

幽門狭窄を伴う根治切除不能進行胃癌に対する内視鏡下胃十二指腸ステント留置術の有効性試験（多施設共同前向き観察研究）	口頭	木村豊、瀧口修司、宮崎安弘、竹野淳、遠藤俊治、西川和宏、高地耕、川田純司、川端良平、藤田淳也、田村茂行、森正樹、土岐祐一郎	第87回日本消化器内視鏡学会総会	2014年5月 於：福岡市	国内
胃癌による胃切除術患者に対する成分栄養剤早期介入の有効性に関する多施設共同無作為化比較試験	口頭	木村豊、今村博司、西川和宏、岸健太郎、井上健太郎、松山仁、赤丸祐介、田村茂行、川瀬朋乃、川田純司、藤原義之、福井淳一、川端良平、下川敏雄	日本外科代謝栄養学会第51回学術集会	2014年7月 於：豊中市	国内
2人体制でおこなう胃癌手術における膈上緑リンパ節郭清のポイント	ビデオプレゼンテーション	西田靖仙、細川正夫	第76回日本臨床外科学会総会	2014年11月 於：郡山市	国内
腹腔鏡下胃切除術の手術成績と予後。	口演	山口和也、久野真史、市川賢吾、棚橋利行、八幡和憲、今井寿、佐々木義之、田中善宏、奥村直樹、松橋延壽、野中健一、高橋孝夫、長田真二、吉田和弘	第86回日本胃癌学会総会	2014年3月：横浜市	国内
当科におけるCY1P0胃癌症例の検討。	口演	奥村直樹、山口和也、棚橋利行、八幡和憲、今井寿、佐々木義之、田中善宏、松橋延壽、高橋孝夫、長田真二、吉田和弘	第86回日本胃癌学会総会	2014年3月：横浜市	国内
80歳以上高齢者における胃癌術後補助化学療法の検討。	口演	棚橋利行、山口和也、八幡和憲、奥村直樹、松橋延壽、吉田和弘	第86回日本胃癌学会総会	2014年3月：横浜市	国内
高度進行胃癌に対するDCS療法の治療成績。	口演	奥村直樹、山口和也、棚橋利行、八幡和憲、今井寿、佐々木義之、田中善宏、松橋延壽、高橋孝夫、長田真二、吉田和弘	第114回日本外科学会定期学術集会	2014年4月：京都市	国内

80歳以上高齢者における胃癌手術症例に対する術後補助化学療法 の現状。	口演	棚橋利行、山口和也、久野真史、市川賢吾、八幡和憲、今井寿、佐々木義之、田中善宏、奥村直樹、松橋延壽、野中健一、高橋孝夫、長田真二、吉田和弘、笹子三津留	第114回日本外科学会定期学術集会	2014年4月：京都市	国内
腹腔鏡下胃切除術におけるリンパ節郭清の術野展開のポイント。	口演	山口和也、棚橋利行、今井寿、佐々木義之、田中善宏、奥村直樹、松橋延壽、高橋孝夫、長田真二、吉田和弘	第69回日本消化器外科学会総会	2014年7月：福島市	国内
進行再発胃癌におけるHER2検査の検討。	口演	奥村直樹、山口和也、棚橋利行、八幡和憲、今井寿、佐々木義之、田中善宏、松橋延壽、長田真二、吉田和弘	第69回日本消化器外科学会総会	2014年7月：福島市	国内
StageIV胃癌におけるDCS療法の治療成績～1次治療でのDoubletとTripletの比較検討～。	口演	奥村直樹、山口和也、棚橋利行、市川賢吾、八幡和憲、山田敦子、今井寿、佐々木義之、田中善宏、松橋延壽、高橋孝夫、長田真二、吉田和弘	第52回日本癌治療学会学術集会	2014年8月：横浜市	国内
腹腔鏡下幽門側胃切除術における術野展開の定型化。	口演	山口和也、田中秀治、深田真宏、兼松昌子、山田敦子、棚橋利行、松井聡、今井寿、佐々木義之、田中善宏、奥村直樹、松橋延壽、高橋孝夫、長田真二、吉田和弘	第27回日本内視鏡外科学会総会	2014年10月：盛岡市	国内
80歳以上高齢者早期胃癌患者に対する腹腔鏡下胃切除術の短期成績。	口演	棚橋利行、山口和也、奥村直樹、田中秀治、深田真宏、兼松昌子、山田敦子、松井聡、今井寿、佐々木義之、田中善宏、松橋延壽、高橋孝夫、長田真二、吉田和弘	第27回日本内視鏡外科学会総会	2014年10月：盛岡市	国内
超音波凝固切開創地を中心とした開腹・鏡視下胃切除術の術野展開。	口演	山口和也、田中秀治、深田真宏、兼松昌子、山田敦子、棚橋利行、松井聡、今井寿、佐々木義之、田中善宏、奥村直樹、松橋延壽、高橋孝夫、長田真二、吉田和弘	第76回日本臨床外科学会総会	2014年11月：郡山市	国内

StageIV胃癌に対する化学療法奏効後外科的治療の検討.	口演	棚橋利行、山口和也、田中秀治、深田真治、兼松昌子、山田敦子、松井聡、今井寿、佐々木義之、田中善宏、奥村直樹、松橋延壽、高橋孝夫、長田真二、吉田和弘	第76回日本臨床外科学会総会	2014年11月：郡山市	国内
当科における腹腔鏡下胃全摘術後の再建法.	口演	奥村直樹、山口和也、棚橋利行、田中秀治、深田真宏、山田敦子、松井聡、佐々木義之、田中善宏、松橋延壽、高橋孝夫、長田真二、吉田和弘	第76回日本臨床外科学会総会	2014年11月：郡山市	国内
胃全摘術	ディベート	布部創也、比企直樹、大橋学、峯真司、渡邊雅之、齋浦明夫、上野雅資、佐野武、山口俊晴	第114回日本外科学会定期学術集会	2014年4月 於：京都	国内
大動脈リンパ節転移を伴う高度進行胃癌に対する術前化学療法+外科切除治療の検討	ポスターセッション	高津有紀子、比企直樹、布部創也、大橋学、佐野武、山口俊晴	第114回日本外科学会定期学術集会	2014年4月 於：京都	国内
食道胃接合部腺癌と下部食道扁平上皮癌の縦隔リンパ節転移に関する検討	ワークショップ	峯真司、渡邊雅之、佐野武、比企直樹、布部創也、大橋学、福長洋介、小西毅、有田淳一、秋吉高志	第69回日本消化器外科学会総会	2014年7月 於：郡山	国内
大動脈リンパ節転移を伴う高度進行胃癌に対する術前化学療法+外科切除治療の検討	ワークショップ	高津有紀子、比企直樹、布部創也、大橋学、佐野武、山口敏晴	第69回日本消化器外科学会総会	2014年7月 於：郡山	国内
進行胃癌に対する腹腔鏡下胃切除術の短期、長期成績と適応拡大への取り組み	シンポジウム	大橋学、比企直樹、布部創也、入野誠之、本多通孝、古川陽菜、山口俊晴	第27回日本内視鏡外科学会総会	2014年10月 於：盛岡	国内

<p>洗浄細胞診陽性胃癌に対する術前化学療法の意義</p>	<p>特別演題・ワークショップ</p>	<p>神谷諭、比企直樹、入野誠之、本多通孝、井田智、辻浦誠浩、速水克、松田達雄、古川陽菜、布部創也、大橋学、佐野武、山口俊晴</p>	<p>第76回日本臨床外科学会総会</p>	<p>2014年11月 於：郡山</p>	<p>国内</p>
<p>ロボット支援下胃切除術における郭清と体腔内手縫い吻合</p>	<p>口頭</p>	<p>西崎正彦 黒田新士 菊地覚次 桑田和也 香川俊輔 白川靖博 藤原俊義</p>	<p>第27回日本内視鏡外科学会総会</p>	<p>2014年10月 於：盛岡市</p>	<p>国内</p>

