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Case Report Pseudomyxoma Peritonei Originating from an Intestinal Duplication Julie Lemahieu, 1 André D Hoore, 2 Stijn Deloose, 3 Raf Sciôt, 1 and Philippe Moerman 1 Department of Imaging and Pathology, University Hospitals Leuven, Minderbroedersstraat, Leuven, Belgium Department of Surgery, University Hospitals Leuven, Belgium

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Name of Journal: *World Journal of Clinical Cases*

Manuscript NO: 61664

Manuscript Type: CASE REPORT

A rare case report of pseudomyxoma peritonei originated from intestinal duplication

pseudomyxoma peritonei derived from intestinal duplication

Abstract

BACKGROUND

Pseudomyxoma peritonei (PMP) is a rare mucinous neoplasm with a relatively low incidence of 1 to 2 per million individuals. It is typically characterized by a type of gelatinous ascites named as "jelly belly". Most cases of PMP occur in association with ruptured mucinous tumors of primary tumors of the appendix (90%). Periodically, PMP can originate from mucinous carcinomas at other sites, including colorectum, gallbladder and pancreas. However, unusual originations can occur as

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