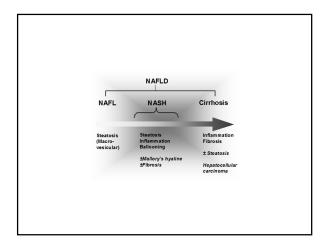
### **Definition of NASH**

Nonalcoholic steatohepatitis (NASH) is the term used to describe the distinct clinical entity in which patients lack a history of significant alcohol consumption but have liver biopsy findings indistinguishable from alcoholic hepatitits.

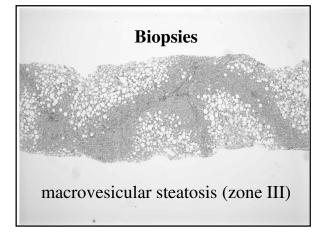


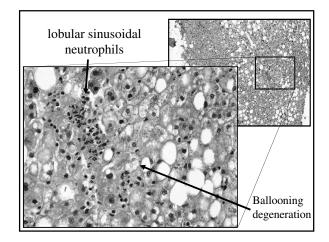
# **Criteria for Dx of NASH**

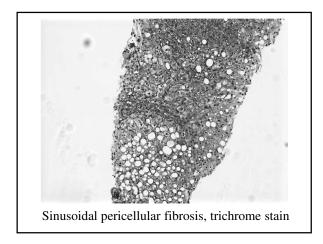
- Liver bx shows macrovesicular fatty change with inflammation and with or without Mallory bodies, fibrosis or cirrhosis.
- Convincing evidence of negligible alcohol consumption (less than 10 g/day of alcohol for women and less than 20 g/day for men).

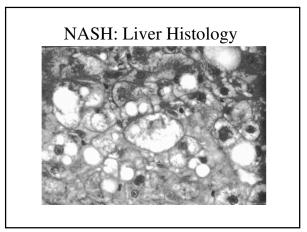
## **Liver Ultrasound Report**

The liver is enlarged. It is diffusely echogenic consistent with fat infiltration of the liver. Other forms of liver disease and more advanced liver disease including early cirrhosis cannot be excluded by this study.



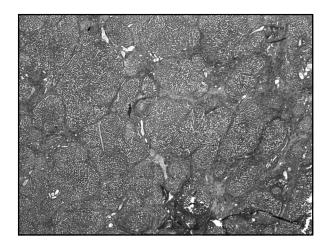






## **Proper Interpretation of the Histology**

...these features are consistent with a Toxic Metabolic etiology.



NASH is Likely a Major Cause of Cryptogenic Cirrhosis

- 74% of 70 consecutive patients with cryptogenic cirrhosis had obesity and/or diabetes
- This percentage is similar to prevalence of obesity and/or diabetes noted in 50 consecutive patients with NASH

## **Epidemiology of NAFLD**

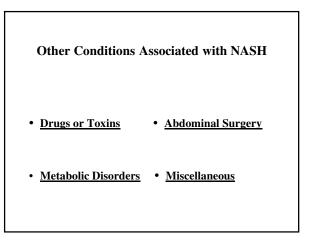
- Estimate: 40 million Americans have NAFLD 5 million are cirrhotic
- Majority of Pts with NASH have metabolic syndrome
- Over age 60 years, 40% have metabolic syndrome

NAFLD is Part of the Metabolic Syndrome Called Syndrome X			
Obesity	Hypertriglyceridemia		
Hyperinsulinemia	Hypertension		
Insulin Resistance	Diabetes		

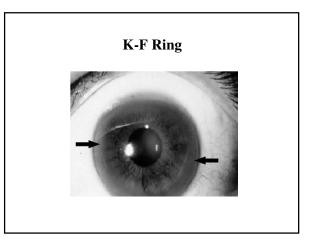
Regular sugar-sweetened beverage consumption is associated with a greater risk of fatty liver disease

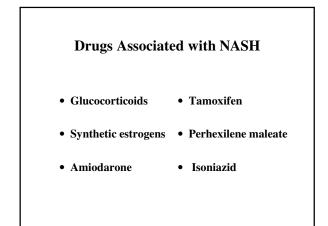
- Examination of cross-sectional association between intake of sugar-sweetened beverages and fatty liver disease in Framingham Heart Study cohorts
- J Ma et al., Journal of Hepatology 2015 May 29

