

World Journal of *Gastrointestinal Endoscopy*

World J Gastrointest Endosc 2011 January 16; 3(1): 1-22





Editorial Board

2009-2013

The *World Journal of Gastrointestinal Endoscopy* Editorial Board consists of 400 members, representing a team of worldwide experts in gastrointestinal endoscopy. They are from 45 countries, including Australia (7), Austria (1), Belgium (6), Brazil (7), Canada (5), Chile (2), China (26), Croatia (2), Cuba (1), Czech Republic (3), Denmark (1), Ecuador (1), Egypt (1), Finland (2), France (10), Germany (27), Greece (11), Hungary (4), India (15), Iran (2), Ireland (2), Israel (6), Italy (37), Japan (62), Lebanon (1), Lithuania (1), Malaysia (2), Mexico (1), Netherlands (6), New Zealand (1), Norway (2), Pakistan (2), Poland (2), Portugal (5), Romania (2), Singapore (2), South Africa (1), South Korea (13), Spain (17), Sweden (3), Thailand (5), Turkey (8), United Arab Emirates (1), United Kingdom (15), and United States (69).

PRESIDENT AND EDITOR-IN-CHIEF

Lian-Sheng Ma, *Beijing*

STRATEGY ASSOCIATE EDITORS-IN-CHIEF

Kazuya Akahoshi, *Iizuka*
William Robert Brugge, *Massachusetts*
Qiang Cai, *Georgia*
Juan J Vila Costas, *Pamplona*
Atsushi Irisawa, *Fukushima*
Andreas Sieg, *Heidelberg*
Gaetana Ilaria Tarantino, *Palermo*
Tony CK Tham, *Northern Ireland*
Konstantinos Triantafyllou, *Haidari*

GUEST EDITORIAL BOARD MEMBERS

Zhong-Ming Bai, *Taipei*
Wai-Keung Chow, *Taichung*
Wei-Hung Chan, *Taipei*
Yang-Yuan Chen, *Changhua*
Yen-Chang Chu, *Taichung*
Hwai-Jeng Lin, *Changhua*
Mei-Yung Tsou, *Taipei*
Bor-Shyang Sheu, *Tainan*
Ming-Yao Su, *Taoyuan*
Deng-Chyang Wu, *Kaohsiung*
Hsiu-Po Wang, *Taipei*
Ming-Shiang Wu, *Taipei*
Sheng-Lei Yan, *Tainan*

MEMBERS OF THE EDITORIAL BOARD



Australia

Hong-Chun Bao, *Victoria*

Michael J Bourke, *Sydney*
Ian C Lawrance, *Western Australia*
Rupert W Leong, *Concord*
Liang Qiao, *Westmead*
Michael Swan, *Victoria*
Rajvinder Singh, *South Australia*



Austria

Christine Kapral, *Linz*



Belgium

Giovanni Dapri, *Brussels*
Pierre Henri Deprez, *Brussels*
Christophe Moreno, *Brussel*
Tom G Moreels, *Antwerp*
Werner Van Steenberghe, *Leuven*
Daniel Urbain, *Brussels*



Brazil

Everson LA Artifon, *São Paulo*
Fátima Figueiredo, *Rio de Janeiro*
Fauze Maluf-Filho, *São Paulo*
Fernando Fornari, *Passo Fundo*
Joaquim PPM Filho, *São Paulo*
José Luiz Sebbas Souza, *São Paulo*
Claudio R Teixeira, *Porto Alegre*



Canada

Majid A Al Madi, *Montreal*

F Douglas Bair, *Ontario*
André Roy, *Québec*
Alan A Weiss, *Vancouver*
Brian Michael Yan, *Alberta*



Chile

Paul Richard Harris, *Marcoleta*
Italo FB Miranda, *Santiago*



China

Annie On On Chan, *Hong Kong*
Philip WY Chiu, *Hong Kong*
Jin Gu, *Beijing*
Simon Law, *Hong Kong*
Fu-Yu Li, *Chengdu*
Ka Ho Lok, *Hong Kong*
Tian-Le Ma, *Shanghai*
Si-Yu Sun, *Shenyang*
Anthony YB Teoh, *Shatin*
Kenneth KY Wong, *Hong Kong*
Jia-Ju Zheng, *Suzhou*
Jiang-Fan Zhu, *Shanghai*



Croatia

Josip Bago, *Zagreb*
Nadan Rustemović, *Zagreb*



Cuba

Damian C Rodriguez, *Havana*



Czech Republic

Marcela Kopacova, *Hradec Kralove*
 Michal Procke, *Prague*
 Miroslav Zavoral, *Prague*



Denmark

Peter Bytzer, *Koege*



Ecuador

Carlos Robles-Medranda, *Portoviejo*



Egypt

Nabil Ali Gad El-Hak, *Mansoura*



Finland

Paulina Salminen, *Turku*
 Lars Mikael Victorzon, *Vaasa*



France

Romain Coriat, *Paris*
 Bernard G Dallemagne, *Strasbourg*
 Gerard Jean Gay, *Vandoeuvre les Nancy*
 Lesur Gilles, *Boulogne*
 René Lambert, *Lyon*
 Sylvain Manfredi, *Rennes*
 Barthet Marc, *Marseille Cedex*
 JF Rey, *Saint Laurent Du Var Cedex*
 José Sahel, *Marseille*
 Nathalie Salles, *Pessac*



Germany

Marcel Binnebösel, *Aachen*
 P Born, *Munich*
 Stefan von Delius, *München*
 Dirk Domagk, *Muenster*
 Christoph Eisenbach, *Heidelberg*
 Ines Gockel, *Mainz*
 Arthur Hoffman, *Mainz*
 Georg FBA Kähler, *Mannheim*
 Günter Kampf, *Hamburg*
 Ralf Kiesslich, *Mainz*
 Andreas Kirschniak, *Tübingen*
 Oliver Pech, *Wiesbaden*
 Michael Pietsch, *Mainz*
 Andreas Probst, *Augsburg*
 Andrea Riphaut, *Bochum*
 Raphael Rosch, *Aachen*
 Claus Schäfer, *Munich*
 Hubert J Scheidbach, *Magdeburg*
 Peter Schemmer, *Heidelberg*
 Hans Scherübl, *Berlin*
 Thomas W Spahn, *Schwerte*
 Holger Sudhoff, *Bielefeld*

Jens Tischendorf, *Aachen*
 Michael Vieth, *Bayreuth*
 Jochen Wedemeyer, *Hannover*
 Uwe Will, *Gera*



Greece

Georgios K Anagnostopoulos, *Athens*
 Anna Eleftheriadou, *Rethymnon*
 Dimitris K Iakovidis, *Lamia*
 Dimitrios Kapetanios, *Thessaloniki*
 John A Karagiannis, *Athens*
 Stefanos Karagiannis, *Kifissia*
 Spiros D Ladas, *Athens*
 Konstantinos A Papadakis, *Heraklion*
 George H Sakorafas, *Athens*
 Elias Xirouchakis, *Areos*



Hungary

Pal Demeter, *Budapest*
 Lujber László, *Pecs*
 Peter Lakatos, *Budapest*
 István Rácz, *Gyor*



India

Ramanathan S Bharathi, *Uttar Pradesh*
 Devendra C Desai, *Mumbai*
 Evan L Fogel, *Indianapolis*
 Uday Chand Ghoshal, *Lucknow*
 Chittor M Habibullah, *Andhra Pradesh*
 Rakesh Kochhar, *Chandigarh*
 Rakesh Kumar, *New Delhi*
 Sri Prakash Misra, *Allahabad*
 Sandeep Nijhawan, *Rajasthan*
 Kaushal Kishor Prasad, *Chandigarh*
 Surinder Singh Rana, *Chandigarh*
 Muthukumaran Rangarajan, *Tamil Nadu*
 D Nageshwar Reddy, *Hyderabad*
 Omar Javed Shah, *Kashmir*
 Virendra Singh, *Chandigarh*



Iran

Tahereh Falsafi, *Tehran*
 Mohammad Rahnvardi, *Tehran*



Ireland

Colm Ó'Moráin, *Dublin*
 Eamonn M Quigley, *Cork*



Israel

Simon Bar-Meir, *Ramat Gan*
 Rami Eliakim, *Haifa*
 Zvi Fireman, *Hadea*
 Irina Hirsh, *Haifa*

Tiberiu Hershcovici, *Jerusalem*
 Jesse Lachter, *Haifa*



Italy

Paola De Angelis, *Rome*
 Paolo G Arcidiacono, *Milan*
 Alberto Arezzo, *Torino*
 Gabrio Bassotti, *San Sisto*
 Giampaolo Bresci, *Pisa*
 Carlo Calabrese, *Bologna*
 Salvatore MA Campo, *Rome*
 Federico Carpi, *Pisa*
 Livio Cipolletta, *Torre del Greco*
 Sandro Contini, *Parma*
 Salvatore Cucchiara, *Rome*
 Gabriele Curcio, *Palermo*
 Luigi Familiari, *Cavalluccio*
 Lorenzo Fuccio, *Bologna*
 Giuseppe Galloro, *Napoli*
 Giovanni B Gasbarrini, *Rome*
 Carlo M Girelli, *Busto Arsizio*
 Mauro Manno, *Baggiovara di Modena*
 Hugo Martines, *Savona*
 Gabriele Masselli, *Rome*
 Emanuele Meroni, *Milan*
 Andrea Moglia, *Pisa*
 Raffaele Pezzilli, *Bologna*
 Venerino Poletti, *Forli*
 Salvatore Pucciarelli, *Padova*
 Franco Radaelli, *Como*
 Marmo Riccardo, *Luigi Curto Polla*
 Maria Elena Riccioni, *Rome*
 Stefania Romano, *Naples*
 Emanuele Rondonotti, *Milano*
 Gianluca Rotondano, *Torre del Greco*
 Vittorio Terruzzi, *Como*
 Cristina Trovato, *Milano*
 Antonio Tucci, *Bologna*
 Maurizio Vecchi, *Milan*
 Maurizio Ventrucci, *Bologna*



Japan

Mitsuhiro Asakuma, *Osaka*
 Hiroki Endo, *Kanagawa*
 Shotaro Enomoto, *Wakayama*
 Kuang-I Fu, *Kashiwa*
 Makoto Hashizume, *Fukuoka*
 Toru Hiyama, *Higashihiroshima*
 Akira Hokama, *Okinawa*
 Akira Horiuchi, *Komagane*
 Kinichi Hotta, *Nagano*
 Atsushi Imagawa, *Kagawa*
 Hiroo Imazu, *Tokyo*
 Haruhiro Inoue, *Yokohama*
 Ryu Ishihara, *Osaka*
 Naoki Ishii, *Tokyo*
 Hajime Isomoto, *Nagasaki*
 Takao Itoi, *Tokyo*
 Satoru Kakizaki, *Gunma*
 Hiroshi Kakutani, *Tokyo*
 Terumi Kamisawa, *Tokyo*
 Yoshihide Kanno, *Sendai*
 Mototsugu Kato, *Sapporo*
 Takashi Kawai, *Tokyo*

Hirofumi Kawamoto, *Okayama*
 Hiroto Kita, *Saitama*
 Koga Komatsu, *Akita*
 Hitoshi Kondo, *Sapporo*
 Hiroaki Kubo, *Fukuoka*
 Keiichiro Kume, *Kitakyusyu*
 Iruru Maetani, *Tokyo*
 Hiroto Miwa, *Hyogo*
 Akihiro Mori, *Aichi*
 Akihiro Mori, *Aichi*
 Yoshihiro Moriwaki, *Yokohama*
 Naoki Muguruma, *Tokushima*
 Shinji Nishiwaki, *Gifu*
 Ichiro Oda, *Tokyo*
 Kazuichi Okazaki, *Osaka*
 Yasuhiro Oono, *Chiba*
 Taro Osada, *Tokyo*
 Yutaka Saito, *Tokyo*
 Yuzo Sakai, *Chiba*
 Naoto Sakamoto, *Tokyo*
 Nobuyuki Sakurazawa, *Tokyo*
 Yasushi Sano, *Hyogo*
 Tomoyuki Shibata, *Toyoake*
 Takashi Shida, *Chiba*
 Atsushi Sofuni, *Tokyo*
 Kazuki Sumiyama, *Tokyo*
 Nobumi Tagaya, *Tochigi*
 Hirokazu Takahashi, *Yokohama*
 Kyosuke Tanaka, *Mie*
 Shinji Tanaka, *Hiroshima*
 Gen Tohda, *Fukui*
 Tomoyuki Tsujikawa, *Shiga*
 Noriya Uedo, *Osaka*
 Shuji Yamamoto, *Kyoto*
 Takayuki Yamamoto, *Yokkaichi*
 Hideo Yanai, *Yamaguchi*
 Kenjiro Yasud, *Kyoto*
 Naohisa Yoshida, *Kyoto*



Lebanon

Kassem A Barada, *Beirut*



Lithuania

Laimas Virginijus Jonaitis, *Kaunas*



Malaysia

Sanjiv Mahadeva, *Kuala Lumpur*
 Sreenivasan Sasidharan, *Pulau Pinang*



Mexico

OT Teramoto-Matsubara, *México*



Netherlands

Marco Bruno, *Rotterdam*
 Dirk Joan Gouma, *Amsterdam*
 Iris Lansdorp-Vogelaar, *Rotterdam*
 Chris JJ Mulder, *Amsterdam*

Vasileios Panteris, *Rotterdam*
 Harald Erwin Vonkeman, *Enschede*



New Zealand

Michael PG Schultz, *Dunedin*



Norway

Magdy El-Salhy, *Stord*
 Odd Helge Gilja, *Bergen*



Pakistan

Syed H Ali Shah, *Karachi*
 Lubna Kamani, *Karachi*



Poland

Stanislaw A Hac, *Gdansk*
 Maciej Michalik, *Pomorskie*



Portugal

Miguel T Coimbra, *Porto*
 Marie I Cremers, *Setúbal*
 Mário Dinis-Ribeiro, *Porto*
 Pedro N Figueiredo, *Coimbra*
 Rui MA da Silva, *Porto*



Romania

Mihai Ciocirlan, *Bucharest*
 Lucian Negreanu, *Bucharest*



Singapore

Zhiwei Huang, *Singapore*
 Surendra K Mantoo, *Singapore*



South Africa

Roland N Ndip, *Alice*



South Korea

Young-Tae Bak, *Seoul*
 Dong Kyung Chang, *Seoul*
 Youn-Seok Cho, *UiJeongbu*
 Seong Woo Jeon, *Daegu*
 Jong-Man Kang, *Seoul*
 Yong Sung Kim, *Gyeonggi-do*
 Hang Lak Lee, *Sungdonggu*
 Suck-Ho Lee, *Cheonan*
 Jong Ho Moon, *Bucheon*
 Dong Kyun Park, *Incheon*
 Dae Kyung Sohn, *Gyeonggi*

Jaekyu Sung, *Daejeon*
 Si-Young Song, *Seoul*



Spain

Jose FN Aguilar, *Palma*
 Adolfo P Blanco, *Asturias*
 Andres Cardenas, *Barcelona*
 Gloria Fernández-Esparrach, *Barcelona*
 Jesús García-Cano, *Cuenca*
 Angels Gines, *Barcelona*
 Angel Lanas, *Zaragoza*
 G Payeras Llodrá, *Madrid*
 Alfredo José Lucendo, *Tomelloso*
 Enrique F Perez-Cuadrado Martinez, *Murcia*
 Luis Rabago, *Madrid*
 Eduardo Redondo-Cerezo, *Cuenca*
 Luis Rodrigo, *Oviedo*
 Jaume Boix Valverde, *Badalona*
 Josep Llach Vila, *Barcelona*
 Santiago Vivas, *León*



Sweden

George Dafnis, *Eskilstuna*
 Per-Ola Park, *Borås*
 Carlos A Rubio, *Stockholm*



Thailand

Somchai Amornytin, *Bangkok*
 Thawatchai Akaraviputh, *Bangkok*
 Udom Kachintorn, *Bangkok*
 Varut Lohsirawat, *Bangkok*
 Rungsun Rerknimitr, *Bangkok*



Turkey

Selcuk Disibeyaz, *Nkara*
 Mehmet Eken, *Istanbul*
 Muammer Kara, *Ankara*
 Taylan Kav, *Ankara*
 Nevin Oruc, *İzmir*
 Burhan Ozdil, *Adana*
 Nurdan Ozmeric, *Emek Ankara*
 Sema Zer Toros, *Istanbul*



United Arab Emirates

Margit Gabriele Muller, *Abu Dhabi*



United Kingdom

Basil J Ammori, *Manchester*
 Simon HC Anderson, *London*
 Adam D Farmer, *London*
 Annette Fritscher-Ravens, *Landon*
 Gianpiero Gravante, *Bristol*
 Abdulzahra Hussain, *London*
 United KV Kodogiannis, *London*
 Seamus J Murphy, *Newry*
 Perminder Phull, *Aberdeen*

Krish Ragnath, *Nottingham*
Jayesh Sagar, *Wishaw*
Reena Sidhu, *Sheffield*
Adrian J Stanley, *Glasgow*
Hu Zhang, *Cambridge*



United States

Maher Aref Abbas, *Los Angeles*
Douglas G Adler, *Utah*
Deepak Agrawal, *Dallas*
Mohammad Al-Haddad, *Indianapolis*
Jamie S Barkin, *Florida*
Pedro W Baron, *Loma Linda*
James Stephen Barthel, *Florida*
Neil Bhattacharyya, *Boston*
Juliane Bingener-Casey, *Rochester*
Cheri Lee Canon, *Birmingham*
Sherman M Chamberlain, *Georgia*
Lawrence B Cohen, *New York*
Lawrence Bruce Cohen, *New York*
Paul G Curcillo II, *Philadelphia*
Kiron M Daskiron, *New Brunswick*
David J Desilets, *Springfield*

John C Deutsch, *Duluth*
Peter Draganov, *Gainesville*
Viktor Ernst Eysselein, *Torrance*
Daniel L Farkas, *Los Angeles*
Ronnie Fass, *Southern Arizona*
Georg Feldmann, *Maryland*
Raja M Flores, *New York*
Catherine T Frenette, *San Francisco*
David Friedel, *New York*
Ronnie Fass, *Tucson*
Seng-Ian Gan, *Seattle*
Denise W Gee, *Massachusetts*
Samuel A Giday, *Maryland*
George F Gowen, *Pottstown*
Sammy Ho, *New York*
Moises Jacobs, *Florida*
Robert Thomas Jensen, *Bethesda*
Michel Kahaleh, *Virginia*
Peter James Kahrilas, *Suite*
Sergey V Kantsevov, *Baltimore*
Christopher Lawrence, *Charleston*
Felix W Leung, *Sepulveda*
Simon K Lo, *California*
Charles Maltz, *New York*
Jeffrey Michael Marks, *Ohio*
Hiroshi Mashimo, *Massachusetts*

Abraham Mathew, *Hershey*
James M Mullin, *Wynnewood*
Harvey J Murff, *Nashville*
Koichi Nagata, *Boston*
Ying-Tian Pan, *Stony Brook*
Jitesh A Patel, *Pittsburgh*
Massimo Raimondo, *Jacksonville*
Amit Rastogi, *Kansas City*
Robert J Richards, *New York*
Praveen Roy, *New Mexico*
David T Rubin, *Chicago*
Enrique Seoane-Vazquez, *Columbus*
Prateek Sharma, *Kansas*
Bo Shen, *Ohio*
Danny A Sherwinter, *Brooklyn*
Andrew Ukleja, *Weston*
Bennie Ray Upchurch, *Ohio*
Shyam Varadarajulu, *Alabama*
Marcelo F Vela, *South Carolina*
Wahid Wassef, *Worcester*
Irving Waxman, *Illinois*
C Mel Wilcox, *Alabama*
Field Farrar Willingham, *Massachusetts*
Timothy A Woodward, *Jacksonville*
Shuhei Yoshida, *Massachusetts*



Contents

Monthly Volume 3 Number 1 January 16, 2011

EDITORIAL

- 1 Endoscopic diagnosis of pancreaticobiliary maljunction
Kamisawa T, Takuma K, Itokawa F, Itoi T

BRIEF ARTICLE

- 6 Mini-laparoscopy in the endoscopy unit: Safety and outcomes in over one thousand patients
Hoffman A, Rahman F, Prengel S, Schuchmann M, Goetz M, Moehler M, Galle PR, Li Z, Kalloo AN, Kiesslich R
- 11 Ultraslim endoscopy with flexible spectral imaging color enhancement for upper gastrointestinal neoplasms
Tanioka Y, Yanai H, Sakaguchi E

CASE REPORT

- 16 Duodenal tuberculosis presenting as gastric outlet obstruction: A case report
Flores HB, Zano F, Ang EL, Estanislao N
- 20 Endoscopic retrieval of a gastric trichobezoar
Konuma H, Fu K, Morimoto T, Shimizu T, Izumi Y, Shiyanagi S, Urao M, Miyazaki A, Watanabe S

Contents

World Journal of Gastrointestinal Endoscopy
Volume 3 Number 1 January 16, 2011

ACKNOWLEDGMENTS I Acknowledgments to reviewers of *World Journal of Gastrointestinal Endoscopy*

APPENDIX I Meetings
I-V Instructions to authors

ABOUT COVER Tanioka Y, Yanai H, Sakaguchi E. Ultraslim endoscopy with flexible spectral imaging color enhancement for upper gastrointestinal neoplasms
World J Gastrointest Endosc 2011; 3(1): 11-15
<http://www.wjgnet.com/1948-5190/full/v3/i1/11.htm>

AIM AND SCOPE *World Journal of Gastrointestinal Endoscopy* (*World J Gastrointest Endosc*, *WJGE*, online ISSN 1948-5190, DOI: 10.4253), is a monthly, open-access, peer-reviewed journal supported by an editorial board of 400 experts in gastrointestinal endoscopy from 45 countries.
The major task of *WJGE* is to report rapidly the most recent results in basic and clinical research on gastrointestinal endoscopy including: gastroscopy, intestinal endoscopy, colonoscopy, capsule endoscopy, laparoscopy, interventional diagnosis and therapy, as well as advances in technology. Emphasis is placed on the clinical practice of treating gastrointestinal diseases with or under endoscopy. Papers on advances and application of endoscopy-associated techniques, such as endoscopic ultrasonography, endoscopic retrograde cholangiopancreatography, endoscopic submucosal dissection and endoscopic balloon dilation are also welcome.

