

Case 5. The Year in Review: An Irresistible Case







Howard J. Sachs, MD  
[www.12DaysinMarch.com](http://www.12DaysinMarch.com)  
Howard@12daysinmarch.com

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	Ref Range & Units	
 <b>WBC</b>	4.3 - 10.8 10 <sup>3</sup> /uL	<b>4.7</b>
 <b>Hemoglobin</b>	13.2 - 17.1 g/dL	<b>13.7</b>
 <b>Hematocrit</b>	39.0 - 52.0 %	<b>39.1</b>
 <b>MCV</b>	80.0 - 100.0 fL	<b>105.3 (H)</b>



Which of the following studies will NOT offer clarity in interpreting this abnormal lab value?

- A. Methylmalonic acid level
- B. Homocysteine level
- C.  $\gamma$ -glutamyl transferase level
- D. Warthin Starry silver stain of antral crypts

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 <b>Platelets</b>	140 - 440 $10^3/\mu\text{L}$	<b>90 (L)</b>




Which of the following would be least likely to explain this abnormal lab value?

- A. Myelodysplastic syndrome
- B. IgG antibody directed against platelet glycoprotein
- C. Splenic sequestration
- D. Rickettsial infection

Ref Range & Units		
 <b>AST</b>	10 - 40 U/L	<b>169 (H)</b>
 <b>ALT</b>	10 - 40 U/L	<b>48 (H)</b>

If liver biopsy on this patient reveals microvesicular steatohepatitis, which of the following would be the least likely cause?

- A. Autoimmune cholangitis
- B. Acetaminophen toxicity
- C. Chronic alcohol ingestion
- D. Aspirin use in 8 y.o. boy with febrile illness
- E. Hypertriglyceridemia

	Ref Range & Units	
 <b>Total Protein</b>	6.0 - 8.0 g/dL	<b>7.0</b>
 <b>Albumin</b>	3.5 - 4.8 g/dL	<b>4.0</b>
 <b>Alkaline Phosphatase</b>	30 - 115 U/L	<b>122 (H)</b>

Which of the following tests would be most useful in determining the etiology of this test abnormality?

- A. Colonoscopy revealing mucosal ulceration
- B. Antibody directed against mitochondrial autoantigens
- C.  $\gamma$ - glutamyl transferase
- D. Endoscopic ultrasound of biliary tree





Patient presented for routine evaluation. He drinks 8 beers per day. He feels well. Never misses work. He is on no medication

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WBC	4.3 - 10.8 $10^3/\mu\text{L}$	4.7
RBC	4.20 - 5.80 $10^6/\mu\text{L}$	3.71 (L)
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MCH	27.0 - 34.0 pg	36.9 (H)
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Total Protein	6.0 - 8.0 g/dL	7.0
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Bilirubin, Total	0.3 - 1.2 mg/dL	1.8 (H)
Bilirubin, Direct	$\leq 0.4$ mg/dL	0.8 (H)
Alkaline Phosphatase	30 - 115 U/L	122 (H)
AST	10 - 40 U/L	169 (H)
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Which of the following most likely explains the basis of his hyperbilirubinemia?

- A. Microvesicular steatohepatitis
- B. Lymphocyte-mediated attack of intralobular bile ducts
- C. Fibrosis and stricture formation of intra- and extra-hepatic bile ducts
- D. Hepatocellular enzyme insufficiency
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H pylori causes Type B Gastritis, not macrocytosis

