

World Journal of *Gastroenterology*

World J Gastroenterol 2017 July 7; 23(25): 4473-4660





EDITORIAL

- 4473 Challenges of modern day transition care in inflammatory bowel disease: From inflammatory bowel disease to biosimilars

Hakizimana A, Ahmed I, Russell R, Wright M, Afzal NA

REVIEW

- 4480 Expression and role of nuclear receptor coregulators in colorectal cancer

Triki M, Lapierre M, Cavailles V, Mokdad-Gargouri R

- 4491 Eubiotic properties of rifaximin: Disruption of the traditional concepts in gut microbiota modulation

Ponziani FR, Zocco MA, D'Aversa F, Pompili M, Gasbarrini A

MINIREVIEWS

- 4500 Localization and role of metabotropic glutamate receptors subtype 5 in the gastrointestinal tract

Ferrigno A, Berardo C, Di Pasqua LG, Siciliano V, Richelmi P, Vairetti M

ORIGINAL ARTICLE

Basic Study

- 4508 Treatment with dimethyl fumarate ameliorates liver ischemia/reperfusion injury

Takasu C, Vaziri ND, Li S, Robles L, Vo K, Takasu M, Pham C, Farzaneh SH, Shimada M, Stamos MJ, Ichii H

- 4517 Enhanced electrogastrography: A realistic way to salvage a promise that was never kept?

Poscente MD, Mintchev MP

- 4529 Glutamine prevents oxidative stress in a model of portal hypertension

Zabot GP, Carvalhal GF, Marroni NP, Licks F, Hartmann RM, da Silva VD, Fillmann HS

- 4538 Hepatitis C virus NS5A region mutation in chronic hepatitis C genotype 1 patients who are non-responders to two or more treatments and its relationship with response to a new treatment

Muñoz de Rueda P, Fuentes Rodríguez JM, Quiles Pérez R, Gila Medina A, Martín Álvarez AB, Casado Ruíz J, Ruíz Extremera Á, Salmerón J

- 4548 Distinct gut microbiota profiles in patients with primary sclerosing cholangitis and ulcerative colitis

Bajer L, Kverka M, Kostovcik M, Macinga P, Dvorak J, Stehlikova Z, Brezina J, Wohl P, Spicak J, Drastich P

- 4559** Anti-inflammatory and anti-apoptotic effects of rosuvastatin by regulation of oxidative stress in a dextran sulfate sodium-induced colitis model

Shin SK, Cho JH, Kim EJ, Kim EK, Park DK, Kwon KA, Chung JW, Kim KO, Kim YJ

- 4569** miR-29a promotes hepatitis B virus replication and expression by targeting SMARCE1 in hepatoma carcinoma

Wu HJ, Zhuo Y, Zhou YC, Wang XW, Wang YP, Si CY, Wang XH

Retrospective Cohort Study

- 4579** Management and outcome of hepatocellular adenoma with massive bleeding at presentation

Klompshouwer AJ, de Man RA, Thomeer MGJ, Ijzermans JNM

Retrospective Study

- 4587** Chronic hepatitis B, nonalcoholic steatohepatitis and physical fitness of military males: CHIEF study

Chen YJ, Chen KW, Shih YL, Su FY, Lin YP, Meng FC, Lin F, Yu YS, Han CL, Wang CH, Lin JW, Hsieh TY, Li YH, Lin GM

- 4595** Comparison of short- and long-term outcomes of laparoscopic vs open resection for gastric gastrointestinal stromal tumors

Ye X, Kang WM, Yu JC, Ma ZQ, Xue ZG

Observational Study

- 4604** Functional lipidomics in patients on home parenteral nutrition: Effect of lipid emulsions

Pironi L, Guidetti M, Verrastro O, Iacona C, Agostini F, Pazzeschi C, Sasdelli AS, Melchiorre M, Ferreri C

- 4615** Cryptogenic multifocal ulcerous stenosing enteritis: Radiologic features and clinical behavior

Hwang J, Kim JS, Kim AY, Lim JS, Kim SH, Kim MJ, Kim MS, Song KD, Woo JY

- 4624** Partners of patients with ulcerative colitis exhibit a biologically relevant dysbiosis in fecal microbial metacommunities

Chen GL, Zhang Y, Wang WY, Ji XL, Meng F, Xu PS, Yang NM, Bo XC

Randomized Controlled Trial

- 4632** Long-term irritable bowel syndrome symptom control with reintroduction of selected FODMAPs

Harvie RM, Chisholm AW, Bisanz JE, Burton JP, Herbison P, Schultz K, Schultz M

EVIDENCE-BASED MEDICINE

- 4644** Anti-apoptotic effect of banhasasim-tang on chronic acid reflux esophagitis

Shin MR, An HJ, Seo BI, Roh SS

SYSTEMATIC REVIEWS

- 4654** Nosocomial spontaneous bacterial peritonitis antibiotic treatment in the era of multi-drug resistance pathogens: A systematic review

Fiore M, Maraolo AE, Gentile I, Borgia G, Leone S, Sansone P, Passavanti MB, Aurilio C, Pace MC

ABOUT COVER

Editorial board member of *World Journal of Gastroenterology*, Ludovico Abenavoli, MD, MSc, PhD, Associate Professor, Health Sciences, University Magna Graecia, Campus Germaneto, Viale Europa, 88100 Catanzaro, Italy

AIMS AND SCOPE

World Journal of Gastroenterology (*World J Gastroenterol*, *WJG*, print ISSN 1007-9327, online ISSN 2219-2840, DOI: 10.3748) is a peer-reviewed open access journal. *WJG* was established on October 1, 1995. It is published weekly on the 7th, 14th, 21st, and 28th each month. The *WJG* Editorial Board consists of 1375 experts in gastroenterology and hepatology from 68 countries.

The primary task of *WJG* is to rapidly publish high-quality original articles, reviews, and commentaries in the fields of gastroenterology, hepatology, gastrointestinal endoscopy, gastrointestinal surgery, hepatobiliary surgery, gastrointestinal oncology, gastrointestinal radiation oncology, gastrointestinal imaging, gastrointestinal interventional therapy, gastrointestinal infectious diseases, gastrointestinal pharmacology, gastrointestinal pathophysiology, gastrointestinal pathology, evidence-based medicine in gastroenterology, pancreatology, gastrointestinal laboratory medicine, gastrointestinal molecular biology, gastrointestinal immunology, gastrointestinal microbiology, gastrointestinal genetics, gastrointestinal translational medicine, gastrointestinal diagnostics, and gastrointestinal therapeutics. *WJG* is dedicated to become an influential and prestigious journal in gastroenterology and hepatology, to promote the development of above disciplines, and to improve the diagnostic and therapeutic skill and expertise of clinicians.

INDEXING/ABSTRACTING

World Journal of Gastroenterology (*WJG*) is now indexed in Current Contents[®]/Clinical Medicine, Science Citation Index Expanded (also known as SciSearch[®]), Journal Citation Reports[®], Index Medicus, MEDLINE, PubMed, PubMed Central and Directory of Open Access Journals. The 2017 edition of Journal Citation Reports[®] cites the 2016 impact factor for *WJG* as 3.365 (5-year impact factor: 3.176), ranking *WJG* as 29th among 79 journals in gastroenterology and hepatology (quartile in category Q2).

