



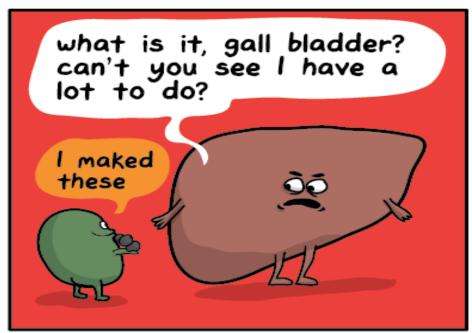
Shahid M. Malik, MD

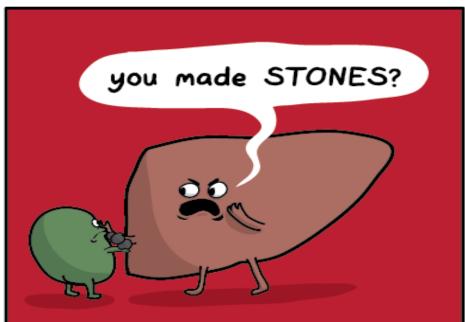
Assistant Professor of Medicine
Program Director, Transplant Hepatology
Division of Gastroenterology, Hepatology & Nutrition
University of Pittsburgh

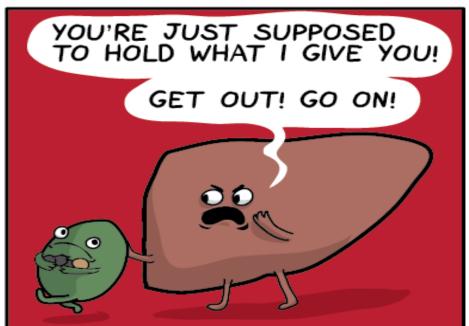
Disclosures

 I have no conflicts of interest regarding the content of my presentation

Cholestasis







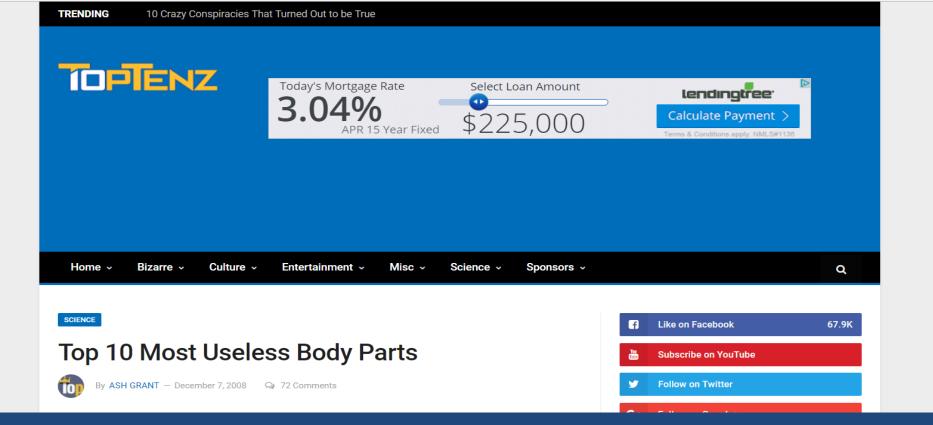


the Awkward Yeti.com

Functions: Liver Vs GB

• LIVER: 500

• GB: 2



- 1. Appendix
- 2. GB
- 3. Male Nipples

Cholestasis

- Greek in origin
- Means bile stoppage
- In its most overt form it presents as jaundice
- In early stages, cholestatic diseases lead to a preferential increase in the cholestatic liver enzymes: AP and GGTP



Differential of Cholestatic Injury Biliary obstruction (stones, strictures, cysts, malignancy) Infiltrative Diseases (Amyloid, Sarcoid, Malignancy, esp: Lymphoma) Drug (Amox/Clav); TPN Outflow (Right sided heart failure, Budd Chiari) Sepsis **Cystic Fibrosis** Cholestatic Hepatitis (HBV/HCV) Rejection; GVHD