2. 学会誌・雑誌等における論文掲載

掲載した論文（発表題目）	発表者氏名	発表した場所 (学会誌・雑誌等 名)	巻号・ページ・年	国内・外 の別
The 10-year follow-up results of a randomised controlled trial comparing left thoracoabdominal and abdominal-transhiatal approaches to total gastrectomy for adenocarcinoma of the oesophagogastric junction or gastric cardia	Kurokawa Y, Sasako M, Sano T, Yoshikawa T, Iwasaki Y, Nashimoto A, Ito S, Kurita A, Mizusawa J, Nakamura K.	<i>Br J Surg</i>	in press	国外
Mediastinal lymph node metastasis and recurrence in adenocarcinoma of the esophagogastric junction.	Kurokawa Y, Hiki N, Yoshikawa T, Kishi K, Ito Y, Ohi M, Wada N, Takiguchi S, Mine S, Hasegawa S, Matsuda T, Takeuchi H.	<i>Surgery</i>	in press	国内
Multicenter large-scale study of prognostic impact of HER2 expression in patients with resectable gastric cancer.	Kurokawa Y, Matsuura N, Kimura Y, Adachi S, Fujita J, Imamura H, Kobayashi K, Yokoyama Y, Shaker MN, Takiguchi S, Mori M, Doki Y.	<i>Gastric Cancer</i>	in press	国内
Prognostic Impact of Major Receptor Tyrosine Kinase Expression in Gastric Cancer.	Kurokawa Y, Matsuura N, Kawabata R, Nishikawa K, Ebisui C, Yokoyama Y, Shaker MN, Hamakawa T, Takahashi T, Takiguchi S, Mori M, Doki Y.	<i>Ann Surg Oncol</i>	21:S584-90, 2014	国外
Validity of response assessment criteria in neoadjuvant chemotherapy for gastric cancer (JCOG0507-A).	Kurokawa Y, Shibata T, Sasako M, Sano T, Tsuburaya A, Iwasaki Y, Fukuda H.	<i>Gastric Cancer</i>	17:514-21, 2014	国内
Phase II study of trastuzumab in combination with S-1 plus cisplatin in HER2-positive gastric cancer (HERBIS-1).	Kurokawa Y, Sugimoto N, Miwa H, Tsuda M, Nishina S, Okuda H, Imamura H, Gamoh M, Sakai D, Shimokawa T, Komatsu Y, Doki Y, Tsujinaka T, Furukawa H.	<i>Br J Cancer</i>	110:1163-8, 2014	国外

Three-year outcomes of a phase II study of adjuvant chemotherapy with S-1 plus docetaxel for stage III gastric cancer after curative D2 gastrectomy.	Fujitani K, Tamura S, Kimura Y, Tsuji T, Matsuyama J, Iijima S, Imamura H, Inoue K, Kobayashi K, Kurokawa Y, Shimokawa T, Tsujinaka T, Furukawa H.	<i>Gastric Cancer</i>	17:348-53, 2014	国内
Phase II study of S-1 monotherapy in patients over 75 years of age with advanced gastric cancer (OGSG0404).	Imamura H, Kishimoto T, Takiuchi H, Kimura Y, Morimoto T, Imano M, Iijima S, Yamashita K, Maruyama K, Otsuji T, Kurokawa Y, Furukawa H.	<i>J Chemother</i>	26 (1) :57-61, 2014	国内
Survival analysis of adjuvant chemotherapy with S-1 plus cisplatin for stage III gastric cancer.	Takahari D, Hamaguchi T, Yoshimura K, Katai H, Ito S, Fuse N, Konishi M, Yasui H, Terashima M, Goto M, Tanigawa N, Shirao K, Sano T, Sasako M	<i>Gastric Cancer</i>	17 - 383-386 - 2014	国外
The earlier the better?	Terashima M	<i>Gastric Cancer</i>	17 - 197-199 - 2014	国外
Influence of endoscopic submucosal dissection on additional gastric resections.	Kawata N, Kakushima N, Tokunaga M, Tanaka M, Sawai H, Takizawa K, Imai K, Hotta K, Yamaguchi Y, Matsubayashi H, Tanizawa Y, Bando E, Kawamura T, Terashima M, Ono H	<i>Gastric Cancer</i>	Epub 2014	国外
Postgastrectomy Syndrome Assessment Scale (PGSAS)-45 and changes in body weight are useful tools for evaluation of reconstruction methods following distal gastrectomy.	Terashima M, Tanabe K, Yoshida M, Kawahira H, Inada T, Okabe H, Urushihara T, Kawashima Y, Fukushima N, Nakada K	<i>Annals of Surgical Oncology</i>	21 - 370-378 - 2014	国外
Early phase II study of robot-assisted distal gastrectomy with nodal dissection for clinical stage IA gastric cancer.	Tokunaga M 徳永正則, Sugisawa N, Kondo J, Tanizawa Y 谷澤豊, Bando E 坂東悦郎, Kawamura T 川村泰一, Terashima	<i>Gastric Cancer</i>	17 - 542-547 - 2014	国外

ロボット支援手術の安全性を評価する臨床第II相試験	寺島雅典、徳永正則、谷澤豊、坂東悦郎、川村泰一、幕内梨恵、三木友一朗、絹笠祐介、上坂克彦	癌の臨床	60・335-339・2014	国内
Perioperative risk assessment for gastrectomy by surgical apgar score.	Miki Y, Tokunaga M, Tanizawa Y, Bando E, Kawamura T, Terashima M	<i>Annals of Surgical Oncology</i>	21・2601-2607・2014	国外
A novel splice variant of XIAP-associated factor 1 (XAF1) is expressed in peripheral blood containing gastric cancer-derived circulating tumor cells.	Hatakeyama K, Yamakawa Y, Fukuda Y, Ohshima K, Wakabayashi-Nakao K, Sakura N, Tanizawa Y, Kinugasa Y, Yamaguchi K, Terashima M, Mochizuki T	<i>Gastric Cancer</i>	Epub 2014	国外
胃癌に対するロボット手術の現状	寺島雅典、徳永正則、谷澤豊、坂東悦郎、川村泰一、幕内梨恵、三木友一朗、絹笠祐介、上坂克彦	臨床外科	69・1382-1388・2014	国内
Efficacy and long-term outcome of pre-emptive endoscopic resection and surgery for multiple synchronous gastric cancers.	Yoshida M, Kakushima N, Tokunaga M, Tanaka M, Takizawa K, Imai K, Hotta K, Matsubayashi H, Tanizawa Y, Bando E, Kawamura T, Terashima M, Ono H	<i>Surgical Endoscopy</i>	Epub 2014	国外
Characteristics and clinical relevance of postgastrectomy syndrome assessment scale (PGSAS)-45: newly developed integrated questionnaires for assessment of living status and quality of life in postgastrectomy patients.	Nakada K, Ikeda M, Takahashi M, Kinami S, Yoshida M, Uenosono Y, Kawashima Y, Oshio A, Suzukamo Y, Terashima M, Kodera Y.	<i>Gastric Cancer</i>	18・147-158・2015	国外
Clinical significance of a papillary adenocarcinoma component in early gastric cancer: a single-center retrospective analysis of 628 surgically resected early gastric cancers.	Sekiguchi M, Kushima R, Oda I, Suzuki H, Taniguchi H, Sekine S, Fukagawa T, Katai H.	<i>J Gastroenterol.</i>	2014 Aug 21. [Epub ahead of print]	国外

