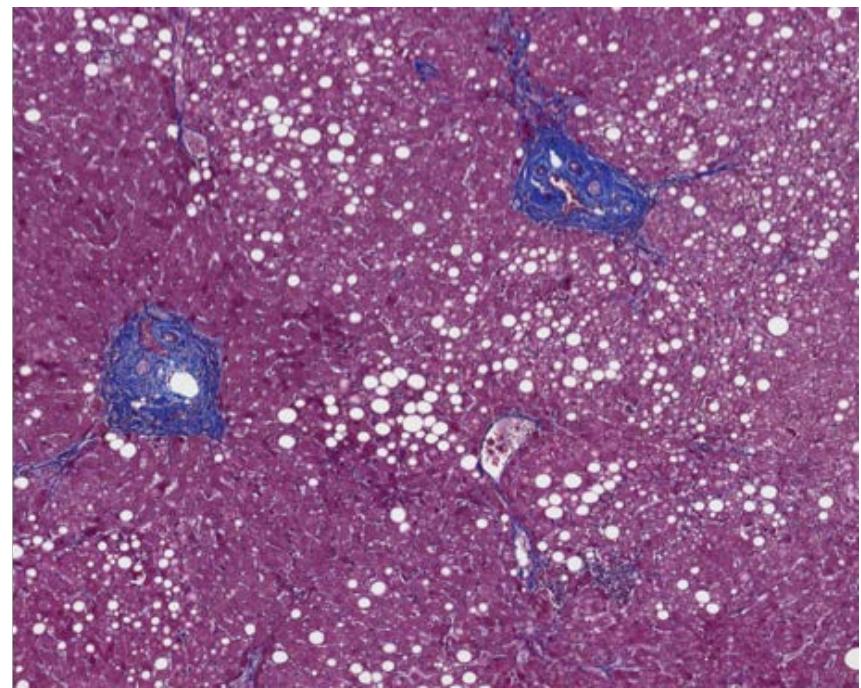


Stéatose

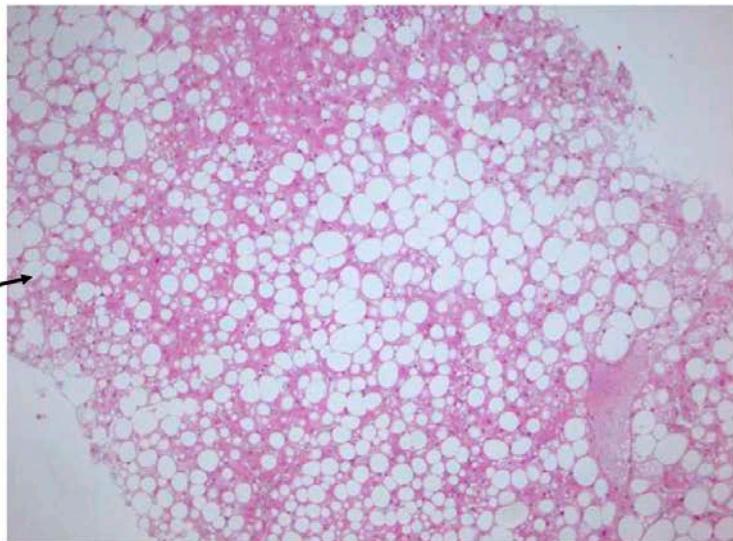
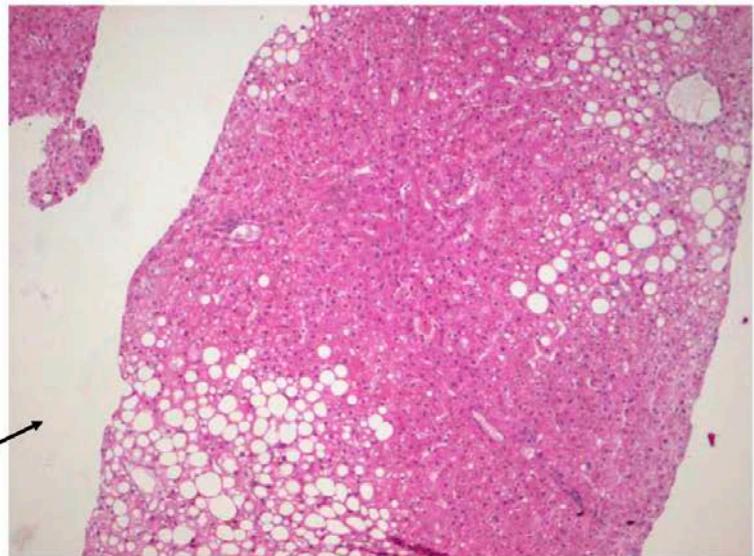
- US : hyperéchogène
- TDM : hypodense
- IRM : hypointense en T1 out of phase
- Stéatose peut être hétérogène



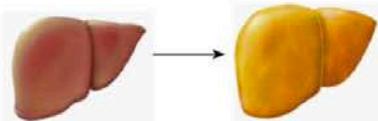


Stéatose

- Gradation : 5 stades
 - 0 % / 1-5 %
 - 6-33 %
 - 34-66 %
 - $\geq 67 \dots$



Normal liver Fatty liver



Stéatose enjeu de l'échographie



hyperéchogénicité :
objective

diffuse



signes écho à partir de 30 %
hyperéchogenicité % teneur en graisse

gradient hépato rénal facile
mais difficile de quantifier et non spécifique

US et maladie de surcharge: stéatose // hémochromatose



Gradient hépato-rénal:

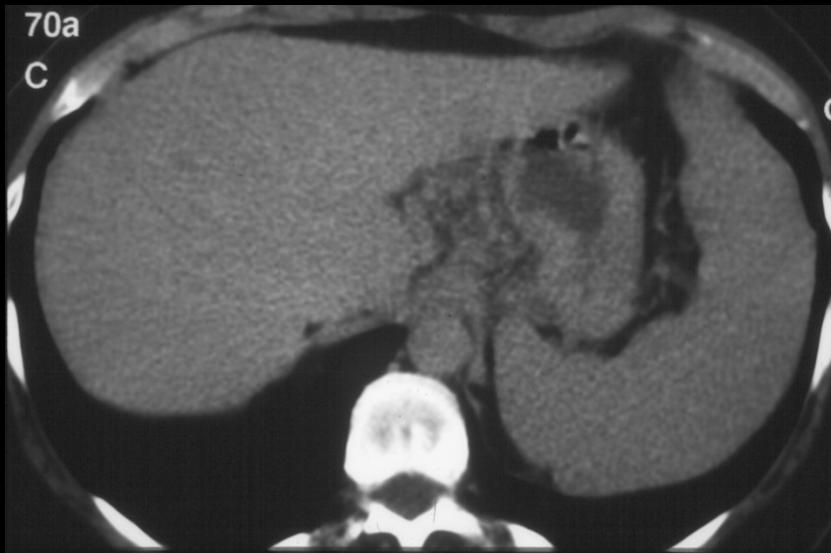
- normal-

- majoré-

- majoré



70a
C



2399/1
.... 80/1
Ax 1800.1

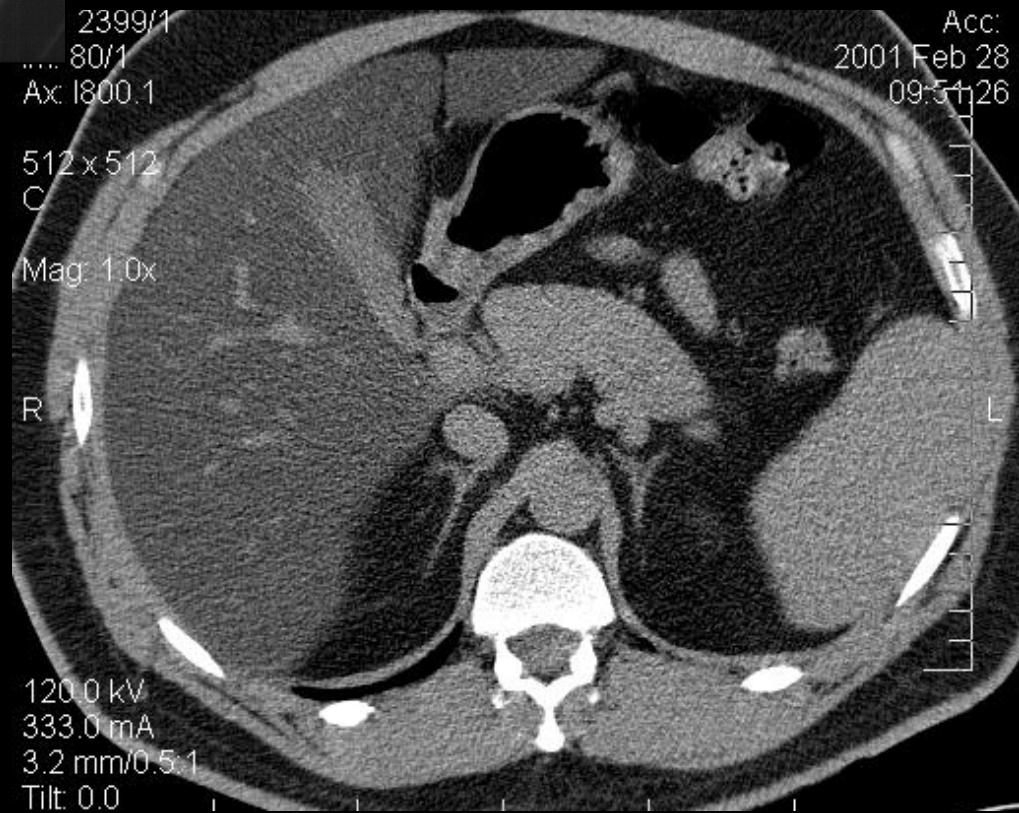
512x512
C

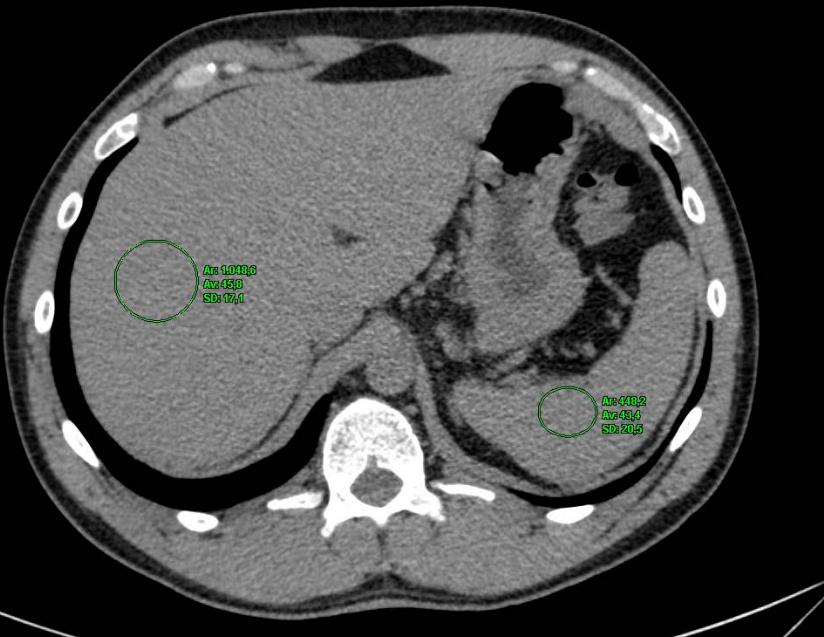
Mag: 1.0x

R

120.0 KV
333.0 mA
3.2 mm/0.5:1
Tilt: 0.0

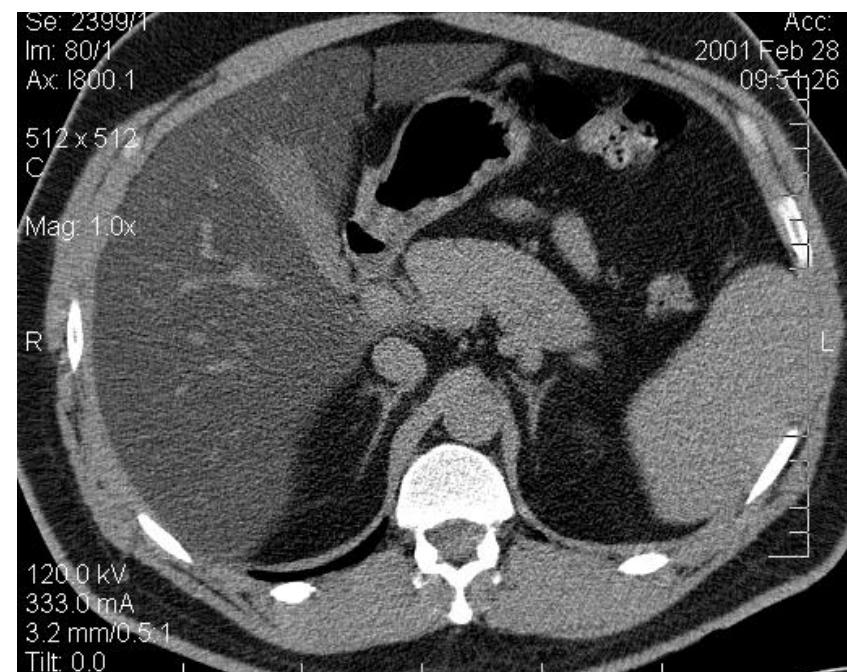
Acc:
2001 Feb 28
09:54:26





CT et maladie de surcharge : stéatose

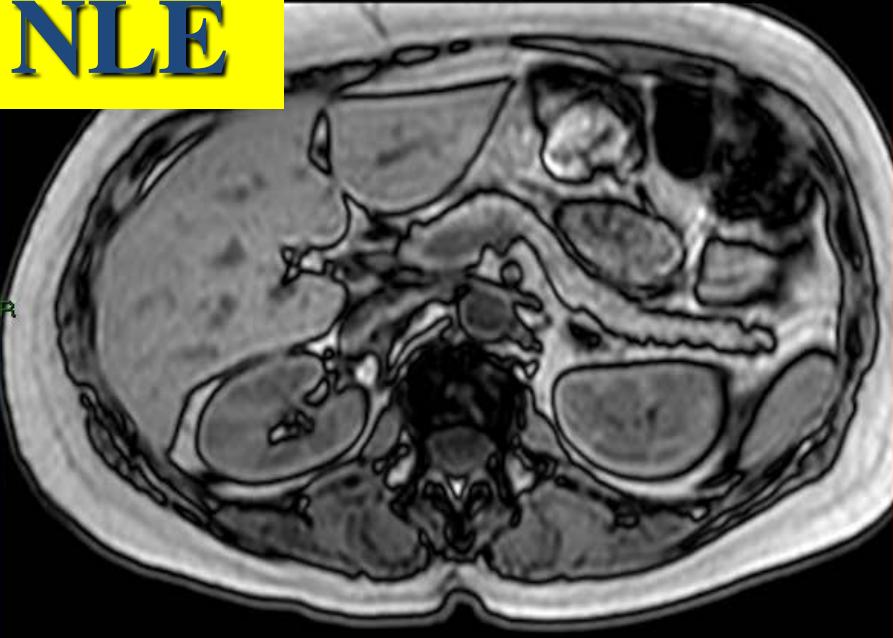
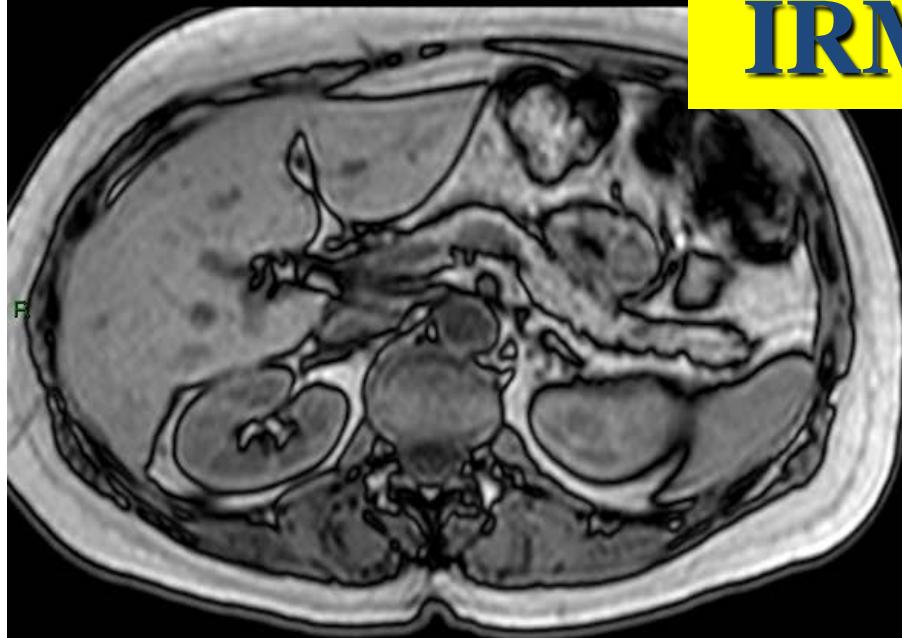
Densité UH foie 46 et rate 43



CT sans injection

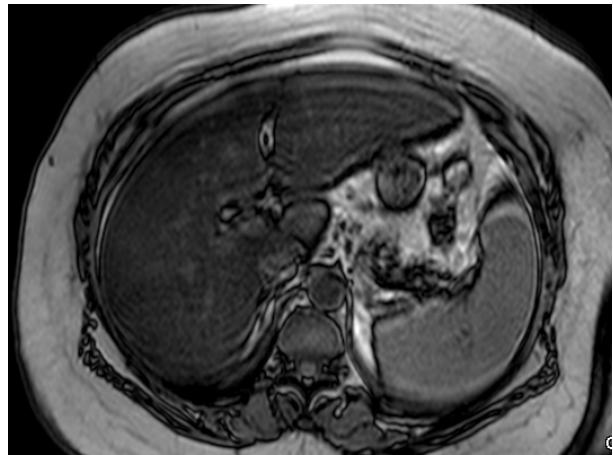
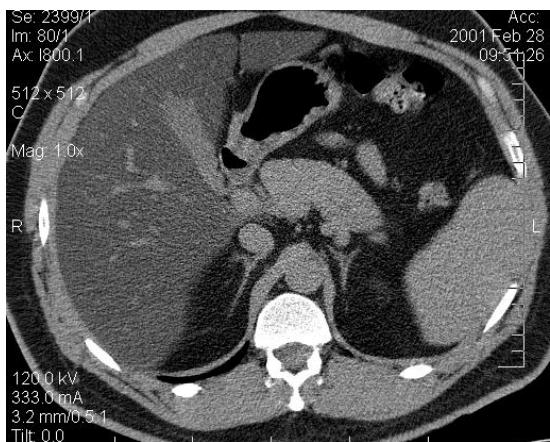
Densité UH foie 27 et rate 40