FLYLEAF I-IV Editorial Board

EDITORS FOR THIS ISSUE

Responsible Assistant Editor: Na Liu
Responsible Electronic Editor: Na Lin
Proofing Editor-in-Chief: Lian-Sheng Ma

Responsible Science Editor: Hai-Ning Zhang
Proofing Editorial Office Director: Hai-Ning Zhang

NAME OF JOURNAL
World Journal of Gastrointestinal Endoscopy

LAUNCH DATE
October 15, 2009

SPONSOR
Beijing Baishideng BioMed Scientific Co., Ltd.,
Room 903, Building D, Ocean International Center,
No. 62 Dongsihuan Zhonglu, Chaoyang District,
Beijing 100025, China
Telephone: +86-10-8538-1892
Fax: +86-10-8538-1893
E-mail: baishideng@wjgnet.com
<http://www.wjgnet.com>

EDITING
Editorial Board of *World Journal of Gastrointestinal Endoscopy*,
Room 903, Building D, Ocean International Center,
No. 62 Dongsihuan Zhonglu, Chaoyang District,
Beijing 100025, China
Telephone: +86-10-5908-0038
Fax: +86-10-8538-1893
E-mail: wjge@wjgnet.com
<http://www.wjgnet.com>

PUBLISHING
Baishideng Publishing Group Co., Limited,
Room 1701, 17/F, Henan Building,
No.90 Jaffe Road, Wanchai,
Hong Kong, China
Fax: 00852-3115-8812
Telephone: 00852-5804-2046
E-mail: baishideng@wjgnet.com
<http://www.wjgnet.com>

SUBSCRIPTION
Beijing Baishideng BioMed Scientific Co., Ltd.,
Room 903, Building D, Ocean International Center,
No. 62 Dongsihuan Zhonglu, Chaoyang District,
Beijing 100025, China
Telephone: +86-10-8538-1892
Fax: +86-10-8538-1893
E-mail: baishideng@wjgnet.com
<http://www.wjgnet.com>

ONLINE SUBSCRIPTION
One-Year Price: 216.00 USD

PUBLICATION DATE
January 16, 2011

CSSN
ISSN 1948-5190 (online)

PRESIDENT AND EDITOR-IN-CHIEF
Lian-Sheng Ma, *Beijing*

STRATEGY ASSOCIATE EDITORS-IN-CHIEF
Kazuya Akahoshi, *Iizuka*
William Robert Brugge, *Massachusetts*
Qiang Cai, *Georgia*
Juan J Vila Costas, *Pamplona*
Atsushi Irisawa, *Fukushima*
Andreas Sieg, *Heidelberg*
Gaetana Ilaria Tarantino, *Palermo*
Tony CK Tham, *Northern Ireland*
Konstantinos Triantafyllou, *Haidari*

EDITORIAL OFFICE
Hai-Ning Zhang, Director
World Journal of Gastrointestinal Endoscopy
Room 903, Building D, Ocean International Center,
No. 62 Dongsihuan Zhonglu, Chaoyang District,
Beijing 100025, China
Telephone: +86-10-5908-0038
Fax: +86-10-8538-1893
E-mail: wjge@wjgnet.com
<http://www.wjgnet.com>

COPYRIGHT
© 2011 Baishideng. All rights reserved; no part of this publication may be commercially reproduced, stored in a retrieval system, or transmitted in any form or by any means, electronic, mechanical, photocopying, recording, or otherwise without the prior permission of Baishideng. Authors are required to grant *World Journal of Gastrointestinal Endoscopy* an exclusive license to publish.

SPECIAL STATEMENT
All articles published in this journal represent the viewpoints of the authors except where indicated otherwise.

INSTRUCTIONS TO AUTHORS
Full instructions are available online at http://www.wjgnet.com/1948-5190/g_info_20100316080002.htm. If you do not have web access please contact the editorial office.

ONLINE SUBMISSION
<http://www.wjgnet.com/1948-5190office/>

Endoscopic diagnosis of pancreaticobiliary maljunction

Terumi Kamisawa, Kensuke Takuma, Fumihide Itokawa, Takao Itoi

Terumi Kamisawa, Kensuke Takuma, Department of Internal Medicine, Tokyo Metropolitan Komagome Hospital, Tokyo 113-8677, Japan

Fumihide Itokawa, Takao Itoi, Department of Gastroenterology, Tokyo Medical University, Tokyo 113-8677, Japan

Author contributions: Kamisawa T and Itoi T wrote the paper; and Kamisawa T, Takuma K, Itoi T and Itokawa F gathered the data.

Correspondence to: Terumi Kamisawa, MD, Department of Internal Medicine, Tokyo Metropolitan Komagome Hospital, 3-18-22 Honkomagome, Bunkyo-ku, Tokyo 113-8677, Japan. kamisawa@cick.jp

Telephone: +81-3-38232101 Fax: +81-3-38241552

Received: October 15, 2010 Revised: December 10, 2010

Accepted: December 17, 2010

Published online: January 16, 2011

Abstract

Pancreaticobiliary maljunction (PBM) is a congenital anomaly defined as a junction of the pancreatic and bile ducts located outside the duodenal wall, usually forming a markedly long common channel. As the action of the sphincter of Oddi does not functionally affect the junction in PBM patients, continuous pancreatobiliary reflux occurs, resulting in a high incidence of biliary cancer. PBM can be divided into PBM with biliary dilatation (congenital choledochal cyst) and PBM without biliary dilatation (maximal diameter of the bile duct \leq 10 mm). The treatment of choice for PBM is prophylactic surgery before malignant changes can take place. Endoscopic retrograde cholangiopancreatography (ERCP) is the most effective examination method for close observation of the pattern of the junction site. When the communication between the pancreatic and bile ducts is maintained, despite contraction of the sphincter on ERCP, PBM is diagnosed. In these patients, levels of pancreatic enzymes in the bile are generally elevated, due to continuous pancreatobiliary reflux *via* a long common channel. Magnetic resonance cholangiopancreatography and 3D-computed tomography can diagnose PBM, based on findings of an anomalous union between the common bile duct and

the pancreatic duct, in addition to a long common channel. Endoscopic ultrasonography and intraductal ultrasonography can demonstrate the junction outside the duodenal wall, and are useful for the diagnosis of associated biliary cancer. Gallbladder wall thickness on ultrasonography can be a screening test for PBM.

© 2011 Baishideng. All rights reserved.

Key words: Pancreaticobiliary maljunction; Pancreatobiliary reflux; Congenital choledochal cyst; Endoscopic retrograde cholangiopancreatography; Endoscopic ultrasonography; Magnetic resonance cholangiopancreatography

Peer reviewers: Omar Javed Shah, Professor, Head, Department of Surgical Gastroenterology, Sher-i-Kashmir Institute of Medical Sciences, Srinagar, Kashmir, India; Sanjiv Mahadeva, MBBS, MRCP, CCST, MD, Associate Professor, Department of Medicine, Faculty of Medicine, University of Malaya, Kuala Lumpur 50603, Malaysia

Kamisawa T, Takuma K, Itokawa F, Itoi T. Endoscopic diagnosis of pancreaticobiliary maljunction. *World J Gastrointest Endosc* 2011; 3(1): 1-5 Available from: URL: <http://www.wjgnet.com/1948-5190/full/v3/i1/1.htm> DOI: <http://dx.doi.org/10.4253/wjge.v3.i1.1>

INTRODUCTION

The main pancreatic duct and the common bile duct open into the duodenum, where they frequently form a common channel, the incidence of which is reported to be from 55%^[1] to 82%^[2]. The length of the common channel ranges from 1 mm to 12 mm, with an average length of 4.4 mm^[3]. The sphincter of Oddi is located at the distal end of the pancreatic and bile ducts; it regulates the outflow of bile and pancreatic juice. A common channel can be so long that the sphincter action does not functionally affect the junction, resulting in two-way regurgitation (pancreatobiliary reflux: regurgitation of pancreatic juice into the common bile duct, and biliopancreatic reflux:

regurgitation of bile juice into the pancreatic duct). Pancreatobiliary maljunction (PBM) forms a markedly long common channel, and is divided into PBM with, and without, biliary dilatation^[4-7]. Pancreatobiliary reflux has been shown to induce carcinogenesis in the biliary tract; similarly, biliopancreatic reflux can induce pancreatitis^[4-6].

PANCREATICOBILIARY MALJUNCTION

PBM is a congenital anomaly defined as a junction of the pancreatic and bile ducts located outside the duodenal wall, usually forming a markedly long common channel. PBM occurs predominantly in females, and is often found in Asian populations. PBM can be divided into PBM with biliary dilatation (congenital choledochal cyst (CCC)) (Figure 1) and PBM without biliary dilatation (Figure 2)^[4-7]. CCC is an anomaly in which the extrapancreatic bile duct, or the extra and intrahepatic bile ducts, are dilated in various ways. The Alonso-Lej classification^[8], which is based on the shape of the dilated bile duct, is notable, and includes a cystic type (Type I), a diverticular type (Type II), and a cyst in the duodenum (choledochal cyst, Type III). Type I is almost always associated with PBM, but Type II and III rarely have PBM. Bile duct diameter greater than 10 mm on a cholangiogram is the most commonly used definition of dilatation^[7].

In PBM patients, since the action of the sphincter of Oddi does not functionally affect the junction, continuous reciprocal reflux between pancreatic juice and bile occurs, resulting in various pathological conditions in the biliary tract and pancreas. As the hydro pressure within the pancreatic duct is usually greater than that in the bile duct, pancreatic juice frequently refluxes into the bile duct in PBM patients, which results in a high incidence of cancer in the biliary tract^[4-7]. In our previous study, bile duct and gallbladder cancers were seen in 14% and 22% of 49 CCC patients, respectively, but, in 70% of 53 PBM patients without biliary dilatation, only gallbladder cancer was detected.

Once PBM is diagnosed, prophylactic flow-diversion surgery (bile duct resection and bilioenteric anastomosis) is performed for CCC. On the other hand, any treatment of PBM without biliary dilatation and without cancer is controversial. Prophylactic cholecystectomy is performed in many institutes, as most biliary cancers that develop in PBM patients without biliary dilatation are gallbladder cancers. However, some surgeons propose excision of the extrahepatic bile duct, together with the gallbladder, for PBM patients without biliary dilatation, because of the risk of bile duct cancer^[9,10].

ENDOSCOPIC RETROGRADE CHOLANGIO-PANCREATOGRAPHY

Endoscopic retrograde cholangiopancreatography (ERCP) is the most effective examination method for close observation of the pattern of the junction site. When the communication between the pancreatic and bile ducts is



Figure 1 Magnetic resonance cholangiopancreatography showing pancreaticobiliary maljunction with biliary dilatation (congenital choledochal cyst).

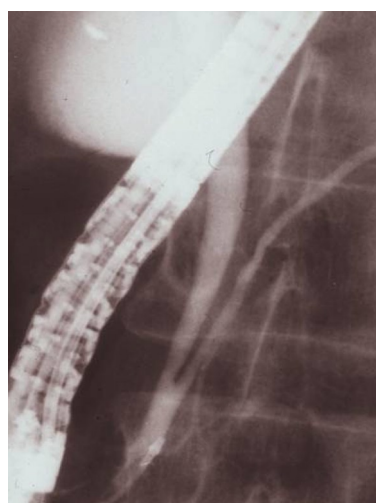


Figure 2 Endoscopic retrograde cholangiopancreatography showing pancreaticobiliary maljunction without biliary dilatation.

maintained despite contraction of the sphincter on ERCP, PBM is diagnosed (Figure 3A and B). The finding can also be assessed with cholangiography, *via* the biliary drainage tube or during operation^[4-7].

Pancreatography *via* the minor duodenal papilla can also demonstrate pancreatobiliary reflux in PBM patients. When injected endoscopically *via* the minor duodenal papilla, the contrast medium is refluxed into the bile duct through a long common channel without outflow into the duodenum^[5].

Biliary levels of pancreatic enzymes, especially amylase, are generally elevated due to continuous pancreatobiliary reflux *via* a long common channel in PBM patients. There are some cases with a relatively long common channel that are not classified as PBM because the sphincter of Oddi includes the pancreaticobiliary ductal junction. We defined a high confluence of pancreaticobiliary ducts (HCPBD) as a common channel length ≥ 6 mm, in which the communication was occluded when the sphincter was contracted (Figure 4A and B), and investigated the clinical significance of a relatively long common channel^[5-7]. In HCPBD patients, the amylase level in the bile was frequently elevated, and hyperplastic change of the gallbladder epithelium was frequently observed. Gallbladder cancer occurred in

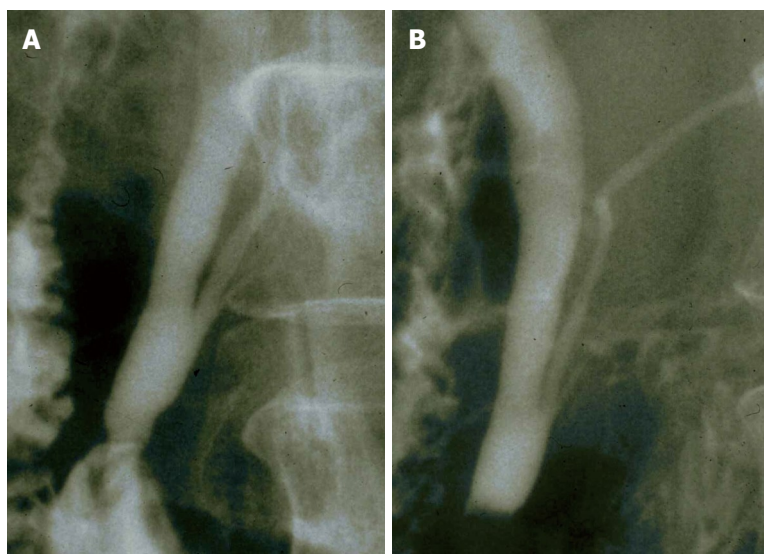


Figure 3 Endoscopic retrograde cholangiopancreatography of a pancreaticobiliary maljunction patient. A: Showing a long common channel; B: The communication between pancreatic and bile ducts was maintained despite contraction of the sphincter.

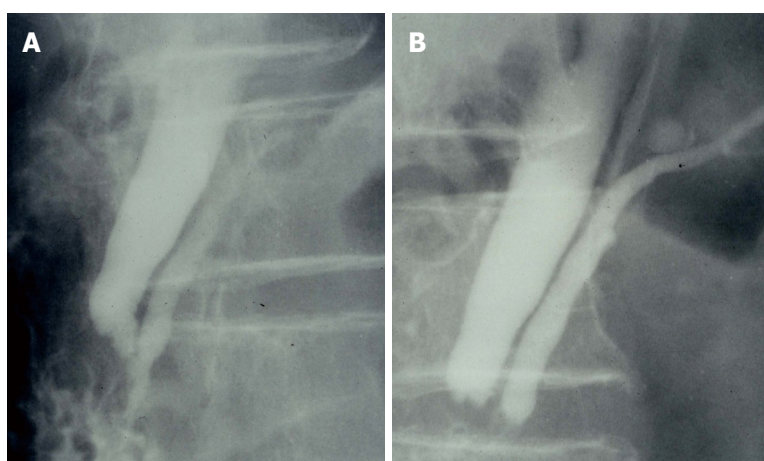


Figure 4 Endoscopic retrograde cholangiopancreatography of a patient with high confluence of pancreaticobiliary ducts. A: A common channel of 9 mm in length; B: The communication between pancreatic and bile ducts was destroyed by sphincter contraction.

11% of the 84 HCPBD patients. These findings suggest that pathophysiological changes similar to PBM occur in HCPBD. However, in HCPBD patients, there was no difference between the sexes, whereas in PBM there was (male:female ratio, 1 : 0.7 in HCPBD and 1 : 3.4 in PBM), and the average age at the time of diagnosis was significantly older in HCPBD patients than in PBM patients without biliary dilatation (average age, 62.0 years *vs* 56.7 years, respectively). The elevated amylase level in the bile in HCPBD patients was lower than that of PBM patients (average 48 665 IU/L *vs* 250 025 IU/L, respectively) and the rate of associated gallbladder cancer was lower in HCPBD patients than in PBM patients. These differences in sex, age at diagnosis, bile amylase level, and rate of associated gallbladder cancer between HCPBD and PBM patients appear to be related to whether pancreatobiliary reflux occurs consistently or intermittently^[5-7].

MAGNETIC RESONANCE CHOLANGIOPANCREATOGRAPHY AND 3D-COMPUTED TOMOGRAPHY

Magnetic resonance cholangiopancreatography (MRCP)

has become a common non-invasive method for obtaining high quality images of the pancreaticobiliary tree.

Reconstruction images on 3D-computed tomography (CT) can also show pancreaticobiliary images. MRCP and 3D-CT can diagnose PBM, based on findings of an anomalous union between the common bile duct and the pancreatic duct, in addition to a long common channel. However, in some cases in which a common channel is not so long and cannot be depicted on MRCP, the MRCP diagnosis of PBM is not possible^[11]. For a definite diagnosis of PBM, ERCP is necessary in order to exclude various false positive or negative results on MRCP or 3D-CT. Diagnostic accuracy can be increased with dynamic MRCP with secretin stimulation or three-dimensional MRCP. Although a large amount of contrast must be injected to evaluate the whole image of CCC on ERCP, it can be achieved with MRCP and 3D-CT.

Pancreatobiliary reflux in PBM patients can be visualized radiologically using secretin-stimulated dynamic MRCP. In normal pancreaticobiliary dynamics, the extrahepatic and intrahepatic bile ducts show no change following secretin injection. On the other hand, in PBM patients, the volume of the extrahepatic bile duct and the gallbladder increases, due to the regurgitation of pan-

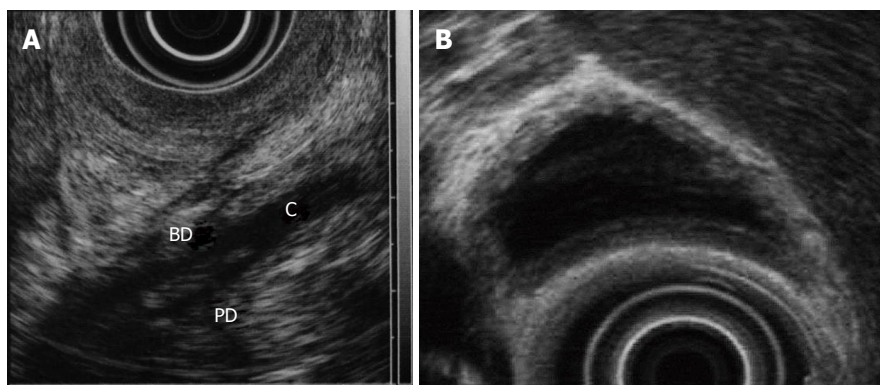


Figure 5 Endoscopic ultrasonography of a pancreaticobiliary maljunction patient. A: The confluence of pancreatic duct and bile duct in the proximal portion of the duodenal wall; B: thickness of inner low echoic layer of the gallbladder. BD: bile duct; PD: pancreatic duct; C: common channel.

