## MINIREVIEWS

**5934** Development of clustered regularly interspaced short palindromic repeats/CRISPR-associated technology for potential clinical applications  
*Huang YY, Zhang XY, Zhu P, Ji L*

**5946** Strategies and challenges in treatment of varicose veins and venous insufficiency  
*Gao RD, Qian SY, Wang HH, Liu YS, Ren SY*

**5957** Diabetes mellitus susceptibility with varied diseased phenotypes and its comparison with phenome interactome networks  
*Rout M, Kour B, Vuree S, Lulu SS, Medicherla KM, Suravajhala P*

## ORIGINAL ARTICLE

### Clinical and Translational Research

**5965** Identification of potential key molecules and signaling pathways for psoriasis based on weighted gene co-expression network analysis  
*Shu X, Chen XX, Kang XD, Ran M, Wang YL, Zhao ZK, Li CX*

**5984** Construction and validation of a novel prediction system for detection of overall survival in lung cancer patients  
*Zhong C, Liang Y, Wang Q, Tan HW, Liang Y*

### Case Control Study

**6001** Effectiveness and postoperative rehabilitation of one-stage combined anterior-posterior surgery for severe thoracolumbar fractures with spinal cord injury  
*Zhang B, Wang JC, Jiang YZ, Song QF, An Y*

### Retrospective Study

**6009** Prostate sclerosing adenopathy: A clinicopathological and immunohistochemical study of twelve patients  
*Feng RL, Tao YP, Tan ZY, Fu S, Wang HF*

**6021** Value of magnetic resonance diffusion combined with perfusion imaging techniques for diagnosing potentially malignant breast lesions  
*Zhang H, Zhang XY, Wang Y*

**6032** Scar-centered dilation in the treatment of large keloids  
*Wu M, Gu JY, Duan R, Wei BX, Xie F*

**6039** Application of a novel computer-assisted surgery system in percutaneous nephrolithotomy: A controlled study  
## Contents

