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Amoxicillin-Clavulanate

Updated: March 30, 2016.

OVERVIEW

Introduction

The combination of amoxicillin and clavulanate is an oral antibiotic widely used in the treatment of mild-tomoderate bacterial infections including sinusitis, bronchitis, otitis media, cellulitis and community acquired pneumonia. Amoxicillin-clavulanate is currently the most common cause of clinically apparent, drug induced acute liver injury both in the United States and Europe.

Background

The combination of amoxicillin and clavulanate is a commonly used antibiotic which is active against many bacterial organisms that cause sinusitis, bronchitis, otitis media, skin and tissue infections and community acquired pneumonia. The combination consists of amoxicillin which is a semisynthetic, third generation penicillin and clavulanate which is a beta lactam that acts as an inhibitor of beta lactamase, the major bacterial enzyme responsible for penicillin resistance. Amoxicillin-clavulanate was approved for use in the United States in 1984 and, currently, 2 to 5 million prescriptions are filled yearly, making it one of the most common antibiotic regimens used. Current indications are for mild-to-moderate bacterial infections due to known or suspected penicillinase resistant gram positive or gram negative organisms. This combination is provided in multiple dose combinations, typically as 250 to 875 mg amoxicillin with 125 mg of clavulanate, given two to three times daily for 7 to 10 days. Amoxicillin-clavulanate is available in multiple generic formulations and under the brand name Augmentin. Common side effects include gastrointestinal upset, diarrhea and rash.

Hepatotoxicity

Amoxicillin-clavulanate has been implicated in hundreds of cases of clinically apparent acute liver injury and this combination is currently the most common cause of drug induced liver disease in most large case series from the United States and Europe. The onset of injury is typically a few days to as long as 8 weeks (average ~3 weeks) after initiation of therapy and often occurs after the course of antibiotic is completed, the delay being a few days to as long as six weeks. The onset is typically with fatigue, low grade fever, nausea and abdominal pain, followed by pruritus and jaundice. The pattern of liver enzyme elevations is typically cholestatic with marked elevations in alkaline phosphatase and gamma glutamyl transpeptidase (Case 1). In some instances, aminotransferase levels are markedly elevated giving a mixed (Case 2) or hepatocellular pattern (Case 3), particularly in younger patients with earlier onset of injury. In children, amoxicillin-clavulanate hepatotoxicity is typically anicteric and presents with nausea, vomiting and abdominal pain rather than jaundice and itching. The pattern of serum enzyme elevations is also much more likely to be hepatocellular in children, but the course of illness is typically benign. Because the liver injury may present days or weeks after stopping therapy, the

association of the liver injury with receipt of amoxicillin/clavulanate may be missed. Immunoallergic features (fever, rash, eosinophilia) can occur, but are not invariably present and are usually not prominent. Autoantibody formation is not common. The hepatic injury is idiosyncratic and is estimated to occur after ~1 in 2,500 prescriptions. The injury is more common in men than women, in the elderly and after multiple courses. Genetic studies indicate a link with HLA types, particularly the extended haplotype: DRB1*15:01-DRB5*01:01-DQB1*06:02.

Likehood score: A (well etablished cause of clinically apparent liver injury).

Mechanism of Injury

The cause of amoxicillin-clavulanate hepatotoxicity is unknown, but is probably immunoallergic in origin. Allergic manifestations can occur and include rash, fever, arthralgias and eosinophilia. Several studies have reported an HLA Class II association with DRB1*15:01 and the extended haplotype DRB1*15:01-DRB1*01:01-DQB1*06:02. An independent HLA Class I association has also been made with HLA-A*02:01. The liver injury appears to be due to the clavulanate rather than amoxicillin, as reexposure to amoxicillin alone has not been associated with recurrence (Case 5), whereas reexposure to the combination is usually followed by a more rapid onset of a more severe hepatic injury, which can include prolonged cholestasis and development of cirrhosis. Other beta lactamase inhibitors (tazobactam and subactam) have not been reported to cause a similar hepatic injury, although it has been reported with other penicillins when combined with clavulanate (ticarcillin/ clavulanate).

Outcome and Management

The liver injury caused by amoxicillin-clavulanate is typically associated with jaundice and can be severe and prolonged (with jaundice lasting 4 to 24 weeks), but rarely results in lasting injury or death. Deaths due to amoxicillin-clavulanate hepatic injury have been described, but largely in patients with other comorbidities including cirrhosis or with multiple exposures. In addition, rare instances of prolonged cholestasis and vanishing bile duct syndrome have been reported after acute amoxicillin-clavulanate injury. Corticosteroids have been used in patients with marked or prolonged cholestasis due to amoxicillin/clavulanate, but their efficacy has not been shown and their use cannot be recommended routinely. Cholestyramine or ursodiol may help alleviate symptoms but probably do not speed recovery. Rechallenge with amoxicillin-clavulanate results in recurrence and should be avoided. Amoxicillin alone, on the other hand, is safe and does not cause recurrence of liver injury except in the rare instance in which the penicillin rather than clavulanate is responsible for the liver injury.

Drug Class: Antiinfective Agents, Aminopenicillins

Other Drugs in the Subclass, Aminopenicillins: Ampicillin, Ampicillin-Sulbactam, Amoxicillin, Bacampicillin, Pivampicillin, Ticarcillin-Clavulanate

CASE REPORTS

Case 1. Cholestatic hepatitis from amoxicillin-clavulanate.

[Modified from a case in the database of the Drug-Induced Liver Injury Network]

A 75 year old man with a history of prostate cancer and regular alcohol use (2 to 3 drinks daily) was given amoxicillin-clavulanate (500 mg/125 mg) for chronic maxillary sinusitis. Because of persistent symptoms, the antibiotic was continued for 31 days. When seen in follow up two weeks later, he complained of jaundice and was admitted to the hospital for evaluation. He had symptoms of dark urine, weakness, and poor appetite. Blood test results showed a total bilirubin of 42.7 mg/dL, ALT 194 U/L, AST 107 U/L, Alk P 257 U/L. Tests for acute

hepatitis A, B, C and E were negative. Ultrasound of the abdomen showed no evidence of biliary obstruction or gallstones. During the hospitalization he developed profound anemia and thrombocytopenia requiring blood and platelet transfusions, and was further treated with corticosteroids and cyclophosphamide. His serum bilirubin peaked at 48.8 mg/dL and remained elevated for several months while aminotransferase and alkaline phosphatase levels were only modestly elevated (Table). The prothrombin time was elevated (INR 1.4 to 1.6) transiently and he developed mild confusion that was believed to be due to hepatic encephalopathy; he was treated with lactulose. During the hospitalization he developed renal failure and required dialysis. A liver biopsy was performed and findings were consistent with amoxicillin/clavulanate hepatotoxicity. He was hospitalized for 2 months and required another 2 months to recover fully. When seen 4 months after onset of the jaundice, he was back to his usual state of health and had normal laboratory tests, including normal aminotransferase and alkaline phosphatase levels, normal serum bilirubin and creatinine, and normal hemoglobin and platelet counts.