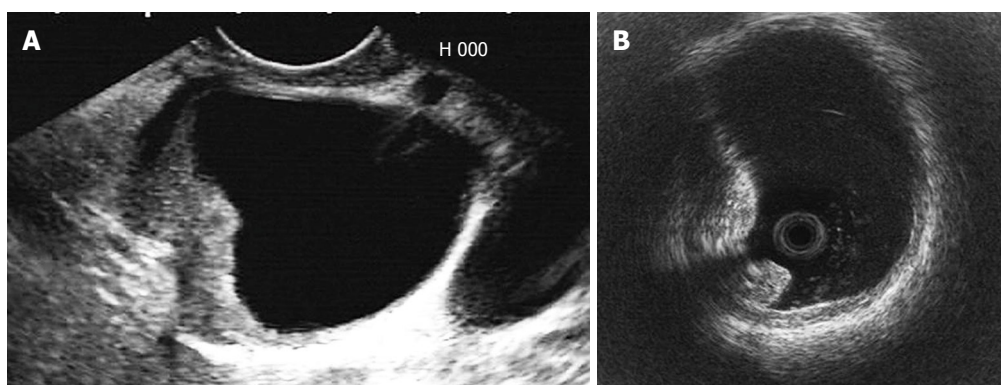


Figure 6 Ultrasonography of a congenital choledochal cyst patient with bile duct cancer in the dilated bile duct. A: Endoscopic ultrasonography; B: Intraductal ultrasonography.

creatic fluid secreted after the injection of secretin into the bile duct^[12].

ENDOSCOPIC ULTRASONOGRAPHY

In PBM, endoscopic ultrasonography (EUS) can detect the confluence of pancreatic duct and bile duct in the proximal portion of the duodenal wall, the so-called common channel (Figure 5A). Therefore, EUS allows definite diagnosis of PBM. When PBM is known about in advance, it may be relatively easy to depict the confluence. PBM often shows a thickness of the inner low echoic layer of the gallbladder (Figure 5B), which means histologically mucosal hyperplasia. Therefore, when we find that sonographic finding by means of EUS, we have to keep the presence of PBM in mind. In fact, several investigators have reported that EUS could confirm the PBM in 4 (2.9%) of 137 patients who underwent screening US^[13].

Bile duct and gallbladder cancers are often seen in PBM. The efficacy of EUS for the diagnosis of gallbladder cancer is well known^[14]. However, it is often difficult, even when using EUS, to distinguish mucosal hyperplasia from early gallbladder cancer to indicate whether the depth of invasion of the cancer is mucosa or muscularis propria of the gallbladder.

EUS is also useful for the diagnosis of bile duct cancer. Since, in particular, congenital bile duct dilation is well-known as a high risk condition for bile duct cancer in the dilated bile duct (Figure 6A), in these cases, EUS should be preoperatively performed for the diagnosis of tumor spreading and staging.

INTRADUCTAL ULTRASONOGRAPHY

Intraductal ultrasonography (IDUS) is performed over-the-guidewire during the ERCP, and is useful for the depiction of the confluence of pancreatic duct and bile duct.

IDUS is also useful for the diagnosis of bile duct cancer (Figure 6B)^[15,16]. However, IDUS has limitations for the diagnosis of bile duct and gallbladder lesions because of shallow US penetration (< 2.0 cm) and maneuverability of passage of probe in case of bile duct stricture or a narrow cystic duct.

Direct choledochoscopy with or without biopsy is useful to confirm the diagnosis of bile duct cancer.

ULTRASONOGRAPHY

Mark dilatation of the common bile duct on ultrasonography (US) suggests CCC. The epithelium of the gallbladder frequently becomes hyperplastic due to exposure to refluxed pancreatic juice. Gallbladder wall thickness on US during medical checkups may serve as an indication of PBM without the need for biliary dilatation, and can serve as a screening test for PBM^[17].

CONCLUSION

When the communication between the pancreatic and bile ducts is maintained despite contraction of the sphincter on ERCP, PBM is diagnosed. MRCP and 3D-CT can be used to diagnose PBM, based on the discovery of an anomalous union between the common bile duct and the

pancreatic duct, in addition to a long common channel. EUS and IDUS can demonstrate the junction outside the duodenal wall, and they are useful for the diagnosis of associated biliary cancer. Gallbladder wall thickness on US can be a screening test for PBM.

REFERENCES

- 1 **Sterling JA.** The common channel for bile and pancreatic ducts. *Surg Gynecol Obstet* 1954; **98**: 420-424
- 2 **Suda K,** Miyano T, Konuma I, Matsumoto M. An abnormal pancreatiko-cholecho-ductal junction in cases of biliary tract carcinoma. *Cancer* 1983; **52**: 2086-2088
- 3 **Dowdy GS Jr,** Waldron GW, Brown WG. Surgical anatomy of the pancreatobiliary ductal system. Observations. *Arch Surg* 1962; **84**: 229-246
- 4 The Japanese Study Group on Pancreaticobiliary Maljunction. Diagnostic criteria of pancreaticobiliary maljunction. *J Hepatobiliary Pancreat Surg* 1994; **1**: 219-221
- 5 **Kamisawa T,** Okamoto A. Biliopancreatic and pancreatobiliary refluxs in cases with and without pancreaticobiliary maljunction: diagnosis and clinical implications. *Digestion* 2006; **73**: 228-236
- 6 **Kamisawa T,** Anjiki H, Egawa N, Kurata M, Honda G, Tsuruta K. Diagnosis and clinical implications of pancreatobiliary reflux. *World J Gastroenterol* 2008; **14**: 6622-6626
- 7 **Kamisawa T,** Takuma K, Anjiki H, Egawa N, Kurata M, Honda G, Tsuruta K, Sasaki T. Pancreaticobiliary maljunction. *Clin Gastroenterol Hepatol* 2009; **7**: S84-S88
- 8 **Alonso-Lej F,** Rever EB Jr, Pessagno DJ. Congenital choledochal cyst. *Int Abstr Surg* 1959; **108**: 1-30
- 9 **Tashiro S,** Imaizumi T, Ohkawa H, Okada A, Katoh T, Kawaharada Y, Shimada H, Takamatsu H, Miyake H, Todani T. Pancreaticobiliary maljunction: retrospective and nationwide survey in Japan. *J Hepatobiliary Pancreat Surg* 2003; **10**: 345-351
- 10 **Funabiki T,** Matsubara T, Miyakawa S, Ishihara S. Pancreaticobiliary maljunction and carcinogenesis to biliary and pancreatic malignancy. *Langenbecks Arch Surg* 2009; **394**: 159-169
- 11 **Kamisawa T,** Tu Y, Egawa N, Tsuruta K, Okamoto A, Kamata N. MRCP of congenital pancreaticobiliary malformation. *Abdom Imaging* 2007; **32**: 129-33
- 12 **Hosoki T,** Hasuike Y, Takeda Y, Michita T, Watanabe Y, Sakamori R, Tokuda Y, Yutani K, Sai C, Mitomo M. Visualization of pancreaticobiliary reflux in anomalous pancreaticobiliary junction by secretin-stimulated dynamic magnetic resonance cholangiopancreatography. *Acta Radiol* 2004; **45**: 375-382
- 13 **Yamao K,** Mizutani S, Nakazawa S, Inui K, Kanemaki N, Miyoshi H, Segawa K, Zenda H, Kato T. Prospective study of the detection of anomalous connections of pancreatobiliary ducts during routine medical examinations. *Hepatogastroenterology* 1996; **43**: 1238-1245
- 14 **Fujita N,** Noda Y, Kobayashi G, Kimura K, Yago A. Diagnosis of the depth of invasion of gallbladder carcinoma by EUS. *Gastrointest Endosc* 1999; **50**: 659-663
- 15 **Tamada K,** Ido K, Ueno N, Kimura K, Ichihama M, Tomiyama T. Preoperative staging of extrahepatic bile duct cancer with intraductal ultrasonography. *Am J Gastroenterol* 1995; **90**: 239-246
- 16 **Noda Y,** Fujita N, Kobayashi G, Ito K, Horaguchi J, Takazawa O, Obana T, Nakahara K, Ishida K, Suzuki T, Hirasawa D, Sugawara T, Ohira T, Onochi K, Harada Y, Tsuchiya T, Sawai T, Uzuki M, Kariya Y. Intraductal ultrasonography before biliary drainage and transpapillary biopsy in assessment of the longitudinal extent of bile duct cancer. *Dig Endosc* 2008; **20**: 73-78
- 17 **Sugai M,** Ishido K, Endoh M, Hada R, Munakata H. Sonographic demonstration of wall thickness of the gallbladder in pediatric patients with pancreatico-biliary maljunction. *J Hepatobiliary Pancreat Sci* 2010; **17**: 345-348

S- Editor Zhang HN L- Editor Herholdt A E- Editor Liu N

Mini-laparoscopy in the endoscopy unit: Safety and outcomes in over one thousand patients

Arthur Hoffman, Fareed Rahman, Sarah Prengel, Marcus Schuchmann, Martin Gotz, Markus Moehler, Peter Robert Galle, Ziping Li, Anthony Nicholas Kalloo, Ralf Kiesslich

Arthur Hoffman, Fareed Rahman, Sarah Prengel, Marcus Schuchmann, Martin Gotz, Markus Moehler, Peter Robert Galle, Ralf Kiesslich, First Department of Internal Medicine, Johannes Gutenberg University Mainz, Mainz 55101, Germany
Ziping Li, Anthony Nicolas Kalloo, Department of Gastroenterology and Hepatology, Johns Hopkins University, Baltimore, MD 21287, United States

Author contributions: Hoffmann A and Rahman F contributed equally to this article; Hoffman A was responsible for study design, performance of mini-laparoscopy and data analysis; Rahman F was responsible for data analysis, preparation of the manuscript and performance of mini-laparoscopy; Prengel S collected and analyzed data; Schuchmann M, Gotz M and Moehler M participated in performance of mini-laparoscopy; Galle PR was the chief of department and was involved with data discussion; Kiesslich R was the chief of the endoscopy branch and was involved with data discussion; and Li Z and Kalloo AN also participated in the study design and discussion.

Correspondence to: Arthur Hoffman, MD, First Department of Internal Medicine, Johannes Gutenberg University of Mainz, Langenbeckstr. 1, Mainz 55101, Germany. ahoff66286@aol.com

Telephone: +49-6131-177299 Fax: +49-6131-175552

Received: September 19, 2010 Revised: December 6, 2010

Accepted: December 13, 2010

Published online: January 16, 2011

Abstract

AIM: To investigate the safety of consecutive mini-laparoscopy guided liver biopsies for the diagnosis and staging of liver diseases.

METHODS: In this study we retrospectively analyzed the safety of mini-laparoscopic liver biopsy performed in an endoscopy unit in 1071 patients. We measured the incidence of bleeding and evaluated the management and outcome of bleeding interventions.

RESULTS: The most common etiologies of liver injury

were viral hepatitis and autoimmune liver disease. 250 patients had macroscopically and histologically proven cirrhosis. 13 patients had no pathological findings. 33% of all patients had bleeding that required argon plasma coagulation of the puncture site during laparoscopy. Significant bleeding occurred more often in patients with liver cirrhosis compared to non-cirrhotic liver diseases but was effectively treated with laparoscopic coagulation.

CONCLUSION: In conclusion, mini-laparoscopy liver biopsy can be performed safely and effectively in high risk patients with advanced liver disease; mini-laparoscopy with liver biopsy can be done safely in an endoscopy unit.

© 2011 Baishideng. All rights reserved.

Key words: Mini-laparoscopy; Cirrhosis; Argon plasma coagulation

Peer reviewers: Oliver Pech, MD, PhD, Attending Physician of Gastroenterology, Vice Director of the Endoscopy Unit, Department of Internal Medicine 2, HSK Wiesbaden, Wiesbaden, Germany; Takashi Shida, MD, PhD, Department of General Surgery, Chiba University Graduate School of Medicine, 1-8-1 Inohana, Chuo-ku, Chiba 260-8670, Japan

Hoffman A, Rahman F, Prengel S, Schuchmann M, Goetz M, Moehler M, Galle PR, Li Z, Kalloo AN, Kiesslich R. Mini-laparoscopy in the endoscopy unit: Safety and outcomes in over one thousand patients. *World J Gastrointest Endosc* 2011; 3(1): 6-10 Available from: URL: <http://www.wjgnet.com/1948-5190/full/v3/i1/6.htm> DOI: <http://dx.doi.org/10.4253/wjge.v3.i1.6>

INTRODUCTION

Although noninvasive techniques for the diagnosis and

staging of liver disease have been developed in recent years, histological evaluation still remains the most accurate method for the assessment of severity and stage of liver diseases^[1,2]. However, in patients with acute or advanced liver disease, there may be an increased risk of bleeding complications with percutaneous liver biopsy, even when done under ultrasound guidance^[3,4]. Furthermore, recent studies have shown that liver histology obtained by blind biopsy underestimates the stage of liver disease and misses the diagnosis of cirrhosis in up to 25% of patients, especially in early and incomplete cirrhosis or macronodular cirrhosis^[5]. To overcome the limitations of safety and diagnostic accuracy in percutaneous liver biopsy, laparoscopic liver biopsy has been touted as a safe and effective diagnostic tool in patients with liver disease^[6,7]. The development of the so-called mini-laparoscopy by using a small diameter laparoscope allows for a minimally invasive procedure for macroscopic evaluation of the peritoneal cavity and safe biopsy of the liver^[8,9]. In this study, we retrospectively analyzed the safety of 1071 consecutive mini-laparoscopy guided liver biopsies, all performed in an endoscopy unit.

MATERIAL AND METHODS

Patients

The medical records of 1071 consecutive patients undergoing diagnostic mini-laparoscopy performed in our endoscopy unit at Johannes Gutenberg University Mainz, Germany from January 2000 until April 2006 were reviewed. All patients gave written informed consent before undergoing mini-laparoscopy. Patients with INR values above 1.5 and platelet counts below 50.000/ μ L received 2 to 4 units of fresh frozen plasma or 1 to 2 units of platelet concentrate immediately before the intervention respectively. Furthermore, every patient received an abdominal ultrasound prior to mini-laparoscopy to exclude ascites. All patients were treated as day cases with post interventional observation. Since this study represents a retrospective analysis, no IRB approval was required.

Technique of mini-laparoscopy

Mini-laparoscopy was performed using intravenous procedural sedation, as described previously^[6]. The abdominal wall was cleaned with Betadine solution and covered with sterile drapes. Puncture site of the abdominal wall was performed at the point of Kalk, located 2 cm left and cephalad of the umbilicus. The puncture itself was performed with a Veress needle with 2.3 mm diameter (Richard Wolf GmbH, Knittlingen, Germany) through a trocar of 2.75 mm diameter (Richard Wolf GmbH, Germany) after local anesthesia of the puncture site with 10 mL of mepivacaine 1% (AstraZeneca, London, UK). After forming a pneumoperitoneum by insufflation of approximately 2 liters nitric oxide, the Veress needle was replaced by the optical instrument (Richard Wolf GmbH, Germany). A xenon light source (Richard Wolf GmbH, Germany) was used for illumination of the abdominal

Table 1 Etiology of liver disease in non-cirrhotic patients

Viral hepatitis <i>n</i> = 319	Autoimmune liver disease <i>n</i> = 141	Liver disease of other origins <i>n</i> = 349
Chronic hepatitis C <i>n</i> = 263 (82%)	Autoimmune hepatitis <i>n</i> = 49 (35%)	Non-alcoholic fatty liver disease <i>n</i> = 99 (28%)
Chronic hepatitis B <i>n</i> = 56 (18%)	Primary biliary cirrhosis <i>n</i> = 48 (34%)	Alcoholic liver disease <i>n</i> = 18 (5%)
	Primary sclerosing cholangitis <i>n</i> = 32 (23%)	Primary/secondary hepatobiliary neoplasia <i>n</i> = 53 (15%)
	Overlap syndrome <i>n</i> = 13 (9%)	Toxic liver injury <i>n</i> = 59 (17%)
		Others <i>n</i> = 120 (34%)

Table 2 Etiology of liver disease in cirrhotic patients

Viral hepatitis <i>n</i> = 119	Autoimmune liver disease <i>n</i> = 20	Liver disease of other origins <i>n</i> = 111
Chronic hepatitis C <i>n</i> = 96 (85%)	Autoimmune hepatitis <i>n</i> = 8 (40%)	Alcoholic liver disease <i>n</i> = 52 (47%)
Chronic hepatitis B <i>n</i> = 23 (15%)	Primary biliary cirrhosis <i>n</i> = 6 (30%)	Others <i>n</i> = 59 (53%)
	Primary sclerosing cholangitis <i>n</i> = 6 (30%)	

cavity. For optimal visibility of the liver, a rotating operation table (Maquet GmbH, Rastatt, Germany) was used and patients were slightly rotated to the left while the upper body was elevated. Besides macroscopic assessment of the liver, the upper abdomen was systematically examined for signs of portal hypertension such as splenomegaly, dilated intra abdominal vessels or peritoneal carcinosis. Liver biopsy was performed under direct laparoscopic visualization with an 18G biopsy needle (Bard Inc., Covington, USA) *via* a second 3 mm incision in the upper right quadrant of the abdomen. If vigorous bleeding was detected immediately after the liver biopsy or the bleeding did not stop after approximately two minutes, a small trocar with a diameter of 3 mm was inserted at the liver biopsy puncture site of the abdominal wall and the bleeding puncture site was treated with argon plasma coagulation (APC) (Söring GmbH, Quickborn, Germany) under direct visualization. If necessary, in patients with high risk of bleeding, the second trocar was inserted in the right upper abdominal quadrant before performance of the liver biopsy in order to rapidly apply APC immediately after the biopsy. Coagulation was considered successful when no further signs of active bleeding could be observed after 2 min.

Statistical analysis

Statistical significance levels were calculated by chi square test.

RESULTS

1071 patients had mini-laparoscopy with the intent of

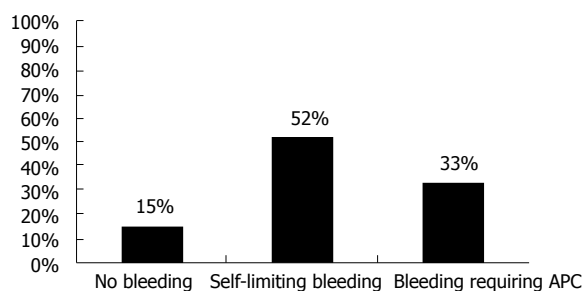


Figure 1 Overall occurrence of bleeding in mini-laparoscopic liver biopsies. APC: argon plasma coagulation.

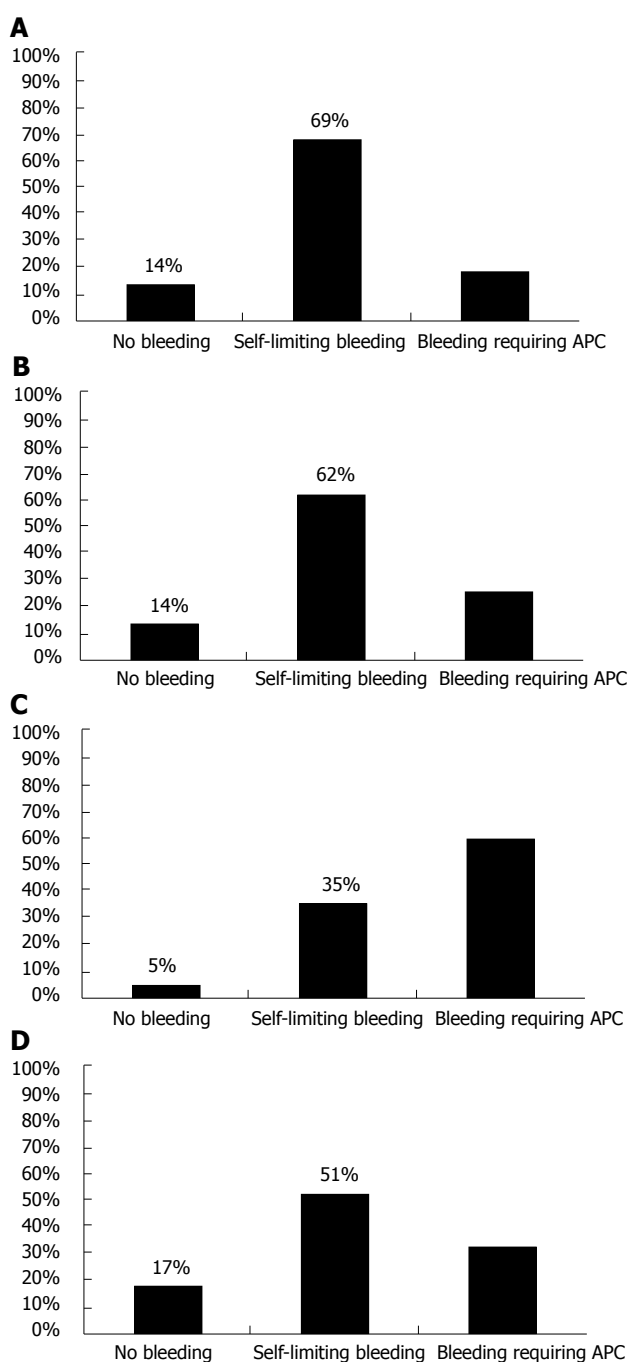


Figure 2 Occurrence of bleeding in mini-laparoscopic liver biopsies in regard to the underlying disease. A: Viral hepatitis; B: Autoimmune hepatitis; C: Cirrhosis; D: Other; APC: argon plasma coagulation.

