

REFERENCES

For references, please visit www.TheAnesthesiaGuide.com.

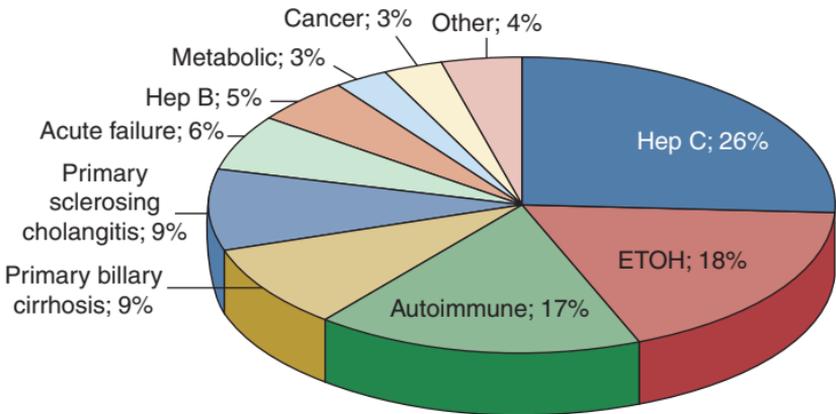
CHAPTER 81

Liver Transplant

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PREOPERATIVE

FIGURE 81-1. Indications for liver transplant



4000–600 liver transplants in the US/year over the last 10 years. Only 300–400 of these are living donor transplants.

CONTRAINDICATIONS**Absolute**

- Pulmonary hypertension: mean pulmonary artery pressure > 45 mm Hg
- Recidivism to alcohol and drug abuse
- Extrahepatic malignancies
- Systemic sepsis

Relative

- Pulmonary hypertension: mean pulmonary artery pressure > 35 mm Hg
- Significant cardiopulmonary disease
- Poor psychosocial support or compliance
- In the case of hepatocellular carcinoma (“Milan criteria”)
 - ▶ 1 tumor >5 cm
 - ▶ >3 tumors <3 cm
 - ▶ Presence of extrahepatic tumors
 - ▶ Presence of vascular invasion

TABLE 81-1 Liver Failure and Associated Morbidities

Neuro	<ul style="list-style-type: none"> • Encephalopathy • Increased intracranial pressure (with fulminant liver failure) • Coma
Respiratory	<ul style="list-style-type: none"> • Hepatopulmonary syndrome <ul style="list-style-type: none"> ▶ Transpulmonary shunts with paradoxical improvement of SaO₂ in supine position (orthodeoxia) • Atelectasis from ascites • Aspiration risk
Cardiac	<ul style="list-style-type: none"> • Hyperdynamic state of cirrhosis <ul style="list-style-type: none"> ▶ Low SVR, high cardiac output, splanchnic vasodilation • Portopulmonary hypertension • Mean PAP >35 mm Hg relative contraindication for liver transplantation
GI	<ul style="list-style-type: none"> • Delayed gastric emptying and increased intra-abdominal pressure (ascites) <ul style="list-style-type: none"> ▶ Rapid sequence induction • Esophageal varices
Renal	<ul style="list-style-type: none"> • Hepatorenal syndrome <ul style="list-style-type: none"> ▶ Type I: doubling of serum creatinine within 2 weeks <ul style="list-style-type: none"> ▪ Mean survival: 1 month ▶ Type II: increase of serum creatinine > 1.5 mg/dL <ul style="list-style-type: none"> ▪ Mean survival: 6 months • Relative hypovolemia • Hyponatremia
ID	<ul style="list-style-type: none"> • Immunosuppression • Spontaneous bacterial peritonitis
Hematology	<ul style="list-style-type: none"> • Myelosuppression <ul style="list-style-type: none"> ▶ Anemia ▶ Leukopenia • Thrombocytopenia from hypersplenism • Coagulopathy of liver disease <ul style="list-style-type: none"> ▶ Frequently hypo-/hypercoagulable disease

ANESTHESIA

Routine setup of liver room at Columbia University Medical Center (courtesy of Dr. Tricia Brentjens)

Drugs

Induction and maintenance

- Propofol
- Etomidate
- Midazolam (have 10 mg available)
- Fentanyl (have 3 mg available)
- Succinylcholine
- Cisatracurium (have 200 mg available)
- Isoflurane (have 2 full bottles available)

Vasoactive drugs

- Epinephrine 100 mcg/mL, 10 mL
- Epinephrine 10 mcg/mL, 10 mL
- Calcium chloride 1 g in 10 mL (have 10 vials available)
- Phenylephrine 40 mcg/mL, 10 mL syringe, and 250 mL bag