**World Journal of Clinical Cases**  
Thrice Monthly Volume 10 Number 18 June 26, 2022

<table>
<thead>
<tr>
<th>Page</th>
<th>Title</th>
<th>Authors</th>
<th>Article Type</th>
</tr>
</thead>
<tbody>
<tr>
<td>6050</td>
<td>Influences of etiology and endoscopic appearance on the long-term outcomes of gastric antral vascular ectasia</td>
<td>Kwon HJ, Lee SH, Cho JH</td>
<td>Randomized Controlled Trial</td>
</tr>
<tr>
<td>6060</td>
<td>Evaluation of the clinical efficacy and safety of TST33 mega hemorrhoidectomy for severe prolapsed hemorrhoids</td>
<td>Tao L, Wei J, Ding XF, Ji LJ</td>
<td>Randomized Controlled Trial</td>
</tr>
<tr>
<td>6069</td>
<td>Sequential chemotherapy and icotinib as first-line treatment for advanced epidermal growth factor receptor-mutated non-small cell lung cancer</td>
<td>Sun SJ, Han JD, Liu W, Wu ZY, Zhao X, Yan X, Jiao SC, Fang J</td>
<td>Randomized Clinical Trial</td>
</tr>
<tr>
<td>6082</td>
<td>Impact of preoperative carbohydrate loading on gastric volume in patients with type 2 diabetes</td>
<td>Lin XQ, Chen YR, Chen X, Cai YP, Lin JX, Xu DM, Zheng XC</td>
<td></td>
</tr>
<tr>
<td>6091</td>
<td>Efficacy and safety of adalimumab in comparison to infliximab for Crohn's disease: A systematic review and meta-analysis</td>
<td>Yang HH, Huang Y, Zhou XC, Wang RN</td>
<td>META-ANALYSIS</td>
</tr>
<tr>
<td>6105</td>
<td>Successful treatment of acute relapse of chronic eosinophilic pneumonia with benralizumab and without corticosteroids: A case report</td>
<td>Izhakian S, Pertzov B, Rosengarten D, Kramer MR</td>
<td>CASE REPORT</td>
</tr>
<tr>
<td>6119</td>
<td>Hepatic epithelioid hemangioendothelioma after thirteen years’ follow-up: A case report and review of literature</td>
<td>Mo WF, Tong YL</td>
<td></td>
</tr>
<tr>
<td>6128</td>
<td>Effectiveness and safety of ultrasound-guided intramuscular lauromacrogol injection combined with hysteroscopy in cervical pregnancy treatment: A case report</td>
<td>Ye JP, Gao Y, Lu LW, Ye YJ</td>
<td></td>
</tr>
<tr>
<td>6136</td>
<td>Carcinoma located in a right-sided sigmoid colon: A case report</td>
<td>Lyu LJ, Yao WW</td>
<td></td>
</tr>
<tr>
<td>Page</td>
<td>Title</td>
<td>Authors</td>
<td></td>
</tr>
<tr>
<td>------</td>
<td>----------------------------------------------------------------------</td>
<td>------------------------------------------------------------------------</td>
<td></td>
</tr>
<tr>
<td>6148</td>
<td>Overlapping syndrome of recurrent anti-N-methyl-D-aspartate receptor</td>
<td>Yin XJ, Zhang LF, Bao LH, Feng ZC, Chen JH, Li BX, Zhang J</td>
<td></td>
</tr>
<tr>
<td></td>
<td>encephalitis and anti-myelin oligodendrocyte glycoprotein demyelinating</td>
<td></td>
<td></td>
</tr>
<tr>
<td></td>
<td>diseases: A case report</td>
<td></td>
<td></td>
</tr>
<tr>
<td>6156</td>
<td>Liver transplantation for late-onset ornithine transcarbamylase</td>
<td>Fu XH, Hu YH, Liao JX, Chen L, Hu ZQ, Wen JL, Chen SL</td>
<td></td>
</tr>
<tr>
<td></td>
<td>deficiency: A case report</td>
<td></td>
<td></td>
</tr>
<tr>
<td>6163</td>
<td>Disseminated strongyloidiasis in a patient with rheumatoid arthritis</td>
<td>Zheng JH, Xue LY</td>
<td></td>
</tr>
<tr>
<td></td>
<td>A case report</td>
<td></td>
<td></td>
</tr>
<tr>
<td>6168</td>
<td>CYP27A1 mutation in a case of cerebrotendinous xanthomatosis: A case</td>
<td>Li ZR, Zhou YL, Jin Q, Xie YY, Meng HM</td>
<td></td>
</tr>
<tr>
<td></td>
<td>report</td>
<td></td>
<td></td>
</tr>
<tr>
<td>6175</td>
<td>Postoperative multiple metastasis of clear cell sarcoma-like tumor in</td>
<td>Huang WP, Li LM, Gao JB</td>
<td></td>
</tr>
<tr>
<td></td>
<td>the gastrointestinal tract in adolescent: A case report</td>
<td></td>
<td></td>
</tr>
<tr>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>6184</td>
<td>Toripalimab combined with targeted therapy and chemotherapy achieves</td>
<td>Liu R, Wang X, Ji Z, Deng T, Li HL, Zhang YH, Yang YC, Ge SH, Zhang L,</td>
<td></td>
</tr>
<tr>
<td></td>
<td>pathologic complete response in gastric carcinoma: A case report</td>
<td>Bai M, Ning T, Ba Y</td>
<td></td>
</tr>
<tr>
<td>6192</td>
<td>Presentation of Boerhaave’s syndrome as an upper-esophageal perforation</td>
<td>Tan N, Luo YH, Li GC, Chen YL, Tan W, Xiang YH, Ge L, Yao D, Zhang MH</td>
<td></td>
</tr>
<tr>
<td></td>
<td>associated with a right-sided pleural effusion: A case report</td>
<td></td>
<td></td>
</tr>
<tr>
<td></td>
<td>cancer patient: A case report and review of literature</td>
<td></td>
<td></td>
</tr>
<tr>
<td>6205</td>
<td>Nontraumatic convexal subarachnoid hemorrhage: A case report</td>
<td>Chen HL, Li B, Chen C, Fan X, Ma WB</td>
<td></td>
</tr>
<tr>
<td>6211</td>
<td>Growth hormone ameliorates hepatopulmonary syndrome and nonalcoholic</td>
<td>Zhang XY, Yuan K, Fang YL, Wang CL</td>
<td></td>
</tr>
<tr>
<td></td>
<td>steatohepatitis secondary to hypopituitarism in a child: A case report</td>
<td></td>
<td></td>
</tr>
<tr>
<td>6218</td>
<td>Vancomycin dosing in an obese patient with acute renal failure: A</td>
<td>Xu KY, Li D, Hu ZJ, Zhao CC, Bai J, Du WL</td>
<td></td>
</tr>
<tr>
<td></td>
<td>case report and review of literature</td>
<td></td>
<td></td>
</tr>
<tr>
<td>6227</td>
<td>Insulinoma after sleeve gastrectomy: A case report</td>
<td>Lobaton-Ginsberg M, Sotelo-González P, Ramirez-Renteria C, Juárez-Aguilar FG, Ferreira-Hermosillo A</td>
<td></td>
</tr>
<tr>
<td>6234</td>
<td>Primary intestinal lymphangiectasia presenting as limb convulsions: A</td>
<td>Cao Y, Feng XH, Ni HX</td>
<td></td>
</tr>
<tr>
<td></td>
<td>case report</td>
<td></td>
<td></td>
</tr>
<tr>
<td>6241</td>
<td>Esophagogastric junctional neuroendocrine tumor with adenocarcinoma:</td>
<td>Kong ZZ, Zhang L</td>
<td></td>
</tr>
<tr>
<td></td>
<td>A case report</td>
<td></td>
<td></td>
</tr>
</tbody>
</table>
Contents

