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1. RAPD Objectives and characteristics: The Revista Andaluza de Patología Digestiva is the official publication of the Andalusian Society of Digestive Pathology (SAPD), which since 2007 has been published in electronic format only, under the name RAPD Online. Its purpose is to disseminate all epidemiological, clinical, basic and sociological aspects of digestive diseases, through the contributions sent to the journal from Andalusia and from the entire scientific community. The official language for the publication of this journal is Spanish, but some contributions may be accepted in the author's original language in English, French or Italian. RAPD Online is published bimonthly, with one of the issues being specially dedicated to the Annual Meeting of the SAPD and the Editorial Board deciding to reserve one or more issues per year for the monographic development of a topic related to the speciality.

All submitted contributions must be original and not be simultaneously under review for publication in another journal. The publication of abstracts or posters is not considered duplicate publication. Manuscripts will be evaluated by expert reviewers, appointed by the editorial board, before being accepted for publication, in a process that will take less than 30 days.

2. RAPD Contents: regular numbers of RAPD Online include defined sections such as:

- Original articles on clinical or basic research.
- Thematic reviews on specific aspects of Gastroenterology.
- Consensus documents.
- Clinical cases.
- Clinical cases with videos or Videoforum.
- Images of the month.
- News and updates on gastroenterology and hepatology.
- Letters to the Editor.

Other contributions that are considered of interest by the Editorial Board, relating to different aspects of clinical practice in the recent past, biographical comments, or other contents of a cultural nature, or related to scientific activities in any territorial area, will be inserted in RAPD Online in sections designed specifically for this purpose.

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4. Writing standard for manuscripts: monographic numbers, thematic reviews, updates and annotated articles will be commissioned by the Editorial Board, but the submission of any of these contributions at the request of an author will be considered by the RAPD Online Management and evaluated with great interest for inclusion in the journal.

All manuscripts will be subject to specific rules, depending on the type of contribution, and to common ethical and legal standards.

A) Specific standard for manuscripts writing

They refer to the recommended length and structure of each type of manuscript. As a basic unit of length for the text, in any of the contributions, a page of 30-31 lines, spaced 1.5 lines apart, with a font size of 12, with 75-80 characters without spaces per line and a total of 400-450 words per page is considered. Texts should be sent spell-checked and in editable format in all their applications (main text, figures, legends or figure captions, tables, graphs, drawings).

Originals: originals can be up to 12 pages long (5,100 words), excluding bibliographical references and captions to figures and tables. It is not advisable to insert more than 10 images, including tables and figures. Colour illustrations and videos will not represent an economic charge for the authors, but the insertion of videos, for technical reasons, will be previously agreed with the editor. However, the editing method of RAPD Online allows, in specific cases, the acceptance of longer manuscripts, or the inclusion of a greater number of images, provided that the characteristics of the material presented so require. It is not advisable to have more than 9 authors, except in the case of collaborative works. In these originals, the first nine participants will be listed at the head of the paper and the rest of the participants will be listed at the end of the first page of the manuscript.

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- a) Introduction and objectives
- b) Material and methods
- c) Results
- d) Conclusions

2° List of abbreviations used in the text.

3° Text: it will include the following sections:

- a) Introduction
- b) Material and methods
- c) Results

- d) Discussion
- e) Conclusions; each of them appropriately headed.

4° Bibliography: according to the specifications established in the group of common standards (See common standards and other supporting documents).

5° Acknowledgements.

6° Figure captions.

7° Tables and figures in text.

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- Main body of the manuscript, containing:

- 1° Structured abstract in Spanish and English. 3-5 key words. The abstract will have a maximum length of 350 words, emphasising the most important aspects of the manuscript.
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- 3° Bibliography: According to the specifications established in the group of common standards (See common standards and other supporting documents).
- 4° Acknowledgements.
- 5° Figure captions
- 6° Tables and Figures in the text.

Consensus documents: texts on Consensus documents are not limited in length in terms of text or images and tables. Exceptionally, the inclusion of videos is allowed. It is not advisable to have more than 10 authors per chapter.

Through the Manuscript Centre, and for the submission of Reviews and Updates, the following information will be required:

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2° Text: Structured according to the criteria of the author(s), for a better understanding of the topic developed.

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4° Acknowledgements

5° Figure captions.

6° Tables and Figures in the text.

Clinical Cases: the manuscripts included in this section will include 1-5 clinical cases, which due to their infrequent or unusual clinical behaviour, or because they provide some diagnostic or therapeutic novelty, deserve to be reported.

The length of the texts in the Clinical Cases section should not exceed 5 pages (2,125 words), excluding bibliographical references and captions to figures and tables, and the number of inserted images should not exceed 5, including tables and figures. However, the RAPD Online editing method allows, in specific cases, the acceptance of longer manuscripts, or the inclusion of a greater number of images, provided that the characteristics of the material presented so require. Colour illustrations and videos will not represent a financial charge for authors, but the insertion of videos, for technical reasons, will be previously agreed with the editor. No more than 5 authors will be admitted, except in specific and reasoned cases.

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- 4° Discussion. To highlight the peculiarities of the case and its consequences.
- 5° Bibliography: According to the specifications established in the group of common standards (See common standards and other supporting documents).
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Clinical Cases with Videos or Videoforum: the manuscripts included in this section will include 1-5 clinical cases, which due to their infrequent or unusual clinical behaviour, or because they provide some diagnostic or therapeutic novelty, deserve to be communicated.

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Videos should be submitted in AVI, MPEG, MP4 OR MOV format, and at a recommended high quality resolution (720p or 1080p). They must not contain personal data of the patients. It is recommended that they be edited to minimise editing time, which should not exceed 10 minutes. If the video includes sound, it must be processed in MP3 format. If the videos to be included are in other formats, please contact the publisher

to verify their validity. They should not exceed 2GB. Through the Manuscript Centre, and for the submission of Clinical Cases - Videoforum, the following information will be required:

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- Main body of the manuscript, containing:

- 1° Structured abstract in Spanish and English. 3-5 key words. The abstract will have a maximum length of 250 words.
- 2° Introduction. To present the clinical problem reported. 3° Description of the clinical case.
- 4° Discussion. To highlight the peculiarities of the case and its consequences.
- 5° Bibliography: According to the specifications established in the group of common standards (See common standards and other supporting documents).
- 6° Acknowledgements. 7° Figure captions.
- 8° Tables and figures in text.
- 9° Videos.

Link tutorial videos: <https://www.sapd.es/videoteca/varios/tutoriales/>

Images of the month: the manuscripts included in this section can take two formats, depending on the authors' preference.