Key Points

Medication:	Amoxicillin-clavulanate 500/125 mg thrice daily for 31 days
Pattern:	Cholestatic (R=1.9)
Severity:	4+ (jaundice, hospitalization, severe thrombocytopenia and acute renal failure)
Latency:	45 days, 14 days after stopping
Recovery:	Slowly over 4 months
Other medications:	Guaifenesin

Laboratory Values

Time After Starting	Time After Stopping	ALT (U/L)	Alk P (U/L)	Bilirubin (mg/dL)	Other		
31 days	0	Amoxicillir	Amoxicillin-clavulanate stopped				
40 days	9 days	Jaundice	Jaundice				
6 weeks	2 weeks	126	195	43.8	Admission, platelets 2,000		
7 weeks	3 weeks	58	165	48.6			
2 months	4 weeks	59	248	48.8			
	5 weeks	44	198	42.4	Acute renal failure		
	7 weeks	40	195	19.4	Liver biopsy		
3 months	8 weeks	40	267	7.4			
98	67	47	242	3.6	Discharged, platelets 376,000		
158	127	11 81 0.6 Creatinine 2.5			Creatinine 2.5		
Normal Values		<42	<115	<1.2			

Comment

This case is an example of severe amoxicillin-clavulanate hepatotoxicity. Liver histology showed central lobular retention of bile with mixed infiltrates of lymphocytes, neutrophils and eosinophils in portal areas and in areas of focal spotty necrosis in the parenchyma. There was minimal steatosis, no fibrosis and no biliary damage or peribiliary fibrosis or edema. The findings were consistent with a drug induced intrahepatic cholestasis. The case was typical of amoxicillin/clavulanate hepatotoxicity in its onset 1 to 2 weeks after stopping therapy in an elderly man without other history or risk factors for liver or biliary disease. The severe thrombocytopenia and anemia were atypical, but similar nonhepatic manifestations of immunoallergic injury have been reported. Important conditions to exclude were biliary obstruction due to malignancy or gallstone disease and viral hepatitis.

Therapy should be limited to symptomatic management of pruritus and avoidance of further hepatic injury. Appropriate management calls for follow up documentation of resolution of liver injury. The patient should be warned to avoid any further exposure to amoxicillin/clavulanate. In view of the severity of the injury, use of amoxicillin alone might also be best avoided.

Case 2. Mixed cholestatic-hepatocellular injury due to amoxicillinclavulanate.

[Modified from a case in the database of the Drug-Induced Liver Injury Network]

A 52 year old man without major medical illnesses developed an upper respiratory infection and was given a 14 day course of amoxicillin-clavulanate. One and two months later, because of similar symptoms of fever and congestion, he was given a 2nd and 3rd 14 day course along with antihistamines and decongestants. Two weeks after the 3rd course of antibiotics, he developed abdominal pain, nausea, poor appetite, and itching, followed soon after by jaundice and dark urine. He was seen and blood tests revealed total bilirubin of 3.9 mg/dL with marked elevations in both ALT and alkaline phosphatase (Table). He was managed as an outpatient, did not undergo liver biopsy, and recovered symptomatically over the next few weeks. When seen one year later, he was asymptomatic and laboratory tests had returned to normal.

Key Points

Medication:	Amoxicillin-clavulanate 500/125 mg twice daily for 42 days
Pattern:	Mixed cholestatic-hepatocellular (R=2.3)
Severity:	2+ (jaundice, never hospitalized)
Latency:	58 days, 16 days after stopping
Recovery:	Six months after stopping
Other medications:	None

Laboratory Values

Time After Starting	Time After Stopping	ALT (U/L)	Alk P (U/L)	Bilirubin (mg/dL)	Other
2 weeks	0	Amoxicillin	Amoxicillin-clavulanate stopped		
4 weeks	16 days	Nausea, jau	ndice, arthral	gia	
	16 days	879	731	3.9	Albumin 4.3 g/dL
5 weeks	3 weeks	629	503	1.7	
7 weeks	5 weeks	141	182	0.8	
10 weeks	8 weeks	90	103	0.8	
14 weeks	3 months	88	95	0.6	INR 0.9
~1 year		57	76	0.7	Albumin 4.5 g/dL
Normal Values		<65	<126	<1.2	

Comment

This case is a typical example of mild-to-moderate amoxicillin-clavulanate hepatotoxicity. The early appearance of itching and high alkaline phosphatase levels (~6 fold elevated) indicated that the injury was "cholestatic," while the high AST and ALT levels (peak at 10-15 fold elevated) provided the evidence that the injury was

"mixed". The repeated courses of the combination within a short period of time may have predisposed to the hepatotoxicity. The jaundice and itching lasted only two weeks, and he eventually had full recovery.

Case 3. Hepatocellular injury due to amoxicillin-clavulanate.

[Modified from the database of the Drug-Induced Liver Injury Network]

A woman in her 20s received a ten day course of amoxicillin-clavulanate (500 mg/125 mg) twice daily for suspected sinusitis. She felt somewhat nauseated during therapy, but was then without symptoms until 8 to 9 weeks after stopping therapy when she developed anorexia, nausea, vomiting and abdominal pain. She noted dark urine and jaundice and went to an emergency room where blood tests were taken showing a total bilirubin of 12.5 mg/dL, ALT 3500 U/L, AST 2554 U/L, and alkaline phosphatase 187 U/L. She was also on birth control pills and had intermittently taken cold and sinus remedies and loratadine for allergic sinusitis during the previous two months. She received a one week course of nabumetone (a nonsteroidal antiinflammatory agent) and prednisone for acute neck spasms one month previously. She denied taking acetaminophen in the recent past and had no history of exposure to viral hepatitis or known risk factors for hepatitis or liver disease. She drank little alcohol and had never received amoxicillin-clavulanate previously. Tests for acute hepatitis A, B, C and E were negative as were tests for antinuclear and smooth muscle antibody. An ultrasound of the abdomen showed no evidence of biliary tract disease or gallstones. She did not undergo liver biopsy. Serum aminotransferase levels decreased while serum bilirubin rose to 27.2 mg/dL before falling (Table). Serum alkaline phosphatase levels were never very elevated. Prothrombin time was abnormal for a short period (peak INR 1.7). She recovered slowly and remained symptomatic with fatigue and abdominal discomfort for several months. When seen six months after onset of jaundice, she had no symptoms and serum bilirubin, aminotransferase and alkaline phosphatase levels had returned to normal.

