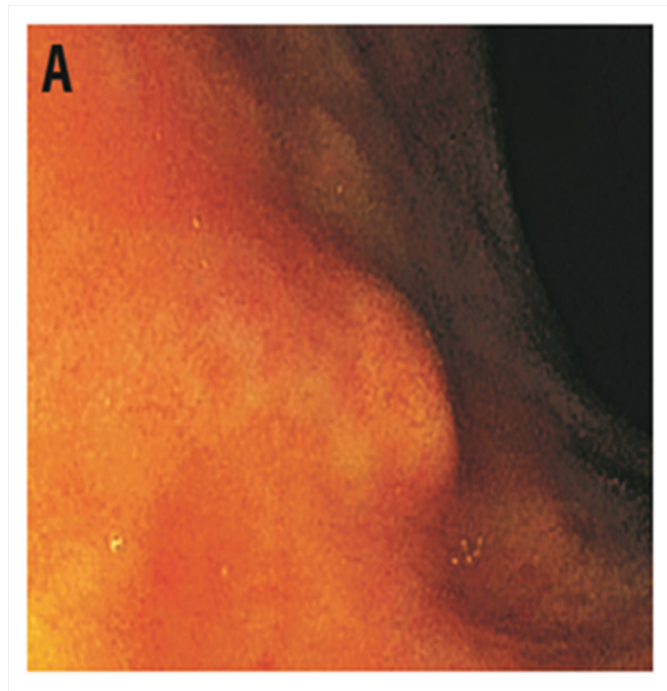


# Rectal carcinoid tumor in adolescent boy

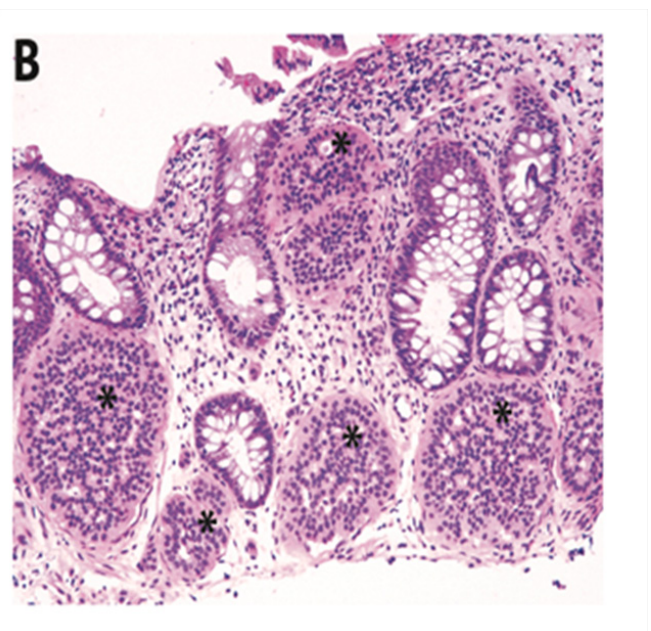
## Case report

The patient is a 17 year old male, who was diagnosed with ulcerative colitis 10 years ago. His treatment was balsalazide. A surveillance colonoscopy was normal except for a small 0.2x 0.2 cms nodular lesion noticed at 15 cms from the anal verge Figure 1(A&B). The nodular lesion was removed by cold biopsy forceps. Biopsies of the terminal ileum were normal. Colonic histology showed chronic minimally active colitis. Histology of the nodular lesion confirmed the diagnosis of rectal carcinoid tumor. Magnetic resonance (MR) enterography and ultrasonography of liver and pancreas were normal. The patient subsequently had an endoscopic ultrasonography with Doppler which did not show any submucosal lesion or rectal wall thickening. There was no indication of residual carcinoid tumor.

Neuroendocrine tumors are rare. The incidence is 2.8 cases per million under the age of 30 years. Carcinoid is the most common neuroendocrine tumor of the GI tract.<sup>1</sup> Tumor size larger than 10 mm, presence of central depression, increasing depth of tumor invasion, lymphatic and venous invasion were significantly associated with a higher incidence of lymph node metastasis.<sup>2</sup> Five year survival rate has been reported to range from 54% to 73% for lymph node positive rectal carcinoid disease even without distant metastases.<sup>3</sup> Most patients are asymptomatic but rectal carcinoids can present with rectal bleeding, abdominal pain, diarrhea and flushing of the skin. Rectal carcinoids are essentially chemo resistant.<sup>4</sup> Surgical resection of metastatic lesions improves survival.<sup>5</sup> Preoperative assessment with EUS for rectal carcinoids is effective to determine depth of invasion.<sup>6</sup>



**Figure 1A** Gross appearance of rectal carcinoid tumor: Colonoscopic appearance of carcinoid showing nodular lesion in rectum.



**Figure 1B** Histology of rectal mucosa showing scattered multiple nests of uniform round cells arranged in lace-like pseudorosetting pattern in lamina propria. H&E; original magnification X100.

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## Conflicts of interest

The author declares, there is no conflict of interest.

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## References

1. Navalkhele P, O Dorisio M, O Dorisio TM. Neuroendocrine tumors in children and young adults incidence, survival, and prevalence in the United States. *Pancreas*. 2010;29:278.
2. Kasuga A, Chino A, Uragami N, et al. Treatment strategy for rectal carcinoids: a clinicopathological analysis of 229 cases at a single cancer institution. *J Gastroenterol Hepatol*. 2012;27(12):1801–1807.
3. Konishi T, Watanabe T, Kishimoto J, et al. Prognosis and risk factors of metastasis in colorectal carcinoids: results of a nationwide registry over 15 years. *Gut*. 2007;56(6):863–868.
4. Bajetta E, Procopio G, Ferrari L, et al. Update on the treatment of neuroendocrine tumors. *Expert Rev. Anticancer Ther*. 2003;3(5):631–642.
5. Niederhuber JE, Fojo T. Treatment of metastatic disease in patients with neuroendocrine tumors. *Surg Oncol Clin N Am*. 2006;15(3):511–533.
6. Ishii N, Horiki N, Itoh T, et al. Endoscopic submucosal dissection and preoperative assessment with endoscopic ultrasonography for the treatment of rectal carcinoid tumors. *Surg Endosc*. 2009;24(6):1413–1419.