

General Surgery V

Editors

Ömer ALABAZ

Alper Sözütek



© Copyright 2026

Printing, broadcasting and sales rights of this book are reserved to Academician Bookstore House Inc. All or parts of this book may not be reproduced, printed or distributed by any means mechanical, electronic, photocopying, magnetic paper and/or other methods without prior written permission of the publisher. Tables, figures and graphics cannot be used for commercial purposes without permission. This book is sold with banderol of Republic of Türkiye Ministry of Culture.

ISBN	Page and Cover Design
978-625-362-004-2	Typesetting and Cover Design by Akademisyen
Book Title	Publisher Certificate Number
General Surgery V	47518
Editors	Printing and Binding
Ömer ALABAZ ORCID iD: 0000-0001-5235-7392 Alper SÖZÜTEK ORCID iD: 0000-0003-1039-9011	Vadi Printingpress
Publishing Coordinator	Bisac Code
Yasin DİLMEN	MED085000
	DOI
	10.37609/akya.4135

Library ID Card

General Surgery V / ed. Ömer Alabaz, Alper Sözütek.
Ankara : Akademisyen Yayınevi Kitabevi, 2026.
153 p. : table. ; 160x235 mm.
Includes References.
ISBN 9786253620042

WARNING

The information contained in this product is only presented as a source for licensed medical workers. It should not be used for any professional medical advice or medical diagnosis. It does not constitute a doctor-patient, therapist-patient and / or any other health-presentation service relationship between the Bookstore and the recipient in any way.

This product is not a synonym or a substitute for professional medical decisions. The Academician Bookstore and its affiliated companies, writers, participants, partners and sponsors are not responsible for injuries and / or damage to humans and devices arising from all applications based on product information.

In the case of prescription of drugs or other chemicals, checking over the current product information for each drug defined by the manufacturer to determine the recommended dose, duration, method and contraindications of the drug is recommended.

It is the physician's own responsibility to determine the optimal treatment an dose for the patient, and to establish a basis for the knowledge and experience of the treating physician about the patient.

The Academician Bookstore is not responsible for any changes to the product, repackaging and customizations made by a third party.

GENERAL DISTRIBUTION

Akademisyen Kitabevi AŞ

Halk Sokak 5 / A Yenışehir / Ankara

Tel: 0312 431 16 33

siparis@akademisyen.com

www.akademisyen.com

PREFACE

Based in Ankara in Turkey, the independent academic publisher, Akademisyen Publishing House, has been publishing books for almost 38 years. As the directors of Akademisyen Publishing House, we are proud to publish more than 4000 books across disciplines so far, especially in Health Sciences. We also publish books in Social Sciences, Educational Sciences, Physical Sciences, and also books on cultural and artistic topics.

Akademisyen Publishing House has recently commenced the process of publishing books in the international arena with the “Scientific Research Book” series in Turkish and English. The publication process of the books, which is expected to take place in March and September every year, will continue with thematic subtitles across disciplines

The books, which are considered as permanent documents of scientific and intellectual studies, are the witnesses of hundreds of years as an information recording platform. As Akademisyen Publishing House, we are strongly committed to working with a professional team. We understand the expectations of the authors, and we tailor our publishing services to meet their needs. We promise each author for the widest distribution of the books that we publish.

We thank all of the authors with whom we collaborated to publish their books across disciplines.

Akademisyen Publishing House Inc.

CONTENTS

Chapter 1	New Strategies for the Treatment of Bile Duct Injuries	1
	<i>Fatih ŞAHİN</i>	
Chapter 2	Which Treatment Method is More Effective for Superficial Squamous Cell Neoplasia Adjacent to Esophageal Varices?	15
	<i>Selim SÖZEN</i>	
Chapter 3	Endoscopic Bariatric Procedures.....	21
	<i>Onur İlkey DİNÇER</i>	
Chapter 4	Change Of Puncture Site İn Right Internal Jugular Central Venous Catheterization – Technical Note.....	41
	<i>Ahmet GÜLTEKİN</i>	
Chapter 5	Decision-Making Processes in General Surgery for Elderly Patients	49
	<i>Yunushan Furkan AYDOĞDU</i>	
Chapter 6	Surgical Treatment of Pulmonary Aspergilloma	61
	<i>Salih Cüneyt AYDEMİR</i>	
Chapter 7	Bile Duct Injuries: Prevention, Diagnosis and Current Management	73
	<i>Muhammet Fatih KEYİF</i>	
Chapter 8	Granulomatous Mastitis.....	83
	<i>Ferdi BOLAT</i>	
Chapter 9	Differences Between Right-Sided And Left-Sided Colon Cancers	97
	<i>Can AYDIN</i> <i>Çağıl KARAEVLİ</i> <i>Sami AÇAR</i>	
Chapter 10	Chylothorax	127
	<i>Mehmet AĞAR</i>	

AUTHORS

Assist. Prof. Muhammet Fatih KEYİF

Department of General Surgery, Bolu Abant İzzet Baysal University Faculty of Medicine

Assoc. Prof. Dr. Sami AÇAR

Department of General Surgery, Faculty of Medicine, Tekirdağ Namık Kemal University

Assist. Prof. Mehmet AĞAR

Firat University, Faculty of Medicine, Thoracic Surgery Clinic

Dr. Salih Cüneyt AYDEMİR

Thoracic Surgeon, Beylikdüzü State Hospital

Res. Assist. Dr. Can AYDIN

Department of General Surgery, Faculty of Medicine, Tekirdağ Namık Kemal University

MD Yunushan Furkan AYDOĞDU

General Surgeon, Department of General Surgery, Ankara Training and Research Hospital

Assist. Prof. Ferdi BOLAT

Department of General Surgery, Faculty of Medicine, Bolu Abant İzzet Baysal University

Dr. Onur İlkay DİNÇER

Antalya Training and Research Hospital

Assoc. Prof. Ahmet GÜLTEKİN

Tekirdağ Namık Kemal University Faculty of Medicine, Department of Anesthesiology and Reanimation

Op. Dr. Çağıl KARAEVLİ

Department of General, Torbalı State Hospital, İzmir, Türkiye

Assoc. Prof. Dr. Selim SÖZEN

Sözen Surgery Clinic

Op. Dr. Fatih ŞAHİN

Hakkari Yuksekova State Hospital

Chapter 1

NEW STRATEGIES FOR THE TREATMENT OF BILE DUCT INJURIES

Fatih ŞAHİN¹

INTRODUCTION

Laparoscopic cholecystectomy has maintained its status as the gold standard surgical approach for the management of gallbladder pathologies since its widespread adoption in the early 1990s. As a minimally invasive technique associated with shorter hospital stays, reduced postoperative pain, and faster recovery, it has become the most frequently performed elective abdominal procedure in the United States, with annual case volumes exceeding 750,000 interventions (1). Despite its well-established benefits, bile duct injury (BDI) remains one of the most serious and feared complications associated with cholecystectomy, given its potential to result in significant morbidity, long-term biliary dysfunction, and decreased quality of life.

The critical nature of BDI in surgical practice is underscored by large-scale epidemiological data. Notably, two major administrative databases encompassing over 850,000 laparoscopic cholecystectomy procedures conducted between 2005 and 2014 in the states of New York and California reported an overall incidence of BDI ranging between 0.1% and 0.2% (3,4). Although this incidence may appear low, the sheer volume of procedures performed annually implies that a substantial number of patients are affected by this complication each year, with important clinical, emotional, and economic consequences.

In this context, the prevention of bile duct injuries represents the most desirable and cost-effective strategy. Preventive efforts rely on adherence to meticulous surgical technique, routine use of intraoperative imaging when indicated, and strict application of anatomical dissection principles such as the critical view of safety. Nevertheless, despite the best preventive strategies, BDIs may still occur due to anatomical variations, inflammation, or technical errors.

¹ Op. Dr., Hakkari Yuksekova State Hospital, dr.fshahin@gmail.com, ORCID iD: 0000-0002-6505-5884

invasive endoscopic interventions, regenerative medicine applications, and next-generation imaging technologies. This evolution is driven by a convergence of evidence underscoring the diagnostic and therapeutic utility of ERCP and MRCP, alongside groundbreaking progress in tissue engineering and 3D bioprinting, which collectively offer novel, potentially paradigm-shifting solutions for bile duct repair and functional restoration.

The integration of these innovative strategies into clinical practice heralds a promising future for the management of biliary tract injuries—enhancing diagnostic precision, reducing procedural morbidity, and offering biologically based, patient-specific treatment modalities. The utilization of bioengineered scaffolds, stem cell therapies, and bioprinted constructs stands to transform reconstructive options, while ongoing refinements in MRCP technology continue to improve non-invasive diagnostic capabilities, reducing dependency on invasive diagnostics and associated risks.

Nonetheless, a number of significant barriers remain. These include the optimization and standardization of regenerative therapies, mitigation of ERCP-related complications such as post-ERCP pancreatitis, and improving accessibility and cost-effectiveness of advanced imaging modalities in diverse healthcare settings. Future research priorities must include the development of evidence-based clinical algorithms, enhancement of biomaterial-host integration, and the rigorous evaluation of long-term clinical outcomes through prospective, controlled trials.

In conclusion, the field of biliary tract injury management is experiencing a transformative shift, driven by the synergistic interplay of endoscopic innovation, regenerative medicine, and advanced imaging. The convergence of these domains holds the potential to revolutionize current standards of care—providing safer, more effective, and individualized treatment pathways for patients affected by these complex and often debilitating injuries. With continued interdisciplinary collaboration and translational research, this evolving landscape promises to redefine the future of hepatobiliary surgery.

REFERENCES

- Jensen, Sofie Anne-Marie Skovbo, et al. “Long-term mortality and intestinal obstruction after laparoscopic cholecystectomy: A systematic review and meta-analysis.” *International Journal of Surgery* 105 (2022): 106841.
- Pisano, Michele, et al. “2020 World Society of Emergency Surgery updated guidelines for the diagnosis and treatment of acute calculus cholecystitis.” *World journal of emergency surgery* 15 (2020): 1-26.

General Surgery V

- Halbert, Caitlin, et al. "Long-term outcomes of patients with common bile duct injury following laparoscopic cholecystectomy." *Surgical endoscopy* 30 (2016): 4294-4299.
- Fong, Zhi Ven, et al. "Diminished survival in patients with bile leak and ductal injury: management strategy and outcomes." *Journal of the American College of Surgeons* 226.4 (2018): 568-576.
- Nakazato, Tetsuya, et al. "Improving attainment of the critical view of safety during laparoscopic cholecystectomy." *Surgical endoscopy* 34 (2020): 4115-4123.
- Michael Brunt, L., et al. "Safe cholecystectomy multi-society practice guideline and state-of-the-art consensus conference on prevention of bile duct injury during cholecystectomy." *Surgical endoscopy* 34 (2020): 2827-2855.
- Wu, Yuhsin V., and David C. Linehan. "Bile duct injuries in the era of laparoscopic cholecystectomies." *Surgical Clinics* 90.4 (2010): 787-802.
- Symeonidis, Dimitrios, et al. "BILE: A literature review based novel clinical classification and treatment algorithm of iatrogenic bile duct injuries." *Journal of Clinical Medicine* 12.11 (2023): 3786.
- Georgiou, Konstantinos, et al. "Intraoperative cholangiography 2020: Quo vadis? A systematic review of the literature." *Hepatobiliary & Pancreatic Diseases International* 21.2 (2022): 145-153.
- Brunt, L. Michael. "Should we utilize routine cholangiography?." *Advances in Surgery* 56.1 (2022): 37-48.
- Desai, Aakash, et al. "Clinical efficacy, timing, and outcomes of ERCP for management of bile duct leaks: a nationwide cohort study." *Endoscopy international open* 9.02 (2021): E247-E252.
- Griffin, Nyree, Geoff Charles-Edwards, and Lee Alexander Grant. "Magnetic resonance cholangiopancreatography: the ABC of MRCP." *Insights into imaging* 3 (2012): 11-21.
- Hekimoglu, Koray, et al. "MRCP vs ERCP in the evaluation of biliary pathologies: review of current literature." *Journal of digestive diseases* 9.3 (2008): 162-169.
- Zong, Chen, et al. "A novel therapy strategy for bile duct repair using tissue engineering technique: PCL/PLGA bilayered scaffold with hMSCs." *Journal of tissue engineering and regenerative medicine* 11.4 (2017): 966-976.
- Sanyal, Arka, and Sourabh Ghosh. "3D bioprinting strategies for recapitulation of hepatic structure and function in bioengineered liver: a State-of-the-art review." *Current Opinion in Biomedical Engineering* 30 (2024): 100526.
- Li, Peilin, et al. "Three-dimensional human bile duct formation from chemically induced human liver progenitor cells." *Frontiers in Bioengineering and Biotechnology* 11 (2023): 1249769.

Chapter 2

WHICH TREATMENT METHOD IS MORE EFFECTIVE FOR SUPERFICIAL SQUAMOUS CELL NEOPLASIA ADJACENT TO ESOPHAGEAL VARICES?

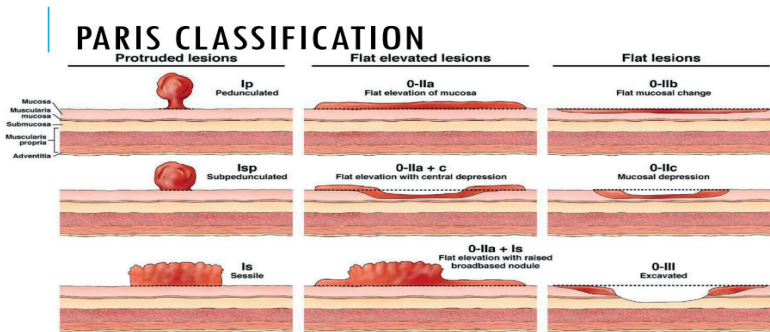
Selim SÖZEN¹

INTRODUCTION

Endoscopic submucosal dissection(ESD) is a minimally invasive technique. ESD has gradually become accepted as an alternative treatment of superficial esophageal squamous-cell neoplasia (SESCN) beside esophagectomy(1). There are American Joint Committee on Cancer (AJCC) system and the Union for International Cancer Control (UICC) for SESCEN staging guidelines(2). Another classification system is the Paris classification for superficial tumors, used in standard endoscopic terminology.(Figure 1)

DISCUSSION

Neoplastic lesions, which are the abnormal growth of tissues, in the gastrointestinal tract are called “superficial” (including dysplasia, which is the abnormal growth of cells in an organ, and adenoma, which is non-cancerous tumours).