- Format A. Images with educational value: these shall include images of any kind, clinical, radiological, endoscopic, anatomopathological, macro and microscopic, which contribute to postgraduate training and therefore deserve to be shown because of their peculiarity, or because they represent a characteristic example.
- Format B. Key images for a diagnosis: These will include images of any kind, clinical, radiological, endoscopic, anatomopathological, macro and microscopic, together with a summarised clinical history, which will provide the possible final diagnostic resolution. This will be presented in a separate section in the same issue of the journal.

The length of the texts in the Images of the Month section must not exceed 1 page (425 words) in the clinical approach to the image presented and 2 pages (850 words), excluding bibliographical references and captions to figures and tables, in the commentary on the image (Format A) or in the diagnostic resolution of the case (Format B). However, the RAPD Online editing method allows, in specific cases, the acceptance of longer manuscripts, or the inclusion of a greater number of images, provided that the characteristics of the material presented so require. Colour illustrations and videos will not represent a financial charge for authors, but the insertion of videos, for technical reasons, will be previously agreed with the editor. No more than 3 authors will be accepted, except in specific and reasoned cases.

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- 2° Description of the image.
- 3° Comments on the image.

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5° Figure captions.

New developments and updates in gastroenterology and hepatology: this section will be devoted to commenting on the scientific and medical developments that have occurred in recent years in the speciality of Gastroenterology and Hepatology.

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- 5° Declaration on the existence or non-existence of a source of funding for the work, or conflicts of interest.

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1° Text of the manuscript.

2° Bibliography: According to the specifications set out in the common standards group (See common standards and other supporting documents).

B) Common standards and other supporting documents

This refers to the set of mandatory standards, both for uniformity in the presentation of manuscripts and for compliance with current legal regulations. In general, the style of manuscripts should follow the guidelines set out in the Vancouver Agreement of the International Committee of Medical Journal Editors. (<http://www.ICMJE.org>).

Units, generic names and abbreviations:

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There is an aid for the conversion of non-international (non-SI) units into international (SI) units. (<http://www.techexpo.com/techdata/techcntr.html>).

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Kandulsky A, Selgras M, Malferteiner P. Helicobacter pylori infection: A Clinical Overview. Dig Liver Dis 2008; 40:619-626.

Alvarez F, Berg PA, Bianchi FB, Bianchi L, Burroughs AK, Cancado EL, et al. International Autoimmune Hepatitis Group Report: review of criteria for diagnosis of autoimmune hepatitis. J Hepatol 1999; 31:929-938.

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Stamatikos M, Sargedi C, Stefanaki C, Safi oleas C, Matthaopoulou I, Safi oleas M. Anthelmintic treatment: An adjuvant therapeutic strategy against Echinococcus granulosus. Parasitol Int (2009), doi:10.1016/j.parint.2009.01.002

Inadomi JM, Somsouk M, Madanick RD, Thomas JP, Shaheen NJ. A cost-utility analysis of ablative therapy for Barrett's esophagus. Gastroenterology (2009), doi: 10.1053/j.gastro.2009.02.062.

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Gurbulak B, Kabul E, Dural C, Citlak G, Yanar H, Gulluoglu M, et al. Heterotopic pancreas as a leading point for small-bowel intussusception in a pregnant woman. JOP (Online) 2007; 8:584-587.

Fishman DS, Tarnasky PR, Patel SN, Rajman I. Management of pancreaticobiliary disease using a new intra-ductal endoscope: The Texas experience. World J Gastroenterol 2009; 15:1353-1358. Available from: URL: <http://www.wjgnet.com/1007-9327/15/1353.asp>. DOI: <http://dx.doi.org/10.3748/wjg.15.1353>

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TREATMENT OF ZENKER'S DIVERTICULUM USING Z-POEM: SHORT-TERM AND LONG-TERM RESULTS.

Boyero Moreno P¹, Muñoz García-Borrueal M¹, Rosón Rodríguez PJ², Jiménez García VA¹, Rodríguez Téllez M¹

VIRGEN MACARENA UNIVERSITY HOSPITAL. SEVILLE.

VITHAS XANIT BENALMÁDENA HOSPITAL. MALAGA.

Abstract

Introduction: Zenker's diverticulum (ZD) is a condition with a low incidence, typically asymptomatic, and when symptoms are present, the most common are food regurgitation and dysphagia. Therapeutic strategies include surgical and endoscopic techniques. Endoscopic techniques are usually preferred, with flexible endoscopic septotomy (FES) being the most common. Peroral endoscopic myotomy (Z-POEM) offers advantages over previous techniques by providing a more controlled and complete myotomy, reducing the risk of complications and recurrence rates.

Material and methods: This is a unicentric retrospective study including patients treated with Z-POEM between January 2022 and July 2024, with a mean follow-up of 1 year. Baseline clinical variables, treatment response, techniques, and safety were analyzed.

Results: A total of 8 patients were included, with a mean age of 69.35 ± 10.11 years. The average diverticulum size was 3.62 ± 1.4 cm. The technical success rate was 100%, the clinical success rate was 87.5% at 3 months, and 42.86% at 12 months. Three patients required reintervention endoscopically. One case of intraprocedural bleeding was recorded, which was resolved endoscopically, and one patient experienced self-limited dysphonia.

Conclusions: Z-POEM may be an effective and safe therapeutic option for the treatment of ZD. In the case of endoscopic reintervention, another Z-POEM or an alternative technique (FES) can be considered.

Keywords: Zenker's diverticulum, Z-POEM, myotomy, endoscopy.

Paula Boyero Moreno
Virgen Macarena University Hospital
paulaboy98@gmail.com

Boyero Moreno P, Muñoz García-Borrueal M, Rosón Rodríguez PJ, Jiménez García VA, Rodríguez Téllez M.
Treatment of Zenker's diverticulum using Z-POEM: short-term and long-term results.
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List of abbreviations

ZD: Zenker's diverticulum, FES: flexible endoscopic septotomy, Z-POEM: Zenker's diverticulum peroral endoscopic myotomy.

Introduction

Zenker's diverticulum (ZD) is a herniation of the posterior pharyngeal wall that occurs in an area of weakness of the inferior pharyngeal constrictor muscle. The orientation of the muscle fibers creates a triangle, known as Killian's triangle, where the diverticulum forms¹. The incidence is around 0.01-0.11% and is more common in elderly male patients². ZDs are usually asymptomatic, and if they do present clinically, the most common symptoms are regurgitation of undigested food and dysphagia, as well as chronic cough, nausea and vomiting, recurrent pneumonia and bronchoaspiration, and weight loss.