IRM NLE



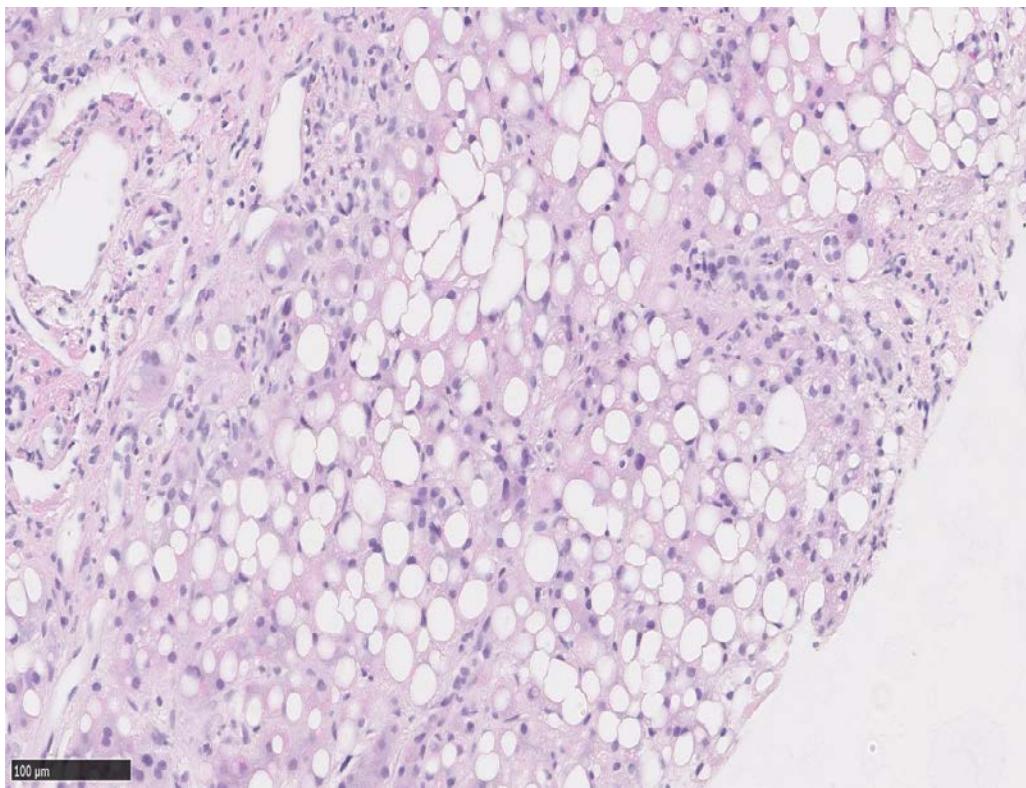
IRM stéatose





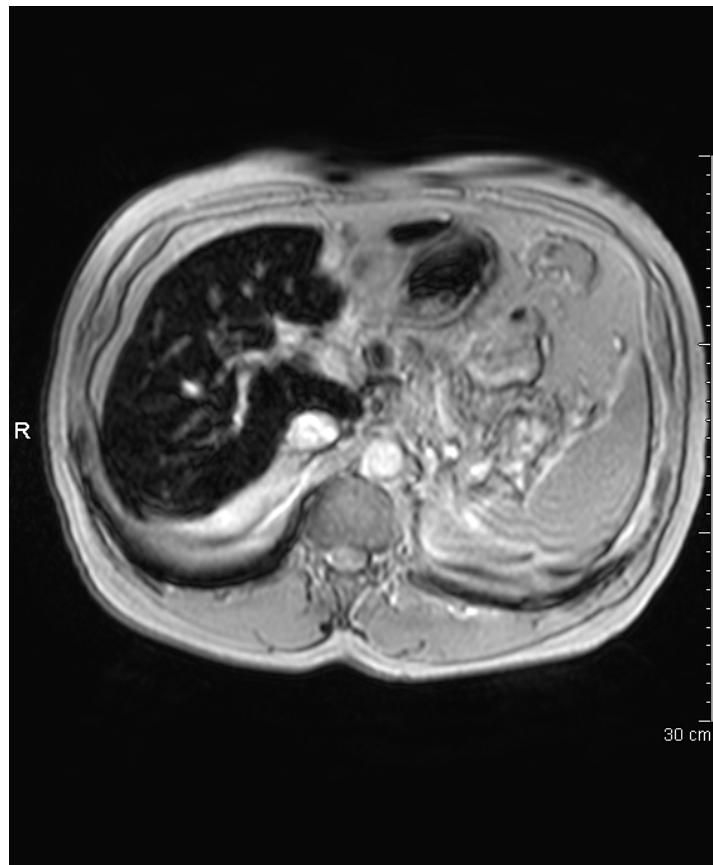


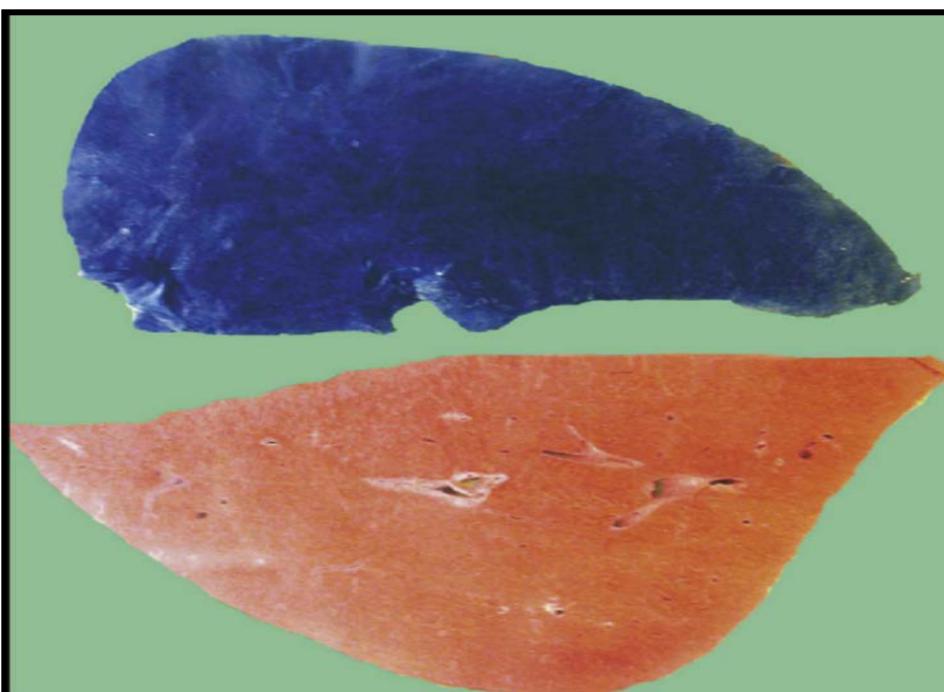
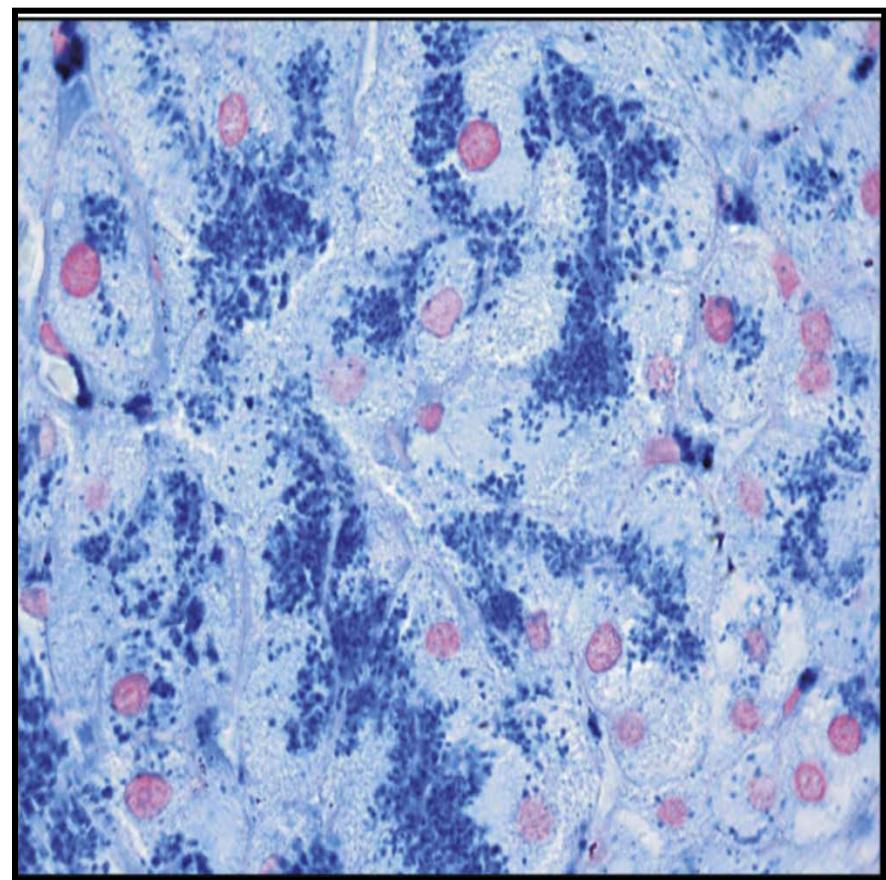
Steatose



Hémochromatose

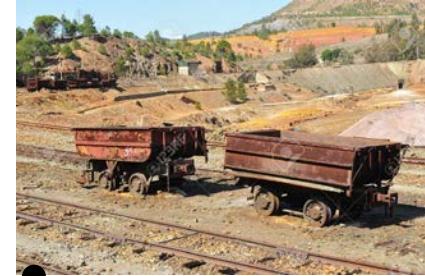
- IRM : hyposignal en particulier en T2 et en écho de gradient
- Méthode de Calcul automatisée