¹ Assoc. Prof. Dr, Sözen Surgery Clinic, selimsozen63@yahoo.com, ORCID iD: 0000-0003-2006-9198

EVL and EIS are commonly applied methods for endoscopic treatment of EV, but only varicose veins with diameters >0.3 cm can be targeted by these two methods(31). APC can effectively seal blood vessels with diameters ≤0.3 cm, thus alleviating the recurrence of varicose veins(32). Sequential APC treatment after EIS is safe and can prevent the recurrence of esophageal varices(33,34). Sequential treatment of esophageal solitary venous dilatation, endoscopic drainage sclerotherapy, and ESD are most effective treatments for esophageal varices with early esophageal cancer(35)

CONCLUSION

In conclusion, Esophageal cancer with esophageal varices could be treated endoscopically safely and effectively(36). APC and ESD are useful treatments in the early superficial cancers with esophageal varices.

REFERENCES

- An W, Liu MY, Zhang J, Cui YP, Gao J, Wang LP, Chen Y, Yang LX, Chen HZ, Jin H, Liu F, Chen J, Li ZS, Wang LW, Shi XG, Sun C. Endoscopic submucosal dissection versus esophagectomy for early esophageal squamous cell carcinoma with tumor invasion to different depths. *Am J Cancer Res.* 2020 Sep 1;10(9):2977-2992. PMID: 33042630; PMCID: PMC7539777.
- Zaidi N, Kelly RJ. The management of localized esophageal squamous cell carcinoma: Western approach. *Chin Clin Oncol.* 2017 Oct;6(5):46. doi: 10.21037/cco.2017.07.07. PMID: 29129086.
- The Paris endoscopic classification of superficial neoplastic lesions: esophagus, stomach, and colon: November 30 to December 1, 2002.
- Mony S, Hu B, Joseph A, et al. Clinical outcomes of endoscopic submucosal dissection for superficial esophageal neoplasia in close proximity to esophageal varices: a multicenter international experience. *Endoscopy.* 2024 Feb;56(2):119-124. DOI: 10.1055/a-2159-2557. PMID: 376116.
- Miller AM, McGill D, Bassett ML. Anticoagulant therapy, anti-platelet agents and gastrointestinal endoscopy. *J Gastroenterol Hepatol* 1999; **14**: 109-113.
- Takahashi H, Arimura Y, Okahara S, Uchida S, Ishigaki S, Tsukagoshi H, Shinomura Y, Hosokawa M. Risk of perforation during dilation for esophageal strictures after endoscopic resection in patients with early squamous cell carcinoma. *Endoscopy* 2011; **43**: 184-189.
- Teoh AY, Chiu PW, Wong SK, Sung JJ, Lau JY, Ng EK. Difficulties and outcomes in starting endoscopic submucosal dissection. *Surg Endosc* 2010; **24**: 1049-1054.
- Akiyama T, Abe Y, Iida H, Endo H, Hosono K, Yoneda K, Takahashi H, Inamori M, Ryo A, Yamanaka S, Inayama Y, Nakajima A. Endoscopic therapy using an endoscopic variceal ligation for minute cancer of the esophagogastric junction complicated with esophageal varices: a case report. *J Med Case Rep* 2010; **4**: 149.
- Wang J, Liu Y, He S, Zhang Y, Dou L, Wang G. Endoscopic submucosal dissection for early esophageal squamous cell carcinoma with esophageal-gastric fundal varices caused by liver cirrhosis: a case report. *Transl Cancer Res.* 2022 Jul;11(7):2433-2437. doi: 10.21037/tcr-21-2624. PMID: 35966322; PMCID: PMC9372251.
- Oyama T, Inoue H, Arima M, et al. Prediction of the invasion depth of superficial squamous cell carcinoma based on microvessel morphology: magnifying endoscopic classification of the Japan Esophageal Society. *Esophagus* 2017; **14**: 105-12. [Crossref].

- Kubota Y, Tanabe S, Ishido K, Yano T, Wada T, Watanabe A, Azuma M, Katada C, Koizumi W. Usefulness of argon plasma coagulation for superficial esophageal squamous cell neoplasia in patients at high risk or with limited endoscopic resectability. *Turk J Gastroenterol* 2020 Jul;31(7):529-537. PMID: 32897227; PMCID: PMC7480201.
- Malick KJ. Clinical applications of argon plasma coagulation in endoscopy. *Gastroenterol Nurs*. 2006; **29**: 386-91.
- Song Y, Feng Y, Sun LH, Zhang BJ, Yao HJ, Qiao JG, Zhang SF, Zhang P, Liu B. Role of argon plasma coagulation in treatment of esophageal varices. *World J Clin Cases*. 2021 Jan 26;9(3):521-527. doi: 10.12998/wjcc.v9.i3.521. PMID: 33553390; PMCID: PMC7829739.
- Zhang X, Feng, X, Linghu E. Argon Plasma Coagulation Is Appropriate for the Treatment of LD0.3Rf Type Esophageal Varices: 2799. *American Journal of Gastroenterology*.2018 Oct; 113():1552-1553.
- Olmos JA**, Marcolongo M, Pogorelsky V, Varela E, Dávalos JR. Argon plasma coagulation for prevention of recurrent bleeding from GI angiodysplasias. *Gastrointest Endosc* 2004; **60**: 881-886.
- van Vilsteren FG**, Pouw RE, Seewald S, Alvarez Herrero L, Sondermeijer CM, Visser M, Ten Kate FJ, Yu Kim Teng KC, Soehendra N, Rösch T, Weusten BL, Bergman JJ. Stepwise radical endoscopic resection versus radiofrequency ablation for Barrett's oesophagus with high-grade dysplasia or early cancer: a multicentre randomised trial. *Gut* 2011; **60**: 765-773.
- Sagawa T**, Takayama T, Oku T, Hayashi T, Ota H, Okamoto T, Muramatsu H, Katsuki S, Sato Y, Kato J, Niitsu Y. Argon plasma coagulation for successful treatment of early gastric cancer with intramucosal invasion. *Gut* 2003; **52**: 334-339.
- Kitamura T**, Tanabe S, Koizumi W, Mitomi H, Saigenji K Argon plasma coagulation for early gastric cancer: technique and outcome. *Gastrointest Endosc* 2006; **63**: 48-54.
- Nomura T, Miyashita M, Makino H, Okawa K, Katsuta M, Tajiri T. Argon plasma coagulation for the treatment of superficial esophageal carcinoma. *J Nippon Med Sch* 2007; **74**: 163-7. [Crossref].
- Min BH, Kim ER, Lee JH, et al. Feasibility and efficacy of argon plasma coagulation for early esophageal squamous cell neoplasia. *Endoscopy* 2013; **45**: 575-8. [Crossref].
- Tahara K, Tanabe S, Ishido K, et al. Argon plasma coagulation for superficial Esophageal squamous-cell carcinoma in high-risk patients. *World J Gastroenterol* 2012; **18**: 5412-7. [Crossref].
- Oyama T, Tomori A, Hotta K, et al. Endoscopic submucosal dissection of early esophageal cancer. *Clin Gastroenterol Hepatol* 2005; **3**: 67-70. [Crossref].
- Ono S, Fujishiro M, Niimi K, Goto O, Yamamichi N, Omata M. Long-term outcomes of endoscopic submucosal dissection for superficial esophageal squamous cell neoplasms. *Gastrointest Endosc* 2009; **70**: 860-6. [Crossref].
- Grund KE, Zindel C, Farin G. Argon plasma coagulation through a flexible endoscope. Evaluation of a new therapeutic method after 1606 uses. *Dtsch Med Wochenschr* 1997; **122**: 432-8.
- Higuchi K, Tanabe S, Azuma M, et al. A phase II study of endoscopic submucosal dissection for superficial esophageal neoplasms (KDOG 0901). *Gastrointest Endosc* 2013 ; **78**: 704-10. [Crossref].
- Ishihara R, Ishii H, Uedo N, et al. Comparison of EMR and endoscopic submucosal dissection for en bloc resection of early esophageal cancers in Japan. *Gastrointest Endosc* 2008; **68**: 1066-72. [Crossref].
- Takahashi H, Arimura Y, Masao H, et al. Endoscopic submucosal dissection is superior to conventional endoscopic resection as a curative treatment for early squamous-cell carcinoma of the esophagus (with video). *Gastrointest Endosc* 2010; **72**: 255-4. [Crossref].
- Repici A, Hassan C, Carlino A, et al. Endoscopic submucosal dissection in patients with early esophageal squamous-cell carcinoma: results from a prospective Western series. *Gastrointest Endosc* 2010; **71**: 715-21. [Crossref].
- Tsujii Y, Nishida T, Nishiyama O, et al. Clinical outcomes of endoscopic submucosal dissection for superficial esophageal neoplasms: a multicenter retrospective cohort study. *Endoscopy* 2015; **47**: 775- 83. [Crossref].

- The Japan Esophageal Society. Guidelines for diagnosis and treatment of carcinoma of the esophagus 2017. Tokyo: Kanehara Co Ltd.
- Yang Q**, Zhao Y, Chen X, Tang P, Li L, Zhao J, Han Y, Wu D, An L, Zhang B, Zhou X, Liu L, Chi YW. Association Between Vein Diameters, Reflux Characteristics, and Clinical Severity in Patients with Chronic Venous Insufficiency in Northwest China. *J Vasc Surg Venous Lymphat Disord* 2020 epub ahead of print [PMID: 32730997 DOI: 10.1016/j.jvsv.2020.07.00].
- Fuccio L**, Zagari RM, Serrani M, Eusebi LH, Grilli D, Cennamo V, Laterza L, Asioli S, Ceroni L, Bazzoli F. Endoscopic argon plasma coagulation for the treatment of gastric antral vascular ectasia-related bleeding in patients with liver cirrhosis. *Digestion* 2009; **79**: 143-150 [PMID: 19329853 DOI: 10.1159/000210087].
- Deguchi H**, Kato J, Maeda Y, Moribata K, Shingaki N, Niwa T, Inoue I, Maekita T, Iguchi M, Tamai H, Ichinose M. Argon plasma coagulation is effective for prevention of recurrent esophageal varices after endoscopic injection sclerotherapy: Single-center case-control study. *Dig Endosc* 2016; **28**: 42-49 [PMID: 26295791 DOI: 10.1111/den.12538].
- Matsui S**, Kudo M, Nakaoka R, Shiomi M, Kawasaki T. Comparison of argon plasma coagulation and paravariceal injection sclerotherapy with 1% polidocanol in mucosa-fibrosing therapy for esophageal varices. *J Gastroenterol* 2004; **39**: 397-399 [PMID: 15168254 DOI: 10.1007/s00535-003-1309-2].
- Xu L, Chen SS, Yang C, Cao HJ. Successful endoscopic treatment of superficial esophageal cancer in a patient with esophageal variceal bleeding: A case report. *World J Clin Cases*. 2024 Sep 26;12(27):6105-6110. doi: 10.12998/wjcc.v12.i27.6105. PMID: 39328865; PMCID: PMC11326098.
- Nakai T, Yoshizaki T, Tanaka S, Yamamoto Y, Sako T, Kitamura Y, Ose T, Ishida T, Ikeda A, Ariyoshi R, Iwatate M, Kawara F, Takao T, Morita Y, Toyonaga T, Kodama Y. Safety and efficacy of endoscopic submucosal dissection for superficial esophageal cancer with esophageal varices. *Esophagus*. 2023 Jul;20(3):515-523. doi: 10.1007/s10388-023-01001-3. Epub 2023 Apr 15. PMID: 37060531.

Chapter 3

ENDOSCOPIC BARIATRIC PROCEDURES

Onur İlkey DİNÇER¹

INTRODUCTION

Obesity is a chronic disease that can be considered a global pandemic with an increasing prevalence. It has been reported that the prevalence of obesity has doubled in 70 countries, and obesity has increased in almost all countries since 1980 (1). Although many different treatment methods such as diet, medication and psychotherapy have been tried in obesity, it has been shown that long-term weight loss cannot be maintained, especially in the morbidly obese patient group, and the psychological and metabolic negative effects of unsuccessful weight loss experiences increase exponentially (2). Studies continue to prove that surgery is the most effective treatment for morbid obesity. Of course, as with any surgical procedure, bariatric procedures are associated with certain complications. Complications of bariatric surgery include a wide range of complications, including venous thromboembolism, anastomotic leaks, infection, bleeding, intestinal obstruction, vitamin deficiencies, and hernias. Although bariatric surgical procedures carry an acceptable risk of mortality and morbidity, some patients are unwilling to take these risks and seek additional treatment options (3). Endoluminal endoscopic procedures are now being used as primary procedures in obese patients with a low body mass index (BMI) (BMI 30-35 kg/m²), as well as in morbidly obese patients who meet traditional bariatric surgery criteria. They are also being used as bridge therapy in superobese patients (3). As in bariatric surgery, the main mechanisms in endoscopic bariatric procedures are of two types: restrictive, which focuses on volume, such as the gastric balloon, and malabsorptive, such as stent and aspiration technologies. Endoscopic primary bariatric procedures will be examined under two headings: gastric interventions and small bowel interventions.

¹ Dr. ,Antalya Training and Research Hospital, onurdncr@hotmail.com, ORCID iD: 0000-0001-8956-351X

DOI: 10.37609/akya.4135.c6719

In a pilot study of 12 patients, the average excess weight loss over 12 weeks was 23.6%. Two patients reported premature device removal due to epigastric pain, nausea, and vomiting. Inflammation was found at the duodenal fixation site in all patients (70). A recent meta-analysis found a 12.6% greater excess weight loss compared with dietary modification, but it failed to significantly impact fasting blood glucose.

CONCLUSION

Obesity will be one of the biggest problems to be overcome worldwide in the coming years due to changing eating habits and lifestyles. Protecting patients from the risks of surgical complications by developing more minimally invasive methods alongside surgical treatment options is a priority. With advancing technology and developing industry, the quality of endoscopic devices and the types and complexity of procedures that can be performed are constantly increasing. Numerous products are currently under development and in the experimental phase in the literature. It is expected that more prominent interventions, due to their minimal intervention and minimal complication risk, will emerge in the coming years.