Key Points

Medication:	Amoxicillin-clavulanate unknown dose twice daily for 14 days
Pattern:	Hepatocellular (R=66)
Severity:	4+ (jaundice, hospitalization and INR >1.5)
Latency:	80 days, 66 days since stopping medication
Recovery:	<2 months after onset of jaundice
Other medications:	Oral contraceptives for 4 years

Laboratory Values

Time After Starting	Time After Stopping	ALT (U/L)	Alk P (U/L)	Bilirubin (mg/dL)	Other
14 days	0	Amoxic	illin-clav	ulanate stop	ped
11 weeks	66 days	Jaundic	e followed	l by fatigue a	and severe itching
12 weeks	75 days	3500	187	12.5	Admission
13 weeks	78 days	955	156	16.4	
14 weeks	84 days	438	159	16.3	
	89 days	308	134	11.6	
15 weeks	91 days	210	145	7.0	Discharge
19 weeks	4 months	21	65	1.9	Asymptomatic
Normal Values		<42	<115	<1.2	

Comment

This case is an example of moderate-to-severe amoxicillin-clavulanate hepatotoxicity. Somewhat atypical was the long incubation period between stopping the antibiotic and onset of jaundice as well as the prominent elevations in serum aminotransferase levels, the heights of which suggested viral hepatitis or acetaminophen overdose. The prominent ALT elevations and symptoms of fatigue and nausea rather than itching indicate a hepatocellular rather than cholestatic pattern of injury. Amoxicillin-clavulanate is typically associated with cholestatic injury but hepatocellular patterns can occur, particularly in younger patients. The potential role of other medications taken at the time (including unacknowledged receipt of acetaminophen) should be considered. While self limited, the course of illness was severe and protracted. Further follow up is warranted to exclude a relapsing, acute onset of autoimmune hepatitis.

Case 4. Mixed cholestatic-hepatocellular injury due to amoxicillinclavulanate.

[Modified from a case in the database of the Drug-Induced Liver Injury Network]

A 60 year old man was hospitalized with cellulitis and was discharged on a 10 day course of amoxicillinclavulanate (500 mg/125 mg) twice daily. At the time of hospitalization, he had normal serum bilirubin, alkaline phosphatase and aminotransferase levels. Five days after stopping the antibiotic, he developed itching followed by poor appetite. After ten days of symptoms, he noted dark urine and pale stools and was seen in an emergency room. He was given intravenous fluids for dehydration and hydroxyzine for itching. When laboratory tests showed a total bilirubin of 7.7 mg/dL, alkaline phosphatase 286 U/L, ALT 387 U/L, and AST 134 U/L, he was admitted for evaluation. Prothrombin time and albumin levels were normal. He had no previous history of liver disease and drank little alcohol. Over the previous week he had taken acetaminophen (up to 4 grams daily) for back pain and itching. On a chronic basis, he took montelukast (Singular) and used inhalers (with albuterol, fluticasone and salmeterol) for asthma; was prescribed losartan (Cozaar) for hypertension; and, used several topical lotions for psoriasis. Tests for acute hepatitis A, B and C were negative. He had IgG antibody to HEV but IgM anti-HEV was negative. Autoantibodies including antinuclear and smooth muscle antibodies were not detected. An abdominal ultrasound showed no evidence of biliary obstruction and no gallstones. A liver biopsy was obtained which showed a mixed hepatocellular-cholestatic pattern of injury compatible with drug induced liver disease. During the next several days serum bilirubin levels rose to 8.1 mg/dL, but then started to decline even as ALT and AST levels rose to 8 to 20 fold above normal and alkaline phosphatase to 2.5 times normal (Table). He was discharged after a week on symptomatic therapy with cholestyramine. He remained symptomatic with itching and fatigue for another month, but on a return visit two months after presentation he felt well and laboratory tests had returned almost to baseline values.

Key Points

Medication:	Amoxicillin-clavulanate 875/125 mg twice daily for 10 days
Pattern:	Mixed cholestatic-hepatocellular (R=3.9)
Severity:	3+ (jaundice, hospitalization)
Latency:	24 days, 14 days after stopping
Recovery:	Within 2 months of stopping
Other medications:	Montelukast, fluticasone-salmeterol, albuterol, losartan, and synthroid chronically

Time After Starting	Time After Stopping	ALT (U/L)	Alk P (U/L)	Bilirubin (mg/dL)	Other
Baseline		44	65	0.6	
10 days	0	Amox	Amoxicillin-clavulanate stopped		
24 days	14 days	371	301	7.8	Pain, itching
4 weeks	19 days	336	286	6.4	Severe itching
5 weeks	23 days	695	244	3.7	
5 weeks	27 days	724	188	2.3	Biopsy
7 weeks	5 weeks	250	142	1.3	
8 weeks	6 weeks	58	101	0.9	
Normal Values		<42	<115		

Laboratory Values

Comment

The history and presentation of this case were typical of moderately severe amoxicillin-clavulanate hepatotoxicity with onset of itching followed by jaundice occurring 1 to 2 weeks after stopping a ten day course of treatment in an older man with no other obvious cause of liver disease. The pattern of serum enzyme results with prominent elevations in serum aminotransferase and alkaline phosphatase levels suggested mixed hepatocellular-cholestatic injury which was also shown by liver histology. While heavy acetaminophen use may have contributed to the initial aminotransferase elevations, the liver biopsy did not show evidence of typical acetaminophen injury. Other medications being taken (montelukast, albuterol, fluticasone, salmeterol, losartan) had been used chronically, have not or only rarely been linked to liver injury and were restarted without recurrence of liver disease. The drug induced liver injury was fully reversible, but symptoms and illness lasted for almost two months.

Case 5. Cholestatic hepatitis due to clavulanate.

