

# Clinical Case

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#### Disclosures

Partner of HEPATOSCAN Medical Services® - Commercial interest in transient liver elastography using FibroScan®.



# Meet the patient

#### Patient details

- Mrs B
- 56 yr; female
- Professional lifestyle
- Recently moved into the area visits GP for fatigue on walking 2-3 blocks, feels depressed and anxious, now referred for abnormal LFTs
- Non smoker, does not drink alcohol
- T2DM x 5 yrs
- Hypertension x 10 yr

#### **Examination**

- BMI 38 kg/m2
- BP 144/92
- Heart- normal heart sounds,
- Chest- clear
- Anterior liver span 5 cm by percussion
- No ascites

# Laboratory investigations

- Hgb: 13 gm/dl
- Platelets: 180000/mm3
- ALT: 24 IU/L;
   AST: 20 IU/L
- LDL-Cholesterol
  - 160 mg/dl, TG 180 mg/dl, HDL-c 38 mg/dl
- GGT- 320 IU/l
- Creatinine 1.02 mg/dl
- Hgb A1C 9.9

# Additional investigations

- Fibroscan-
  - TE 12 kPa
  - CAP 350 db/m

Echo: Increased E/E'
Biventricular hypertrophy

O2 sat 90% room air



#### - Evaluation

FIB-4: 1.27 - < F3

NAFLD Fibrosis Score: - 1.4 - < F2



- A) Should I perform liver biopsy? Patient with the possibility of steatohepatitis (NASH).
- B) Lifestyle-modifying measures and repeat ultrasound in 6 months?
- C) Initiation of liraglutide? Obese and diabetic patient probable with steatohepatitis. Another drug for weight loss or NAH?



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#### - 6 months later...

Physical exam: BMI = 37 Kg/m<sup>2</sup>

Abdominal ultrasound - severe steatosis and hepatomegaly.

#### - Lab Results

Platelets: 158.000

A1C: 7.2%

ALT: 48 IU/l (5-40 IU/l)

AST: 50 IU/l (5-40 IU/l)

Albumin 3,8 g/L



#### - Evaluation

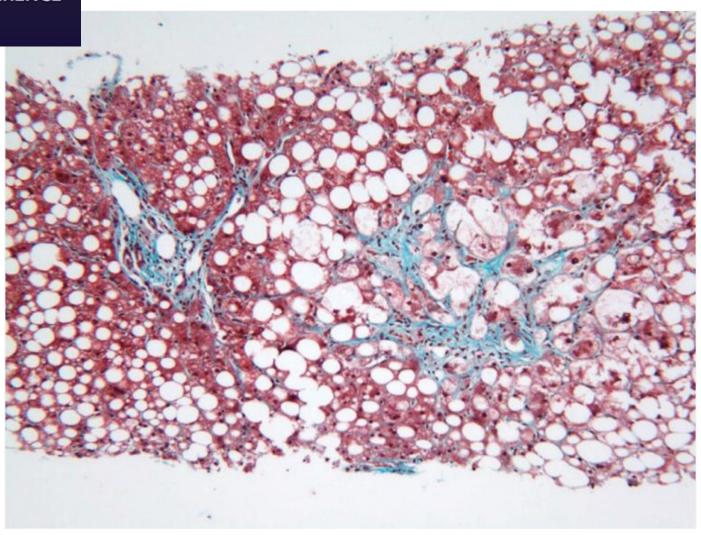
NAFLD Fibrosis Score: 2.6 - F3-F4 FIB - 4: 2.56 - gray zone

TE (XL): 10,3 kPa/ IQR 18%/ SR 100%/ CAP 350 dB/m/ IQR 40 - F3



- A) Should I perform liver biopsy?
- B) Reinforce lifestyle-modifying measures and repeat ultrasound/ NITs in 6 months?
- C) Initiate liraglutide? Another drug?







- A) Continue to follow US/NITs and lifestyle modification?
- B) Start vitamin E and new evaluation at 6 months?
- C) Start pioglitazone and new evaluation at 6 months?

# Liver and diabetes Cause or consequence?



### NAFLD - Nonalcoholic Fatty Liver Disease

N = 132,377 35-79y FU = 8y Non diabetic individuals

✓ A total of 6,555 incident diabetes (3,734 men and 2,821 women) were identified, on average, over 5.8 years of follow-up.

✓ The risk of incident diabetes was significantly associated with NAFLD [HR=2.08 (men) and 2.65 (women)].

✓ Elevated ALT, AST, GGT and ALP were also significantly associated with the increased risk of diabetes.

# NAFLD - Nonalcoholic Fatty Liver Disease and diabetes

- ✓ In longitudinal studies with histological assessment, diabetes was a risk factor for NAFLD fibrosis progression.
- ✓ Higher mortality from cardiovascular events and complications related to chronic liver disease have been reported in patients with NAFLD and diabetes.

Adams, et al. Gastroenterology 2005; Ekstedt, et al. Hepatology 2006 Rafiq, et al. Clin Gastroenterol Hepatol 2009 Targer, et al. Diabetes Care 2007 Stepanova, et al. Dig Dis Sci 2013

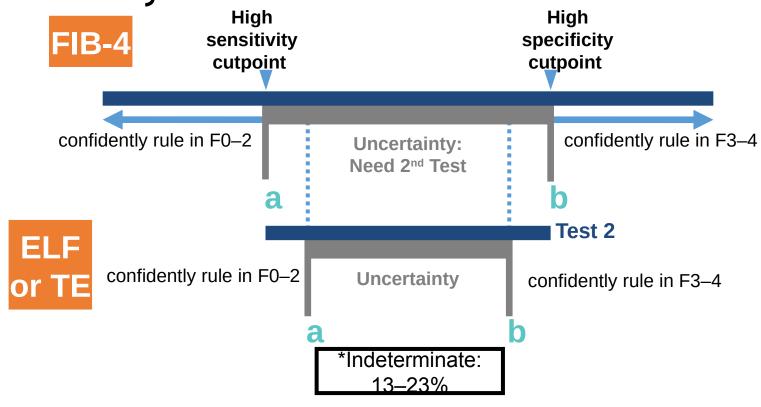
# What is the role of ALT and/or AST in the diagnosis of NASH and fibrosis in DM2 patients?

In patients with DM2 and steatohepatitis aminotransferases were elevated only in 15-38%.

Leite et al. Liver Int 2011 Prashanth et al. J Assoc Physicians India 2009.

Repeatedly normal ALT and/or AST do not exclude higher degrees of inflammatory activity or advanced stages of fibrosis.

Leite et al. Liver Int 2011 Williamson RM. QJM 2012 Portillo Sanchez et al. J Clin Endocrinol Metab 2015 Combination of two NITs to discriminate advanced fibrosis due to NASH increases accuracy



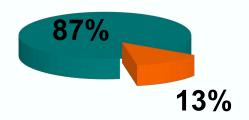
Reduces the indeterminate zone by mimicking clinical practice with initial use of a simple test (e.g. FIB-4), followed by a specialty test (e.g. ELF or LS by TE)

# Evaluation of fibrosis in our cohort of diabetics with NAFLD

- 303 exams
- 61% female; 59 ± 9 years.
- BMI: 31 ±5 Kg/m<sup>2</sup>

Advanced fibrosis (TE ≥9.6 kPa)

Studies with TE: prevalences 7.3-18.8% of advanced fibrosis



Sporea I et al. J Gastrointestin Liver Dis. 2016; Kwok et al. Gut 2015; Roulot et al. Liver Int 2017