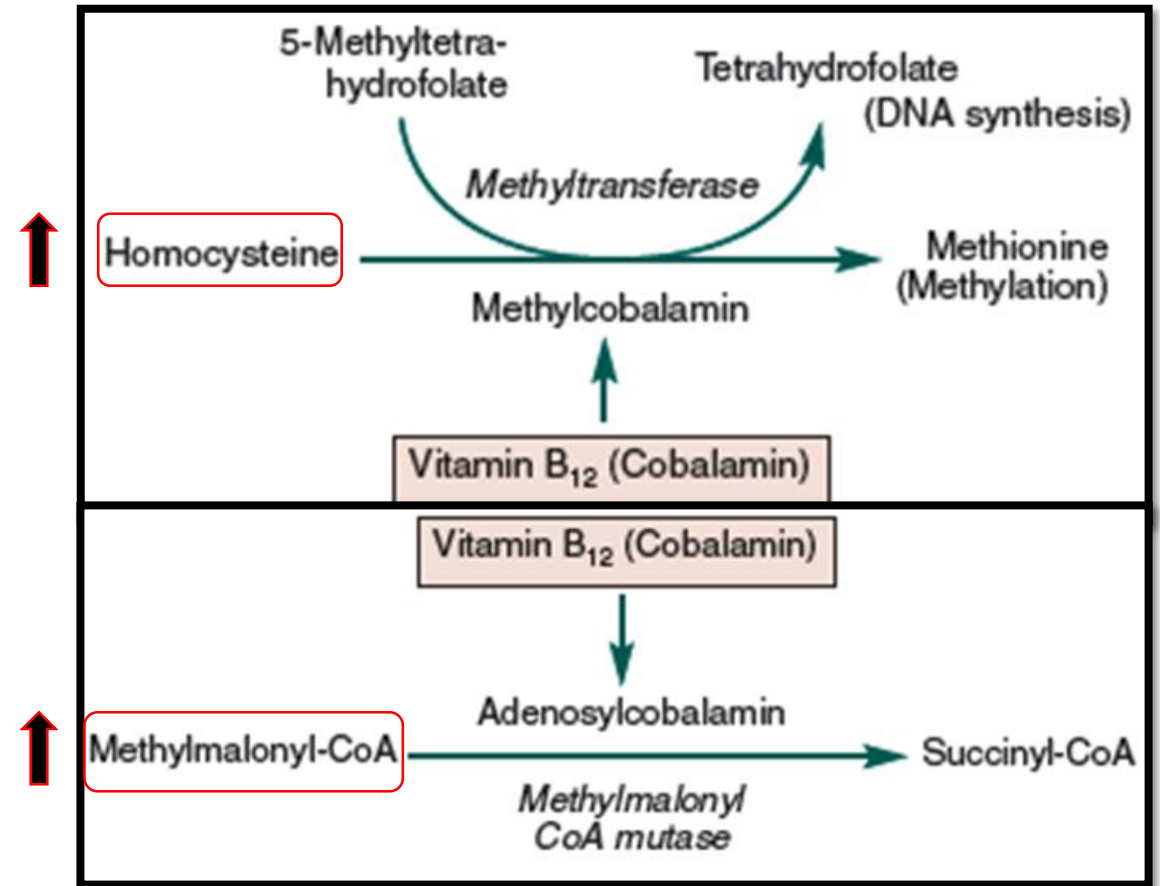




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These tests would identify causes of macrocytosis:

- A. *Methylmalonic acid level*
- B. *Homocysteine level*
- C.  $\gamma$ -glutamyl transferase level
- D. Warthin Starry silver stain of antral crypts