FLYLEAF

I-IX Editorial Board

EDITORS FOR THIS ISSUE

Responsible Assistant Editor: *Xiang Li*
Responsible Electronic Editor: *Fen-Fen Zhang*
Proofing Editor-in-Chief: *Lian-Sheng Ma*

Responsible Science Editor: *Ze-Mao Gong*
Proofing Editorial Office Director: *Jin-Lei Wang*

NAME OF JOURNAL
World Journal of Gastroenterology

ISSN
 ISSN 1007-9327 (print)
 ISSN 2219-2840 (online)

LAUNCH DATE
 October 1, 1995

FREQUENCY
 Weekly

EDITORS-IN-CHIEF
Damian Garcia-Olmo, MD, PhD, Doctor, Professor, Surgeon, Department of Surgery, Universidad Autonoma de Madrid; Department of General Surgery, Fundacion Jimenez Diaz University Hospital, Madrid 28040, Spain

Stephen C Strom, PhD, Professor, Department of Laboratory Medicine, Division of Pathology, Karolinska Institutet, Stockholm 141-86, Sweden

Andrzej S Tarnawski, MD, PhD, DSc (Med), Professor of Medicine, Chief Gastroenterology, VA Long Beach Health Care System, University of California, Irvine, CA, 5901 E. Seventh Str., Long Beach,

CA 90822, United States

EDITORIAL BOARD MEMBERS
 All editorial board members resources online at <http://www.wjgnet.com/1007-9327/editorialboard.htm>

EDITORIAL OFFICE
 Jin-Lei Wang, Director
 Yuan Qi, Vice Director
 Ze-Mao Gong, Vice Director
World Journal of Gastroenterology
 Baishideng Publishing Group Inc
 7901 Stoneridge Drive, Suite 501,
 Pleasanton, CA 94588, USA
 Telephone: +1-925-2238242
 Fax: +1-925-2238243
 E-mail: editorialoffice@wjgnet.com
 Help Desk: <http://www.fpublishing.com/helpdesk>
<http://www.wjgnet.com>

PUBLISHER
 Baishideng Publishing Group Inc
 7901 Stoneridge Drive, Suite 501,
 Pleasanton, CA 94588, USA
 Telephone: +1-925-2238242
 Fax: +1-925-2238243
 E-mail: bpoffice@wjgnet.com
 Help Desk: <http://www.fpublishing.com/helpdesk>

<http://www.wjgnet.com>

PUBLICATION DATE
 July 7, 2017

COPYRIGHT
 © 2017 Baishideng Publishing Group Inc. Articles published by this Open-Access journal are distributed under the terms of the Creative Commons Attribution Non-commercial License, which permits use, distribution, and reproduction in any medium, provided the original work is properly cited, the use is non commercial and is otherwise in compliance with the license.

SPECIAL STATEMENT
 All articles published in journals owned by the Baishideng Publishing Group (BPG) represent the views and opinions of their authors, and not the views, opinions or policies of the BPG, except where otherwise explicitly indicated.

INSTRUCTIONS TO AUTHORS
 Full instructions are available online at <http://www.wjgnet.com/bpg/gerinfo/204>

ONLINE SUBMISSION
<http://www.fpublishing.com>

Observational Study

**Cryptogenic multifocal ulcerous stenosing enteritis:
Radiologic features and clinical behavior**

Jiyoung Hwang, Jin Sil Kim, Ah Young Kim, Joon Seok Lim, Se Hyung Kim, Min Ju Kim, Mi Sung Kim,
Kyoung Doo Song, Ji Young Woo

Jiyoung Hwang, Department of Radiology, Soonchunhyang University Seoul Hospital, Seoul 04401, South Korea

Jin Sil Kim, Ah Young Kim, Department of Radiology and Research Institute of Radiology, University of Ulsan College of Medicine, Asan Medical Center, Seoul 05505, South Korea

Joon Seok Lim, Department of Radiology, Research Institute of Radiological Science, Yonsei University College of Medicine, Seoul 03722, South Korea

Se Hyung Kim, Department of Radiology, Seoul National University Hospital, Seoul 03080, South Korea

Min Ju Kim, Department of Radiology, Anam Hospital, Korea University College of Medicine, Seoul 15355, South Korea

Mi Sung Kim, Department of Radiology, Sungkyunkwan University School of Medicine, Kangbuk Samsung Hospital, Seoul 03181, South Korea

Kyoung Doo Song, Department of Radiology and Center for Imaging Science, Samsung Medical Center, Sungkyunkwan University School of Medicine, Seoul 06351, South Korea

Ji Young Woo, Department of Radiology, Hallym University College of Medicine, Kangnam Sacred Heart Hospital, Seoul 07441, South Korea

Author contributions: Hwang J and Kim JS contributed equally to this work; Kim AY, Lim JS, Kim SH, Kim MJ, Kim MS, Song KD and Woo JY collected the data; Kim AY managed the study; Hwang J, Kim JS and Kim AY contributed to the analysis and wrote the paper; all authors have read and approved the final version to be published.

Institutional review board statement: This study was approved by the institutional review board of Asan medical center, Seoul, South Korea (number, 2016-0235).

Conflict-of-interest statement: The authors declare no conflicts

of interests related to the publication of this study.

Data sharing statement: No additional data are available.

Open-Access: This article is an open-access article which was selected by an in-house editor and fully peer-reviewed by external reviewers. It is distributed in accordance with the Creative Commons Attribution Non Commercial (CC BY-NC 4.0) license, which permits others to distribute, remix, adapt, build upon this work non-commercially, and license their derivative works on different terms, provided the original work is properly cited and the use is non-commercial. See: <http://creativecommons.org/licenses/by-nc/4.0/>

Manuscript source: Invited manuscript

Correspondence to: Ah Young Kim, MD, PhD, Professor, Department of Radiology and Research Institute of Radiology, University of Ulsan College of Medicine, Asan Medical Center, 88, Olympic-ro 43-gil, Songpa-gu, Seoul 05505, South Korea. aykim@amc.seoul.kr
Telephone: +82-2-30104361
Fax: +82-2-4764719

Received: February 15, 2017

Peer-review started: February 19, 2017

First decision: March 16, 2017

Revised: March 23, 2017

Accepted: May 19, 2017

Article in press: May 19, 2017

Published online: July 7, 2017

Abstract**AIM**

To investigate the characteristic radiologic findings of cryptogenic multifocal ulcerous stenosing enteritis (CMUSE) which can be differentiated from other similar bowel disease and to assess their clinical behavior.

METHODS

Twenty pathologically and clinically confirmed CMUSE patients (males:females = 8:12; mean age: 40.4 years) between March 2002 and August 2015 from seven academic centers in South Korea were retrospectively reviewed. We evaluated small bowel series (SBS; $n = 25$), computed tomography (CT) enterography ($n = 21$), magnetic resonance (MR) enterography ($n = 2$), and abdominopelvic CT ($n = 18$) images, focusing on enteric and perienteric manifestations. Any change in radiologic features during follow-up period was recorded. We evaluated clinical data including presenting symptoms, laboratory finding and presence of relapse from electronic medical records. Histopathologic findings were also evaluated.