Patients with NASH				
	Normal ALT	Increased ALT	p.value	
Fibrosis Stage 2 or greater	22%	34%	NS	
		Fracanzani AL, et al. Hepatology 2008:48;792		

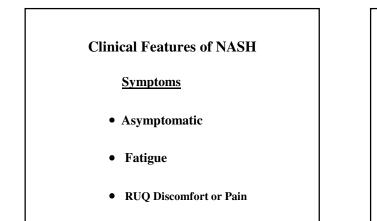


One disorder that is **critical** to exclude in young individuals is **Wilson's disease** 



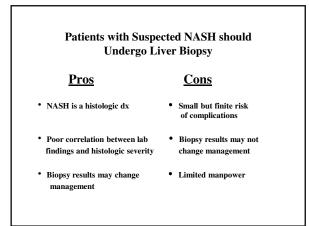


Progression to Cirrhosis	10 Yr Survival
38 -50 %	20%
8-26%	60%



## Laboratory Features of NASH

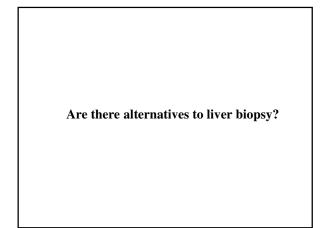
- ALT and AST 2 to 4 fold elevated in most patients
- Alk phos mildly elevated in a third of patients
- Albumin, PT, bilirubin most often normal
- Serum ferritin elevated in half the patients

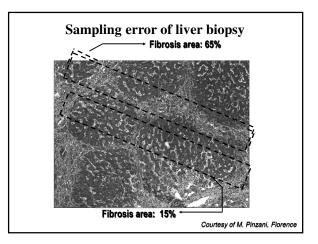


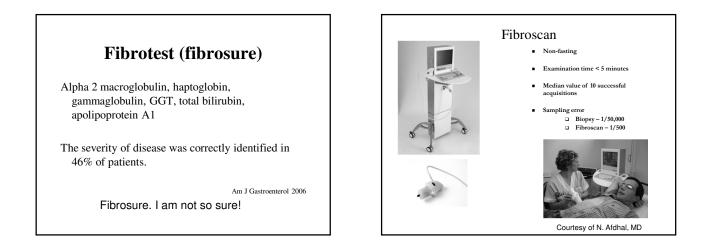
#### Suspected NASH: Reasonable to perform liver biopsy

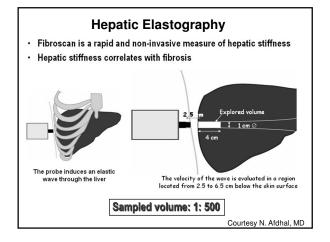
If any of following present

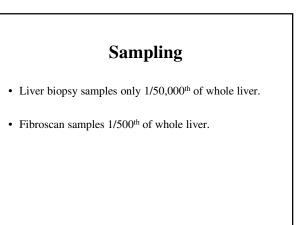
- Peripheral stigmata of chronic liver disease
- Splenomegaly
- Cytopenia
- Abnormal iron studies
- Diabetes and/or significant obesity in a patient over 45 years of age with elevated ALT