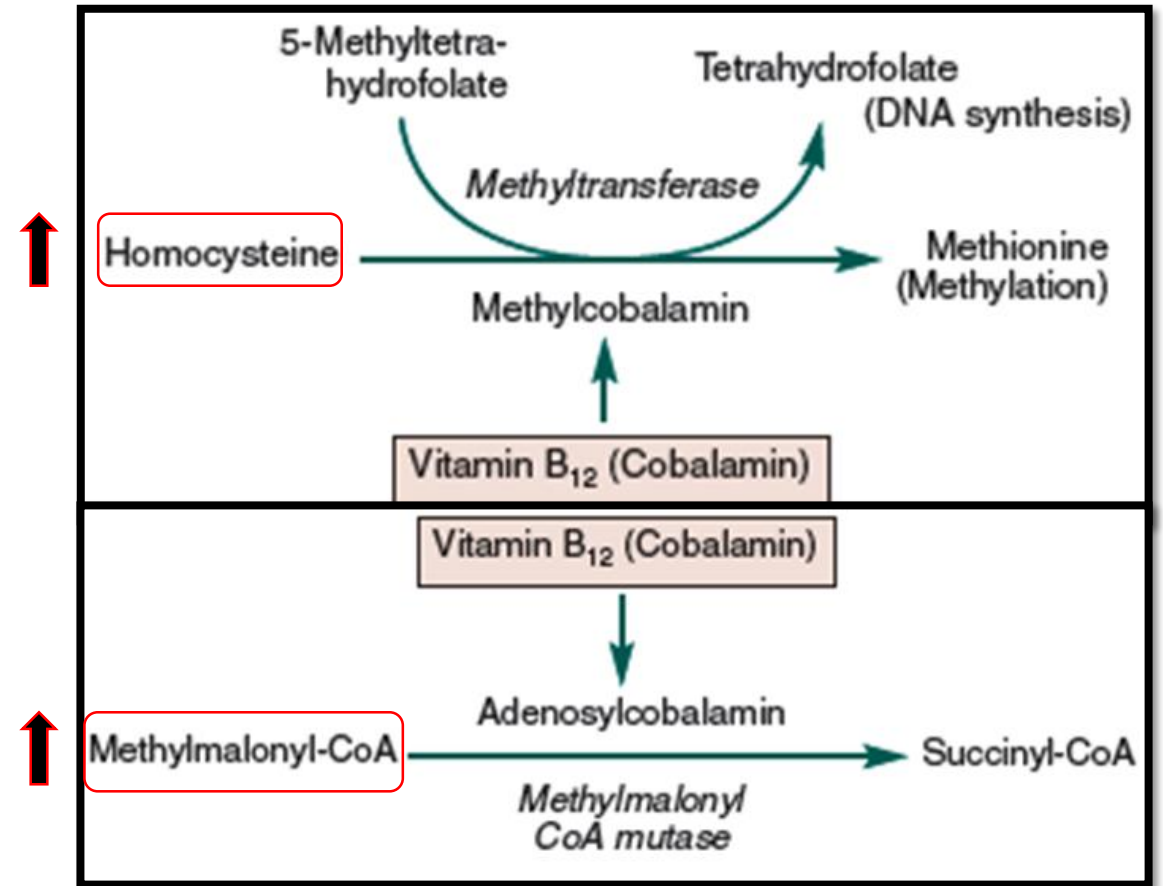
*MMA elevated in B-12 deficiency only*  
*Homocysteine elevated in B-12 AND folate deficiency*

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These tests would identify causes of macrocytosis:

- A. Methylmalonic acid level
- B. Homocysteine level
- C.  $\gamma$ -glutamyl transferase level – **induced by alcohol**
- D. Warthin Starry silver stain of antral crypts

Chronic alcoholic liver disease leads to macrocytosis  
*Mechanism: 'membrane lipid abnormalities'*