REFERENCES

1. GBD 2015 Obesity Collaborators, Afshin A et. al. Health Effects of Overweight and Obesity in 195 Countries over 25 Years. *N Engl J Med*. 2017 Jul 6;377(1):13-27. doi: 10.1056/NEJ-Moa1614362.
2. Crujeiras AB, Goyenechea E, Abete I, et al. Weight regain after a diet-induced loss is predicted by higher 94 baseline leptin and lower ghrelin plasma levels. *Clin Endocrinol Metab* 2010; 95:5037-44. doi: 10.1210/jc.2009-2566.
3. Scerbo, M.H., Felinski, M.M., Bajwa, K.S., et al. (2020). Endoscopic Primary Bariatric Procedures. In: Nguyen, N., Brethauer, S., Morton, J., Ponce, J., Rosenthal, R. (eds) *The ASMBS Textbook of Bariatric Surgery*. Springer, Cham. https://doi.org/10.1007/978-3-030-27021-6_36.
4. DeBaakey M, Ochsner A. Bezoars and concretions: a comprehensive review of the literature with an analysis of 303 collected cases and a presentation of 8 additional cases. *Surgery*. 1938;4:934-63.
5. Gleysteen JJ. A history of intragastric balloons. *Surg Obes Relat Dis*. 2016;12:430-5. doi: 10.1016/j.soard.2015.10.074.
6. Benjamin SB, Maher KA, Cattau EL Jr, et al. Double-blind controlled trial of Garren-Edwards gastric bubble: an adjunctive treatment for exogenous obesity. *Gastroenterol*. 1988;95:581-8. doi: 10.1016/s0016-5085(88)80001-5.
7. Lindor KD, Hughes RW Jr, Ilstrup DM, et al. Intragastric balloons in comparison with standard therapy for obesity—a random double-blind trial. *Mayo Clin Proc* 1987;62:992-6. 6. Hogan RB, Johnston JH, Long BW, et al. A double-blind, randomized, sham-controlled trial of the gastric bubble for obesity. *Gastrointest Endosc* 1989;35:381-5.
8. Ramhanadany EM, Fowler J, Baird IM. Effect of gastric balloon versus sham procedure on weight loss in obese subjects. *Gut*. 1989;30:1054-7.

9. Galloro G, De Palma GD, Catanzano C, et al. Preliminary endoscopic technical report of a new silicone intragastric balloon in the treatment of morbid obesity. *Obes Surg.* 1999;9:68–71. doi: 10.1381/09608929976553827.
10. Ponce J, Lutfi RE. Intra-gastric Balloon Therapy. Nguyen NT, Brethauer A, Morton JM, Ponce J, Rosenthal RJ (eds.) *The ASMBS Textbook of Bariatric Surgery*. 2nd ed. Switzerland: Springer; 2020. p. 375–381.
11. Abu Dayyeh BK, Maselli DB, Rapaka B, et al. Adjustable intragastric balloon for treatment of obesity: a multicentre, open-label, randomised clinical trial. *Lancet.* 2021 Nov 27;398(10315):1965–1973. doi: 10.1016/S0140-6736(21)02649-0.
12. Neto MG, Silva LB, Grecco E, et al. Brazilian intragastric balloon consensus statement (BIBC): practical guidelines based on experience of over 40,000 cases. *Surg Obes Relat Dis.* 2018;14(2):151–159. doi: 10.1016/j.soard.2017.09.528.
13. Mion F, Napoleon B, Roman S, et al. Effects of intragastric balloon on gastric emptying and plasma ghrelin levels in non-morbid obese patients. *Obes Surg.* 2005;15(4):510–516. doi: 10.1381/0960892053723411.
14. Su HJ, Kao CH, Chen WC, et al. Effect of intragastric balloon on gastric emptying time in humans for weight control. *Clin Nucl Med.* 2013;38:863–868. doi: 10.1097/RLU.0000000000000224.
15. Courcoulas A, Abu Dayyeh BK, Eaton L, et al. Intra-gastric balloon as an adjunct to lifestyle intervention: a randomized controlled trial. *Int J Obes.* 2017;41(3):427–433. doi: 10.1038/ijo.2016.229.
16. Silva LB, Neto MG. Intra-gastric balloon. *Minim Invasive Ther Allied Technol.* 2022 Apr;31(4):505–514. doi: 10.1080/13645706.2021.1874420.
17. Moore RL, Seger MV, Garber SM, et al. Clinical safety and effectiveness of a swallowable gas-filled intragastric balloon system for weight loss: consecutively treated patients in the initial year of U.S. commercialization. *Surg Obes Relat Dis.* 2019;15(3): 417–423. doi: 10.1016/j.soard.2018.12.007.
18. Machytka E, Klvana P, Kornbluth A, et al. Adjustable intragastric balloons: a 12-month pilot trial in endoscopic weight loss management. *Obes Surg.* 2011;21(10):1499–1507. doi: 10.1007/s11695-011-0424-z.
19. Vantanasiri K, Matar R, Beran A, et al. The efficacy and safety of a procedureless gastric balloon for weight loss: a systematic review and meta-analysis. *OBES Surg.* 2020;30(9):3341–3346. doi: 10.1007/s11695-020-04522-3.
20. Lecumberri E, Krekshi W, Matía P, et al. Effectiveness and safety of air-filled balloon Heliosphere BAG® in 82 consecutive obese patients. *Obes Surg.* 2011 Oct;21(10):1508–12. doi: 10.1007/s11695-010-0314-9.
21. Giardiello C, Borrelli A, Silvestri E, et al. Air-filled vs water-filled intragastric balloon: a prospective randomized study. *Obes Surg.* 2012;22(12):1916–1919. doi: 10.1007/s11695-012-0786-x.
22. *ORBERA™ Intra-gastric Balloon System* – P140008. <https://www.fda.gov/MedicalDevices/ProductsandMedicalProcedures/DeviceApprovalsandClearances/Recently-ApprovedDevices/ucm457416.htm>. Accessed 20 Dec 2024.
23. *Obalon Balloon System* – P160001. <https://www.fda.gov/MedicalDevices/Product sandMedicalProcedures/DeviceApprovalsandClearances/Recently-ApprovedDevices/ucm520741.htm>. Accessed 20 Dec 2024.
24. *The FDA alerts health care providers about potential risks with fluid-filled intragastric balloons.* <https://www.fda.gov/medicaldevices/resourcesforyou/healthcareproviders/ucm540655.htm>. Accessed 20 Dec 2024.
25. Usuy E, Brooks J. Response Rates with the Spatz3 Adjustable Balloon. *Obes Surg* 2018;28:1271–6. doi: 10.1007/s11695-017-2994-x.
26. Ramai D, Singh J, Mohan BP, et al. Influence of the Elipse Intra-gastric Balloon on Obesity and Metabolic Profile: A Systematic Review and Meta-Analysis. *J Clin Gastroenterol.* 2021 Nov-Dec 01;55(10):836–841. doi: 10.1097/MCG.0000000000001484.

27. Keren D, Rainis T. Intra-gastric Balloons for Overweight Populations-1 Year Post Removal. *Obes Surg* 2018;28:2368-73. doi: 10.1007/s11695-018-3167-2.
28. Zurawinski W, Sokolowski D, Krupa-Kotara K, et al. Evaluation of the results of treatment of morbid obesity by the endoscopic intra-gastric balloon implantation method. *Wideochir Inne Tech Maloinwazyjne* 2017;12:37-48. doi: 10.5114/wiitm.2017.66856.
29. Almeghaiseb ES, Ashraf MF, Alamro RA, et al. Efficacy of intra-gastric balloon on weight reduction: Saudi perspective. *World J Clin Cases*. 2017 Apr 16;5(4):140-147. doi: 10.12998/wjcc.v5.i4.140.
30. Gollisch KSC, Raddatz D. Endoscopic intra-gastric balloon: a gimmick or a viable option for obesity? *Ann Transl Med*. 2020 Mar;8(Suppl 1):S8. doi: 10.21037/atm.2019.09.67. doi: 10.21037/atm.2019.09.67.
31. Sallet JA, Marchesini JB, Paiva DS, et al. Brazilian multicenter study of the intra-gastric balloon. *Obes Surg*. 2004;14(7):991-998. doi: 10.1381/0960892041719671.
32. Matar ZS, Mohamed AA, Abukhater M, et al. Small bowel obstruction due to air-filled intra-gastric balloon. *Obes Surg*. 2009;19(12):1727-1730. doi: 10.1007/s11695-009-9880-0.
33. Vanden Eynden F, Urbain P. Small intestine gastric balloon impaction treated by laparoscopic surgery. *Obes Surg*. 2001;11(5):646-648. doi: 10.1381/09608920160556913.
34. Abu Dayyeh BK, Rajan E, Gostout CJ. Endoscopic sleeve gastropasty: a potential endoscopic alternative to surgical sleeve gastrectomy for treatment of obesity. *Gastrointest Endosc*. 2013 Sep;78(3):530-5. doi: 10.1016/j.gie.2013.04.197.
35. Devière J, Ojeda Valdes G, Cuevas Herrera L, et al. Safety, feasibility and weight loss after transoral gastroplasty: first human multicenter study. *Surg Endosc*. 2008;22(3):589-98. doi: 10.1007/s00464-007-9662-5.
36. Fogel R, De Fogel J, Bonilla Y, et al. Clinical experience of transoral suturing for an endoluminal vertical gastroplasty: 1-year follow-up in 64 patients. *Gastrointest Endosc*. 2008;68:51-58. doi: 10.1016/j.gie.2007.10.061.
37. Moreno C, Closset J, Dugardeyn S, Baréa M, et al. Transoral gastroplasty is safe, feasible, and induces significant weight loss in morbidly obese patients: results of the second human pilot study. *Endoscopy*. 2008;40(5):406-13. doi: 10.1055/s-2007-995748.
38. Familiari P, Costamagna G, Bléro D, et al. Transoral gastroplasty for morbid obesity: a multicenter trial with a 1-year outcome. *Gastrointest Endosc*. 2011;74(6):1248-58. doi: 10.1016/j.gie.2011.08.046.
39. Espinós JC, Turró R, Moragas G, Bronstone A, et al. Gastrointestinal physiological changes and their relationship to weight loss following the POSE procedure. *Obes Surg*. 2016;26(5):1081-9. doi: 10.1007/s11695-015-1863-8.
40. López-Nava G, Bautista-Castaño I, Jimenez A, et al. The primary obesity surgery Endoluminal (POSE) procedure: one-year patient weight loss and safety outcomes. *Surg Obes Relat Dis*. 2015;11(4):861-5. doi: 10.1016/j.soard.2014.09.026.
41. Miller K, Turró R, Greve JW, et al. MILEPOST multicenter randomized controlled trial: 12-month weight loss and satiety outcomes after pose (SM) vs. medical therapy. *Obes Surg*. 2017;27(2):310-22. doi: 10.1007/s11695-016-2295-9.
42. Sullivan S, Swain JM, Woodman G, Antonetti M, et al. Randomized sham-controlled trial evaluating efficacy and safety of endoscopic gastric plication for primary obesity: The ESSENTIAL trial. *Obesity (Silver Spring)*. 2017 Feb;25(2):294-301. doi: 10.1002/oby.21702.
43. Cummings DE, Schauer PR. Endoscopic gastric plication for obesity: Where might it fit in the scheme of things? *Obesity (Silver Spring)*. 2017 Feb;25(2):284-285. doi: 10.1002/oby.21766.
44. Ahishali E. Endoscopic sleeve gastroplasty for obesity: a multicenter study of 248 patients with 24 months follow-up. *Turk J Gastroenterol*. 2018;29(3):373-4. doi: 10.5152/tjg.2018.180501.
45. Lopez-Nava G, Sharaiha RZ, Vargas EJ, et al. Endoscopic sleeve gastroplasty for obesity: a multicenter study of 248 patients with 24 months follow-up. *Obes Surg*. 2017;27(10):2649-55. doi: 10.1007/s11695-017-2693-7.

46. Sharaiha RZ, Kumta NA, Saumoy M, et al. Endoscopic sleeve gastropasty significantly reduces body mass index and metabolic complications in obese patients. *Clin Gastroenterol Hepatol*. 2017;15(4):504–10. doi: 10.1016/j.cgh.2016.12.012.
47. Kumar N, Abu Dayyeh BK, Lopez-Nava Breviere G, et al. Endoscopic sutured gastropasty: procedure evolution from first-in-man cases through current technique. *Surg Endosc*. 2018;32(4):2159–64. doi: 10.1007/s00464-017-5869-2.
48. Graus Morales J, Crespo Pérez L, Marques A, et al. Modified endoscopic gastropasty for the treatment of obesity. *Surg Endosc*. 2018;32(9):3936–42. doi: 10.1007/s00464-018-6133-0.
49. Sartoretto A, Sui Z, Hill C, et al. Endoscopic sleeve gastropasty (ESG) is a reproducible and effective endoscopic bariatric therapy suitable for widespread clinical adoption: a large, international multicenter study. *Obes Surg*. 2018;28(7):1812–21. doi: 10.1007/s11695-018-3135-x.
50. Abu Dayyeh BK, Acosta A, Camilleri M, et al. Endoscopic sleeve gastropasty alters gastric physiology and induces loss of body weight in obese individuals. *Clin Gastroenterol Hepatol*. 2017;15(1):37–43.e1. doi: 10.1016/j.cgh.2015.12.030.
51. Kumar N, Sullivan S, Thompson CC. The role of endoscopic therapy in obesity management: intragastric balloons and aspiration therapy. *Diabetes Metab Syndr Obes*. 2017 Jul 6;10:311–316. doi: 10.2147/DMSO.S95118.
52. Sullivan S, Edmundowicz SA, Thompson CC. Endoscopic bariatric and metabolic therapies: new and emerging technologies. *Gastroenterology*. 2017;152(7):1791–801. doi: 10.1053/j.gastro.2017.01.044.
53. Lee HL. Role of Restrictive Endoscopic Procedures in Obesity Treatment. *Clin Endosc*. 2017 Jan;50(1):17–20. doi: 10.5946/ce.2017.022.
54. Choi YI, Kim KO. Experimental Gastric Non-Balloon Devices. *Clin Endosc*. 2018 Sep;51(5):420–424. doi: 10.5946/ce.2018.150.
55. Thompson CC, Abu Dayyeh BK, Kushner R, et al. Percutaneous Gastrotomy Device for the Treatment of Class II and Class III Obesity: Results of a Randomized Controlled Trial. *Am J Gastroenterol*. 2017 Mar;112(3):447–457. doi: 10.1038/ajg.2016.500.
56. Yen YA, Wang CC, Sung WW, et al. Intragastric injection of botulinum toxin A for weight loss: A systematic review and meta-analysis of randomized controlled trials. *J Gastroenterol Hepatol*. 2022 Jun;37(6):983–992. doi: 10.1111/jgh.15847.
57. Foschi D, Corsi F, Lazzaroni M, et al. Treatment of morbid obesity by intraparietogastric administration of botulinum toxin: a randomized, double-blind, controlled study. *Int J Obes*. 2007 Apr;31(4):707–12. doi: 10.1038/sj.ijo.0803451.
58. Sánchez Torralvo FJ, Vázquez Pedreño L, Gonzalo Marín M, Endoscopic intragastric injection of botulinum toxin A in obese patients on bariatric surgery waiting lists: A randomised double-blind study (IntraTox study). *Clin Nutr*. 2021 Apr;40(4):1834–1842. doi: 10.1016/j.clnu.2020.10.008.
59. Osio M, Mailland E, Muscia F, et al. Botulinum neurotoxin-A does not spread to distant muscles after intragastric injection: A double-blind single-fiber electromyography study. *Muscle Nerve*. 2010 Aug;42(2):165–9. doi: 10.1002/mus.21662.
60. Mittermair R, Keller C, Geibel J. Intragastric injection of botulinum toxin A for the treatment of obesity. *Obes Surg*. 2007 Jun;17(6):732–6. doi: 10.1007/s11695-007-9135-x.
61. de Moura EGH, Ribeiro IB, Frazão MSV, et al. EUS-Guided Intragastric Injection of Botulinum Toxin A in the Preoperative Treatment of Super-Obese Patients: A Randomized Clinical Trial. *Obes Surg*. 2019 Jan;29(1):32–39. doi: 10.1007/s11695-018-3470-y.
62. Chang PC, Jhou HJ, Chen PH, et al. Intragastric Botulinum Toxin A Injection Is an Effective Obesity Therapy for Patients with BMI > 40 kg/m²: a Systematic Review and Meta-analysis. *Obes Surg*. 2020 Oct;30(10):4081–4090. doi: 10.1007/s11695-020-04842-4.
63. Arabpour E, Golmoradi H, Tape PMK, et al. Intragastric botulinum toxin injection for weight loss: current trends, shortcomings and future perspective. *Clin Endosc*. 2025 Jan;58(1):10–24. doi: 10.5946/ce.2024.153.