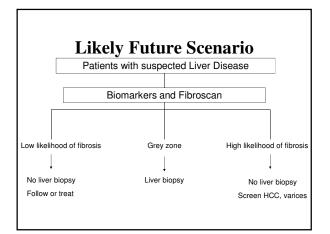






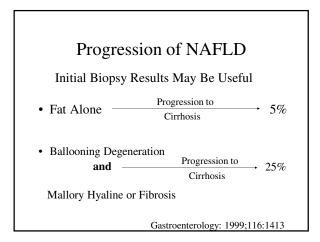


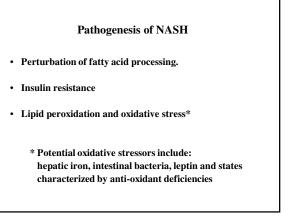


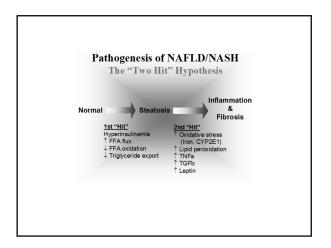


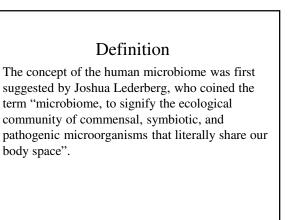
Transient elastography (Fibroscan) is accurate in most patients with NAFLD. With high negative predictive value and modest positive predictive value, Fibroscan is useful as a screening test to exclude advances fibrosis.

> Wong V W-S, et al. Hepatology 2010; 51:454-462













Welcome to Boston Population: 650,000

Welcome to Gut Population: 100 trillion

# The Gut Microbiota and NAFLD

- Microbiota in addition to regulating body fat gain and insulin resistance:
  - Change gene expression
  - · Increase energy harvest from diet
  - Produce ethanol
  - · Affect inflammation and immunity

E Lau et al. Gut Microbiota: Association with NAFLD and Metabolic Disturbances. Biomedical Research International. 2015.

## Prevention and Treatment of NASH

- Prevention of obesity and metabolic syndrome
- Treatment of metabolic syndrome
- Coffee ?
- Bariatric Surgery if appropriate

Pioglitazone therapy over a 12 month period in nondiabetic NASH patients resulted in improvement in biochemical, metabolic and histological parameters (including fibrosis).

> Aithal GP, et al. Gastroenterology 2008: 135;1176

#### NASH: Weight Loss is Beneficial !

- 1. 9% or greater weight loss resulted in:
- · Biochemical improvement
- Histologic improvement (steatosis, ballooning, inflamm<sup>n</sup>)
- Improvement in Insulin resistance
- · Higher Adiponectin levels

Harrison SA, et al. Hepatology 2009;49:80

2. Following bariatric surgery, Hepatic Fibrosis improved or was reversed in 66%.

Furuya CK Jr., et al. J Gastro Hepatol 2007;22:510

#### Vitamin E May Have Benefit

- 247 Adults with NASH (without diabetes) randomly assigned to pioglitazone (30 mg. daily) Vitamin E (800 IU daily) or placebo for 96 weeks\*.
- Vitamin E group had significant improvement in global histology scores compared with placebo (43% vs 19%).
- Concerns regarding Vitamin E and increased mortality have led many Hepatologists to not recommend Vitamin E or to use 400 IU daily.

\* Sanyal, A, et al. Pioglitazone, vitamine E, or placebo for nonalcoholic steatohepatitis. N Engl J Med 2010

## Other Pharmacological Treatment Modalities

Pioglitazone
 Ursodeoxycholic acid (UDCA)

Losartan

• Omega-3 Fatty Acids

- Metformin
- Liraglutide
- Orlistat
  Atorvastatin
- Probucol
  Pentoxifylline
- Betaine

Of limited or no proven efficacy

 Who is this person and how old was he when he died? Voltaire (1694 – 1778) 83 years

2. How many cups of coffee did he drink every day?

50-72 !

# Mechanism of Protective Effect of Coffee Unknown

- Caffeine, cafestol and kahweol protective in experimental studies
- Antioxidant effect
- Insulin sensitizing effect
- Coffee drinkers have higher levels of plasma adiponectin

More than 1 Billion People in the World Have Chronic Liver Disease

Consuming two cups of coffee per day reduces hospitalization rate and mortality from chronic liver disease by more than 50%

Ruhl CE, et al. Gastroenterology 2005;129:1928

## References: Coffee and Liver Disease, 2012 Sanjiv Chopra, MD, MACP

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2. Casiglia E, et al. Unexpected effects of coffee consumption on liver enzymes. Eur J Epidemiol. 1993;9:293.

3. Tverdal A, Skurveit S. Coffee intake and mortality from liver cirrhosis. Ann Epidemiol. 2003;13:419.

4. Shimazu T, et al. Coffee consumption and the risk of primary liver cancer; pooled analysis of two prospective studies in Japan. Int J Cancer. 2005;10:150.