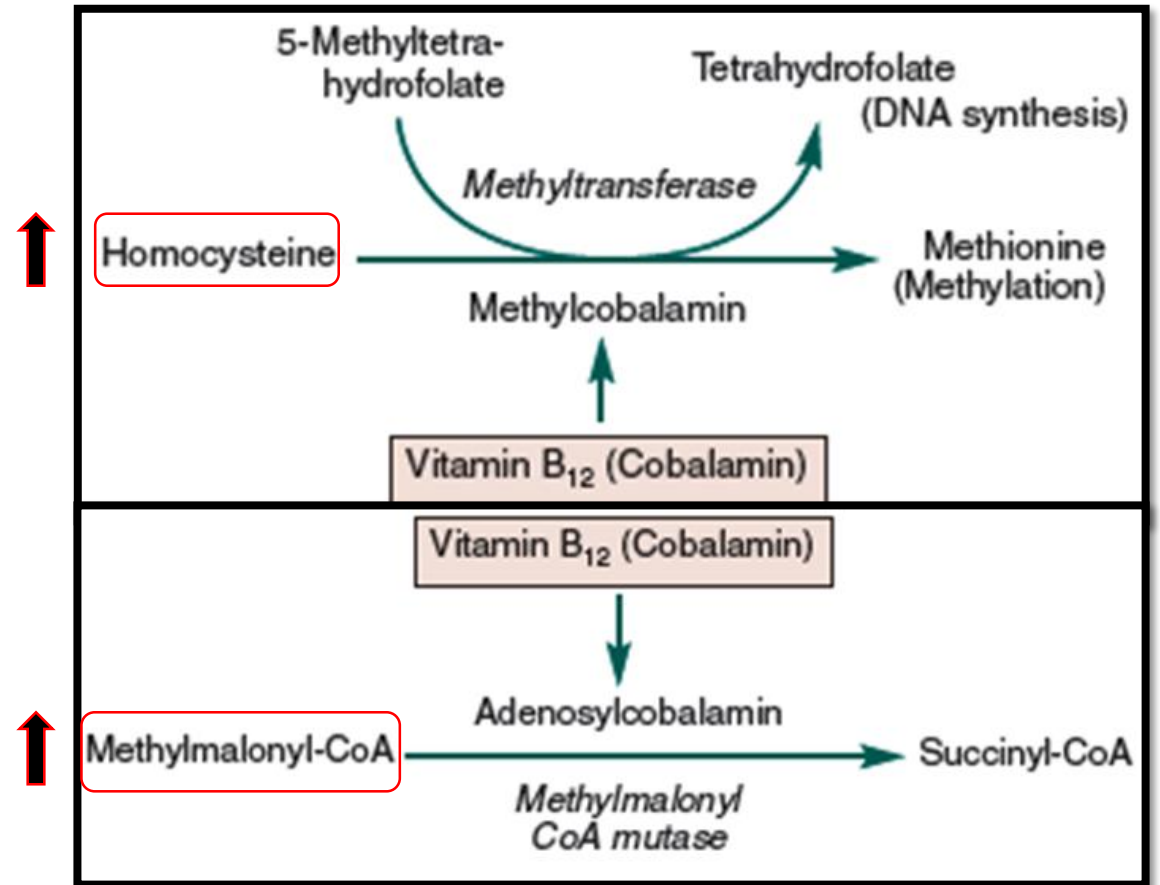
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Possible mechanism: defective cholesterol esterification



