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Rupture of small cystic pancreatic neuroendocrine tumor with many microtumors

Ryota Sagami, Hidefumi Nishikiori, Shoichiro Ikuyama, Kazunari Murakami

Abstract

Pancreatic neuroendocrine tumors (pNETs) are particularly rare. The various forms of PNETs, such as cystic degeneration, make differentiation from other similar pancreatic lesions difficult. We can detect small lesions by endoscopic ultrasound (EUS) and obtain preoperative pathological diagnosis by EUS-guided fine needle aspiration (FNA).We describe ~~a~~ an interesting case of pNET in a 42-year-old woman with no family history. Computed tomography

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