Autoimmune Cholangitis

Secondary sclerosing diseases

ICP, BRIC, PFIC

Paraneoplastic

Idiopathic

Primary Biliary Cholangitis &

Primary Sclerosing Cholangitis

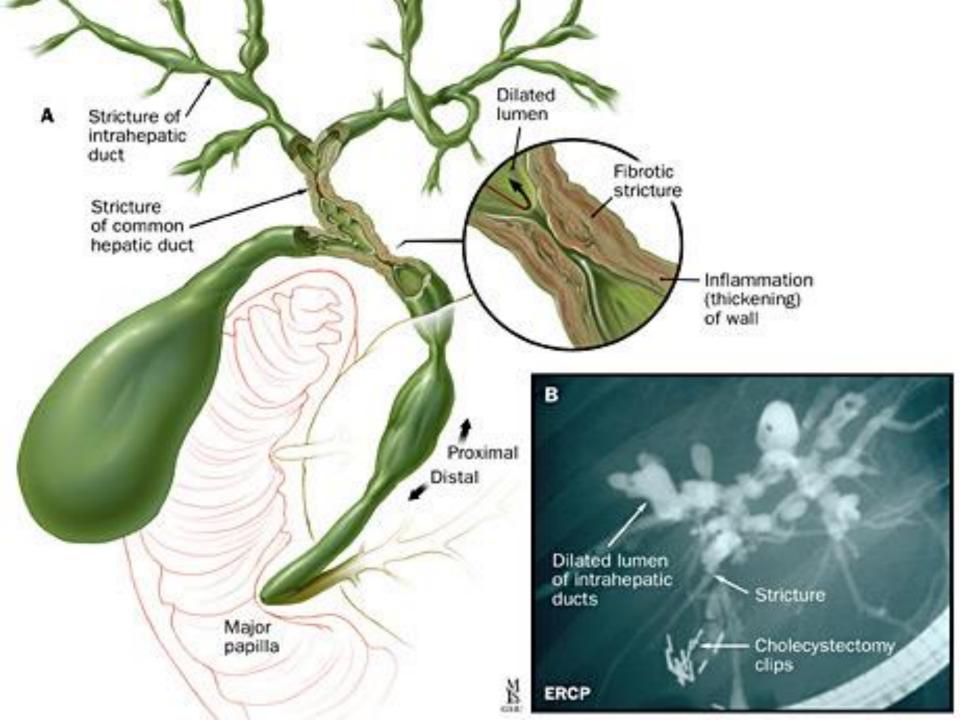
 18 papers and 4 abstracts all published within last year with relatively high IF

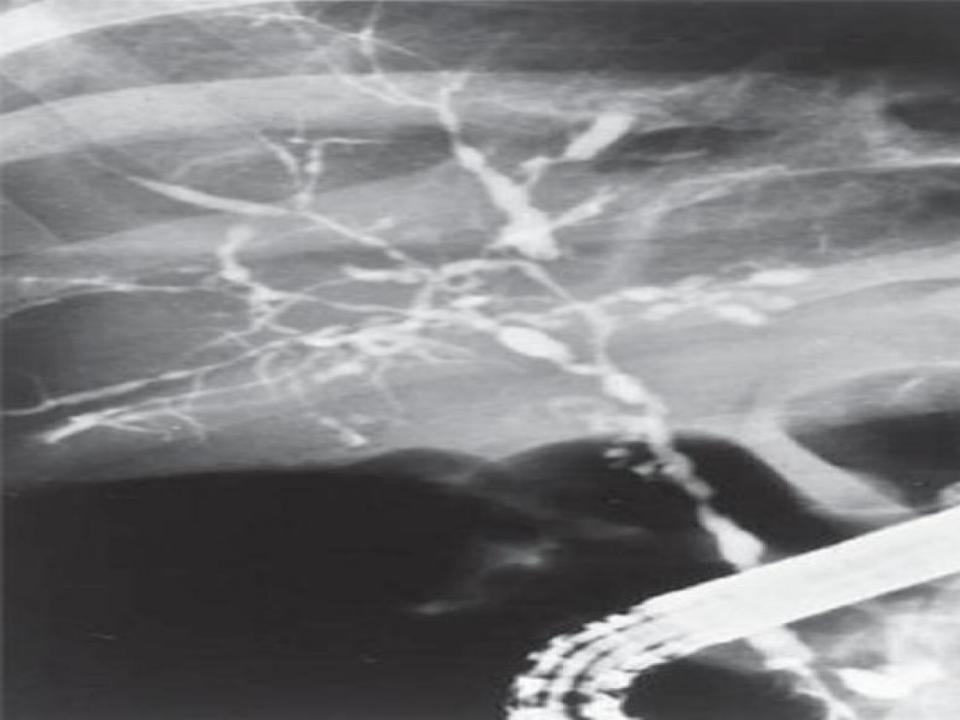
Primary Sclerosing Cholangitis

- Chronic progressive disorder of unknown etiology (?Immune mediated) that is characterized by inflammation, fibrosis, and stricturing of medium size and large ducts in the intrahepatic and extrahepatic biliary tree
- Dx made on cholangiogram
 - MRCP > ERCP
- 40K cases in US
- 70% are male; average age dx 40
- ~ 80% of patients have concomitant IBD (UC)
 - Increased risk of colon cancer

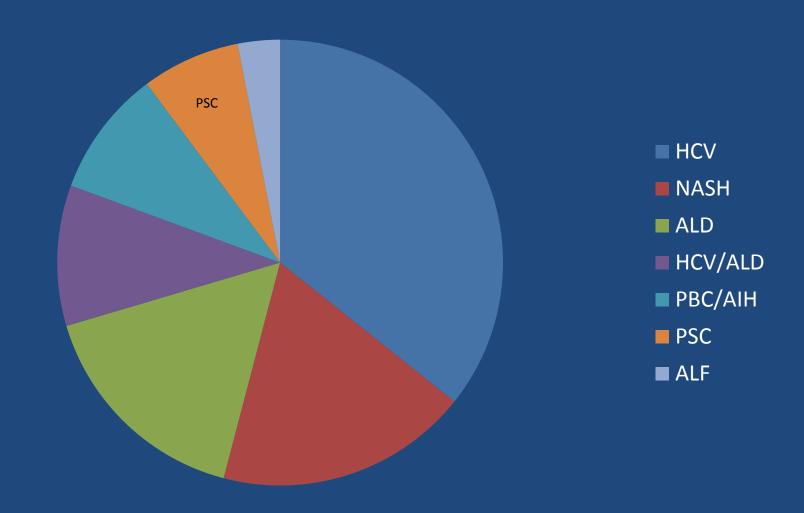
PSC

- Increased risk of cholangiocarcinoma
- Increased risk of GB cancer
- No proven medical treatment
- ? 50% of patients will require OLTx at 15 years