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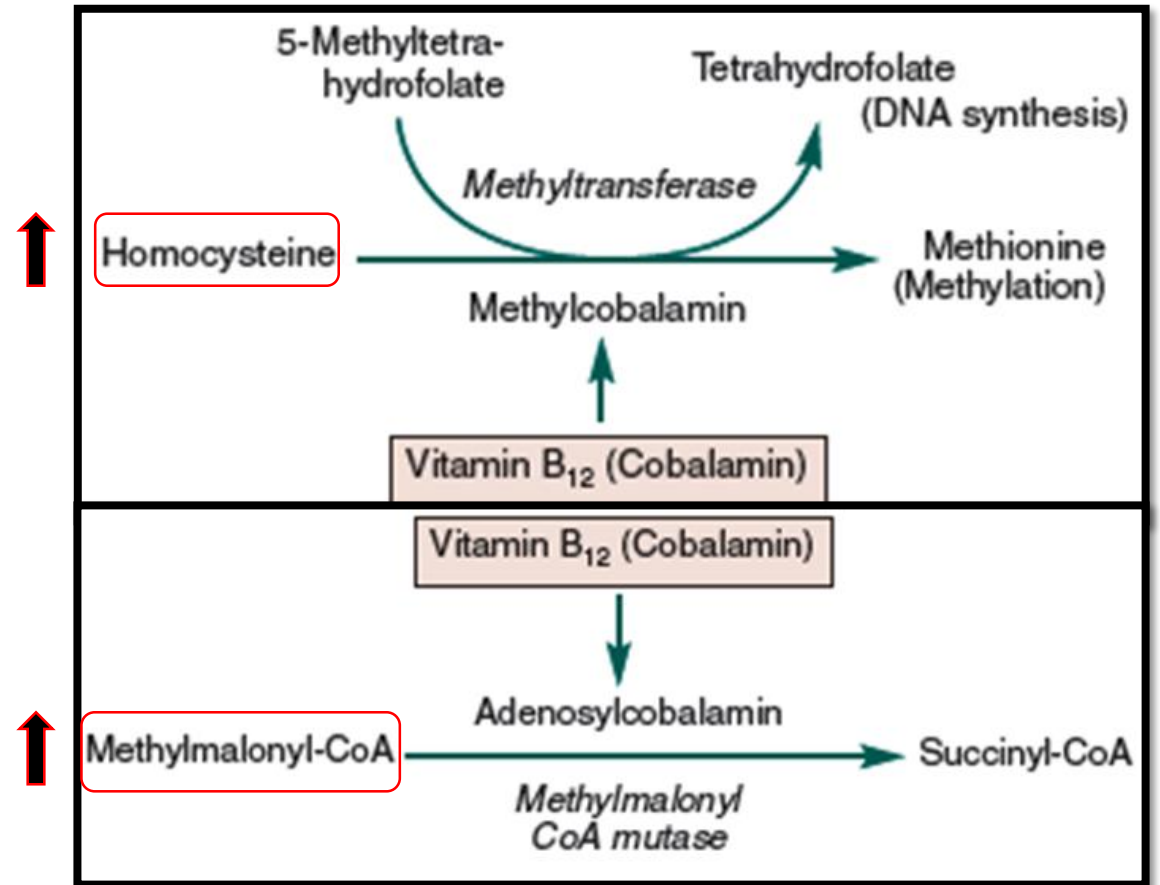
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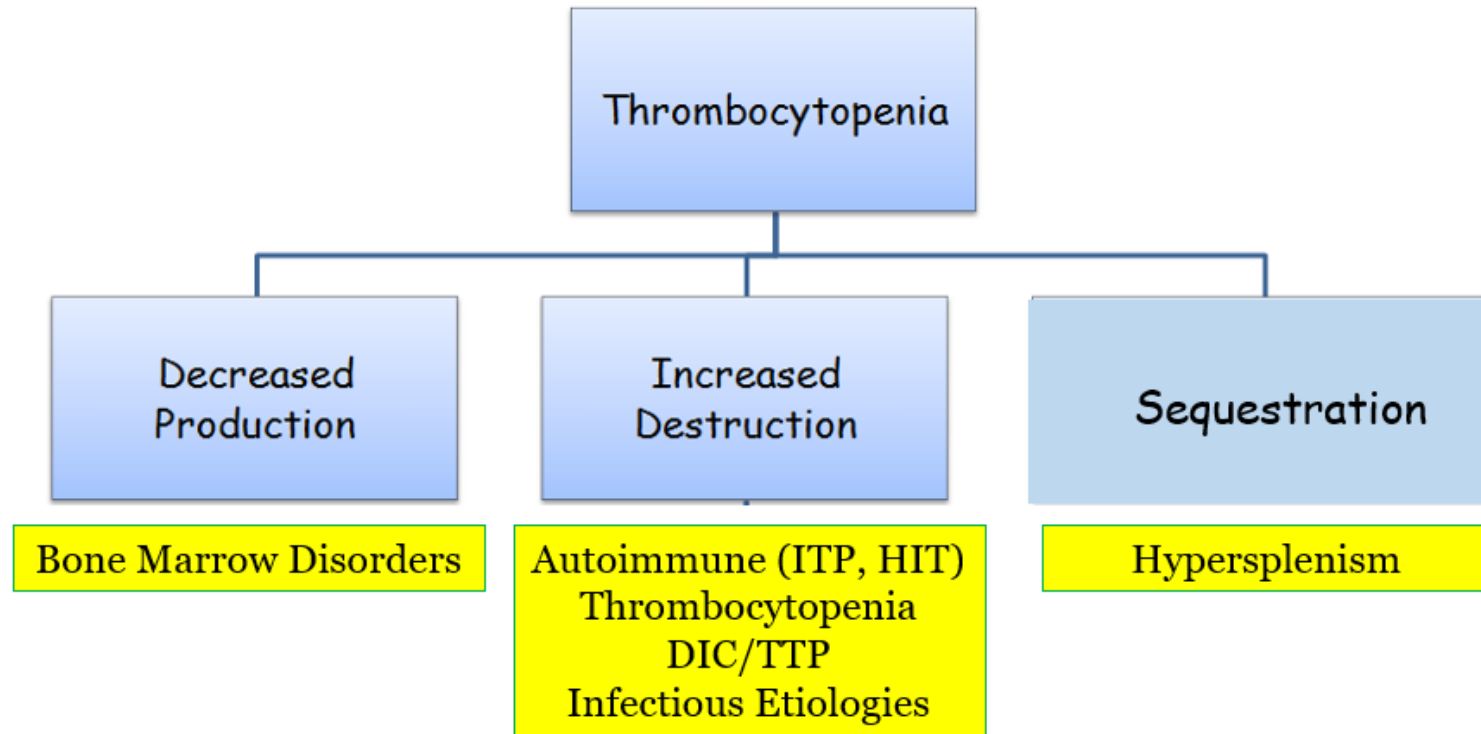
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*'Fatty Liver, Fatty RBC Membranes'*

