

TRANSPLANT HEPATOLOGY Blueprint

Maintenance of Certification Examination (MOC)

Purpose of the exam

The exam, which is developed jointly by the ABIM and the American Board of Pediatrics, is designed to evaluate the knowledge, diagnostic reasoning, and clinical judgment skills expected of the certified transplant hepatologist in the broad domain of the discipline. The ability to make appropriate diagnostic and management decisions that have important consequences for patients will be assessed. The exam may require recognition of common as well as rare clinical problems for which patients may consult a certified transplant hepatologist.

Exam content

Exam content is determined by a pre-established blueprint, or table of specifications, which is reviewed annually and updated as needed for currency. Trainees, training program directors, and certified practitioners in the discipline are surveyed periodically to provide feedback and inform the blueprinting process.

The primary medical content categories of the blueprint are shown below, with the percentage assigned to each for a typical exam:

Medical Content Category	% of Exam
Pretransplant	45%
Perioperative	20%
Post-transplant	25%
Transplant Immunology	5%
Miscellaneous	5%
	100%

ABIM is committed to working toward health equity and believes that board-certified physicians should have an understanding of health care disparities. Therefore, health equity content that is

clinically important to each discipline will be included in assessments, and the use of gender, race, and ethnicity identifiers will be re-evaluated.

Exam format

The exam is composed of up to 240 single-best-answer multiple-choice questions, predominantly describing patient scenarios. All candidates will see up to 180 common questions. ABIM candidates will see up to 60 additional questions specific to adult content areas; American Board of Pediatrics candidates will see up to 60 additional questions specific to Pediatric content areas. Questions ask about the work done (that is, tasks performed) by physicians in the course of practice:

- Making a diagnosis
- Ordering and interpreting results of tests
- Recommending treatment or other patient care
- Assessing risk, determining prognosis, and applying principles from epidemiologic studies
- Understanding the underlying pathophysiology of disease and basic science knowledge applicable to patient care

Clinical information presented may include patient photographs, radiographs, electrocardiograms, recordings of heart or lung sounds, and other media to illustrate relevant patient findings. More information on how exams are developed can be found at: https://www.abim.org/about/exam-information/exam-development.aspx.

Examinees will have access to an external resource (e.g., UpToDate®) for the exam. A tutorial including examples of ABIM exam question format can be found at https://www.abim.org/maintenance-of-certification/assessment-information/transplanthepatology/exam-tutorial.aspx.

The blueprint can be expanded for additional detail as shown below. Each of the medical content categories is listed there, and below each major category are the content subsections and specific topics that *may* appear in the exam. <u>Please note:</u> actual exam content may vary.

Pretransplant 45% of Exam



Biliary atresia (pediatrics only)	2%
Genetic liver disease	4%
Cholestatic syndromes (including progressive familial	
intrahepatic cholestasis [PFICs])	
Hereditary hemorrhagic telangiectasia	
(internal medicine only)	
Wilson disease	
Alpha-1-antitrypsin deficiency	
Iron overload syndromes	
Mitochondrial defect	
Urea cycle defect (pediatrics only)	
Cystic fibrosis	
Fibrocystic diseases (including Caroli disease	
and choledochal cysts)	
Familial amyloid polyneuropathy (FAP) (internal medicine only)	
Other genetic liver diseases (including erythropoietic	
protoporphyria)	
Autoimmune disorders	3%
Primary biliary cholangitis (internal medicine only)	
Hepatitis	
Overlap syndrome (including autoimmune cholangiopathy)	
Primary sclerosing cholangitis (including IgG4 cholangiopathy)	
Sarcoidosis (internal medicine only)	
Celiac disease	
Viral hepatitis	7%
Hepatitis A (HAV)	
Hepatitis B (HBV)	
Hepatitis C (HCV)	
Hepatitis D (HDV)	
Hepatitis E (HEV)	
Other viruses (including Epstein-Barr virus,	
cytomegalovirus, herpes simplex virus)	
Budd-Chiari syndrome, sinusoidal obstruction syndrome, and	
congestive hepatopathy <2% Growth failure (pediatrics only)	2%
Portal hypertension	2%