RESULTS

The main symptoms were abdominal pain ($n = 12$) and anemia ($n = 10$). All patients showed small bowel strictures ($n = 52$, mean: 2.6 per patient) on initial CT/MR, located in the ileum ($n = 47$) or jejunum ($n = 5$). Strictures showed short-length (mean: 10.44 mm) and circumferential bowel wall thickening (mean: 5.56 mm) with layered enhancement ($n = 48$) that were also noted on initial SBS ($n = 36$) with shallow ulcers ($n = 10$). Some ulcerative lesions or wall thickening progressed into strictures on follow-up SBS/CT, and some strictures revealed recurrent ulceration on follow-up SBS. There were no penetrating disease features like fistula or abscess and no gastrointestinal tract involvement except the small bowel. Nine patients experienced disease recurrence (median relapse-free period: 32 mo) even post-operatively. Histopathologic features of surgically resected specimens were characterized as multiple superficial ulcerations confined to mucosa or submucosa and multiple strictures.

CONCLUSION

Under characteristic radiologic findings with multiple short-segmental strictures and/or shallow ulcers of the small intestine, CMUSE should be considered when assessing patients with recurrent abdominal pain and anemia.

Key words: Cryptogenic multifocal ulcerous stenosing enteritis; Small intestine; Computed tomography; Small bowel series; Diagnosis

© **The Author(s) 2017.** Published by Baishideng Publishing Group Inc. All rights reserved.

Core tip: The patients with multiple unexplained strictures and recurrent abdominal pain are challenging to most physician in daily practice. Because cryptogenic multifocal ulcerous stenosing enteritis (CMUSE) is a rare disease, this disease entity is frequently misdiagnosed as other small bowel diseases and often undergoes unnecessary surgery. Then, we focused on the characteristic radiologic and clinical findings of CMUSE that can differentiate the disease from other similar bowel diseases. In our study, characteristic

radiologic features of CMUSE were multiple short strictures and/or shallow ulcers of the small intestine without significant bowel obstruction. Some strictures had recurrent shallow ulceration that might progress to more severe strictures. Under these radiologic findings with relapsing episodes, CMUSE should be considered when assessing patient with recurrent abdominal pain and anemia.

Hwang J, Kim JS, Kim AY, Lim JS, Kim SH, Kim MJ, Kim MS, Song KD, Woo JY. Cryptogenic multifocal ulcerous stenosing enteritis: Radiologic features and clinical behavior. *World J Gastroenterol* 2017; 23(25): 4615-4623 Available from: URL: <http://www.wjgnet.com/1007-9327/full/v23/i25/4615.htm> DOI: <http://dx.doi.org/10.3748/wjg.v23.i25.4615>

INTRODUCTION

Cryptogenic multifocal ulcerous stenosing enteritis (CMUSE) is a rare disease characterized by chronic and recurrent ileus resulting from multiple unexplained strictures and superficial ulcerations of the small intestine^[1,2]. At present, CMUSE is diagnosed mainly based on clinical and macroscopic imaging features of the small intestine. However, CMUSE is frequently misdiagnosed as other ulcerative small bowel diseases, such as Crohn's disease (CD) or nonsteroidal anti-inflammatory drugs (NSAIDs)-induced enteropathy, because of similar clinical symptoms and nonspecific laboratory findings^[3,4].

Although double-balloon endoscopy seems to be the most important diagnostic tool, it is still difficult to diagnose CMUSE even after ruling out the above-mentioned diseases due to limited diagnostic tools and lack of characteristic radiologic features on computed tomography (CT) or small bowel series (SBS). Consequently, most patients with CMUSE have suffered from improper diagnosis and management, including unnecessary surgery.

To our knowledge, most studies about CMUSE have focused on clinical findings and vague radiologic findings such as the presence or absence of strictures and ulcers^[3,4]. Hence, this study aims to investigate the characteristic findings of CMUSE that can differentiate the disease from other similar bowel diseases and to assess their clinical behavior through analysis of serial radiologic imagings including SBS, routine abdominopelvic CT (APCT), CT enterography (CTE), and/or magnetic resonance (MR) enterography (MRE) as part of routine abdominal examination in a clinical setting.

MATERIALS AND METHODS

Study population

Between March 2002 and August 2015, 36 patients with suspected CMUSE were included from seven

academic centers in South Korea. The diagnostic criteria for CMUSE were based on previous studies^[1,5]: (1) unexplained small bowel strictures; (2) superficial ulcer in the mucosa and submucosa; (3) chronic or relapsing ulcerative stenosis and abdominal pain; (4) no signs of systemic inflammation; and (5) persistent and occult blood loss from the gastrointestinal (GI) tract except during bowel rest or postoperative period. Patients' records were reviewed to confirm CMUSE as final diagnosis. Twelve patients were excluded due to lack of any pathologic proof suggesting superficial mucosal or submucosal disease. Four patients were also excluded because their final histopathology showed transmural inflammation ($n = 2$) or panmural inflammation with bowel perforation ($n = 2$). Patients (8 males, 12 females) were aged 15-64 years (mean, 40.4 ± 14.8) when CMUSE was diagnosed. All patients underwent at least one radiologic examination including SBS, APCT, CTE, or MRE. This study was approved by the IRB of Asan medical center, Seoul, Korea (number, 2016-0235).

Clinical and laboratory data

Clinical data included sex, age, presenting symptoms, period from onset to hospital visit, the initial diagnosis, and laboratory findings [hemoglobin (Hb), albumin, C-reactive protein (CRP), and white blood cell count (WBC)] at first admission. Moreover, information on follow-up duration, presence of relapse, relapsing interval, treatment methods, treatment outcome, histopathologic findings, total number of each radiologic examination, and total number of surgical resection were collected from the electronic medical records. The type, number, and descriptions of endoscopic inspections were also recorded.

Remission was defined as removal of the presenting symptoms such as bleeding, abdominal pain, or anemia, and relapse as recurrence of bleeding, abdominal pain, or anemia; need for additional medication; or evidence of a recurrent imaging finding by any imaging after remission.

Radiologic examinations

All analysis images were available on a picture archiving and communications system (PACS). SBS were performed in 16 patients, after they fasted for at least 8 h, using a barium and/or methylcellulose solution. A single-contrast examination was performed to evaluate the entire small intestine, followed by a double-contrast examination after peroral administration of an effervescent agent in 9 patients.

APCT and CTE were performed in 11 and 13 patients, respectively, using multidetector CT scanners (Light Speed plus, Light Speed 16 or Light Speed VCT 64, GE Healthcare, Milwaukee, WI, United States; Somatom sensation 16 or Somatom Definition, Siemens Medical Systems, Erlangen, Germany; Toshiba Aquilion 64, Toshiba Medical System, Tokyo,

Japan) with intravenous contrast material (Ultravist 370, Schering, Berlin, Germany; Omnipaque, Amersham Health, Inc., Princeton, NJ, United States). APCT consisted of a non-contrast enhanced image and a portal phase image with a fixed 60-70-s delay following the injection of intravenous contrast media.