Treatment is indicated when symptoms are present, and therapeutic strategies include surgical and endoscopic techniques. Surgical techniques are performed by transcervical surgery, are more invasive, and have higher complication rates; therefore, endoscopic techniques are usually the treatment of choice.

Endoscopic techniques can be performed with a rigid endoscope or a flexible endoscope. Rigid endoscopic septostomy requires the patient's neck to be placed in hyperextension, requires general anesthesia, and is usually performed by otolaryngologists. Flexible endoscopic septostomy (FES) is the most common technique and usually has fewer complications compared to rigid endoscopy³.

Flexible endoscopic septotomy (FES) is performed with the aid of a diverticuloscope, which is an over-tube with a longer portion at one end that is positioned in the esophagus and a shorter portion that is placed toward the diverticulum, increasing the stability of the position, exposing the septum, and protecting the diverticular and esophageal walls. Once properly positioned, septotomy is performed using kneedle knives or scissor-type scalpels, and the mucosal defect is closed with clips.

Peroral endoscopic myotomy on Zenker's diverticulum, called Z-POEM, is a third-space technique derived from esophageal POEM, and the materials required are similar to those used in that procedure. It consists of performing a mucosotomy on the septum after creating a submucosal wheal with saline solution or other colloidal substances and

subsequent submucosal tunneling on both sides of the septum (diverticular and esophageal). Once the cricopharyngeal muscle is fully exposed, the myotomy is performed, extending up to 2-3 cm distal to the bottom of the diverticulum. The mucosotomy is then closed with clips, and it is recommended to dissect the mucosal flap in large diverticula.

FES offers technical success rates of around 90% and clinical success rates of between 56% and 90%, with recurrence rates of 10-30% and adverse effects of 11%⁴. Z-POEM has been reported to have technical success rates of 90-100%, clinical success rates of around 90%, and recurrence rates of 0-14.7% with complications occurring in 6.7-11% of cases⁴. This latter technique offers advantages over the previous ones as it allows for a more controlled and complete myotomy, reduces the risk of perforation and mediastinitis by preserving the mucosa, and has lower recurrence rates⁵.

The most common complications are emphysema and perforation. However, pneumoperitoneum, pneumomediastinum, aspiration pneumonia, fever, and bleeding may also occur⁵.

According to the classification proposed by Kaminski *et al.*⁶ the size of the diverticulum could guide the choice of the most appropriate therapeutic strategy. In cases of small diverticula, which are those less than 2 cm in size, because they tend to have a short, thick septum with a shallow diverticulum that compromises maneuverability and prevents or hinders the use of the diverticuloscope, the literature suggests that Z-POEM is usually the preferred option. In diverticula between 2 and 5 cm, which are more common and usually have a septum of intermediate thickness, it is suggested that both therapies, Z-POEM and FES, can be considered valid. In large diverticula (greater than 5 cm), which usually have thin and long septa, it is suggested that Z-POEM be performed with an intentional incision (mucosal resection) of the redundant mucosal flap to reduce the remaining mucosal septum and avoid symptoms after treatment.

The objective of our study was to evaluate the efficacy and safety of Z-POEM as an endoscopic treatment for Zenker's diverticulum.

Materials and methods

This is a single-center retrospective study that includes patients treated with Z-POEM between January 2022 and July 2024, with a mean follow-up of 1 year. Baseline clinical variables, treatment response variables, technical variables, and safety variables were analyzed.

Results

A total of 8 patients with Zenker's diverticulum treated with Z-POEM were included, with a mean age of 69.35 ± 10.11 years, 62.5% of whom were women. The mean size of the diverticulum was 3.62 ± 1.4 cm. Seven patients had never received treatment, and one had previously been treated with FES with Ligasure and SB-Knife.

All procedures were performed under orotracheal intubation and with prophylactic antibiotic therapy in the operating room and a short course after the procedure.

With regard to the technique, a Hybrid-knife® type T scalpel was used in all cases except one, which was type I. The diathermy source used was VIO3 (ERBE), with Endocut I 2-2 and Precise Sect 4-4.5 modes. The myotomy extended 2 to 3 cm below the diverticular base, and the mucosotomy was closed with 11 mm hemoclips in all cases, requiring an average of 5 clips.

As immediate complications, there was one case of bleeding during the esophageal submucosal tunnel, which was resolved endoscopically with hemostatic forceps. A case of self-limiting dysphonia was described as a late complication.

Technical success was 100%. Clinical success, assessed as the complete disappearance of symptoms, was 87.5% at 3 months, 75% at 6 months, and 42.86% at 12 months. Three patients required endoscopic reintervention, one of them with a new Z-POEM and the other two with SB-Knife; one of the patients treated with SB-Knife and the other treated with the second Z-POEM responded clinically.

Discussion

The results obtained in our study show a technical success rate of 100%, which is consistent with the high effectiveness of the Z-POEM technique reported in the literature. However, the clinical success observed in our series was lower than that described in previous studies, with complete resolution of symptoms in 87.5% of patients at three months, but with a decrease in this percentage at six and twelve months (75% and 42.86%, respectively). It is important to note that the definition of clinical success varies considerably between different studies. In many studies, clinical success is measured using dysphagia scales and assessing symptom improvement, whereas in our research we defined it as the total absence of symptoms, which is a stricter criterion and may therefore have influenced the results observed.

In addition, the small sample size of our study may have influenced the generalizability of the results. Although clinical efficacy in terms of complete resolution of symptoms is slightly lower than reported in other studies, the Z-POEM technique proved to be safe and effective for the treatment of Zenker's diverticulum, with only one case of bleeding during the procedure and one case of self-limiting dysphonia as a late complication.

According to current literature, Z-POEM is described as a more controlled, less invasive option with results that are not inferior to FES, positioning it as an excellent alternative for the treatment of this pathology. Although ZD is rare, when it presents symptoms, it can significantly affect patients' quality of life, highlighting the importance of effective and safe techniques such as Z-POEM.

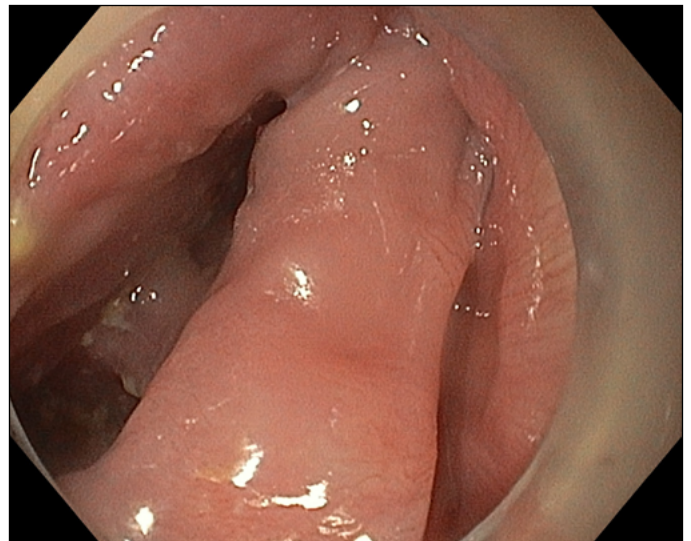


Figure 1. Diverticular septum.