Hémochromatose-Stéatose

- Hémochromatose :
 - isolée : pas de signe de surcharge en fer visible en écho
 - compliquée d'une stéatose : on voit la stéatose en écho
 - Évolution vers la fibrose puis la cirrhose et ses complications
- Hyperferritinémie
 - NAFLD et AFLD



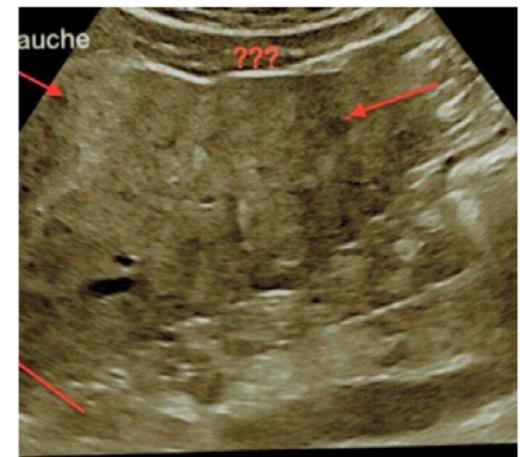
Cirrhose

- Atrophie en particulier du lobe droit
- Hypertrophie du lobe caudé et du lobe gauche
- Contours nodulaires
- Hypertension portale
 - Splénomégalie
 - Ascite
 - Circulation veineuse portosystémique
 - Diminution de la vitesse dans la veine porte
- TIPSS

cirrhose



Nodule de régénération



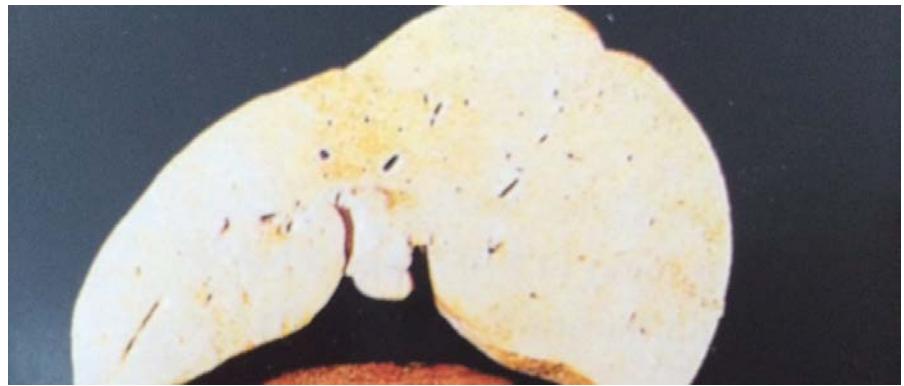
Nodule dysplasique

Carcinome hépatocellulaire

Il semble exister un continuum évolutif entre ces trois entités, ces transitions s'accompagnant d'une augmentation de la taille, de la cellularité et de la néoangiogenèse de ces différents nodules



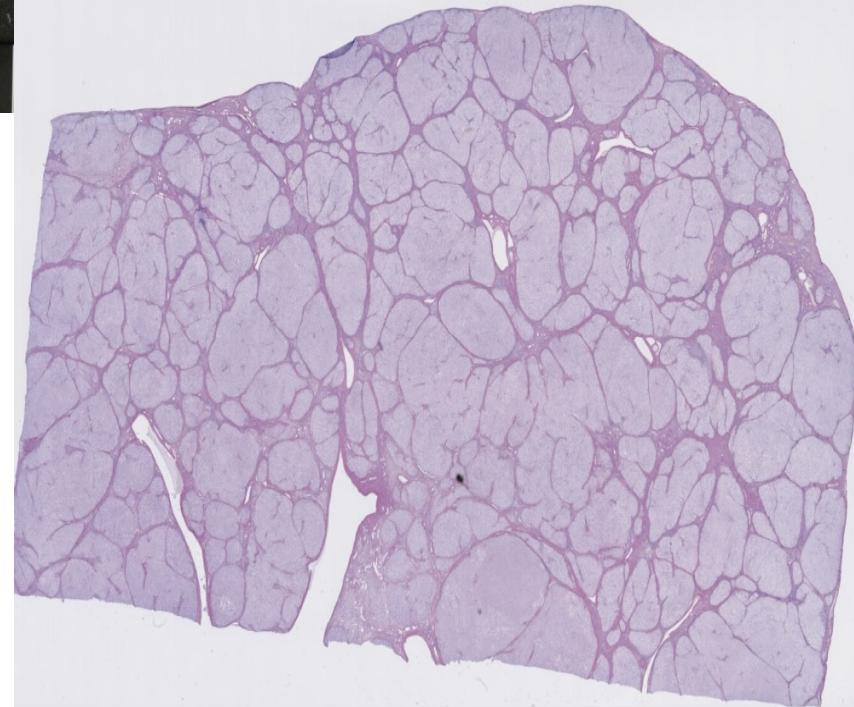
Foie normal



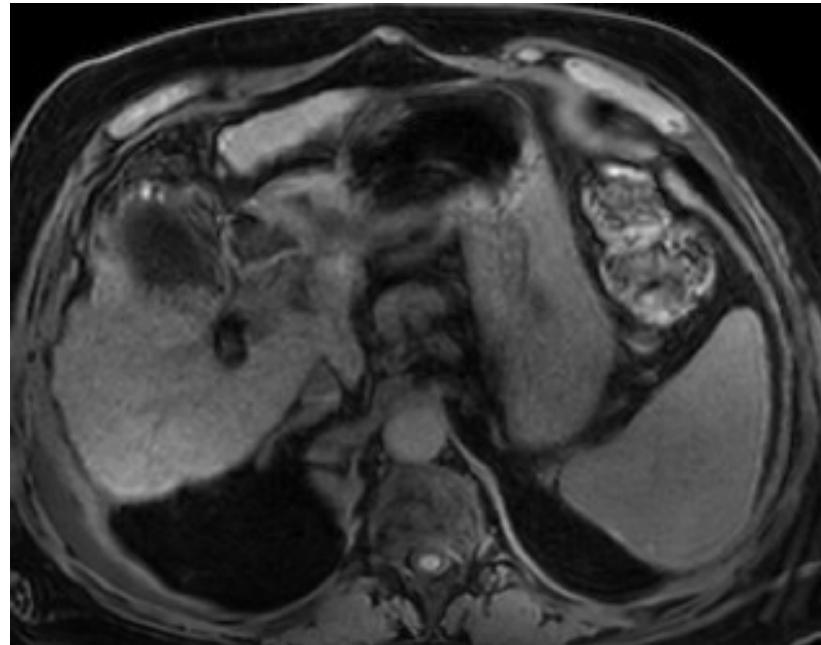
Foie stéatosique



Foie cirrhotique

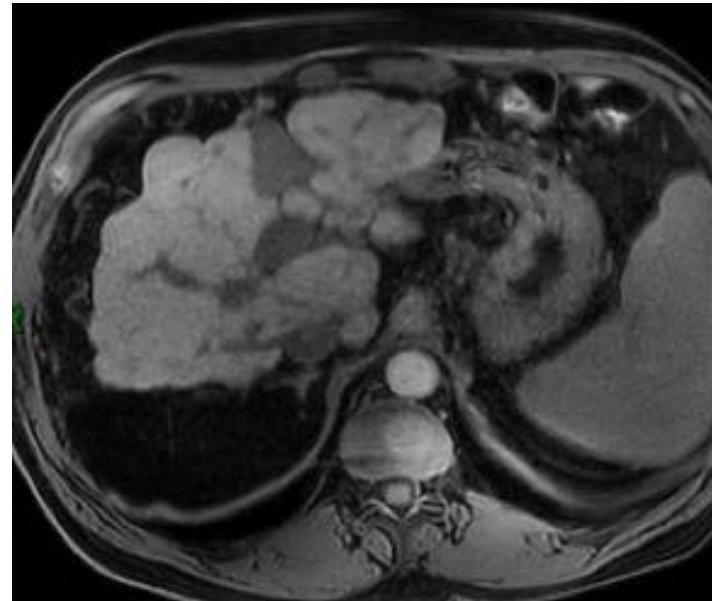
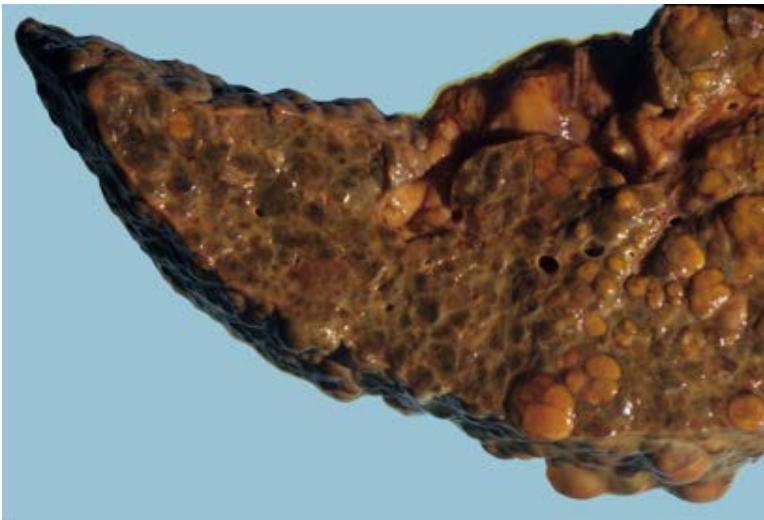