Development and external validation of a nomogram for overall survival after curative resection in serosa-negative, locally advanced gastric cancer.	Hirabayashi S, Kosugi S, Isobe Y, Nashimoto A, Oda I, Hayashi K, Miyashiro I, Tsujitani S, Kodera Y, Seto Y, Furukawa H, Ono H, Tanabe S, Kaminishi M, Nunobe S, Fukagawa T, Matsuo R, Nagai T, Katai H, Wakai T, Akazawa K.	<i>Ann Oncol.</i>	2014 Jun;25(6):1179	国外
画像による胃癌のリンパ節転移診断	深川剛生	<i>臨床外科</i>	69(13):1438-1442(2014)	国内
残胃のがん	深川剛生、大橋真記真記、森田信司、片井均	<i>胃外科のすべて</i>	242-251、(2014.3)	国内
Efficacy of endoscopic gastroduodenal stenting for gastric outlet obstruction due to unresectable advanced gastric cancer: A prospective multicenter study	Endo S, Takiguchi S, Miyazaki Y, Nishikawa K, Kawabata R, Takachi K, Kimura Y, Takeno A, Tamura S, Mori M, Doki Y	<i>J Surg Oncol</i>	109(3) - 208-212 - 2014	国外
Phase II study of S-1 monotherapy in patients over 75 years of age with advanced gastric cancer (OGSG0404)	Imamura H, Kishimoto T, Takiuchi H, Kimura Y, Morimoto T, Imano M, Iijima S, Yamashita K, Maruyama K, Otsuji T, Kurokawa Y, Furukawa H	<i>J Chemo</i>	26(1) - 57-61 - 2014	国外
Accuracy of CT staging of locally advanced gastric cancer after neoadjuvant chemotherapy - cohort evaluation within the randomized phase II study	Toshikawa T, Tanabe K, Nishikawa K, Ito Y, Matsui T, Kimura Y, Hirabayashi N, Mikata S, Iwahashi M, Fukushima R, Takiguchi N, Miyashiro I, Morita S, Miyashita Y, Tsuburaya A, Sakamoto STanabe K, Nishikawa K, Ito Y, Matsui T, Kimura Y, Hasegawa S, Aoyama T, Hayashi T, Morita S, Miyashita Y	<i>Ann Surg Oncol</i>	21(Suppl 3) - S385-S389 - 2014	国外
Incidence of metachronous gastric cancer in the remnant stomach after synchronous multiple cancer surgery.	Nozaki I, Hato S, Kobatake T, Ohta K, Kubo Y, Nishimura R, Kurita A.	<i>Gastric Cancer</i>	17: 61-66, 2014	国外



胃癌ESD後穿孔に対して腹腔鏡下修復術が有用であった1例	羽藤慎二, 浅野博昭, 井野英男, 内藤 稔	日外科系連会誌	39(4):697-702, 2014	国内
Addition of docetaxel to S-1 without platinum prolongs survival of patients with advanced gastric cancer: a randomized study (START).	Koizumi W, Kim YH, Fujii M, Kim HK, Imamura H, Lee KH, Hara T, Chung HC, Satoh T, Cho JY, Hosaka H, Tsuji A, Takagane A, Inokuchi M, Tanabe K, Okuno T, Ogura M, Yoshida K, Takeuchi M, Nakajima T/ The JACCRO and KCSG Study Group.	J Cancer Res Clin	140: 319-328 2014.	国外
Sequential paclitaxel followed by tegafur and uracil (UFT) or S-1 versus UFT or S-1 monotherapy as adjuvant chemotherapy for T4a/b gastric cancer (SAMIT): a phase 3 factorial randomised controlled trial.	Tsuburaya A, Yoshida K, Kobayashi M, Yoshino S, Takahashi M, Takiguchi N, Tanabe K, Takahashi N, Imamura H, Tatsumoto N, Hara H, Nishikawa K, Fukushima R, Nozaki I, Kojima H, Miyashita Y, Oba K, Buyse M, Morita M, Sakamoto J.	Lancet Oncology	15:886-93 2014	国外
胃癌-高齢者の胃癌治療-	奥村直樹、棚橋利行、山口和也、吉田和弘	消化器外科	37 ( 9 ) :1409-1415, 2014	国内
消化器系 縫合不全 (上部消化管)	田中善宏、吉田和弘	臨床外科	69(11):212-215, 2014	国内
胃に特有の術前術後ケア	棚橋利行、奥村直樹、山口和也、吉田和弘	消化器外科 NURSING	19(11):1081-1085, 2014	国内
D1+郭清を伴う自律神経温存幽門保存胃切除術-「起点」「受け」「底」の概念を用いて-	二宮基樹、丁田泰宏、金澤卓、三宅総一郎、戸嶋俊明、加藤卓也	手術	471-477, 2014	国内

<p>Development and external validation of a nomogram for overall survival after curative resection in serosa-negative, locally advanced gastric cancer.</p>	<p>Hirabayashi S, Kosugi S, Isobe Y, Nashimoto A, Oda I, Hayashi K, Miyashiro I, Tsujitani S, Kodera Y, Seto YO, Furukawa H, Ono H, Tanabe S, Kaminishi M, <u>Nunobe S</u>, Fukagawa T, Matsuo R, Nagai T, Katai H, Wakai T, Akazawa K.</p>	<p><i>Ann Oncol.</i></p>	<p>25(6):1179-84. 2014,</p>	<p>国外</p>
<p>Prognostic significance of complications after curative surgery for gastric cancer.</p>	<p>Kubota T, Hiki N, Sano T, Nomura S, <u>Nunobe S</u>, Kumagai K, Aikou S, Watanabe R, Kosuga T, Yamaguchi T.</p>	<p><i>Ann Surg Oncol.</i></p>	<p>21(3):891-8. 2014,</p>	<p>国外</p>
<p>Long-term Survival Outcomes of Advanced Gastric Cancer Patients Who Achieved a Pathological Complete Response with Neoadjuvant Chemotherapy: A Systematic Review of the Literature.</p>	<p>Cho H, Nakamura J, Asaumi Y, Yabusaki H, Sakon M, Takasu N, Kobayashi T, Aoki T, Shiraishi D, Kishimoto H, <u>Nunobe S</u>, Yanagisawa S, Suda T, Ueshima S, Matono S, Maruyama H, Tatsumi M, Seya T, Tanizawa Y, Yoshikawa T.</p>	<p><i>Ann Surg Oncol.</i></p>	<p>[Epub ahead of print], 2014 .</p>	<p>国外</p>

#### IV. 研究成果の刊行物・別刷

# Impact of reconstruction method on visceral fat change after distal gastrectomy: Results from a randomized controlled trial comparing Billroth I reconstruction and Roux-en-Y reconstruction

Koji Tanaka, MD,<sup>a</sup> Shuji Takiguchi, MD, PhD,<sup>a</sup> Isao Miyashiro, MD, PhD,<sup>b</sup> Motohiro Hirao, MD, PhD,<sup>c</sup> Kazuyoshi Yamamoto, MD, PhD,<sup>c</sup> Hiroshi Imamura, MD, PhD,<sup>d</sup> Masahiko Yano, MD, PhD,<sup>b</sup> Masaki Mori, MD, PhD,<sup>a</sup> Yuichiro Doki, MD, PhD,<sup>a</sup> and Osaka University Clinical Research Group for Gastroenterological Study,<sup>e</sup> Osaka, Japan

**Background.** Visceral fat is one of the causes of metabolic syndrome. Among the various types of bariatric surgery, duodenal-jejunal bypass is one of the most common procedures. However, the effect of duodenal bypass on fat changes is not completely understood. We examined the effect of duodenal bypass on visceral fat changes by comparing Billroth I (BI) and roux-en Y (RY) reconstruction in distal gastrectomy.

**Methods.** This retrospective study used data from 221 patients registered for a prospective randomized trial that compared BI to RY in distal gastrectomy with lymphadenectomy to treat gastric cancer. With a software package, we first quantified the visceral fat area (VFA) on cross-sectional computed tomography scans obtained at the level of the umbilicus before and 1 year after surgery, and then determined the impact of duodenal bypass on visceral fat changes.

**Results.** Clinicopathological background data did not differ between BI and RY. Rates of BMI reduction for BI and RY also did not differ. The VFA reduction rate for RY ( $47.2 \pm 25.5\%$ ) was greater than for BI ( $36.8 \pm 34.2\%$ ,  $P = .0104$ ). Adjuvant chemotherapy (chemotherapy versus no chemotherapy,  $P = .0136$ ), type of reconstruction (BI versus RY,  $P < .0001$ ), and pathologic stage ( $p$  stage I versus  $p$  stage II–IV,  $P = .0468$ ) correlated significantly with postoperative visceral fat loss. Multivariate logistic regression analysis identified reconstruction (BI versus RY,  $P = .0078$ ) as a significant determinant of visceral fat loss.