[Modified from: Peroux JL, Peroux E, Jais F, Philit F, Chichmanian RM. [Augmentin hepatotoxicity: responsibility of clavulanic acid? Apropos of a case] Gastroenterol Clin Biol 1992; 16: 102-3. French. *PubMed Citation*]

A 20 year old woman developed jaundice 10 days after starting an intravenous regimen of amoxicillinclavulanate (2 grams daily) for bacterial pneumonia and pleurisy. She had no history of liver disease, alcohol abuse or risk factors for viral hepatitis. Serum bilirubin, alkaline phosphatase and aminotransferase levels were normal on admission, but rose gradually during the hospitalization and antibiotic therapy (Table). When jaundice was first noted laboratory testing showed a total bilirubin of 2.6 mg/dL (direct 1.9 mg/dL), ALT 112 U/L, AST 58 U/L, Alk P 425 U/L, and GGT 212 U/L. The amoxicillin-clavulanate was discontinued, but antibiotic therapy was still needed and she was treated with amoxicillin (3 grams daily) and metronidazole (1.5 grams daily). On this regimen, her pulmonary symptoms and fever improved rapidly and liver tests gradually improved and were in the normal range 30 days after stopping amoxicillin with clavulanate.

Key Points

Medication:	Amoxicillin-clavulanate intravenously for 10 days
Pattern:	Cholestatic (R=0.6)
Severity:	3+ (jaundice, hospitalization prolonged)

Table continued from previous page.

Medication:	Amoxicillin-clavulanate intravenously for 10 days
Latency:	10 days, still on therapy
Recovery:	Within 1 month of stopping
Other medications:	Amoxicillin, metronidazole

Laboratory Values

Time After Starting	Time After Stopping		Alk P (U/L)	Bilirubin (mg/dL)	Other
0	0	39	112	0.5	Admission
		Intrav	enous a	moxicillin-clavulan	ate started
1 day	0	96	146	0.9	
2 days	0	75	206		
5 days	0	117	417		
7 days	0	112	425	2.6	
10 days	0	Amoxicillin-clavulanate stopped			
11 days	1 day	58	285	2.2	
14 days	4 day	27	296	2.1	Amoxicillin started
17 days	7 day	17	222	1.5	
6 weeks	4 weeks	10	95	0.5	
Normal Values	Normal Values		<133	<1.2	

Comment

The history and presentation of this case were typical of mild amoxicillin-clavulanate hepatotoxicity with onset within a week of starting therapy. Of interest was that routine laboratory testing demonstrated a rapid onset of hepatic injury that was subclinical for the first week. The rapidity of onset suggests previous exposure to amoxicillin-clavulanate, although no such history was obtained. Once the antibiotic combination was stopped, recovery was prompt. Some form of antibiotic therapy was considered necessary and the attending physicians felt comfortable using amoxicillin as the injury was thought to be due to clavulanate rather than the penicillin. The subsequent course supported their conclusions. Several other cases in the literature have supported the belief that the injury in typical amoxicillin-clavulanate hepatotoxicity is due to clavulanate. However, in an individual case, there is always the possibility that the hepatic injury is actually due to the amoxicillin; although in that situation the clinical syndrome should be dominated by signs and symptoms of hypersensitivity (rash, fever, eosinophilia) usually arising within 1 to 2 weeks of starting therapy, features that can occur with typical amoxicillin-clavulanate liver injury but are generally mild and self limited.

PRODUCT INFORMATION

REPRESENTATIVE TRADE NAMES

Amoxicillin-Clavulanate - Augmentin®

DRUG CLASS

Box continued from previous page.

Antiinfective Agents

COMPLETE LABELING

Product labeling at DailyMed, National Library of Medicine, NIH

CHEMICAL FORMULAS AND STRUCTURES

DRUG	CAS REGISTRY NO	MOLECULAR FORMULA	STRUCTURE
Amoxicillin- Clavulanate	79198-29-1	C16-H19-N3-O5-S.C8-H9-N- O5	
Amoxicillin Anhydrous	26787-78-0	C16-H19-N3-O5-S	
Clavulanic Acid	58001-44-8	C8-H9-N-O5	

ANNOTATED BIBLIOGRAPHY

References updated: 30 March 2016

- Zimmerman HJ. Hepatic injury from the treatment of infectious and parasitic diseases. In, Zimmerman HJ. Hepatotoxicity: the adverse effects of drugs and other chemicals on the liver. 2nd ed. Philadelphia: Lippincott, 1999, pp. 589-638.
- (Expert review of hepatotoxicity published in 1999; amoxicillin-clavulanate is capable of causing acute liver injury with jaundice that is typically cholestatic in pattern but can be hepatocellular, onset is delayed as long as 35 days after the drug is stopped, etiology is idiosyncrasy likely to be immunologic, most likely due to the clavulanate, recovery in 30 to 74 days, rarely causes fatality).
- Moseley RH. Hepatotoxicity of antimicrobials and antifungal agents. In, Kaplowitz N, DeLeve LD, eds. Druginduced liver disease. 3rd ed. Amsterdam: Elsevier, 2013, pp. 463-82.
- (*Review of hepatotoxicity of antibiotics; amoxicllin-clavulanate typically causes a cholestatic or mixed pattern of injury with a preponderance in elderly men and with consecutive uses; features of hypersensitivity occur in up*

to 60% of patients; fatal, severe and prolonged cases have been described; mechanism is likely immunologic and due to the clavulanate rather than amoxicillin).

