

# The Utility of Cross-Sectional Imaging in Predicting Etiology of Abnormal Liver Tests in **Early Post-Liver Transplant Patients**

## Introduction

There is a growing number of patients who require liver transplantation (LT). Within the first three months post-transplantation, there can be fluctuations in a patient's enzymes which may be indicative of multiple potential pathologic processes . In order to protect the lifespan of the recently transplanted liver, we must understand the etiology these lab abnormalities.

Cross-sectional imaging through computed tomography (CT) magnetic resonance cholangiopancreatograpy (MRCP) is typic among first line workup. Endoscopic retrograde cholangiopancreatography (ERCP) may be needed to interven on pathology within the liver bile ducts.

This study plans to address the current ambiguity regarding the reliability of MRCPs and CTs in context of recent surgical anatom through the evaluation of performance characteristics of MRC and/or CT in accurately predicting biliary etiology for elevated liver tests as compared to ERCP.

## Methods

- This is a retrospective analysis of patients undergoing liver transplant at KECK Medicine of USC from 2005 to 2023
- Inclusion criteria: patient ages 18 to 70, abnormal liver test within 3 months of transplant and with cross sectional imaging, follow-up for at least 1 year post-transplant
- Exclusion criteria: patients with normal liver tests within th first 3 months post-transplant, less than 1 year follow-up
- Sub-group analysis performed comparing patients who underwent CT/MRI and (ERCP)
- Primary outcome: comparison of findings on cross-section imaging and determine concordance or discordance with
- Secondary outcomes: rate of upstaging diagnosis at time of ERCP as compared to imaging
- Statistical analysis: Frequencies and percentages will be measured for categorical variables; mean and standard deviation were calculated for continuous variables

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	Table 1: Patient Der	m
	$\Delta g_{\Theta} (v_{\Theta} arg_{+} \pm SD)$	
	Male gender; n (%)	
	Ethnicity; n (%)	
liver	Hispanic	
	African American	
	Other	
. (	Etiology of Liver Disease; n (%)	
OT	Alcohol-related liver disease (NAFLD)	
	Hepatitis B (HBV)	
	Autoimmune hepatitis (AIH)	
or	Other*	
Or	Multiple etiologies	
cally	Type of liver transplant (years <u>+</u> SD)	
	Cadaveric	
	Living donor Type of anastomosis: n (%)	
IE	Duct-to-duct	
	Roux hepaticojejunostomy	
	SD = standard deviation; *Other diagnoses including primary sclerosing cholangitis (PSC); po and cryptogenic cir	lyc rho
he	Table 2: Characteristics in LT Patients	W
.OTTy		
CP	Days to Abnormal Liver Tests; days SD	
d	AST (U/L)	
	ALT (U/L)	
	Total Bilirubin (mg/dL)	
	CT Findings; n (%)	
	Extrahepatic Duct Dilation	
	Duct Stenosis Filling Defect	
	Unremarkable biliary findings	
	Intrahepatic Duct Dilation	
	Extrahepatic Duct Dilation	
	Filling Defect	
	Unremarkable biliary findings FRCP: n (%)	
-		
	Table 3: Comparison of Imaging a	ar
sts	Age (years, $\pm$ SD)	
	Type of liver transplantation; n (%)	
	Living donor	
ho	Type of anastomosis; n (%)	
	Roux hepaticojejunostomy	
	Days to Abnormal Liver Tests; days <u>+</u> SD Liver Tests (mean + SD)	
	AST (U/L)	
	ALT (U/L) Alkaline Phosphatase (IU/L)	
	Total Bilirubin (mg/dL)	
	ERCP Findings; n (%)	
ERCP	Extrahepatic Duct Dilation	
of	Duct Stenosis Filling Defect	
	Unremarkable biliary findings	
	Discordance of Findings; n (%)	_
	Between CT and ERCP	
	Upstaged Findings based on ERCP	
	Anastomotic stricture not seen on imaging Multifocal intrahepatic strictures not seen on imaging	
	Liver Biopsy Performed Prior to ERCP; n (%)	
	Final Diagnosis consistent with Liver Biopsy*	<u> </u>

ographics
N = 39 patients
63.4 (23.3)
28 (72.0)
16 (41.0)
10 (25.6)
2 (5.1)
4 (10.3)
7 (17.9)
6 (15.4)
6 (15.4)
1 (2.6)
6 (15.4)
2 (5.1)
6 (15.4)
12 (30.8)
56.4 (13.4)
35 (89.7)
4 (10.3)
75 (64-1)

23 (04.1) 14 (35.9)

ystic liver disease, primary biliary cirrhosis (PBC), alpha-1 antitrypsin disease

ith Acute Liver 7	<b>Test Elevation</b>

	_
N = 39 patients	
32.8 (31.1)	
114.2(257.9)	
126.7 (157.6)	
216.4 (137.6)	
3.3 (5.1)	
27 (69.2)	
0 (0.0)	
1 (2.6)	
0 (0.0)	
1 (2.6)	
14 (35.9)	
20 (51.3)	
7 (17.9)	
3 (7.7)	
1 (2.6)	
1 (2.6)	
9 (23.1)	
10 (25.6)	

nd Endoscopic Findings	
N = 10	
59.8 (16.3)	
9 (90.0)	
1 (10.0)	
9 (90.0)	
1 (10.0)	
36.5 (30.0)	
70.4 (64.5)	
96.6 (83.2)	
271.0 (113.3)	
4.1 (6.6)	
2 (2.0)	
3 (3.0)	
4 (40.0)	
0 (0.0)	
3 (3.0)	
3 (3.0)	
5 (5.0)	
3 (3.0)	
2 (2.0)	
1 (1.0)	
2 (20.0)	
2 (100.0)	

ERCP showing anastomotic stricture Unremarkable MRCP Results • 39 patient charts were analyzed of which the mean age was 63.4 years and 72% were male. • The most common etiology of liver disease was NAFLD (15.4%), HCV (15.4) and combination of multiple liver pathologies (for example NALFD and HCC, etc) (Table 1) Most patients (89.7%) underwent orthotopic cadaveric liver transplant and had duct to duct biliary anastomosis (64.1%). • Multiple imaging modalities with both CT and MRCP was performed in 41% of patients, of which 43.8% had discordant liver or biliary findings. In 71.4% of these patients, MRCP identified either a intraductal filling defect or stricture not seen on CT. CT did identify patients with hematomas not seen on MRCP. • An ERCP was performed in 10 (25.6%) patients following CT/MRCP. There was discordant findings between MRCP and ERCP in 3 (30%) patients and between CT and ERCP in 5 (50%) patients. A biliary diagnosis was upstaged during ERCP in 3 (30%) patients. Summary In patients with acute liver test elevation within 3 months of liver transplantation, cross-sectional imaging may not be a reliable source to identify potential biliary etiologies as seen in 30% of patients having upstaged diagnoses based on ERCP findings. In addition, cross-sectional imaging findings between CT and MRCP appear inconsistent with 43.8% of patients having discordant findings between tests. Acknowledgements Thank you Dr. Jennifer Phan for providing me with this amazing opportunity, Dr. Will Minteer and Dr. Kurt Hong for constantly guiding and supporting me throughout my research, CTSI for providing a patient list, and the department of transplant hepatology for assisting us in clinical research.

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\*Liver biopsy diagnosis includes reperfusion injury and acute cellular rejection



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