Liver Transplants US Last Decade



British Society Guidelines on PSC

Gut August 2019 Chapman et al

- 1. Recommend that UDCA not be routinely used for new dx of PSC
- 2. ERCP should be performed with expert multidisciplinary assessment
 - Mandatory that dominant strictures be sampled
- Imaging should be prompted by change in symptoms or biochemical abnormalities
 - Recommend against routine monitoring of Ca 19-9
 - No recommendations on routine cholangio surveillance
- GB US yearly
- In patients with developing cirrhosis US +/- AFP per guidelines

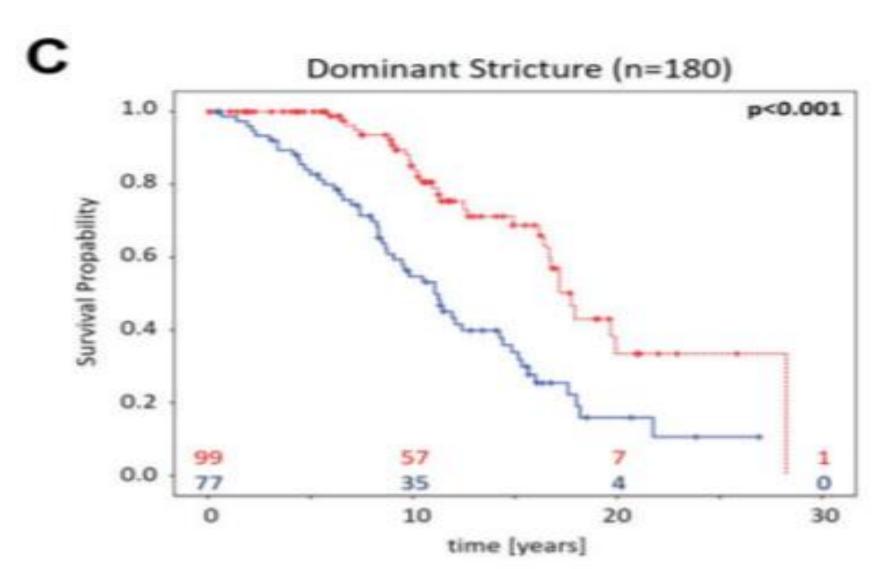
AGA Clinical Practice Update on HB Ca in PSC: Expert Review

Bowlus, Lim, Lindor CGH Nov 2019

- 1. survey for cholangio and GB Ca in all adult patients with PSC regardless of disease stage especially in first year of dx and in patients with concomitant UC
- 2. US/CT or MRI with or without Ca 19-9 q6-12 months
- 3. no surveillance in patients less than 20 or those with small duct disease
- 4. strongly consider chole in pts with gb polyps
 8mm or greater

Scheduled ERCP dilation of dominant stricture in patients with PSC

Gut, Dec 2019



ERCP in Decompensated Cirrhosis

J Clin Exp Hepatol Sept 2019; Jagtap et al

India, January 2012 to December 2016

•	Probability of Mortality			54 y/o ETOH Cirrhosis undergoing CABG						
	7 days		30 days		90 days		1 year		5 years	
•	2.009	%	7.99	%	12.585	%	25.172	%	55.2	%

- One month mortality 8.5%
 - Bleeding, cholangitis, post ercp pancreatitis, cardiac, sbp, variceal bleed (3), pse
- MELD > 18, OR 5.6 (adverse events)

Validation of Amsterdam-Oxford model for PSC

JOH, Nov 2019 Goet et al

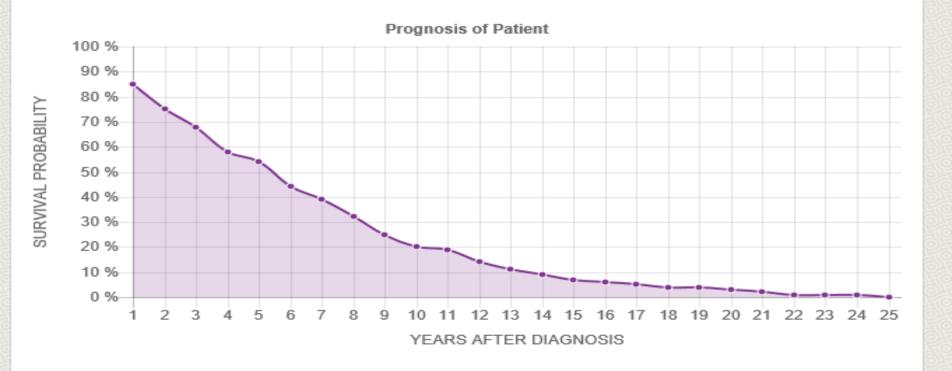
- 534 Patients with PSC
- Mean age 39 years; large duct in 87%
- 66% male; 92% on UDCA
- 60% UC
- 8 year f/u
- 7 parameters: large/small duct; age, albumin,
 AP, AST, TBr and platelets