- Petri WA Jr. Penicillins, cephalosporins, and other β-lactam antibiotics. In, Brunton LL, Chabner BA, Knollman BC, eds. Goodman & Gilman's the pharmacological basis of therapeutics. 12th ed. New York: McGraw-Hill, 2011, pp. 1477-1504.
- (Textbook of pharmacology and therapeutics).
- Brogden RN, Carmine A, Heel RC, Morley PA, Speight TM, Avery GS. Amoxycillin/ clavulanic acid: a review of its antibacterial activity, pharmacokinetics and therapeutic use. Drugs 1981; 22: 337-62. PubMed PMID: 7037354.
- (Review of structure, pharmacology, mechanism of action, therapeutic efficacy and safety of oral amoxicillinclavulanate: "there have been occasional reports of mild increases in serum transaminase levels during treatment").
- van den Broek JW, Buennemeyer BL, Stricker BH. [Cholestatic hepatitis caused by a combination of amoxicillin and clavulanic acid (Augmentin)]. Ned Tijdschr Geneeskd 1988; 132: 1495-7. Dutch. PubMed PMID: 3173514.
- (66 year old man developed fatigue and jaundice shortly after a 2nd course of amoxicillin-clavulanate [bilirubin 7.1 rising to 35 mg/dL, ALT 325 U/L, GGT 272 U/L], resolving in 3 months and recurring 4 days after restarting the combination but not with two courses of amoxicillin alone).
- Verhamme M, Ramboer C, Van de Bruaene P, Inderadjaja N. Cholestatic hepatitis due to an amoxycillin/ clavulanic acid preparation. J Hepatol 1989; 9: 260-4. PubMed PMID: 2809168.
- (Two cases: 60 and 53 year old men developed jaundice and pruritus 3 weeks after starting 10 day course of amoxicillin-clavulanate [bilirubin 13.0 and 13.8 mg/dL, ALT 356 and 314 U/L, Alk P 647 and 273 U/L], resolving within 2 to 3 months).
- Schneider JE, Kleinman MS, Kupiec JW. Cholestatic hepatitis after therapy with amoxicillin/clavulanate potassium. N Y State J Med 1989; 89: 355-6. PubMed PMID: 2739957.
- (44 year old woman developed abdominal pain and jaundice 20 days after starting a 10 day course of amoxicillinclavulanate [bilirubin 1.9 rising to 4.6 mg/dL, AST 129 U/L, Alk P 259 U/L], resolving within 2 months).
- Dowsett JF, Gillow T, Heagerty A, Radcliffe M, Toadi R, Isle I, Russell RCG. Amoxycillin/clavulanic acid (Augmentin)-induced intrahepatic cholestasis. Dig Dis Sci 1989; 34: 1290-3. PubMed PMID: 2752877.
- (75 year old man developed nausea followed by jaundice ~1 week after finishing a 10 day course of amoxicillin/ clavulanate [bilirubin 4.7 rising to 16 mg/dL, ALT 225 U/L, Alk P 316 U/L, eosinophilia], biopsy showing cholestatic hepatitis, resolving in 14 weeks).
- Reddy KR, Brillant P, Schiff ER. Amoxicillin-clavulanate potassium-associated cholestasis. *Gastroenterology* 1989; 96: 1135-41. PubMed Citation
- (Between 1984-87, 18 cases of possible amoxicillin-clavulanate hepatotoxicity with adequate documentation were reported to the sponsor, including 7 cholestatic, 6 mixed and 4 hepatocellular cases; 16 with jaundice, 14 pruritus, 2 fever; eosinophilia in 60%; latency 2-43 days, 14 occuring after stopping; bilirubin 2.7-44 mg/dL, ALT 43-440 U/L, Alk P 63-1330 U/L, resolving in 30-74 days, no fatalities).
- Stricker BH, Van den Broek JW, Keuning J, Eberhardt W, Houben HGJ, Johnson M, Blok APR. Cholestatic hepatitis due to antibacterial combination of amoxicillin and clavulanic acid (augmentin).
- *Dig Dis Sci. 1989; 34: 1576-80. PubMed Citation (5 cases of suspected amoxicillin-clavulanate hepatotoxicity; all were men with jaundice arising 6 to 25 days after starting antibiotic [bilirubin 9-32 times ULN, ALT 2 to 6.3 times ULN, Alk P 1 to 2.6 times ULN], two were rechallenged and both had recurrence).*

- Cleau D, Jobard JM, Alves T, Gury S, Rey B, Vuillemard M, Noirot A, et al. [Cholestatic hepatitis induced by the amoxicillin-clavulanic acid combination. A case and review of the literature]. Gastroenterol Clin Biol 1990; 14: 1007-9. French. PubMed PMID: 2289658.
- (50 year old man developed pruritus and jaundice a week after completing a 7 day course of amoxicillinclavulanate [bilirubin 14.2 mg/dL, ALT 76 U/L, Alk P 526 U/L], resolving within 2 months).
- Pelletier G, Ink O, Fabre M, Hagège H. [Hepatic cholestasis probably due to the combination of amoxicillin and clavulanic acid]. Gastroenterol Clin Biol 1990; 14: 601. French. PubMed PMID: 2397873.
- (79 year old woman developed pruritus and jaundice 4 weeks after starting an 8 day course of amoxicillinclavulanate [bilirubin 12.2 mg/dL, ALT 5 times ULN, Alk P 8.5 times ULN], resolving within 4 months).
- Reddy KR, Schiff ER. Hepatitis and augmentin. Dig Dis Sci 1990; 35: 1045-6. PubMed Citation
- (Letter in response to Sticker commenting on the rarity of the injury in children despite wide scale use).
- Michielsen PP, Van Outryve MJ, Van Marck EA, De Maeyer MH, Pelckmans PA, Van Maercke YM. Amoxycillin/clavulanic acid induced cholestasis. J Hepatol 1990; 11: 392. PubMed PMID: 2290034.
- (24 year old man developed jaundice 23 days after starting 7 day course of amoxicillin-clavulanate [bilirubin 6.8 mg/dL, ALT 119 U/L, Alk P 275 U/L], resolving within 2 months).
- Escallier F, Dalac S, Caillot D, Boulitrop C, Collet E, Lambert D. [Erythema multiforme, aplasia, cholestatic hepatitis during treatment with Augmentin (amoxicillin + clavulanic acid)]. Rev Med Interne 1990; 11: 73-5. French. PubMed PMID: 2326558.
- (82 year old man developed rash 5 days after starting amoxicillin-clavulanate, which was continued for 3 more days when he developed fever, erythema multiforme, bone marrow aplasia, [platelets 13,000, leukocytes 200] and liver test abnormalities [ALT 88 U/L, Alk P 367 U/L], with complex course but ultimate recovery within next six months).
- Alexander P, Roskams T, Van Steenbergen W, Peetermans W, Desmet V, Yap SH. Intrahepatic cholestasis induced by amoxicillin/clavulanic acid (Augmentin): a report on two cases. *Acta Clin Belg 1991; 46: 327-32. PubMed Citation*
- (Two cases: 61 year old man developed pruritus and jaundice at end of 28 day course of amoxicillin-clavulanate [bilirubin 3.8 mg/dL, ALT 1.5 times ULN, Alk P 4 times ULN, eosinophils 9%], resolving within 12 weeks; 78 year old woman developed pruritus and jaundice 2 weeks after a course of amoxicillin-clavulanate [bilirubin 18.8 mg/dL, ALT 60 U/L, Alk P 682 U/L], resolving within 8 weeks).
- Wong FS, Ryan J, Dabkowski P, Dudley FJ, Sewell RB, Smallwood RA. Augmentin-induced jaundice. Med J Aust 1991; 154: 698-701. PubMed PMID: 2034154.
- (8 cases of amoxicillin-clavulanate associated jaundice, ages 45 to 77 years, 6 men, 2 women; onset 0.5 to 6 weeks after stopping [peak bilirubin 6.4 to 34.7 mg/dL, ALT 77 to 921 U/L, Alk P 188 to 1420 U/L], resolving in 2-21 weeks).
- Silvain C, Fort E, Levillain P, Labat-Labourdette J, Beauchant M. Granulomatous hepatitis due to combination of amoxicillin and clavulanic acid. Dig Dis Sci 1992; 37: 150-2. PubMed PMID: 1728522.
- (79 year old man developed jaundice 20 days after finishing 14 day course of amoxicillin-clavulanate [bilirubin 16.9 mg/dL, ALT 350 U/L, Alk P 1870 U/L, eosinophils 610], resolving within 11 weeks).
- Ryan J, Dudley FJ. Cholestasis with ticarcillin-potassium clavulanate (Timentin). Med J Aust 1992; 156: 291. PubMed PMID: 1738336.