- Ephedrine 5 mg/mL, 10 mL
- Atropine 100 mcg/mL, 10 mL
- Sodium bicarbonate 50 mEq (have 10 vials available)

Others

- Magnesium sulfate 2 g (4 mL of the 0.5 g/mL solution): Not drawn up!
- Methylprednisolone sodium 500 mg
- Mannitol 12.5 g/20 mL: Not drawn up
- Acetylcysteine: discuss with your attending physician
 - Load: 150 mg/kg in 200 cc over 1 hour
 - Maintenance: 50 mg/kg in 500 cc D5W over 4 hours
 - Then 100 mg/kg in 1 L D5W over 16 hours

Mandatory infusions

- Norepinephrine 4 mg/250 mL N saline
- Vasopressin 100 U/100 mL N saline
- Furosemide 100 mg/100 mL N saline

Have available

- Dopamine 200 mg/250 mL premix bag
- Dobutamine 250 mg/250 mL premix bag
- Nitroglycerin 50 mg/250 mL premix bottle
- Aminocaproic acid (Amicar) — antifibrinolytic: various protocols

Monitors

Routine monitors

- EKG 3 channels: I, II, V
- SpO₂
- NIBP cuff of medium, large, extra large sizes

Hemodynamic monitoring

- Do not heparinize pressure lines/bags!
- Three pressure transducers on the right: for right femoral artery, PA, CVP
- One pressure transducer on the left: for left radial artery
- Pediatric CVL kit [20 g] for femoral arterial line
- Assorted 20 G cannulas for radial arterial line placement
- Double lumen Cordis for IJ placement
- Oximetric PA catheter (needs to be calibrated prior to insertion)

IV setup

- All IVs are primed with PlasmaLyte except when otherwise indicated
- Rapid transfuser FMS2000 primed
- Two IV lines on warmers
- Venous infusion port (VIP) on microdrip with six ports for IV drips
- Call the blood bank and check on available products:
 - 10 PRBCs, 10 FFP, and 12 platelets

Others

- Point of care arterial blood gas (ABG) analyzer with at least 20 cartridges
- Multiple test tubes for complete blood count (CBC) and coagulation studies
- Two forced air warming machines with upper and lower body blankets
- For CVVHD: high-flow set for the blood warmer on the left side of the OR table to warm the returning limb of CVVH

MONITORING

- Arterial catheter (femoral and radial)
 - Radial artery catheter often dampens during anhepatic phase
- PA catheter
 - Maximize cardiac output
 - Measure PA pressures
- Pulse pressure variation (PPV)
 - Allows estimate of volume responsiveness
- Point of care arterial blood gas analyzer
 - Measure pH, acidemia, hematocrit, potassium
- Point of care coagulation test: thrombelastography or rotational thromboelastometry (ROTEM®)
 - Allows estimation of global *in vivo* hemostasis including coagulation factor defects, platelet function, abnormal fibrinogen polymerization, anticoagulant defects and hyperfibrinolysis

INDUCTION

- Rapid sequence induction for almost all patients (ascites)
- Patients are hyperdynamic with low SVR and maintain cardiac output with increased endogenous catecholamines: beware of hypotension after induction
- Most induction drugs will need a higher dose and have a delayed onset (larger volume of distribution) but have a prolonged half-life time (decreased hepatic metabolism)

MAINTENANCE

Surgical technique

Complete caval clamping

Vena cava is clamped completely above and below the liver and the donor cava is anastomosed end-end to the recipient.

Advantages:

- Fast
- Simpler (and possibly better) caval anastomosis

Disadvantages:

- Severe hypotension due to loss of preload during anhepatic phase
- Renal outflow obstruction and renal injury

Veno-venous bypass

Cannulation of portal vein and femoral vein with return of blood through a bypass pump either to a cannula in the axillary vein or the internal jugular vein. Anticoagulation is rarely required if heparin coated circuits are used.

Advantages:

- Hemodynamic stability during anhepatic phase
- Adequate renal venous drainage
- Less intestinal engorgement from portal occlusion

Disadvantages:

- Cumbersome
- Technical complications from access

Piggyback technique

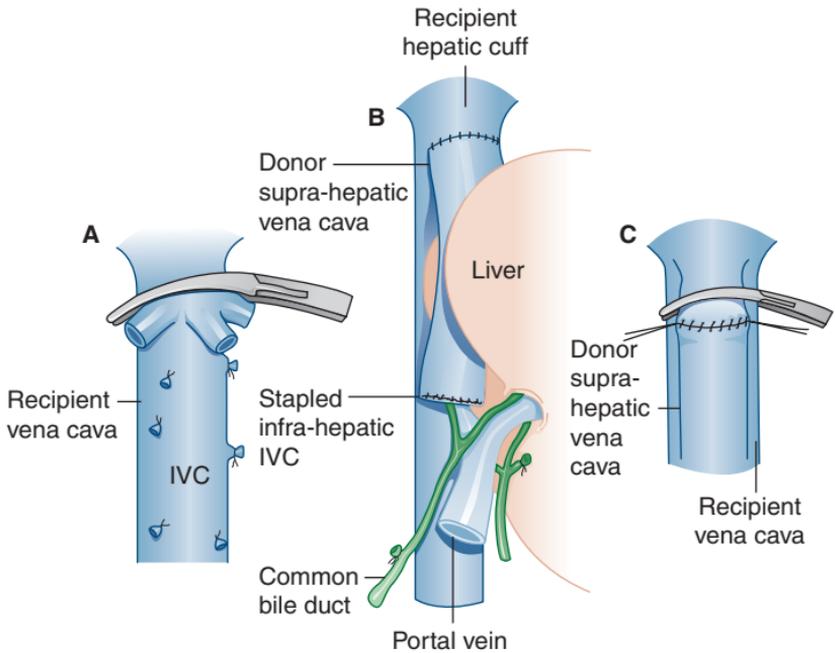
Instead of replacing the vena cava with a typical bicaval approach, the recipient vena cava is preserved, and an anastomosis is created between the donor suprahepatic cava and a cuff of the recipient hepatic veins. Partial occlusion of the vena cava at the level of the hepatic veins. Partial caval blood flow is maintained throughout.

Advantages:

- Less hemodynamic instability
- Renal venous drainage is maintained

Disadvantages:

- Caval anastomosis is technically more difficult
- Portal occlusion may still cause intestinal engorgement (may be alleviated with temporal portocaval shunt)

FIGURE 81-2. Piggyback technique of liver transplantation

The recipient vena cava is preserved, and an anastomosis is created between the donor suprahepatic vena cava and a cuff of the recipient hepatic veins. Reproduced from Minter RM: *Current Procedures: Surgery*. Figure 32-7A-C. Available at: www.accesssurgery.com. Copyright © The McGraw-Hill Companies, Inc. All rights reserved.

TABLE 81-2 Four Phases of Liver Transplantation

Pre-anhepatic/ dissection phase	<ul style="list-style-type: none"> • Bleeding due to dissection in area of portal hypertension • Correct coagulopathy, anemia, and optimize volume status
Anhepatic	<ul style="list-style-type: none"> • Low preload and cardiac output despite low SVR • Acidosis: unable to metabolize lactate
Reperfusion	<ul style="list-style-type: none"> • Possibly sudden <ul style="list-style-type: none"> ▶ Acidosis ▶ Hyperkalemia ▶ Pulmonary hypertension ▶ Right ventricular distension and failure ▶ Cardiac arrest
Neohepatic phase	<ul style="list-style-type: none"> • If the liver functions well: within hours improvement of <ul style="list-style-type: none"> ▶ Acidosis ▶ Coagulopathy ▶ Vasodilatory state

PRE-ANHEPATIC/DISSECTION PHASE

Bleeding due to dissection in area of portal hypertension

- Treat only clinical bleeding, not laboratory numbers:
 - Increased INR can be associated with hypercoagulability (deficiency of liver derived protein C and S and other anticoagulant factors)
 - Thrombelastography/rotational thrombolastography (ROTEM) may detect specific coagulation defects
- Maintain isovolemia and optimize fluid status:
 - Occult blood loss from retroperitoneal dissection or bleeding from collateral veins
 - Acute fluid loss when ascites is drained
- Do not transfuse platelets unless platelet count low, clinical bleeding evident and after consultation with the surgery team
 - Platelets sequester in the spleen and platelet transfusion is associated with increased thrombotic complications
- Often furosemide is required to maintain urine output and create intravascular “room” for transfusion of coagulation factors (plasma)

ANHEPATIC

Low preload and cardiac output despite low SVR

- Requires adequate preload especially with caval cross-clamping
 - Goal CVP 12–15
- The anesthesiologist needs to be confident that he/she will be able to maintain adequate hemodynamics during the anhepatic phase before the liver is removed
- Often requires high doses of vasopressors
 - Norepinephrine 2–10 mcg/min
 - Vasopressin 1–4 units/h
 - Goal: MAP 60 mm Hg or within 20% of preoperative BP
 - No urine output during anhepatic phase may indicate worse renal outcome
- Acidosis: unable to clear lactate during anhepatic phase
 - Treat aggressively with sodium bicarbonate (50–100 mEq over 20 minutes) as acidosis will worsen with reperfusion
- Large fluid administration during the anhepatic phase will cause volume overload and possible right heart failure with reperfusion
 - Use preferentially vasopressors to maintain blood pressure
- We routinely give 2 g magnesium sulfate IV over 10 minutes during anhepatic phase to prevent arrhythmias during reperfusion
- Check arterial blood gas 20 minutes prior to reperfusion to detect treatable abnormalities: acidosis, hyperkalemia, hypocalcemia