**Conclusion.** Visceral fat loss after distal gastrectomy was greater for RY than for BI, and duodenal bypass may be associated with reduction of visceral fat. (*Surgery* 2014;155:424-31.)

From the Department of Gastroenterological Surgery, Graduate School of Medicine,<sup>a</sup> Osaka University; Department of Surgery,<sup>b</sup> Osaka Medical Center for Cancer and Cardiovascular Diseases; Department of Surgery,<sup>c</sup> National Hospital Organization Osaka National Hospital; Department of Surgery,<sup>d</sup> Sakai Municipal Hospital; and Department of Gastroenterological Surgery,<sup>e</sup> Osaka University, 2-2 Yamadaoka, Suita city, Osaka, Japan

This study was registered with clinical trial identification number UMIN000000878.

Accepted for publication August 12, 2013.

Reprint requests: Shuji Takiguchi, MD, PhD, Department of Gastroenterological Surgery, Graduate School of Medicine, Osaka University, Osaka, Japan. E-mail: stakiguti@gesurg.med.osaka-u.ac.jp.

0039-6060/\$ - see front matter

© 2014 Mosby, Inc. All rights reserved.

<http://dx.doi.org/10.1016/j.surg.2013.08.008>

424 SURGERY

IT HAS BEEN PROVEN that gastric bypass surgery affects the release of gastrointestinal hormones<sup>1</sup> and induces malabsorption,<sup>2</sup> but there are no conclusive data about the effects of duodenal bypass on visceral fat changes. There are various types of bariatric procedures, including gastric banding, sleeve gastrectomy, roux-en Y bypass, biliopancreatic diversion with duodenal switch, and duodenal-jejunal bypass. However, to the best of our knowledge, no authors have evaluated fat reduction specifically caused by duodenal bypass because the size of the remnant stomach and the

length of the jejunal bypass differ among the various procedures.

The selection of the reconstruction method, either Billroth I (BI) or roux-en Y (RY), after distal or subtotal gastrectomy is still controversial. A large, multi-institutional, randomized controlled trial was conducted by the Osaka University Clinical Research Group for Gastroenterological Study (Japan)<sup>3,4</sup> to address this problem. The primary endpoint of this study was to compare body weight loss 1 year after surgery between the BI and RY groups. Secondary endpoints were postoperative complications, nutritional state, and quality of life. This trial gave us an opportunity to prospectively evaluate data about the effects of BI and RY on visceral and subcutaneous fat loss because patients whose remnant stomachs were large enough so that either technique could be performed were assigned randomly intraoperatively to undergo either BI and RY, and the reconstruction methods were prescribed by the protocol.

Visceral fat areas (VFAs) estimated from a single computed tomography (CT) scan at the level of the umbilicus are known to correlate with the total volume of visceral fat.<sup>5,6</sup> On the basis of this knowledge, a practical, standardized technique has been developed to determine the VFA from a single CT scan.<sup>7</sup>

In the present study, we used CT and a software package to quantify the VFA of patients before and 1 year after surgery. We then determined the impact of the type of reconstructive procedure on visceral fat changes in patients with gastric cancer who underwent distal gastrectomy with lymphadenectomy.

## METHODS

**Patients.** Between May 2004 and October 2009, a total of 332 patients with gastric cancer were registered in the original study. After completion of the informed consent process, patients were included in the study if they met the eligibility criteria.<sup>4</sup> After initial laparotomy, the location of the tumor was confirmed to be in the middle or lower third of the stomach and the proportion of residual stomach was regulated as one-third of the original stomach. The operator also checked the length of the residual stomach to confirm that either reconstruction procedure could be performed after distal gastrectomy. The surgeon confirmed the eligibility and exclusion criteria immediately after the initial laparotomy, and patients were then randomized intraoperatively to either the BI group or the RY group. Randomization was performed by the

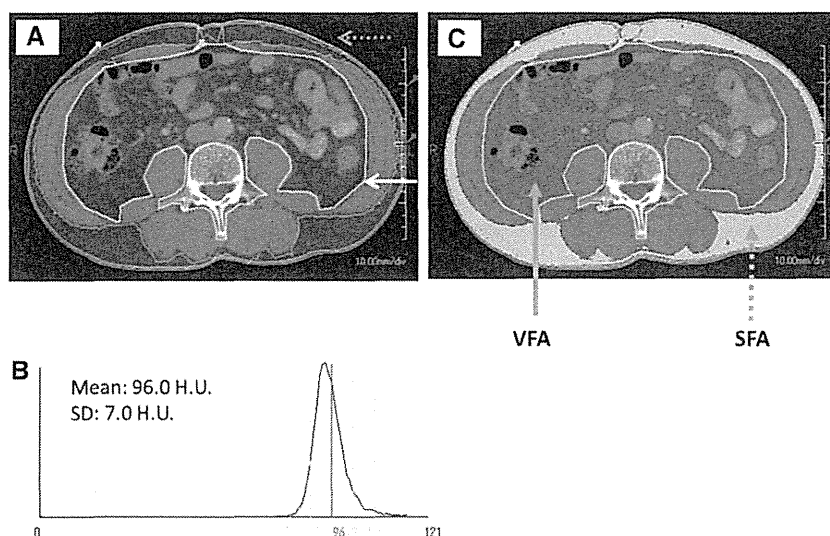
minimization method according to the patient's body mass index (BMI) (<25 or  $\geq 25$  kg/m<sup>2</sup>) and institution.

To evaluate visceral fat changes, we collected CT scans both before and 1 year after surgery. A total of 221 patients, whose CT scans at the umbilicus level both before and 1 year after surgery could be obtained, were retrospectively analyzed in this study. Information about the patients' backgrounds and clinicopathological data were extracted from the data collected by the original study. This study was approved by the institutional review boards of all participating hospitals and was conducted in accordance with the Declaration of Helsinki.

**Operative procedure.** Patients underwent gastrectomy with systematic lymphadenectomy at 18 high-volume institutions in Osaka, Japan. All 18 institutions were participants in the surgical study group "Osaka University Clinical Research Group for Gastroenterological Study." Overall, more than 50 gastrectomies were performed each year in these 18 hospitals. All operations were performed or supervised by senior surgeons who were members of the Japanese Gastric Cancer Association. During the planning of the study, all participating surgeons reached an agreement concerning the technical details of the reconstructive procedures.

Endotracheal general anesthesia and standard laparotomy or laparoscopic operations were used for all patients in each institution. Gastric tumors located in the lower or middle third of the stomach were treated with distal gastrectomy. Lymphadenectomy approaches were categorized as D1–3, as defined by the Japanese Classification for Standard Dissection.<sup>8</sup> D1 involves dissecting the paragastric nodes, whereas D2 also includes dissection of the nodes along the left gastric, common hepatic, and celiac arteries. D3 includes the nodes dissected in D1 and D2, as well as dissection of the hepatoduodenal and retropancreatic nodes, the nodes along the superior mesenteric vein, and the para-aortic nodes between the level of the celiac axis and the inferior mesenteric artery.

For BI reconstruction, the duodenum and remnant stomach were sutured. For RY reconstruction, the jejunum was divided 20 cm distal to the ligament of Treitz, and the portion of the jejunum closest to the patient's head was closed, followed by the remaining gastric pouch, which was anastomosed to the jejunum. The oral portion of the jejunum was then anastomosed to the mid-jejunum 30 cm distal to the gastrojejunostomy.



**Fig 1.** Illustration of method used to determine abdominal fat distribution on a CT scan obtained at the umbilicus level. (A) White line (*solid arrow*) outlines the intraperitoneal area. Gray line (*dotted arrow*), drawn with a cursor automatically or manually, outlines the subcutaneous fat layer, in which attenuation is measured. (B) Histogram of the CT numbers (in Hounsfield units) in the lesion outlined in (A) (mean  $\pm$  2 SD). (C) Region defined as visceral fat tissue (*solid arrow*). Total fat area was calculated from the region outlining the circumference of the abdominal wall. The VFA was subtracted, and the remainder was regarded as the SFA (*dotted arrow*).