- (75 year old man developed pruritus and jaundice 31 days after completing a 9 day course of intravenous ticarcillin/ clavulanate and shortly after a 28 day course of induction chemotherapy for lymphoma [bilirubin 8.1 mg/dL, ALT 448 U/L, Alk P 1330 U/L], with persistent jaundice and death 4 weeks later while liver tests were improving).
- Cabelleria Rovira E, Masso Ubeda RM, Arago López JV, Sanchís Closa A. [Cholestatic hepatitis from amoxicillin-clavulanic acid]. An Med Interna 1992 Jul; 9(7): 360-1. Spanish. PubMed PMID: 1633249.
- (71 year old man developed pruritus and jaundice which resolved in 3 weeks and then recurred after a 3 day course of amoxicillin-clavulanate [bilirubin 17 mg/dL, ALT 126 U/L, Alk P 577 U/L], at which point a history of taking this antibiotic before the first episode was obtained).
- Larrey D, Vial T, Micaleff A, Babany G, Morichau-Beauchant M, Michel H, Benhamou JP. Hepatitis associated with amoxycillin-clavulanic acid combination report of 15 cases.
- *Gut 1992; 33: 368-71. PubMed Citation (15 cases of amoxicillin-clavulanate hepatotoxicity reported to French regional centers including 12 men [80%], ages 39-82 years [mean=64], with latency of 7-60 days [mean=18], 11 arising after stopping, all had jaundice and 12 pruritus; no fatalities).*
- Hebbard GS, Smith KG, Gibson PR, Bhathal PS. Augmentin-induced jaundice with a fatal outcome. *Med J Aust* 1992; 156: 285-6. *PubMed Citation*
- (81 year old man developed jaundice and pruritus 1 week after completing a 4 week course of amoxicillinclavulanate [bilirubin 21.9 mg/dL, ALT 97 U/L, Alk P 231 U/L], with prolonged jaundice [bilirubin 35.8 mg/ dL], drowsiness and death 5 weeks later).
- Peroux JL, Peroux E, Jais F, Philit F, Chichmanian RM. [Augmentin hepatotoxicity: responsibility of clavulanic acid? Apropos of a case] Gastroenterol Clin Biol 1992; 16: 102-3. French. PubMed PMID: 1537473.
- (20 year old woman developed jaundice 7 days after starting amoxicillin-clavulanate [bilirubin 3.7 mg/dL, ALT 112 U/L, Alk P 425 U/L], resolving within 30 days despite continuing treatment with amoxicillin).
- Yap I, Gwee KA, Wee A. Augmentin-induced cholestatic jaundice--a case report. Singapore Med J 1993; 34: 464-5. PubMed PMID: 8153703.
- (55 year old man with alcoholic pancreatitis developed jaundice after 6 days of amoxicillin-clavulanate [bilirubin 4.0 mg/dL, AST 56 U/L, Alk P 324 U/L], resolving within 5 weeks).
- Horsmans Y; Geubel AP. Amoxycillin-clavulanic acid-erythromycin cross-liver toxicity: a case report. J Hepatol 1994; 21: 911-2. PubMed PMID: 7890912.
- (67 year old man developed fatigue and pruritus 6 days after completing 5 day course of amoxicilllin-clavulanate [bilirubin unclear, ALT 185 U/L, Alk P 235 U/L], and enzyme elevations later rose upon starting erythromycin [twice]).
- Hanssens M, Mast A, Van Maele V, Pauwels W. [Cholestatic jaundice caused by amoxicillin-clavulanic acid in 4 patients]. Ned Tijdschr Geneeskd 1994; 138: 1481-3. Dutch. PubMed PMID: 8052321.
- (4 cases of amoxicillin-clavulanate hepatotoxicity: 2 men and 2 women, ages 72 to 84 years, treated for 7 to 10 days with onset after 25 to 35 days with jaundice, fever and pruritus or abdominal pain [bilirubin 7.2-18.8 mg/dL, ALT 77-320 U/L, Alk P 195-630 U/L], resolving in 40 to 60 days).
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- (Analysis of 34 cases of amoxicillin-clavulanate associated jaundice compared to 136 controls found male sex [65% vs 46%], older age [60 vs 40 years], longer treatment [5.7 vs 7 days], but not dose or other medical conditions, were more frequent in cases with liver injury).

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- 58 year old man developed jaundice and pruritus 36 days after starting a 10 day course of amoxicillin-clavulanate, resolving within 2 months).
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- (25 year old man developed abdominal pain and jaundice 5 weeks after starting 6 day course of amoxicillinclavulanate [bilirubin 7.0 mg/dL, ALT 268 U/L, Alk P 2287 U/L, amylase 878 U/L], resolving within 1 month).
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- Hautekeete ML, Brenard R, Horsmans Y, Henrion J, Verbist L, Derue G, Druez P, et al. Liver injury related to amoxycillin-clavulanic acid: interlobular bile-duct lesions and extrahepatic manifestations. J Hepatol 1995; 22: 71-7. PubMed PMID: 7751590.
- (Analysis of liver histology from 7 cases of amoxicillin-clavulanate hepatotoxicity; intralobular bile duct abnormalities found in all biopsies but no ductopenia).
- Reddy KR, Schiff ER. Hepatotoxicity of antimicrobial, antifungal, and antiparasitic agents. *Gastroenterol Clin North Am 1995; 24: 923-36. PubMed Citation*
- (*Review of hepatotoxicity of antibiotics including amoxicillin-clavulanate; histologically centrizonal cholestasis is characteristic as well as mild-to-moderate inflammation and bile duct abnormalities*).
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- (73 year old woman developed jaundice 5 days after starting amoxicillin-clavulanate [bilirubin 6.2 mg/dL, ALT 140 U/L, Alk P 1783 U/L], resolving within one month).
- Bralet MP, Zafrani ES.[Hepatitis caused by the amoxicillin-clavulanic acid combination. An example of druginduced biliary hepatotoxicity] Ann Pathol 1996; 16: 425-9. French. PubMed PMID: 9090930.
- (Five cases of amoxicillin-clavulanate hepatotoxicity, biopsies showing centrolobular cholestasis in all and bile duct injury without ductopenia in 4).