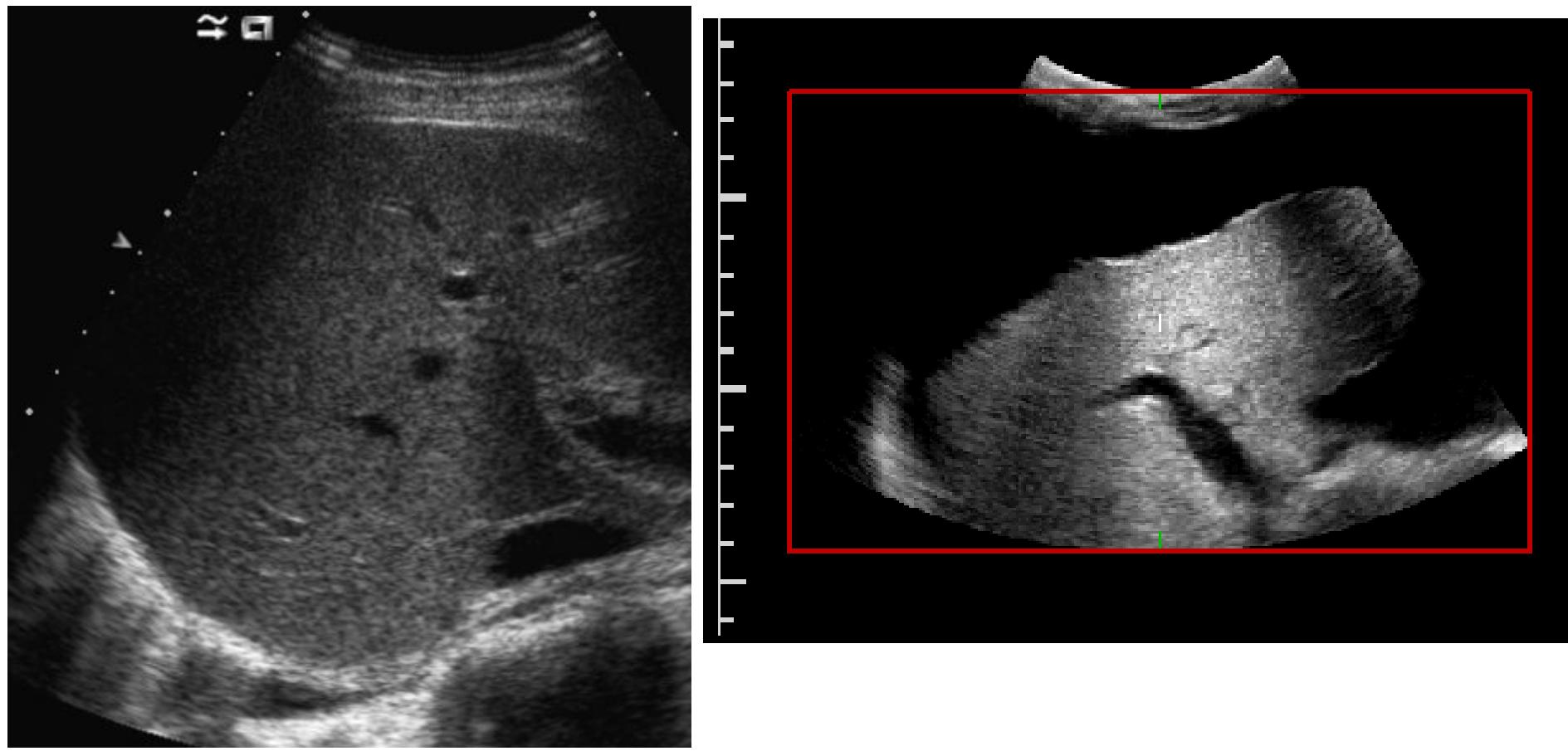
cirrhose





cirrhose





US et Cirrhose décompensée:

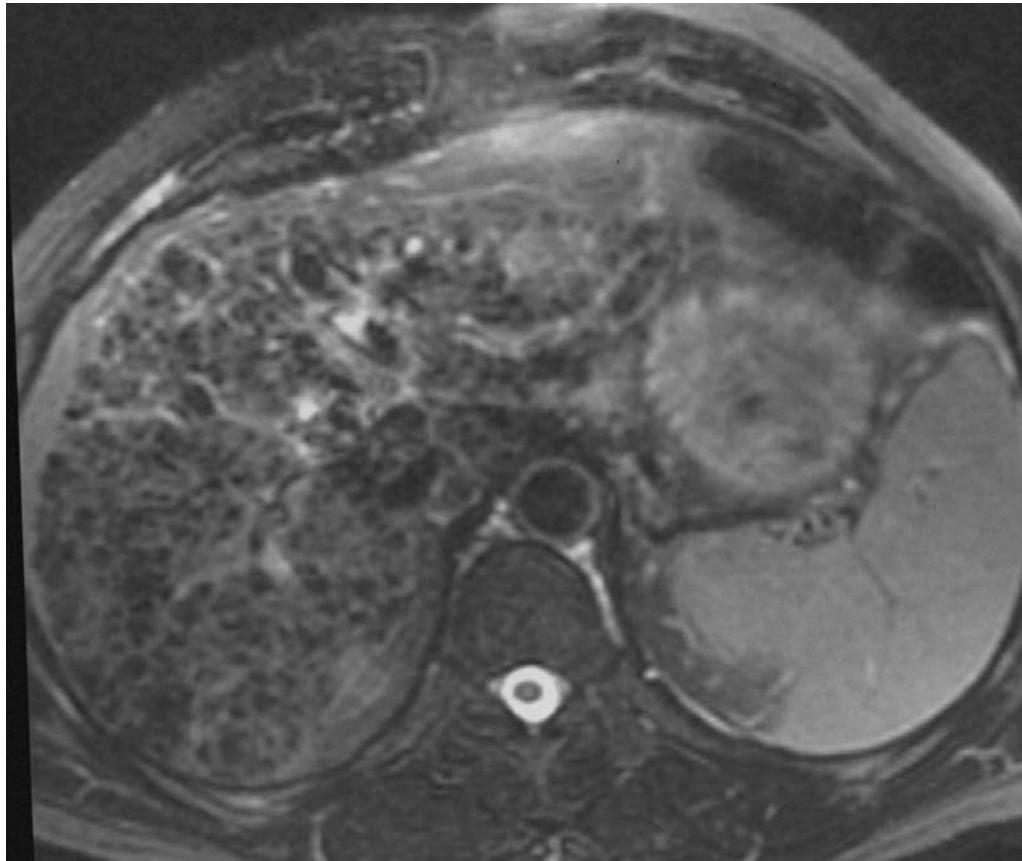
- Veine porte ? Ascite ?
- Hépatocarcinome ?

-43 ABD >



CT et cirrhose décompensée:

- Veine porte ?
- Hépatocarcinome ?
- Autre problème : intestin (ischémie, perforation)



IRM et cirrhose décompensée:

- Veine porte ?
- Hépatocarcinome ?