General Surgery V

64. Hennen C, Demir S, Dafsari HS, et al. Botulism after intragastric botulinum toxin injections for weight reduction. *Eur J Neurol*. 2023 Dec;30(12):3979-3981. doi: 10.1111/ene.16040.
65. Sandler BJ, Rumbaut R, Swain CP, et al. Human experience with an endoluminal, endoscopic, gastrojejunal bypass sleeve. *Surg Endosc*. 2011;25(9):3028-33. doi: 10.1007/s00464-011-1665-6.
66. Sandler BJ, Rumbaut R, Swain CP, et al. One-year human experience with a novel endoluminal, endoscopic gastric bypass sleeve for morbid obesity. *Surg Endosc*. 2015 Nov;29(11):3298-303. doi: 10.1007/s00464-015-4081-5.
67. Na HK, De Moura DTH; Study Group for Endoscopic Bariatric and Metabolic Therapies of the Korean Society of Gastrointestinal Endoscopy. Various Novel and Emerging Technologies in Endoscopic Bariatric and Metabolic Treatments. *Clin Endosc*. 2021 Jan;54(1):25-31. doi: 10.5946/ce.2021.021.
68. Rubino F, Forgione A, Cummings DE, et al. The mechanism of diabetes control after gastrointestinal bypass surgery reveals a role of the proximal small intestine in the pathophysiology of type 2 diabetes. *Ann Surg*. 2006 Nov;244(5):741-9. doi: 10.1097/01.sla.0000224726.61448.1b.
69. Dixon JB, le Roux CW, Rubino F, et al. Bariatric surgery for type 2 diabetes. *Lancet*. 2012 Jun 16;379(9833):2300-11. doi: 10.1016/S0140-6736(12)60401-2.
70. Rodriguez-Grunert L, Galvao Neto MP, Alamo M, et al. First human experience with endoscopically delivered and retrieved duodenal-jejunal bypass sleeve. *Surg Obes Relat Dis*. 2008 Jan-Feb;4(1):55-9. doi: 10.1016/j.soard.2007.07.012.

Chapter 4

CHANGE OF PUNCTURE SITE IN RIGHT INTERNAL JUGULAR CENTRAL VENOUS CATHETERIZATION – TECHNICAL NOTE

Ahmet GÜLTEKİN¹

INTRODUCTION

The rationale for optimizing puncture site and catheter depth will be addressed in this technical note. Forssmann performed the first cardiac catheterization on himself; however, his findings were rejected and ridiculed by his colleagues, preventing him from continuing his research. His pioneering work was later carried forward in the United States by André Frédéric Cournand and Dickinson Woodruff Richards Jr. In 1956, the three physicians jointly received the Nobel Prize in Medicine for their studies on the vascular and cardiac systems. The ingenuity and perseverance of these clinicians paved the way for one of the most commonly performed hospital procedures today: the insertion of peripheral and central venous catheters (1). A central venous catheter (CVC) is typically defined as one whose tip is positioned in the proximal segment of the superior vena cava, within the right atrium, or in the inferior vena cava. These catheters may be inserted through peripheral or centrally located veins, with the internal jugular, subclavian, and femoral veins being the most common access sites (2).

BACKGROUND

Historically, central venous catheterization has been performed using landmark-based methods, which depend on anatomical familiarity and the palpation of nearby arterial structures. Yet, such approaches do not account for anatomical variability at the intended puncture site. Numerous studies have shown that considerable deviations from ‘typical anatomy’ may occur in many patients, particularly involving the internal jugular, subclavian, and femoral veins (3).

¹ Assoc. Prof., Tekirdağ Namık Kemal University Faculty of Medicine, Department of Anesthesiology and Reanimation, ahmetgultekin82@yahoo.com, ORCID iD: 0000-0003-4570-8339

puncture point at the right internal jugular vein to the right margin of the sternal angle, measured via the right sternoclavicular joint (19).

CONCLUSION

When directly observed optimal catheter depths were compared to the formula-based values, it was found that formulas incorporating the distance from the puncture site to the right sternal angle are preferred. This is because the distal movement of the ultrasound probe facilitates easier and more precise CVC placement, making height-based formulas less accurate in these circumstances.

REFERENCES

1. Sette P, Dorizzi RM, Azzini AM. Vascular access: an historical perspective from Sir William Harvey to the 1956 Nobel prize to Andre F. Courmand, Werner Forssmann, and Dickinson W. Richards. *J Vasc Access*. 2012;13(2):137-44.
2. Smith RN, Nolan JP. Central venous catheters. *BMJ*. 2013;347:f6570.
3. Saugel B, Scheeren TWL, Teboul JL. Ultrasound-guided central venous catheter placement: a structured review and recommendations for clinical practice. *Crit Care*. 2017;21(1):225.
4. Hilty WM, Hudson PA, Levitt MA, Hall JB. Real-time ultrasound-guided femoral vein catheterization during cardiopulmonary resuscitation. *Ann Emerg Med*. 1997;29(3):331-6; discussion 7.
5. Lamperti M, Bodenham AR, Pittiruti M, Blaivas M, Augoustides JG, Elbarbary M, et al. International evidence-based recommendations on ultrasound-guided vascular access. *Intensive Care Med*. 2012;38(7):1105-17.
6. Dietrich CF, Horn R, Morf S, Chiorean L, Dong Y, Cui XW, et al. Ultrasound-guided central vascular interventions, comments on the European Federation of Societies for Ultrasound in Medicine and Biology guidelines on interventional ultrasound. *J Thorac Dis*. 2016;8(9):E851-E68.
7. Troianos CA, Hartman GS, Glas KE, Skubas NJ, Eberhardt RT, Walker JD, et al. Special articles: guidelines for performing ultrasound guided vascular cannulation: recommendations of the American Society of Echocardiography and the Society Of Cardiovascular Anesthesiologists. *Anesth Analg*. 2012;114(1):46-72.
8. Blaivas M, Brannam L, Fernandez E. Short-axis versus long-axis approaches for teaching ultrasound-guided vascular access on a new inanimate model. *Acad Emerg Med*. 2003;10(12):1307-11.
9. Chittoodan S, Breen D, O'Donnell BD, Iohom G. Long versus short axis ultrasound guided approach for internal jugular vein cannulation: a prospective randomised controlled trial. *Med Ultrason*. 2011;13(1):21-5.
10. Brass P, Hellmich M, Kolodziej L, Schick G, Smith AF. Ultrasound guidance versus anatomical landmarks for internal jugular vein catheterization. *Cochrane Database Syst Rev*. 2015;1(1):CD006962.
11. Bodenham Chair A, Babu S, Bennett J, Binks R, Fee P, Fox B, et al. Association of Anaesthetists of Great Britain and Ireland: Safe vascular access 2016. *Anaesthesia*. 2016;71(5):573-85.
12. Peres PW. Positioning central venous catheters--a prospective survey. *Anaesth Intensive Care*. 1990;18(4):536-9.
13. Hodzic S, Golic D, Smajic J, Sijercic S, Umihanic S, Umihanic S. Complications Related to Insertion and Use of Central Venous Catheters (CVC). *Med Arch*. 2014;68(5):300-3.

General Surgery V

14. Bannon MP, Heller SF, Rivera M. Anatomic considerations for central venous cannulation. *Risk Manag Healthc Policy*. 2011;4:27-39.
15. Kim MC, Kim KS, Choi YK, Kim DS, Kwon MI, Sung JK, et al. An estimation of right- and left-sided central venous catheter insertion depth using measurement of surface landmarks along the course of central veins. *Anesth Analg*. 2011;112(6):1371-4.
16. Jayaraman J, Shah V. Bedside prediction of the central venous catheter insertion depth - Comparison of different techniques. *J Anaesthesiol Clin Pharmacol*. 2019;35(2):197-201.
17. Wirsing M, Schummer C, Neumann R, Steenbeck J, Schmidt P, Schummer W. Is traditional reading of the bedside chest radiograph appropriate to detect intraatrial central venous catheter position? *Chest*. 2008;134(3):527-33.
18. Barnwal NK, Dave ST, Dias R. A comparative study of two techniques (electrocardiogram- and landmark-guided) for correct depth of the central venous catheter placement in paediatric patients undergoing elective cardiovascular surgery. *Indian J Anaesth*. 2016;60(7):470-5.
19. Ju H, Sun X, Feng Y. Determination and prediction of the appropriate depth of right internal jugular vein catheterization via the middle approach in adults using transesophageal echocardiography. *Echocardiography*. 2019;36(8):1496-500.

Chapter 5

DECISION-MAKING PROCESSES IN GENERAL SURGERY FOR ELDERLY PATIENTS

Yunushan Furkan AYDOĞDU¹

INTRODUCTION

Increased life expectancy and advances in healthcare have led to older individuals encountering diseases requiring surgical treatment more frequently. Today, a significant proportion of patients evaluated in general surgery practice are aged 65 and over, and this proportion is increasing (1-6). Despite advances in Surgical techniques, anesthesia practices, and perioperative care, postoperative complication rates in elderly patients are still higher than in younger patients, making surgical decision-making processes significantly more complex (2,3,7-11).

One of the most important challenges in managing elderly surgical patients is the heterogeneity of this patient group. Two individuals of the same chronological age may have significant differences in functional capacity, burden of comorbidities, level of independence, and physiological reserve. Therefore, assessing surgical risk based solely on age is insufficient and can often be misleading (2,12-16). The increasingly accepted approach today is that the primary factor determining surgical outcomes is the patient's physiological reserve capacity rather than chronological age (1,13,12,17,18).

The decline in physiological reserve in the elderly is characterized by functional deterioration in multiple organ systems and reduced tolerance to stress factors. This situation causes elderly patients to become more vulnerable in the face of major physiological stress, such as surgical intervention. Reserve loss, particularly in the cardiovascular, pulmonary, and renal systems, is among the key factors that increase the risk of complications developing in the postoperative period (1,4,9,19,20). In this context, the concept of "physiological frailt" has come to the

¹ MD, General Surgeon, Department of General Surgery, Ankara Training and Research Hospital, yfaydogdu92@gmail.com, ORCID iD: 0000-0002-2418-2393

Chapter 6

SURGICAL TREATMENT OF PULMONARY ASPERGILLOMA

Salih Cüneyt AYDEMİR¹

Aspergillus is a fungal species widely distributed in nature. The most encountered pathogenic species in humans is *Aspergillus fumigatus*. *Aspergillus* spores acquired via inhalation may cause various diseases. The most frequently encountered of these is aspergilloma, which represents a saprophytic infection.

Aspergillomas are mass-forming lesions created by the settlement of heavily inhaled *Aspergillus* spores into pre-existing cavities within the host, where fungal hyphae combine with blood elements and necrotic tissue. Frequently observed in tuberculous cavities, aspergillomas may develop in any cystic or cavitory substrate, including bronchiectasis, bronchogenic cysts, or cavitating bronchogenic carcinoma.

The most common symptom in patients is hemoptysis. Minor hemoptysis episodes that resolve spontaneously may be observed, as well as life-threatening fatal hemorrhages. Cough, chest pain, dyspnea, and fever are other symptoms that may be seen in patients with aspergilloma.

On conventional chest radiographs and computed tomography scans, a mass formation demonstrating the air-crescent sign is characteristic of aspergilloma. It is generally seen in the upper lobes and is usually unilateral. The significance of isolating *Aspergillus* species in sputum cultures for diagnosis is uncertain. Detection of precipitating antibodies against *Aspergillus* in serum constitutes both a specific and sensitive finding. Definitive diagnosis is established by histological demonstration of *Aspergillus* in biopsy material and by culture of the organism.