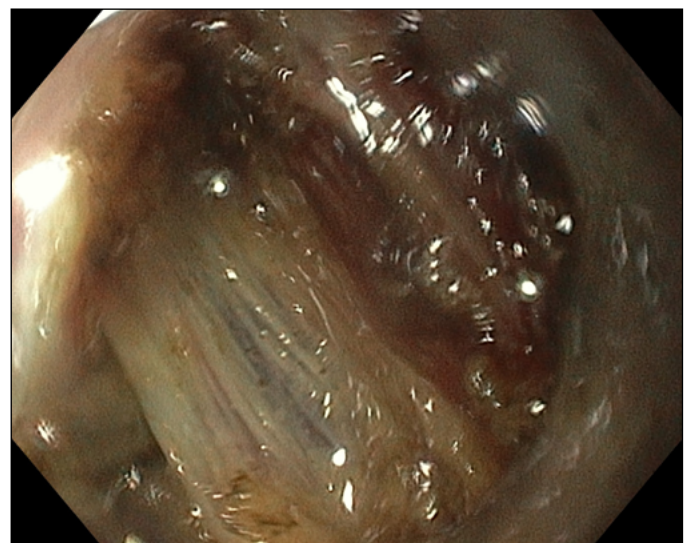


Figure 2. Z-POEM: myotomy.

In our sample, less than 50% of patients required endoscopic reintervention, suggesting that the technique, although effective, does not guarantee a permanent solution

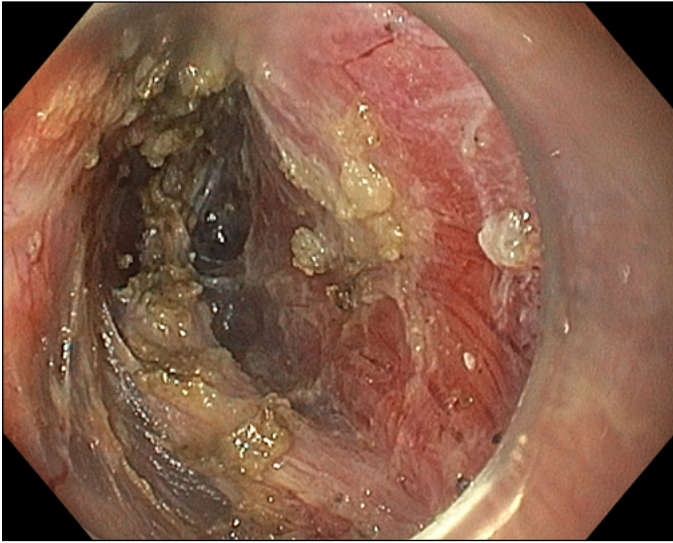


Figure 3. Z-POEM: complete myotomy.

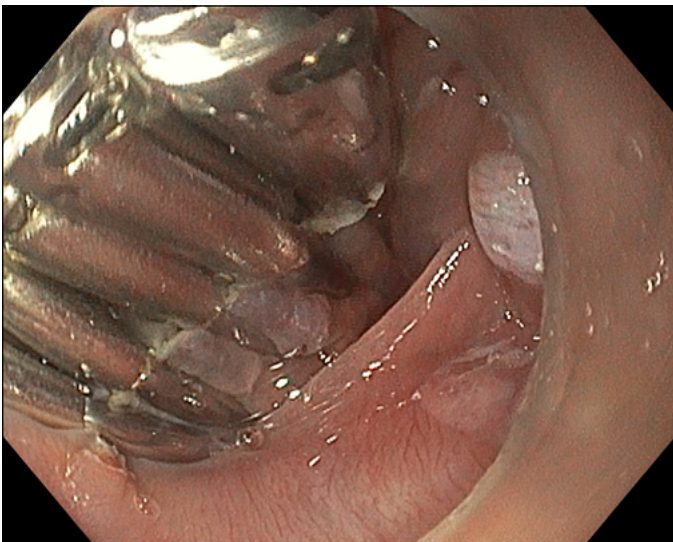


Figure 4. Z-POEM: closure of mucosotomy with clips.

in all cases. This need for reintervention could be related to the nature of the pathology, the variability in the size and characteristics of the diverticulum, factors that do not always allow for a definitive solution in a single procedure. Currently, our group has slightly modified the technique, performing longer tunneling and myotomies, as well as final dissection of the mucosal flap, obtaining more satisfactory results. However, further studies with a larger sample size and long-term follow-up are required to evaluate the durability of the results and