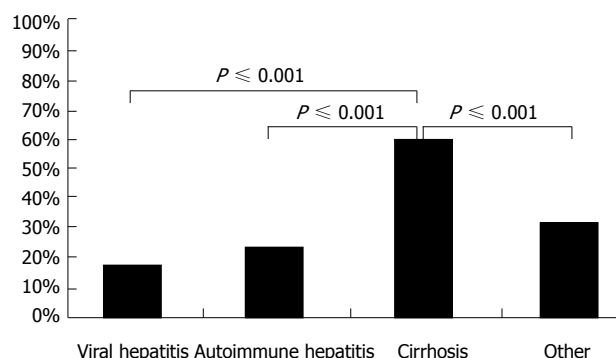


Figure 3 Occurrence of bleeding requiring argon plasma coagulation in regard to underlying disease.

liver biopsy. Indication for mini-laparoscopy and liver biopsy was for diagnosis of acute or chronic liver disease of unknown origin and for staging of cirrhosis. Patients with a history of abdominal surgery were not considered for mini-laparoscopy due to potential intra abdominal adhesions. Gender distribution was 55% males versus 45% females and the median age was 48 (± 14) years and 50 (± 14) years respectively. The etiology of the underlying liver disease is shown in Tables 1 and 2. Five hundred and fifty seven (52%) patients displayed minor self-limiting bleeding after the puncture of the liver and 161 (15%) showed no bleeding at all. In 353 cases (33%) of all mini-laparoscopic liver biopsies, prolonged bleeding occurred that required APC of the puncture site (Figure 1).

In order to determine whether the underlying liver injury influenced the risk and severity of bleeding, patients were stratified into four groups by type of liver disease: viral and autoimmune hepatitis, the most frequent disease entities; liver cirrhosis, the category with the highest expected risk of bleeding; and liver diseases of other origins. There were no significant differences in the severity of the bleeding between patients with viral and autoimmune hepatitis. Both groups displayed either no bleeding in 14% or self-limiting bleeding in 69% and 62% respectively. APC was used to treat bleeding in 17% of patients with viral hepatitis and in 24% of patients with autoimmune mediated liver disease (Figure 2A and B). In patients with liver cirrhosis, 5% of cases showed no signs of bleeding and 35% had only minor bleeding after liver biopsy (Figure 2C); APC was used in 60% of all cases. Patients with liver diseases of different origins than the ones mentioned above had no or mild bleeding in 17% and 51% respectively. APC was required in 32% of all cases (Figure 2D). Significant bleeding with the need for APC occurred with greater frequency in patients with liver cirrhosis compared to patients without advanced liver disease ($P \leq 0.001$, Figure 4).

DISCUSSION

Diagnostic laparoscopy is a valuable tool in the diagnosis of a variety of gastrointestinal illnesses. It is useful in the staging of upper GI-tract malignancies such as gastric

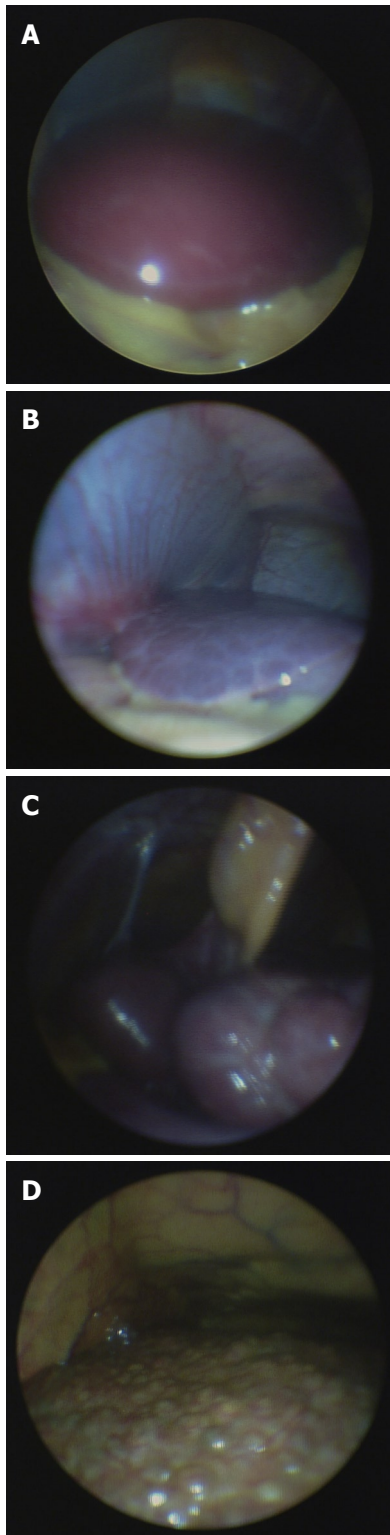


Figure 4 Laparoscopic images of different stages of liver disease. A: Healthy liver; B: Fibrotic liver; C: Macronodular cirrhosis; D: Micronodular cirrhosis.

and pancreatic cancer and sensitive for the diagnosis of peritoneal carcinosis^[10-13]. When used for the evaluation of liver disease, it allows for a macroscopic inspection of the liver and the ability to perform targeted biopsies of focal lesions on the liver surface. Splenic biopsies can also be performed *via* a diagnostic laparoscopy^[14]. For these diagnostic non-surgical procedures, the minimally invasive technique of mini-laparoscopy which requires smaller insertions to the abdominal wall than conventional laparoscopy has been shown to be useful^[15].

In our study, we evaluated the occurrence and immediate management of bleeding after liver biopsy during mini-laparoscopy performed in an endoscopy unit. The primary reason for choosing mini-laparoscopy over percutaneous biopsy was to have the ability to treat potential bleeding complications and perform macroscopic evaluation of the liver since the severity of liver injury can be underestimated if based only on the histological result of the biopsy material^[16]. Besides the additional macroscopic information, mini-laparoscopic liver biopsy enables the histological assessment of liver injury in patients with advanced cirrhosis where percutaneous biopsy would be contraindicated due to the high risk of a bleeding complication^[17]. In this study, we demonstrate that bleeding from liver biopsy occurred significantly more frequently in patients with cirrhosis than non-cirrhotic patients, resulting in an increased need for APC of the biopsy site. Presumably as a result of APC, there were no major post-interventional bleeding complications in any patients. Liver biopsy was performed safely even in patients with decompensated Child C cirrhosis with portal hypertension and marked coagulopathy after administration of fresh frozen plasma and/or platelets.

Some recently published studies describe a lower risk than that of previous reports, indicating that the safety of this procedure has improved^[18,19].

These findings confirm previously published data^[20,21] where mini-laparoscopy showed similar safety data compared to percutaneous biopsy but mini-laparoscopy was more sensitive in assessing the severity of the liver disease.

In conclusion, mini-laparoscopy guided liver biopsy performed in an endoscopy unit is a safe and effective technique for the evaluation of patients with liver diseases. It enables the macroscopic assessment of the liver and the ability to perform liver biopsy in patients with high risk of bleeding, allowing for the management of complications and enabling histological diagnosis in patients with advanced liver disease or cirrhosis.

COMMENTS

Background

Histological assessment of liver injury still represents the most important diagnostic tool in the evaluation of liver diseases. However, percutaneous liver biopsy may be associated with bleeding complications in patients with advanced liver disease. Thus, a safe method for obtaining liver histology in this high risk patient population is crucial.

Research frontiers

Due to non-invasive assessment of liver fibrosis with fibroscan® technology or radiographic imaging, liver histology may not be necessary in some patients. However, to date, liver biopsy seems crucial in most cases. Whether non-invasive techniques will replace liver biopsy in the future remains to be determined by further research.

Innovations and breakthroughs

The authors demonstrated that the assessment of liver histology is possible even in critically ill patients with compromised liver function.

Applications

Mini-laparoscopy, not only a valuable tool for liver biopsy in patients with advanced liver disease, can also be used for tumor staging in patients with intra abdominal malignancies. Especially, peritoneal carcinosis can be detected in an early stage before visualization with radiological techniques is possible.

Peer reviews

Overall, this is an interesting and well written retrospective study about mini-laparoscopy and liver biopsy in a large series of 1071 consecutive patients. A prospective randomized control study of mini-laparoscopy versus percutaneous biopsy was recommended. However, patients with advanced liver disease or compromised liver function or blood coagulation should not be included in such a study as it is known that these patients have a higher risk of bleeding complications after percutaneous biopsy.

REFERENCES

- 1 **Wong GL**, Wong VW, Choi PC, Chan AW, Chum RH, Chan HK, Lau KK, Chim AM, Yiu KK, Chan FK, Sung JJ, Chan HL. Assessment of fibrosis by transient elastography compared with liver biopsy and morphometry in chronic liver diseases. *Clin Gastroenterol Hepatol* 2008; **6**: 1027-1035
- 2 **Desmet VJ**, Gerber M, Hoofnagle JH, Manns M, Scheuer PJ. Classification of chronic hepatitis: diagnosis, grading and staging. *Hepatology* 1994; **19**: 1513-1520
- 3 **Grant A**, Neuberger J. Guidelines on the use of liver biopsy in clinical practice. British Society of Gastroenterology. *Gut* 1999; **45** Suppl 4: IV1-IV11
- 4 **Buckley A**, Petrunia D. Practice guidelines for liver biopsy. Canadian Association of Gastroenterology. *Can J Gastroenterol* 2000; **14**: 481-482
- 5 **Helmreich-Becker I**, Schirmacher P, Denzer U, Hensel A, Meyer zum Büschenfelde KH, Lohse AW. Minilaparoscopy in the diagnosis of cirrhosis: superiority in patients with Child-Pugh A and macronodular disease. *Endoscopy* 2003; **35**: 55-60
- 6 **Helmreich-Becker I**, Meyer zum Büschenfelde KH, Lohse AW. Safety and feasibility of a new minimally invasive diagnostic laparoscopy technique. *Endoscopy* 1998; **30**: 756-762
- 7 **Poniachik J**, Bernstein DE, Reddy KR, Jeffers LJ, Coelho-Little ME, Civantos F, Schiff ER. The role of laparoscopy in the diagnosis of cirrhosis. *Gastrointest Endosc* 1996; **43**: 568-571
- 8 **Hulscher JB**, Nieveen van Dijkum EJ, de Wit LT, van Delden OM, van Lanschot JJ, Obertop H, Gouma DJ. Laparoscopy and laparoscopic ultrasonography in staging carcinoma of the gastric cardia. *Eur J Surg* 2000; **166**: 862-865
- 9 **Reddy KR**, Levi J, Livingstone A, Jeffers L, Molina E, Kligerman S, Bernstein D, Kodali VP, Schiff ER. Experience with staging laparoscopy in pancreatic malignancy. *Gastrointest Endosc* 1999; **49**: 498-503
- 10 **Conlon KC**. Staging laparoscopy for gastric cancer. *Ann Ital Chir* 2001; **72**: 33-37
- 11 **Arnold JC**, Schneider AR, Zöpf T, Neubauer HJ, Jakobs R, Benz C, Riemann JF. [Laparoscopic tumor staging in gastrointestinal carcinomas: significance of internal medicine laparoscopy]. *Z Gastroenterol* 2001; **39**: 19-23
- 12 **Barreiro CJ**, Lillemoe KD, Koniaris LG, Sohn TA, Yeo CJ, Coleman J, Fishman EK, Cameron JL. Diagnostic laparoscopy for periampullary and pancreatic cancer: what is the true benefit? *J Gastrointest Surg* 2002; **6**: 75-81
- 13 **Denzer U**, Hoffmann S, Helmreich-Becker I, Kauczor HU, Thelen M, Kanzler S, Galle PR, Lohse AW. Minilaparoscopy in the diagnosis of peritoneal tumor spread: prospective controlled comparison with computed tomography. *Surg Endosc* 2004; **18**: 1067-1070
- 14 **Denzer U**, Helmreich-Becker I, Galle PR, Lohse AW. Minilaparoscopy-guided spleen biopsy in systemic disease with splenomegaly of unknown origin. *Endoscopy* 2002; **34**: 495-498
- 15 **Weickert U**, Jakobs R, Siegel E, Eickhoff A, Schilling D, Riemann JF. [Complications of diagnostic laparoscopy]. *Dtsch Med Wochenschr* 2005; **130**: 16-20
- 16 **Gebo KA**, Herlong HF, Torbenson MS, Jenckes MW, Chandler G, Ghanem KG, El-Kamary SS, Sulkowski M, Bass EB. Role of liver biopsy in management of chronic hepatitis C: a systematic review. *Hepatology* 2002; **36**: S161-S172
- 17 **Denzer U**, Helmreich-Becker I, Galle PR, Lohse AW. Liver assessment and biopsy in patients with marked coagulopathy: value of mini-laparoscopy and control of bleeding. *Am J Gastroenterol* 2003; **98**: 893-900
- 18 **Seeff LB**, Everson GT, Morgan TR, Curto TM, Lee WM, Ghany MG, Shiffman ML, Fontana RJ, Di Bisceglie AM, Bonkovsky HL, Dienstag JL. Complication rate of percutaneous liver biopsies among persons with advanced chronic liver disease in the HALT-C trial. *Clin Gastroenterol Hepatol* 2010; **8**: 877-883
- 19 **West J**, Card TR. Reduced mortality rates following elective percutaneous liver biopsies. *Gastroenterology* 2010; **139**: 1230-1237
- 20 **Inabnet WB**, Deziel DJ. Laparoscopic liver biopsy in patients with coagulopathy, portal hypertension, and ascites. *Am Surg* 1995; **61**: 603-606
- 21 **Denzer U**, Arnoldy A, Kanzler S, Galle PR, Dienes HP, Lohse AW. Prospective randomized comparison of minilaparoscopy and percutaneous liver biopsy: diagnosis of cirrhosis and complications. *J Clin Gastroenterol* 2007; **41**: 103-110

S- Editor Zhang HN L- Editor Roemmele A E- Editor Liu N

Ultrastim endoscopy with flexible spectral imaging color enhancement for upper gastrointestinal neoplasms

Yukari Tanioka, Hideo Yanai, Eiki Sakaguchi

Yukari Tanioka, Hideo Yanai, Eiki Sakaguchi, Department of Gastroenterology and Hepatology, Department of Clinical Research, National Hospital Organization Kanmon Medical Center, Shimonoseki, Yamaguchi 752-8510, Japan

Author contributions: Tanioka Y performed esophago-gastro-duodenoscopy and prepared the manuscript; Yanai H made the study plan, performed ultrastim endoscopy and endoscopic sub-mucosal dissection; and Sakaguchi E analyzed the results.

Correspondence to: Hideo Yanai, MD, PhD, Department of Clinical Research, National Hospital Organization Kanmon Medical Center, 1-1 Sotoura, Chofu, Shimonoseki, Yamaguchi 752-8510, Japan. yanaih@simonoseki2.hosp.go.jp

Telephone: +81-83-2411199 Fax: +81-83-2411301

Received: August 17, 2010 Revised: December 12, 2010

Accepted: December 19, 2010

Published online: January 16, 2011

Abstract

AIM: To conduct a preliminary study on the effect of flexible spectral imaging color enhancement (FICE) used in combination with ultrastim endoscopy by focusing on the enhanced contrast between tumor and non-tumor lesions.

METHODS: We examined 50 lesions of 40 patients with epithelial tumors of the upper gastrointestinal tract before endoscopic submucosal dissection using ultrastim endoscopy with conventional natural color imaging and with FICE imaging. We retrospectively investigated the effect of the use of FICE on endoscopic diagnosis in comparison with normal light.

RESULTS: Visibility of the epithelial tumors of the upper gastrointestinal tract with FICE was superior to normal light in 54% of the observations and comparable to normal light in 46% of the observations. There was no lesion for which visibility with FICE was inferior to that with normal light. FICE visualized 69.6% of hyperemic lesions and 58.8% of discolored lesions better than conventional endoscopy with natural color imaging. FICE

significantly improved the visibility of lesions with hyperemia or discoloration compared with normocolored lesions.

CONCLUSION: This study suggests that the use of FICE would improve the ability of ultrastim endoscopy to detect epithelial tumors of the upper gastrointestinal tract.

© 2011 Baishideng. All rights reserved.

Key words: Ultrastim endoscopy; Upper gastrointestinal neoplasms; Flexible spectral imaging color enhancement

Peer reviewer: Hongchun Bao, PhD, Research Fellow, The Center for Micro-Photonics, Faculty of Engineering & Industrial Sciences, Swinburne University of Technology, PO Box 218, Hawthorn, Victoria 3122, Australia

Tanioka Y, Yanai H, Sakaguchi E. Ultrastim endoscopy with flexible spectral imaging color enhancement for upper gastrointestinal neoplasms. *World J Gastrointest Endosc* 2011; 3(1): 11-15 Available from: URL: <http://www.wjgnet.com/1948-5190/full/v3/i1/11.htm> DOI: <http://dx.doi.org/10.4253/wjge.v3.i1.11>

INTRODUCTION

Recent practical application of ultrastim endoscopy has led to the rapid spread of transnasal endoscopy as a screening method that causes less pain than peroral endoscopy for the patient during esophago-gastro-duodenoscopy (EGD)^[1,2]. Garcia *et al*^[3] reported that ultrastim transnasal endoscopy required a significantly shorter recovery time with significantly lower costs for recovery rooms, personnel expenses, intravenous access devices and oxygen monitors compared to conventional peroral endoscopy with a sedated patient. In addition, transnasal ultrastim endoscopy is appropriate for patients with trismus, those who cannot accept insertion for peroral

endoscopy without sedation, and patients with stenosis of the upper gastrointestinal tract in the pharynx or lower regions^[4].

However, there are some disadvantages caused by the downsizing of scopes. Compared with conventional endoscopy, ultraslim endoscopy provides less resolution due to the smaller number of pixels and less illumination. Lower power for supplying air and water necessitates a longer time for yielding good images and the smaller diameters of forceps channels limit the usable instruments, resulting in smaller specimens and less information from tissues from a biopsy. For these reasons, ultraslim endoscopy is mainly used for screening purposes rather than detailed close examination. Thus, the ability of ultraslim endoscopy for detection and early diagnosis of gastric cancer should be studied in detail.

Flexible spectral imaging color enhancement (FICE) is one of the diagnostic methods using specific light spectra based on spectral image processing technology (Fujinon Corporation, Saitama, Japan). Currently, it is mainly employed for detailed diagnostic workups for superficial lesions in the gastrointestinal tract using high-resolution magnifying endoscopy and is gradually spreading. Visible light consists of wavelengths from red to purple. A spectral image, the image captured by each wavelength, of a specific wavelength is electrically amplified and reconstructed to create a FICE image. FICE provides comparison of spectral images of diseased and surrounding normal areas for enhancement of the contrast by combining wavelengths with greater differences in signals^[5,6].

In the present study, we focused on the enhanced color contrast between tumor and non-tumor areas in FICE images and conducted a preliminary retrospective study on the effect of FICE used in combination with ultraslim endoscopy for observation of superficial epithelial tumors of the upper gastrointestinal tract.