CTE consisted of a non-contrast image, enteric phase, and portal venous phase image. The enteric- and portal- phase scans were obtained with a 20-25-s delay using a bolus-triggering method or a fixed 40-s delay, and a fixed 70-80-s delay after injection, respectively. CTE or MRE was performed after ingestion of 1200 mL of neutral oral contrast agent using a polyethylene glycol electrolytic solution (Colyte, Taejun Pharmacy, Seoul, South Korea) or 2.5% sorbitol solution.

MRE was acquired with a 3.0-T system (Ingenia, Philips Healthcare, Best, The Netherlands) after intravenous injection of a spasmolytic agent (Buscopan, Boehringer Ingelheim, Ingelheim, Germany). T2-weighted images, diffusion-weighted images (with b -factors of 0 and 900 s/mm²), and dynamic contrast-enhanced images were obtained. Coronal dynamic fat suppressed spoiled gradient-echo T1-weighted sequences, including unenhanced imaging and enteric and portal venous phase sequences, were performed with 0.2 mL/kg body weight gadoterate meglumine (Dotarem; Guerbet, Villepinte, France). Enteric phase was obtained 7 s after the contrast material first arrived at the iliac bifurcation after injection. Subsequent phase imaging was performed continuously after completion of the prior phase with a brief pause to let the patient breathe.

Image analysis

All radiologic images were reviewed by three board-certified radiologists (Kim AY, > 10 years of experience in abdominal imaging; Hwang J, 5 years; and Kim JS, 2 years) in consensus.

In the first stage, all initial radiologic examinations obtained during the initial hospital visit were analyzed, focusing on ulcer, stricture, bowel wall thickening, and other ancillary findings. Radiologic jejunal involvement was regarded as involvement of the proximal two-fifths of the intact small bowel^[6].

For strictures, defined as upstream dilatation of small bowel > 3 cm in diameter, the number, location, lesion length, and interval between each lesion were recorded during all radiologic studies. Small bowel obstruction was also evaluated with a 4-grading scale (absent, low-grade, high-grade, or complete obstruction). While a low-grade obstruction was defined as dilatation of proximal small bowel 3-4 cm in diameter, high-grade obstruction was defined as proximal small bowel dilatation of > 4 cm in diameter. Complete obstruction was defined as high-grade obstruction without distal passage of contrast media. Any deep penetrating disease features such as sinus,

fistula, or abscess were recorded.

On APCT/CTE/MRE, thickness (one-side wall thickness), enhancement pattern (layered or transmural), enhancement degree (mild, similar enhancement to adjacent normal bowel; moderate, more than normal bowel wall enhancement; strong, marked enhancement compared with normal bowel) of stricture were recorded respectively. In addition, the presence of segmental bowel wall thickening (> 10 cm in length) was recorded, noting the length, thickness, enhancement pattern, and enhancement degree. Mesenteric manifestations such as mesenteric lymphadenopathy, mesenteric hypervascularity or infiltration, and ascites were recorded. In addition, the number, location, lesion length, shape (eccentric, circumferential, linear or aphthoid) of ulcer, and interval between each lesion were evaluated on SBS.

In the second stage, follow-up images (11 SBS, 7 APCT, 13 CTE, and 1 MRE) were assessed focusing on any change from the previous image findings for each patient. Follow-up images without any presenting symptom or image findings changes were excluded from analysis.

RESULTS

Clinical characteristics and laboratory findings

Clinical and laboratory findings are summarized in Table 1. The median (range) period from initial onset of symptoms to hospital visit was 8 (0.1-25) years. The main symptoms were abdominal pain (60%, 12/20) and anemia (50%, 10/20); four patients presented with GI bleeding (20%, 4/20). On initial laboratory test, the mean Hb level was 9.9 ± 2.4 g/dL and mean albumin level was 3.4 ± 2.2 g/dL. WBC and CRP levels were within normal ranges for most patients or slightly increased in two and three patients, respectively.

The median (range) follow-up period was 4 (0-13.5) years. The initial diagnosis before confirmative diagnosis included CD, intestinal tuberculosis, post-ischemic stricture, CMUSE, vasculitis, protein-losing enteropathy, and non-granulomatous jejunoileitis.

Endoscopic findings

Double-balloon enteroscopy (DBE) or capsule endoscopy (CE) was performed in 10 patients (Table 1). All patients showed multiple ulcers on endoscopic examination: jejunal ($n = 1$), ileal ($n = 6$), and both ($n = 3$). Ulcerative lesions revealed discrete margin and various shapes including linear, circular, geographic, or irregular. Some showed transverse or circular alignment. The intervening mucosa appeared normal. Nine patients also showed multiple strictures: jejunal ($n = 2$), ileal ($n = 6$), and both ($n = 1$). All strictures showed short segmental involvement and were often accompanied by ulcers. Full small intestine evaluation failed in five patients with DBE (5/9, 55.5%) and in two patients with CE (2/5, 40%) due to stricture.

Histopathologic findings

A total of 15 biopsy specimens were available from 11 patients who underwent endoscopic examination. All lesions revealed chronic, nonspecific inflammation with/without ulceration.

Sixteen patients underwent surgical resection of small intestine. Histopathologic features of these resected specimens were characterized as multiple superficial ulcerations confined to mucosa or submucosa and multiple strictures. Some strictures were ulcerated. Ulcerative lesions showed linear or circular configuration and some were associated with submucosal thickening by fibrosis. Some ulcerative lesions were accompanied by chronic crypt inflammation, pyloric gland metaplasia, or lymphoid aggregates. Inflammatory change was not seen in the small bowel segment between ulcers. There was no giant cell granuloma, transmural inflammation, fistula or fissural ulceration, or vasculitis features.

Radiological findings

A total of 21 CTE, 18 APCT, 2 MRE, and 25 SBS were finally included in the analysis. Among them, 8 CTE, 11 APCT, 1 MRE, and 14 SBS were performed during the patients' initial hospital visit.

On initial CT or MRI, all patients revealed strictures with a total of 52 (mean number per patient, 2.6; Table 2 and Figure 1) analyzed. Most strictures were located in the ileum (90.4%), followed by jejunum (9.6%). While most patients showed only ileal strictures (17/20, 85%), two had only jejunal strictures and one had both. Mean stricture length and thickness was 10.44 ± 3.95 mm and 5.56 ± 1.58 mm, respectively. On contrast-enhanced CT or MRI, most strictures showed moderate (75%) or strong (9.6%) enhancement with a layered pattern (92.3%; Figure 1). The median (range) interval length between strictures was 4 (1.5-20) cm.

Four out of 20 patients showed segmental bowel wall thickening on the initial CT or MRI (Figures 2 and 3). Mean length and thickness of this bowel wall thickening was 36.25 ± 13.77 cm and 7.75 ± 1.26 mm, respectively (Table 2).

Despite of multiple strictures, high-grade small bowel obstruction was noted in only two patients on CT or MR images. Most patients showed no or low-grade small bowel obstruction.

