The Characteristics of Video Capsule Endoscopy in Pediatric Henoch–Schönlein Purpura with Gastrointestinal Symptoms

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Abstract

**Background** Henoch–Schönlein purpura (HSP) is a systemic small-vessel vasculitis that commonly affects gastrointestinal tract. The video capsule endoscopy (VCE) characteristics of pediatric HSP patients were rarely investigated.

**Methods** Patients diagnosed with HSP by VCE examination at our hospital from February 2010 to January 2019 are analyzed. The clinical features, laboratory findings, and the characteristics of VCE findings are studied.

**Results** There are 30 patients enrolled in this investigation from February 2010 to January 2020. The mean age of these patients is 96.9±35.8 months, and the most frequent finding of VCE is mucosal erosion, which account for 69 % of the patients, and followed by mucosal erythema or petechia accounted for 79.3 % of the patients. Regarding to the disease location detected by endoscopy, jejunum is the most common involved part of the gastrointestinal tract in pediatric HSP patients. All the patients had jejunum involved except in one patient the VCE did not pass through the pylorus. One third of the patients involved the descent part of duodenum. No side effects are observed in this study.

**Conclusions** VCE is safe and effective in the diagnosis of gastrointestinal involved HSP patients with or without typical skin purpura. Jejunum is the most common involve location in gastrointestinal.

**Background**

Henoch–Schönlein purpura (HSP) is a systemic small-vessel vasculitis that often affects children and occasionally affect adults. It is relatively common in the Asian population than the Western population. HSP mainly involves the small vessels of skin, joints, gastrointestinal (GI) tract, and kidney. About 50% to 85% of the HSP patients have gastrointestinal symptoms [1]. GI symptoms include acute abdominal pain, nausea, vomiting, hematochezia or melena, and diarrhea. But endoscopic evaluations are not performed in all HSP patients with GI symptoms. Among the patients with GI symptoms, there is a rare portion of patients without skin manifestations, which need endoscopy evaluation for diagnosis and differential diagnosis. HSP most affects the descent part of the duodenum in adult patients, and the typical indication of endoscopy of HSP are erythema,
petechia, erosion and ulceration [1, 2]. Although esophagogastrroduodenoscopy (EGD) is useful in the
diagnosis of majority HSP patients affect GI tract, there is still a portion of patients who could not be
diagnosed due to the location of disease that could not be reached by EGD which need video capsule
endoscopy (VCE) examination.
There are several literatures revealed the traits of endoscopy findings of adult HSP patients. However,
there is no report focuses on the diagnosis value of VCE in pediatric HSP patients. Here we report a
cohort of pediatric HSP patients who performed VCE examination. All the patients are diagnosed with
HSP and had an atypical manifestation of HSP or with severe GI symptoms. Here, we summarize the
clinical and VCE features of these patients.
Materials And Methods
We enrolled in 30 cases of HSP patients who performed VCE examination at Children’s Hospital,
Zhejiang University School of Medicine from February 2020 to January 2019. The clinical features,
laboratory findings, and the findings of VCE were reviewed. This study was approved by the Ethics
Committee in Children’s hospital, Zhejiang University School of Medicine.
Diagnosis of HSP
The diagnosis of HSP based on typical clinical manifestations of the disease: palpable purpura without
thrombocytopenia or coagulopathy, and two or three of the remaining clinical features: arthritis or
arthralgia, abdominal pain, and renal disease.
Indications for VEC examination and device
The indications for VEC examination were as following: 1. Patients who were suspected HSP without
typical skin palpable purpura and EGD did not reveal characteristic findings of HSP. 2. HSP patients
with GI symptoms but were not wholly response to steroids treatment or dependent on steroids. 3.
Patients who were suspected HSP but needed to be differentiated with other small intestinal diseases.
Patients with massive intestinal bleeding could not tolerate VCE examination or with acute surgical
indications were excluded. The device used for VCE was OMOM (Chongqing, China). The capsule was
either swallowed by the patient or was delivered by EGD in patients who cannot swallow the capsule.
Statistical analysis
The continuous variables with normal distribution were presented as mean ± SD, otherwise were presented with median ± interquartile range (IQR); and, discontinuous variables were presented as number or percentage. The statistical analyses were conducted with SPSS 22.0 statistical software (SPSS Inc., IBM Corp., Armonk, NY, United States).