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- (47 year old man developed pruritus and jaundice one month after starting amoxicillin-clavulanate [bilirubin 11.6 mg/dL, ALT 159 U/L, Alk P 195 U/L, 4% eosinophils], resolving within 3 months).
- Rene JM, Buenestado J, Pinol MC. [Hepatotoxicity caused by amoxicillin-clavulanic acid] Gastroenterol Hepatol 1997; 20: 337-8. Spanish. PubMed PMID: 9296855.
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- (31 year old man developed pruritis and jaundice the day after finishing a 5 day course of amoxicillin-clavulanate [bilirubin 2.9 mg/dL, ALT 291 U/L, Alk P 509 U/L], yet had tolerated amoxicillin alone and later had recurrence after 2 days of reexposure to amoxicillin-clavulanate [bilirubin 3.8 mg/dL, ALT 396 U/L, Alk P 616 U/L], resolving within 3 months of stopping).
- de Haan F, Stricker BH. [Liver damage associated with the combination drug amoxicillin-clavulanic acid (Augmentin)]. Ned Tijdschr Geneeskd 1997; 141: 1298-301. Dutch. PubMed PMID: 9380177.
- (Summary of 40 cases of amoxicillin-clavulanate hepatotoxicity reported to Dutch Drug Safety Unit between 1982-96, mean age 61 years, 70% men, latency mean=3 weeks from starting [maximum 55 days], resolved in all [mean=6 weeks, range 10 to 168 days], 5 had recurrence on reexposure).
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- Two cases: 68 year old man developed jaundice and pruritus 7 days after finishing 10 day course of amoxicillinclavulanate [bilirubin 14.8 mg/dL, ALT 378 U/L, Alk P 542 U/L], resolving in 2 months; 95 year old man developed abdominal pain and jaundice 33 days after starting 2 week course [bilirubin 7.7 mg/dL, ALT 225 U/L, Alk P 1229 U/L], resolving within 2 months).
- Nathani MG, Mutchnick MG, Tynes DJ, Ehrinpreis MN. An unusual case of amoxicillin/clavulanic acid-related hepatotoxicity. Am J Gastroenterol 1998; 93: 1363-5. PubMed PMID: 9707067.
- (40 year old woman developed rash and jaundice 4 weeks after finishing 18 day course of amoxicillin-clavulanate [bilirubin 10.3 mg/dL, ALT 199 U/L, Alk P 362 U/L, eosinophils 7%], resolving and then recurring within 5 days of restarting [bilirubin 1.2 mg/dL, ALT 199 U/L, Alk P 362 U/L], resolving in 4 weeks).

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- (21 year old man developed itching and jaundice 8 days after stopping a 10 day course of amoxicillin-clavulanate [bilirubin 11.8 rising to 33 mg/dL, ALT 840 U/L, Alk P 390 U/L], resolving within 10 weeks of onset).
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- (73 year old developed jaundice 5 weeks after starting 2 week course of amoxicillin-clavulanate [bilirubin 20.1 mg/dL, ALT 12.5 times ULN, Alk P 3.5 times ULN], resolving within 6 weeks)
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- (72 year old man developed jaundice and pruritus 36 days after starting 20 day course of amoxicillin-clavulanate [bilirubin 4.3 rising to 16 mg/dL, ALT 470 U/L, Alk P 400 U/L], resolving in 11 weeks with apparent clinical response to ursodiol therapy).
- Hautekeete ML, Horsmans Y, Van Waeyenberge C, Demanet C, Henrion J, Verbist L, Brenard R, et al. HLA association of amoxicillin-clavulanate--induced hepatitis. Gastroenterology 1999; 117: 1181-6. PubMed PMID: 10535882.
- (35 patients with amoxicillin-clavulanate hepatotoxicity were tested for HLA A and B class associations: DRB1*1501-DRB5*0101-DQB1*0602 haplotype found in 57% of cases vs 15% of controls; less common with hepatocellular injury pattern).
- Limauro DL, Chan-Tompkins NH, Carter RW, Brodmerkel GJ Jr, Agrawal RM. Amoxicillin/clavulanateassociated hepatic failure with progression to Stevens-Johnson syndrome. Ann Pharmacother 1999; 33: 560-4. PubMed PMID: 10369618.
- (37 year old man developed jaundice, rash and pruritus 3 weeks after stopping 10 day course of amoxicillinclavulanate [bilirubin 27.7 mg/dL, ALT 53 U/L, Alk P 562 U/L, eosinophils 10%], with progression of rash to Stevens Johnson syndrome, septicemia and death).
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- (22 cases of amoxicillin-clavulanate hepatotoxicity tested for HLA class II associations: DRB1*1501 found in 70% vs 20% of 134 controls: all had extended haplotype of DRB1*1501-DRB5*0101-DQA1*0102-DQB1*0602).