http://www.fcbkapp.nl/psc/8/

PSC Subtype Large Duct Age at PSC Diagnosis 24 Years reset to original reference values Albumin lower limit of normal (LLN) 4.1 Alkaline Phosphatase (ALP) upper limit of normal (ULN) 400 120 U/L µkat/L Aspartate Aminotransferase (AST) upper limit of normal (ULN) 72 35 U/L µkat/L Billirubin upper limit of normal (ULN) 0.4 Platelets lower limit of normal (LLN) 240 x10°/L	Input Parameters					
Age at PSC Diagnosis 24 reset to original reference values Albumin lower limit of normal (LLN) 4.1 Alkaline Phosphatase (ALP) upper limit of normal (ULN) 400 120 U/L µkat/L Aspartate Aminotransferase (AST) upper limit of normal (ULN) 72 35 U/L µkat/L Bilirubin upper limit of normal (ULN) 0.4 Platelets lower limit of normal (LLN)	PSC Subtype					
reset to original reference values Albumin lower limit of normal (LLN) 4.1	Large Duct	-				
reset to original reference values Albumin lower limit of normal (LLN) 4.1	Age at PSC Diagnosis	to at DSC Diagnosis				
reset to original reference values Albumin lower limit of normal (LLN) 4.1	Age at 1 30 Diagnosis					
Albumin lower limit of normal (LLN) 4.1	24	4				
4.1		reset to origin	nal reference values			
Alkaline Phosphatase (ALP) upper limit of normal (ULN) 400	Albumin lower limit of normal (LLN)					
Alkaline Phosphatase (ALP) upper limit of normal (ULN) 400	4.1		g/I g/dI			
400	7.1	33	g, e g, de			
Aspartate Aminotransferase (AST) upper limit of normal (ULN) 72 35 U/L µkat/L Bilirubin upper limit of normal (ULN) 0.4 reference 17 µmol/L mg/dL Platelets lower limit of normal (LLN)	Alkaline Phosphatase (ALP) upper limit	t of normal (ULN)				
Aspartate Aminotransferase (AST) upper limit of normal (ULN) 72 35 U/L µkat/L Bilirubin upper limit of normal (ULN) 0.4 reference 17 µmol/L mg/dL Platelets lower limit of normal (LLN)	400		U/I West II			
72 Bilirubin upper limit of normal (ULN) 0.4 Platelets lower limit of normal (LLN) reference 17 µmol/L mg/dL	400	120	U/L μκατ/L			
72 Bilirubin upper limit of normal (ULN) 0.4 Platelets lower limit of normal (LLN) reference 17 μmol/L mg/dL	Aspartate Aminotransferase (AST)	upper limit of normal (UI	LN)			
Bilirubin upper limit of normal (ULN) 0.4		reference				
0.4 reference 17 µmol/L mg/dL Platelets lower limit of normal (LLN)	72	35	U/L µkat/L			
0.4 reference 17 µmol/L mg/dL Platelets lower limit of normal (LLN)	Pilipubin upos limit of pormal (III N)					
Platelets lower limit of normal (LLN)	Difficulti upper limit of normal (OLN)	reference				
reference	0.4	17	µmol/L mg/dL			
reference						
	Platelets lower limit of normal (LLN)					
240 x10°/L	240					
	240	150	X10°/L			

Output Parameters

Amsterdam-Oxford PSC Score	3.55
5-year transplant-free estimated survival	53.70%
10-year transplant-free estimated survival	20.25%
15-year transplant-free estimated survival	7.03%



Patients with PSC and Development of Validation of a Risk Scoring System

Hepatology May 2019; Goode Et al

- UK-PSC research cohort
- 1001 pts!

Uk-PSC.com

- Age
- TBr dx and year 2
- Platelets dx and year 2
- Hg at dx
- AP dx and year 2
- Extrahepatic disease at dx
- Variceal bleed by year 2
- My same patient:
- 5 year transplant free survival ~ 46% (vs 53% with Amsterdam)

Farnesoid X Receptor Agonist Cilofexor (GS 9674) in PSC OCA in PSC: Aesop trial;

Trauner et al Hepatology Sept 2019

OCA in PSC: Aesop trial; AASLD 2017

- 1. Phase II Double Blind Placebo Controlled
- 2. 13 sites; large duct, non cirrhotic
- 52 patients randomized (age 43, 58% male, 60% IBD; 46% UDCA)
- Baseline AP 348 (288-439)
- 21% reduction in AP Cilofexor at 12 weeks
 (.029 vs placebo), independent of UDCA use
- Well tolerated

Improved Outcomes of Patients With Primary Sclerosing Cholangitis

CGH, Stokkeland et al August 2019

- 1. Sweden 2005-2014; 2914 patients!
- 61% male, 41 years of age
 - 60% of patients on UDCA
 - 75% 5-ASA
 - 75% Azathioprine
 - 34% 6MP
- 14% statin
- Statin associated with reduction in all cause mortality: HR 0.68 (CI 0.54-0.88) and the need for LT
 - Aza also with similar decrease in all cause mortality and LT
- UDCA did not affect mortality



THE BEST OF THE LIVER MEETING® 2019

Cholestatic and Autoimmune Liver Diseases



Prospective validation of the prognostic value of liver stiffness (LS) (FibroScan®) in PSC: the FICUS study

Aim:

To assess prospectively the prognostic value of liver stiffness (LS) as evaluated by FibroScan® in patients with PSC