Systemic or topical antifungals are ineffective in the treatment of aspergilloma except in limited cases. Bronchial artery embolization, applied to control hemoptysis episodes, is also generally unsuccessful due to the difficulty in identifying the bleeding vascular structures and the development of extensive

¹ Dr., Thoracic Surgeon, Beylikdüzü State Hospital, dr.caydemir@gmail.com,
ORCID iD: 0000-0002-9552-3255

REFERENCES

1. Marr K.A, Patterson T, Denning D. Aspergillosis; pathogenesis, clinical manifestations and therapy. *Infect Dis Clin North Am* 2002; 16: 875-94.
2. Rippon J.W. *Medical Mycology: The Pathogenic Fungi and the Pathogenic Actinomycetes*, 3th ed Philadelphia, WB Saunders, 1988: 582.
3. Bardana E.J. The clinical spectrum of aspergillosis-Part II: Clasification and description of saprophytic, allergic and invasive variants of human diseases. *Crit Rev Clin Lab Sci* 1981; 13: 85-159.
4. Pennington J.E. Aspergillus lung disease. *Med Clin North Am* 1980; 64: 475-90.
5. British Tuberculosis and Thoracic Association. Aspergilloma and residual tuberculosis cavities: the result of a resurvey. *Tubercle* 1970; 51: 227-45.
6. Glimp R.A, Bayer A.S. Pulmonary aspergilloma: diagnostic and therapeutic considerations. *Arch intern med* 1983; 143: 303-8.
7. Stevens D.A, Kan V.L, Judson M.A, Morrison V.A, Dummer S, Denning D.W, Bennet J.E, Walsh T.J, Patterson T.F, Pankey G.A. Practice guidelines for diseases caused by aspergillus. *Clin Infect Dis* 2000; 30: 696-709.
8. Editorial. Quandary about treatment of aspergillomas persists. *Lancet* 1996; 347: 1640.
9. Massard G, Roeslin N, Wihlm J.M, Dumont P, Witz J.P, Morand G. Pleuro-pulmonary aspergilloma: clinical spectrum and results of surgical treatment. *Ann Thorac Surg* 1992; 54: 1159-64.
10. Kaplan J, Johns C.J. Mycetomas in pulmonary sarcoidosis: nonsurgical management. *J Hopkins Med J* 1979; 145: 157-61.
11. Remy J, Arnaud A, Fardou H, Giraud R, Voisin C. Treatment of hemoptysis by embolization of bronchial arteries. *Radiology* 1977; 122: 33-7.
12. Jewkes J, Kay P.H, Paneth M, Citron K.M. Pulmonary aspergilloma: analysis of prognosis in relation to haemoptysis and survey of treatment. *Thorax* 1983; 38: 572-8.
13. Kaestel M, Meyer W, Mittelmeier H.O, Gebhardt C. Pulmonar aspergilloma: clinical findings and surgical treatment. *Thorac Cardiovasc Surg* 1999; 47: 340-5.
14. Conlan A.A, Abramor E, Moyes D.G. Pulmonary aspergilloma-indications for surgical intervention: an analysis of 22 cases. *S Afr Med J* 1987; 71: 285-8.
15. Oakley R.E, Petrou M, Goldstraw P. Indications and outcome of surgery for pulmonary aspergilloma. *Thorax* 1997; 52: 813-5.
16. Rinaldi M. Invasive aspergillosis. *Rev Infect Dis* 1983; 5:1061-77
17. Onul B. Yedinci Bölüm – Muhtelif Mantar Hastalıkları – Aspergillosis. *Tibbi Mikoloji. AÜ Tıp Fak Yayınları. Ankara* 1950; 17: 282-8.
18. Khurana S, Thaker K, Kane G.C. Pulmonary aspergillosis, part 1: allergic disease and mycetomas; recognizing the pattern of illness is key. *J Res Dis* 2002; 23(5): 300-7.
19. Garvey J, Crastnopol P, Weitsz D, Khan F. The surgical treatment of pulmonary aspergillomas. *Thorac Cardiovasc Surg* 1977; 74(4): 542-7.
20. Dar M.A, Ahmad M, Weinstein A.J, Mehta A.C, Golish J.A. Thoracic aspergillosis. *Cleve Clin Q* 1984; 51:615-30.
21. Unat E.K. Aspergilloz. *Tibbi Mikoloji Ders Kitabı 2.Baskı. İstanbul* 1962: 138-44.
22. Hinson K.F.W, Moon A.J, Plummer N.S: Bronchopulmonary aspergillosis. Review and report of eight cases. *Thorax* 1952; 7: 317-33.
23. Levinson W, Jawetz E. Fırsatçı Mikroplar – Aspergillus. *Tibbi Mikoloji ve İmmünoloji* 4. Baskı 1997; 50: 281-2.
24. Conant N.F, Smith D.T, Baker R.D, Callaway J.L, Martin D.S, Aspergillosis. *Manual of clinical mycology. Chap 10. WB Saunders Company. Philadelphia* 1969: 203-12.
25. Schaffner A. Macrophage-Aspergillus interactions. *Immunology* 1994; 60: 545-52.
26. Waldorf A.R, Levitz S.M, Diamond R.D. In vivo bronchoalveolar macrophage defense against *Rhizopus oryzae* and *Aspergillus fumigatus*. *J Infect Dis* 1984; 150: 752-60.

27. Schaffner A, Douglas H, Braude A. Selective protection against candida by mononuclear and against mycelia by polymorphonuclear phagocytes in resistance to *Aspergillus*. *J Clin Invest* 1982; 69: 617-31.
28. Cenci E, Mencacci A, Fe d'Ostiani C, Del Sero G, Mosci P, Montagnoli C, Bacci A, Romani L. Cytokine- and T helper-dependent lung mucosal immunity in mice with invasive pulmonary aspergillosis. *J Infect Dis* 1998;178:1750-60.
29. Albilda S.M, Talbot G.H. Pulmonary aspergillosis. In Fishman AP (ed): *Pulmonary diseases and disorders*, vol.2, 2nd ed. New York, McGraw-Hill, 1988.
30. Latge J.P. *Aspergillus fumigatus* and aspergillosis. *Clin Microbiol Rev* 1999; 12: 310-50.
31. Kumar R, Gaur S.N. Prevalence of allergic pulmonary aspergillosis in patients with bronchial asthma. *Asian Pac J Allergy Immunol* 2000; 18: 181-5.
32. Addis B. Pulmonary mycotic disease. *Spencer's Pathology of the Lung*. 5th Ed. Ch 8 Philip S. Hasleton 1996;257-304.
33. Patterson R, Greenberger P.A, Radin R.C, Robert M. Allergic bronchopulmonary aspergillosis: staging as an aid to management. *Ann Intern Med* 1982; 96: 286-91.
34. Patterson R, Greenberger P.A, Halwig J.M, Liotta J.L, Roberts M. Allergic bronchopulmonary aspergillosis: natural history and classification of early disease by serologic and roentgenographic studies. *Arch Intern Med* 1986; 146: 916-8.
35. Katzenstein A, Liebow A, Friedman P. Bronchocentric granulomatosis, mucoid impaction and hypersensitivity reaction to fungi. *Am Rev Respir Dis* 1975; 111: 497-508.
36. Rippon J.W. Mycetoma, in *Medical Mycology: The Pathogenic Fungi and the Pathogenic Actinomycetes*, ed 3. Philadelphia, WB Saunders, 1988: 80.
37. Irwin A. Radiology of the aspergilloma. *Clin Radiol* 1966; 18: 432-8.
38. Kathuria S.K, Rippon J. Non-aspergillus aspergilloma. *Am J Clin Path* 1982;78:870-3.
39. Nime F.A, Hutchins G.M. Oxalosis caused by aspergillus infection. *John Hopkins Med J* 1973; 133: 183-94.
40. Kaplan W. : Direct fluorescent antibody tests for the diagnosis of mycotic diseases. *Ann Clin Lab Sci* 1976;3:25-9.
41. Cook D.J, Achong M.R, King D.E. Disseminated aspergillosis in an apparently healthy patient. *Am J Med* 1990; 88: 74-6.
42. Denning D.W, Follansbee S.E, Scolaro M, Norris S, Edelstein H, Stevens D.A. Pulmonary aspergillosis in the acquired immunodeficiency syndrome. *N Eng J Med* 1991; 324: 654-62.
43. Bech A.O. Diffuse bronchopulmonary aspergillosis. *Thorax* 1961; 16:144-52.
44. Binder R, Faling J, Pugatch R, Mahasaen C, Snider G. Chronic necrotizing pulmonary aspergillosis: a discrete clinical entity. *Medicine* 1982; 61: 109-24.
45. Sharma O.P, Chwogule R. Mant faces of pulmonary aspergillosis. *Eur Resp J* 1998; 12: 705-15.
46. Soubani A.O, Chandrasekar P.H. The clinical spectrum of pulmonary aspergillosis. *Chest* 2002; 121: 1988-99.
47. Rafferty P, Biggs B, Crompton G.K, Grant I. What happens to patients with pulmonary aspergilloma? *Thorax* 1983; 38: 579-83.
48. Smith R.L, Morelli M.J, Aranda C.P. Pulmonary aspergilloma diagnosed by fiberoptic bronchoscopy. *Chest* 1987; 92: 948-9.
49. Cockrill B.A, Hales C.A. Allergic bronchopulmonary aspergillosis. *Annu Rev Med* 1999; 50: 303-16.
50. Stevens DA, Schwartz H.J, Lee J.H, Moskovitz B.L, Jerome D.C, Catanzaro A, Bamberger D.M, Weinmann A.J, Tuazon C.U, Judson M.A, Thomas A.E, Platts-Mills T.A, DeGraff A.C. A randomized trial of itraconazole in allergic bronchopulmonary aspergillosis. *N Engl J Med* 2000; 342: 756-62.
51. Jennings T, Hardin T. Treatment of aspergillosis with itraconazole. *Ann Pharmacother* 1993; 27: 1206-11.

52. Henderson A, Pearson J. Treatment of pulmonary aspergillosis with observation on the use of natamycin. *Torax* 1996; 23: 519-23.
53. Uflacker R, Kaemmerer A, Picon P.D, Rizzon C.F, Neves C.M, Oliveira E.S, Oliveira M.E, Azevedo S.N, Ossanai R. Bronchial artery embolization in the management of hemoptysis: technical aspects and long-term results. *Radiology* 1985; 157: 637-44.
54. Koşar A, Uzun C, Kırıl H, Ürek Ş, Örki A, Aydemir C, Dudu C.Ş, Arman B. Postpnömonektomik bronkoplevral fistüllerde cerrahi tedavi yaklaşımları. *Heybeliada Tıp Bülteni* 2001; 7(2): 18-23.
55. McPherson P. Pulmonary asoergilliosis in Argyll. *Br J Dis Chest* 1965;59:148.
56. The Research Committee of the British Tuberculosis Association. Aspergillus in persistent lung cavities after tuberculosis. *Tubercle* 1968;49:1-4.
57. Türker H, Karakurt Z, Akın H, Erdem E, Uysal E, Kapaklı N, Sulu E, Atasalihi A. Pulmonary aspergilloma in a Turkish hospital population. *Turkish Respiratory Journal* 2002; 3(1): 7-14.
58. Regnard J-F, Icard P, Nicolosi M, Spaggiari L, Magdeleinat P, Jauffret B, Levasseur P. Aspergilloma: a series of 89 surgical cases. *Ann Thorac Surg* 2000; 69: 898-903.
59. Thomas D.A, Gonzalez-Rothi R.J. Aspergilloma in an open chest cavity. *Chest* 1989; 95: 1156-8.
60. Israel H.L, Lenchner G.S, Atkinson G.W. Sarcoidosis and aspergilloma- the role of surgery. *Chest* 1982; 82: 430-2.
61. Tomlinson J.R, Sahn S.A. Aspergilloma in sarcoid and tuberculosis. *Chest* 1987; 92: 505-8.
62. Babatasi G, Massetti M, Chapelier A, Fadel E, Macchiarini P, Khayat A, Dartevelle P. Surgical treatment of pulmonary aspergilloma: current outcome. *J Thorac Cardiovasc Surg* 2000; 119: 906-12.
63. Daly R.C, Pairolero P.C, Piehler J.M, Trastek V.F, Payne W.S, Bernatz P.E. Pulmonary aspergilloma: results of surgical treatment. *J Thorac Cardiovasc Surg* 1986; 92: 981-8.
64. Park C.K, Jheon S. Results of surgical treatment for pulmonary aspergilloma. *European Journal of Cardio-thoracic Surgery* 2002; 21: 918-23.
65. Solit R.W, McKeown J.J, Smullens S, Fraimow W. The surgical implications of intracavitary mycetomas (fungus balls). *J Thorac Cardiovasc Surg* 1971; 62: 411-22.
66. Jewkes J, Kay P.H, Paneth M, Citron K.M. Pulmonary aspergilloma: analysis of prognosis in relation to haemoptysis and survey of treatment. *Thorax* 1983; 38: 572-8.
67. Massard G, Roeslin N, Wihlm J.M, Dumont P, Witz J.P, Morand G. Pleuropulmonary aspergilloma: clinical spectrum and results of surgical treatment. *Ann Thorac Surg* 1992; 54: 1159-64.
68. Uflacker R, Kaemmerer A, Neves C, Picon P.D. Management of massive hemoptysis by bronchial artery embolization. *Radiology* 1983; 146: 627-34.
69. Lucke J.C. Thoracic Mycotic and actinomycotic infections of the lung – aspergillosis. *General Thoracic Surgery* (5th ed). Shields T.W, LoCicero III J, Ponn R.B. Lippincott Williams & Wilkins. Philadelphia, PA 2000: 1090-5.
70. Chen J, Chang Y, Luh S, Surgical treatment for pulmonary aspergilloma: a 28 year experience. *Thorax* 1997; 52: 810-3.
71. Libshitz H.I, Atkinson G.W, Israel H.L. Pleural thickening as a manifestation of aspergillus superinfection. *Am J Radiol* 1974; 120: 883-6.
72. Battaglini J.W, Murray G.F, Keagy B.A, Starek P.J, Wilcox B.R. Surgical management of symptomatic pulmonary aspergilloma. *Ann Thorac Surg* 1985; 39: 512-6.
73. Paterson TF, Thompson GR, Denning DW, et al. Practice Guidelines for the Diagnosis and Management of Aspergillosis: 2016 Update by the Infectious Diseases Society of America. *Clinical Infectious Diseases*. 2016;63(4): e1–e60. doi:10.1093/cid/ciw326.
74. Denning DW, Cadranel J, Beigelman-Aubry C, et al. Chronic pulmonary aspergillosis: rationale and clinical guidelines for diagnosis and management. *European Respiratory Journal*. 2016;47(1): 45–68. doi:10.1183/13993003.00583-2015.
75. Ullmann AJ, Aguado JM, Arikan-Akdagli S, et al. Diagnosis and management of Aspergillus diseases: executive summary of the 2017 ESCMID-ECMM-ERS guideline. *Clinical Microbiology and Infection*. 2018;24(Suppl 1): e1–e38. doi:10.1016/j.cmi.2018.01.002.