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- (24 year old woman developed pruritus at the end of a 10 day course of amoxicillin-clavulanate and jaundice 10 days later [bilirubin 10.5 mg/dL, ALT 30 U/L, Alk P 314 U/L], resolving within 2 months).
- Chawla A; Kahn E; Yunis EJ; Daum F. Rapidly progressive cholestasis: An unusual reaction to amoxicillin/ clavulanic acid therapy in a child.J Pediatr 2000; 136: 121-3. PubMed PMID: 10636987.
- (2 year old boy developed rash 1 day after a 10 day course of amoxicillin-clavulanate with jaundice 3 days later [bilirubin 3.5 rising to 11 mg/dL, ALT 246 U/L, Alk P 630 U/L, GGT 555 U/L], with persistence of cholestasis requiring liver transplant 8 months later, explant showed vanishing bile duct syndrome).
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- (Two cases: 71 year old man developed jaundice 10 days after 7 day course of amoxicillin-clavulanate [bilirubin 6.4 mg/dL, ALT 77 U/L, Alk P 258 U/L], with slow resolution and apparent reponse to ursodiol; 81 year old man developed jaundice 1 week after a 1 week course [bilirubin 18.6 mg/dL, ALT 87 U/L, Alk P 427 U/L], resolving in 3 months and apparent response to ursodiol).
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- (Systematic review of reports of adverse reactions to amoxicillin-clavulanate found 208 cases of liver injury, 158 could be evaluated: 70% men, mean age 60 years, mean duration of therapy 2 weeks, latency to onset averaging 25 days and mean time to resolution 12 weeks; 55% had mixed pattern of injury; 3 fatalities).
- Berg P, Hahn EG. Hepatotoxic reactions induced by beta-lactamase inhibitors. Eur J Med Res 2001; 6: 535-42. PubMed PMID: 11772541.
- (Systematic review of hepatotoxicity of beta lactamase inhibitors: convincing evidence for hepatotoxicity found only for combinations with clavulanate: case report of 55 year old man developing jaundice 2 weeks after a 14 day course of amoxicillin/clavulanate [bilirubin rising to ~22 mg/dL, Alk P ~800 U/L]).
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- (Two cases: 62 year old man developed pruritus 2.5 weeks after finishing a 1 week course of amoxicillin-clavulanate [bilirubin 29.0 mg/dL, ALT 470 U/L, Alk P 271 U/L, eosinophils 8%], resolving within 5 weeks; 58 year old man developed pruritus 3 weeks after finishing 1 week course of therapy [bilirubin 5.5 mg/dL, ALT 248 U/L, Alk P 255 U/L, no eosinophilia], resolving within 6 weeks).
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- (61 year old woman with cirrhosis developed jaundice 54 days after finishing a 21 day course of amoxicillinclavulanate [bilirubin 27.1 mg/dL, ALT 73 U/L, Alk P normal], with progressive liver failure and death 11 days later).

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- (42 year old man developed pruritus and jaundice 5 weeks after starting 4 day course of amoxicillin-clavulanate, with prolonged course and worsening upon reexposure, liver biopsies showing progression to cirrhosis).
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- (Review of antibiotic induced liver injury, including amoxcillin-clavulanate).
- Zaidi SA. Hepatitis associated with amoxicillin/clavulanic acid and/or ciprofloxacin. Am J Med Sci 2003; 325: 31-3. (PubMed PMID: 12544082.
- 80 year old man developed rash 8 days after starting amoxicillin-clavulanate and 5 days after starting ciprofloxacin and found to have abnormal liver tests 5 days later [bilirubin 0.8 rising to 1.8 mg/dL, ALT 154 to 972 U/L, Alk P 120 to 358 U/L], resolving in 1 month).
- Martí J. [Cholestatic hepatitis due to amoxicillin-clavulanic acid with positive re-exposure]. Enferm Infecc Microbiol Clin 2003; 21: 322-3. Spanish. PubMed PMID: 12809593.
- (86 year old man developed jaundice after an 8 day course of amoxicillin-clavulanate [bilirubin 8.5 mg/dL, ALT 388 U/L, Alk P 337 U/L], resolving within 4 weeks; had a previous history of similar response).
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- (Summary of 25 years of adverse drug reaction reporting in Sweden identified 103 cases of druginduced acute liver failure: none were attributed to amoxicillin-clavulanate).
- Fontana RJ, Shakil AO, Greenson JK, Boyd I, Lee WM. Acute liver failure due to amoxicillin and amoxicillin/ clavulanate. Dig Dis Sci 2005; 50: 1785-90.
- *PubMed Citation* (Patient found to have abnormal liver tests [ALT 141 U/L, Alk P 98 U/L, normal bilirubin] 19 days after starting a 12 day course of amoxicillin-clavulanate, and admitted with acute liver failure 1 month later with bilirubin 27.6 mg/dL, ALT 620 U/L, Alk P 170 U/L, INR 2.0; dying 10 days later).
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- (At least 6 months follow up on 406 patients with drug induced liver injury in Spanish database identified 28 with evidence of chronic liver injury; amoxicillin-clavulanate accounted for 4 cases, rate of 6%; 3 had cholestatic enzyme pattern).
- Lucena MI, Andrade RJ, Fernández MC, Pachkoria K, Pelaez G, Durán JA, Villar M, et al.; Spanish Group for the Study of Drug-Induced Liver Disease (Grupo de Estudio para las Hepatopatías Asociadas a Medicamentos (GEHAM)). Determinants of the clinical expression of amoxicillin-clavulanate hepatotoxicity: a prospective series from Spain. Hepatology 2006; 44: 850-6. PubMed PMID: 17006920.
- (Series of 69 patients with amoxicillin-clavulanate hepatotoxicity which represented ~14% of drug induced liver injury; mean time to onset of jaundice 16 days [range 1-71], 2-55 days after stopping; 52% males; mean age 56 years; 31% cholestatic, 33% mixed and 36% hepatocellular; 1 patient died, 4 developed chronicity; hepatocellular cases were younger in age and had lower bilirubin and higher ALT levels).
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- Chalasani N, Fontana RJ, Bonkovsky HL, Watkins PB, Davern T, Serrano J, Yang H, Rochon J; Drug Induced Liver Injury Network (DILIN). Causes, clinical features, and outcomes from a prospective study of drug-induced liver injury in the United States. Gastroenterology 2008; 135: 1924-34. PubMed PMID: 18955056.
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- (36 year old former alcoholic developed jaundice 10 days after stopping fluconazole and 10 days after stopping 3 day course of amoxicillin-clavulanate [bilirubin 18.7 mg/dL, ALT 128 U/L, AST 125 U/L, Alk P 539 U/L, 26% eosinophils], with subsequent worsening jaundice and ascites but eventual recovery in 2 months: acute on chronic injury).
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- (76 year old man developed fever, pruritus and jaundice 4 weeks after starting a 3 week course of amoxicillinclavulanate [bilirubin 2.2 rising to 50.1 mg/dL, ALT 352 U/L, Alk P 266 U/L, eosinophils 8%], with prompt clinical response to prednisone [1 mg/kg/day] and ursodiol and recovery in 10 weeks).
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- (62 year old man with a hepatic hydatid cyst and previous episode of unexplained jaundice, redeveloped jaundice and pruritus 20 days after starting amoxicillin-clavulanate [bilirubin 12 mg/dL, ALT and Alk P normal]).
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- (Among 1038 cases of drug induced liver injury enrolled in a US database between 2004 and 2014, 117 were attributed to amoxicillin-clavulanate [9%]; mean age 60 years, 62% men, mean latency 29 days, mostly jaundiced and cholestatic [median bilirubin 6 mg/dL, ALT 362 U/L, Alk P 288 U/L, R value 2.7], 3 underwent liver transplant with resolution in almost all cases, although sometimes delayed).