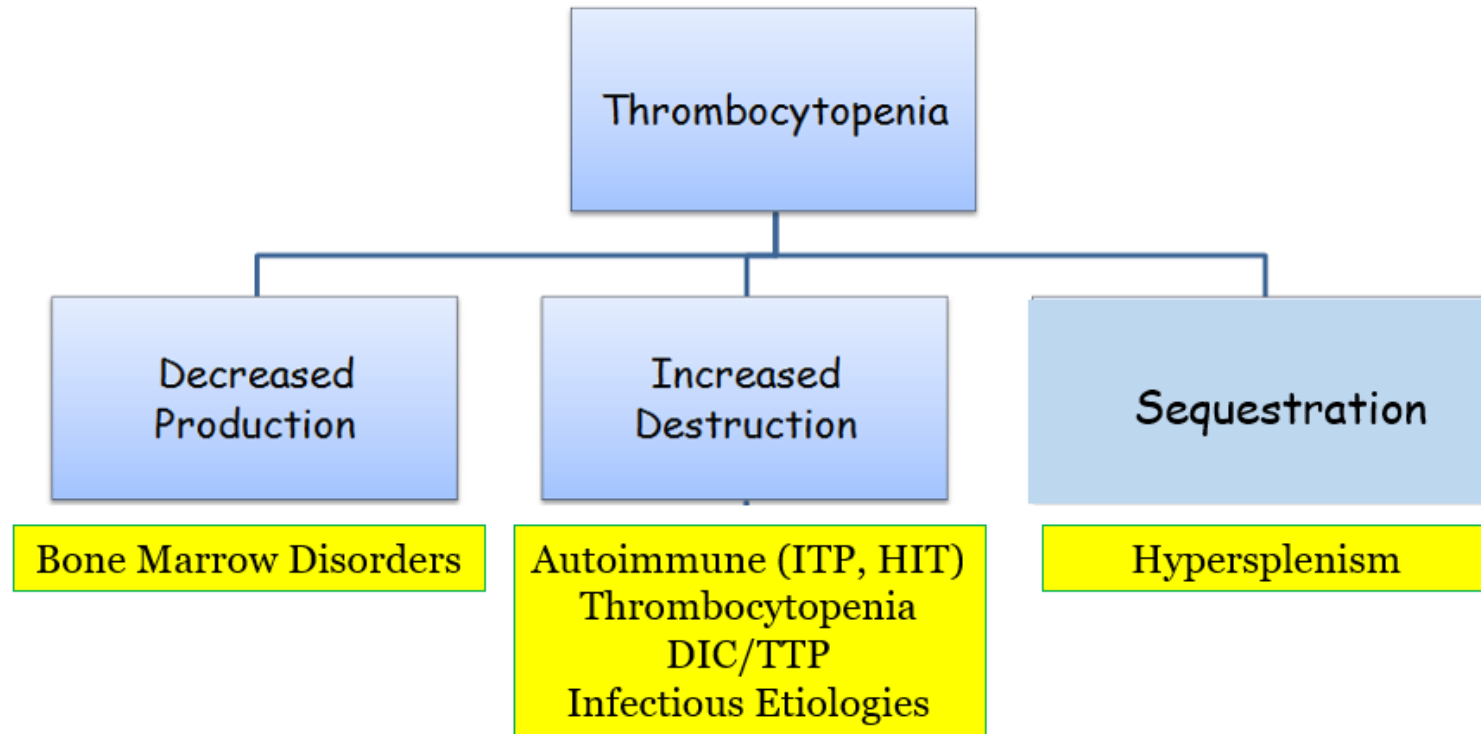




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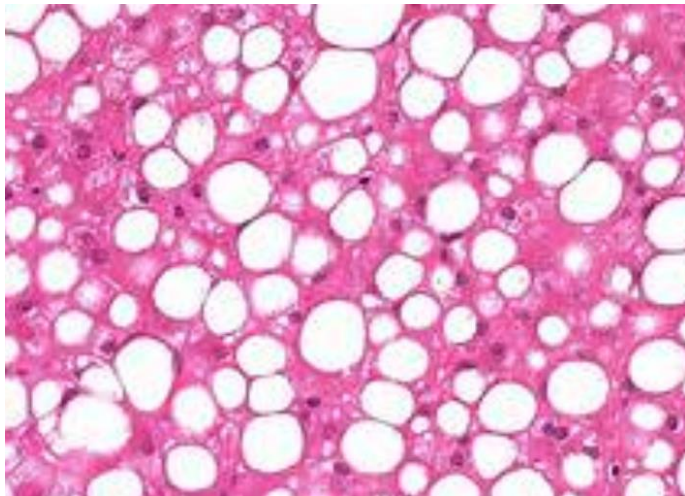


Other disorders associated with thrombocytopenia:




- B. IgG antibody directed against platelet glycoprotein: autoimmune thrombocytopenia
- C. Splenic sequestration (i.e. hypersplenism): portal HTN
- D. Rickettsial infection: RMSF (platelets are consumed, probably on an immunologic basis?)

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Step One (elevated alk  $\Phi$ ): Is it Bone or Liver?



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OK kids, how do you put this case together?



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Macrocytosis in a heavy drinker = chronic liver disease

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Alkaline phosphatase = may be elevated in alcoholic liver disease (but not on the USMLE Step One)