Results
The geography features of the patients
There were 31 patients performed VCE examination and was suspected HSP. One patient who had recurrent abdominal pain for three years and did not have any other typical symptoms of HSP was excluded. There were 30 patients enrolled in this study from February 2010 to January 2020. The geography information of these patients was showed in Table 1. All the patients had GI symptoms, including abdominal pain, vomiting, and intestinal bleeding. The symptoms of these patients were also shown in Table 1. The complications included hypertension, appendicitis, acute pancreatitis and acute intestinal perfusion. A patient with intestinal perfusion had surgery. Five patients had a history of HSP.

| Items                                | Results                        |
|--------------------------------------|--------------------------------|
| Age (M)                              | 96.9 ± 35.8                    |
| Male/Female                          | 19/11                          |
| Duration of disease admitted (D)     | 12.5 (IQR: 6.5–20.0)           |
| Symptoms                             |                                |
| Purpura, n (%)                       | 13 (43.3)                      |
| Abdominal pain, n (%)                | 30 (100)                       |
| Vomiting, n (%)                      | 9 (30)                         |
| Intestinal bleeding, n (%)           | 8 (26.7)                       |
| Arthralgia, n (%)                    | 4 (13.3)                       |
| Hematuresis/proteinuria, n (%)       | 3 (10.0)                       |
| WBC (× 10E9/L)                       | 17.4 ± 7.1                     |
| Hb (g/L)                             | 128.4 ± 11.7                   |
| PLT (× 10E12/L)                      | 372.3 ± 111.2                  |
| CRP (mg/L)                           | 5.5 (IQR: 0.8–20.8)            |
| ESR (mm/h, n = 24)                   | 8 (IQR: 6.0–20.8)              |
| Serum albumin (g/L, n = 23)          | 36.4 ± 6.8                     |
| Plasma D-dimers (mg/L, n = 22)       | 2.2 (IQR: 0.5–8.4)             |

HSP: Henoch–Schönlein purpura; WBC: white blood cell; Hb: hemoglobin; PLT: platelet; CRP: C-reactive protein; ESR: erythrocyte sedimentation rate.

The Laboratory Findings Of The Patients
The laboratory results of these HSP patients were listed in Table 1. Mean count of white blood cells and serum plasm D-dimers levels were elevated, and media CRP and ESR levels were normal.

Images Of Hsp Patients
Fourteen patients had magnetic resonance hydrography (MRE) or computed tomography (CT) scan of abdominal, and ten patients revealed thickening of the small bowl.
The Features Of Endoscopy

All the patients had an EGD examination. Among them, 27 patients had first EGD in our hospital, three patients had EGD in other hospitals before admission, and one patient had second EGD in our hospital. No typical traits of HSP were detected in the EGD examination of 19 patients. One patient revealed duodenum ulcers by first EGD examination at the acute stage of diseases, while the second EGD examination in our hospital five months later was normal. The most frequent findings of EGD were mucosal ecchymosis, petechiae, erosion, and multiple ulcers. EGD revealed typical traits of HSP in the decent part of the duodenum in nine patients. Two patients had the whole stomach involved, and one also had the lower part of the esophagus involved. Thirteen patients performed a colonoscopy, and two patients have detected ulcers in the terminal ileum, and one patient has detected a polyp in the colon, which considered as comorbidity.

