

# Solid-cystic pseudopapillary neoplasm of pancreas: An increasingly diagnosed entity

Sir,

Solid-cystic pseudopapillary neoplasm (SPN) of the pancreas is a rare neoplasm, accounting for about 0.17% and 2.7% of all nonendocrine tumors of the pancreas.<sup>[1]</sup> It is most commonly seen in young women.<sup>[1,2]</sup>

Recently, SPN is getting more frequently diagnosed in the Indian subcontinent as there is better diagnostic preevaluation and increased awareness about this uncommon tumor.<sup>[3,4]</sup> Although it is predominantly seen in females, occasionally it can be seen in males.<sup>[1,3]</sup> Ultrasound-guided fine-needle aspiration (FNA) is useful in providing preoperative diagnosis.<sup>[1]</sup> Whenever possible, trucut biopsy of the lesion should be done which can provide a histopathological diagnosis. Histopathologically, the main differential diagnoses of SPN are well-differentiated neuroendocrine neoplasm and acinar cell carcinomas.<sup>[2,3]</sup> A panel of immunohistochemical markers along with clinical, imageological, and morphological findings may help in arriving at a conclusive diagnosis. The typical paranuclear positivity of CD99 aids in the diagnosis of SPN.<sup>[3,5]</sup>

Although SPN is usually benign, malignant SPNs are known and metastasis also is described.<sup>[2]</sup> Complete surgical resection is associated with long-term survival even in the presence of metastasis, thereby making preoperative diagnosis important.<sup>[2]</sup>

To conclude, although SPN of the pancreas is a rare neoplasm, awareness and clinical suspicion about this uncommon are required to consider this tumor especially in young females. Ultrasound-guided FNA and/or trucut biopsy can help in arriving at preoperative diagnosis. Most of the SPNs are benign, and complete surgical resection remains the treatment of choice.

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## Conflicts of interest

There are no conflicts of interest.

**Rashmi Patnayak, Venkatarami Reddy<sup>1</sup>,  
Amitabh Jena<sup>2</sup>, Thota Asha**

Departments of Pathology, <sup>1</sup>Surgical Gastroenterology and <sup>2</sup>Surgical Oncology, Sri Venkateswara Institute of Medical Sciences, Tirupati, Andhra Pradesh, India

**Correspondence to:** Dr. Amitabh Jena,  
Department of Surgical Oncology, Sri Venkateswara Institute of Medical Sciences, Tirupati - 517 507, Andhra Pradesh, India.  
E-mail: dramitabh2004@yahoo.co.in

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