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*Gilbert's Syndrome* (unconjugated hyperbilirubinemia)  
 $0.8/1.8 = 0.4$

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All the other causes would be associated with conjugated hyperbilirubinemia (direct/total ratio >50%)

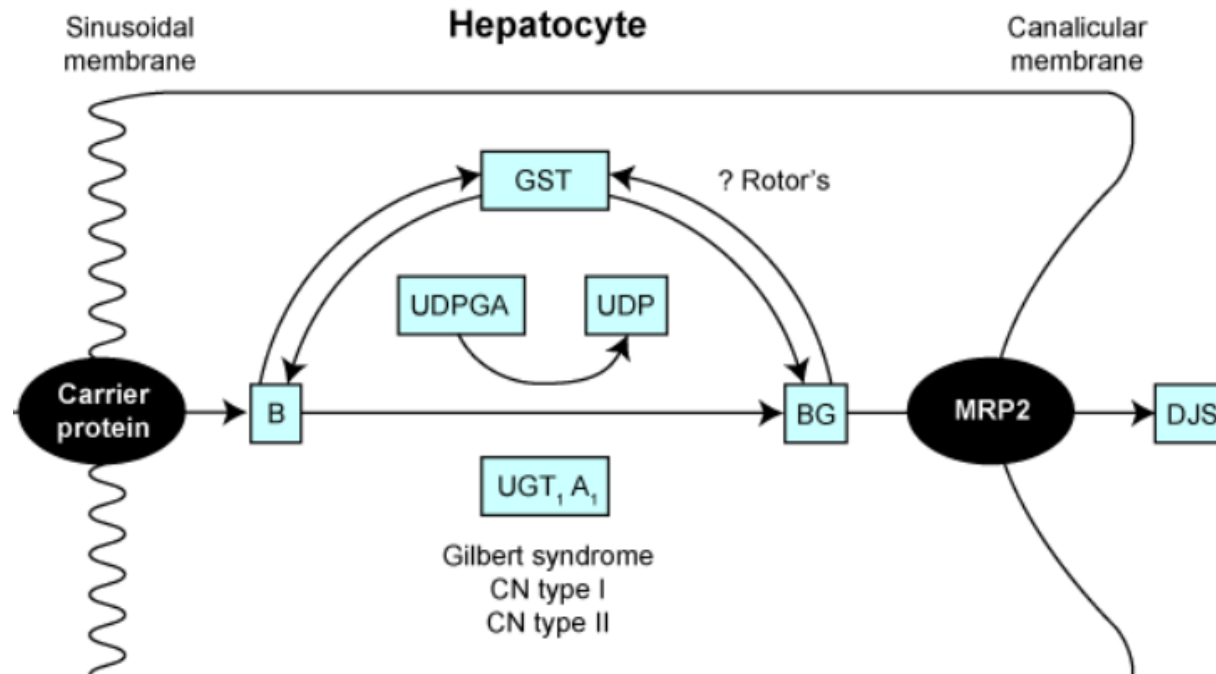
- A. Microvesicular steatohepatitis: Alcoholic liver disease (usually described with Mallory bodies)
- B. Lymphocyte-mediated attack of intralobular bile ducts: Primary Biliary Cholangitis
- C. Fibrosis and stricture formation of intra- and extra-hepatic bile ducts: Sclerosing Cholangitis
- D. **Hepatocellular enzyme insufficiency**
- E. Ground-glass hepatocytes with fine, cytoplasmic granular inclusions: Hepatitis B (with surface antigen inclusions)