Chapter 7

BILE DUCT INJURIES: PREVENTION, DIAGNOSIS AND CURRENT MANAGEMENT

Muhammet Fatih KEYIF¹

1) DEFINITION AND CLINICAL SIGNIFICANCE

Bile duct injury (BDI) refers to the broad clinical spectrum of iatrogenic damage associated with cholecystectomy, ranging from early bile leak, biloma, and sepsis to late ductal stricture, recurrent cholangitis and secondary liver injury. (1,2) Reducing this spectrum to a single complication heading is often insufficient; because under the same clinical title, there are conditions ranging from a minimal leak to complete transection and complex injuries close to the hilum, for which the treatment strategy changes completely. (1,3) In clinical practice, bile leak becomes more visible in the early period through bile drainage output, biloma, and signs of infection, whereas major ductal injuries, in the presence of delayed diagnosis or inadequate initial management, tend to follow a course with more severe morbidity, longer hospitalization and a higher need for repeat intervention. (4) This difference arises not only from the anatomical severity of the injury, but also from the capacity of the approach applied in the early period either to enlarge the problem or bring it under control. (3,5) Evaluating BDI only as a problem resolved in the early postoperative days may be misleading, because the main burden often accumulates through strictures that emerge during long-term follow-up, recurrent cholangitis attacks and the associated loss of quality of life. (2,6) From the perspective of long-term outcomes, even if postoperative recovery appears complete for the patient, the tendency of the biliary system to develop fibrosis and stenosis over time prolongs the clinical story. Indeed, the association of bile leak and ductal injury with adverse long-term outcomes in some series suggests that these complications are a rare but high-impact clinical problem. (2,4) At this point, the main goal is to establish the correct balance between the

¹ Assist. Prof., Department of General Surgery, Bolu Abant Izzet Baysal University Faculty of Medicine, drfatihkeyif16@gmail.com, ORCID iD: 0000-0001-7346-1041

REFERENCES

- Pesce A, Palmucci S, Greca GL, et al. Iatrogenic bile duct injury: impact and management challenges. *Clinical and Experimental Gastroenterology*. 2019;12:121-128. doi:10.2147/CEG.S169492
- Schreuder AM, Busch OR, Besselink MG, et al. Long-term impact of iatrogenic bile duct injury. *Digestive Surgery*. 2019;37(1):10-21. doi:10.1159/000496432
- de'Angelis N, Catena F, Memeo R, et al. 2020 WSES guidelines for the detection and management of bile duct injury during cholecystectomy. *World Journal of Emergency Surgery*. 2021;16(1):30. doi:10.1186/s13017-021-00369-w
- Fong ZV, Pitt HA, Strasberg SM, et al. Diminished survival in patients with bile leak and ductal injury: management strategy and outcomes. *Journal of the American College of Surgeons*. 2018;226(4):568. doi:10.1016/j.jamcollsurg.2017.12.023
- Wang X, Yu WL, Fu XH, et al. Early versus delayed surgical repair and referral for patients with bile duct injury: a systematic review and meta-analysis. *Annals of Surgery*. 2020;271(3):449. doi:10.1097/SLA.0000000000003448
- Booij KAC, Coelen RJ, de Reuver PR, et al. Long-term follow-up and risk factors for strictures after hepaticojejunostomy for bile duct injury: an analysis of surgical and percutaneous treatment in a tertiary center. *Surgery*. 2018;163(5):1121-1127. doi:10.1016/j.surg.2018.01.003
- Calabrese EC, Slater BJ, Awad Z, et al. SAGES-AHPBA 2025 guideline for the surgical management of bile duct injury following cholecystectomy. *Surgical Endoscopy*. 2026;40(1):1-17. doi:10.1007/s00464-025-12352-6
- Wakabayashi G, Iwashita Y, Hibi T, et al. Tokyo Guidelines 2018: surgical management of acute cholecystitis: safe steps in laparoscopic cholecystectomy for acute cholecystitis (with videos). *Journal of Hepato-Biliary-Pancreatic Sciences*. 2018;25(1):73-86. doi:10.1002/jhbp.517
- Brunt LM, Deziel DJ, Telem DA, et al. Safe cholecystectomy multi-society practice guideline and state-of-the-art consensus conference on prevention of bile duct injury during cholecystectomy. *Surgical Endoscopy*. 2020;34(7):2827-2855. doi:10.1007/s00464-020-07568-7
- Takahashi H, Raj R, Hughes A, et al. A systematic review of gallbladder anomalies. *Journal of the Society of Laparoendoscopic Surgeons*. 2026;30(1):e2025.00102. doi:10.4293/JLS.2025.00102. PubMed PMID: 41584374; PubMed Central PMCID: PMC12828709.
- Kazi IA, Siddiqui MA, Thimmappa ND, et al. Post-operative complications of cholecystectomy: what the radiologist needs to know. *Abdominal Radiology (New York)*. 2025;50(1):109-130. doi:10.1007/s00261-024-04387-5. PubMed PMID: 38940909; PubMed Central PMCID: PMC11711778.
- Tenorio-Flores E, Sanchez-Rodriguez IG, Garcia-Blanco MC, et al. Comprehensive imaging insights into post-cholecystectomy complications for enhanced clinical practice. *Clinical Medicine Research*. 2024;22(4):206-214. doi:10.3121/cmr.2025.1985. PubMed PMID: 39993831; PubMed Central PMCID: PMC11849967.
- Symeonidis D, Tepetes K, Tzovaras G, et al. BILE: a literature review based novel clinical classification and treatment algorithm of iatrogenic bile duct injuries. *Journal of Clinical Medicine*. 2023;12(11). doi:10.3390/jcm12113786
- Emara MH, Ahmed MH, Radwan MI, et al. Post-cholecystectomy iatrogenic bile duct injuries: emerging role for endoscopic management. *World Journal of Gastrointestinal Surgery*. 2023;15(12):2709-2718. doi:10.4240/wjgs.v15.i12.2709. PubMed PMID: 38222007; PubMed Central PMCID: PMC10784825.
- Tringali A, Costa D, Ramai D. Endoscopic management of biliary leaks: where are we now? *World Journal of Gastrointestinal Endoscopy*. 2025;17(7):107587. doi:10.4253/wjge.v17.i7.107587. PubMed PMID: 40677582; PubMed Central PMCID: PMC12264820.
- Esparham A, Calabrese EC, Ganescu O, et al. SAGES-AHPBA 2025: a systematic review and meta-analysis on the surgical management of bile duct injury following cholecystectomy. *Surgical Endoscopy*. 2025;39(12):7905-7925. doi:10.1007/s00464-025-12357-1

General Surgery V

- Mercado MA, Vilatoba M, Domínguez-Rosado I, et al. Evolution of the repair of bile duct injury in a high-volume center in Latin America. *Journal of Gastrointestinal Surgery*. 2025;29(12):102233. doi:10.1016/j.gassur.2025.102233
- Pirriianu C, Toma EA, Enciu O, et al. Management of iatrogenic bile-duct injury after cholecystectomy, 1995-2025: systematic review and meta-analysis. *Life*. 2025;15(12):1858. doi:10.3390/life15121858. PubMed PMID: 41465797; PubMed Central PMCID: PMC12734171.
- Vincenzi P, Mocchegiani F, Nicolini D, et al. Bile duct injuries after cholecystectomy: an individual patient data systematic review. *Journal of Clinical Medicine*. 2024;13(16):4837. doi:10.3390/jcm13164837. PubMed PMID: 39200979; PubMed Central PMCID: PMC11355347.

Bölüm 8

GRANULOMATOUS MASTITIS

Ferdi BOLAT¹

1. DEFINITION AND HISTORICAL PERSPECTIVE

Granulomatous mastitis (GM) is a heterogeneous group of diseases characterized by granulomatous inflammation of the mammary gland. Idiopathic granulomatous mastitis (IGM), the most common form of this group, is a diagnosis of exclusion made after all known etiological causes have been ruled out; systemic or infectious conditions such as sarcoidosis, tuberculosis, Wegener's granulomatosis, and foreign body reactions may also cause granulomatous breast inflammation (1,2). From a clinical standpoint, IGM is arguably one of the most deceptive benign breast conditions encountered in surgical practice, as its ability to closely mimic malignancy — both at the bedside and on imaging — frequently leads to unnecessary patient anxiety and diagnostic uncertainty.

The disease was first described by Kessler and Wolloch in 1972; although significant advances have been made over the past half century regarding its etiology, diagnosis, and treatment, clinical management remains controversial. High long-term recurrence rates, the absence of a single pathogenetic mechanism, and the scarcity of randomized controlled trials continue to make IGM an active area of research (2). Awareness of this entity is therefore as important as familiarity with its management, and clinicians who encounter inflammatory breast masses regularly should maintain a consistently high index of suspicion.

2. CLASSIFICATION

Granulomatous mastitis is classified into two main categories based on its etiological basis: infectious GM and non-infectious GM. This distinction is of critical importance to the clinician, as it directly determines the treatment approach. In practice, the classification of GM begins with a thorough clinical

¹ Assist. Prof. Department of General Surgery, Faculty of Medicine, Bolu Abant İzzet Baysal University, drferdibolat@gmail.com, ORCID iD: 0000-0002-3012-2362

in particular should be re-evaluated. Follow-up visits also serve a function beyond disease surveillance: they provide a structured opportunity to assess the psychological impact of a chronic, relapsing condition on a predominantly young female patient population, and clinicians should be attentive to signs of anxiety or depression during these encounters and refer appropriately when needed.

11. LIMITATIONS OF CURRENT EVIDENCE AND FUTURE DIRECTIONS

The idiopathic granulomatous mastitis literature consists largely of retrospective series, case reports, and methodologically heterogeneous systematic reviews. The absence of randomized controlled trials prevents answers to fundamental clinical questions such as the optimal corticosteroid dose and duration, the appropriate timing of immunomodulatory agents, and the precise definition of surgical indications.

In the coming period, research priorities should be shaped around the following themes: prospective microbiome studies aimed at clarifying the true pathogenetic role of *Corynebacterium*; randomized trials comparing corticosteroid dosing protocols; identification of biomarkers that can be used to predict recurrence; and determination of safe treatment options during pregnancy and lactation. It is also worth calling attention to the need for patient-reported outcome measures in future IGM research; most existing studies define success in terms of radiological resolution or recurrence rates, yet what matters most to patients — quality of life, cosmetic outcome, return to normal daily activities, and psychological wellbeing — is rarely systematically captured, and incorporating these endpoints into future trial designs would bring the evidence base closer to the lived reality of the disease.

The establishment of international databases and multicenter cohorts will be critical in overcoming the sample size limitations created by the rarity of the disease.

REFERENCES

- Barreto DS, Sedgwick EL, Nagi CS, et al. Granulomatous mastitis: Etiology, imaging, pathology, treatment, and clinical findings. *Breast Cancer Res Treat.* 2018;171(3):527–534. doi:10.1007/s10549-018-4870-3
- Benlghazi A, Messaoudi H, Belouad M, et al. Idiopathic granulomatous mastitis: A challenging case report and comprehensive review of the literature. *Int J Surg Case Rep.* 2024;118:109555. doi:10.1016/j.ijscr.2024.109555
- Jiao Y, Chang K, Jiang Y, et al. Identification of periductal mastitis and granulomatous lobular mastitis: A literature review. *Ann Transl Med.* 2023;11(3):158. doi:10.21037/atm-22-6473

- Wang X, He X, Liu J, et al. Immune pathogenesis of idiopathic granulomatous mastitis: From etiology toward therapeutic approaches. *Front Immunol.* 2024;15. doi:10.3389/fimmu.2024.1295759
- Co M, Cheng VCC, Wei J, et al. Idiopathic granulomatous mastitis: A 10-year study from a multi-centre clinical database. *Pathology.* 2018;50(7):742–747. doi:10.1016/j.pathol.2018.08.010
- Güven F. Non-infectious mastitis: Idiopathic granulomatous mastitis. *Turk J Radiol Semin.* 2023;11(3):213–228. doi:10.4274/trs.2023.2310111
- Chen W, Zhang D, Zeng Y, et al. Clinical characteristics and microbiota analysis of 44 patients with granulomatous mastitis. *Front Microbiol.* 2023;14. doi:10.3389/fmicb.2023.1175206
- Velidedeoglu M, Umman V, Kilic F, et al. Idiopathic granulomatous mastitis: Introducing a diagnostic algorithm based on 5 years of follow-up of 152 cases from Turkey and a review of the literature. *Surg Today.* 2022;52(4):668–680. doi:10.1007/s00595-021-02367-6
- Ong SS, Ho PJ, Liow JJK, et al. A meta-analysis of idiopathic granulomatous mastitis treatments for remission and recurrence prevention. *Front Med.* 2024;11. doi:10.3389/fmed.2024.1346790
- Shanbhag NM, Ameri MA, Shanbhag SN, et al. Diagnostic challenges and insights into granulomatous mastitis: A systematic review. *Cureus.* 2024;16. doi:10.7759/cureus.75733
- Granulomatous lobular mastitis: Imaging, diagnosis, and treatment. *AJR Am J Roentgenol.* 2019. doi:10.2214/AJR.08.1528
- Yuan QQ, Xiao SY, Farouk O, et al. Management of granulomatous lobular mastitis: An international multidisciplinary consensus (2021 edition). *Mil Med Res.* 2022;9(1):20. doi:10.1186/s40779-022-00380-5
- Sarmadian R, Safi F, Sarmadian H, et al. Treatment modalities for granulomatous mastitis, seeking the most appropriate treatment with the least recurrence rate: A systematic review and meta-analysis. *Eur J Med Res.* 2024;29(1):164. doi:10.1186/s40001-024-01761-3
- Toman D, Prokop J, Kubala O, et al. Granulomatous mastitis treatment options and our experience. *Rozhl Chir.* 2021;100(4):192–175. doi:10.33699/PIS.2021.100.4
- Kapoor NS, Blair SL. ASO author reflections: A symptom-based algorithm for management of granulomatous mastitis in the United States. *Ann Surg Oncol.* 2024;31(11):7405–7406. doi:10.1245/s10434-024-15807-7
- Zhou Y, Xu L. Clinical efficacy of different methods for treatment of granulomatous lobular mastitis: A systematic review and network meta-analysis. *PLoS One.* 2025;20(2):e0318236. doi:10.1371/journal.pone.0318236
- Xu G, Limaye S. Idiopathic granulomatous mastitis. *Ann Breast Surg.* 2023;7. doi:10.21037/abs-22-57

Chapter 9

DIFFERENCES BETWEEN RIGHT-SIDED AND LEFT-SIDED COLON CANCERS

Can AYDIN¹
Çağıl KARAEVLİ²
SAMİ AÇAR³

1- EMBRYOLOGICAL DEVELOPMENT AND ANATOMICAL DIFFERENCES

The colon develops from two parts of the gastrointestinal system: the midgut and the hindgut. The cecum, ascending colon, and the proximal two-thirds of the transverse colon are named as the right colon and originate from the midgut. The distal one-third of the transverse colon, descending colon, sigmoid colon, and proximal rectum are called the left colon and originate from the hindgut. These differences influence their blood flow, nerves, tissues, and function, essentially making the right and left colon behave differently.