Methods:

- 13 institutions, across 11 countries
- 616 PSC patients included, analysis at 2 years in 514 with valid LS at inclusion

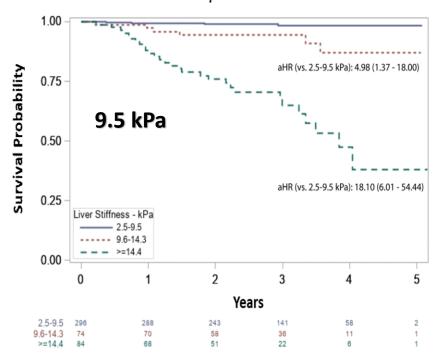
Main Findings:

Strong and independent (adjusted for bilirubin, ALP, and Mayo Risk Score) predictive value of LS

Conclusions:

This study validates the prognostic value of LS (FibroScan®) in PSC and supports the role of LS as a risk stratifying tool and potential surrogate endpoint in clinical trials.

Transplant-free Survival



Chazouilleres O, et al., Abstract 47





Bile acid profiles predict hepatic decompensation in primary sclerosing cholangitis

Aim:

- Patients with PSC often demonstrate increased plasma bile acid (BA) concentrations due to ongoing cholestasis.
- We aim to establish whether bile acid profiles are predictive of hepatic decompensation (HD): ascites, variceal hemorrhage, or encephalopathy.

Methods:

Plasma BA profiles and alkaline phosphatase levels of 425 patients with PSC were measured using clinically available assays and gradient boosting was used to build multivariable models predicting development of HD.

Main Findings:

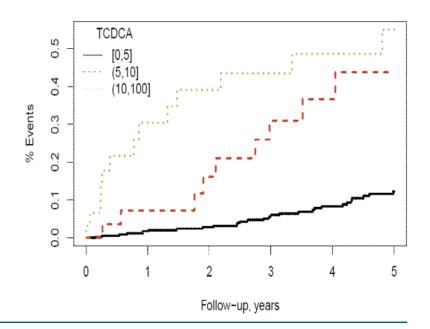
Using a machine learning technique, BA profiles are independently capable of predicting future HD events in PSC patients over 5 years.

Conclusions:

BA profiles have prognostic value and should be considered in disease management and as an exploratory endpoint in future clinical trials.

Event Rates of Hepatic Decompensation Stratified by the Range of TCDCA Concentration

TCDCA.gp4	N	Events	Time	Event rate	CI (rate)
[0,5]	365	29	1228	0.024	0.016 - 0.034
(5,10)	29	9	82	0.110	0.050 - 0.208
(10,100]	31	13	66	0.196	0.104 - 0.335



Mousa OY, et al., Abstract 188





PBC

Primary Biliary Cholangitis

formerly known as Primary Biliary Cirrhosis (2015)

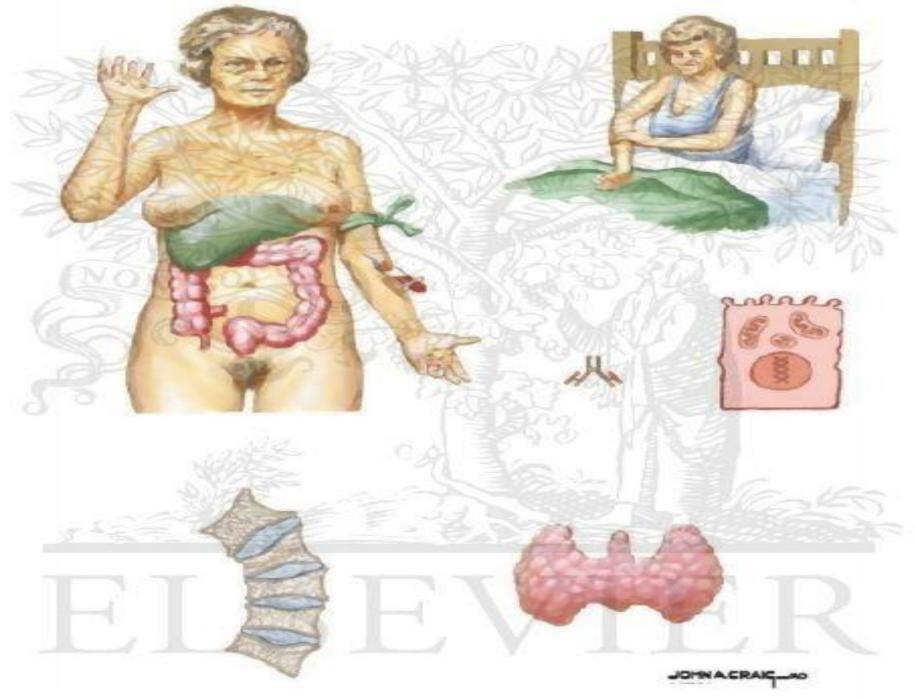
- Chronic liver disease resulting from progressive immunologic destruction of the small intralobular bile ducts that may result in cirrhosis and liver failure
- 95% of patients are female
- Diagnosis typically made 4th/5th decades of life
- Fatigue, itching (pruritus), abdominal discomfort (less common)
- 50% of pts are asymptomatic

