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Manuscript Type: CASE REPORT

Peutz-Jeghers syndrome with mesenteric fibromatosis: A case report and review of literature

Huai-Jie Cai, Han Wang, Nan Cao, Wei Wang, Xi-Xi Sun, Bin Huang

Abstract

BACKGROUND

Peutz-Jeghers syndrome (PJS) and mesenteric fibromatosis (MF) are rare

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Peutz-Jeghers syndrome with mesenteric fibromatosis: A case

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Peutz-Jeghers syndrome (PJS) is a rare **familial disorder**, with an incidence of 1 in 12-30,000 live births . It is an autosomal dominant condition with incomplete penetrance [2]. Nonsense, **frameshift** and missense mutations inactivating the LKB1 gene on chromosome 19p13.3 have been implicated as the underlying abnormality [3].

Cited by: 8 **Author:** Sayantan Bhattacharya, Sunondo R Mah...
Publish Year: 2010

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Published in: Journal of Coloproctology · 2012
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https://jmedicalcasereports.biomedcentral.com/articles/10.1186/1752-1947-4-44

Feb 08, 2010 · **Peutz-Jeghers syndrome** (PJS) is a rare **familial disorder** characterised by mucocutaneous pigmentation, gastrointestinal and extragastrointestinal hamartomatous polyps and an increased risk of **malignancy**. **Peutz-Jeghers polyps** in the **bowel** may result in intussusception.

Cited by: 8 **Author:** Sayantan Bhattacharya, Sunondo R Mah...

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Case Presentation. We **report** the **case** of a 24-year-old Caucasian male who presented with **melaena**. Pigmentation of the **buccal mucosa** was noted but he was **pain-free and** examination of the abdomen was unremarkable. **Upper gastrointestinal endoscopy** revealed multiple polyps.

Cited by: 8 **Author:** Sayantan Bhattacharya, Sunondo R Mah...
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Peutz-Jeghers **syndrome** with intermittent upper intestinal obstruction: **A case report and review** of the **literature**. Duan SX(1), Wang GH, Zhong J, Ou WH, Fu MX, Wang FS, Ma SH, Li JH.

Cited by: 2 **Author:** Shou-Xing Duan, Guang-Huan Wang, Jun...
Publish Year: 2017

Peutz-Jeghers syndrome: Case report and literature review

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[PDF] Adult Intussusception in Patients with Peutz-Jeghers ...

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Peutz–Jeghers syndrome

Autosomal Dominant Genetic Disorder

Peutz–Jeghers syndrome is an autosomal dominant genetic disorder characterized by the development of benign hamartomatous polyps in the gastrointestinal tract and hyperpigmented macules on the lips and oral mucosa. This syndrome can be classed as one of various hereditary intestinal polyposis syndromes and one of various hamartomatous polyposis syndromes. It has an incidence of approximately 1 in 25,000 to 300,000 births.

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