Between weeks 6–10, the midgut temporarily herniates and rotates 270° counterclockwise. As a result of this rotation, the cecum moves from the upper quadrant to the right lower quadrant. The hindgut remains relatively fixed and joins the pelvic structures to form the distal rectum [1,2].

The splenic flexure is the embryological boundary. It is the point between the end of the transverse colon and the start of the descending colon. This area is a transition zone, influenced by both the midgut and hindgut. It is also known as a “watershed” area because it receives blood supply from both the superior mesenteric artery (SMA) and the inferior mesenteric artery (IMA). Perfusion in this region is relatively weaker; therefore, ischemic colitis is more likely to occur

¹ Res. Assist. Dr., Department of General Surgery, Faculty of Medicine, Tekirdağ Namık Kemal University, cnaydinn12@gmail.com, ORCID iD: 0000-0003-0712-2994

² Op. Dr., Department of General, Torbalı State Hospital, İzmir, Türkiye, ckrvl77@gmail.com, ORCID iD: 0000-0002-4280-7430

³ Assoc. Prof. Dr., Department of General Surgery, Faculty of Medicine, Tekirdağ Namık Kemal University, acarsami@gmail.com, ORCID iD: 0000-0003-4096-3963

Regarding treatment response, primary tumor localization is currently accepted as a predictive biomarker. While RAS/BRAF wild-type left colon tumors significantly benefit from anti-EGFR-based therapies, the efficacy of anti-EGFR treatments is limited in right colon tumors despite having the exact same biological profile [78,79]. For this reason, current ESMO and NCCN guidelines define tumor localization as a clear decision point in the selection of treatment for metastatic disease [52,60].

On the other hand, the introduction of immunotherapy into clinical practice has radically changed the treatment paradigm, especially for MSI-H/dMMR right colon cancers. In this patient group, which was previously considered to have a poor prognosis, long-lasting responses and a significant survival advantage have been achieved with PD-1 inhibitors [34,68]. Thus, the poor prognostic biology in right colon cancers has turned into a therapeutic advantage with proper patient selection.

Evaluated from a surgical standpoint, localization continues to impact the extent of resection, the scope of lymph node dissection, and perioperative risks [70]. However, modern oncological surgical principles (e.g., complete mesocolic excision) have the potential to improve survival independently of localization [53].

In conclusion, the question of “right or left?” in colon cancer management is no longer just an anatomical description; it is a fundamental clinical parameter that predicts the disease’s biology, treatment response, and long-term prognosis. The current approach necessitates a personalized, multidisciplinary, and guideline-based management strategy that considers tumor localization, molecular features, and patient factors collectively [52,60]. This perspective will pave the way for the development of more refined risk classifications and targeted treatment algorithms for colon cancer in the future.

REFERENCES

- Sadler TW. Langman’s Medical Embryology. 14th ed. Philadelphia: Wolters Kluwer; 2019.
- Standring S, editor. Gray’s Anatomy: The Anatomical Basis of Clinical Practice. 42nd ed. Elsevier; 2020.
- Chaouch MA, Abid N, Gabbouj S, et al. Comparative efficacy and long-term oncological safety of extended right hemicolectomy versus standard right hemicolectomy for splenic flexure cancers: a meta-analysis. *Front Oncol.* 2024; 13:1244693.
- Zenger S, Gürbüz B, et al. Differences Between Right and Left Colon Cancers in Terms of Clinicopathological Features. *Turk J Colorectal Dis.* 2020;30(4):253–60.
- Mik M, Berut M, Dziki L, et al. Right- and left-sided colon cancer – clinical and pathological differences of the disease entity in one organ. *Arch Med Sci.* 2017;13(1):157–62.

- Kalantzis I, Nonni A, et al. Clinicopathological differences and correlations between right and left colon cancer. *World J Clin Cases*. 2020;8(8):1424–36.
- Lee MS, Menter DG, Kopetz S. Right Versus Left Colon Cancer Biology: Integrating the Consensus Molecular Subtypes. *J Natl Compr Canc Netw*. 2017;15(3):411–9.
- Viana EF, et al. Anatomy and pathology of the mesocolon and implications for oncological resection. *Front Cell Infect Microbiol*. 2020; 10:498502.
- Hsu HC, Chen YC, et al. Clinicopathological and molecular differences in colorectal cancer according to location. *JGH Open*. 2019;3(6):242–250
- Japanese Society for Cancer of the Colon and Rectum (JSCCR) Guidelines 2022 for the treatment of colorectal cancer. *Int J Clin Oncol*. 2023;28(1):1–43.
- Benedix F, et al. Comparison of 17,641 patients with right- and left-sided colon cancer: differences in epidemiology, perioperative course, histology, and survival. *Dis Colon Rectum*. 2010;53(1):57–64.
- Lee GH, Malietzis G, Askari A, Bernardo D, Al-Hassi HO, Clark SK. Is right-sided colon cancer different to left-sided colorectal cancer? – a systematic review. *Eur J Surg Oncol*. 2015;41(3):300–8.
- Yahagi M, Okabayashi K, Hasegawa H, Tsuruta M, Kitagawa Y. The Worse Prognosis of Right-Sided Compared with Left-Sided Colon Cancers: A Systematic Review and Meta-analysis. *J Gastrointest Surg*. 2016;20(3):648–55.
- Petrelli F, Tomasello G, Borronovo K, et al. Prognostic Survival Associated With Left-Sided vs Right-Sided Colon Cancer. *JAMA Oncol*. 2017;3(2):211–9.
- Weiss JM, Pfau PR, O'Connor ES, et al. Mortality by stage for right- versus left-sided colon cancer: analysis of surveillance, epidemiology, and end results–Medicare data. *J Clin Oncol*. 2011;29(33):4401–9.
- Papagiorgis PC, Oikonomou A, Zampetoglou M, et al. Tumor location may affect colorectal cancer morphology and prognosis. *Tech Coloproctol*. 2010;14(1):51–4.
- Ziranu P, Pretta A, Pozzari M, et al. CDX2 expression correlates with clinical outcomes in MSI-H metastatic colorectal cancer patients receiving immune checkpoint inhibitors. *Sci Rep*. 2023;13:4397. doi:10.1038/s41598-023-31497-2.
- Osmond B, Zhang C, Dinh V, Boman BM, Facey COB. HOX Genes and Cancer Stem Cells: Update on HOX expression, DNA methylation, biomarker status, and genetic changes in colorectal cancer. *J Stem Cell Res Dev Ther*. 2020;6(4):045. doi:10.24966/SRDT-2060/100045.
- Lee MS, McGuffey EJ, Morris JS, et al. Association of CpG island methylator phenotype and EREG/AREG methylation and expression in colorectal cancer. *Br J Cancer*. 2016;114(12):1352–61.
- Baran B, Ozupek NM, Tetik NY, et al. Difference Between Left-Sided and Right-Sided Colorectal Cancer: A Focused Review of Literature. *Gastroenterol Res*. 2018;11(4):264–273.
- Toyota M, Ahuja N, Ohe-Toyota M, et al. CpG island methylator phenotype in colorectal cancer. *Proc Natl Acad Sci U S A*. 1999;96(15):8681–6.
- Kane MF, Loda M, Gaida GM, et al. Methylation of the hMLH1 promoter correlates with lack of expression of hMLH1 in sporadic colon tumors. *Cancer Res*. 1997;57(5):808–11.
- Juo YY, Johnston FM, Zhang DY, et al. Prognostic value of CpG island methylator phenotype among colorectal cancer patients: a systematic review and meta-analysis. *Ann Oncol*. 2014;25(12):2314–27.
- Kaz AM, Wong CJ, Dzieciatkowski S, et al. Patterns of DNA methylation in the normal colon vary by anatomical location, gender, and age. *Epigenetics*. 2014;9(4):492–502.
- Guinney J, Dienstmann R, Wang X, et al. The consensus molecular subtypes of colorectal cancer. *Nat Med*. 2015;21(11):1350–6.
- Yamauchi M, Morikawa T, Kuchiba A, et al. Assessment of colorectal cancer molecular features along bowel subsites challenges the conception of distinct dichotomy of proximal versus distal colorectum. *Gut*. 2012;61(6):847–54.
- Liu Y, Sethi NS, Hinoue T, et al. Comparative molecular analysis of gastrointestinal adenocarcinomas. *Cancer Cell*. 2018;33(4):721–735.e8.

- De Rook W, Claes B, Bernasconi D, et al. Effects of KRAS, BRAF, NRAS, and PIK3CA mutations on the efficacy of cetuximab plus chemotherapy in chemotherapy-refractory metastatic colorectal cancer: a retrospective consortium analysis. *Lancet Oncol.* 2010;11(8):753–62.
- Loree JM, Bailey AM, Johnson AM, et al. Molecular landscape of ERBB2/ERBB3-mutated metastatic colorectal cancer. *J Natl Cancer Inst.* 2018;110(12):1409–17.
- Missiaglia E, Jacobs B, D'Ario G, et al. Distal and proximal colon cancers differ in terms of molecular, pathological, and clinical features. *Ann Oncol.* 2014;25(10):1995–2001.
- Sveen A, Bruun J, Eide PW, et al. Colorectal cancer consensus molecular subtypes translated to preclinical models uncover potentially targetable cancer cell dependencies. *Clin Cancer Res.* 2018;24(4):794–806.
- Dienstmann R, Vermeulen L, Guinney J, et al. Consensus molecular subtypes and the evolution of precision medicine in colorectal cancer. *Nat Rev Cancer.* 2017;17(2):79–92.
- Benson AB, Venook AP, Al-Hawary MM, et al. Colon Cancer, Version 2.2021, NCCN Clinical Practice Guidelines in Oncology. *J Natl Compr Canc Netw.* 2021;19(3):329–59.
- Overman MJ, McDermott R, Leach JL, et al. Nivolumab in patients with metastatic DNA mismatch repair-deficient or microsatellite instability-high colorectal cancer (CheckMate 142): an open-label, multicentre, phase 2 study. *Lancet Oncol.* 2017;18(9):1182–91.
- National Comprehensive Cancer Network (NCCN). Clinical Practice Guidelines in Oncology: Colon Cancer. Version 2.2023.
- Yoshino T, Arnold D, Taniguchi H, et al. Panitumumab versus Bevacizumab in RAS Wild-Type Metastatic Colorectal Cancer in the PARADIGM Trial: An Open-Label, Multicenter, Phase III Randomized Controlled Trial. *Lancet.* 2023;402(10396):1115–29.
- Heinemann V, von Weikersthal LF, Decker T, et al. FOLFIRI plus cetuximab versus FOLFIRI plus bevacizumab as first-line treatment for patients with metastatic colorectal cancer (FIRE-3): a randomised, open-label, phase 3 trial. *Lancet Oncol.* 2014;15(10):1065–75.
- Van Cutsem E, Köhne CH, Láng I, et al. Cetuximab plus irinotecan, fluorouracil, and leucovorin as first-line treatment for metastatic colorectal cancer: updated analysis of overall survival according to tumor KRAS and BRAF mutation status. *J Clin Oncol.* 2011;29(15):2011–9.
- Kopetz S, Grothey A, Yaeger R, et al. Encorafenib, Binimetinib, and Cetuximab in BRAF V600E-Mutated Colorectal Cancer. *N Engl J Med.* 2019;381(17):1632–43.
- Gopalakrishnan V, Helmink BA, Spencer CN, Reuben A, Wargo JA. The influence of the gut microbiome on cancer, immunity, and cancer immunotherapy. *Cancer Cell.* 2018;33(4):570–80.
- Purcell RV, Visnovska M, Biggs PJ, Schmeier S, Frizelle FA. Distinct gut microbiome patterns associate with consensus molecular subtypes of colorectal cancer. *Sci Rep.* 2017;7(1):11590.
- Mima K, Nishihara R, Qian ZR, Cao Y, Sukawa Y, Nowak JA, et al. *Fusobacterium nucleatum* in colorectal carcinoma tissue and patient prognosis. *Gut.* 2016;65(12):1973–80.
- Wu S, Rhee KJ, Albesiano E, Rabizadeh S, Wu X, Yen HR, et al. A human colonic commensal promotes colon tumorigenesis via activation of T helper type 17 T cell responses. *Nat Med.* 2009;15(9):1016–22.
- Becht E, de Reyniès A, Giraldo NA, Pilati C, Buttard B, Lacroix L, et al. Immune and stromal classification of colorectal cancer is associated with molecular subtypes and relevant for precision immunotherapy. *Clin Cancer Res.* 2016;22(16):4057–66.
- Yanova M, Maltseva D, Tonevitsky A. Sidedness matters: single-cell perspectives on left- and right-sided colorectal cancer. *Front Cell Dev Biol.* 2025; 13:1720996.
- Zhang L, Zhao Y, Dai Y, et al. Immune landscape of colorectal cancer tumor microenvironment from different primary tumor location. *Front Immunol.* 2018;9:1578
- Zhong M, Xiong Y, Ye Z, et al. Microbial community profiling distinguishes left-sided and right-sided colon cancer. *Front Cell Infect Microbiol.* 2020;10:498502
- Lakemeyer L, Wittau M, Henne-Bruns D, Kornmann M, Lemke J. Diagnostic and Prognostic Value of CEA and CA19-9 in Colorectal Cancer. *Diseases.* 2021;9(1):21.