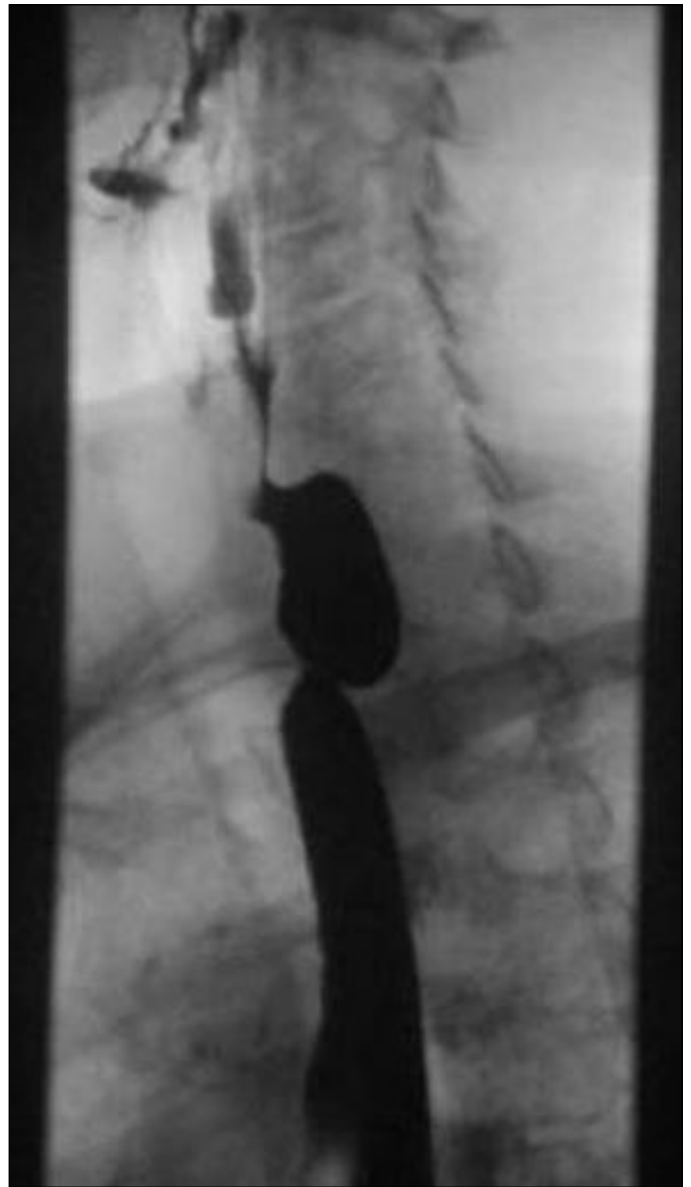


Figure 5. Barium esophagram, lateral view, prior to Z-POEM.

determine predictive factors that may influence the success of the treatment.

Conclusions

Z-POEM may be an effective and safe therapeutic option for the treatment of Zenker's diverticulum.

In the event of endoscopic reoperation, another Z-POEM or an alternative technique (FES) may be performed.



Figure 6. Barium esophagram, lateral view, after Z-POEM.

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BEYOND HELICOBACTER PYLORI: GASTRIC T-CELL LYMPHOMA AS AN UNUSUAL CAUSE OF GASTROINTESTINAL BLEEDING.

Plaza Fernández A, Fernández Carrasco M, Rodríguez Mateu A

Abstract

Primary gastric T-cell lymphoma is a rare and aggressive neoplasm with nonspecific clinical presentation. We report the case of a 75-year-old patient who presented with upper gastrointestinal bleeding secondary to a gastric mass, with histological analysis confirming a high-grade T-cell lymphoma. This case highlights the importance of considering this entity in the differential diagnosis of hemorrhagic gastric lesions.

Keywords: gastric T-cell lymphoma, gastrointestinal bleeding.

Introduction

Primary gastric lymphomas represent a small percentage of gastric neoplasms, and among them, B-cell lymphomas, such as MALT lymphoma and diffuse large B-cell lymphoma, are the most common. In contrast, primary gastric T-cell lymphoma is extremely rare and poses a diagnostic challenge due to its nonspecific clinical presentation. We present a clinical case that presented initially as gastrointestinal bleeding.

Clinical case

A 75-year-old patient with a history of *Helicobacter pylori* infection eradicated years ago and resected urothelial carcinoma currently in remission. The patient had no known autoimmune diseases or viral infections. In the months prior to admission, the patient reported progressive asthenia and weight loss of approximately 6 kg, with no other relevant digestive symptoms. He went to the emergency department due to an episode of hematemesis associated with syncope. Laboratory tests revealed anemia in the transfusion range, and an urgent gastroscopy was performed, which revealed a large ulcerated infiltrative mass in the gastric body at the level of the greater curvature, with irregular and friable edges and active bleeding. Hemostatic control was achieved by sclerotherapy with adrenaline and the application of hemostatic powders. Subsequently, the patient presented a new episode of hematemesis with hemodynamic instability, so a computed tomography (CT) angiogram was performed, which revealed active bleeding from the gastric lesion. Given the instability, urgent surgery was indicated, and a partial gastrectomy was performed with control of the hemorrhagic focus using

CLINICAL CASE

hemostatic sutures. Subsequently, during admission, review of the previous CT images revealed infiltration of the tail of the pancreas and splenic artery, as well as perigastric and retroperitoneal lymphadenopathy, with no distant involvement (Figures 1 and 2). Histological analysis showed diffuse high-grade T-cell lymphoid proliferation, with positivity for CD3 and CD7 and negativity for B markers, confirming the diagnosis of gastric T-cell lymphoma. The patient was referred to Hematology and started treatment with systemic chemotherapy (modified CHOP). The initial course was favorable, with a good clinical response and no new bleeding episodes during short-term follow-up.



Figure 1. CT image showing gastric neoplasm with transmurial extension and loss of the fatty separation plane with the tail of the pancreas and splenic artery due to infiltration of the same.

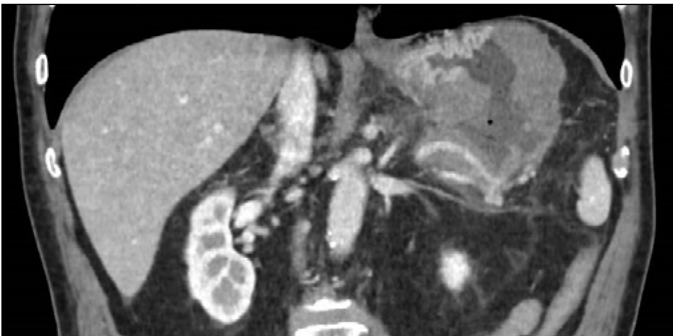


Figure 2. Abdominal CT (coronal section) showing the gastric neoplasm in close contact with the splenic artery and the tail of the pancreas.

Discussion

Primary gastric T-cell lymphoma is an extremely rare neoplasm, with a very low incidence compared to B-cell lymphomas, among which MALT lymphoma is the most common subtype^{1,2}. Unlike these, its association with *Helicobacter pylori* is not well established, although it has been linked to viral infections such as HTLV-1, HBV, HCV, or HIV, and to autoimmune diseases^{2,3}.

The clinical presentation is usually nonspecific, with symptoms such as abdominal pain, weight loss, and asthenia predominating. Gastrointestinal bleeding, as in our case, is an unusual form of onset⁴. Endoscopically, these lymphomas can take many forms: infiltrating masses, ulcers, or thickening of folds⁵.

Diagnosis requires biopsy with immunohistochemical study, with the CD3+ and CD20– phenotype being characteristic. Echoendoscopy allows assessment of transmural involvement and locoregional lymphadenopathy, while CT is essential for assessing distant disease. Treatment is usually based on chemotherapy, with regimens such as CHOP, although the course is often aggressive and the prognosis is poor^{2,4}.

In conclusion, primary gastric T-cell lymphoma is a rare entity with a complex diagnosis. Its atypical presentation, such as gastrointestinal bleeding, requires a high index of suspicion for timely diagnosis and treatment.

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ESOPHAGITIS DISSECANS SUPERFICIALIS IN AN HIV PATIENT WITH A HISTORY OF SUBSTANCE ABUSE

León Sanjuan GF, García Martínez A, Benavente Oyega MA

JEREZ HOSPITAL. CÁDIZ.