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 Klatsky AL, Morton C, Udaltsova N, Friedman G. Coffee, cirrhosis, and transaminase enzymes. Arch Intern Med. 2006;166:1190.

7. Hu G, Tuomilehto J, Pukkala E, Hakulinen T, Antikainen R, Vartiainen E, Jousilahti P. Joint effects of coffee consumption and serum gamma-glutamyltransferase on the risk of liver cancer. Hepatology. 2008;48(1):129-36.

8. Larsson SC, Wolk A. Coffee consumption and risk of liver cancer: a meta-analysis. Gastroenterology. 2007;132(5):1740-5.

 Corrao G, Zambon A, Bagnardi V, D'Amicis A, Klatsky A; Collaborative SIDECIR Group. Coffee, caffeine, and the risk of liver cirrhosis. Ann Epidemiol. 2001;11(7):458-65.

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 Schilter B, Perrin I, Cavin C, Huggett AC. Placental glutathione Stransferase (GST-P) induction as a potential mechanism for the anticarcinogenic effect of the coffee-specific components cafestol and kahweol. Carcinogenesis. 1996;17(11):2377-84.

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 Kalthoff S, Ehmer U, Freiberg N, Manns MP, Strassburg CP. Coffee induces expression of glucuronosyltransferases by the aryl hydrocarbon receptor and Nrf2 in liver and stomach. Gastroenterology. 2010;139(5):1699-710.

15. Freedman ND et al. Coffee consumption is associated with response to peginterferon and ribavirin therapy in patients with chronic hepatitis C. Gastroenterology 2011; 140: 161-69.

 Molloy JW, Calcagno CJ, Williams CD, Jones FJ, Torres DM, Harrison SA. Association of coffee and caffeine consumption with fatty liver disease, nonalcoholic steatohepatitis, and degree of hepatic fibrosis. Hepatology. 2012;55(2):429-36.

Large prospective study; Coffee consumption inversely associated with total and cause-specific mortality.

 Freedman, ND Ph.D., Park, Y Sc.D., Abnet, CC Ph.D., et al. Association of Coffee Drinking with Total and Cause-Specific Mortality N Engl J Med 2012; 366:1891-1904 "Coffee is so good, the infidels should not have exclusive use of it."

**Pope Vincent III** 

**Potential Approaches to Treatment in the Future** 

- Will likely include combination therapy and life-style changes.
- Experimental study in a rat model of NASH combining angiotensin II receptor blocker with an oral iron chelator attenuated progression.
- Moderate exercise and coffee consumption likely of benefit.

#### Patients with Elevated Transaminases are <u>not</u> at Higher Risk for Statin Hepatotoxicity

		Mild-Modera <u>Elevations</u>	te Severe <u>Elevations</u>
•	1439 with normal transaminases prescribed a statin	1.9%	0.2%
•	342 with elevated transaminases prescribed a statin	4.7%	0.6 %
ŀ	2245 with elevated transaminases <u>not</u> prescribed a statin	6.4%	0.4%
		Chalasani et al: G	astroenterology 2004;126

## **Obeticholic Acid (OCA) Treatment**

- OCA is a semisynthetic bile acid analogue and a Farsenoid X Receptor (FXR\*) ligand
- 283 patients with NASH enrolled in a OCA 25 mg p.o. vs. Placebo trial for 72 weeks
- In January 2014, trial stopped on basis of interim efficacy analysis

\*FXR is a key player in control of multiple metabolic pathways – bile acids, glucose, lipids

### Quiz Answer True or False

- 1. NAFLD is the most common hepatic disorder in the U.S.
- 2. Serum ferritin is elevated in 50% of pts with NASH.
- 3. NASH has been reported in children.
- 4. Progression to cirrhosis occurs in 15-20% of pts.
- 5. NASH is likely the leading cause of cryptogenic cirrhosis.

Quiz (Continued) Answer True or False

- 6. The histologic features of NASH maybe seen in Wilson's disease.
- 7. Both Amiodarone and Tamoxifen can cause NASH.
- 8. Primary hepatocellular carcinoma has been reported in patients with NASH and cirrhosis.