MATERIAL AND METHODS

Our institution introduced the ultraslim endoscopy system for EGD in April 2007 and had examined 380 patients, mainly for screening, by February 2008. During this period, we examined 53 lesions in 42 patients with epithelial tumors of the upper gastrointestinal tract who underwent endoscopic submucosal dissection (ESD). The examination was conducted using ultraslim endoscopy with conventional natural color imaging and with FICE imaging before ESD, after obtaining the consent of the patient. Based on the observation of these 53 lesions, we retrospectively investigated the effect of the use of FICE on the visibility of upper gastrointestinal tumorous lesions. The lesions consisted of 3 superficial carcinomas of the esophagus, 3 gastric non neoplastic polyps, 19 gastric adenomas and 28 early gastric cancers.

We used the EG-530N2 (tip diameter 5.9 mm, forceps channel diameter 2.0 mm, four-way bending; Fujinon Corporation, Saitama, Japan) for ultraslim endoscopy of the upper gastrointestinal tract. For FICE imaging, we

Table 1 Five-point scale for evaluation of observation with flexible spectral imaging color enhancement

Point	Evaluation of observation with FICE
1	FICE fails to visualize lesions detectable with conventional images
2	FICE is slightly inferior to conventional images
3	FICE is comparable to conventional images
4	FICE is superior to conventional images
5	FICE allows detection of lesions not easily detected with conventional images or clearer visualization of areas poorly defined with conventional images

FICE: flexible spectral imaging color enhancement.

Table 2 Results of evaluation of visibility during observation with flexible spectral imaging color enhancement

Evaluation of FICE	1	2	3	4	5	Subtotal
Superficial carcinoma of the esophagus	0	0	2	1	0	3
Gastric non neoplastic polyp	0	0	3	0	0	3
Gastric adenoma	0	0	8	11	0	19
Early gastric cancer	0	0	11	17	0	28
	0	0	24	29	0	53

FICE: flexible spectral imaging color enhancement.

selected a wavelength set of 525 nm (4), 495 nm (5) and 495 nm (4) for R, G and B respectively, which provides optimal illumination and highlights hyperemic changes commonly seen in epithelial tumors.

We used a five-point scale (Table 1) to evaluate the visibility of lesions which is related to the ability of screening to detect lesions, based on comparison of conventional images and FICE images mainly in long- and middle-distance views.

In addition, the lesions were classified as discolored, hyperemic or normocolored (no color differences from surrounding mucosa) for comparison of visibility in the observation with FICE. Fisher's exact probability test was used for statistical analysis.

RESULTS

No lesions were graded 1 or 2, inferior to conventional images, for visibility with FICE. In 24 of the 53 lesions (45.3%), visibility with FICE was graded 3, comparable to conventional images. In 29 of the 53 lesions (54.7%), visibility with FICE was graded 4, superior to conventional images (Table 2, Figures 1-3).

Regarding color hue changes in the lesions, 17 of 25 lesions (68.0%) with hyperemic changes were better visualized with FICE than conventional images (FICE 4, Figures 1-3). Eleven of 18 discolored lesions (61.1%) were also better visualized with FICE. However, nine of ten normocolored lesions (90%) detected with conventional images were similarly visualized with FICE (Table 3). In short, FICE significantly improved the visibility of lesions with hyperemia ($P = 0.0027$) or discoloration ($P = 0.0159$) compared with normocolored lesions.

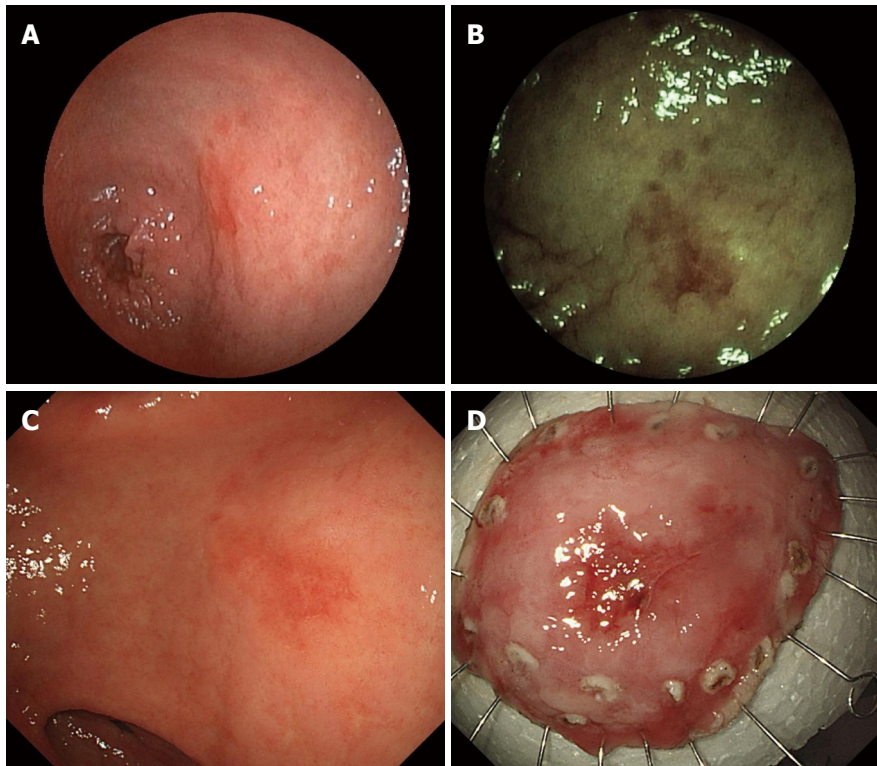


Figure 1 A slightly depressed lesion with hyperemia was found on the surface of the posterior wall of the lower body of the stomach. The hyperemia of the lesion is highlighted and the boundary is more clearly visualized with flexible spectral imaging color enhancement (FICE). The visibility with FICE was graded 4. Endoscopic submucosal dissection (ESD) was performed for local complete resection of the moderately differentiated adenocarcinoma limited to within the mucosa. A: Conventional image with ultrastim endoscopy; B: Image with FICE: The hyperemia of the lesion is highlighted and the boundary is more clearly visualized with FICE (FICE 4); C: Conventional image with conventional esophagogastroduodenoscopy; D: Dissected section after ESD.

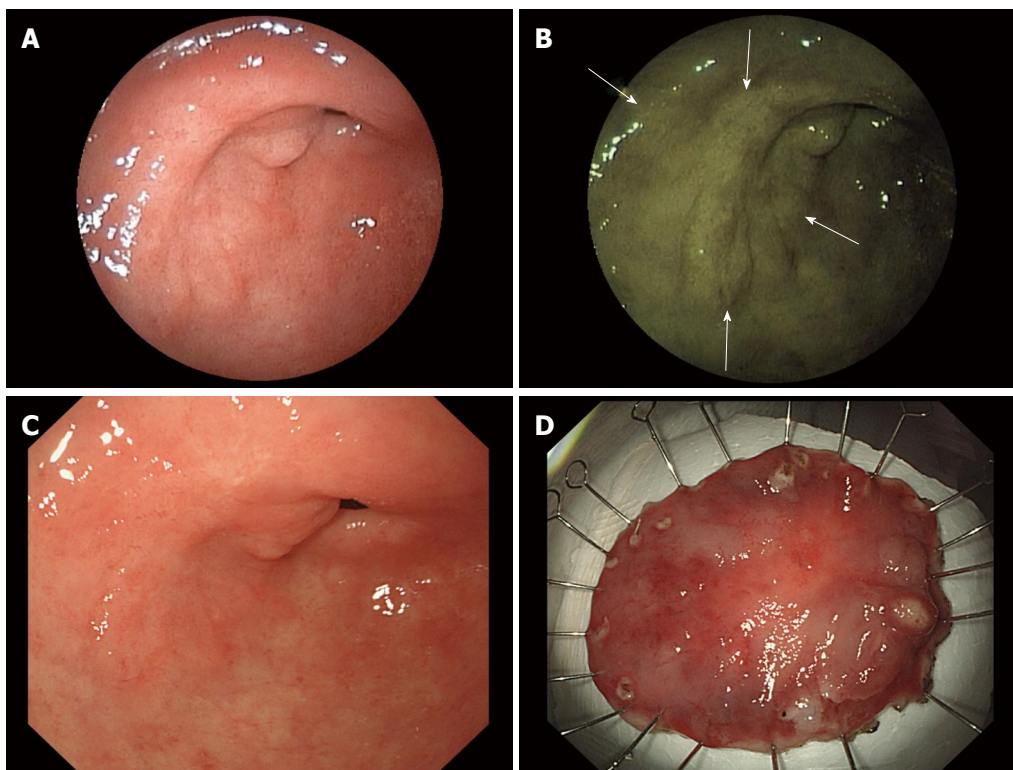


Figure 2 A discolored change was found in the anterior wall and surrounding areas of the gastric antrum. Color contrast between the lesion and surrounding mucosa with atrophic changes is highlighted and the boundary is more clearly visualized with flexible spectral imaging color enhancement (FICE). The visibility with FICE was graded 4. Endoscopic submucosal dissection (ESD) was performed for local complete resection of the well-differentiated adenocarcinoma limited to within the mucosa. A: Conventional image with ultrastim endoscopy; B: Image with FICE: Color contrast between the lesion and surrounding mucosa is highlighted and the boundary is more clearly visualized (FICE 4); C: Conventional image with conventional esophagogastroduodenoscopy; D: Dissected section after ESD.

DISCUSSION

To aid examination and diagnosis using conventional endoscopy, conventional chromoendoscopy with the spraying of a dye such as indigo carmine highlights the surface irregularities of lesions and is a common and very useful method for defining lesions. However, there are problems: additional costs for spraying the dye, the time

involved and the inability to highlight the capillary patterns which is important for the early diagnosis of cancer^[7].

Virtual chromoendoscopy systems were developed to correct these problems^[5]. FICE uses spectral estimation technology to pick up different given wavelengths from all wavelength components from the CCD and produces images using arithmetical processing. FICE provides real-time switching at the flick of a switch. In addition,

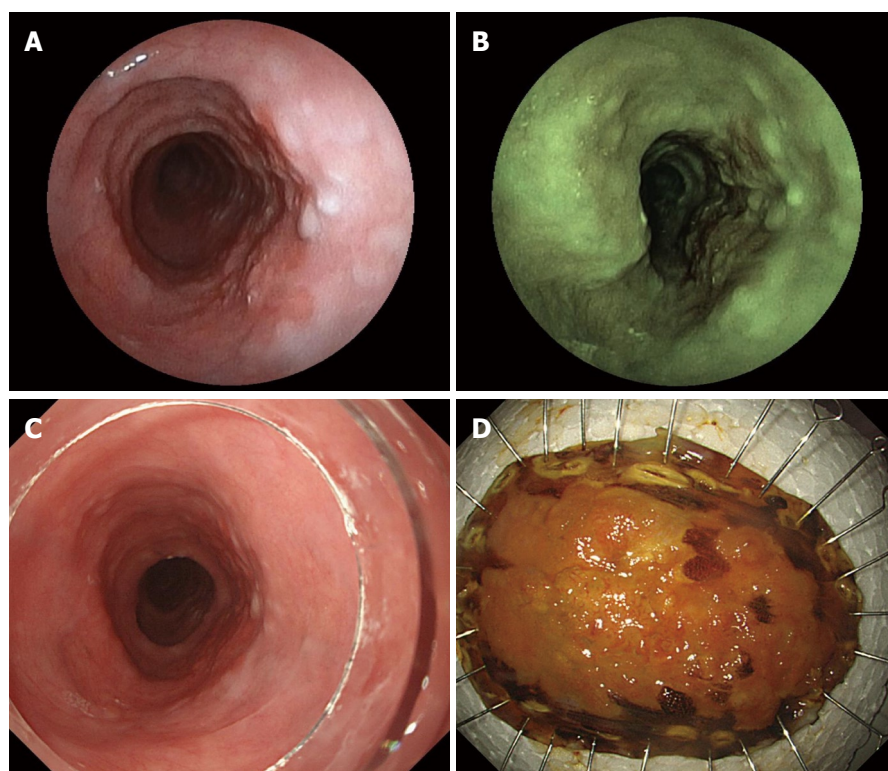


Figure 3 A rough, slightly elevated squamous lesion with hyperemia in a semicircle is found 25 cm distal from the superior incisor line. The hyperemia of the lesion is highlighted and the boundary is more clearly visualized with flexible spectral imaging color enhancement (FICE). The visibility was graded 4. Endoscopic submucosal dissection (ESD) was performed for local complete resection of the squamous cell carcinoma including the muscularis mucosae. A: Conventional image with ultraslim endoscopy: A rough, elevated squamous lesion with hyperemia in a semicircle is found; B: Image with FICE: The hyperemia of the lesion is highlighted and the boundary is more clearly visualized with FICE (FICE 4); C: Conventional image with conventional esophagogastroduodenoscopy; D: Dissected section sprayed with iodine after ESD.

Table 3 Evaluation of color hue changes and visibility of lesions with flexible spectral imaging color enhancement observation

Evaluation of FICE	1	2	3	4	5	Subtotal
Hyperemic ^b	0	0	8	17	0	25
Discolored ^a	0	0	7	11	0	18
Normocolored	0	0	9	1	0	10
	0	0	24	29	0	53

^a*P* = 0.0159 *vs* Normocolored; ^b*P* = 0.0027 *vs* Normocolored; FICE: flexible spectral imaging color enhancement.

as FICE electrically amplifies spectral images of given wavelengths and reconstructs the images, it offers brighter images with subtle highlighted color changes and hyperemic areas on the surface of the mucosa^[8]. Therefore, we obtained sufficient illumination for long-distance observation with FICE in all cases in this study.

FICE observation of superficial epithelial tumors of the gastrointestinal tract resulted in superior visibility for 54.7% and comparable visibility for 45.3% compared with conventional images. These results suggested that FICE would be useful as a diagnostic aid for ultraslim endoscopy which is becoming more common. FICE significantly improved the visibility of lesions with hyperemia (68.0%) and discoloration (61.1%) compared with conventional images. The results showed that, due to the characteristics of FICE, observation by ultraslim endoscopy with FICE more clearly visualized the color contrast between diseased and normal lesions than observation with conventional images. FICE was also expected to improve the visibility of lesions with color contrast in the mucosa in

long-distance observation by lower-resolution ultraslim endoscopy.

Yoshida reported that a review of endoscopic observation of gastritis-like early gastric cancer diagnosed as benign by conventional endoscopy revealed discolored lesions in 29 of 132 cases (22.0%), hyperemic lesions in 83 cases (62.9%) and normocolored lesions in 20 cases (15.2%)^[7]. These lesions, not easily diagnosed by conventional endoscopy, may be overlooked in observations with conventional ultraslim endoscopy. Of these lesions, discolored and hyperemic lesions accounted for 84.9%. FICE was considered to prevent overlooking lesions with such color changes.

This study suggests that the use of FICE would improve the ability of ultraslim endoscopy to detect epithelial tumors of the upper gastrointestinal tract under conditions without the spraying of a dye or a biopsy.

ACKNOWLEDGEMENTS

This study was conducted in collaboration with Fujinon Corporation (Saitama, Japan).

COMMENTS

Background

The reduction in the endoscope diameter would improve the acceptance of unsedated endoscopy. However, ultraslim endoscopy provides less resolution due to smaller number of pixels and less illumination. Flexible spectral imaging color enhancement (FICE) is one of the diagnostic methods using specific light spectra based on spectral image processing technology (Fujinon Corporation, Saitama, Japan). We focused on the effect of FICE used in combination with ultraslim endoscopy for observation of superficial epithelial tumors of the upper gastrointestinal tract.

Research frontiers

The analysis and enhancement of diagnostic accuracy of ultrathin endoscopy is a new frontier for upper gastrointestinal cancer surveillance.

Innovations and breakthroughs

FICE provides comparison of spectral images of diseased and surrounding normal areas for enhancement of the contrast by combining wavelengths with greater differences in signals.

Applications

Our study suggests that the use of FICE would improve the ability of ultrathin endoscopy to detect epithelial tumors of the upper gastrointestinal tract under conditions without the spraying of a dye or a biopsy.

Terminology

Ultrathin endoscopes: a shaft diameter of 6 mm or less which allows them to be passed through the nose or mouth. Flexible spectral Imaging Color Enhancement (FICE): Visible light consists of wavelengths from red to purple. A spectral image, the image captured by each wavelength, of a specific wavelength is electrically amplified and reconstructed to create a FICE image.

Peer review

Overall, the manuscript is good. The authors show flexible spectral imaging color enhancement could improve ultrathin endoscopy in detection of epithelial tumors of the upper gastrointestinal tract. Therefore I could recommend publication if several points were discussed.

REFERENCES

- 1 Birkner B, Fritz N, Schatke W, Hasford J. A prospective randomized comparison of unsedated ultrathin versus standard esophagogastroduodenoscopy in routine outpatient gastroenterology practice: does it work better through the nose? *Endoscopy* 2003; **35**: 647-651
- 2 Preiss C, Charton JP, Schumacher B, Neuhaus H. A randomized trial of unsedated transnasal small-caliber esophagogastroduodenoscopy (EGD) versus peroral small-caliber EGD versus conventional EGD. *Endoscopy* 2003; **35**: 641-645
- 3 Garcia RT, Cello JP, Nguyen MH, Rogers SJ, Rodas A, Trinh HN. Unsedated ultrathin EGD is well accepted compared with conventional sedated EGD: a multicenter randomized trial. *Gastroenterology* 2003; **125**: 1606-1612
- 4 Abe K, Miyaoka M. Trial of transnasal esophagogastroduodenoscopy. *Digestive Endoscopy* 2006; **18**: 212-217
- 5 Pohl J, May A, Rabenstein T, Pech O, Ell C. Computed virtual chromoendoscopy: a new tool for enhancing tissue surface structures. *Endoscopy* 2007; **39**: 80-83
- 6 Osawa H, Yoshizawa M, Yamamoto H, Kita H, Satoh K, Ohnishi H. Optimal band imaging system can facilitate detection of changes in depressed-type early gastric cancer. *Gastrointest Endosc* 2008; **67**: 226-234
- 7 Yoshida S. Endoscopy, Gastric Cancer. In: Sugimura T, Sasako M, editors. Oxford University Press: Tokyo, 1997: 168-188
- 8 Mouri R, Yoshida S, Tanaka S, Oka S, Yoshihara M, Chayama K. Evaluation and validation of computed virtual chromoendoscopy in early gastric cancer. *Gastrointestinal Endoscopy* 2009; **69**: 1052-1058

S- Editor Zhang HN L- Editor Roemmele A E- Editor Liu N

Duodenal tuberculosis presenting as gastric outlet obstruction: A case report

Haydee Buluran Flores, Felix Zano, Ena Lyn Ang, Norberto Estanislao

Haydee Buluran Flores, Felix Zano, Ena Lyn Ang, Norberto Estanislao, Philippine General Hospital, Manila City 1000, Philippines

Author contributions: Flores HB wrote the manuscript and contributed to the literature search; Zano F supervised the manuscript writing; and Ang EL and Estanislao N contributed to the literature search and editing the manuscript.

Correspondence to: Haydee Buluran Flores, MD, Gastrointestinal Clinic, Philippine General Hospital, Taft Avenue, Manila City 1000, Philippines. wishbonemydog@yahoo.com

Telephone: +63-2-9228604530 Fax: +63-2-5672983

Received: February 9, 2010 Revised: December 13, 2010

Accepted: December 20, 2010

Published online: January 16, 2011

Peer reviewers: Noriya Uedo, Director, Endoscopic Training and Learning Center; Vice Director, Department of Gastrointestinal Oncology, Osaka Medical Center for Cancer and Cardiovascular Diseases, Osaka, Japan; David Friedel, MD, Gastroenterology, Winthrop University Hospital, 222 Station Plaza North, Suite 428, Mineola NY 11501, United States

Flores HB, Zano F, Ang EL, Estanislao N. Duodenal tuberculosis presenting as gastric outlet obstruction: A case report. *World J Gastrointest Endosc* 2011; 3(1): 16-19 Available from: URL: <http://www.wjgnet.com/1948-5190/full/v3/i1/16.htm> DOI: <http://dx.doi.org/10.4253/wjge.v3.i1.16>

Abstract

Gastric outlet obstruction is commonly associated with malignancies and peptic ulcer disease. However, when no malignancy is seen and the patient is non-responsive to conventional peptic ulcer treatment, other etiologies need to be explored. We report a case of gastric outlet obstruction due to duodenal tuberculosis. The patient is a 31 year old male who presented with 1 year history of recurrent epigastric pain and an acute episode of vomiting. Endoscopy revealed duodenal stricture. Computed tomography scan showed pyloric antral thickening. The patient was referred to the surgery service and underwent an exploratory laparotomy and gastrojejunostomy. A duodenal mass and calcified lymph nodes were noted on exploration and biopsy revealed a tuberculous origin. The patient was started on anti-tuberculosis medications and had improved on discharge. Gastroduodenal tuberculosis is rare and pyloric stenosis resulting from tuberculosis is even rarer. This, however, should be considered in patients who come from areas where the disease is endemic.