The basic anastomotic procedures, such as sutures made by hand or machine and standard laparotomy or laparoscopic operations, were not prescribed in detail by the protocol.

Of the 118 patients in the RY group, gastrojejunostomy was performed by hand in 8 patients, by circular stapler in 82 patients, and by linear stapler in 28 patients. The Roux-en-Y limb was ascended through the retrocolic route in 71 patients and the antecolic route in 47 patients.

**Quantification of VFAs and subcutaneous fat areas (SFAs).** The VFA was measured with "FatScan," which was described previously,<sup>7</sup> on one cross-sectional CT scan obtained at the level of the umbilicus. Figure 1 illustrates the method used to determine the fat tissue area on a CT scan. First, the intraperitoneal area was defined by tracing its contour manually on the scan. Thereafter, a region of interest on the subcutaneous fat layer was defined by tracing its contour on each scan either automatically or manually; then, the attenuation range of the CT numbers (in Hounsfield units) for fat tissue was calculated (Fig 1, A). A histogram for fat tissue was computed based on the mean attenuation  $\pm$  2 SD (Fig 1, B). Within the region outlined in Fig 1, A, the tissue with attenuation within the mean  $\pm$  2 SD was considered to be the VFA. Pixels with attenuation values in the selected attenuation range are depicted. The total fat area was calculated in

the region outlining the circumference of the abdominal wall. The VFA (solid arrow) was subtracted, and the remainder was regarded as the SFA (dotted arrow) (Fig. 1, C).

**Statistical analysis.** Differences between groups were examined for statistical significance with the Student *t*-test with Yates' correction,  $\chi^2$  test, Fisher's exact probability test, or Wilcoxon rank-sum test. Statistical analysis was performed with JMP version 9.0 (SAS Institute, Cary, NC). Univariate analysis was performed to identify the factors associated with visceral fat loss. The identified variables were subsequently entered into multivariate analysis, and logistic regression analysis was used to identify independent factors that influence visceral fat loss.

## RESULTS

**Comparison of characteristics of patients who underwent BI or RY.** Table I compares the background characteristics of patients who underwent BI or RY. Age, sex, preoperative BMI, preoperative VFA, preoperative SFA, preoperative serum albumin levels, preoperative lymphocyte counts, preoperative prognostic nutritional index values,<sup>9</sup> operative approach, and lymphadenectomy were not significantly different between the two groups. With regard to operative factors, such as operative approach (ie, the proportion of patients who underwent laparoscopy versus

**Table I.** Patient demographics, tumor characteristics, and operative details

	BI group, n = 103	RY group, n = 118	P value
Age, y*	64.1 ± 9.2	64.1 ± 10.3	.9765
Men/women	65/38	85/33	.1563
Preoperative BMI,* kg/m <sup>2</sup>	22.4 ± 3.2	22.7 ± 3.0	.3846
Preoperative total fat area, cm <sup>2*</sup>	204.0 ± 73.9	215.6 ± 90.5	.3023
Preoperative VFA, cm <sup>2*</sup>	83.9 ± 38.9	92.6 ± 43.6	.1175
Preoperative SFA, cm <sup>2*</sup>	120.1 ± 54.8	122.9 ± 62.5	.7243
Preoperative serum albumin, mg/dL	4.12 ± 0.39	4.12 ± 0.51	.9864
Preoperative lymphocyte count	1,846 ± 699	1,924 ± 596	.3966
Preoperative PNI*	50.6 ± 5.7	51.2 ± 6.1	.4862
Operative approach (laparoscopy/laparotomy)	24/79	28/90	.9404
Lymphadenectomy (D1/D2+D3)	38/57	46/62	.9503
Adjuvant chemotherapy (yes/no)	13/90	16/102	.8368
Recurrence (yes/no)	5/98	4/114	.7372

\*Data are mean ± SD. Comparisons between BI and RY groups with the Student *t* test. Other parameters were compared with  $\chi^2$  or Fisher exact test.

BI, Billroth I reconstruction; BMI, body mass index; PNI, prognostic nutritional index; RY, roux-en Y reconstruction; SFA, subcutaneous fat area; VFA, visceral fat area.

laparotomy) and field of lymphadenectomy, there were no significant differences between the groups. There were also no significant differences between the groups with regard to adjuvant chemotherapy and cancer recurrence. Information about the composition of food consumed after surgery was collected by questionnaire. Most of the patients who underwent BI (90.9%) and RY (86.9%) consumed a normal diet, whereas 9.1% of BI and 13.1% of RY patients consumed a soft or liquid diet ( $P = .3824$ ). The mean intervals for when the follow-up CT was performed after surgery were  $376 \pm 111$  days for BI and  $374 \pm 77$  days for RY ( $P = .9980$ ).

**Comparison of postoperative nutritional states of patients who underwent BI or RY.** Table II lists comparative data for BMI, VFA, and SFA. Postoperative BMI, postoperative SFA, postoperative serum albumin levels, postoperative lymphocyte counts, postoperative prognostic nutritional index values, and the rate of reduction of BMI ( $\Delta$ BMI%) were not substantially different between the BI

**Table II.** Comparison of postoperative nutritional status of patients in the BI and RY groups

	BI group, n = 103	RY group, n = 118	P value
Postoperative BMI, kg/m <sup>2</sup>	20.3 ± 2.8	20.5 ± 2.4	.6106*
Postoperative total fat area, cm <sup>2</sup>	139.3 ± 63.2	127.2 ± 61.2	.1497*
Postoperative VFA, cm <sup>2</sup>	50.0 ± 27.3	43.9 ± 22.2	.0821*
Postoperative SFA, cm <sup>2</sup>	89.7 ± 46.6	83.4 ± 47.8	.3239*
Postoperative serum albumin, mg/dL	4.21 ± 0.34	4.18 ± 0.42	.5789
Postoperative lymphocyte count	1,891 ± 625	1,908 ± 575	.8429
Postoperative PNI*	51.2 ± 5.1	51.3 ± 5.6	.8379
$\Delta$ BMI%	8.9 ± 6.6	9.5 ± 7.1	.2634†
$\Delta$ Total fat area %	29.6 ± 25.8	37.0 ± 25.4	.0117†
$\Delta$ VFA%	36.8 ± 34.2	47.2 ± 25.5	.0032†
$\Delta$ SFA%	22.2 ± 28.4	27.3 ± 32.8	.0732†

\*Student *t* test.

†Wilcoxon rank-sum test.

Data are mean ± SD.

BI, Billroth I reconstruction; BMI, body mass index; PNI, prognostic nutritional index;  $\Delta$  Total fat area %, rate of reduction of total fat area;  $\Delta$ BMI%, rate of reduction of BMI;  $\Delta$ SFA%, rate of reduction of SFA;  $\Delta$ VFA%, rate of reduction of VFA; RY, roux-en Y reconstruction; SFA, subcutaneous fat area; VFA, visceral fat area.

and RY groups. The postoperative VFA of the RY group ( $43.9 \pm 22.2$  cm<sup>2</sup>) was smaller than that of the BI group ( $50.0 \pm 27.3$  cm<sup>2</sup>), but the difference was not clinically important ( $P = .0821$ ). The rate of reduction of the VFA ( $\Delta$ VFA%) in the RY group ( $47.2 \pm 25.5\%$ ) was greater than in the BI group ( $36.8 \pm 34.2\%$ ;  $P = .0032$ ). For the muscle reduction rate, there was no difference between the BI ( $2.5 \pm 18.1\%$ ) and RY ( $3.1 \pm 16.8\%$ ;  $P = .7970$ ) groups. Figure 2 shows the correlation between preoperative BMI and  $\Delta$ VFA% according to the reconstruction method used. The  $\Delta$ VFA% for the RY group was greater in patients with greater BMI than in patients with lesser BMI. In contrast, the  $\Delta$ VFA% for the BI group was similar between patients with greater and lesser BMI. Patients were divided into two BMI groups according to the median of the preoperative BMI ( $22.5$  kg/m<sup>2</sup>). Tables III and IV show postoperative data for the BMI  $\geq 22.5$  kg/m<sup>2</sup> and the BMI  $< 22.5$  kg/m<sup>2</sup> groups. In the BMI  $\geq 22.5$  kg/m<sup>2</sup> group, preoperative BMI, postoperative BMI, and  $\Delta$ BMI% were not different between the BI and RY groups. The postoperative VFA of the RY group ( $50.4 \pm 21.9$  cm<sup>2</sup>) was less than that of the BI group ( $61.9 \pm 30.0$  cm<sup>2</sup>;  $P = .0218$ ). The  $\Delta$ VFA% of the RY group ( $52.1 \pm 19.5\%$ ) was also greater than that of the BI group ( $35.4 \pm 42.9\%$ ;