On initial SBS, 36 strictures were detected in 13 patients (mean number of stricture per patient, 1.8; Table 3, Figures 1 and 4). Most strictures were located in the ileum (97.2%), and the mean stricture length was 9.81 ± 4.65 mm. Median (range) interval length between strictures was 4.5 (1.5-20.4) cm. Seven patients also showed 10 ulcers on SBS; nine were located in the ileum and showed eccentric alignment (90%), and six ulcerative lesions were accompanied by stricture (Figure 4).

Follow-up imaging (11 SBS, 7 APCT, 13 CTE, and 1 MRE) was performed in 14 patients. While there was

Table 1 Clinical characteristics, diagnostic tools, and treatment outcomes of patients with cryptogenic multifocal ulcerous stenosing enteritis

Patient No./sex/age (yr)	Duration until hospital visit (yr) ¹	Presenting symptom at admission	Hb (g/dL)	SBS	APCT/CTE/MRE	DE/CE	Treatment	Treatment response (No. of relapse)	Relapse-free period (yr)	No of surgery	Follow up period (yr) ²
1/M/23	9.6	Abdominal pain	13.3	1	3/-/-	1/-	Surgery	Relapse (1)	1.1	1	2
2/F/56	22.9	Abdominal pain	11.4	ND	1/-/-	-/-	Steroid, 5-ASA, surgery	Remission		1	3
3/M/49	10.4	Abdominal pain	13.7	ND	1/1/-	-/1	Surgery	Remission		1	1.5
4/F/61	25	Anemia	10.5	1	1/-/-	-/-	Surgery	Remission		1	2.9
5/F/29	1	Anemia	7.4	2	6/1/-	1/1	Steroid, 5-ASA, surgery	Relapse (1)	1	1	9.9
6/F/37	8.6	Anemia	8	2	1/-/-	-/-	Steroid, 5-ASA, surgery, others (albumin)	Relapse (2)	2.7, 6.4	1	13.4
7/F/45	7.5	Anemia, GI bleeding	7.6	3	-/3/-	2/-	Steroid, surgery	Relapse (2)	0.5, 3.7	1	9.4
8/F/49	5.1	Abdominal pain	12.9	ND	-/2/1	2/-	Steroid, surgery	Relapse (3)	1.4, 6, 1.7	3	9.8
9/M/54	3.5	Abdominal pain, anemia	10.4	1	-/1/-	-/-	Surgery	Remission		1	2.5
10/M/30	4.8	Abdominal pain	8.5	5	1/4/-	-/1	5-ASA, surgery	Relapse (2)	7.9, 0.7	2	10
11/F/16	14.6	Abdominal pain	7.7	ND	-/1/-	-/-	Surgery	Remission		1	0.5
12/M/24	0.9	Abdominal pain, anemia, GI bleeding	11.3	1	-/2/-	-/-	Surgery, others (conservative treatment)	Relapse (1)	2.5	1	3.6
13/M/41	12	Abdominal pain	9	1	-/1/-	1/-	Steroid	Remission		0	0
14/F/64	23	Abdominal pain	12.3	ND	-/-/1	-/-	Surgery	Remission		1	0
15/F/39	10.3	Abdominal pain	9	1	-/3/-	-/-	5-ASA, surgery	Relapse (1)	2	1	6.2
16/M/60	10	Anemia, GI bleeding	9.1	ND	-/1/-	-/1	Steroid, 5-ASA	Remission		0	1.1
17/F/15	4	Abdominal pain, anemia	7.7	4	1/-/-	-/-	Steroid, 5-ASA, surgery	Relapse (3)	3, 4, 4, 3	4	13.5
18/F/34	0.1	Anemia	7.6	1	1/1/-	1/-	Steroid, tuberculosis drug	Remission		0	10.6
19/M/33	1.2	Anemia	6.1	1	1/-/-	1/1	Steroid	Remission		0	4.5
20/F/48	0.2	GI bleeding	13.4	1	1/-/-	-/-	Surgery	Remission		1	4.5

¹Time interval between the symptom onset and hospital visit (yr); ²Time interval between prior remission and recurrence (yr). SBS: Small bowel series; APCT: Abdominopelvic CT; CTE: CT enterography; MRE: MR enterography; DE: Double-balloon endoscopy; CE: Capsule endoscopy; ND: Not done; 5-ASA: 5-aminosalicylic acid; GI: Gastrointestinal.

no significant interval change in radiologic features for three of the patients, new strictures and/or ulcers were detected at another site with a progressive nature in seven patients, even post-operatively. On follow-up SBS examinations, ulcers were accompanied with stricture and these ulcerative strictures progressed into more severe strictures over time (Figure 4). Segmental bowel wall thickening newly developed in three patients on follow-up CT. Finally, seven patients had segmental bowel wall thickening on the initial or follow-up CT. In these patients, this radiologic feature completely improved on the next follow-up CT, but eventually, strictures manifested within this thickened bowel segment during the follow-up period (Figures 2

and 3). Disease recurrence at the surgical anastomotic site was noted in only one patient.

There was no penetrating disease features, including fissure, fistula, or abscess, by CT/MRI or SBS. GI involvement other than small intestine was not seen.

Treatment outcomes

Sixteen patients underwent surgical resection (80%), and three received reoperation due to recurred disease (median surgery number, 1). In eight patients, surgery was performed as initial treatment on first admission. Five patients underwent surgery due to ineffective medical therapy during hospitalization. Only four

