

## Solitary Juvenile Polyp at a Rural Ugandan Hospital Presenting with Recurrent Rectal Bleeding

Gideon K. Kurigamba<sup>1\*</sup>, Vivian V. Akello<sup>1</sup>, Asaph Owamukama<sup>2</sup>, Irene Nanyanga<sup>3</sup>

<sup>1</sup>Department of Surgery, Uganda Christian University, Kampala, Uganda; <sup>2</sup>Department of Surgery, Bwindi Community Hospital, Buhoma, Uganda; <sup>3</sup>Department of Pediatrics and Child Health, Makerere University, Kampala, Uganda

### ABSTRACT

Juvenile Polyps (JP) are rare but important causes of acute gastrointestinal symptoms in children. They are a recognized cause of painless rectal bleeding in preschool age children and also the most common intraluminal disorder of the colon in children. They are often solitary, pedunculated and small in size but may occasionally grow to large sizes or occur in great numbers, as in juvenile polyposis syndrome. Histologically juvenile polyps are similar to inflammatory polyps with irregular dilated glands, lamina propria expansion and granulation tissue expansion. Sporadic juvenile polyps of the colon occur in up to 2 percent of children under the age of 10 years, are usually solitary, and are not associated with an increased cancer risk.

**Keywords:** Juvenile polyps; Rectal bleeding; Polypectomy; Uganda

### DESCRIPTION

The etiology, diagnosis, clinical presentation, and management of these intestinal polyps depend on the type of polyp or polyposis syndrome. A change in bowel habits, abdominal pain, rectal bleeding, rectal prolapse, and even intussusception may be the initial presentation in children. In addition to a careful history, including a detailed family history, a physical examination, contrast studies, and endoscopic examination are vital diagnostic tools. Juvenile polyps may also present with prolapse of the polyp from the anus, abdominal pain due to intussusception or may even be asymptomatic. All such polyps should be removed by colonoscopy or transanal resection [1]. To share the experience of Juvenile polyposis, at Bwindi community hospital Uganda. These are the children second degree relatives, who presented with recurrent rectal bleeding for 4 months after every passage of stool, with prolapse of a mass and were managed at Bwindi community hospital. These patients were all admitted for a total of two days in Hospital. Average hemoglobin was 11.5 mg/dl. Both had a pre-operative colonoscopy that revealed a solitary polyp in the rectal canal. They both underwent Examination Under Anesthesia (EUA) and transanal polypectomy. Juvenile Polyps (JP) is rare, but a key cause of rectal bleeding in children and in low resource settings, transanal polypectomy remains the treatment of choice.

A 3-year-old girl was admitted to our surgical department with an on and off prolapsing anal mass for 4 months associated with rectal bleeding. There was no history of diarrhea, vomiting, constipation, abdominal distention, fevers or weight loss. Patient family history was negative for colorectal polyps or colorectal cancers. The Current weight was 15 kg and height of 34 inches [2]. Abdominal examination was normal. Rectal exam revealed a pedunculated smooth round mass in the rectum at 12 O'clock with some visible fresh blood on the anal verge originating approximately 5 cm from the anal verge. The Colonoscopy done revealed a 2 cm pedunculated polyp in the rectum while the rest of the large bowel and the terminal ileum were normal. Hemoglobin level was 12.1 mg/dl, HIV test negative. Transanal polypectomy was done; the child recovered uneventfully and was discharged at day 1 post-operative with no recurrent episode of rectal bleeding at follow up. Histology of the resected polyp was compatible with juvenile polyp with no signs of dysplasia.

A 9-year-old boy presented to the surgical OPD with recurrent pinkish prolapsing anal mass on passage of stool since 2 years of age and occasional blood stained well-formed stools. He had no history of vomiting, constipation, abdominal distention, fevers or weight loss. There was no history of a similar illness in the first degree relatives. However his cousin was recently treated at our facility with similar symptoms for an anal mass. EUA plus trans anal polypectomy was done and the patient improved on postoperative analgesia and metronidazole, discharged on day 1

**Correspondence to:** Gideon K. Kurigamba, Department of Surgery, Uganda Christian University, Kampala, Uganda, Tel: 256775581231; E-mail: gideonkurigamba@gmail.com

**Received date:** December 06, 2021; **Accepted date:** December 20, 2021; **Published date:** December 27, 2021

**Citation:** Kurigamba GK, Akello VV, Owamukama A, Nanyanga I (2021) Solitary Juvenile Polyp at a Rural Ugandan Hospital Presenting with Recurrent Rectal Bleeding. J Blood Disord Transfus. 12:490.