Abstract

Esophagitis dissecans superficialis is a rare and often underdiagnosed entity, whose diagnosis relies on characteristic endoscopic findings.

We present the case of a 58-year-old man with stage C3 human immunodeficiency virus (HIV) infection and a history of daily inhaled cannabis, cocaine, and heroin use. He was admitted with pneumococcal pneumonia, oropharyngeal candidiasis, and cachexia. Chest CT revealed thickening of the distal esophageal wall. Endoscopy demonstrated a circumferential desquamative and friable lesion consistent with superficial esophageal dissecans. Biopsies ruled out malignancy and viral infection.

The compatible diagnosis was esophagitis dissecans superficialis, most likely secondary to chronic toxic substance use.

Keywords: esophagitis dissecans superficialis, immunosuppression, toxic substances.

Introduction

Esophagitis dissecans superficialis (EDS) is characterised by circumferential desquamation of the oesophageal mucosa, causing striking lesions that usually have a benign course. We present the case of a patient with HIV infection and substance abuse.

Gloria Francisca León Sanjuan
Jerez Hospital. Jerez de la Frontera, Cádiz.
glorialeon36@gmail.com

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Esophagitis dissecans superficialis in an HIV patient with a history of substance abuse.
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CLINICAL CASE

Clinical case

A 58-year-old male, daily user of cannabis, cocaine, and inhaled heroin. Diagnosed in 2004 with human immunodeficiency virus (HIV) infection, currently stage C3, with regular adherence to treatment for years. History of hepatitis C virus infection, treated and cured with direct-acting antivirals.

He was admitted to the Infectious Diseases Unit for asthenia, pneumococcal pneumonia, oropharyngeal candidiasis, and cachexia. In this context, a chest CT scan was performed, showing evidence of thickening of the oesophageal wall in the distal third. Therefore, an oral endoscopy was requested, which identified a circumferential area of desquamative mucosa with friability on rubbing, compatible with oesophagitis, approximately 32 cm from the dental arch (Figure 1).

Biopsies were taken from the surrounding mucosa for microbiology and anatomopathological study, with no evidence of malignancy or viral induction, as well as negative immunohistochemistry for cytomegalovirus. No other pathological findings were evident on endoscopy.

Given the clinical context, exposure to multiple toxins, and the endoscopic and histological data, a compatible diagnosis was established: Esophagitis dissecans superficialis probably related to damage to the oesophageal mucosa caused by toxins.

Discussion

Superficial dissecting oesophagitis is a rare and often underdiagnosed condition. Its diagnosis is based on the finding of characteristic endoscopic lesions such as diffuse mucosal desquamation and the presence of 'whitish membranes', given that it can often be asymptomatic and histological examination is not specific in this condition^{1,2}.

Although the available literature on this subject is limited, this condition has been associated with factors such as physical and chemical trauma, smoking, autoimmune bullous dermatoses, consumption of toxic substances, and certain drugs such as bisphosphonates, NSAIDs, doxycycline, and apixaban.

Despite its striking endoscopic appearance, it has a benign course, achieving complete healing after removal of the causative agent, if it is a drug or toxin, or through control of the underlying disease³.

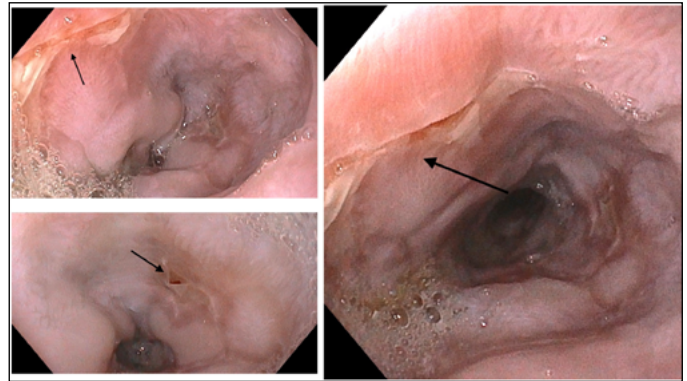


Figure 1. Oesophageal endoscopic image showing areas of desquamated oesophageal mucosa, with the appearance of a partially detached membrane, without ulceration or active bleeding, findings consistent with esophagitis dissecans superficialis.

In the case we present, the findings are consistent with esophagitis dissecans superficialis associated with the consumption of toxic substances such as cocaine and heroin, in the context of possible mucosal fragility due to HIV.

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GASTROINTESTINAL MANIFESTATIONS OF BLUE RUBBER BLEB NEVUS SYNDROME: A CASE REPORT AND LITERATURE REVIEW.

Lorente Martínez MA, Baute Trujillo EA, Moreno Barrueco M, Martín Navas MA, Candel Erenas JM

UNIVERSITY CLINICAL HOSPITAL SAN CECILIO. GRANADA.

Abstract

Blue Rubber Bleb Nevus Syndrome (BRBNS) is a rare disorder characterized by multiple vascular nodules affecting the skin and gastrointestinal tract. Gastrointestinal involvement is highly variable, ranging from chronic anemia and iron deficiency to episodes of acute gastrointestinal bleeding. Endoscopic techniques play a key role in diagnosis, evaluation of disease extent, and therapeutic management of vascular lesions.

Keywords: Blue Rubber Bleb Nevus syndrome, gastrointestinal hemorrhage, vascular malformation.

Introduction

Blue Rubber Bleb Nevus Syndrome (BRBNS) is a rare condition characterized by the development of multiple vascular nodules on the skin and in the gastrointestinal tract. Although congenital forms exist, most cases are secondary to somatic mutations of genes involved in the regulation of angiogenesis, mainly the TEK gene that encodes the angiopoietin-1 receptor. Its mutation causes permanent activation of the receptor, independent of its ligand, and triggers the uncontrolled proliferation of endothelial cells, responsible for the development of multiple venous malformations¹.

Miguel Ángel Lorente Martínez
University Clinical Hospital San Cecilio. Granada.
lorenmtznmiguel@gmail.com

Lorente Martínez MA, Baute Trujillo EA, Moreno Barrueco M, Martín Navas MA, Candel Erenas JM.
Gastrointestinal manifestations of blue rubber bleb nevus syndrome: a case report and literature
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CLINICAL CASE

Many patients are diagnosed during childhood, however, they can debut in adulthood in the form of chronic anemia, iron deficiency, or even acute gastrointestinal bleeding.

Clinical case

We present the case of a 40-year-old male with no relevant family history who was being monitored by dermatology due to multiple blue nevi on his skin and was referred to gastroenterology due to a positive fecal occult blood test. The patient described intermittent melena-like stools and occasional episodes of hematemesis. On examination, the abdominal evaluation showed no pathological findings, and the rectal examination was not consistent with melena.