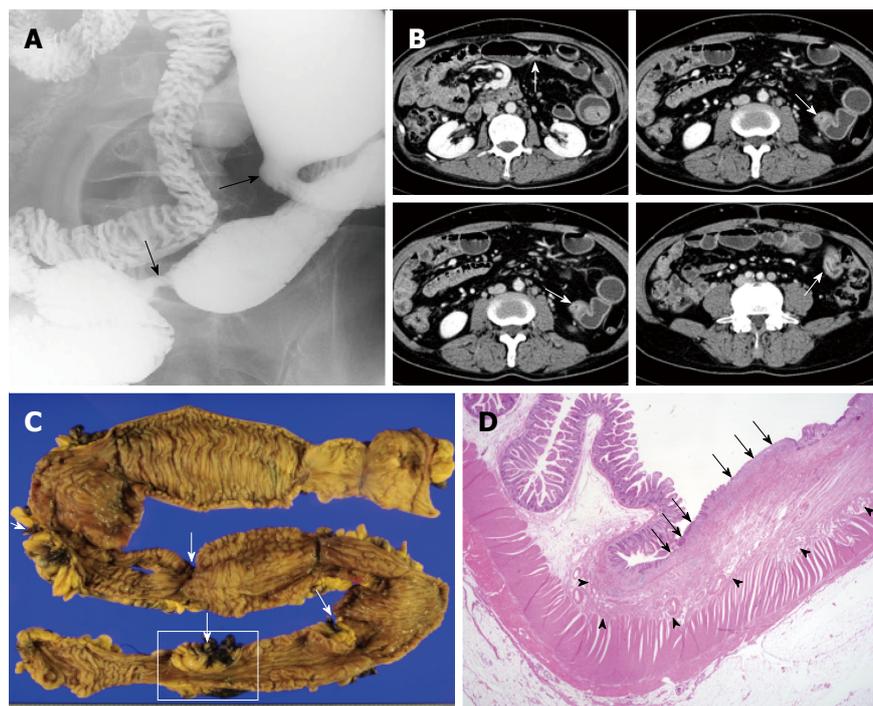


Figure 1 A 54-year-old man with abdominal pain and anemia. A: Small bowel series spot compression image demonstrates multiple short strictures along the ileum (arrows); B: On axial images of contrast-enhanced computed tomography (CT) enterography, short-segmental strictures (arrows) show moderate and layered bowel wall enhancement with mild dilatation of intervening segment between strictures; C: Gross specimen of resected small intestine shows multiple short segmental strictures (arrows) with mild dilatation of intervening bowel segment, which corresponds with CT images (arrows in B); D: On the low-power H&E staining of surgical specimen (boxed area in Figure 1C) there is a superficial ulcer (arrows) and submucosal fibrosis (arrow heads) without evidence of transmural inflammation or granulomatous lesion (magnification $\times 10$).

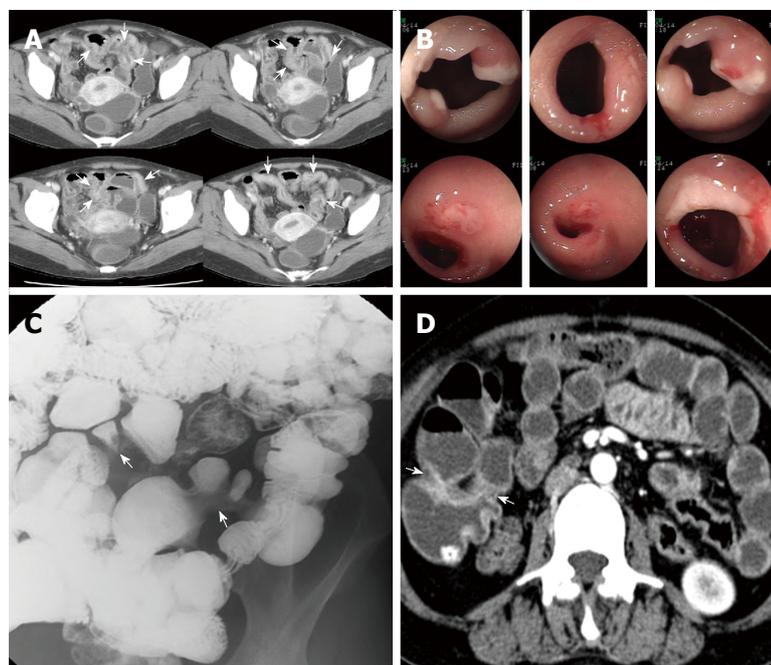


Figure 2 A 34-year-old woman with anemia. A: Axial images of contrast-enhanced computed tomography (CT) enterography show segmental bowel wall thickening (arrows) with suspicious strictures along the distal ileum; B: On retrograde double-balloon endoscopic examination at the same time period there are multiple sharply demarcated ulcers at or near the strictures of the ileum; C: Small bowel series spot radiograph obtained two months later reveals segmental luminal narrowing (arrows) along the distal ileum, without overt ulceration; D: On axial images of contrast-enhanced CT enterography after two years, previous bowel wall thickening of distal ileum has improved and prominent strictures are noted at the corresponding ileal segment.

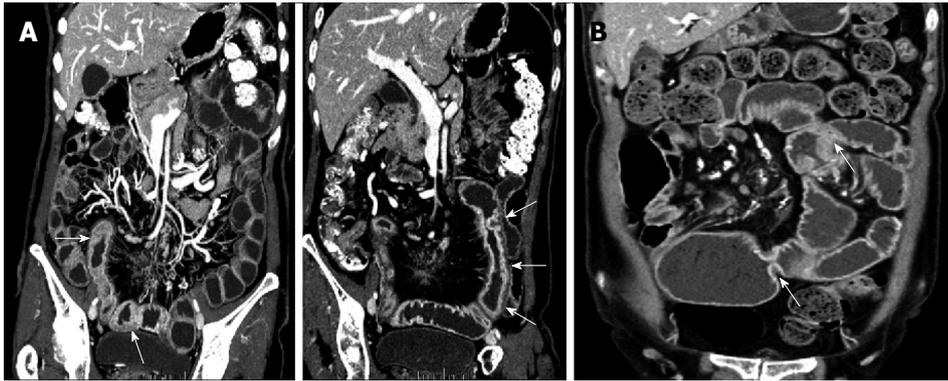


Figure 3 A 49-year-old woman with abdominal pain. A: Coronal reformatted images of computed tomography (CT) enterography show long segmental bowel wall thickening with layered enhancement pattern, involving distal jejunum and proximal ileum. B: This bowel wall thickening of the small intestine is not noted on the coronal reformatted image of CT enterography four years later, and instead, several short-segmental strictures (arrows) have developed in the same bowel loop.

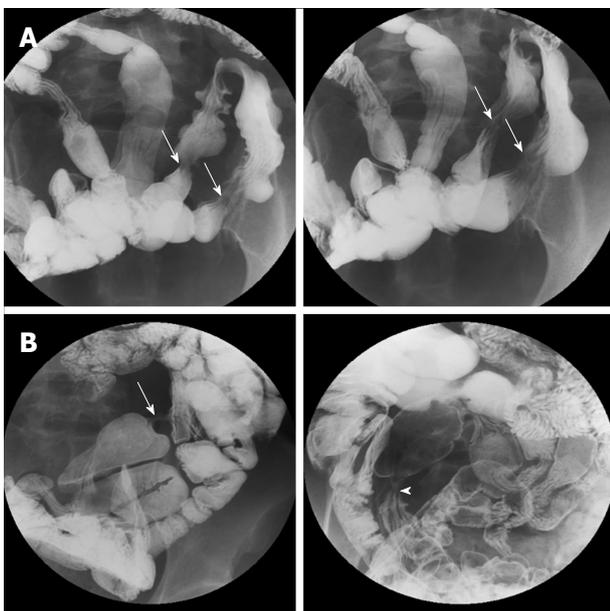


Figure 4 A 45-year-old woman with gastrointestinal bleeding and anemia. A: Small bowel series (SBS) spot compression radiographs show shallow ulcers (arrows) with mucosal fold thickening and subtle luminal narrowing at the ileum; B: After four years, double-contrast SBS spot radiographs demonstrate a severe stricture (arrow) and another stricture with shallow ulceration (arrowhead), suggestive of worsening of previous strictures.

patients showed remission with medical treatment: steroid only ($n = 2$), steroid with oral 5-aminosalicylic acid ($n = 1$), and steroid with tuberculosis drugs ($n = 1$). None of them had recurrence.

As the final outcome, nine patients (45%) revealed disease recurrence (median relapse-free period, 2.6 years; range, 0.5-7.9). The median (range) relapse number was 2 (1-3).