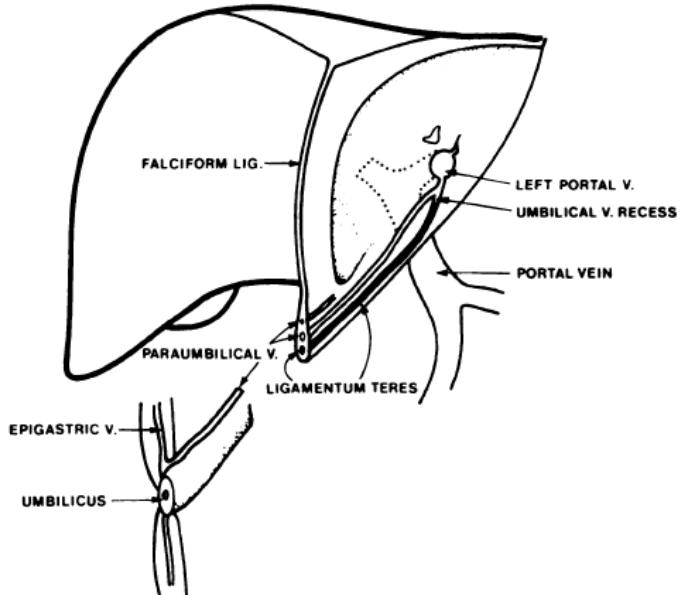


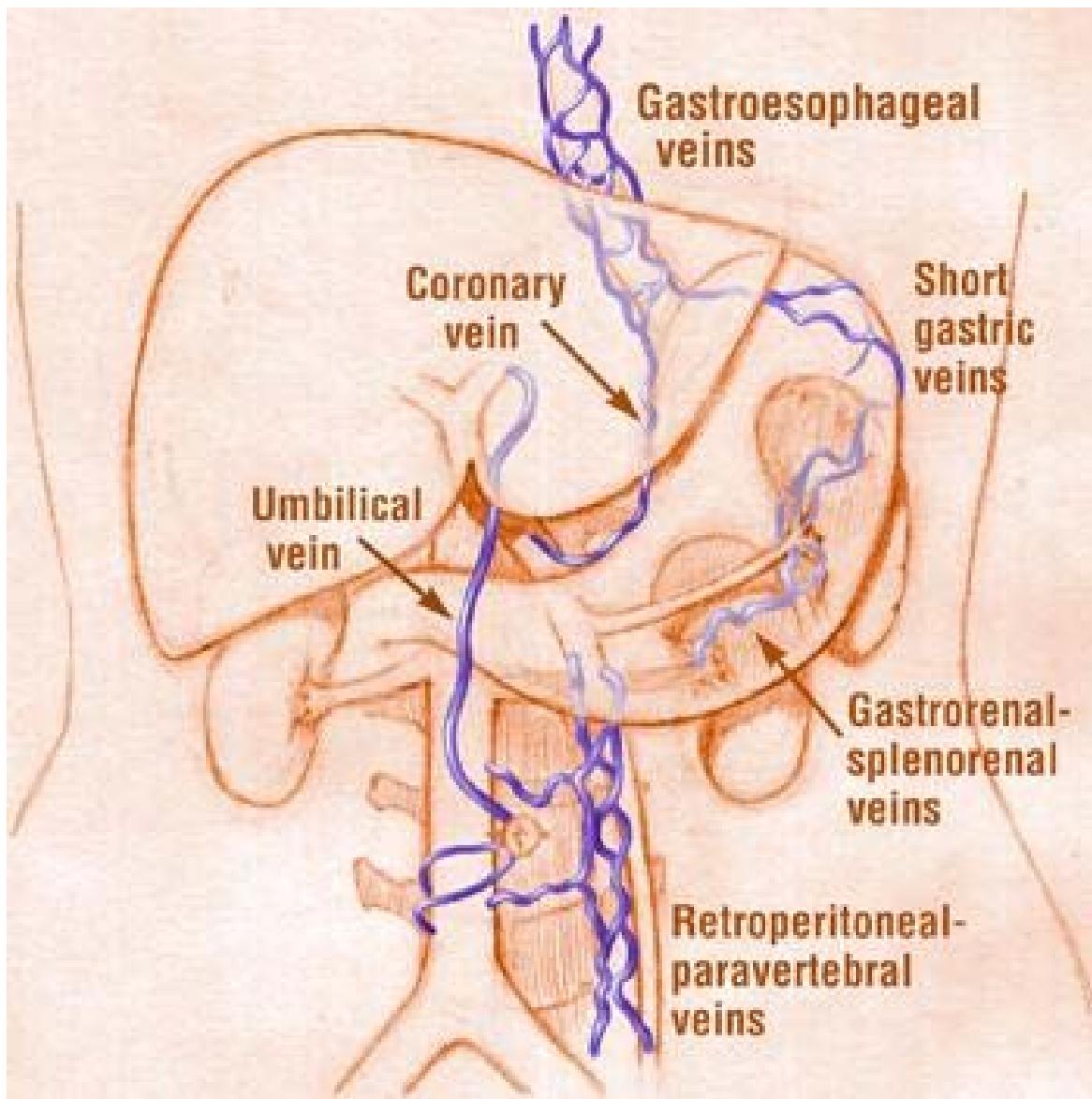
Fig. 1.—Anatomy of falciform ligament showing umbilical vein recess (Baumgarten recess) and its relation to paraumbilical vein.

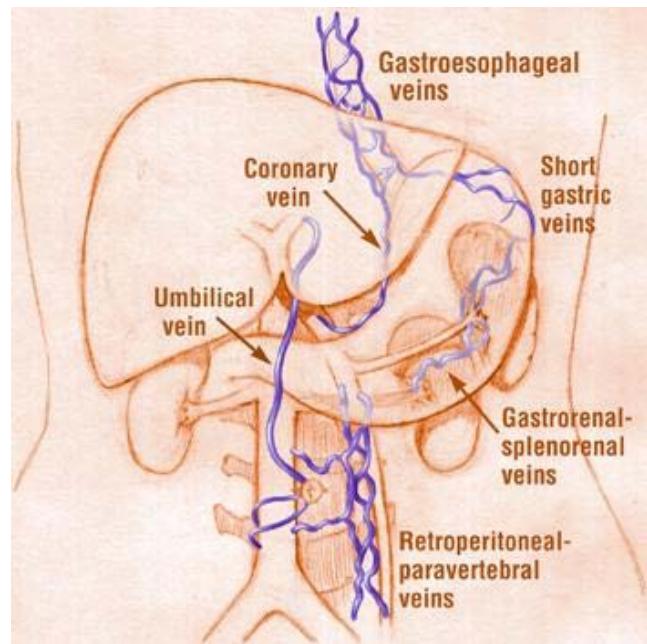
The Recanalized Umbilical Vein in Portal Hypertension: A Myth

Michel Lafortune¹
André Constantin¹
Guy Breton¹
André G. Légaré¹
Pierre Lavoie²

The demonstration of a vessel in the falciform ligament, traditionally presumed to be a reopened umbilical vein, is an important sonographic sign of portal hypertension. This vessel was sought in 200 umbilicoporphographies (all portal hypertensive) and in 41 autopsy-dissected falciform ligaments (34 normal and seven cirrhotic). The normal falciform ligament contained one to three tiny collapsed paraumbilical veins. In cirrhotics, the number and caliber of paraumbilical veins increased. A reopened umbilical vein was never found. The authors conclude that the umbilical vein does not recanalize in portal hypertension. The vessel involved is actually an enlarged paraumbilical vein.

AJR 144:549-553, March 1985
0361-803X/85/1443-0549
© American Roentgen Ray Society