2011 Baishideng. All rights reserved.

Key words: Duodenal tuberculosis; Tuberculosis; Gastric outlet obstruction; Duodenal stricture

INTRODUCTION

Tuberculosis is a major health problem worldwide. In the Philippines, it is endemic. The most common manifestation is a pulmonary disease but involvement of the gastrointestinal tract is not uncommon. Gastrointestinal tuberculosis often involves the ileocecal region. The stomach as well as the duodenum are rare sites for tuberculosis and are usually a result of secondary spread from a primary pulmonary disease. An autopsy series reported an incidence of only 0.5%^[1]. A primary case of gastroduodenal tuberculosis is an even rarer disease and only a few cases are reported in the literature^[1-5].

CASE REPORT

A 31 year old male was admitted to our institution with a 1 year history of recurrent epigastric pain and an acute episode of vomiting. Epigastric pain was characterized as intermittent, mild and gnawing in character. The patient also reported a slight undocumented weight loss. No other associated signs and symptoms were noted. No medications had been taken and he had not consulted medical staff. The patient had an acute episode of vomiting, prompting a consultation and subsequent admission.

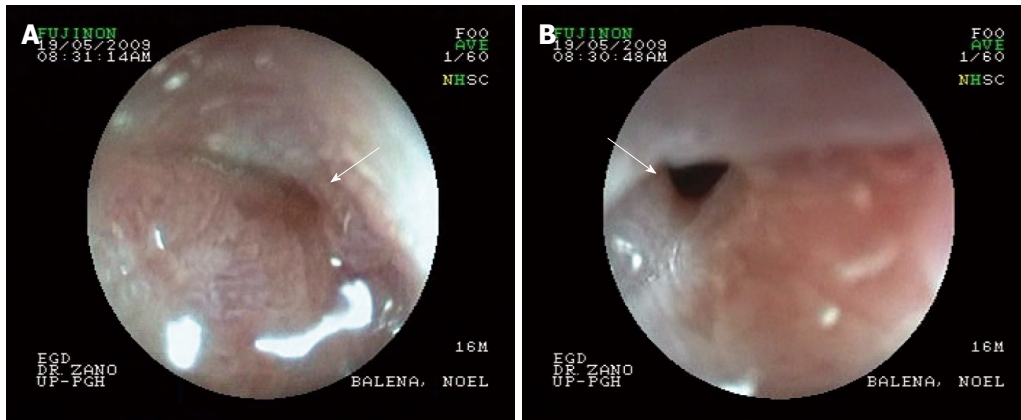


Figure 1 Endoscopic view of the 2nd part of the duodenum of the patient showing narrowing of lumen/stricture (white arrow). The gastroscop was unable to pass beyond this point.

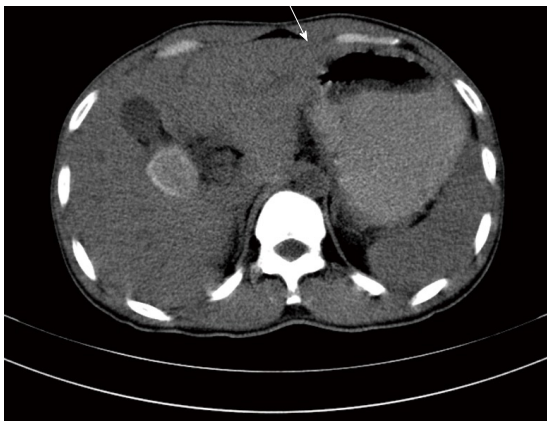


Figure 2 Abdominal computed tomography scan showing pyloroantral thickening and distended stomach.

The patient had no known co-morbidities and no history of hospitalization. He had no history of tuberculosis and no known exposure to the disease. The patient's family history was also unremarkable.

Physical examination only revealed direct tenderness in the epigastrium. There was no note of lymphadenopathies and the rest of the physical exam was unremarkable. Body mass index was 21. Complete blood count was within normal limits. Chest x-ray was normal. Esophago gastroduodenoscopy revealed duodenal stricture at the 1st part of the duodenum (Figure 1). Computed tomography of the whole abdomen showed pyloroantral thickening and a distended stomach (Figure 2). The impression then was gastric outlet obstruction secondary to duodenal stricture, probably secondary to peptic ulcer cicatrization to rule out duodenal malignancy.

The patient was referred to the surgical service. He underwent exploratory laparotomy and gastrojejunostomy. A duodenal mass and calcified lymph nodes were noted intraoperatively. Excision of the mass was done as well as biopsy of the calcified lymph nodes. Post-operative recovery was uneventful. On histopathological examination, a chronic granulomatous inflammation with Langhans-

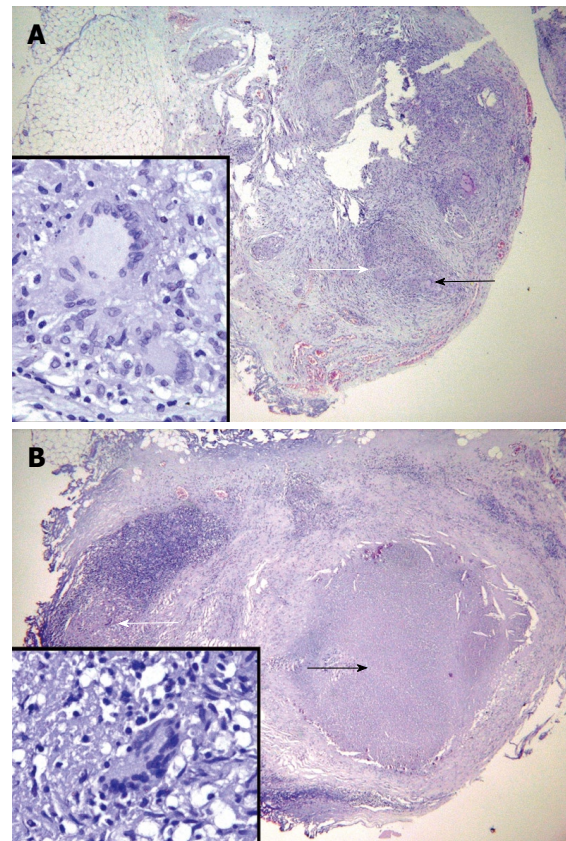


Figure 3 Histopathological examination image. A: Duodenal mass showing chronic granulomatous inflammation (white arrow) with Langhans-type giant cell (black arrow); B: Calcified lymph node showing caseation necrosis (black arrow) and a multinucleated giant cell (inset, white arrow).

type giant cell in the duodenal wall and the calcified lymph node showed caseation necrosis and multi-loculated giant cells were noted (Figure 3A and B). The patient was then diagnosed as having duodenal stricture secondary to primary duodenal tuberculosis.

The patient was started on quadruple anti-tuberculous medication and had improved on discharge. The patient was symptom free 3 mo later.

DISCUSSION

Gastric outlet obstruction is commonly associated with malignancies and peptic ulcer disease. However, when no malignancy is seen and a patient is non-responsive to conventional peptic ulcer treatment, other etiologies need to be explored. In this case, our patient presented with gastric outlet obstruction and a diagnosis of ulcer cicatrization was first made since it is one of the more common causes of duodenal strictures. Also, his young age made the diagnosis of malignancy less likely. Since the patient already presented with gastric outlet obstruction, surgery was deemed necessary for the patient and the diagnosis of gastroduodenal tuberculosis was made post-operatively. In other reported cases of gastroduodenal tuberculosis, medical management was a sufficient treatment modality^[2]. However, for those presenting with obstruction, surgery may be necessary to relieve the obstruction and make the diagnosis.

The diagnosis of this disease is difficult and is often made post-operatively. There are no pathognomonic clinical features. A review of 23 consecutive cases of gastroduodenal tuberculosis (15 year span) in India noted that vomiting (60.8%) and epigastric pain (56.5%) are the most common presenting symptoms. Other symptoms noted are weight loss, upper GI bleeding and fever. The mean age at presentation in this series was 34.4 years with the duration of symptoms varying from 2 d to 15 years^[1]. Most reported cases of duodenal tuberculosis come from areas with high prevalence of tuberculosis such as India and Africa. Hence, a high index of suspicion is needed when a patient is from a place endemic for tuberculosis. Another emerging concern is the increasing prevalence of Human Immunodeficiency Virus (HIV) infection. The annual risk of developing active tuberculosis when co-infected with HIV is 20-30 times the risk in non-HIV infected individual^[6]. In the Philippines there is a low prevalence of HIV infection^[7]. Thus, the HIV test was not done for this case since the patient had no high risk behavior. However, for those patients with risk factors (e.g. multiple sexual partners, men who have sex with men, intravenous drug users) and those from areas with high prevalence of HIV infection, testing for HIV co-infection may be beneficial. However, there is a lack of data on changes in the frequency or clinical manifestations of abdominal tuberculosis^[6].

The radiological features of duodenal tuberculosis are also non-specific. On barium studies, patients were found to have either one or a combination of mucosal ulcerations, luminal narrowing, extrinsic compression and proximal dilatations^[2]. Endoscopy may not be diagnostic and biopsies may only show nonspecific inflammation^[8]. In our case, biopsy was not performed during endoscopy since the patient was already deemed to require surgery due to the obstruction. Most case reports also diagnosed duodenal tuberculosis post-operatively. Diagnosis is made through histopathological findings of caseation necrosis and Langhans type giant cell.

Management of duodenal tuberculosis is still primarily medical. Studies have shown that if the diagnosis is made prior to surgery, most lesions improve with appropriate treatment^[9]. Even in patients with strictures, balloon dilatation has been shown to work together with medication^[10]. In this case, no trial of medication was done since biopsy was not performed pre-operatively. Performing biopsy pre-operatively may have changed the management if the biopsy proved to be diagnostic. Also, performing a tuberculosis polymerase chain reaction test may have helped in the diagnosis if tuberculosis was made as part of the differentials. However, the cost-effectiveness of this study may need to be assessed.

In conclusion, gastroduodenal tuberculosis is rare and pyloric stenosis resulting from tuberculosis is even rarer. There are no specific signs or symptoms and no characteristic endoscopic findings. It is our recommendation that among patients with a similar presentation who come from areas endemic for tuberculosis, every effort should be made to confirm the diagnosis to avoid unnecessary surgeries.

REFERENCES

- 1 **Rao YG**, Pande GK, Sahni P, Chattopadhyay TK. Gastro-duodenal tuberculosis management guidelines, based on a large experience and a review of the literature. *Can J Surg* 2004; **47**: 364-368
- 2 **Chavhan GE**, Ramakantan R. Duodenal tuberculosis: radiological features on barium studies and their clinical correlation in 28 cases. *J Postgrad Med* 2003; **49**: 214-217
- 3 **Gheorghe L**, Băncilă I, Gheorghe C, Herlea V, Vasilescu C, Aposteanu G. Antro-duodenal tuberculosis causing gastric outlet obstruction--a rare presentation of a protean disease. *Rom J Gastroenterol* 2002; **11**: 149-152
- 4 **Baqai MT**. Duodenal Tuberculosis: delays and difficulties in diagnosis. *J R Coll Physicians Edinb* 2005; **35**:330-331. Available from: URL: http://www.rcpe.ac.uk/journal/issue/journal_35_4/baqai.pdf
- 5 **Agrawal S**, Shetty SV, Bakshi G. Primary hypertrophic tuberculosis of the pyloroduodenal area: report of 2 cases. *J Postgrad Med* 1999; **45**: 10
- 6 **Sinkala E**, Gray S, Zulu I, Mudenda V, Zimba L, Vermund S, Drobniowski F, Kelly P. Clinical and ultrasonographic features of abdominal tuberculosis in HIV positive adults in Zambia. *BMC Infectious Diseases* 2009; **9**: 44
- 7 **Philippine HIV**, Registry A. Available from: URL: http://www.doh.gov.ph/files/NEC_HIV_Dec-AIDSreg2008.pdf
- 8 **Agashe P**, Dixit T. Duodenal tuberculosis a case report and review of literature. Available from: URL: http://www.bhj.org/journal/special_issue_tb/SP_4.HTM
- 9 **Anand BS**, Nanda R, Sachdev GK. Response of tuberculous stricture to antituberculous treatment. *Gut* 1988; **29**: 62-69
- 10 **Vij JC**, Ramesh GN, Choudhary V, Malhotra V. Endoscopic balloon dilation of tuberculous duodenal strictures. *Gastro-intest Endosc* 1992; **38**: 510-511
- 11 **Lamberty G**, Pappalardo E, Dresse D, Denoël A. Primary duodenal tuberculosis: a case report. *Acta Chir Belg* 2008; **108**: 590-591
- 12 **Sandeep N**, Prachis A, Saket K, Harsh U. Abdominal tuberculosis; Presentation, diagnosis and management. *Hungarian Med J* 2008; **2**: 201-214
- 13 **Khan R**, Abid S, Jafri W, Abbas Z, Hameed K, Ahmad Z. Diagnostic dilemma of abdominal tuberculosis in non-HIV patients: an ongoing challenge for physicians. *World J Gastroenterol* 2006; **12**: 6371-6375

- 14 **Manzelli A**, Stolfi VM, Spina C, Rossi P, Federico F, Canale S, Gaspari AL. Surgical treatment of gastric outlet obstruction due to gastroduodenal tuberculosis. *J Infect Chemother* 2008; **14**: 371-373
- 15 **Kshirsagar AY**, Kanetkar SR, Langade YB, Potwar SS, Shekhar N. Duodenal stenosis secondary to tuberculosis. *Int Surg* 2008; **93**: 265-267
- 16 **Tan KK**, Chen K, Sim R. The spectrum of abdominal tuberculosis in a developed country: a single institution's experience over 7 years. *J Gastrointest Surg* 2009; **13**: 142-147

S- Editor Zhang HN **L- Editor** Roemmele A **E- Editor** Liu N

Endoscopic retrieval of a gastric trichobezoar

Hironori Konuma, Kuangi Fu, Takashi Morimoto, Takayoshi Shimizu, Yuko Izumi, Satoko Shiyonagi, Masahiko Urao, Akihisa Miyazaki, Sumio Watanabe

Hironori Konuma, Kuangi Fu, Takashi Morimoto, Takayoshi Shimizu, Yuko Izumi, Akihisa Miyazaki, Department of Gastroenterology, Juntendo University Nerima Hospital, Tokyo 177-0033, Japan

Satoko Shiyonagi, Masahiko Urao, Department of Pediatric Surgery, Juntendo University Nerima Hospital, Tokyo 177-0033, Japan

Sumio Watanabe, Department of Gastroenterology, Juntendo University School of Medicine, Tokyo 177-0033, Japan

Author contributions: Konuma H and Fu KI supplied the data about this case report; Morimoto T, Shimizu T, Izumi Y, Shiyonagi S, Urao M, Miyazaki A, and Watanabe S analyzed the data of the patient; and Konuma H and Fu KI wrote the paper.

Correspondence to: Kuangi Fu, MD, PhD, Department of Gastroenterology, Juntendo University Nerima Hospital, 3-1-10 Nerimatakanodai, Nerima, Tokyo 177-0033, Japan. fukuangi@hotmail.com

Telephone: +81-3-59233111 Fax: +81-3-59233111

Received: September 13, 2010 Revised: October 22, 2010

Accepted: October 29, 2010

Published online: January 16, 2011

34 cm in length and 100 g in weight. The patient was discharged uneventfully 5 d thereafter. She was advised to visit a psychiatrist to avoid suffering from a relapse. Follow-up EGD showed no trichobezoar, and the patient's frontal hair grew back.

© 2010 Baishideng. All rights reserved.

Key words: Gastric bezoar; Trichobezoar; Endoscopic retrieval; Grasper; Retrieval net

Peer reviewers: Takayuki Yamamoto, MD, PhD, Inflammatory Bowel Disease Center, Yokkaichi Social Insurance Hospital, 10-8, Hazuyamacho, Yokkaichi 510-0016, Japan; Philip Wai Yan Chiu, Associate Professor, Department of Surgery, Institute of Digestive Disease, Chinese University of Hong Kong, Prince of Wales Hospital, 30-32 Ngan Shing Street, Shatin, N.T., Hong Kong, China

Konuma H, Fu K, Morimoto T, Shimizu T, Izumi Y, Shiyonagi S, Urao M, Miyazaki A, Watanabe S. Endoscopic retrieval of a gastric trichobezoar. *World J Gastrointest Endosc* 2011; 3(1): 20-22 Available from: URL: <http://www.wjgnet.com/1948-5190/full/v3/i1/20.htm> DOI: <http://dx.doi.org/10.4253/wjge.v3.i1.20>

Abstract

A 9-year-old girl presented with a chief complaint of abdominal pain. Esophagogastroduodenal endoscopy (EGD) identified a long and large gastric trichobezoar extending into the duodenum. We attempted endoscopic retrieval after informed consent was obtained from the patient's mother. Initially, a grasper with 5-prolongs, commonly used for retrieval of endoscopically excised polyps, failed to remove the whole trichobezoar. When a net was used instead, it proved impossible to remove the trichobezoar completely. Therefore, we withdrew the scope from the mouth, leaving the net grasping the trichobezoar firmly in the stomach. Subsequently, we were able to retrieve about 70% of the trichobezoar manually by grasping the snare part of the net directly. A second pass found no deep laceration or perforation endoscopically. The remaining trichobezoar was completely retrieved with the net. The procedure was completed within 15 min. The retrieved specimens were

INTRODUCTION

Bezoars are collections of indigestible materials that accumulate to form a mass in the gastrointestinal tract. Most are commonly found in the stomach, and they can be categorized into five major groups: (1) phytobezoars; (2) pharmacobezoars; (3) trichobezoars; (4) lactobezoars; and (5) foreign body bezoars, according to their composition. Trichobezoars, composed of hair, usually occur in young women and cases which extend throughout the small bowel into the cecum, are known as the Rapunzel syndrome^[1]. Gastric trichobezoars are generally more difficult to remove endoscopically and, thus, most reported cases require surgery^[2]. Herein we report a case of a gastric trichobezoar successfully retrieved with endoscopy.

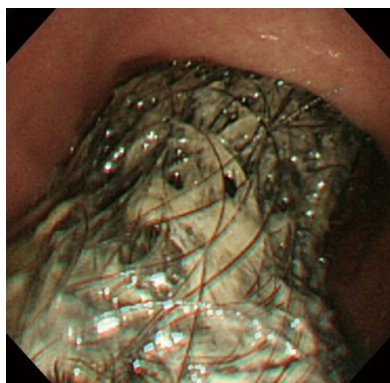


Figure 1 A transnasal esophagogastroduodenal endoscopy identified a long and large gastric trichobezoar extending into the duodenum.



Figure 2 A standard gastroscope (GIFQ260; Olympus, Tokyo, Japan) was used for retrieval. A grasper with 5-prolongs (Olympus, Tokyo, Japan), commonly used for retrieval of polyps excised endoscopically, was used to retrieve the gastric trichobezoar.

CASE REPORT

A 9-year-old girl presented with a chief complaint of abdominal pain. She had experienced bullying at her elementary school for two years. Over the same period, her mother noticed her habit of trichotillomania (hair pulling) and trichophagia (hair eating). The girl was unconscious of this habit but had tried to vomit the eaten hair when it was noticed or she was warned. She had no prior history of medical problems or mental disturbance. At first, she visited a nearby hospital because of abdominal pain. Physical examination showed a healthy 9-year-old girl with a body weight of 33 Kg and a height of 1.33 m. There was frontal balding and mild tenderness without defense or rigidity in the epigastric region, although no abdominal mass was palpable. An abdominal computed tomography (CT) revealed an inhomogenous mass in the stomach. Therefore, she was sent to our hospital for further investigation and treatment. Laboratory data were within normal limits except for a slight elevation of C-reactive protein (CRP; 1.3 g/dL).

Endoscopic technique

A transnasal esophagogastroduodenal endoscopy (EGD) identified a long and large gastric trichobezoar extending into the duodenum (Figure 1). We attempted endoscopic retrieval after informed consent had been obtained from



Figure 3 A net for collecting the removed colorectal polyps was used instead of a grasper for endoscopic retrieval.

the patient's mother. A standard gastroscope (GIFQ260; Olympus, Tokyo, Japan) was used after intravenous administration of midazolam (7 mg). To avoid bowel movement and to relax the lower esophageal sphincter, scopolamine butylbromide (20 mg) was administered intravenously before endoscopic removal. Carbon dioxide (CO₂) was administered using a CO₂ regulator (Olympus UCR; Olympus, Tokyo, Japan) connected to the endoscope supply tube for insufflation during the endoscopic procedure.

