Henoch-Schönlein Purpura with hematochezia in an adult patient

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ABSTRACT

Henoch-Schönlein purpura (HSP) is a self-limiting vasculitic disease, commonly seen in children. It is characterized by palpable purpura, arthritis, hematuria and abdominal pain. Hematochezia is an uncommon symptom. The patient being reported is an adult male who presented with vomiting, abdominal pain and hematochezia, followed by appearance of palpable purpura and arthritis. He was diagnosed to have HSP. Hematochezia as one of the initial presentations of HSP in adults is an uncommon scenario.

Key words: Henoch-Schönlein purpura; Vasculitis; Hematochezia; Purpura; Arthritis; Leukocytoclastic vasculitis

INTRODUCTION

Henoch-Schönlein purpura (HSP) is a common form of childhood vasculitis, characterized by palpable purpura, arthritis, colicky abdominal pain and nephritis. It is a multi-systemic IgA mediated small vessel vasculitis, with 90% patients being younger than 10 years of age.¹² It is an immune mediated response triggered by an infection or by certain drugs; and occurs more often during spring, winter and autumn seasons.³ The infections that may precede HSP include group A Streptococcus, infectious mononucleosis, hepatitis, Mycoplasma, Campylobacter, Helicobacter pylori, Yersinia, Shigella, Salmonella, Brucella, Legionella, Parvovirus, Adenovirus, Varicella and Rotavirus. Vaccinations against typhoid, measles, yellow fever and cholera may also lead to the development of HSP. Some of the implicated drugs include penicillin, ampicillin, erythromycin, quinine, losartan and catarabine.

Majority of the patients present with palpable purpura. Abdominal symptoms as presenting complaint prior to appearance of purpura are uncommon.⁴ Our patient, an adult, has abdominal symptoms of vomiting, pain and hematochezia, followed by purpura and arthritis.

CASE PRESENTATION

A 20 year old male presented with complaints of severe diffuse abdominal pain, non-projectile vomiting and hematochezia since 2 days. He had consulted a local practitioner who prescribed him oral pantoprazole and dicyclomine. The following day he noticed purplish spots all over his upper and lower limbs, and buttocks. On the day of presentation to our hospital, he had painful swelling of right elbow and ankle. He had mild throat pain about 4 days ago, for which he resorted to warm salt water gargle. There was no history of fever. He had not taken any drugs or vaccinations recently.

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On examination, he was conscious and oriented, with stable vitals. His right elbow and ankle joints were swollen and tender, with local rise in temperature. He had purpuric lesions over his upper (Figure 1) and lower (Figure 2) limbs, and gluteal regions. His abdomen was mildly distended, with no tenderness on palpation. Other systemic examinations were normal. There were no signs of meningeal irritation. His throat examination was also normal. From his history and clinical presentation, the probable diagnosis of HSP was made.

His blood investigations showed leucocytosis (12200 cells/cumm with 74% neutrophils, 20% lymphocytes, eosinophils 6%), elevated erythrocyte sedimentation rate (30 mm/hour) and C-reactive protein (47 mg/L), and hypoalbuminemia (2.8 mg/dL). Antistreptolysin O (ASO) titers were elevated (440). His renal and liver functions, C3 and C4 complement levels and electrolytes were normal. Anti-nuclear antibody and anti-double stranded DNA were negative. Urine microscopy did not show any haematuria. Epstein-Barr viral capsid antigen IgM and IgG, cytomegalovirus IgM and IgG, HIV, hepatitis B virus surface antigen, anti-hepatitis C virus were negative. Blood cultures were sterile. Ultrasound abdomen showed mild to moderate ascites. Contrast enhanced computerized tomography (CT) of the abdomen revealed long segment symmetrical circumferential continuous wall thickening in the ileal loops with moderate luminal narrowing, and mild to moderate ascites (Figure 3). Multiple erosions were seen in the rectum, sigmoid, descending, transverse and ascending colon on colonoscopy (Figure 4). Upper gastrointestinal (GI) endoscopy was normal. Histopathologic examination of skin biopsy from the pretilial region showed leukocytoclastic vasculitis, and that from the ileal and colonic mucosa showed edema with mild lymphoplasmacytic infiltration and lymphoid aggregates. Echocardiography and chest Xray were also normal.

