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Case Report

Contribution of the Clinical and Histopathological Features in the Positive Diagnosis of the Juvenile Polyposis Syndrome -

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ABSTRACT

Juvenile Polyposis Syndrome (JPS) is a rare genetic disease characterized by the presence of the juvenile hamartomatous polyps. The condition is caused by germline mutation in the BMPR1A or the SMAD4 genes and it is inherited in an autosomal dominant manner. It predisposes affected persons to a high risk of malignant tumors, mainly colorectal and stomach cancers. The confirmation of the diagnosis is based on genetic analysis. But at first, family history, pedigree, clinical criteria and histopathological analysis guide to an inherited disease. We present here a Congolese patient with suggestive clinical and histopathological features which lead to the JPS.

Keywords: Juvenile polyposis syndrome; Clinic; Histopathology; Genetic Disease; Hereditary Cancer

INTRODUCTION

Juvenile Polyposis Syndrome (JPS) is a rare condition characterized by the presence of the juvenile hamartomatous polyps mostly located in the gastrointestinal tract [1]. It predisposes affected individuals to an increased risk of Colorectal Cancer (CRC) and less frequently to extra gastrointestinal malignancies, especially in adulthood [1-4]. The clinical manifestation of the disease appears after birth, especially in young subjects. The germline mutation occurs in BMPR1A (Bone Morphogenetic Protein Receptor Type-1A) gene mapped on chromosome band 10q23.2 or in SMAD4 (Mothers against decapentaplegic homolog 4) located on chromosome band 18q21.2 [2,5,6]. Both genes are tumor suppressors and are involved in the Transforming Growth Factor-Beta (TGF-β) signaling pathways [4,6]. The challenge in this condition is the rapid recognition of the disease and the detection of the mutation in the index patient and the family members in order to perform radiologic, endoscopic examinations and rapid polypectomy and to prevent partially cancer [6]. But the search of mutation is initially guided by elements in favor of the hereditary character of the affection such as in lynch syndrome (other predisposed CRC affection) [7]. The interrogatory in search of familial history of polyps or cancers, the pedigree, the endoscopic examination and the clinical criteria, are the first guiding elements for JPS positive diagnosis [4]. The histological analysis in research of typical features of JPS also helps the clinicians to recognize the condition [8,9]. We report here a clinical observation whose suggestive endoscopic and clinic histological phenotype in favor of JPS have motivated the research of the mutations in BMPR1A and SMAD4 genes.

PATIENT PRESENTATION

He is a 25-year-old congolese man, single, with a father who died from a CRC (Figure 1). Its disease begins at the age of 15 years-old by recurrent diarrhea, rectal bleeding, persistent abdominal pains, abdominal bloating followed by cachexia and occlusive syndrome. The barium enema and the colonoscopy showed numerous polyps along the colon. The tumor macroscopic examination found approximately more than 50 sessile and pedunculated polyps of different sizes (Figure 2). Histological analysis (17H255A2) exhibited evocative lesions of JPS and a slight dysplasia (Figure 3). Direct sequencing (performed in Human Genetic Center of University hospital of Leuven, Belgium) identified a heterozygous mutation in SMAD4 defined as « c.1229_1230delAG », exon 10 (ENSE00003694477) with the pathogenic variant « p.Ser411Leufs*17 » (of the abnormal predicted protein). We concluded to a juvenile polyposis coli form with a slight dysplasia in young man carrying SMAD4 germline mutation. The patient underwent a surgical cure (partial colectomy). He has been doing well for two years and is followed by a multidisciplinary team.

DISCUSSION

Juvenile polyposis syndrome appears in three clinical forms: juvenile infantile polyposis, generalized juvenile Polyposis and juvenile colorectal polyposis [5]. The approximate annual incidence is 1:16,000 to 1:100,000 in the population [2,5,10,11]. Family history is found in 20 to 50% of the affected patients [1,4,5,9]. Clinical criteria defined by Jass et al. in 1998 and revised in 2000 by World Health Organization (WHO) are used to facilitate the recognition of JPS (Table 1) [3,5,6,11,12]. Diagnosis may be suspected in presence of one criterion. Our patient met the three clinical criteria guidelines (Table 1), and presented the juvenile colonic polyposis form. According to publications data, JPS is manifested by constant presence of juvenile hamartomatous polyps [5,9]. The number of polyps varies from 1 to over 100, and they are most frequently located in the distal colon (such as in our patient) and the rectum [2,4,11]. Others gastrointestinal locations are stomach, duodenum, small intestine [3,4,13]. Polyps can appear at any age (from childhood through adulthood), most often during the adolescence (around 12-14 years-old) [11,13,14]. JPS

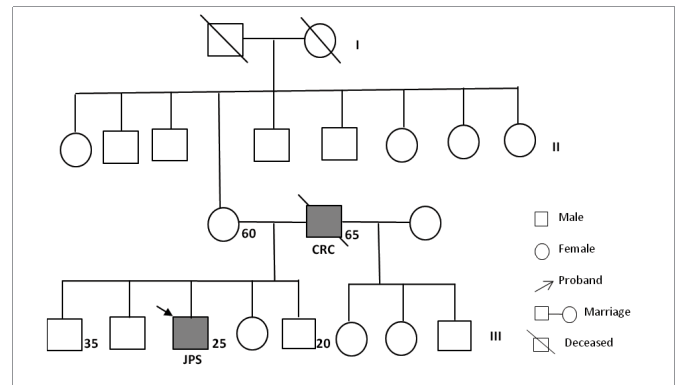


Figure 1: Family pedigree of patient. Proband (arrow) with JPS having a father died for colorectal cancer.