**Copyright:** © 2021 Kurigamba GK, et al. This is an open-access article distributed under the terms of the Creative Commons Attribution License, which permits unrestricted use, distribution, and reproduction in any medium, provided the original author and source are credited.

post-operative without any episode of bleeding. Histology of the polyp was compatible with juvenile polyp [3].

Juvenile polyp refers to the type of polyp (hamartomatous, usually limited to the colon), not the patient age. Solitary juvenile polyps are among the most common causes of lower gastrointestinal bleeding in child hood. The rectal bleeding is recurrent in nature, like the case was with our second patient. None of our patients had anemia or stunted growth which are some of the presenting symptoms in these children [4]. Besides anaemia, infants can present with rectal prolapse too. Previous studies found rectal bleeding as the commonest symptom in 97.7% and 78.5% as well as the highest incidence of polyps was between ages 2 and 10 years (85.1%). Both of the patients were all under the age of 10 and rectal bleeding was one of the main symptoms (100%) though none had any preoperative anaemia as evidenced by the CBC. Our two patients presented with per rectal bleeding and prolapse of a mass, but the pre-operative hemoglobin levels done showed no evidence of anemia. Coburn et al. has found that 86.7% juvenile polyps located in the recto sigmoid area [5]. Three percent of cases had a positive family history. Our two cases all had rectal location of the polyps but with no previous noted family history. Colonoscopy was done in these children and no other polyps were not seen. Colonoscopy is necessary in children since up to 60% of juvenile polyps are thought to occur in more proximal locations than the recto sigmoid junction and multiple polyps are associated with an increased risk of malignancy [6].

Both of our patients under went examination under anesthesia and transanal polypectomy. A lone star retractor was necessary for the second older child. In a low resource setting like our study site trans anal polypectomy is a feasible and effective technique especially for polypectomy in low rectal polyps with minimal complications. This is a treatment of choice also when advanced endoscopic skills of polypectomy are not available. Patients with numerous polyps in the colon may benefit from proctocolectomy with ileo-anal anastomosis, which is required if adenomatous polyps with epithelial dysplasia are found. Solitary polyps should removed by colonoscopic polypectomies or through transanal resection. In our patients trans anal resection was done [7].

## CONCLUSION

Laparotomy to perform enterotomies and polypectomies may be needed. If there are clusters of polyps in isolated areas, limited segmental resection may be appropriate. Juvenile Polyps (JP) are

rare, but a key cause of rectal bleeding in children and a high index of suspicion is key in diagnosis.

## RECOMMENDATIONS

Colonoscopic evaluation of children with rectal bleeding is essential in the diagnosis of juvenile polyps. Transanal excision of the polyps in low resource settings is sufficient for the polyps accessible through the anus.

## DISCLOSURE OF INTERESTS

All authors report no potential conflicts of interest.

## CONTRIBUTION TO AUTHORSHIP

All authors participated in the editing of this manuscript.

## DETAILS OF ETHICS APPROVAL

Ethical clearance from the Hospital scientific committee was obtained.

## FUNDING

The study had no external source of finding.

## REFERENCES

1. Nagasaki A, Yamanaka K, Toyohara T, Ohgami H, Aoki T, Sueishi K. Management of colorectal polyps in children. *Acta Paediatr Jpn.* 1993;35(1):32-35.
2. Wu CT, Chen CA, Yang YJ. Characteristics and diagnostic yield of pediatric colonoscopy in Taiwan. *Pediatr Neonatol.* 2015;56(5): 334-338.
3. Wiseman J, Emil S. Minimal access surgical management of large juvenile polyps in children. *J Pediatr Surg.* 2009;44(9):e9-e13.
4. Hagh Ashtiani MT, Monajemzadeh M, Motamed F, Moradi Tabriz H, Mahjoub F, Karamian H, et al. Colorectal polyps: A clinical, endoscopic and pathologic study in iranian children. *Med Princ Pract.* 2009;18(1):53-56.
5. Brosens LAA, van Hattem A, Hylind LM, Iacobuzio-Donahue C, Romans KE, Axilbund J, et al. Risk of colorectal cancer in juvenile polyposis. *Gut.* 2007;56(7):965-967.
6. Coburn MC, Pricolo VE, DeLuca FG, Bland KI. Malignant potential in intestinal juvenile polyposis syndromes. *Ann Surg Oncol.* 1995;2(5):386-391.
7. Lee BG, Shin SH, Lee YA, Wi JH, Lee YJ, Park JH. Juvenile polyp and colonoscopic polypectomy in childhood. *Pediatr Gastroenterol Hepatol Nutr.* 2012;15(4):250-255.