Thirty patients had a VCE examination. The Capsule did not pass through pylorus in one patient, and the others all went through the whole small bowel. The mean time of the VCE examination was 21.0 days (IQR: 13.8 to 36.0) after the initial symptoms of HSP appeared. VCE detected multiple mucosal ecchymosis, erosion, and irregular superficial ulcers, which resembled the findings of EGD in 27 patients. Moreover, some patients with massive intestinal bleeding tend to have diffuse erosion and large areas of ulcers (Fig. 1). The numbers and percentage of different lesions identified by VCE were listed in Table 2. One male patient had massive intestinal bleeding, and intestinal perfusion was treated with surgery and followed with oral methotrexate (MTX). He had a VCE examination to assess the recovery of intestinal lesion eight months after the onset of disease onset, and the VCE only detected mucosal congestion in the jejunum. The disease location of patients detected by endoscopy was shown in Table 3. The most frequently involved disease location in this cohort was jejunum, which account for 96.7% of the patients, and followed by the descent part of the duodenum which was accounts for 33.3%. None of these patients affected colon. There was no side effect observed in this study.
Table 2
The characteristics of VCE at small intestinal in pediatric HSP patients

| VCE findings          | Number (Percentage) |
|-----------------------|---------------------|
| Edema/Congestion      | 12 (41.4%)          |
| Mucosal erythema or petechia | 20 (69.0%)   |
| Mucosal erosion       | 23 (79.3%)          |
| Multiple ulcers       | 17 (58.6%)          |

VCE: Video capsule endoscopy; HSP: Henoch–Schönlein purpura

Table 3
The disease location detected by video capsule endoscopy.

| Disease location       | Number (Percentage) |
|------------------------|---------------------|
| Esophagus              | 0 (0.0%)            |
| Stomach                | 3 (10.0%)           |
| Duodenum bulb          | 6 (20.0%)           |
| Descent of duodenum    | 10 (33.3%)          |
| Jejunum                | 29 (96.7%)          |
| Ileum                  | 13 (44.3%)          |
| Total                  | 30                  |

The EGD, VCE, and colonoscopy findings of HSP patients with or without skin purpura were shown in Table 4.

Table 4
Number of patients detected typical lesions of HSP in patients with or without skin purpura by EGD, colonoscopy and VCE.

|                      | With skin purpura N = 13 | Without skin purpura N = 17 | Total |
|----------------------|--------------------------|-----------------------------|-------|
| EGD                  | 8                        | 4                           | 12    |
| Colonoscopy          | 0a                       | 2                           | 2     |
| VCE                  | 12b                      | 17                          | 29    |

HSP: Henoch–Schönlein purpura; EGD: Esophagogastroduodenoscopy; VCE: Video capsule endoscopy.
a. Colonoscopy detected polyp by colonoscopy in one patient is not considered related to HSP; b. VCE did not pass the gastric pylorus in one patient.

Treatment

90% of the patients were initially treated by steroids, and another three patients received PPI or montelukast for unremarkable gastrointestinal symptoms. Four patients were treated by immunoglobulin combined with steroids, and seven patients were treated by immunosuppressants, because they were not completely response to steroids or dependent on steroids.

Immunosuppressants including vincristine, cyclophosphamide, and methotrexate. One patient with intestinal perfusion had surgery and then treated with methotrexate for two months.

Discussion

VCE was approved to be used in pediatrics over two years of age by the US FDA in 2009 [3]. The most frequent indications for VCE in children is inflammatory bowel disease (IBD), obscure gastrointestinal bleeding (OGIB), malabsorption, protein-losing enteropathies, abdominal pain, small bowel polyps and tumors [4]. We have widely applied VCE in the diagnosis and evaluation of Crohn’s disease, abdominal pain, small intestinal bleeding, small intestinal polyps and chronic diarrhea in our hospital. And the application of VCE is rare except retention observed in our experience.
Diagnosis of HSP usually does not need endoscopy examination. However, patients with the following indications need perform endoscopy: 1). Abdominal pain without typical purpura, and could not be explained by other acute abdominal diseases; 2). HSP with massive intestinal bleeding; 3). HSP with recurrent intestinal symptoms or steroid- dependent; 4). Chronic abdominal pain needs to be differentiated from other gastrointestinal diseases, especially Crohn’s disease. Under these circumstances, it is crucial to perform an endoscopy to confirm the diagnosis and understand the extent of the disease. Furthermore, perform endoscopy is also beneficial to determine the proper medical treatment and duration of treatment. It is dependent on the clinical manifestation to perform the EGD, colonoscopy, or VCE.