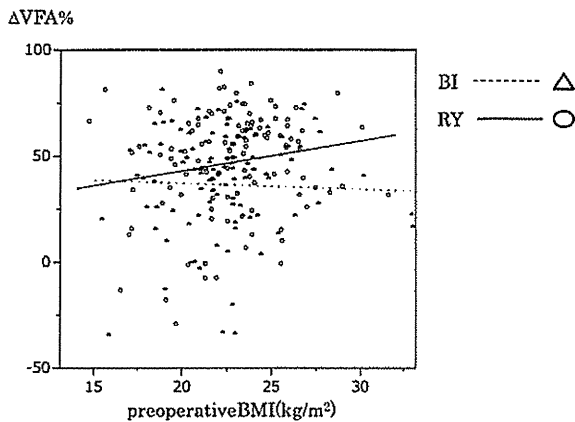


Fig 2. Comparison between BI and RY groups of changes in visceral fat according to preoperative BMI.

Table III. Postoperative BMI and fat areas of the BMI  $\geq 22.5$  kg/m<sup>2</sup> group

High BMI group	BI group, n = 47	RY group, n = 62	P value
Preoperative BMI, kg/m <sup>2</sup>	24.9 ± 2.5	24.9 ± 1.9	.9607*
Postoperative BMI, kg/m <sup>2</sup>	21.4 ± 2.9	21.3 ± 2.4	.7931*
ΔBMI%	10.2 ± 6.7	12.2 ± 5.6	.1012*
Preoperative VFA, cm <sup>2</sup>	104.7 ± 40.7	112.3 ± 42.0	.3461*
Postoperative VFA, cm <sup>2</sup>	61.9 ± 30.0	50.4 ± 21.9	.0218*
ΔVFA%	35.4 ± 42.9	52.1 ± 19.5	.0041†
Preoperative SFA, cm <sup>2</sup>	143.0 ± 59.2	152.0 ± 63.2	.4510*
Postoperative SFA, cm <sup>2</sup>	110.2 ± 50.9	104.5 ± 51.7	.5663*
ΔSFA%	20.0 ± 28.6	29.6 ± 21.9	.0930†

\*Student t-test.

†Wilcoxon rank-sum test.

Data are mean ± SD.

BI, Billroth I reconstruction; BMI, body mass index; ΔBMI%, rate of reduction of BMI; ΔSFA%, rate of reduction of SFA; ΔVFA%, rate of reduction of VFA; RY, roux-en Y reconstruction; SFA, subcutaneous fat area; VFA, visceral fat area.

$P = .0041$ ). In the BMI  $< 22.5$  kg/m<sup>2</sup> group, there were no significant differences between the BI and RY groups in terms of postoperative BMI, ΔBMI%, postoperative VFA, and ΔVFA%.

**Determinants of postoperative visceral fat loss.**

Before the factors associated with visceral fat loss were analyzed, the study population was divided into a high ΔVFA% group and a low ΔVFA% group by the median ΔVFA% (48.5%). Univariate analysis was used to identify significant factors associated with visceral fat loss. As shown in Table V, among the

Table IV. Postoperative BMI and fat area of the BMI  $< 22.5$  kg/m<sup>2</sup> group

Low BMI group	BI group, n = 56	RY group, n = 56	P value
Preoperative BMI (kg/m <sup>2</sup> )	20.2 ± 1.8	20.2 ± 2.0	.8691*
Postoperative BMI (kg/m <sup>2</sup> )	19.4 ± 2.3	19.3 ± 1.8	.7537*
ΔBMI%	7.7 ± 6.2	6.4 ± 7.4	.2949†
Preoperative VFA (cm <sup>2</sup> )	66.4 ± 27.1	70.9 ± 34.1	.4384*
Postoperative VFA (cm <sup>2</sup> )	39.4 ± 19.8	36.7 ± 20.4	.4720*
ΔVFA%	38.0 ± 25.2	41.7 ± 30.1	.4729†
Preoperative SFA (cm <sup>2</sup> )	100.9 ± 42.6	90.7 ± 43.1	.2117*
Postoperative SFA (cm <sup>2</sup> )	72.4 ± 34.6	60.0 ± 28.9	.0410*
ΔSFA%	24.0 ± 28.2	24.7 ± 41.7	.3594†

\*Student t-test.

†Wilcoxon rank-sum test.

Data are mean ± SD.

BI, Billroth I reconstruction; BMI, body mass index; ΔBMI%, rate of reduction of BMI; ΔSFA%, rate of reduction of SFA; ΔVFA%, rate of reduction of VFA; RY, roux-en Y reconstruction; SFA, subcutaneous fat area; VFA, visceral fat area.

clinicopathologic factors that we examined, adjuvant chemotherapy (performed versus not performed,  $P = .0046$ ), type of reconstruction (BI versus RY,  $P = .0087$ ), and p stage (p stage I versus p stage II–IV,  $P = .0468$ ) correlated with postoperative visceral fat loss. No deaths occurred during the course of this study. There was no significant difference in morbidity/postoperative complications between the low (10/111; 9.0%) and high (11/110; 10%) VFA groups ( $P = .8017$ ) when patients were divided by the median of the preoperative VFA value. Multivariate logistic regression analysis that included the above factors identified reconstruction (BI versus RY,  $P = .0078$ ) and adjuvant chemotherapy (performed versus not performed,  $P = .0172$ ) as significant predictors of visceral fat loss.

**DISCUSSION**

Gastrectomy usually leads to body weight loss. The mechanisms of postgastrectomy weight loss include impaired food intake and malabsorption.<sup>10-12</sup> In previous studies authors reported that body weight loss is mainly caused by loss of body fat.<sup>13,14</sup> With respect to anatomical localization, body fat is divided into subcutaneous fat and visceral fat. To our knowledge, there is little information on the changes that take place in visceral and subcutaneous fat after gastrectomy. We found that visceral fat loss after distal gastrectomy was greater in patients who underwent



**Table V.** Univariate and multivariate analysis of risk factors for visceral fat loss

<i>Factors</i>	<i>High/low</i>	<i>Univariate P value</i>	<i>Odds ratio</i>	<i>95% CI</i>	<i>Multivariate P value</i>
Reconstruction		.0087	2.0965	1.2142–3.6573	.0078
RY	69/49				
BI	42/61				
Sex		.6323			
Men	77/73				
Women	34/37				
Lymphadenectomy		.3324			
D2 or D3	64/71				
D1	46/39				
Operative approach		.2182			
Laparotomy	81/88				
Laparoscopy	30/22				
Adjuvant chemotherapy		.0046	4.6106	1.3056–17.9177	.0172
Yes	22/7				
No	89/103				
Recurrence		1.0000			
Yes	5/4				
No	105/106				
Location of tumor		.6364			
M	37/40				
L	74/70				
Age, y		.1582			
≥65	64/53				
<65	47/57				
Pathologic stage		.0468	1.26856	0.4556–3.7129	.6501
II or III	28/16				
I	83/94				
Postoperative complications		.1656			
Yes	10/4				
No	101/106				

BI, Billroth I reconstruction; L, lower third of stomach; M, middle third of stomach; RY, roux-en Y reconstruction.