6247 Foreign body granuloma in the tongue differentiated from tongue cancer: A case report
   Jiang ZH, Xu R, Xia L

6254 Modified endoscopic ultrasound-guided selective N-butyl-2-cyanoacrylate injections for gastric variceal hemorrhage in left-sided portal hypertension: A case report
   Yang J, Zeng Y, Zhang JW

6261 Management of type IIIb dens invaginatus using a combination of root canal treatment, intentional replantation, and surgical therapy: A case report
   Zhang J, Li N, Li WL, Zheng XY, Li S

6269 Clivus-involved immunoglobulin G4 related hypertrophic pachymeningitis mimicking meningioma: A case report
   Yu Y, Lv L, Yin SL, Chen C, Jiang S, Zhou PZ

6277 De novo brain arteriovenous malformation formation and development: A case report
   Huang H, Wang X, Guo AN, Li W, Duan RH, Fang JH, Yin B, Li DD

6283 Coinfection of Streptococcus suis and Nocardia asiatica in the human central nervous system: A case report
   Chen YY, Xue XH

6289 Dilated left ventricle with multiple outpouchings — a severe congenital ventricular diverticulum or left-dominant arrhythmogenic cardiomyopathy: A case report
   Zhang X, Ye RY, Chen XP

6298 Spontaneous healing of complicated crown-root fractures in children: Two case reports
   Zhou ZL, Guo L, Sun SK, Li HS, Zhang CD, Kou WW, Xu Z, Wu LA

6307 Thyroid follicular renal cell carcinoma excluding thyroid metastases: A case report
   Wu SC, Li XY, Liao BJ, Xie K, Chen WM

6314 Appendiceal bleeding: A case report
   Zhou SY, Guo MD, Ye XH

6319 Spontaneous healing after conservative treatment of isolated grade IV pancreatic duct disruption caused by trauma: A case report
   Mei MZ, Ren YF, Mou YP, Wang YY, Jin WW, Lu C, Zhu QC

6325 Pneumonia and seizures due to hypereosinophilic syndrome—organ damage and eosinophilia without synchronisation: A case report
   Ishida T, Murayama T, Kobayashi S

6333 Creutzfeldt-Jakob disease presenting with bilateral hearing loss: A case report
   Na S, Lee SA, Lee JD, Lee ES, Lee TK

LETTER TO THE EDITOR

6338 Stem cells as an option for the treatment of COVID-19
   Cuevas-González MV, Cuevas-González JC
ABOUT COVER
Editorial Board Member of World Journal of Clinical Cases, Cristina Tudoran, PhD, Assistant Professor, Department VII, Internal Medicine II, Discipline of Cardiology, "Victor Babes" University of Medicine and Pharmacy Timisoara, Timisoara 300041, Timis, Romania. cristina13.tudoran@gmail.com

AIMS AND SCOPE
The primary aim of World Journal of Clinical Cases (WJCC, World J Clin Cases) is to provide scholars and readers from various fields of clinical medicine with a platform to publish high-quality clinical research articles and communicate their research findings online.