- Lo C-M, Jiang J-K, Lin C-C, et al. Detecting microsatellite instability in colorectal cancer using Transformer-based colonoscopy image classification and retrieval. *Diagnostics*. 2024;14(5):976.
- Hamid MA, Pammer LM, Oberparleiter S, Günther M, Amann A, Gruber RA, et al. Multidimensional differences of right- and left-sided colorectal cancer and their impact on targeted therapies. *NPJ Precision Oncology*. 2025 Apr 22;9(1):116. doi:10.1038/s41698-025-00892-y.
- Vogel JD, Felder SI, Bhama AR, Hawkins AT, Langenfeld SJ, Shaffer VO, et al. The American Society of Colon and Rectal Surgeons clinical practice guidelines for the management of colon cancer. *Dis Colon Rectum*. 2022;65(2):148-177. doi:10.1097/DCR.0000000000002323.
- Argilés G, Tabernero J, Labianca R, Hochhauser D, Salazar R, Iveson T, et al.; ESMO Guidelines Committee. Localised colon cancer: ESMO Clinical Practice Guidelines for diagnosis, treatment and follow-up. *Ann Oncol*. 2020;31(10):1291–1305. doi:10.1016/j.annonc.2020.06.022.
- Hohenberger W, Weber K, Matzel K, Papadopoulos T, Merkel S. Standardized surgery for colonic cancer: complete mesocolic excision and central ligation—technical notes and outcome. *Colorectal Dis*. 2009 May;11(4):354-364; discussion 364-365. doi:10.1111/j.1463-1318.2008.01735.x.
- Kim MK, Lee IK, Kye B-H, Kim J-G. Procedural difficulty differences according to tumor location do not compromise the clinical outcome of laparoscopic complete mesocolic excision for colon cancer: a retrospective analysis. *Oncotarget*. 2017;8:64509–64519. doi:10.18632/oncotarget.19780.
- Stamos MJ, Brady MT. Anastomotic leak: are we closer to eliminating its occurrence? *Ann Laparosc Endosc Surg*. 2018;3:66. doi:10.21037/ales.2018.07.07
- Green BL, Marshall HC, Collinson F, Quirke P, Guillou PJ, Jayne DG, et al. Long-term follow-up of the Medical Research Council CLASICC trial of conventional versus laparoscopically assisted resection in colorectal cancer. *Br J Surg*. 2013 Jan;100(1):75-82. doi:10.1002/bjs.8945.
- Moghadamyeghaneh Z, Masoomi H, Mills SD, Carmichael JC, Pigazzi A, Nguyen NT, et al. Outcomes of conversion of laparoscopic colorectal surgery to open surgery: analysis of the National Inpatient Sample (2009–2012). *JSLs*. 2014;18(4):e2014.00230. doi:10.4293/JSLs.2014.00230.
- Nfonsam V, Aziz H, Pandit V, Khalil M, Jandova J, Joseph B. Analyzing clinical outcomes in laparoscopic right vs. left colectomy in colon cancer patients using the NSQIP database. *Cancer Treat Commun*. 2016;8:1–4. doi:10.1016/j.ctrc.2016.03.006.
- Torun M, Uzun O, Duman M, Polat E, Senger AS, Dinçer M, et al. Comparison of oncological outcomes after curative resection for right-side colon cancer and left-side colon cancer: a retrospective observational study. *Turk J Colorectal Dis*. 2024;34(4):123-129. doi:10.4274/tjcd.galenos.2024.2024-7-3
- National Comprehensive Cancer Network (NCCN). NCCN Clinical Practice Guidelines in Oncology: Colon Cancer. Version 5.2025. NCCN; 2025.
- Grothey A, Sobrero AF, Shields AF, et al. Duration of adjuvant chemotherapy for stage III colon cancer. *N Engl J Med*. 2018;378(13):1177-1188.
- Baxter NN, Morris AM. Adjuvant Therapy for Stage II Colon Cancer. *J Clin Oncol*. 2022;40(16):1785-1794.
- Platt JR, Brown G, Seligmann JF, et al. FOXTROT2: innovative trial design to evaluate the role of neoadjuvant chemotherapy in colon cancer. *ESMO Open*. 2023;8(1):100276.
- Taieb J. FOXTROT: Are We Ready to Dance? *J Clin Oncol*. 2023;41(3):469-472.
- Chalabi M, Fanchi LF, Dijkstra KK, et al. Neoadjuvant immunotherapy leads to pathological responses in mismatch repair-deficient or microsatellite instability-high early-stage colon cancers (NICHE). *Nat Med*. 2020;26(4):566-576.
- ESMO. Metastatic colorectal cancer: ESMO Clinical Practice Guideline.
- Le DT, Durham JN, Smith KN, et al. Mismatch repair deficiency predicts response of solid tumors to PD-1 blockade. *Science*. 2017;357(6349):409-413.
- André T, Shiu KK, Kim TW, et al. Pembrolizumab in Microsatellite-Instability-High Advanced Colorectal Cancer. *N Engl J Med*. 2020;383(23):2207-2218.

General Surgery V

- Chalabi M, et al. Neoadjuvant Immunotherapy in Locally Advanced dMMR Colon Cancer (NICHE-2). *N Engl J Med*. 2024.
- Vogel JD, Eskicioglu C, Weiser MR, et al. The American Society of Colon and Rectal Surgeons Clinical Practice Guidelines for the Treatment of Colon Cancer. *Dis Colon Rectum*. 2017;60(10):999-1017.
- Lim DR, Kuk JK, Kim T, et al. Oncologic outcomes of right- vs left-sided colon cancer. *Medicine (Baltimore)*. 2017;96(42):e8241.
- Jaruhathai S, Santidamrongkul A, Wiwitkeyoonwong J, Duangnapa B. The impact of colon cancer sites on survival rates after curative resection and adjuvant chemotherapy: a single-center study. *Asian Pac J Cancer Care*. 2022;7(4):615-620. doi:10.31557/apjcc.2022.7.4.615-620.
- Asghari-Jafarabadi M, Wilkins S, Plazzer JP, Yap R, McMurrick PJ. Prognostic factors and survival disparities in right-sided versus left-sided colon cancer. *Sci Rep*. 2024;14:12306. doi:10.1038/s41598-024-63143-3. PMID:38811769.
- Sawayama H, Miyamoto Y, Hiyoshi Y, Ogawa K, Kawai K, Baba H, et al. Overall survival after recurrence in stage I–III colorectal cancer patients in accordance with the recurrence organ site and pattern. *Ann Gastroenterol Surg*. 2021 Jul;5(6):813–822. doi:10.1002/ags3.12483. PMCID: PMC8560596.
- Ribic CM, Sargent DJ, Moore MJ, et al. Tumor microsatellite instability status and benefit from fluorouracil-based adjuvant chemotherapy. *N Engl J Med*. 2003;349(3):247-257.
- Masoomi H, Buchberg B, Dang P, Carmichael JC, Mills S, Stamos MJ. Outcomes of right vs. left colectomy for colon cancer. *J Gastrointest Surg*. 2011 Nov;15(11):2023–2028. doi:10.1007/s11605-011-1655-y. PMID:21845511.
- Sargent DJ, Goldberg RM, Jacobson SD, et al. A pooled analysis of adjuvant chemotherapy for colon cancer. *J Clin Oncol*. 2009;27(12):1948-1956.
- Tejpar S, Stintzing S, Ciardiello F, et al. Prognostic and predictive relevance of primary tumor location. *JAMA Oncol*. 2017;3(2):194-201.
- Venook AP, Niedzwiecki D, Innocenti F, et al. Impact of primary tumor location. *J Clin Oncol*. 2017;35(30):3503-3511.
- Overman MJ, Lonardi S, Wong KYM, et al. Nivolumab plus ipilimumab in dMMR/MSI-H CRC. *J Clin Oncol*. 2018;36(8):773–779.
- Bufill JA. Colorectal cancer: evidence for distinct genetic categories based on proximal or distal tumor location. *Ann Intern Med*. 1990;113(10):779-788.
- Wirbel J, Pyl PT, Kartal E, et al. Meta-analysis of fecal metagenomes reveals global microbial signatures of colorectal cancer. *Nat Med*. 2019;25(4):679-689.
- Chen ZK, et al. Gut microbiota and the colorectal cancer tumor microenvironment: recent advances and perspectives. *World J Gastroenterol*. 2026;32(4):XXX-XXX. doi:10.3748/wjg.v32.i4.XXX. PMCID: PMC12836184.

Chapter 10

CHYLOTHORAX

Mehmet AĞAR¹

INTRODUCTION

Chylothorax is the pathological accumulation of a fluid called “chyle” (lymphatic fluid) in the pleural cavity as a result of obstruction, injury of the thoracic duct, or decreased lymphatic drainage. Chyle is a milky-appearing body fluid formed in the intestinal lacteal system and contains high amounts of triglycerides, chylomicrons, T-lymphocytes, immunoglobulins, enzymes, and fat-soluble vitamins (A, D, E, K).

Chylothorax was first described by Bartloet in 1633 (1). In the following years, Pecquet described the “cisterna chyli” (Pecquet’s cistern) in 1651 (2). In the field of surgical management, Lampson demonstrated in 1948 that chylothorax can be successfully treated by direct ligation of the thoracic duct (3).

Although chylothorax is rare, it is a potentially life-threatening condition when not managed in a timely and effective manner. It may lead to serious clinical outcomes such as nutritional and metabolic disorders, immunological deficiency, and respiratory failure. In untreated cases, the mortality rate can reach up to 50%; however, with modern and aggressive treatment approaches, this rate has decreased to 10-16% in postoperative cases (4).

The management of chylothorax should be carried out by a multidisciplinary team depending on the etiology and the amount of drainage. Pleural drainage, dietary modification, and pharmacological treatment usually constitute the first-line therapy, while interventional radiological methods and surgical treatment are considered advanced treatment options.

¹ Assist. Prof., Firat University, Faculty of Medicine, Thoracic Surgery Clinic, md.mehmetagar@gmail.com, ORCID iD: 0000-0002-4129-766X

For diagnosis, a pleural fluid triglyceride level above 110 mg/dL or the demonstration of chylomicrons by lipoprotein electrophoresis is considered the gold standard. Today, the most common cause of chylothorax is iatrogenic injury related to thoracic surgery, while malignancies, especially lymphoma, are the leading cause in non-traumatic cases. The most important factor guiding treatment is the drainage volume, and conservative management has a lower success rate in high-output leaks exceeding 1000 mL per day.

Loss of chyle may result in immunosuppression due to lymphocyte and immunoglobulin depletion, malnutrition due to protein and vitamin loss, and electrolyte imbalances. Therefore, the management of chylothorax requires a multidisciplinary approach. Advances in imaging, minimally invasive interventions, and surgical techniques have improved treatment outcomes.

In conclusion, chylothorax is a complex clinical condition that requires early diagnosis and appropriate management, as delayed treatment is associated with high mortality.

REFERENCES

- Peralta R, Ramzee AF, Bakhsh ZK, et al. Utility of an alpha-1 adrenergic agonist in the management of chylothorax: a case series and management algorithm. *European Journal of Case Reports in Internal Medicine*. 2024;11(8): 004705. doi:10.12890/2024_004705
- Santos LLD, Santos CLD, Hu NKT, et al. Outcomes of chylothorax nonoperative management after cardiothoracic surgery: a systematic review and meta-analysis. *Brazilian Journal of Cardiovascular Surgery*. 2023;38(6): e20220326. doi:10.21470/1678-9741-2022-0326
- V D Jr, Arshad AM, Ayub II, et al. A case of chylothorax in non-Hodgkin lymphoma. *Cureus*. 2024;16(10): e71957. doi:10.7759/cureus.71957
- Ahn HY, I H. Non-conservative management of chylothorax. *Journal of Chest Surgery*. 2021;54(4): 325–329. doi:10.5090/jcs.21.056
- Son J, Kim S, Son BS, et al. Outcomes of conservative management for chylothorax following minimally invasive lung cancer surgery. *Journal of Cardiothoracic Surgery*. 2025;20(1): 416. doi:10.1186/s13019-025-03618-0
- Nair SK, Petko M, Hayward MP. Aetiology and management of chylothorax in adults. *European Journal of Cardio-Thoracic Surgery*. 2007;32(2): 362–369. doi:10.1016/j.ejcts.2007.04.024
- Agrawal V, Doelken P, Sahn SA. Pleural fluid analysis in chylous pleural effusion. *Chest*. 2008;133(6): 1436–1441. doi:10.1378/chest.07-2232
- Rahman NM, Chapman SJ, Davies RJ. Pleural effusion: a structured approach to care. *British Medical Bulletin*. 2005;72: 31–47. doi:10.1093/bmb/ldh040
- Boffa DJ, Sands MJ, Rice TW, et al. A critical evaluation of a percutaneous diagnostic and treatment strategy for chylothorax after thoracic surgery. *European Journal of Cardio-Thoracic Surgery*. 2008;33(3): 435–439. doi:10.1016/j.ejcts.2007.11.028
- Lee EW, Shin JH, Ko HK, et al. Lymphangiography to treat postoperative lymphatic leakage: a technical review. *Korean Journal of Radiology*. 2014;15(6): 724–732. doi:10.3348/kjr.2014.15.6.724
- Papoulidis P, Vidanapathirana P, Dunning J. Chylothorax: new insights in treatment. *Journal of Thoracic Disease*. 2018;10(Suppl 33): S3976–S3977. doi:10.21037/jtd.2018.09.94
- Rudrappa M, Paul M. Chylothorax. In: *StatPearls*. Treasure Island (FL): StatPearls Publishing; 2026.

General Surgery V

- Ur Rehman K, Sivakumar P. Non-traumatic chylothorax: diagnostic and therapeutic strategies. *Breath*. 2022;18(2): 210163. doi:10.1183/20734735.0163-2021
- Thamkittikun C, Tovichien P. Clinical approach for pulmonary lymphatic disorders. *World Journal of Clinical Cases*. 2024;12(27): 6020–6026. doi:10.12998/wjcc.v12.i27.6020
- Chalret du Rieu M, Baulieux J, Rode A, et al. Management of postoperative chylothorax. *Journal of Visceral Surgery*. 2011;148(5): e346–e352. doi:10.1016/j.jvisc Surg.2011.09.006
- Wasmuth-Pietzuch A, Hansmann M, Bartmann P, et al. Congenital chylothorax: lymphopenia and high risk of neonatal infections. *Acta Paediatrica*. 2004;93(2): 220–224. doi:10.1080/08035250310007312