He was given pulse doses of methylprednisolone (20 mg/kg/day) for 3 days, and was kept nil per oral (NPO) with adequate hydration and intravenous pantoprazole (40 mg twice daily). By day 2 of admission, his arthritis disappeared and purpuric lesions started subsiding. He was started on soft oral diet but had vomiting and abdominal
pain. There were no further episodes of hematochezia. He was again kept NPO with adequate hydration. Intravenous methylprednisolone was continued at reduced dosage (0.8 mg/kg/day). By day 6 of admission, he was asymptomatic. Oral diet was started again, which he tolerated. He was discharged on day 7 on oral prednisolone (1 mg/kg/day), which was tapered and stopped over 1 month. He was also advised strict rest from physical activities for 10 days. He was reviewed every week for 1 month and then monthly for next 3 months; and continued to be asymptomatic.

**DISCUSSION**

HSP was first reported by Wilan in 1808. Eduard Heinrich Henoch found a coexistence between purpura and GI haemorrhage in 1874. Johannes Lucas Schonlein, in 1837, reported an association between purpura and arthritis. HSP is less common among adults, and have more of renal involvement. The diagnosis of HSP requires the presence of at least 2 of the following (87.1% sensitivity and 87.7% specificity):  
- Palpable purpura  
- Age < 20 years at the onset of symptoms  
- Bowel angina  
- Biopsy showing leukocytoclastic vasculitis.

The GI symptoms are seen in 10 to 20% of cases. These include colicky abdominal pain (86%), occult blood loss (66%), vomiting (40%), diarrhoea (20%) and massive colorectal bleeding (20%). These symptoms are caused by vasculitis resulting in subserosal and submucosal haemorrhages and the accumulation of fluid in the bowel wall. The duodenum (2nd portion), stomach and colon are the commonly involved areas on endoscopic evaluation. The GI lesions are usually irregular ulcers or nodules or hematoma-like protrusions. The CT scan findings include intramural gas in the bowel wall along with decreased or absent bowel wall enhancements, intestinal wall thickening, dilatation of lumen and the presence of intra-abdominal fluid. Ultrasound scan of the abdomen us used to identify other abdominal pathologies like intussusception and bowel perforation, and abdominal Xray is useful to detect perforation. Arteriography is the gold standard investigation for intestinal ischemia.

The use of corticosteroids in the treatment of HSP is controversial. Positive outcomes have been reported in cases with GI and renal involvement. In a randomized trial, oral prednisone was found to be effective in reducing abdominal pain. Persistent severe abdominal pain even after initiating corticosteroid therapy is uncommon and should raise the probability of GI complications like bowel ischemia, intussusception and perforation. Intravenous immunoglobulin (IVIG) is an alternative for treatment of steroid-resistant cases. Other treatment options for refractory GI cases include plasma exchange and immunosuppressive drugs like azathioprine, cyclophosphamide and cyclosporine. Analgesics like acetaminophen, ibuprofen and naproxen, are useful for joint pain and fever. Ranitidine, an H2 blocker, may be used in patients with moderate GI involvement. Patients are also advised rest and adequate hydration. Resting with the legs raised may help reduce the development of rash. This is because the petechiae and purpuric lesions tend to develop in dependent areas of the body such as the legs. On exercising the temperature regulation mechanisms within the calf muscles break down, leading to reduced venous return and blood stasis. The end result is inflammation and vascular injury. Moreover, rest is also important for relief of joint pain.

Our patient was an adult, who presented with initial complaints of abdominal pain, vomiting and hematochezia. He later developed purpuric lesions and arthritis. His ASO titers were elevated. His skin biopsy showed leukocytoclastic vasculitis. His abdominal CT was suggestive of HSP and colonic erosions were seen on colonoscopy. He did not have any renal complications. He was treated with corticosteroids, and improved.

**CONCLUSION**

HSP is generally a self-limited, multi-systemic IgA mediated small vessel vasculitic disease, commonly affecting children. Adults usually face renal complications. Hematochezia, as one of the initial presentations, is an uncommon. Early diagnosis and treatment of HSP can prevent surgical intervention for GI complications like bowel ischemia, intussusception and perforation.

**REFERENCES**

https://doi.org/10.1016/j.cgh.2010.02.011


https://doi.org/10.1002/art.1780400513

https://doi.org/10.1016/S0022-3468(05)80009-0


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