To our best knowledge, there is no study focused on the VCE examination of pediatric HSP patients. This study is the largest cohort of pediatric HSP patients who have a VCE examination so far. The typical finding of VCE in the small intestinal is similar to the discovery by EGD in other reports from the adult cohort [5], presenting with mucosal edema, congestion, erosion, sporadic purpura or diffuse purpura, and usually with multiple irregular superficial ulcers. In our cohort, the most frequently involved part of the GI tract was jejunum. Almost all the patients had jejunum involved except one did not pass through pylorus. It was different from the reported literature by Eon Jeong Nam et al. [2]. In their report with a series of adult HSP patients, the second part and the terminal ileum were the most frequently involved parts of HSP patients, and colon was frequently involved as well. However, the VCE or small intestinal endoscopy was not performed in this study; thus, the jejunum was not assessed. On the contrary, in our cohort, none of the patients had colon involved. The difference between the studies could partially be explained by the age of patients in the study. The mean age of patients was 96.7 months in our cohort while they were adults in the study of Eon Jeong Nam et al..

And also, the location and extent of disease depends on the time of performing endoscopy and the severity of disease. The media time of our patients had VCE was around three weeks. It is accessible to diagnosis HSP with typical symptom and skin purpura according to the diagnostic criteria [6]. HSP is reported to share similar clinical manifestation and sometimes has a colonoscopy appearance that resembles ulcerative colitis[7]. Application of VCE in these patients could help to
make the diagnosis, thus avoid unnecessary prolong use of steroids and also the side effects of steroids. Although HSP is self-limited, we observed that the inflammation of small intestinal is rather long in our study. The most prolonged patients observed inflammation in the small bowel more than eight months.

MRE and CT were also performed in 14 patients, and 71.4% of them detected thickening of the small intestinal wall. Compared with VCE, the MRE and CT finding of these patients are not typical, and could not identify superficial ulcers of mucosa or evaluate the condition of bleeding.

This study has some limitations. First, this was a retrospective study with a small cohort of patients. The study only included patients diagnosed by VCE as HSP, not all the HSP patients with GI symptoms were evaluated. Second, all of the patients had EGD and VCE in this study. But not all patients had a colonoscopy examination. Thus, the lesions in the colon may be omitted in some patients.

Conclusions
In conclusion, this is the first cohort study focused on the VCE finding of pediatric HSP patients. The typical discovery of endoscopy could make the diagnosis of HSP with or without palpable purpura. VCE is a safe and effective method in the diagnosis of HSP. It is more sensitive compared with MRE or CT scan. VCE could be recognized as the first-line examination for patients suspecting HSP with disease location in the small bowel.

List Of Abbreviations
Henoch–Schönlein purpura: HSP; Gastrointestinal: GI; Esophagogastroduodenoscopy: EGD; Video capsule endoscopy: VCE; Interquartile range: IQR; Magnetic resonance hydrography: MRE; Computed tomography: CT; Inflammatory bowel disease: IBD; Obscure gastrointestinal bleeding: OGIB.

Declarations
Ethics approval and consent to participate:
This study was approved by the Ethics Committee of the Children’s Hospital of Zhejiang University School of Medicine.

Consent for publication:
Written informed consent was obtained from the parents of the patient for the publication.

Availability of data and materials:
All data generated or analysed during this study are included in this published article.
Competing interests:
The authors declare that they have no competing interests.

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Authors' contributions:
YF collected and analyzed clinical and laboratory data of the patients and write the manuscript. HZ and KP interpreted the data of video capsule endoscopy. JC designed and was a major contributor in writing the manuscript. All authors read and approved the final manuscript.

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Figures

![Image of endoscopy showing mucosal edema, congestion, erythema, petechia, diffuse erosion and multiple irregular superficial ulcers at small bowel in pediatric HSP patients](image-url)

**Figure 1**

The video capsule endoscopy showed mucosal edema, congestion, erythema, petechia, diffuse erosion and multiple irregular superficial ulcers at small bowel in pediatric HSP patients.