DISCUSSION

To our knowledge, no study focused on the radiologic features of CMUSE to assist in the differentiation from other similar bowel diseases. Even radiologic information previously reported was relatively simple

Table 2 Radiologic features on computed tomography/magnetic resonance imaging of patients with cryptogenic multifocal ulcerous stenosing enteritis ($n = 20$)

Stricture	$n = 52$
Total No. (mean No. per patient, range)	52 (2.6, 1-6)
Location (Jejunum/Ileum)	5/47
Stricture length (mm), mean \pm SD	10.44 \pm 3.95
Stricture thickness (mm), mean \pm SD	5.56 \pm 1.58
Enhancement pattern (layered/transmural)	48/4
Degree of enhancement (mild/moderate/strong)	8/39/5
Interval between strictures (cm), median (range)	4 (1.5-20)
Segmental bowel wall thickening	$n = 4$
Location (Jejunum/Ileum/both)	0/3/1
Wall thickening length (cm), mean \pm SD	36.25 \pm 13.77
Wall thickening thickness (mm), mean \pm SD	7.75 \pm 1.26
Wall thickening pattern (circumferential/eccentric)	2/2
Enhancement pattern (layered/transmural)	3/1
Degree of enhancement (mild/moderate/strong)	1/2/1
Small bowel obstruction	
Absent/low grade/high grade/complete	13/5/2/0
Mesenteric manifestation	
Mesenteric lymphadenopathy	6
Mesenteric hypervascularity or infiltration	4

descriptions and mostly confined to SBS features^[1,4,5,7,8].

In this study, the main radiologic manifestations of CMUSE patients were multiple strictures and/or ulcers of the small intestine without significant bowel obstruction. These results were similar to those previously reported. However, bowel wall strictures in this study were short and thin with a layered enhancement pattern (< 2 cm long and < 1 cm thick) with layered enhancement pattern on axial imaging tools such as CT or MRI. Thus, short, thin strictures with layered enhancement may represent relatively superficial disease process, accounting for no or low-grade small bowel obstruction. Histopathologic proof might provide a more convincing explanation as the stricture segment of a surgical specimen revealed fibrosis confined only to submucosa (Figure 1).

The distinctive radiologic finding was the presence of long-segmental bowel wall thickening on CT. Overall, seven patients showed this feature on initial and/or

Table 3 Radiologic features on small bowel series of patients with cryptogenic multifocal ulcerous stenosing enteritis ($n = 14$)

Stricture	$n = 36$
Total No. (mean No. per patient, range)	36 (1.8, 1-5)
Location (Jejunum/Ileum)	1/35
Stricture length (mm), mean \pm SD	9.81 \pm 4.65
Interval between strictures (cm), median (range)	4.5 (1.5-20.4)
Ulcer	$n = 10$
Total No. (mean No. per patient, range)	10 (0.5, 1-2)
Location (Jejunum/Ileum)	1/9
Ulcer length (mm), mean \pm SD	11.3 \pm 5.58
Ulcer pattern (eccentric/circumferential/linear/aphthous)	9/1/0
Small bowel obstruction	
Absent/low grade/high grade/complete	7/5/2/0

follow-up CT. Three of them underwent DBE at the same time and showed multiple, shallow ulcerations of the small intestine in accordance with CT results. On follow-up CT, this segmental bowel wall thickening progressed to multiple strictures with improved thickening (Figures 2 and 3), confirmed by follow-up endoscopic examination. Thus, this segmental bowel wall thickening on CT might represent the presence of multiple, superficial ulcerations, although there was no direct correlation between these two diagnostic tools in all patients.

One different point of our study results is that ulcers were less frequently depicted on SBS with shallow ulcerations mainly located in the ileum, contrary to previous endoscopic reports of multiple ulcers mainly in jejunum and proximal ileum^[5]. We think that the relatively small numbers of ulcers in our radiologic examinations might be due to delayed hospital visits during the healing stage of the ulcer with stricture formation. Early in CMUSE disease progression, more ulcers might be detected and more strictures noted in advanced stages. Considering the long median period from initial onset of symptoms to the hospital visits in our study (8 years), less frequent ulcer depiction can be predicted. Another assumption is the relatively high failure rate of endoscopic examination due to strictures. In this study, the endoscopic failure rate was 55.5% in DBE and 40% of CE; hence, the endoscopic examination failure rate might be relevant to the larger numbers of clinically reported jejunal ulcers.

More importantly, during follow-up studies, these ulcers tended to progress to strictures with ulceration healing (Figure 4). Conversely, strictures accompanied by shallow ulceration during disease recurrence progressed to more severe strictures. Perhaps shallow ulcers and their healing, in the repeated disease process, lead to more severe strictures that required surgical intervention. Furthermore, recurrent ulcers or strictures rarely involved the anastomosis site of the small intestine after surgical resection. Strictures and ulcers showed no consistent direction such as

the mesenteric border lining. Like previous reports, there were no penetrating disease features, including fissure, fistula formation, or abscess^[1,3,7]. Mesenteric lymphadenopathy or mesenteric infiltration was rarely seen and there was no other GI involvement except the small intestine. Hence, we think that these radiologic features might be helpful to differentiate CMUSE from other inflammatory bowel disease, especially CD.

The main clinical manifestations of the patients in this study were abdominal pain and anemia, resulting from chronic or relapsing subileus episodes associated with small intestinal strictures and small intestinal occult blood loss, respectively. In addition, most patients did not show biological signs of systemic inflammatory reaction. These clinical features largely met the description of previous reports^[1,3,5,9,10].

Contrary to previous major reports^[1,3,5,9,10], it was difficult to establish the efficacy of steroid therapy in this study. Only four patients were treated only with steroids; others were treated with combination of steroid with oral 5-aminosalicylic acid or tuberculosis drugs. Furthermore, the efficacy of steroid therapy is debatable. Matsumoto *et al.*^[5] reported steroid, oral 5-aminosalicylic acid, and oral azathioprine failed to induce mucosal healing or prevent recurrence of small intestine ulcers. Kim *et al.*^[11] reported on steroid-refractory disease cases. Similarly, in this study, two out of four patients treated with steroid only required surgery due to disease relapse and were no longer steroid-sensitive.

As a final outcome, the major therapeutic choice of this study population was surgery. Eight patients underwent surgery during the first hospital visit and eight on subsequent visits, with three patients requiring two or more surgical treatments. Another cause of surgical treatment was retention of capsule endoscope in the small intestine due to stricture ($n = 2$). With the long median pre-diagnostic period (8 years) and high disease relapse rate (45%), higher choice of surgery rather than medicine could be acceptable in this study due to the frequent manifestation of strictures rather than ulcers. Perhaps the conflicting responsiveness to steroid use in CMUSE patients is linked closely to different disease stages, *i.e.*, early stage manifesting as multiple shallow ulcers only and the advanced stage as severe strictures. In addition, some investigators reported significant steroid dependency^[1,7]. Hence, further investigation with more collective data is necessary regarding proper CMUSE treatment.