Laboratory tests showed no anemia, iron deficiency, or elevated urea levels. A study was initiated with gastroscopy, colonoscopy (Figure 1), and subsequently capsule endoscopy (Figures 2 and 3) which revealed multiple nevi associated with angiodysplasias throughout the digestive tract.



Figure 1. Colonic nevus.

Given the presence of multiple venous malformations in different areas, the etiological study was completed with genetic analysis (TEK, VHL, etc.), which did not reveal any pathological alterations. The multidisciplinary clinical evaluation ruled out the main diagnostic alternatives (Von Hippel-Lindau syndrome, Osler-Weber-Rendu disease, etc.), establishing the diagnosis of BRBNS.

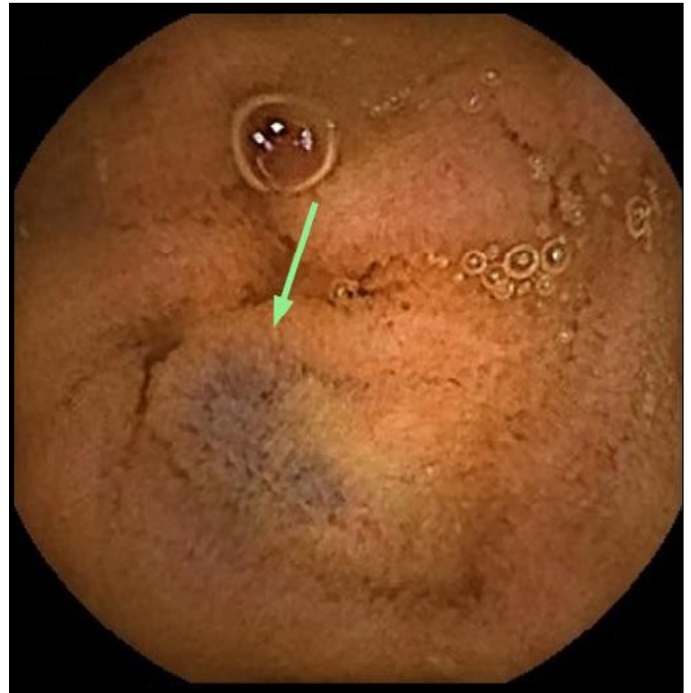


Figure 2. 7 mm nevus at the level of the jejunum.



Figure 3. 10 mm vascular malformation occupying half the circumference, suggestive of a blue nevus.

The patient is currently under close follow-up by the gastroenterology department, maintaining a wait-and-see approach given the absence of clinical and hematimetric repercussions of the gastrointestinal lesions.

Discussion

The diagnosis of BRBNS requires the collaboration of different medical specialties, based on clinical evaluation, endoscopic findings, and genetic studies, although the latter are not essential for diagnosis.

Endoscopic techniques are essential for diagnosing and assessing the extent of the disease and applying therapeutic measures to vascular lesions.

The current lack of scientific evidence makes therapeutic abstention a valid option in mild cases or cases without active bleeding. In cases of bleeding, current studies suggest prioritizing endoscopic therapy (argon plasma coagulation, polypectomy loop excision)² over systemic treatments (sirolimus), which are reserved as a second line due to controversy over their safety and management^{3,4}.

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RECURRENT DIGESTIVE HEMORRHAGE DUE TO FUNDIC VARICES SECONDARY TO LEFT-SIDED PORTAL HYPERTENSION AFTER DISTAL PANCREATECTOMY: SPLENECTOMY AS DEFINITIVE TREATMENT.

Fernandez Carrasco M, Plaza Fernández A, Sánchez Tripiana M, Diéguez Castillo C

TORRECARDENAS UNIVERSITY HOSPITAL. ALMERIA.

Abstract

Extrahepatic portal hypertension (EPH) is defined as an increase in pressure within the portal venous system without the presence of chronic liver disease. Left-sided portal hypertension (LSPH) is a rare form of extrahepatic portal hypertension, typically associated with pancreatic pathology. The formation of collateral circulation and the development of gastric varices are the main associated findings, which can lead to potentially severe upper gastrointestinal hemorrhages (UGIH)¹.

We present the case of a patient who underwent a distal pancreatectomy and subsequently developed recurrent UGIH

due to fundic varices as a consequence of altered splenic circulation.

Keywords: left-sided portal hypertension, gastric varices, splenectomy.

Clinical Case

A 50-year-old woman with a history of distal pancreatectomy for unilocular acinar cell cystadenoma. She consulted for sudden onset of hematemesis, associated with dizziness and arterial hypotension. Initial laboratory

Marta Fernandez Carrasco
Torrecardenas University Hospital. Almeria.
mfcarrasco16@gmail.com

Fernandez Carrasco M, Plaza Fernández A, Sánchez Tripiana M, Diéguez Castillo C.
Recurrent digestive hemorrhage due to fundic varices secondary to left-sided portal
hypertension after distal pancreatectomy: splenectomy as definitive treatment.
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CLINICAL CASE

tests showed anemia (Hb 7.1 g/dL) and normal coagulation parameters. Fluid replacement and red blood cell transfusion were initiated.

Upper gastrointestinal endoscopy (UGE) revealed the presence of variceal conglomerate in the gastric fundus as the source of the symptoms, and endoscopic administration of cyanoacrylate was performed. Computed tomography (CT) was performed, showing post-surgical changes due to corporocoid pancreatectomy, numerous splenic venous collaterals, and varices in the fundus and lesser curvature of the stomach.

The patient was readmitted one month later due to recurrent bleeding with a new episode of hematemesis and anemia. UGE showed persistent fundic varices. The surgical committee decided to perform a splenectomy as definitive treatment. The patient underwent laparoscopic splenectomy without intraoperative complications. Her progress was favorable, with no new episodes of bleeding during the one-year follow-up period and no increased rate of infections.

Discussion

Segmental extrahepatic portal hypertension can be caused by thrombosis or obstruction of the splenic vein (SV), often due to complications of chronic pancreatitis. Other causes include pancreatic carcinoma, pseudocysts, fibrosis, retroperitoneal tumors, and surgeries such as distal pancreatectomy with spleen preservation, which interrupts splenic venous drainage. Venous obstruction diverts the return from the spleen to vessels with lower pressure, generating collateral circulation and perigastric varices. Ligation of the SV without venous reconstruction increases the risk of increased pressure in this area, with the consequent formation of varices¹.