Initially, we used a grasper with 5-prolongs, commonly used for retrieval of endoscopically excised polyps, (Olympus, Tokyo, Japan). However we could only remove completely the portion of the trichobezoar present in the stomach as passage of esophagogastric junction (EGJ) was impossible (Figure 2). We then attempted retrieval using a net, also commonly used for retrieval of excised colorectal polyps (Figure 3). The trichobezoar did pass EGJ in part but we could not retrieve the whole of it with the endoscope. Therefore, we withdrew the scope from the mouth leaving the net grasping the trichobezoar firmly in the stomach. Subsequently, we were able to retrieve about 70% of the trichobezoar manually by grasping the snare part of the net directly. A second pass found mild erosion at the EGJ, although no deep laceration or perforation was detected endoscopically. The remaining trichobezoar was completely retrieved with the net. The whole procedure was electronically recorded and was completed within 15 min. The retrieved specimens were 1.8 cm × 3.2 cm × 34 cm in length and 100 g in weight (Figure 4). The patient was discharged uneventfully 5 d after endoscopic retrieval. She was advised to visit a psychiatrist to avoid suffering from a relapse. Follow-up EGD showed no trichobezoar, and her frontal hair grew back.

DISCUSSION

Endoscopic retrieval or surgical removal are chosen for bezoar removal based on the size and composition. As gastric trichobezoars are generally more difficult to remove endoscopically, most of the reported cases required surgery^[2]. Laparoscopic removal is cosmetically



Figure 4 The retrieved specimens were 1.8 cm × 3.2 cm × 34 cm in length and 100 g in weight.

superior to open laparotomy^[3,4]. However, if possible, endoscopic removal is less invasive and can save time, cost and abdominal damage. To the best of our knowledge, however, there are only two successful cases reported in the English literature^[5,6]. Soehendra used a Nd: YAG laser to disrupt the bezoar followed by the retrieval of its fragments, requiring more than 100 passages of the endoscope in three sessions of 2 to 3 h^[5]. This procedure is, therefore, troublesome and needs special equipment. Meanwhile, Saeed carried out successful retrieval using a two-channel endoscope, an overtube, and a grasping forceps with the patient under intubation^[6].

In our case, we used a single channel endoscope, a grasper, and a net to remove the trichobezoar completely in one session (two passages of endoscope) with the patient under sedation and without intubation. We conducted the endoscopic procedure under sedation instead of general anesthesia after discussion with the anesthetists and pediatric surgeons in our hospital. We would have performed the procedure under general anesthesia if we could not have completed the endoscopic retrieval safely within 30 min. The shape of the trichobezoar in our case was most important for successful endoscopic retrieval. If the maximal diameter had been too large to

pass the esophagogastric junction, we would not have been able to retrieve it endoscopically. Considering the whole procedure retrospectively, the grasping power of a grasper with 5-prolongs was not as great as that of a net, although the trichobezoar could not be removed with the endoscope even used together with a net. If we had used a two channel endoscope backloaded with an overtube according to Saeed's suggestion, the bezoar would have been grasped more firmly, enabling the passage of the EGJ more easily. However, holding the snare part of the net directly following withdrawing the endoscope from the mouth, eventually enabled us to retrieve the grasped trichobezoar successfully. Furthermore, we used CO₂ insufflation to avoid esophageal perforation associated with laceration caused by endoscopic retrieval.

In conclusion, we have reported a case of a gastric trichobezoar successfully retrieved endoscopically using a net made for polyp retrieval. Endoscopic removal of a gastric trichobezoar is less invasive and cost effective than surgical removal and this procedure should, therefore, be considered as an option for treatment with various endoscopic accessories.

REFERENCES

- 1 **Vaughan ED Jr**, Sawyers JL, Scott HW Jr. The Rapunzel syndrome. An unusual complication of intestinal bezoar. *Surgery* 1968; **63**: 339-343
- 2 **De Backer A**, Van Nooten V, Vandenplas Y. Huge gastric trichobezoar in a 10-year-old girl: case report with emphasis on endoscopy in diagnosis and therapy. *J Pediatr Gastroenterol Nutr* 1999; **28**: 513-515
- 3 **Nirasawa Y**, Mori T, Ito Y, Tanaka H, Seki N, Atomi Y. Laparoscopic removal of a large gastric trichobezoar. *J Pediatr Surg* 1998; **33**: 663-665
- 4 **Kanetaka K**, Azuma T, Ito S, Matsuo S, Yamaguchi S, Shirono K, Kanematsu T. Two-channel method for retrieval of gastric trichobezoar: report of a case. *J Pediatr Surg* 2003; **38**: e7
- 5 **Soehendra N**. Endoscopic removal of a trichobezoar. *Endoscopy* 1989; **21**: 201
- 6 **Saeed ZA**, Ramirez FC, Hepps KS, Dixon WB. A method for the endoscopic retrieval of trichobezoars. *Gastrointest Endosc* 1993; **39**: 698-700

S- Editor Zhang HN L- Editor Hughes D E- Editor Liu N



Acknowledgments to reviewers of *World Journal of Gastrointestinal Endoscopy*

Many reviewers have contributed their expertise and time to the peer review, a critical process to ensure the quality of *World Journal of Gastrointestinal Endoscopy*. The editors and authors of the articles submitted to the journal are grateful to the following reviewers for evaluating the articles (including those published in this issue and those rejected for this issue) during the last editing time period.

Hongchun Bao, PhD, Research Fellow, The Center for Micro-Photonics, Faculty of Engineering & Industrial Sciences, Swinburne University of Technology, PO Box 218, Hawthorn, Victoria 3122, Australia

Philip Wai Yan Chiu, Associate Professor, Department of Surgery, Institute of Digestive Disease, Chinese University of Hong Kong, Prince of Wales Hospital, 30-32 Ngan Shing Street, Shatin, N.T., Hong Kong, China

David Friedel, MD, Gastroenterology, Winthrop University Hospital, 222 Station Plaza North, Suite 428, Mineola NY 11501, United States

Lesur Gilles, MD, Hopital Ambroise Paré, 9 avenue Charles de Gaulle, Boulogne 92100, France

Carlo M Girelli, MD, 1st Department of Internal Medicine, Ser-

vice of Gastroenterology and Digestive Endoscopy, Hospital of Busto Arsizio, Via Arnaldo da Brescia, Busto Arsizio, VA 121052, Italy

Dimitrios Kapetanios, MD, Gastroenterology Department, George Papanikolaou Hospital, Exohi, Thessaloniki 57010, Greece

Sanjiv Mahadeva, MBBS, MRCP, CCST, MD, Associate Professor, Department of Medicine, Faculty of Medicine, University of Malaya, Kuala Lumpur 50603, Malaysia

Oliver Pech, MD, PhD, Attending Physician of Gastroenterology, Vice Director of the Endoscopy Unit, Department of Internal Medicine 2, HSK Wiesbaden, Wiesbaden, Germany

Omar Javed Shah, Professor, Head, Department of Surgical Gastroenterology, Sher-i-Kashmir Institute of Medical Sciences, Srinagar, Kashmir, India

Takashi Shida, MD, PhD, Department of General Surgery, Chiba University Graduate School of Medicine, 1-8-1 Inohana, Chuo-ku, Chiba 260-8670, Japan

Noriya Uedo, Director, Endoscopic Training and Learning Center; Vice Director, Department of Gastrointestinal Oncology, Osaka Medical Center for Cancer and Cardiovascular Diseases, Osaka, Japan

Takayuki Yamamoto, MD, PhD, Inflammatory Bowel Disease Center, Yokkaichi Social Insurance Hospital, 10-8, Hazuyamacho, Yokkaichi 510-0016, Japan

Meetings

Events Calendar 2011

January 14-15, 2011
AGA Clinical Congress of
Gastroenterology and Hepatology:
Best Practices in 2011
Miami, FL 33101, United States

January 20-22, 2011
Gastrointestinal Cancers Symposium
2011
San Francisco, CA 94143,
United States

January 28-29, 2011
9. Gastro Forum München
Munich, Germany

February 04-05, 2011
13th Duesseldorf International
Endoscopy Symposium
Duesseldorf, Germany

February 13-27, 2011
Gastroenterology: New Zealand
CME Cruise Conference
Sydney, NSW, Australia

February 24-26, 2011
Inflammatory Bowel Diseases
2011-6th Congress of the European
Crohn's and Colitis Organisation
Dublin, Ireland

February 24-26, 2011
2nd International Congress on
Abdominal Obesity
Buenos Aires, Brazil

February 26-March 1, 2011
Canadian Digestive Diseases Week
Westin Bayshore, Vancouver
British Columbia, Canada

March 03-05, 2011
42nd Annual Topics in Internal
Medicine
Gainesville, FL 32614,
United States

March 14-17, 2011
British Society of Gastroenterology
Annual Meeting 2011
Birmingham, England, United
Kingdom

March 17-19, 2011
41. Kongress der Deutschen
Gesellschaft für Endoskopie und
Bildgebende Verfahren e.V.
Munich, Germany

March 17-20, 2011
Mayo Clinic Gastroenterology &
Hepatology 2011
Jacksonville, FL 34234, United States

March 25-27, 2011
MedicReS IC 2011 Good Medical
Research
Istanbul, Turkey

April 07-09, 2011
International and Interdisciplinary
Conference Excellence in Female
Surgery
Florence, Italy

April 15-16, 2011
Falk Symposium 177, Endoscopy
Live Berlin 2011 Intestinal Disease
Meeting, Stauffenbergstr. 26
Berlin 10785, Germany

April 18-22, 2011
Pediatric Emergency Medicine:
Detection, Diagnosis and Developing
Treatment Plans
Sarasota, FL 34234, United States

April 20-23, 2011
9th International Gastric Cancer
Congress, COEX, World Trade
Center, Samseong-dong
Seoul 135-731, South Korea

April 25-27, 2011
The Second International Conference
of the Saudi Society of Pediatric
Gastroenterology, Hepatology &
Nutrition
Riyadh, Saudi Arabia

April 28-30, 2011
4th Central European Congress of
Surgery
Budapest, Hungary

May 07-10, 2011
Digestive Disease Week
Chicago, IL 60446, United States

May 12-13, 2011
2nd National Conference Clinical
Advances in Cystic Fibrosis
London, England, United Kingdom

May 21-24, 2011
22nd European Society of
Gastrointestinal and Abdominal
Radiology Annual Meeting and
Postgraduate Course
Venice, Italy

May 25-28, 2011
4th Congress of the Gastroenterology
Association of Bosnia and

Herzegovina with international
participation, Hotel Holiday Inn
Sarajevo, Bosnia and Herzegovina

June 11-12, 2011
The International Digestive Disease
Forum 2011
Hong Kong, China

June 13-16, 2011
Surgery and Disillusion XXIV Spige
II ESYS, Napoli, Italy

June 22-25, 2011
ESMO Conference: 13th World
Congress on Gastrointestinal Cancer
Barcelona, Spain

September 10-11, 2011
New Advances in Inflammatory
Bowel Disease
La Jolla, CA 92093, United States

September 10-14, 2011
ICE 2011-International Congress of
Endoscopy, Los Angeles Convention
Center, 1201 South Figueroa Street
Los Angeles, CA 90015, United
States

September 30-October 1, 2011
Falk Symposium 179, Revisiting
IBD Management: Dogmas to be
Challenged, Sheraton Brussels Hotel
Brussels 1210, Belgium

October 19-29, 2011
Cardiology & Gastroenterology
Tahiti 10 night CME Cruise
Papeete, French Polynesia

October 22-26, 2011
19th United European
Gastroenterology Week
Stockholm, Sweden

October 28-November 02, 2011
ACG Annual Scientific Meeting &
Postgraduate Course
Washington, DC 20001, United
States

November 11-12, 2011
Falk Symposium 180, IBD 2011:
Progress and Future for Lifelong
Management, ANA Interconti Hotel,
1-12-33 Akasaka, Minato-ku
Tokyo 107-0052, Japan

December 01-04, 2011
2011 Advances in Inflammatory
Bowel Diseases/Crohn's & Colitis
Foundation's Clinical & Research
Conference
Hollywood, FL 34234, United States



Instructions to authors

GENERAL INFORMATION

World Journal of Gastrointestinal Endoscopy (*World J Gastrointest Endosc*, *WJGE*, online ISSN 1948-5190, DOI: 10.4253), is a monthly, open-access (OA), peer-reviewed online journal supported by an editorial board of 400 experts in gastrointestinal endoscopy from 45 countries.

The biggest advantage of the OA model is that it provides free, full-text articles in PDF and other formats for experts and the public without registration, which eliminates the obstacle that traditional journals possess and usually delays the speed of the propagation and communication of scientific research results.

Maximization of personal benefits

The role of academic journals is to exhibit the scientific levels of a country, a university, a center, a department, and even a scientist, and build an important bridge for communication between scientists and the public. As we all know, the significance of the publication of scientific articles lies not only in disseminating and communicating innovative scientific achievements and academic views, as well as promoting the application of scientific achievements, but also in formally recognizing the "priority" and "copyright" of innovative achievements published, as well as evaluating research performance and academic levels. So, to realize these desired attributes of *WJGE* and create a well-recognized journal, the following four types of personal benefits should be maximized. The maximization of personal benefits refers to the pursuit of the maximum personal benefits in a well-considered optimal manner without violation of the laws, ethical rules and the benefits of others. (1) Maximization of the benefits of editorial board members: The primary task of editorial board members is to give a peer review of an unpublished scientific article via online office system to evaluate its innovativeness, scientific and practical values and determine whether it should be published or not. During peer review, editorial board members can also obtain cutting-edge information in that field at first hand. As leaders in their field, they have priority to be invited to write articles and publish commentary articles. We will put peer reviewers' names and affiliations along with the article they reviewed in the journal to acknowledge their contribution; (2) Maximization of the benefits of authors: Since *WJGE* is an open-access journal, readers around the world can immediately download and read, free of charge, high-quality, peer-reviewed articles from *WJGE* official website, thereby realizing the goals and significance of the communication between authors and peers as well as public reading; (3) Maximization of the benefits of readers: Readers can read or use, free of charge, high-quality peer-reviewed articles without any limits, and cite the arguments, viewpoints, concepts, theories, methods, results, conclusion or facts and data of pertinent literature so as to validate the innovativeness, scientific and practical values of their own research achievements, thus ensuring that their articles have novel arguments or viewpoints, solid evidence and correct conclusion; and (4) Maximization of the benefits of employees: It is an iron law that a first-class journal is unable to exist without first-class editors, and only first-class editors can create a first-class academic journal. We insist on strengthening our team cultivation and construction so that every employee, in an open, fair and transparent environment, could contribute their wisdom to edit and publish high-quality articles, thereby realizing the maximization of the personal benefits of editorial board members, authors and readers, and yielding the greatest social and economic benefits.

Aims and scope

The major task of *WJGE* is to report rapidly the most recent results in basic and clinical research on gastrointestinal endoscopy including: gastroscopy, intestinal endoscopy, colonoscopy, capsule endoscopy, laparoscopy, interventional diagnosis and therapy, as well as advances in technology. Emphasis is placed on the clinical practice of treating gastrointestinal diseases with or under endoscopy. Papers on advances and application of endoscopy-associated techniques, such as endoscopic ultrasonography, endoscopic retrograde cholangiopancreatography, endoscopic submucosal dissection and endoscopic balloon dilation are also welcome.

Columns

The columns in the issues of *WJGE* will include: (1) Editorial: To introduce and comment on major advances and developments in the field; (2) Frontier: To review representative achievements, comment on the state of current research, and propose directions for future research; (3) Topic Highlight: This column consists of three formats, including (A) 10 invited review articles on a hot topic, (B) a commentary on common issues of this hot topic, and (C) a commentary on the 10 individual articles; (4) Observation: To update the development of old and new questions, highlight unsolved problems, and provide strategies on how to solve the questions; (5) Guidelines for Basic Research: To provide guidelines for basic research; (6) Guidelines for Clinical Practice: To provide guidelines for clinical diagnosis and treatment; (7) Review: To review systemically progress and unresolved problems in the field, comment on the state of current research, and make suggestions for future work; (8) Original Article: To report innovative and original findings in gastrointestinal endoscopy; (9) Brief Article: To briefly report the novel and innovative findings in gastrointestinal endoscopy; (10) Case Report: To report a rare or typical case; (11) Letters to the Editor: To discuss and make reply to the contributions published in *WJGE*, or to introduce and comment on a controversial issue of general interest; (12) Book Reviews: To introduce and comment on quality monographs of gastrointestinal endoscopy; and (13) Guidelines: To introduce consensus and guidelines reached by international and national academic authorities worldwide on basic research and clinical practice in gastrointestinal endoscopy.

Name of journal

World Journal of Gastrointestinal Endoscopy

CSSN

ISSN 1948-5190 (online)

Indexed and Abstracted in

PubMed Central, PubMed

Published by

Baishideng Publishing Group Co., Limited

SPECIAL STATEMENT

All articles published in this journal represent the viewpoints of the authors except where indicated otherwise.

Biostatistical editing

Statistical review is performed after peer review. We invite an expert in Biomedical Statistics from to evaluate the statistical method used in the paper, including *t*-test (group or paired comparisons), chi-

Instructions to authors

squared test, Ridit, probit, logit, regression (linear, curvilinear, or stepwise), correlation, analysis of variance, analysis of covariance, *etc.* The reviewing points include: (1) Statistical methods should be described when they are used to verify the results; (2) Whether the statistical techniques are suitable or correct; (3) Only homogeneous data can be averaged. Standard deviations are preferred to standard errors. Give the number of observations and subjects (*n*). Losses in observations, such as drop-outs from the study should be reported; (4) Values such as ED50, LD50, IC50 should have their 95% confidence limits calculated and compared by weighted probit analysis (Bliss and Finney); and (5) The word 'significantly' should be replaced by its synonyms (if it indicates extent) or the *P* value (if it indicates statistical significance).

Conflict-of-interest statement

In the interests of transparency and to help reviewers assess any potential bias, *WJGE* requires authors of all papers to declare any competing commercial, personal, political, intellectual, or religious interests in relation to the submitted work. Referees are also asked to indicate any potential conflict they might have reviewing a particular paper. Before submitting, authors are suggested to read "Uniform Requirements for Manuscripts Submitted to Biomedical Journals: Ethical Considerations in the Conduct and Reporting of Research: Conflicts of Interest" from International Committee of Medical Journal Editors (ICMJE), which is available at: http://www.icmje.org/ethical_4conflicts.html.

Sample wording: [Name of individual] has received fees for serving as a speaker, a consultant and an advisory board member for [names of organizations], and has received research funding from [names of organization]. [Name of individual] is an employee of [name of organization]. [Name of individual] owns stocks and shares in [name of organization]. [Name of individual] owns patent [patent identification and brief description].

Statement of informed consent

Manuscripts should contain a statement to the effect that all human studies have been reviewed by the appropriate ethics committee or it should be stated clearly in the text that all persons gave their informed consent prior to their inclusion in the study. Details that might disclose the identity of the subjects under study should be omitted. Authors should also draw attention to the Code of Ethics of the World Medical Association (Declaration of Helsinki, 1964, as revised in 2004).

Statement of human and animal rights

When reporting the results from experiments, authors should follow the highest standards and the trial should conform to Good Clinical Practice (for example, US Food and Drug Administration Good Clinical Practice in FDA-Regulated Clinical Trials; UK Medicines Research Council Guidelines for Good Clinical Practice in Clinical Trials) and/or the World Medical Association Declaration of Helsinki. Generally, we suggest authors follow the lead investigator's national standard. If doubt exists whether the research was conducted in accordance with the above standards, the authors must explain the rationale for their approach and demonstrate that the institutional review body explicitly approved the doubtful aspects of the study.

Before submitting, authors should make their study approved by the relevant research ethics committee or institutional review board. If human participants were involved, manuscripts must be accompanied by a statement that the experiments were undertaken with the understanding and appropriate informed consent of each. Any personal item or information will not be published without explicit consents from the involved patients. If experimental animals were used, the materials and methods (experimental procedures) section must clearly indicate that appropriate measures were taken to minimize pain or discomfort, and details of animal care should be provided.