In summary, multiple short strictures and/or shallow ulcers of the small intestine without significant bowel obstruction are characteristic radiologic features of CMUSE on CT, MRI, or SBS. The strictures are short, thin, mainly located in the ileum, and show layered enhancement. Some have recurrent shallow ulceration that may progress to more severe

strictures. Considering that routine clinical practice involves the use of CT and SBS abdominal imaging tools, as well as the risk of endoscopic failure due to strictures, comprehensive understanding regarding the characteristic radiologic features on CT/MRI and SBS would be helpful to diagnose CMUSE and prevent unnecessary surgery. Therefore, under these radiologic findings with relapsing episodes, CMUSE should be considered when assessing patient with recurrent abdominal pain and anemia.

ACKNOWLEDGMENTS

The authors thank the Korean Society of Abdominal Radiology for their support in data collection.

COMMENTS

Background

Cryptogenic multifocal ulcerous stenosing enteritis (CMUSE) is a rare disease characterized by chronic and recurrent ileus resulting from multiple unexplained strictures and superficial ulcerations of the small intestine. At present, CMUSE is diagnosed mainly based on clinical and macroscopic imaging features of the small intestine. However, CMUSE is frequently misdiagnosed as other ulcerative small bowel diseases, because of similar clinical symptoms and nonspecific laboratory findings. Knowledge about characteristic radiologic features of CMUSE and their clinical course would be very useful in differentiating this disease entity from other similar bowel diseases.

Research frontiers

Because this disease is very rare, previous studies about CMUSE have mainly focused on clinical manifestations. Even radiologic information previously reported was relatively simple description and mostly confined to small bowel series (SBS) feature such as the presence or absence of stricture.

Innovations and breakthroughs

Multiple short strictures and/or shallow ulcers of the small intestine without significant bowel obstruction are characteristic radiologic features of CMUSE on computed tomography (CT), MRI, or SBS. The strictures are short, thin, mainly located in the ileum, and show layered enhancement. Some have recurrent shallow ulceration that may progress to more severe strictures. This comprehensive understanding regarding the characteristic radiologic features on CT/MRI and SBS would be helpful to diagnose CMUSE and prevent unnecessary surgery.

Applications

Under characteristic radiologic findings with multiple short-segmental strictures and/or shallow ulcers of the small intestine, CMUSE should be considered when assessing patients with recurrent abdominal pain and anemia.

Terminology

CT enterography or MR enterography is, basically, specialized imaging technique for small bowel evaluation, which needs mandatory requirement of optimal bowel distention, proper suppression of active bowel movement, and multiplanar imaging analysis. Bowel enhancement timing is also important in

these imaging modalities.

Peer-review

It's a well-written manuscript, the characteristic radiologic features on CT/MRI and SBS would be helpful to diagnose CMUSE and prevent unnecessary surgery.

REFERENCES

- 1 **Perlemuter G**, Guillevin L, Legman P, Weiss L, Couturier D, Chaussade S. Cryptogenetic multifocal ulcerous stenosing enteritis: an atypical type of vasculitis or a disease mimicking vasculitis. *Gut* 2001; **48**: 333-338 [PMID: 11171822]
- 2 **Chang DK**, Kim JJ, Choi H, Eun CS, Han DS, Byeon JS, Kim JO. Double balloon endoscopy in small intestinal Crohn's disease and other inflammatory diseases such as cryptogenic multifocal ulcerous stenosing enteritis (CMUSE). *Gastrointest Endosc* 2007; **66**: S96-S98 [PMID: 17709044 DOI: 10.1016/j.gie.2007.01.016]
- 3 **Chung SH**, Park SU, Cheon JH, Kim ER, Byeon JS, Ye BD, Keum B, Shim KN, Jung SA, Kim JO, Jeon SR, Song HJ, Moon JS, Chang DK. Clinical Characteristics and Treatment Outcomes of Cryptogenic Multifocal Ulcerous Stenosing Enteritis in Korea. *Dig Dis Sci* 2015; **60**: 2740-2745 [PMID: 25708899 DOI: 10.1007/s10620-015-3595-y]
- 4 **Kwon SO**, Kim YS, Kim SY, Hong SW, Lee HK, Moon JS. A case of cryptogenic multifocal ulcerous stenosing enteritis: differential diagnosis from Crohn's disease. *J Gastrointest Liver Dis* 2012; **21**: 309-312 [PMID: 23012673]
- 5 **Matsumoto T**, Iida M, Matsui T, Yao T. Chronic nonspecific multiple ulcers of the small intestine: a proposal of the entity from Japanese gastroenterologists to Western enteroscopists. *Gastrointest Endosc* 2007; **66**: S99-S107 [PMID: 17709045 DOI: 10.1016/j.gie.2007.01.004]
- 6 **Park SK**, Yang SK, Park SH, Park SH, Kim JW, Yang DH, Jung KW, Kim KJ, Ye BD, Byeon JS, Myung SJ, Yu CS, Kim JH. Long-term prognosis of the jejunal involvement of Crohn's disease. *J Clin Gastroenterol* 2013; **47**: 400-408 [PMID: 23269310 DOI: 10.1097/MCG.0b013e3182705f9e]
- 7 **Kohoutová D**, Bures J, Tycová V, Bártošová J, Tachecí I, Rejchrt S, Vacek Z, Repák R, Kopáčová M. Severe cryptogenic multifocal ulcerous stenosing enteritis. A report of three cases and review of the literature. *Acta Medica (Hradec Kralove)* 2010; **53**: 25-29 [PMID: 20608229]
- 8 **Perlemuter G**, Chaussade S, Soubrane O, Degoy A, Louvel A, Barbet P, Legman P, Kahan A, Weiss L, Couturier D. Multifocal stenosing ulcerations of the small intestine revealing vasculitis associated with C2 deficiency. *Gastroenterology* 1996; **110**: 1628-1632 [PMID: 8613071]
- 9 **Debray C**, besancon F, hardouin JP, martin E, marche C, khoury K. [Cryptogenetic plurifocal ulcerative stenosing enteritis]. *Arch Mal Appar Dig Mal Nutr* 1964; **53**: 193-206 [PMID: 14140712]
- 10 **Matsumoto T**, Iida M, Matsui T, Yao T, Watanabe H, Yao T, Okabe H. Non-specific multiple ulcers of the small intestine unrelated to non-steroidal anti-inflammatory drugs. *J Clin Pathol* 2004; **57**: 1145-1150 [PMID: 15509673 DOI: 10.1136/jcp.2003.015735]
- 11 **Kim CW**, Yu CS, Yoon YS, Yoon SN, Lim SB, Kim JC. Steroid-refractory cryptogenic multifocal ulcerous stenosing enteritis. *Am J Surg* 2011; **202**: e48-e51 [PMID: 21824599 DOI: 10.1016/j.amjsurg.2010.09.020]

P- Reviewer: Freeman HJ, Wani IA **S- Editor:** Gong ZM
L- Editor: A **E- Editor:** Zhang FF





Published by **Baishideng Publishing Group Inc**
7901 Stoneridge Drive, Suite 501, Pleasanton, CA 94588, USA
Telephone: +1-925-223-8242
Fax: +1-925-223-8243
E-mail: bpgoffice@wjgnet.com
Help Desk: <http://www.f6publishing.com/helpdesk>
<http://www.wjgnet.com>



ISSN 1007-9327