Symptoms generally do not occur and it is detected incidentally. In some cases (4-17%), it can cause gastrointestinal bleeding due to rupture of gastric varices. Diagnosis is based on endoscopy, ultrasound, and CT, which identifies splenic obstruction, splenomegaly, and collateral circulation².

Initial treatment aims to stabilize the patient. Alcohol sclerotherapy, band ligation, and cyanoacrylate are endoscopic options for controlling bleeding. In classic PH, endovascular techniques include transjugular intrahepatic portosystemic shunting (TIPS), which reduces portal pressure, and balloon retrograde transvenous obliteration (BRTO), which is more effective in cardiopulmonary varices³.

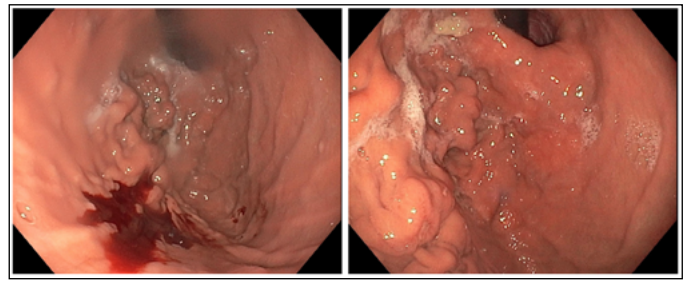


Figure 1. Upper gastrointestinal endoscopy image showing active upper gastrointestinal bleeding secondary to variceal cluster in the gastric fundus.



Figure 2. Abdominal CT image showing splenic collaterals and fundic varices.

The non-indication of TIPS in this context is based on the fact that the procedure is not effective in segmental portal hypertension, since the overall portal pressure gradient is not elevated and the cause of gastric varices is localized obstruction of the splenic vein. TIPS diverts the main portal flow to the systemic circulation, but does not resolve the splenic venous congestion or localized hypertension that causes gastric varices in this scenario¹⁵.

Therefore, splenectomy is considered the definitive treatment, as it eliminates the source of venous congestion and prevents hemorrhagic recurrence in patients with segmental portal hypertension secondary to thrombosis or compression of the splenic vein. In patients at high surgical risk, embolization of the splenic artery may be considered. This latter option is less invasive but carries a risk of complications such as splenic infarction or abscesses³.

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TYPE IVA CHOLEDOCHAL CYST DIAGNOSED IN ADULTHOOD

Cano de la Cruz JD, Sánchez Sánchez MI, Diego Martínez R, Bravo Aranda AM

REGIONAL UNIVERSITY HOSPITAL OF MALAGA. MALAGA.

Abstract

Choledochal cysts are congenital dilations of the intrahepatic and extrahepatic bile ducts, most commonly diagnosed during childhood, with adult diagnosis being less frequent. We present a case of a large type IVA choledochal cyst, incidentally diagnosed in adulthood through imaging studies.

Keywords: choledochal cyst, type IVA, Todani.

Clinical Case

We present the case of a 33-year-old woman admitted for pain in the right hypochondrium and fever. Laboratory tests reveal abnormal liver function (cytolysis and dissociated cholestasis).

An abdominal ultrasound is performed, which shows dilation of the intrahepatic and extrahepatic bile ducts, with a common bile duct measuring up to 15 mm. An abdominal CT scan and MR cholangiography were performed, revealing marked cystic dilatation of the common bile

duct (90 x 57 x 58 mm), associated with cystic dilatation of the intrahepatic bile duct, predominantly on the left side, findings suggestive of a type IVA common bile duct cyst (Figures 1-2). Clinical improvement was observed after administration of analgesia and empirical antibiotic therapy. It was decided to perform a cephalic duodenopancreatectomy (CDP) due to the intrapancreatic component of the lesion and the potential risk of malignancy, which was carried out without complications. After anatomopathological analysis of the surgical specimen (Figure 3), malignant degeneration of the lesion was ruled out.

Discussion

Choledochal cysts are rare entities consisting of a congenital dilation of the bile duct. They are mostly diagnosed in children, although in recent years there has been an increase in their incidence in the adult population. According to Todani's classification¹, the most common (80-90%) are type I (dilatation of the extrahepatic bile duct), while type IVA cysts involve dilatation of both the extrahepatic and intrahepatic bile ducts

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José David Cano de la Cruz
Regional University Hospital of Malaga
jose.davidc@hotmail.com
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Cano de la Cruz JD, Sánchez Sánchez MI, Diego Martínez R, Bravo Aranda AM.
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Figure 1. MR cholangiography. Coronal section - Marked cystic dilatation of the common bile duct (90 x 57 x 58 mm) associated with cystic dilatation of the intrahepatic bile duct, predominantly on the left side.



Figure 2. Abdominal CT. Axial section - Cystic dilatation of the common bile duct associated with cystic dilatation of the intrahepatic bile duct, predominantly on the left side, findings suggestive of a type IVA choledochal cyst according to Todani's classification.

and are extremely rare (1-2%). In 70-80% of cases, they are related to an abnormal pancreatico-biliary junction (AUPJ), predisposing to reflux of pancreatic secretions, with proteolytic activity on the common bile duct².

In adults, they manifest as pain in the right hypochondrium, jaundice, pancreatitis, or cholangitis, with diagnosis primarily based on imaging tests, particularly MR cholangiography due to its high sensitivity (90-100%).

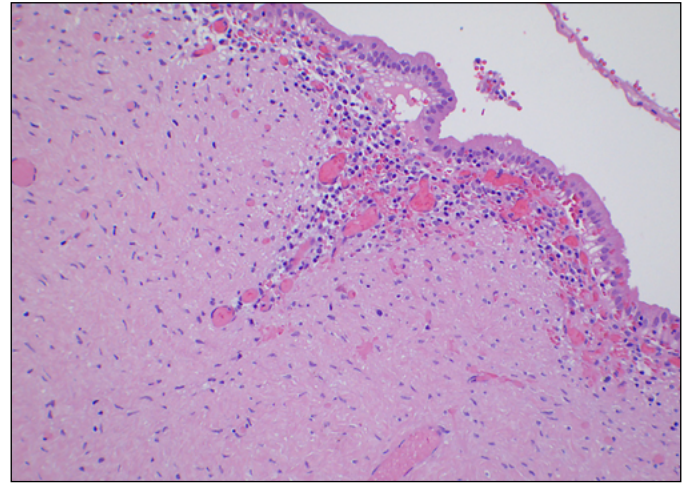


Figure 3. Pathological anatomy. Hematoxylin-eosin staining - Columnar-type cyst lining epithelium, with areas of chronic subepithelial inflammation.

The treatment of choice will always be surgical, with complete removal of the cyst, given its potential risk of malignancy³.

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