PBC diagnosis: 2 of the following 3

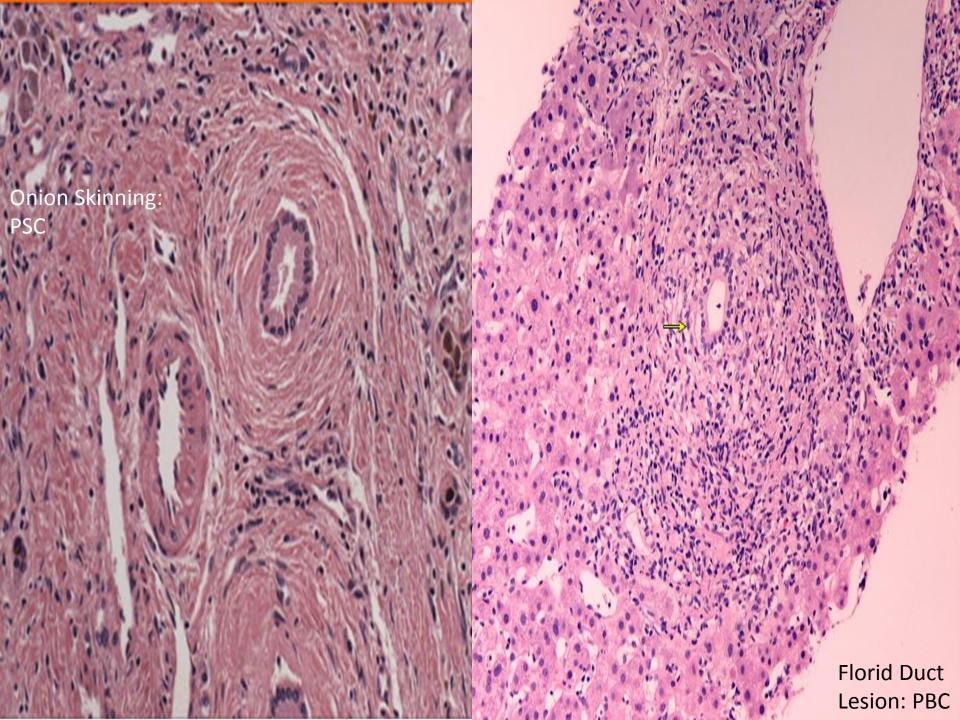
- Elevation in LFTs specifically AP/GGTP (cholestatic enzymes)
- Antimitochondrial antibodies (AMA) are the serologic hallmark; present in 95% of patients
- Liver biopsy classically shows 'florid duct lesion'

PBC

- PBC used to be one of the more common indications for liver transplant
- But since the approval of UDCA, the prognosis
 of pts with PBC (diagnosed before onset of
 cirrhosis) is quite good



© ELSEVIER, INC. - NETTERIMAGES.COM





I DON'T OFTEN HATE, BUT WHEN UPREMERTIO HATE THE NEW ENGLAND PATRIOTS.







Talken Tie Talk

STREET, STREET, SQUA

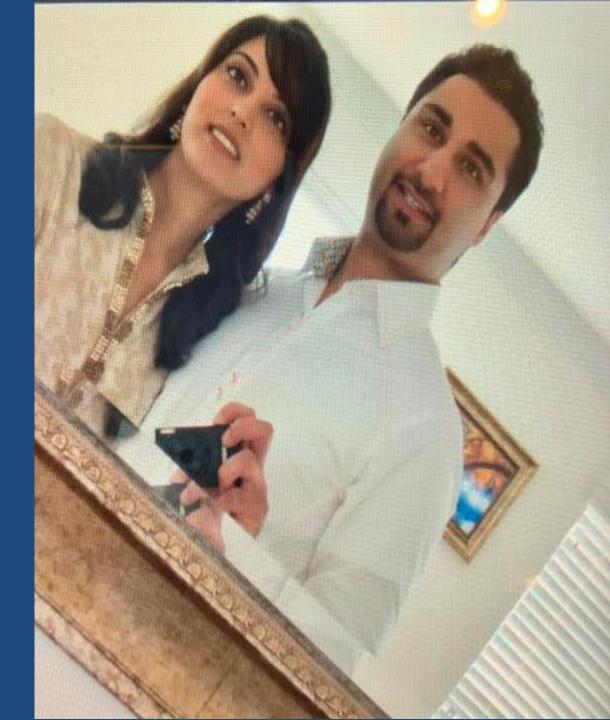


Boston 2009

- Wife: So...what do you do?
- Me: I'm a hepatologist
- Wife: What the heck is that
- Me: I'm a liver physician
- Wife: a liver what?
- Me: I'm a doctor
- Wife: I DO!