RY compared with those who underwent BI. In a previous study investigators reported that visceral fat reduction is greater after RY gastric bypass compared to vertical banded gastroplasty.<sup>14</sup> Our results are comparable with other reports in the field of bariatric surgery.

However, in previous reports there were differences, such as the size of the remnant stomach and the length of the jejunal bypass, between the operative procedures. To the best of our knowledge, this was the first study to focus on the specific impact of duodenal bypass on visceral fat loss. Because the jejunal bypass was made as short as possible (the afferent limb was as close as 20 cm) and the size of the remnant stomach was equivalent between the BI and RY groups, variations in malabsorption between the groups were minimized. Thus, we believe that this study was also the first to evaluate prospectively collected data to determine the specific effects of duodenal bypass on visceral and subcutaneous fat loss in a population in which the remnant stomach was of a similar size.

It is assumed that the number of patients with gastric cancer who are obese is increasing because of the high prevalence of obesity among the general population. The number of patients diagnosed with early gastric cancer is increasing as the result of earlier detection of cancer, and the 5-year survival rate for patients with early gastric cancer (most often treated with radical resection) is approximately 95%.<sup>15</sup> Consequently, death by causes other than cancer is the most common cause of death among patients with early gastric cancer. Cerebrovascular disorders, cardiac disease, and respiratory disease are reported to be common causes of death in patients with early gastric cancer.<sup>16</sup> When treating these patients, we should therefore consider the most effective means of reducing the risk of death due to causes other than cancer. In recent years, visceral fat accumulation has been identified as one of the underlying causes of metabolic syndrome. This syndrome is characterized by glucose intolerance, obesity, hypertension, and dyslipidemia. Many

studies have demonstrated that body fat distribution is associated with the development of metabolic disorders, and that excessive abdominal fat, especially intra-abdominal visceral fat, is associated with various obesity-related complications and poor prognosis.<sup>17,18</sup> Visceral fat is becoming a target for the treatment of obesity-related complications such as hypertension, dyslipidemia, diabetes mellitus, and cardiovascular disease.<sup>19</sup>

Our study revealed that duodenal bypass in addition to gastrectomy promoted visceral fat loss, especially in obese patients. Previous studies of bariatric surgery have reported that the decrease in absolute BMI in lower BMI groups is less than that of the groups with greater BMI 1 year after RY bypass operation.<sup>20</sup> This finding is consistent with our results. RY reconstruction might be a better choice for obese patients who require distal gastrectomy to treat gastric cancer. Our results also suggest that duodenal bypass is a useful procedure for nonobese patients with metabolic syndrome-associated conditions such as diabetes mellitus, hypertension, and hyperlipidemia, because the reduction in visceral fat was greater after this procedure.

There have been a few reports about the effect on diabetes of rearrangements of gastrointestinal anatomy after surgery for gastric cancer.<sup>21-23</sup> Lanzarini et al<sup>21</sup> reported that gastrectomy with RY reconstruction (60–70 cm limb) in type 2 diabetes patients who underwent operation mainly for gastric cancer correlated with remission of diabetes in 65% and improvement in 30.4% of patients. Another study reported that patients who underwent duodenal bypass had significantly improved diabetes compared with those who did not.<sup>23</sup>

The mechanism by which duodenal bypass reduces visceral fat could not be elucidated in this study. However, previous studies of bariatric operation have reported that visceral fat reduction is greater after RY gastric bypass than after vertical banded gastroplasty.<sup>14</sup> Although the mechanisms of fat reduction or improvement in insulin resistance are not understood completely within the context of bariatric surgery, gut hormones are thought to play a critical role. Among the various gut hormones, gastric inhibitory polypeptide (GIP) is reported to regulate fat metabolism. GIP is released from the duodenal endocrine K cells immediately after the absorption of fat or glucose.<sup>24</sup> Furthermore, fat intake induces hypersecretion of GIP, which increases nutrient uptake and triglyceride accumulation in adipocytes.<sup>25</sup> Korner et al<sup>26</sup> reported lower GIP levels after RY gastric bypass compared

to adjustable gastric banding, and concluded that blunted GIP secretion after RY may contribute to the greater weight loss and improved glucose homeostasis compared to adjustable gastric banding. Fat malabsorption may be another factor; clinical tests after RY revealed significantly lower fat absorption than after BI and double-tract reconstruction, in which the passage of food through the duodenum is accommodated.<sup>27</sup>

Our study has several limitations. First, we could not elucidate the mechanism of greater reduction of visceral fat after duodenal bypass, because data about gut hormones were not acquired. In addition, the long-term results are unknown, because we examined CT data only 1 year after surgery. In studies of the long-term results of bariatric surgery, compared with nonsurgical control patients, the use of RY gastric bypass operation in severely obese patients was associated with a greater rate of diabetes remission and a lesser risk of cardiovascular disease and other poor health outcomes after 6 years. On the other hand, there are some reports of recurrence or worsening of diabetes mellitus, especially in non-obese patients, after RY gastric bypass.<sup>28,29</sup> Further investigations will be necessary to provide long-term follow-up data and to understand how duodenal bypass markedly decreases fat.

We thank Toshimitsu Hamasaki, Associate Professor of the Department of Biomedical Statistics at Osaka University Graduate School of Medicine, who provided us with advice regarding statistical analysis for this article.

## REFERENCES

1. Korner J, Inabnet W, Conwell IM, Taveras C, Daud A, Olivero-Rivera L, et al. Differential effects of gastric bypass and banding on circulating gut hormone and leptin levels. *Obesity* (Silver Spring, Md) 2006;14:1553-61.
2. Kumar R, Lieske JC, Collazo-Clavell ML, Sarr MG, Olson ER, Vrtiska TJ, et al. Fat malabsorption and increased intestinal oxalate absorption are common after Roux-en-Y gastric bypass surgery. *Surgery* 2011;149:654-61.
3. Takiguchi S, Yamamoto K, Hirao M, Imamura H, Fujita J, Yano M, et al. A comparison of postoperative quality of life and dysfunction after Billroth I and Roux-en-Y reconstruction following distal gastrectomy for gastric cancer: results from a multi-institutional RCT. *Gastric Cancer* 2012;15:198-205.
4. Hirao M, Takiguchi S, Imamura H, Yamamoto K, Kurokawa Y, Fujita J, et al. Comparison of Billroth I and Roux-en-Y reconstruction after distal gastrectomy for gastric cancer: one-year postoperative effects assessed by a multi-institutional RCT. *Ann Surg Oncol* 2013;20:1591-7.
5. Tokunaga K, Matsuzawa Y, Ishikawa K, Tarui S. A novel technique for the determination of body fat by computed tomography. *Int J Obes* 1983;7:437-45.
6. Kvist H, Chowdhury B, Sjostrom L, Tylen U, Cederblad A. Adipose tissue volume determination in males by computed tomography and 40K. *Int J Obes* 1988;12:249-66.

