An unusual case of upper gastrointestinal bleeding in an adolescent female

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Short communication

A 12-year-old female patient presented in paediatric emergency with history of upper gastrointestinal bleeding for 2 days with severe anaemia and associated mucocutaneous pigmentation. Two units of blood transfusion was given to the patient. Upper GI Endoscopy revealed well defined polyps in pyloric antrum, larger one prolapsing into pyloric antrum from duodenal bulb suggestive of Peutz-Jeghers Hamartomatous polyp (Figure 1). On single contrast barium meal examination a small well defined lobulated filling defect is found in pyloric antrum of stomach. Another large filling defect is seen in duodenal bulb causing expansion of it and protruding into antropyloric region (Figure 2).

Peutz-Jeghers syndrome is a rare, AD disease linked to a mutation of the STK 11 gene which is characterized by the development of benign hamartomatous polyps in the gastrointestinal tract in association with a hyperpigmentation on the lips and oral mucosa. These polyps are also at risk of acute gastrointestinal bleeding, intussusception and bowel obstruction.1 In PJS polyps can be located anywhere in the gastrointestinal tract, most commonly seen in the small bowel (70-90%), the colon (50%) and the stomach (25%).2 Solitary Peutz-Jeghers polyps are generally have their origin in the small bowel, duodenum, colon, rectum and are extremely rare in the stomach, with only 8 well documented cases being reported to date.3 PJS also known as periorificial lentiginosis with an incidence of 1/200,000 live-borns. The most frequent complication at young age is recurrent intussusception due to multiple hamartomatous polyps, usually in the small intestine.4

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Conflict of interest

Author declares that there is no conflict of interest.

References

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