring paracentesis:

Predicted interval _____ weeks

Day unit appointment booked for _____

Or information given to patient to contact Day Unit at Freeman Hospital

Renal function			
Have the following been documented in the discharge letter:			
Discharge creatinine	Y	N	
Frequency of U&Es monitoring in the community	Y	N	
Once ascites is controlled that diuretics can be reduced to the lowest effective dose	Y	N	

Hepatic encephalopathy			
Encephalopathy present	Y	N	
Lactulose	Y	N	
Rifaximin	Y	N	

Lactulose and Rifaximin are recommended for patients with persistent or a previous unprovoked episode of encephalopathy, unless contraindicated.

Portal hypertension			
Varices	Y	N	
Grade of varices	1	2	3
Previous variceal bleed?	Y	N	
Is a repeat OGD required?	Y	N	
If so, date booked for _____			
Prophylaxis			
Is patient on a B Blocker (carvedilol preferred)	Y	N	
If not, why not? _____			
Has advice been given about titrating dose? (aim HR 60/min and SBP >100)	Y	N	
Variceal band ligation	Y	N	

A repeat OGD is recommended at 4 weeks for those who have had variceal banding. Non-selective β Blockers are recommended as primary prophylaxis for medium/large varices and for small varices with red signs or Childs C cirrhosis.

Substance / alcohol misuse			
Alcohol misuse	Y	N	
Input from alcohol liaison team	Y	N	
Community follow up plans	Y	N	
Thiamine prescribed	Y	N	

Treatment plan			
If treatment limitations or palliative care have been decided, has this been detailed in the discharge letter and does the patient have an appropriate Treatment Escalation Plan or Emergency Health Care Plan?	Y	N	

Communication with patient			
Have the following been explained to the patient and/or family?			
The diagnosis of chronic liver disease	Y	N	
The importance of abstinence (if applicable)	Y	N	
Current medications and reasons for taking them	Y	N	
Patient given the cirrhosis management toolkit leaflet	Y	N	

Fig. 3. Decompensated cirrhosis discharge bundle.³⁷

Laboratory-based prognostic scores help determine disease severity and guide management.³² The most important determinant of long-term outcome is abstinence from alcohol – optimising nutrition, treating concomitant infections and supportive treatment for complications (including ascites, HE, AVH, AKI) are the mainstay of in-hospital treatment.³³

Corticosteroids (prednisolone 40 mg od) are thought to be beneficial for a subgroup of patients with severe AH.³² Infection and bleeding should be excluded prior to starting steroids. If bilirubin starts to fall in this period, corticosteroids may not be required. If corticosteroids are started, response should be assessed at day 4–7 using the Lille score,^{34,35} and treatment stopped in those who are not responding.³² In those who respond, prednisolone is continued for 28 days.

The deteriorating patient

Patients with AD who develop acute on chronic liver failure (ACLF), a severe form of decompensation involving organ failure and systemic inflammation, and are suitable for treatment escalation should be referred to critical care.² Patients under 60 years old with no contraindications to transplant and, in the case of ARLD, abstinent for >3 months who develop three-organ failure can now be referred for consideration of urgent liver transplant assessment. A UK pilot demonstrated that transplant prioritisation improved survival for this group.²⁰

In patients who deteriorate despite treatment, and in whom escalation is considered futile, early palliative care should be considered. Palliative care in advanced liver disease is covered elsewhere and is beyond the scope of this article. However, advanced liver disease carries a significant physical and psychosocial burden, particularly in the last year of life, and parallel planning with early palliative care improves quality of life and even prognosis.³⁶

Discharge

Patients admitted with alcohol-related liver disease (AD) have high readmission rates. Ensuring appropriate follow-up on discharge is essen-

tial. Patients should receive clear self-management advice, contact details for the liver team, and support for alcohol abstinence where needed.

Use of a cirrhosis discharge bundle (Fig. 3), has been shown to improve follow-up and reduce readmissions.³⁷ Early discharge clinics can also reduce readmissions and length of stay, and improve outcomes.³⁸

Advanced care planning should also be considered, discussed with the patient and family, shared with primary care, and include community palliative input where ongoing support is required.

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CRediT authorship contribution statement

Dina Mansour: Writing – review & editing, Writing – original draft, Conceptualization.

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