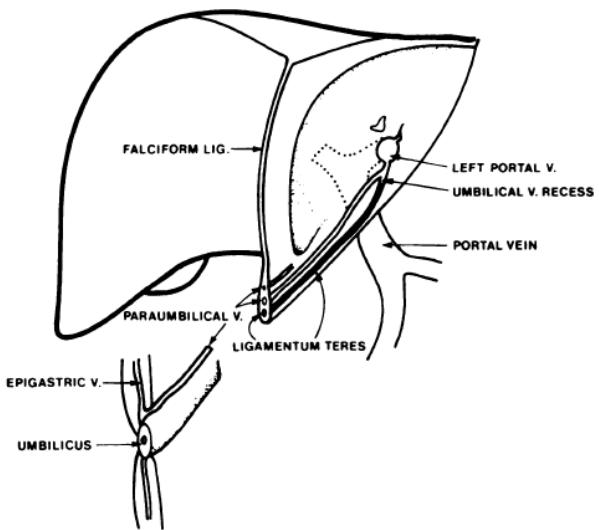
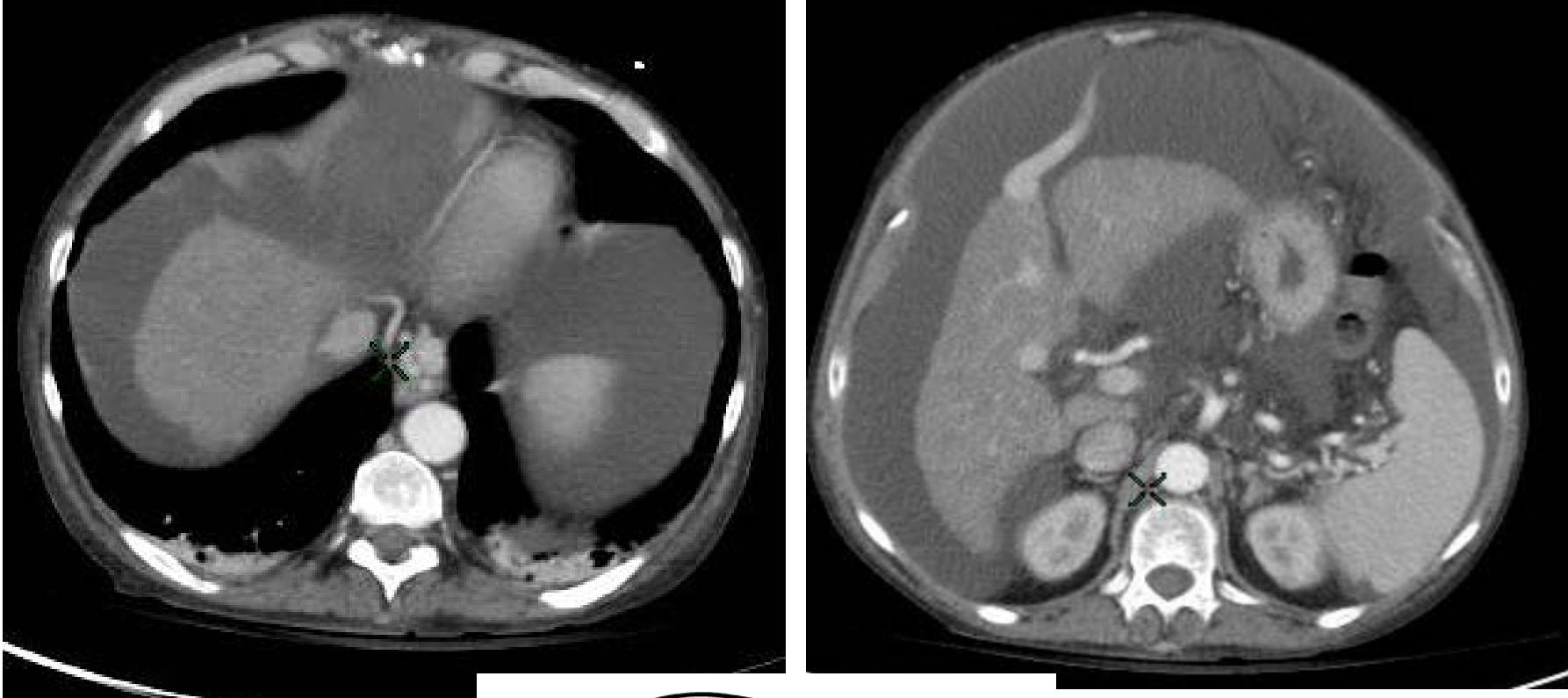
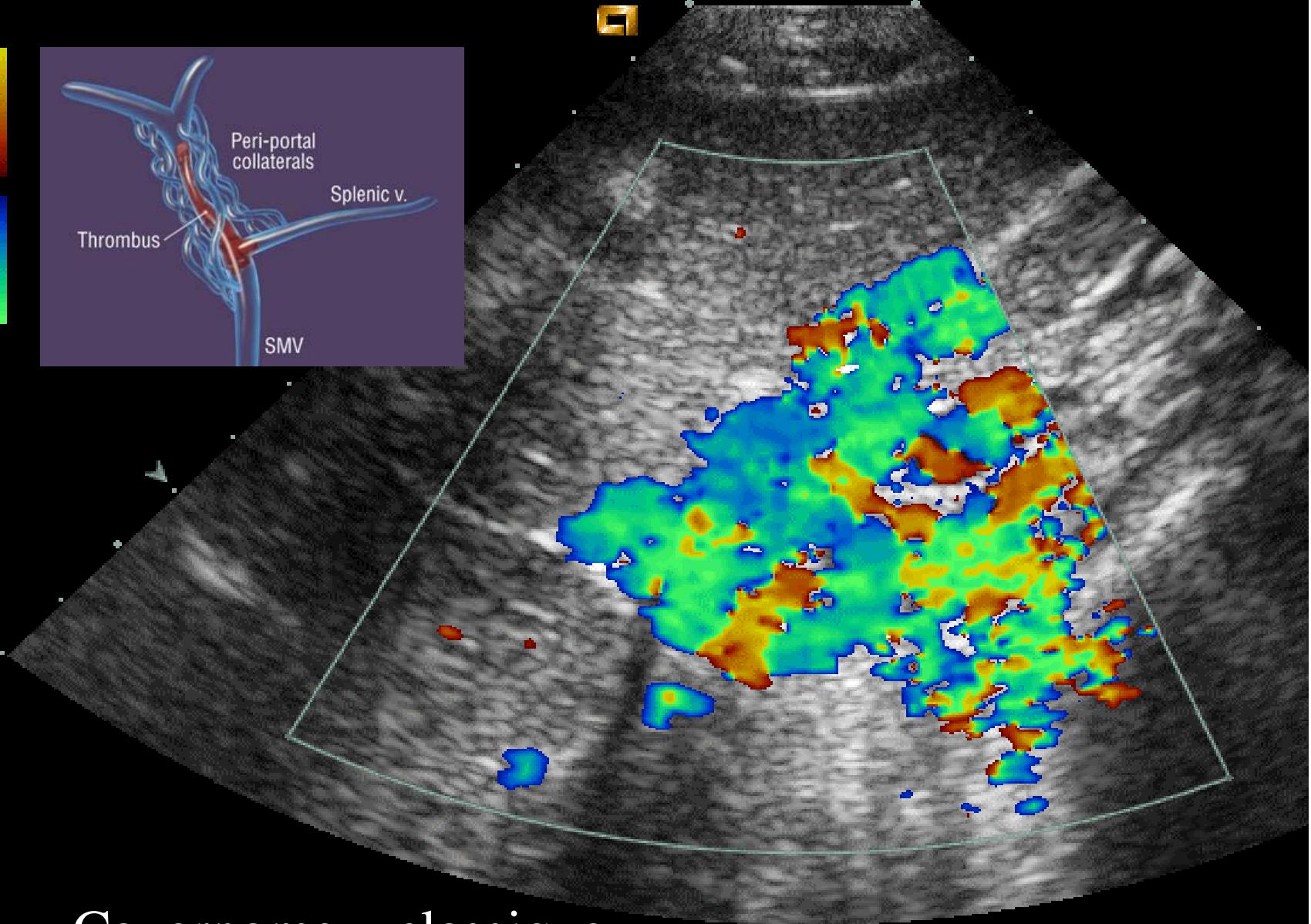
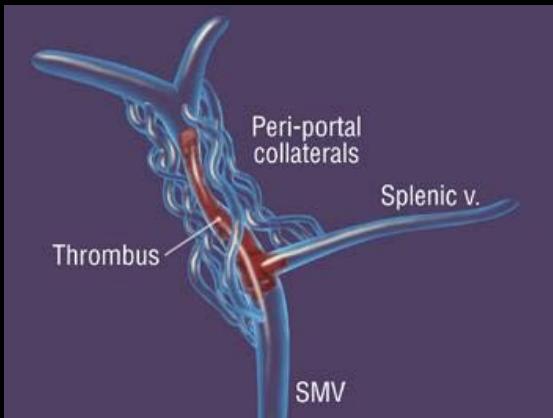
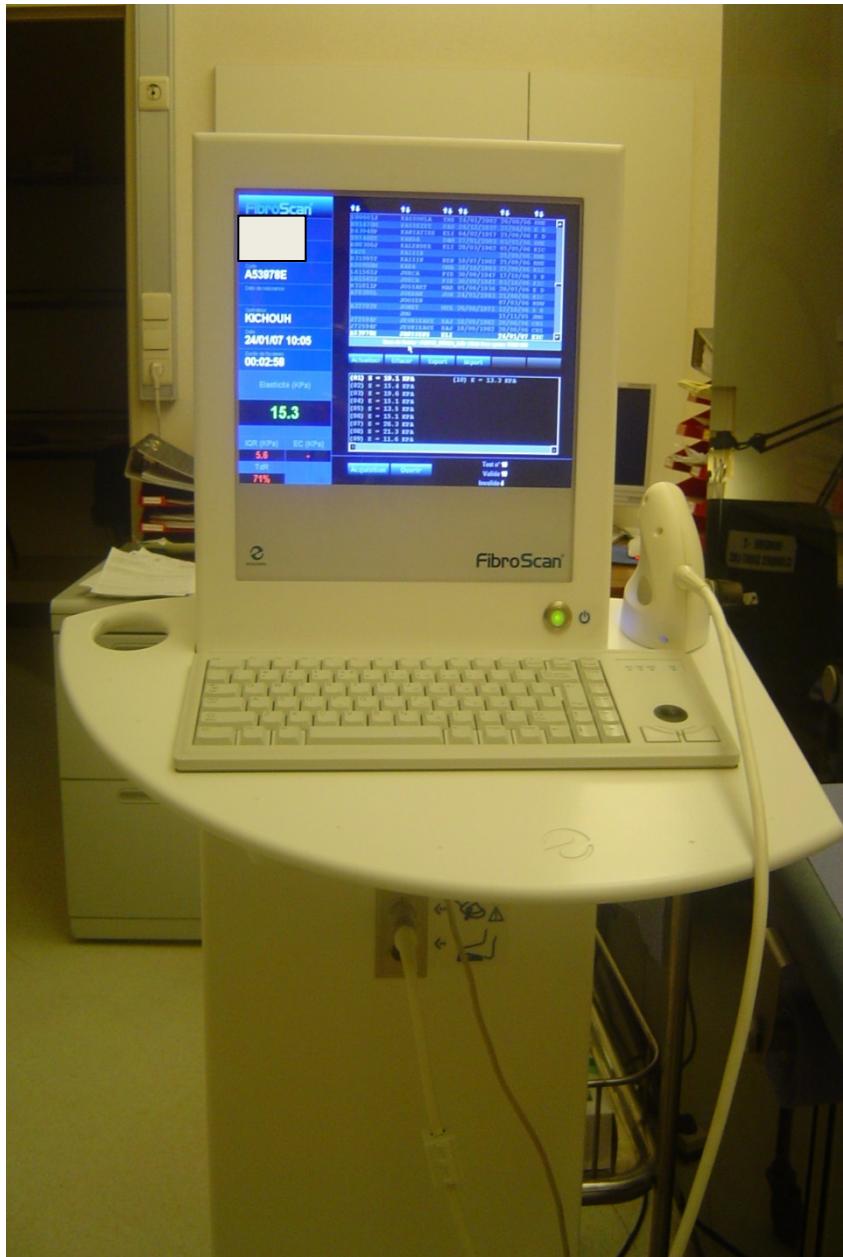


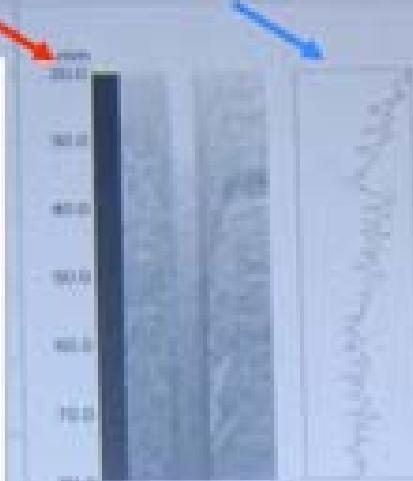
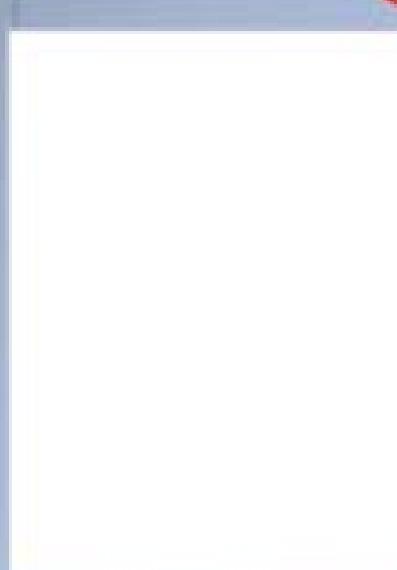
Fig. 1.—Anatomy of falciform ligament showing umbilical vein recess (Baumgarten recess) and its relation to paraumbilical vein.

.12



Cavernome « classique »





Elasticité (KPa)

11.6

11.1 0 = 11.6 KPa
11.1 0 = 11.6 KPa
11.0 0 = 11.6 KPa

IQR (KPa) EC (KPa)

3.1

13.5

TdR

100%

Tissu n° 1

Véhicule 10

Immunofluo

Le Fibroscan® est un dispositif médical d'aide au diagnostic. L'examen doit être réalisé par un opérateur certifié.
Les résultats doivent être interprétés par un médecin radiologue ou gastroentérologue (GECI) et/ou un clinicien.