7. Yoshizumi T, Nakamura T, Yamane M, Islam AH, Menju M, Yamasaki K, et al. Abdominal fat: standardized technique for measurement at CT. *Radiology* 1999;211:283-6.
8. Japanese Gastric Cancer A. Japanese Classification of Gastric Carcinoma. 2nd English Edition. *Gastric Cancer* 1998;1:10-24.
9. Onodera T, Goseki N, Kosaki G. Prognostic nutritional index in gastrointestinal surgery of malnourished cancer patients [in Japanese]. *Nihon Geka Gakkai Zasshi* 1984; 85:1001-5.
10. Armbrecht U, Lundell L, Lindstedt G, Stockbruegger RW. Causes of malabsorption after total gastrectomy with Roux-en-Y reconstruction. *Acta Chir Scand* 1988;154:37-41.
11. Bradley EL 3rd, Isaacs J, Hersh T, Davidson ED, Millikan W. Nutritional consequences of total gastrectomy. *Ann Surg* 1975;182:415-29.
12. Friess H, Bohm J, Muller MW, Glasbrenner B, Riepl RL, Malferteiner P, et al. Maldigestion after total gastrectomy is associated with pancreatic insufficiency. *Am J Gastroenterol* 1996;91:341-7.
13. Adams JF. The clinical and metabolic consequences of total gastrectomy. I. Morbidity, weight, and nutrition. *Scand J Gastroenterol* 1967;2:137-49.
14. Olbers T, Bjorkman S, Lindroos A, Maleckas A, Lonn L, Sjostrom L, et al. Body composition, dietary intake, and energy expenditure after laparoscopic Roux-en-Y gastric bypass and laparoscopic vertical banded gastroplasty: a randomized clinical trial. *Ann Surg* 2006;244:715-22.
15. Isobe Y, Nashimoto A, Akazawa K, Oda I, Hayashi K, Miyashiro I, et al. Gastric cancer treatment in Japan: 2008 annual report of the JGCA nationwide registry. *Gastric Cancer* 2011;14:301-16.
16. Kunisaki C, Akiyama H, Nomura M, Matsuda G, Otsuka Y, Ono H, et al. Significance of long-term follow-up of early gastric cancer. *Ann Surg Oncol* 2006;13:363-9.
17. Fujioka S, Matsuzawa Y, Tokunaga K, Tarui S. Contribution of intra-abdominal fat accumulation to the impairment of glucose and lipid metabolism in human obesity. *Metabolism* 1987;36:54-9.
18. Marcus MA, Murphy L, Pi-Sunyer FX, Albu JB. Insulin sensitivity and serum triglyceride level in obese white and black women: relationship to visceral and truncal subcutaneous fat. *Metabolism* 1999;48:194-9.
19. Sjostrom L, Peltonen M, Jacobson P, Sjostrom CD, Karason K, Wedel H, et al. Bariatric surgery and long-term cardiovascular events. *JAMA* 2012;307:56-65.
20. Lee WJ, Wang W, Lee YC, Huang MT, Ser KH, Chen JC. Effect of laparoscopic mini-gastric bypass for type 2 diabetes mellitus: comparison of BMI>35 and <35 kg/m2. *J Gastrointest Surg* 2008;12:945-52.
21. Lanzarini E, Csendes A, Lembach H, Molina J, Gutierrez L, Silva J. Evolution of type 2 diabetes mellitus in non morbid obese gastrectomized patients with Roux en-Y reconstruction: retrospective study. *World J Surg* 2010; 34:2098-102.
22. Lee W, Ahn SH, Lee JH, Park DJ, Lee HJ, Kim HH, et al. Comparative study of diabetes mellitus resolution according to reconstruction type after gastrectomy in gastric cancer patients with diabetes mellitus. *Obes Surg* 2012;22:1238-43.
23. Kim JW, Cheong JH, Hyung WJ, Choi SH, Noh SH. Outcome after gastrectomy in gastric cancer patients with type 2 diabetes. *World J Gastroenterol* 2012;18:49-54.
24. Falko JM, Crockett SE, Cataland S, Mazzaferri EL. Gastric inhibitory polypeptide (GIP) stimulated by fat ingestion in man. *J Clin Endocrinol Metab* 1975;41:260-5.
25. Miyawaki K, Yamada Y, Ban N, Ihara Y, Tsukiyama K, Zhou H, et al. Inhibition of gastric inhibitory polypeptide signaling prevents obesity. *Nat Med* 2002;8:738-42.
26. Korner J, Bessler M, Inabnet W, Taveras C, Holst JJ. Exaggerated glucagon-like peptide-1 and blunted glucose-dependent insulinotropic peptide secretion are associated with Roux-en-Y gastric bypass but not adjustable gastric banding. *Surg Obes Relat Dis* 2007;3:597-601.
27. Takase M, Sumiyama Y, Nagao J. Quantitative evaluation of reconstruction methods after gastrectomy using a new type of examination: digestion and absorption test with stable isotope <sup>13</sup>C-labeled lipid compound. *Gastric Cancer* 2003; 6:134-41.
28. DiGiorgi M, Rosen DJ, Choi JJ, Milone L, Schrope B, Olivero-Rivera L, et al. Re-emergence of diabetes after gastric bypass in patients with mid- to long-term follow-up. *Surg Obes Relat Dis* 2010;6:249-53.
29. Arterburn DE, Bogart A, Sherwood NE, Sidney S, Coleman KJ, Haneuse S, et al. A multisite study of long-term remission and relapse of type 2 diabetes mellitus following gastric bypass. *Obes Surg* 2013;23:93-102.

WJG 20<sup>th</sup> Anniversary Special Issues (8): Gastric cancer

## Current status of function-preserving surgery for gastric cancer

Takuro Saito, Yukinori Kurokawa, Shuji Takiguchi, Masaki Mori, Yuichiro Doki

Takuro Saito, Yukinori Kurokawa, Shuji Takiguchi, Masaki Mori, Yuichiro Doki, Department of Gastroenterological Surgery, Osaka University Graduate School of Medicine, Osaka 565-0871, Japan

Author contributions: All authors contributed to conception and design, acquisition of data, or analysis and interpretation of data.

Correspondence to: Yukinori Kurokawa, MD, PhD, Department of Gastroenterological Surgery, Osaka University Graduate School of Medicine, 2-2-E2, Yamadaoka, Suita, Osaka 565-0871, Japan. [ykurokawa@gesurg.med.osaka-u.ac.jp](mailto:ykurokawa@gesurg.med.osaka-u.ac.jp)

Telephone: +81-6-68793251 Fax: +81-6-68793259

Received: May 27, 2014 Revised: July 16, 2014

Accepted: September 5, 2014

Published online: December 14, 2014

### Abstract

Recent advances in diagnostic techniques have allowed the diagnosis of gastric cancer (GC) at an early stage. Due to the low incidence of lymph node metastasis and favorable prognosis in early GC, function-preserving surgery which improves postoperative quality of life may be possible. Pylorus-preserving gastrectomy (PPG) is one such function-preserving procedure, which is expected to offer advantages with regards to dumping syndrome, bile reflux gastritis, and the frequency of flatus, although PPG may induce delayed gastric emptying. Proximal gastrectomy (PG) is another function-preserving procedure, which is thought to be advantageous in terms of decreased duodenogastric reflux and good food reservoir function in the remnant stomach, although the incidence of heartburn or gastric fullness associated with this procedure is high. However, these disadvantages may be overcome by the reconstruction method used. The other important problem after PG is remnant GC, which was reported to occur in approximately 5% of patients. Therefore, the reconstruction technique used with PG should facilitate postoperative

endoscopic examinations for early detection and treatment of remnant gastric carcinoma. Oncologic safety seems to be assured in both procedures, if the preoperative diagnosis is accurate. Patient selection should be carefully considered. Although many retrospective studies have demonstrated the utility of function-preserving surgery, no consensus on whether to adopt function-preserving surgery as the standard of care has been reached. Further prospective randomized controlled trials are necessary to evaluate survival and postoperative quality of life associated with function-preserving surgery.

© 2014 Baishideng Publishing Group Inc. All rights reserved.

**Key words:** Gastric cancer; Function preserving surgery; Quality of life; Pylorus preserving surgery; Proximal gastrectomy

**Core tip:** We reviewed the current status of two function-preserving surgeries for gastric cancer (GC), pylorus-preserving surgery and proximal gastrectomy (PG). Although both procedures appear to be oncologically safe for early GC, issues regarding postoperative quality of life remain, especially with PG. The effect of the reconstruction method after PG on postoperative quality of life was analyzed, including the novel double tract reconstruction method, which is expected to overcome disadvantages associated with esophagogastrectomy and jejunal interposition reconstruction. Although some reports showed a benefit with function-preserving surgery, further randomized trials are needed.

Saito T, Kurokawa Y, Takiguchi S, Mori M, Doki Y. Current status of function-preserving surgery for gastric cancer. *World J Gastroenterol* 2014; 20(46): 17297-17304 Available from: URL: <http://www.wjgnet.com/1007-9327/full/v20/i46/17297.htm> DOI: <http://dx.doi.org/10.3748/wjg.v20.i46.17297>