12.06.09





10 year anniversary

Reynolds Wrap Heavy Duty Aluminum Foil - 55 sq ft

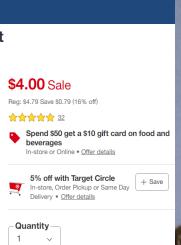
Shop all Reynolds













Age and Sex Response to UDCA and Tx Free Survival in PBC

Cheung et al, CGH Sept 2019

- Global PBC Study Group
- Retrospective study of 4355 patients with PBC
- 1961-2014
- Male patients were older at the start of treatment (58 versus 54)
- Younger pts (male and female; 45 years or less) were less likely to respond to therapy
- Risk of LT or death decreased significantly with advancing age
- Patients less than 35 especially (HR 14.59)

Incidence, Prevalence and Outcome of PBC in Sweden

Marschall et al; Sci Rep August 2019

- 1. 1987-2015
- 2. 5K patients with PBC matched to 10 controls
- Prevalence of PBC increased from 5/100K to 37 between 87 and 2014
- Female male ratio 4:1
- PBC individuals aged 15-39 had a substantially high risk of death (HR 12.7); risk was highest in men

Factors associated with progression of Early Stage PBC

CGH; August 2019 Gatselis et al

- 1. 1615 pateints, mean age 55 with early stage
 PBC (based on albumin and br)
- Median f/u 8 years
- Of the patients: 904 developed 'moderate' disease and 201 'advanced' disease over the study period
- 236 patients had a 'clinical event'
- UDCA and LFTs are biggest predictors!

BEZAFIBRATE AND PBC

- Placebo controlled trial of bezafibrate in PBC NEJM June 2018
- Phase III trial, Paris
- 100 patients who had inadequate response to UDCA
- 50/50 bezafibrate (500mg) vs placebo
- AP normalized 67% of pts in study arm versus
 2% placebo

Bezafibrate

- is a fibrate drug used as a lipid-lowering agent to treat hyperlipidaemia. It helps to lower LDL cholesterol and triglyceride in the blood, and increase HDL.
- It was patented in 1971 and approved for medical use in 1978
 - Not available in US
 - Gemfibrozil and fenofibrate

Bezafibrate and long term outcomes in PBC

Hepatology, Honda et al Feb 2019

- Bezafibrate is 2nd line agent in Japan for PBC
- From a cohort of 873 patients with PBC
- Enrolled 118 patients treated with combo UDCA and bezafibrate
- Combo therapy significantly decreased prognostic scores (GLOBE and UK-PBC) and long term prognosis (Tx free survival and liver related death)— especially in those with early PBC

Pruritus is common and undertreated in PBC in the UK

CGH June 2019 Hegade et al

- 1. > 2000 patients, self reported data
- 73% experienced pruritus
- Severe in 12%
- Younger age and AP elevation were associated with pruritus
- Less than 50% patients received medical therapy (cholestyramine, rifampin, naltrexone)

Smoking and risk of PBC: systemic review and Meta analysis

J Gastrointestin Liver Dis June 2019 Wijarnpreecha et al

- 1. Nine case control studies; 21K participants
- Smoking OR 1.31 risk of PBC versus non smokers

Trends in LT for PBC in Europe

Ailment Pharmacol Ther Feb 2019; Harms et al

- Patients undergoing LT from 1986 to 2015 in European LT Registry
- 6029 pts with PBC (5.3% of all transplants)
- Indication dropped from 20% in 1986 to only 4% of all transplants in 2015
- PBC was the only indication showing consistent decrease throughout al decades

Outcomes of LT for PBC; SRTR Database

DDS, Sayiner et al August 2019

- 1. 1994-2016
- 223,000 LT; 8K for PBC (3.6%)
- Mean age 55; 80% white; 86% female
- Mean MELD 21
- 6.6% retx
- Waitlist mortality 18%
- Tx rate was 57.7% (versus 53% HCV)
- Waitlist dropout 25%
- Mortality and graft loss were significantly lower compared to HCV

Recurrence of PBC after LT

Gastro Jan 2019 Montano-Loza et al

- 785 (89% female) patients with PBC who underwent LT;
 mean age 54 years; mean f/u 7 y ears
- 1983-June 2016; USA and Europe
- Recurrence occurred in 22% of patients at 5 years and 36% at 10 years
- Factors associated with recurrence:
 - 1. age < 50 at diagnosis of PBC</p>
 - Age of LT less than 60 years
 - Use of FK (HR 2.31)
 - Increased AP and TBr 6 months post LT
 - Recurrence was associated with increase risk of graft loss.
 - We were not in this study, but for the record UPMC alone has done 600 PBC transplants! This study was the composite of 16 European centers



THE BEST OF THE LIVER MEETING® 2019

Cholestatic and Autoimmune Liver Diseases



Bezafibrate is superior to placebo in improving pruritus in chronic cholestatic liver disease: the FITCH trial

Hypothesis:

Bezafibrate may relieve cholestasis-associated pruritus by alleviating hepatobiliary inflammation and thereby reducing secretion of a biliary itch factor (*Hepatology* 2014;60:399-407).

Methods:

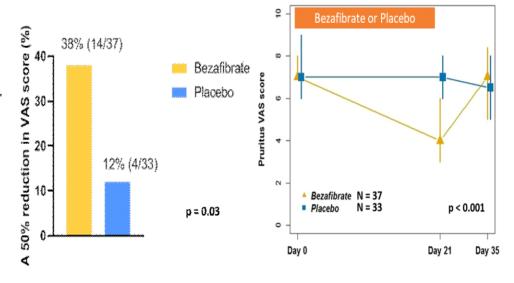
Double-blind, randomized, placebo-controlled trial in patients with PSC (n=44), PBC (24), or secondary sclerosing cholangitis (2) with moderate to severe cholestasis-associated pruritus.

Main Findings:

Bezafibrate (400 mg daily) led in 38% of the patients to \geq 50% reduction of pruritus whereas patients treated with placebo reached the primary endpoint in 12% (p=0.03; see figure).