TIS0.9 MI 1.3

C5-1
65Hz
RP

2D
61%
Dyn R 55
P Med
Gen

ElastQ
Liver EQI 9

EQI Avg 12.2 kPa
EQI Med 12.0 kPa
EQI IQR/Med 16 %

Liver EQI 10

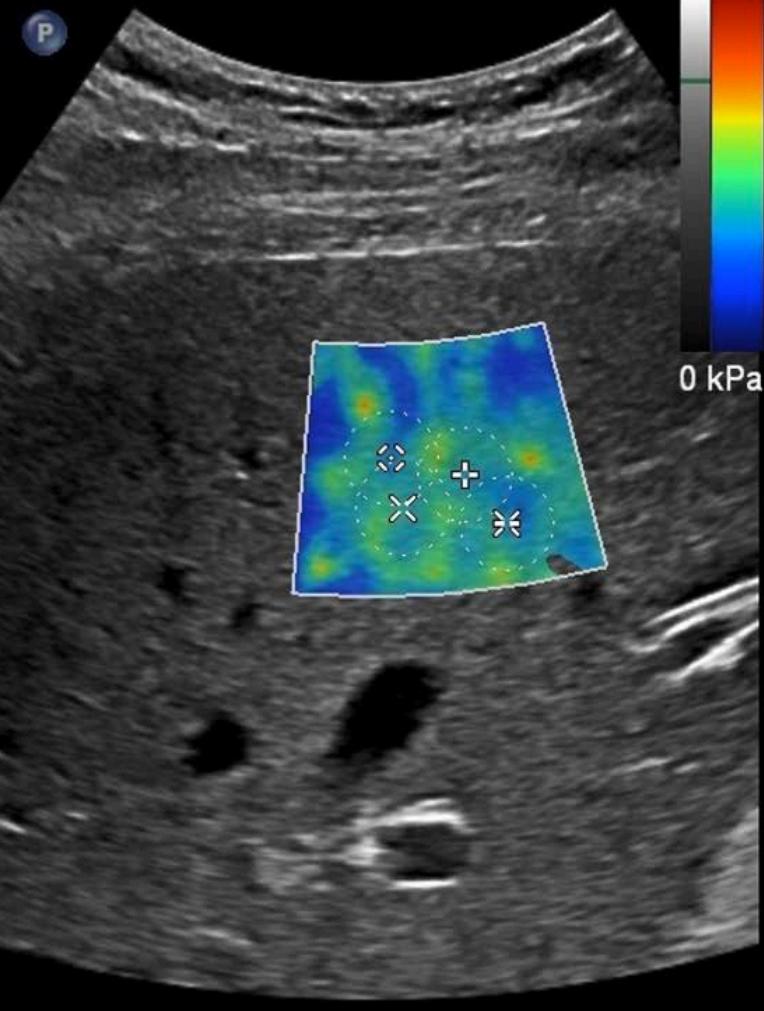
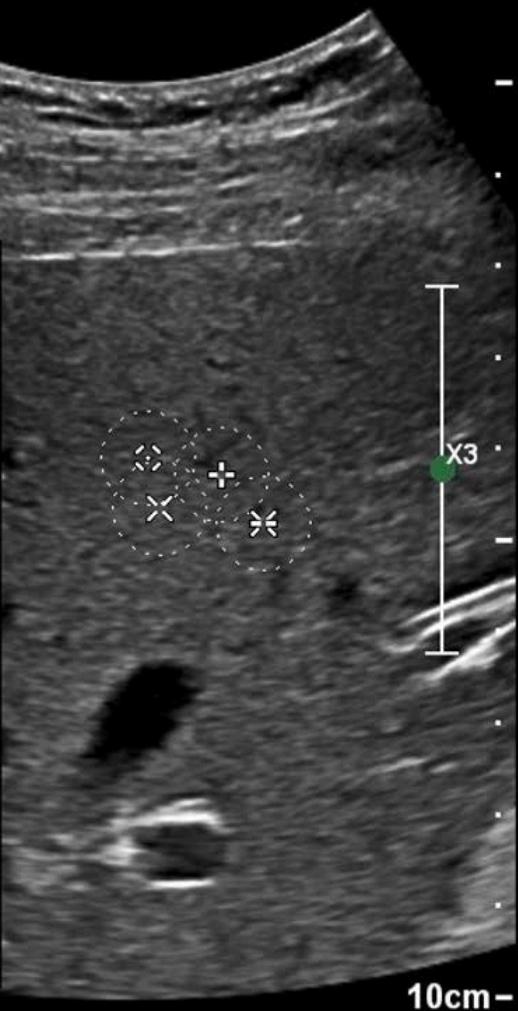
EQI Avg 12.2 kPa
EQI Med 12.0 kPa
EQI IQR/Med 12 %

Liver EQI 1

EQI Avg 11.6 kPa
EQI Med 11.6 kPa
EQI IQR/Med 17 %

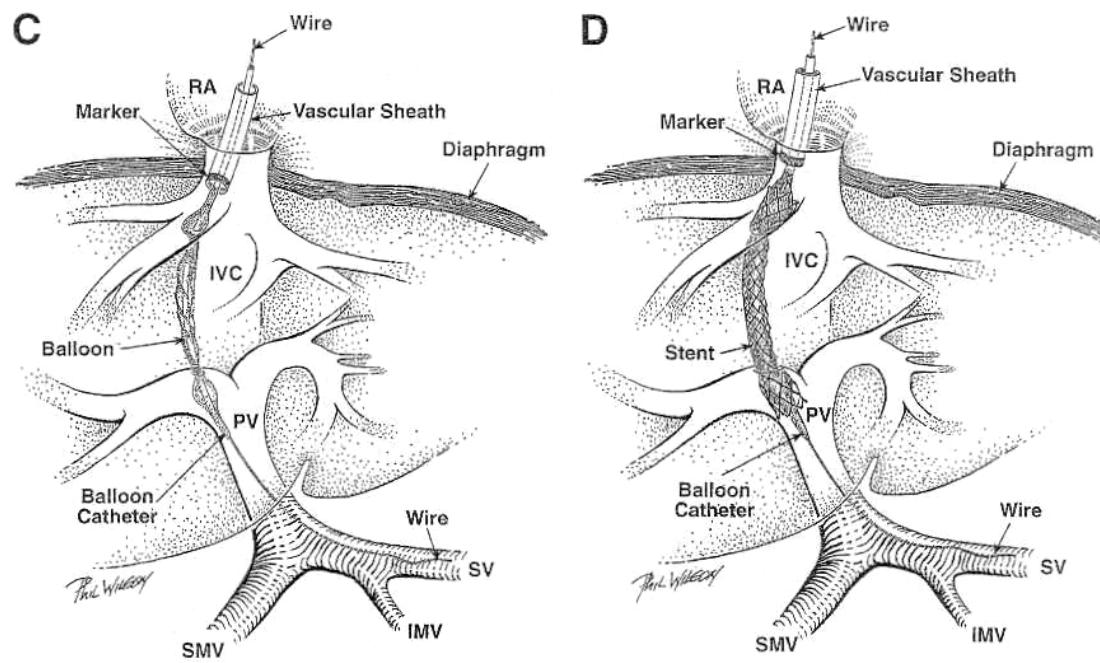
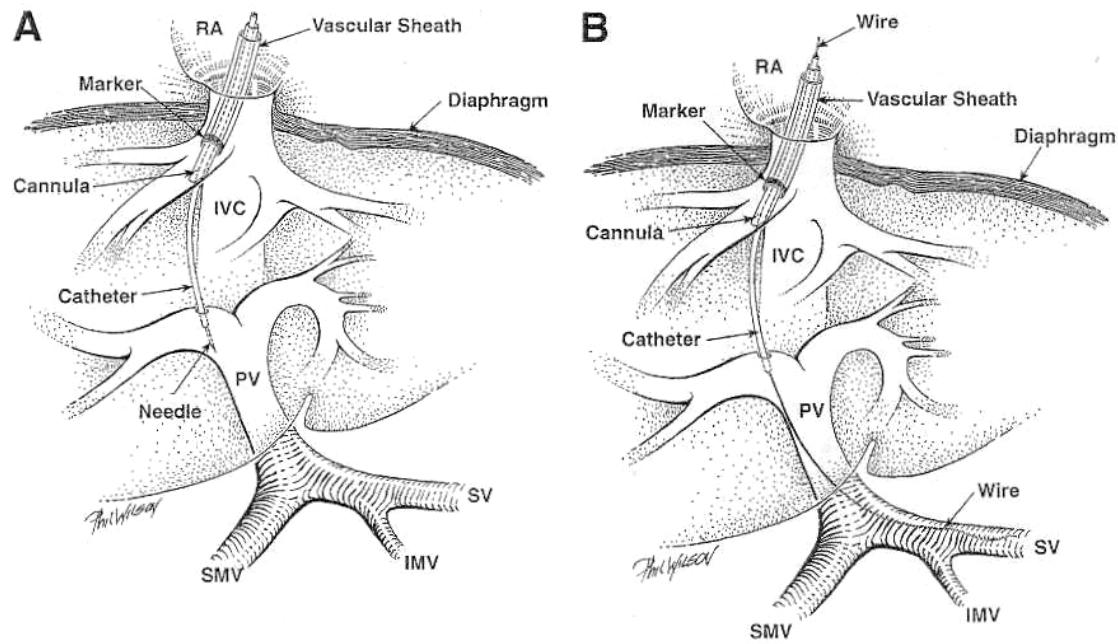
Liver EQI 2

EQI Avg 11.0 kPa
EQI Med 11.1 kPa
EQI IQR/Med 17 %



Shunts porto-caves intra-hépatiques par voie transjugulaire (TIPS)

- Indications:
 - Ascites médico-résistantes
 - Hémorragies par rupture de VO récidivantes
 - Budd-Chiari





51:43



52:50

