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Primary sclerosing cholangitis and cholangiocarcinoma

https://pubmed.ncbi.nlm.nih.gov/16496232

Primary sclerosing cholangitis (PSC), a cholestatic liver disease characterized by fibrosing inflammatory damage of the biliary tree, is a risk factor for **cholangiocarcinoma** (CCA). Indeed, the prevalence of CCA i...

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Cholangiocarcinoma (CCA) is the most common malignancy in patients with **primary sclerosing cholangitis** (PSC) and carries a high rate of mortality. Although the pathogenesis of CCA in PSC is largel...

Cited by: 11 **Author:** Junmin Song, Junmin Song, Yang Li, Christ...

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Purpose of review: **Primary sclerosing cholangitis** (PSC) is a rare cholestatic liver disease characterized by progressive fibroinflammatory destruction of the intrahepatic and/or extrahepatic bile ducts. It is...

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Primary Sclerosing Cholangitis

Medical Condition

Chronic liver disease involving inflammation, scarring and narrowing of bile ducts.

- Very rare (Fewer than 20,000 cases per year in US)
- Requires lab test or imaging
- Treatable by a medical professional
- Can last several months or years

The cause is not clearly understood. The symptoms include fatigue, pain in the upper right abdomen, night sweats, enlarged spleen, enlarged liver, chills, weight loss, jaundice and itching. The known cure is liver transplant. It can be treated by managing the symptoms using drugs such as antibiotics, opioid antagonists, bile acid sequestrants and ursodeoxycholic acid (UDCA).

Symptoms

The most common symptoms are fatigue and intense itching. Other symptoms appear as the disease progresses and include:

- Pain in the upper right abdomen
- Jaundice
- Enlarged spleen and/or liver
- Night sweats, chills
- Malabsorption of fat leading to fatty stools
- Weight loss

Treatments

The known cure is liver transplant. It can be treated by managing the symptoms using drugs such as antibiotics, opioid antagonists, bile acid sequestrants and ursodeoxycholic acid (UDCA).

Medication

Name of Journal: *World Journal of Gastrointestinal Oncology*

Manuscript NO: 66486

Manuscript Type: REVIEW

Review of incidence and outcomes of treatment of cholangiocarcinoma in patients with primary sclerosing cholangitis

Saffioti F *et al*. Cholangiocarcinoma in primary sclerosing cholangitis

Francesca Saffioti, Vasileios K Mavroeidis

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Cholangiocarcinoma (CCA) is the most common malignancy **in patients with primary sclerosing cholangitis** (PSC) and carries a high rate of mortality. Although the pathogenesis of CCA in PSC is largely unknown, inflammation-driven carcinogenesis concomitant with various genetic and epigenetic abnormalities are underlying factors.

Cited by: 16 **Author:** Junmin Song, Junmin Song, Yang Li, Chri...
Publish Year: 2020

[PDF] Cholangiocarcinoma - The Lancet

https://www.thelancet.com/pdfs/journals/lancet/PIIS0140-6736(13)61903-0.pdf

The lifetime **incidence of cholangiocarcinoma** in this patient population ranges between 5% and 10%.23–26 About 50% of **patients with primary sclerosing cholangitis** who develop **cholangiocarcinoma** are diagnosed with **cholangio-carcinoma** within 24 months of diagnosis of **primary sclerosing cholangitis**.23,27 The risk of cholangio-

People also ask

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How old do you have to be to get cholangiocarcinoma?

What kind of cancer is mixed hepatocellular cholangiocarcinoma?

Feedback

Primary sclerosing cholangitis and cholangiocarcinoma