Conclusions:

Bezafibrate is superior to placebo in improving pruritus in chronic cholestatic liver diseases such as PSC and PBC.



Vries de ES, et al., Abstract 13





Durable response in the markers of cholestasis through 5 years of OLE study of OCA in PBC patients

Objective:

Open-label extension (OLE) of the POISE phase 3, randomized, double-blind, placebo-controlled, study in patients with PBC to assess long-term safety of obeticholic acid (OCA) and durability of effects on serum markers of cholestasis

Methods:

Following the 1-year double-blind phase, patients on placebo started OCA and were then pooled with OCA-treated patients to evaluate the efficacy and safety of up to 6 years of OCA treatment.

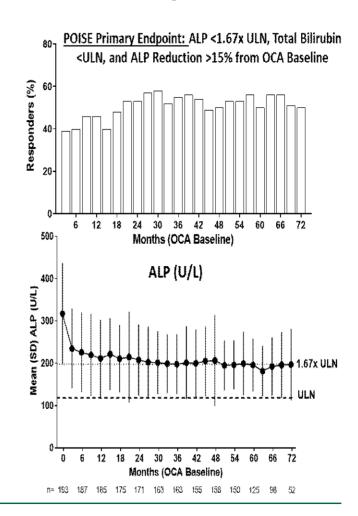
Main Findings:

- ALP, AST, GGT, and key inflammatory markers were significantly reduced; reductions were sustained for the duration of treatment.
- Total bilirubin was maintained within the normal range.

Conclusions:

Six years of treatment with OCA resulted in sustained improvements in markers of cholestasis and inflammation and in stabilization of liver stiffness with no new safety observations.

Nevens F, et al., Abstract LO6





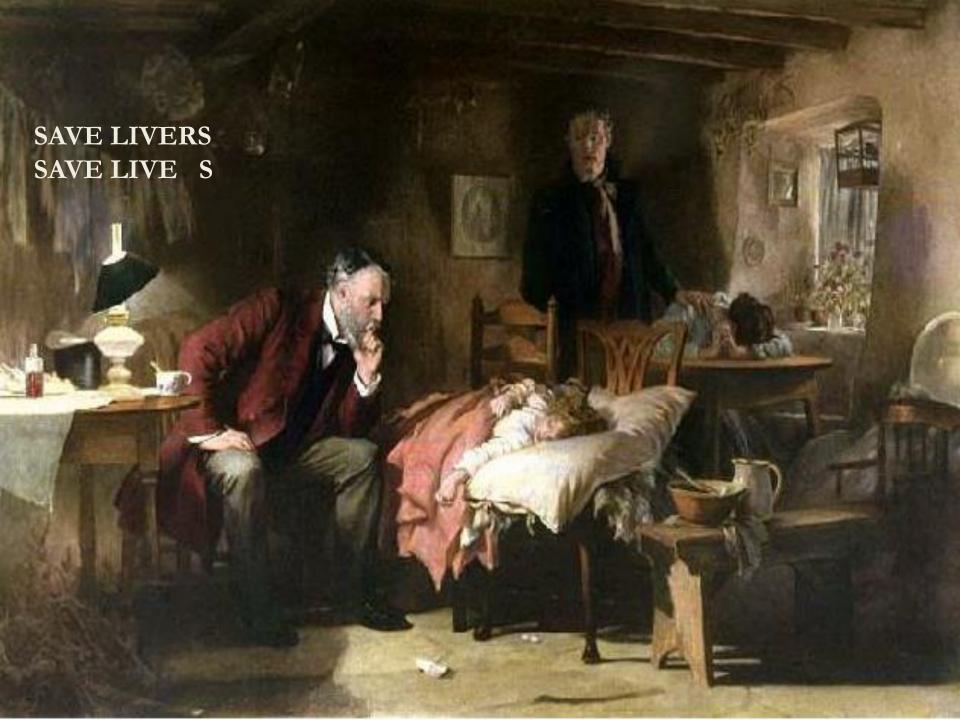


Eleven Take Home Points

- 1. Scheduled dilation of dominant stricture may have mortality benefit in patients with PSC
- 2. ERCP should be performed with expert multi-disciplinary assessment
 - ERCP in pts with cirrhosis (esp B/C; MELD > 18) carries a one month mortality near 10%
- 3. The Amsterdam-Oxford and UK-PSC online validation scores offer prognostic information with readily available clinical data
- 4. FibroScan also has very good prognostic value in PSC
- 5. Statins may have mortality benefit in pts with PSC.
 - Ongoing study of FXR Agonist.

Take home points (cont)

- 6. Younger patients (less than 35) dx with PBC have a more aggressive course
- 7. Although many patients with PBC will respond to UDCA up to a 1/3rd will not and AP is probably the best short term clinical predictor
- 8. Role of Bezafibrate in mortality benefit and sx relief
- 9. OCA has durable (biochemical) response at 5 years in patients with PBC
- 10. PBC as an indication for LT has dropped nearly five fold; Recurrence of PBC post LT is not uncommon
- 11. Probably should think twice before agreeing to give a talk on a Saturday the day following your ten year anniversary



2019 Annual Update in Medical Hepatology

Latest Advances in Cholestatic Diseases



Shahid M. Malik, MD Assistant Professor of Medicine Program Director Transplant Hepatology Fellowship