REPERFUSION

Washout of graft often associated with sudden:

- Acidosis
- Hyperkalemia: closely watch T waves
- Pulmonary hypertension
- Right ventricular distension and failure
- Cardiac arrest
 - Reperfusion will be worse with prolonged cold or warm ischemic time, steatotic grafts or with organs after donation after cardiac death (DCD)
 - Consider requesting from surgeon to flush the graft with portal vein blood prior to finishing the caval anastomosis, or slow reopening of portal vein
 - Aggressively treat:
 - Hyperkalemia: Insulin/glucose, bicarbonate, CaCl₂
 - Hypocalcemia
 - Acidosis: bicarbonate and hyperventilation
 - Hypotension: vasopressors, epinephrine in 10–30 mcg intravenous boluses

NEO-HEPATIC

If the liver functions well: within hours, improvement of

- Acidosis
- Coagulopathy
- Vasodilatory state
 - ▶ Excessive fibrinolysis possible due to graft reperfusion and release of tissue plasminogen activator (tPA): consider transfusion of cryoprecipitate
 - ▶ Avoid platelet transfusions especially if hepatic artery or portal vein anastomosis is precarious. Discuss with surgical team

POSTOPERATIVE

Extubation in the operating room is possible but little benefit

TABLE 81-3 Early Complications after Liver Transplant

0–24 h	<ul style="list-style-type: none"> • Bleeding • Graft failure <ul style="list-style-type: none"> ▶ Hepatic artery thrombosis ▶ Portal vein thrombosis ▶ Early graft failure/delayed graft function • Vasodilatory shock
1–5 days	<ul style="list-style-type: none"> • Acute kidney injury • Infection • Bile leak • Prolonged respiratory failure
>5 days	<ul style="list-style-type: none"> • Rejection • Malnutrition • Prolonged hepatic failure • “Failure to thrive”

Diagnostic features and therapy of early complications

Bleeding

Diagnosis:

- Increased drain output (consider checking drain fluid hematocrit)
- Increased abdominal girth (measure hourly)
- Increased intra-abdominal pressure (measure hourly)

Rx:

Correction of coagulopathy

If bleeding persists or substantial bleeding: re-exploration

Hepatic artery or portal vein thrombosis

Diagnosis

- Persistent
 - ▶ Metabolic (lactate) acidosis
 - ▶ Elevated total bilirubin
 - ▶ Coagulopathy
 - ▶ Ascites
 - ▶ Vasodilatory state and high vasopressor requirements
 - ▶ Low urine output and worsening acute kidney injury

- ▶ Late and severe cases:
 - Persistent hypoglycemia
 - Depressed mental status/encephalopathy
 - Hypothermia
- Diagnosis confirmed by Doppler ultrasound
- Elevated transaminases are due to (ischemic) injury to the liver and not necessarily a sign of dysfunction

Differential diagnosis

- Early allograft dysfunction due to
 - ▶ Primary non-function
 - ▶ Small for size syndrome

Rx:

- Re-exploration
- Rarely amenable to endovascular intervention

Bile leak

Diagnosis

- Persistent
 - ▶ Bilious fluid from drains
 - ▶ Increased total bilirubin, alkaline phosphatase (AP) and gamma-glutamyltranspeptidase (γ -GT)
 - ▶ Fever, malaise
- Diagnosis confirmed by ultrasound (intra-abdominal fluid) and drain fluid total bilirubin concentration

Rx:

- Re-exploration (breakdown of biliary anastomosis often due to ischemic injury to the bile) and possibly Roux-en-Y revision
- Percutaneous drainage
- Endoscopic biliary stent placement

Acute rejection

Diagnosis

- Days after surgery
 - ▶ Worsening liver function
 - ▶ Increasing total bilirubin and transaminases
 - ▶ Malaise and fever
- Diagnosis confirmed by biopsy

Rx:

- Increased immunosuppression
- Pulse steroids
- OKT-3 (muromonab-CD3 antibody)

PEARLS AND TIPS

- Magnesium (2g IV over 20 minutes) during the anhepatic phase may stabilize membranes and ameliorate cardiac arrhythmias of reperfusion
- Avoid platelet transfusion postoperative if there is concern about the patency of the hepatic artery anastomosis

REFERENCES

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