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**Gilbert's Syndrome** (unconjugated hyperbilirubinemia): *insufficiency* of UGT involved in conjugation



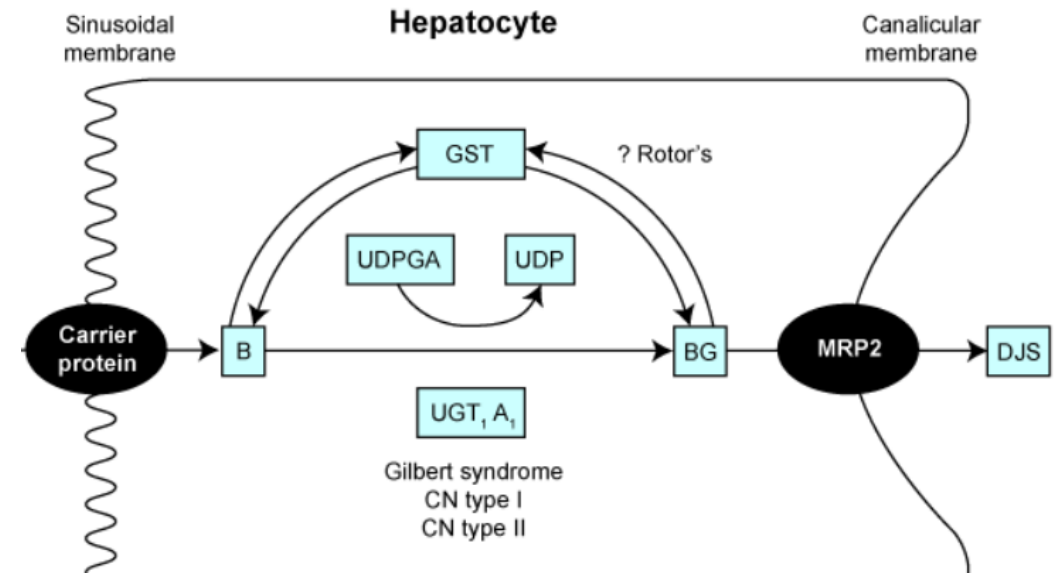
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 $0.8/1.8 = 0.4$

	Ref Range & Units	Value
Total Protein	6.0 - 8.0 g/dL	7.0
Albumin	3.5 - 4.8 g/dL	4.5
Bilirubin, Total	0.3 - 1.2 mg/dL	1.7 (H)
Bilirubin, Direct	<=0.4 mg/dL	0.2
Alkaline Phosphatase	30 - 115 U/L	60

Repeat fasting value following 6 weeks of alcohol cessation



Case 5. The Year in Review: An Irresistible Case



Howard J. Sachs, MD  
[www.12DaysinMarch.com](http://www.12DaysinMarch.com)  
Howard@12daysinmarch.com