	Varices	
	Ascites	
	Encephalopathy Spontaneous bacterial peritonitis Noncirrhotic portal hypertension	
	Hepatic hydrothorax	
	Hepatopulmonary syndrome and portopulmonary hypertension	Hepatorenal
	syndrome	
Liver t	umors	4%
	Hepatocellular carcinoma	
	Hepatoblastoma (pediatrics only)	
	Cholangiocarcinoma (internal medicine only)	
	Other tumors (including benign, hemangioendothelioma, and neuroendocrine)	
Selecti	ion and evaluation for transplantation	9%
	Pediatric End-Stage Liver Disease (PELD) and Model for	
	End-Stage Liver Disease (MELD) scoring systems, including	
	psychosocial issues	
	Contraindications to transplantation	
	Exceptions to PELD and MELD systems	
	Live donor selection	
	Impact of active infection, malignancy, and malnutrition	
	on outcome	
	Multiorgan (liver, kidney) recipients	
	Co-morbidities (including human immunodeficiency virus [HIV] infection)	
Acute	liver failure	5%
	Epidemiology	
	Etiology	
	Pathobiology	
	Assessment	
	Prognostic indicators	
	Treatment	



Indications for transplantation

Outcome as a function of age and diagnosis

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Metabolic dysfunction-associated steatotic liver disease,	
including metabolic dysfunction-associated steatohepatitis	<2%
Liver diseases of pregnancy	<2%
Transfer of care	<2%
Drug-induced liver disease	<2%
operative	20% of Exam
Donor selection	3%
Extended-criteria donors	
Steatosis	
Viral infection	
Domino (sequential) (internal medicine only)	
Surgical options, complications specific to	
graft and donor types (including ABO blood type)	3%
Perioperative complications	6%
Initial poor function or primary nonfunction	
Vascular complications	
Infections (viral, bacterial, and fungal) Hepatitis	
B and C antiviral therapy	
Biliary complications	
Allograft rejection	
Metabolic complications (including neurotoxicity	
and nephrotoxicity)	
Drug hepatotoxicity	2%
Nutritional support	2%
Living donor	<2%
Small for size syndrome	
Donor complications	
Recipient complications	
Donor transmission of disease	<2%
Donation after circulatory death	<2%

Alcohol-associated liver disease (internal medicine only)



<2%

2%

Split graft transplantation

ransplant	25% of Exam
Immune complications	4%
Rejection	
Graft-versus-host disease	
Alloimmune and autoimmune diseases (de novo)	
Nonimmune complications	5%
Diabetes mellitus	
Renal	
Bone	
Growth and development (pediatrics only)	
Cardiovascular complications	
Vascular complications	
Viral infections (cytomegalovirus, Epstein-Barr virus, and human herpesvirus)	5%
Bacterial infections	
Fungal infections	
Emerging infections Recurrence of disease (including hepatitis C, cancer, PBC, AIH)	3%
Post-transplant malignancy	3% 2%
Post-transplantation lymphoproliferative disorder (PTLD) Surfor malignancy	
Indications for retransplantation	2%
Adherence to medical regimen	2%
Quality of life	2%
lant Immunology	5% of Exam



2%

Basic immunology

Innate and adaptive immune system Immune response Tolerance Mechanism of action and pharmacokinetics (PK) of 2% immunosuppressive medications Cyclosporine and tacrolimus Mycophenolate mofetil (MMF), mycophenolic acid (MPA), and azathioprine Sirolimus and everolimus Antibody therapy **Drug-drug interactions** Corticosteroids Short-term immune and nonimmune toxicity of immunosuppressive medications <2%

Miscellaneous 5% of Exam 2% **Statistics** Kaplan-Meier (KM) Cox proportional hazards Relative risk Odds ratio Receiver operating characteristic curves **Ethics** 2% Psychosocial evaluation Living donor transplantation Transplant tourism Clinical trial participation Managed care and reimbursement issues <2% <2% **Regulatory issues** Policy implications of organ shortage Regulation

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