SUBMISSION OF MANUSCRIPTS

Manuscripts should be typed in 1.5 line spacing and 12 pt. Book

Antiqua with ample margins. Number all pages consecutively, and start each of the following sections on a new page: Title Page, Abstract, Introduction, Materials and Methods, Results, Discussion, Acknowledgements, References, Tables, Figures, and Figure Legends. Neither the editors nor the publisher are responsible for the opinions expressed by contributors. Manuscripts formally accepted for publication become the permanent property of Baishideng Publishing Group Co., Limited, and may not be reproduced by any means, in whole or in part, without the written permission of both the authors and the publisher. We reserve the right to copy-edit and put onto our website accepted manuscripts. Authors should follow the relevant guidelines for the care and use of laboratory animals of their institution or national animal welfare committee. For the sake of transparency in regard to the performance and reporting of clinical trials, we endorse the policy of the International Committee of Medical Journal Editors to refuse to publish papers on clinical trial results if the trial was not recorded in a publicly-accessible registry at its outset. The only register now available, to our knowledge, is <http://www.clinicaltrials.gov> sponsored by the United States National Library of Medicine and we encourage all potential contributors to register with it. However, in the case that other registers become available you will be duly notified. A letter of recommendation from each author's organization should be provided with the contributed article to ensure the privacy and security of research is protected.

Authors should retain one copy of the text, tables, photographs and illustrations because rejected manuscripts will not be returned to the author(s) and the editors will not be responsible for loss or damage to photographs and illustrations sustained during mailing.

Online submissions

Manuscripts should be submitted through the Online Submission System at: wjge@wjgnet.com. Authors are highly recommended to consult the ONLINE INSTRUCTIONS TO AUTHORS (http://www.wjgnet.com/1948-5190/g_info_20100316080002.htm) before attempting to submit online. For assistance, authors encountering problems with the Online Submission System may send an email describing the problem to <http://www.wjgnet.com/1948-5190office/>, or by telephone: +86-10-59080038. If you submit your manuscript online, do not make a postal contribution. Repeated online submission for the same manuscript is strictly prohibited.

MANUSCRIPT PREPARATION

All contributions should be written in English. All articles must be submitted using word-processing software. All submissions must be typed in 1.5 line spacing and 12 pt. Book Antiqua with ample margins. Style should conform to our house format. Required information for each of the manuscript sections is as follows:

Title page

Title: Title should be less than 12 words.

Running title: A short running title of less than 6 words should be provided.

Authorship: Authorship credit should be in accordance with the standard proposed by International Committee of Medical Journal Editors, based on (1) substantial contributions to conception and design, acquisition of data, or analysis and interpretation of data; (2) drafting the article or revising it critically for important intellectual content; and (3) final approval of the version to be published. Authors should meet conditions 1, 2, and 3.

Institution: Author names should be given first, then the complete name of institution, city, province and postcode. For example, Xu-Chen Zhang, Li-Xin Mei, Department of Pathology, Chengde Medical College, Chengde 067000, Hebei Province, China. One author may be represented from two institutions, for example, George Sgourakis, Department of General, Visceral, and

Transplantation Surgery, Essen 45122, Germany; George Sgourakis, 2nd Surgical Department, Korgialenio-Benakio Red Cross Hospital, Athens 15451, Greece

Author contributions: The format of this section should be: Author contributions: Wang CL and Liang L contributed equally to this work; Wang CL, Liang L, Fu JF, Zou CC, Hong F and Wu XM designed the research; Wang CL, Zou CC, Hong F and Wu XM performed the research; Xue JZ and Lu JR contributed new reagents/analytic tools; Wang CL, Liang L and Fu JF analyzed the data; and Wang CL, Liang L and Fu JF wrote the paper.

Supportive foundations: The complete name and number of supportive foundations should be provided, e.g., Supported by National Natural Science Foundation of China, No. 30224801

Correspondence to: Only one corresponding address should be provided. Author names should be given first, then author title, affiliation, the complete name of institution, city, postcode, province, country, and email. All the letters in the email should be in lower case. A space interval should be inserted between country name and email address. For example, Montgomery Bissell, MD, Professor of Medicine, Chief, Liver Center, Gastroenterology Division, University of California, Box 0538, San Francisco, CA 94143, United States. montgomery.bissell@ucsf.edu

Telephone and fax: Telephone and fax should consist of +, country number, district number and telephone or fax number, e.g., Telephone: +86-10-59080039 Fax: +86-10-85381893

Peer reviewers: All articles received are subject to peer review. Normally, three experts are invited for each article. Decision for acceptance is made only when at least two experts recommend an article for publication. Reviewers for accepted manuscripts are acknowledged in each manuscript, and reviewers of articles which were not accepted will be acknowledged at the end of each issue. To ensure the quality of the articles published in *WJGE*, reviewers of accepted manuscripts will be announced by publishing the name, title/position and institution of the reviewer in the footnote accompanying the printed article. For example, reviewers: Professor Jing-Yuan Fang, Shanghai Institute of Digestive Disease, Shanghai, Affiliated Renji Hospital, Medical Faculty, Shanghai Jiaotong University, Shanghai, China; Professor Xin-Wei Han, Department of Radiology, The First Affiliated Hospital, Zhengzhou University, Zhengzhou, Henan Province, China; and Professor Anren Kuang, Department of Nuclear Medicine, Huaxi Hospital, Sichuan University, Chengdu, Sichuan Province, China.

Abstract

There are unstructured abstracts (no more than 256 words) and structured abstracts (no more than 480). The specific requirements for structured abstracts are as follows:

An informative, structured abstracts of no more than 480 words should accompany each manuscript. Abstracts for original contributions should be structured into the following sections. AIM (no more than 20 words): Only the purpose should be included. Please write the aim as the form of "To investigate/study/...; MATERIALS AND METHODS (no more than 140 words); RESULTS (no more than 294 words): You should present *P* values where appropriate and must provide relevant data to illustrate how they were obtained, e.g. 6.92 ± 3.86 vs 3.61 ± 1.67 , $P < 0.001$; CONCLUSION (no more than 26 words).

Key words

Please list 5-10 key words, selected mainly from *Index Medicus*, which reflect the content of the study.

Text

For articles of these sections, original articles, rapid communication and case reports, the main text should be structured into the

following sections: INTRODUCTION, MATERIALS AND METHODS, RESULTS and DISCUSSION, and should include appropriate Figures and Tables. Data should be presented in the main text or in Figures and Tables, but not in both. The main text format of these sections, editorial, topic highlight, case report, letters to the editors, can be found at: http://www.wjgnet.com/1948-5190/g_info_20100316080002.htm.

Illustrations

Figures should be numbered as 1, 2, 3, *etc.*, and mentioned clearly in the main text. Provide a brief title for each figure on a separate page. Detailed legends should not be provided under the figures. This part should be added into the text where the figures are applicable. Figures should be either Photoshop or Illustrator files (in tiff, eps, jpeg formats) at high-resolution. Examples can be found at: <http://www.wjgnet.com/1007-9327/13/4520.pdf>; <http://www.wjgnet.com/1007-9327/13/4554.pdf>; <http://www.wjgnet.com/1007-9327/13/4891.pdf>; <http://www.wjgnet.com/1007-9327/13/4986.pdf>; <http://www.wjgnet.com/1007-9327/13/4498.pdf>. Keeping all elements compiled is necessary in line-art image. Scale bars should be used rather than magnification factors, with the length of the bar defined in the legend rather than on the bar itself. File names should identify the figure and panel. Avoid layering type directly over shaded or textured areas. Please use uniform legends for the same subjects. For example: Figure 1 Pathological changes in atrophic gastritis after treatment. A: ...; B: ...; C: ...; D: ...; E: ...; F: ...; G: ...*etc.* It is our principle to publish high resolution-figures for the printed and E-versions.

Tables

Three-line tables should be numbered 1, 2, 3, *etc.*, and mentioned clearly in the main text. Provide a brief title for each table. Detailed legends should not be included under tables, but rather added into the text where applicable. The information should complement, but not duplicate the text. Use one horizontal line under the title, a second under column heads, and a third below the Table, above any footnotes. Vertical and italic lines should be omitted.

Notes in tables and illustrations

Data that are not statistically significant should not be noted. ^a*P* < 0.05, ^b*P* < 0.01 should be noted (*P* > 0.05 should not be noted). If there are other series of *P* values, ^c*P* < 0.05 and ^d*P* < 0.01 are used. A third series of *P* values can be expressed as ^e*P* < 0.05 and ^f*P* < 0.01. Other notes in tables or under illustrations should be expressed as ¹F, ²F, ³F; or sometimes as other symbols with a superscript (Arabic numerals) in the upper left corner. In a multi-curve illustration, each curve should be labeled with ●, ○, ■, □, ▲, △, *etc.*, in a certain sequence.

Acknowledgments

Brief acknowledgments of persons who have made genuine contributions to the manuscript and who endorse the data and conclusions should be included. Authors are responsible for obtaining written permission to use any copyrighted text and/or illustrations.

REFERENCES

Coding system

The author should number the references in Arabic numerals according to the citation order in the text. Put reference numbers in square brackets in superscript at the end of citation content or after the cited author's name. For citation content which is part of the narration, the coding number and square brackets should be typeset normally. For example, "Crohn's disease (CD) is associated with increased intestinal permeability^[1,2]". If references are cited directly in the text, they should be put together within the text, for example, "From references^[19,22-24], we know that..."

When the authors write the references, please ensure that the order in text is the same as in the references section, and also

Instructions to authors

ensure the spelling accuracy of the first author's name. Do not list the same citation twice.

PMID and DOI

Please provide PubMed citation numbers to the reference list, e.g. PMID and DOI, which can be found at <http://www.ncbi.nlm.nih.gov/sites/entrez?db=pubmed> and <http://www.crossref.org/SimpleTextQuery/>, respectively. The numbers will be used in E-version of this journal.

Style for journal references

Authors: the name of the first author should be typed in bold-faced letters. The family name of all authors should be typed with the initial letter capitalized, followed by their abbreviated first and middle initials. (For example, Lian-Sheng Ma is abbreviated as Ma LS, Bo-Rong Pan as Pan BR). The title of the cited article and italicized journal title (journal title should be in its abbreviated form as shown in PubMed), publication date, volume number (in black), start page, and end page [PMID: 11819634 DOI: 10.3748/wjg.13.5396].

Style for book references

Authors: the name of the first author should be typed in bold-faced letters. The surname of all authors should be typed with the initial letter capitalized, followed by their abbreviated middle and first initials. (For example, Lian-Sheng Ma is abbreviated as Ma LS, Bo-Rong Pan as Pan BR) Book title. Publication number. Publication place: Publication press, Year: start page and end page.

Format

Journals

English journal article (list all authors and include the PMID where applicable)

- 1 **Jung EM**, Clevert DA, Schreyer AG, Schmitt S, Rennert J, Kubale R, Feuerbach S, Jung F. Evaluation of quantitative contrast harmonic imaging to assess malignancy of liver tumors: A prospective controlled two-center study. *World J Gastroenterol* 2007; **13**: 6356-6364 [PMID: 18081224 DOI: 10.3748/wjg.13.6356]

Chinese journal article (list all authors and include the PMID where applicable)

- 2 **Lin GZ**, Wang XZ, Wang P, Lin J, Yang FD. Immunologic effect of Jianpi Yishen decoction in treatment of Pixu-diarhoea. *Shijie Huaren Xiaobua Zazhi* 1999; **7**: 285-287

In press

- 3 **Tian D**, Araki H, Stahl E, Bergelson J, Kreitman M. Signature of balancing selection in Arabidopsis. *Proc Natl Acad Sci USA* 2006; In press

Organization as author

- 4 **Diabetes Prevention Program Research Group**. Hypertension, insulin, and proinsulin in participants with impaired glucose tolerance. *Hypertension* 2002; **40**: 679-686 [PMID: 12411462 PMCID: 2516377 DOI:10.1161/01.HYP.0000035706.28494.09]

Both personal authors and an organization as author

- 5 **Vallancien G**, Emberton M, Harving N, van Moorselaar RJ; Alf-One Study Group. Sexual dysfunction in 1, 274 European men suffering from lower urinary tract symptoms. *J Urol* 2003; **169**: 2257-2261 [PMID: 12771764 DOI:10.1097/01.ju.0000067940.76090.73]

No author given

- 6 21st century heart solution may have a sting in the tail. *BMJ* 2002; **325**: 184 [PMID: 12142303 DOI:10.1136/bmj.325.7357.184]

Volume with supplement

- 7 **Geraud G**, Spierings EL, Keywood C. Tolerability and safety of frovatriptan with short- and long-term use for treatment of migraine and in comparison with sumatriptan. *Headache* 2002; **42** Suppl 2: S93-99 [PMID: 12028325 DOI:10.1046/j.1526-4610.42.s2.7.x]

Issue with no volume

- 8 **Banit DM**, Kaufer H, Hartford JM. Intraoperative frozen section analysis in revision total joint arthroplasty. *Clin Orthop Relat Res* 2002; **(401)**: 230-238 [PMID: 12151900 DOI:10.1097/00003086-200208000-00026]

No volume or issue

- 9 Outreach: Bringing HIV-positive individuals into care. *HRSA Careaction* 2002; 1-6 [PMID: 12154804]

Books

Personal author(s)

- 10 **Sherlock S**, Dooley J. Diseases of the liver and biliary system. 9th ed. Oxford: Blackwell Sci Pub, 1993: 258-296

Chapter in a book (list all authors)

- 11 **Lam SK**. Academic investigator's perspectives of medical treatment for peptic ulcer. In: Swabb EA, Azabo S. Ulcer disease: investigation and basis for therapy. New York: Marcel Dekker, 1991: 431-450

Author(s) and editor(s)

- 12 **Breedlove GK**, Schorfheide AM. Adolescent pregnancy. 2nd ed. Wicczorek RR, editor. White Plains (NY): March of Dimes Education Services, 2001: 20-34

Conference proceedings

- 13 **Harnden P**, Joffe JK, Jones WG, editors. Germ cell tumours V. Proceedings of the 5th Germ cell tumours Conference; 2001 Sep 13-15; Leeds, UK. New York: Springer, 2002: 30-56

Conference paper

- 14 **Christensen S**, Oppacher F. An analysis of Koza's computational effort statistic for genetic programming. In: Foster JA, Lutton E, Miller J, Ryan C, Tettamanzi AG, editors. Genetic programming. EuroGP 2002: Proceedings of the 5th European Conference on Genetic Programming; 2002 Apr 3-5; Kinsdale, Ireland. Berlin: Springer, 2002: 182-191

Electronic journal (list all authors)

- 15 Morse SS. Factors in the emergence of infectious diseases. Emerg Infect Dis serial online, 1995-01-03, cited 1996-06-05; 1(1): 24 screens. Available from: URL: <http://www.cdc.gov/ncidod/eid/index.htm>

Patent (list all authors)

- 16 **Pagedas AC**, inventor; Ancel Surgical R&D Inc., assignee. Flexible endoscopic grasping and cutting device and positioning tool assembly. United States patent US 20020103498. 2002 Aug 1

Statistical data

Write as mean \pm SD or mean \pm SE.

Statistical expression

Express *t* test as *t* (in italics), *F* test as *F* (in italics), chi square test as χ^2 (in Greek), related coefficient as *r* (in italics), degree of freedom as *v* (in Greek), sample number as *n* (in italics), and probability as *P* (in italics).

Units

Use SI units. For example: body mass, *m* (B) = 78 kg; blood pressure, *p* (B) = 16.2/12.3 kPa; incubation time, *t* (incubation) = 96 h; blood glucose concentration, *c* (glucose) 6.4 ± 2.1 mmol/L; blood CEA mass concentration, *p* (CEA) = 8.6 ± 24.5 μ g/L; CO₂ volume fraction, 50 mL/L CO₂, not 5% CO₂; likewise for 40 g/L formaldehyde, not 10% formalin; and mass fraction, 8 ng/g, etc. Arabic numerals such as 23, 243, 641 should be read 23 243 641.

The format for how to accurately write common units and quantities can be found at: <http://www.wjgnet.com/wjg/help/15.doc>.

Abbreviations

Standard abbreviations should be defined in the abstract and on first mention in the text. In general, terms should not be abbreviated unless they are used repeatedly and the abbreviation is helpful to the reader. Permissible abbreviations are listed in Units, Symbols and Abbreviations: A Guide for Biological and Medical Editors and

Authors (Ed. Baron DN, 1988) published by The Royal Society of Medicine, London. Certain commonly used abbreviations, such as DNA, RNA, HIV, LD50, PCR, HBV, ECG, WBC, RBC, CT, ESR, CSF, IgG, ELISA, PBS, ATP, EDTA, mAb, can be used directly without further explanation.

Italics

Quantities: *t* time or temperature, *c* concentration, *A* area, *l* length, *m* mass, *V* volume.

Genotypes: *gyrA*, *arg 1*, *c myc*, *c fos*, etc.

Restriction enzymes: *EcoRI*, *HindI*, *BamHI*, *Kbo I*, *Kpn I*, etc.

Biology: *H. pylori*, *E. coli*, etc.

Examples for paper writing

Editorial: http://www.wjgnet.com/1948-5190/g_info_20100316080004.htm

Frontier: http://www.wjgnet.com/1948-5190/g_info_20100313155344.htm

Topic highlight: http://www.wjgnet.com/1948-5190/g_info_20100316080006.htm

Observation: http://www.wjgnet.com/1948-5190/g_info_201007124105.htm

Guidelines for basic research: http://www.wjgnet.com/1948-5190/g_info_20100313155908.htm

Guidelines for clinical practice: http://www.wjgnet.com/1948-5190/g_info_20100313160015.htm

Review: http://www.wjgnet.com/1948-5190/g_info_201007124313.htm

Original articles: http://www.wjgnet.com/1948-5190/g_info_201007133454.htm

Brief articles: http://www.wjgnet.com/1948-5190/g_info_20100313160645.htm

Case report: http://www.wjgnet.com/1948-5190/g_info_201007133659.htm

Letters to the editor: http://www.wjgnet.com/1948-5190/g_info_201007133856.htm

Book reviews: http://www.wjgnet.com/1948-5190/g_info_20100313161146.htm

Guidelines: http://www.wjgnet.com/1948-5190/g_info_20100313161315.htm

SUBMISSION OF THE REVISED MANUSCRIPTS AFTER ACCEPTED

Please revise your article according to the revision policies of *WJGE*. The revised version including manuscript and high-resolution image figures (if any) should be copied on a floppy or compact disk. The author should send the revised manuscript,

along with printed high-resolution color or black and white photos, copyright transfer letter, and responses to the reviewers by courier (such as EMS/DHL).

Editorial Office

World Journal of Gastrointestinal Endoscopy

Editorial Department: Room 903, Building D,

Ocean International Center,

No. 62 Dongsihuan Zhonglu,

Chaoyang District, Beijing 100025, China

E-mail: wjge@wjgnet.com

<http://www.wjgnet.com>

Telephone: +86-10-59080038

Fax: +86-10-85381893

Language evaluation

The language of a manuscript will be graded before it is sent for revision. (1) Grade A: priority publishing; (2) Grade B: minor language polishing; (3) Grade C: a great deal of language polishing needed; and (4) Grade D: rejected. Revised articles should reach Grade A or B.

Copyright assignment form

Please download a Copyright assignment form from http://www.wjgnet.com/1948-5190/g_info_201007134847.htm.

Responses to reviewers

Please revise your article according to the comments/suggestions provided by the reviewers. The format for responses to the reviewers' comments can be found at: http://www.wjgnet.com/1948-5190/g_info_201007134601.htm.

Proof of financial support

For paper supported by a foundation, authors should provide a copy of the document and serial number of the foundation.

Links to documents related to the manuscript

WJGE will be initiating a platform to promote dynamic interactions between the editors, peer reviewers, readers and authors. After a manuscript is published online, links to the PDF version of the submitted manuscript, the peer-reviewers' report and the revised manuscript will be put on-line. Readers can make comments on the peer reviewer's report, authors' responses to peer reviewers, and the revised manuscript. We hope that authors will benefit from this feedback and be able to revise the manuscript accordingly in a timely manner.

Science news releases

Authors of accepted manuscripts are suggested to write a science news item to promote their articles. The news will be released rapidly at EurekAlert/AAAS (<http://www.eurekalert.org>). The title for news items should be less than 90 characters; the summary should be less than 75 words; and main body less than 500 words. Science news items should be lawful, ethical, and strictly based on your original content with an attractive title and interesting pictures.

Publication fee

Authors of accepted articles must pay a publication fee. EDITORIAL, TOPIC HIGHLIGHTS, BOOK REVIEWS and LETTERS TO THE EDITOR are published free of charge.