Figure 2: Macroscopic aspect of resected colon tumor Presence of multiples pedunculated and sessile polyps (> 50) of different sizes in the colon.

is the most common hereditary form of polyposis in the childhood, the protein losing-enteropathy and the prolapse of rectal polyps are frequently observed in children (Genetic home reference 2013) [4,6,10,13]. JPS is also accompanied by other nonspecific symptoms among which the rectal bleeding and the recurrent abdominal pains are the most common revealing signs [6,13]. We have in (Table 2) reported the main clinical manifestations [2,14,15]. The histological analysis which is practiced without difficulties in African hospitals is essential in the approach of positive diagnosis. Concerning our patient, the exhibited tumor histological profile (Figure 3) was consistent with literature data [1,8,15]. Typically, the microscopic aspects show the juvenile polyps (Figure 3A) with a characteristic architecture: Glands are more often spherical, separated, dilated, cystic and rich in mucus (Figure 3B, C) [3,8,11]. They are associated with lamina propia edema, massive infiltrating inflammatory cells (Figure 3B, C) [2,4,11]. The stroma is dense with dystrophic blood vessels [1,3,6]. Juvenile hamartomatous polyps are identified by histological examination in 88-100% of cases [13]. Genetically, JPS is a Mendelian pathology [3,6,16-19]. The condition is inherited from one parent (as in our Report) in an autosomal dominant manner (in 75% of patients), except in cases of de novo mutation reported in 25% of JPS (Genetic home reference 2013) [9,14]. BMPR1A and SMAD4 mutations are detected approximately in 40 to 60% of patients [14]. Both genes are TGF-β signaling proteins which initiate cell cycle arrest and growth inhibition [14,16]. Polyps in JPS can degenerate and the approximate risk of developing CRC varies between 38 to 50%. In case of multiple polyps, especially in adults [3,5,11,14]. Hamartomatous polyposis including JPS is responsible approximately for 1% of all CRC [5]. The occurrence cancer risk in the JPS (when the mutation is known in the family) requires the early screening of asymptomatic family members and the polypectomy is highly recommended. Regular endoscopy

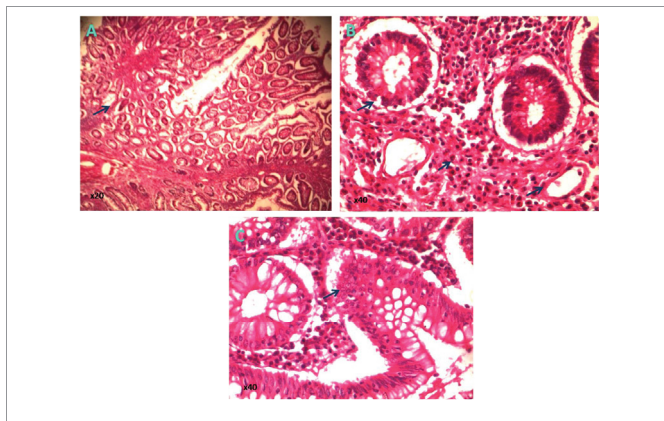


Figure 3: Image of histological section of juvenile polyps (Hematoxylin and Eosin-stained)
 A) Juvenile polyps (examination No 17H255A2) showing increased of cellular elements, lamina propia and proliferative glands
 B) and C) Excess of small, cystical and spherical proliferative glands with abundant mucus. Lamina propia with peri-glandular edema, inflammatory cells and dystrophic blood vessels.

Number	Clinical criteria
1	Multiple juvenile polyps in the colorectum, more than five.
2	Patient with a family history of juvenile polyposis syndrome, regardless the number of juvenile polyps and their site.
3	Multiple juvenile polyps throughout the intestinal tract with exclusion of the other causes of digestive polyposis.

Table 2: Natural history and phenotype features in juvenile polyposis syndrome.

Juvenile polyposis syndrome (JPS)	Discovery and Features	Authors
History	In 1964, first description of clinical features and pathology by McColl et al.	Veale et al. 1966 [15] Cichy et al. 2014 [16]
	In 1998, identification of mutations in SMAD4 gene associated with familial JPS. In 2001, discovery of BMPR1A gene also causative of JPS.	Howe et al. 1998 [17] Houlston et al. 1998 [18] Howe et al. 2001 [19]
Clinic	Rectal bleeding. Recurrent abdominal pains. Diarrhea. Prolapse of rectal polyps. Anemia. Protein losing-enteropathy. Cachexia. Occlusive syndrome. One or over 100 Polyps in GIT. The disease exhibits three clinical forms	Larsen Haidle et al. 2017 [2] Andrade et al. 2015 [13] Veale et al. 1966 [15]
Histopathology	Juvenile hamartomatous polyps	Cauchin et al. 2015 [9] Jelsig et al. 2014 [11]
Genes SMAD4 (18q21.2) BMPR1A(10q23.2)	SMAD4 is a member of SMAD family of signal transduction proteins. BMPR1A belongs to BMP, a family of transmembrane serine-threonine kinases	Cichy et al. 2014 [16] (UCSC-2013GRCh38/hg38)

TGF-B: Transforming Growth Factor-Beta; GIT: Gastrointestinal Tract.

monitoring (every 2 to 3 years) of persons carrying a BMPR1A or a SMAD4 germline mutation is recommended starting at 10 to 15 years-old [3,5,9,16].

CONCLUSION

The combination of endoscopy, pedigree, clinical criteria and histological analyzes is at first a good approach to spot JPS. It allows to guide towards a pathology inherited and especially to signify the presence of polyps (and cancer) for a rapid surgical management.

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AUTHORS' CONTRIBUTIONS

GD: Endoscopic examination; GD, BBL, JFP, MK, LZA: Review and data analysis; HP: Data analysis and manuscript preparation.

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