WJCC mainly publishes articles reporting research results and findings obtained in the field of clinical medicine and covering a wide range of topics, including case control studies, retrospective cohort studies, retrospective studies, clinical trials studies, observational studies, prospective studies, randomized controlled trials, randomized clinical trials, systematic reviews, meta-analysis, and case reports.

INDEXING/ABSTRACTING
The WJCC is now indexed in Science Citation Index Expanded (also known as SciSearch®), Journal Citation Reports/Science Edition, Scopus, PubMed, and PubMed Central. The 2021 Edition of Journal Citation Reports® cites the 2020 impact factor (IF) for WJCC as 1.337; IF without journal self cites: 1.301; 5-year IF: 1.742; Journal Citation Indicator: 0.33; Ranking: 119 among 169 journals in medicine, general and internal; and Quartile category: Q3. The WJCC’s CiteScore for 2020 is 0.8 and Scopus CiteScore rank 2020: General Medicine is 493/793.
Insulinoma after sleeve gastrectomy: A case report

Miry Lobaton-Ginsberg, Pilar Sotelo-González, Claudia Ramirez-Renteria, Fany Gabriela Juárez-Aguilar, Aldo Ferreira-Hermosillo

Abstract

BACKGROUND
Laparoscopic sleeve gastrectomy (LSG) has been proposed as an effective and durable treatment for severe obesity and glucose metabolism disorders, and its prevalence has increased from 5% to 37% since 2008. One common complication after bariatric surgery is a postprandial hyperinsulinemic hypoglycemic state. While rare, insulinomas can cause this state, where symptoms are more common in the fasting state; thus, evaluation of insulin secretion is needed. Until now, there have been no reports of insulinoma after LSG.

CASE SUMMARY
We describe the case of a 43-year-old woman who was referred to the obesity clinic 2 years after LSG was performed. She had symptoms of hypoglycemia predominantly in the fasting state and documented hypoglycemia of less than 30 mg/dL, which are compatible with Whipple’s triad. Initially, dumping syndrome was suspected, but after a second low fasting plasma glucose was documented, a 72-h fasting test was performed that tested positive. Computed tomography and endoscopic ultrasound were performed, identifying the presence of a homogeneous hypoechoic semi-oval tumoral lesion in the pancreas. The diagnosis was compatible with insulinoma. After laparoscopic enucleation of the insulinoma, the symptoms and hypoglycemia disappeared. The histopathological report described a well-differentiated grade 2 neuroendocrine tumor with positive chromogranin and synaptophysin and Ki67 immunopositivity in 4% of the neoplastic cells.
CONCLUSION
Insulinoma after LSG is a rare condition, and clinicians must be aware of it, especially if the patient has hypoglycemic symptoms during the fasting state.

Key Words: Insulinoma; Hypoglycemia; Bariatric surgery; Glucagon-like peptide 1 amide; Neuroendocrine tumors; Case report

©The Author(s) 2022. Published by Baishideng Publishing Group Inc. All rights reserved.

Core Tip: Neuroglycopenic symptoms compatible with Whipple’s triad were identified in a woman 2 years after laparoscopic sleeve gastrectomy, predominantly occurring in the fasting state. After discarding late dumping syndrome, a 72-h fasting test was performed and tested positive. Imaging techniques documented the presence of a tumoral lesion in the pancreas, compatible with insulinoma. After laparoscopic enucleation of the insulinoma, the symptoms were relieved. When hypoglycemia occurs after bariatric surgery, evaluation of insulin secretion is needed to conduct a correct diagnostic approach. Follow-up must be performed by a multidisciplinary team.

INTRODUCTION
The obesity pandemic has become a great topic of interest due to its implications for quality of life, comorbidities, increasing mortality, and the economic impact on health services worldwide[1]. Bariatric surgery (BS) is an effective and durable treatment for severe obesity and glucose metabolism disorders, with laparoscopic Roux-Y gastric bypass surgery (RYGB) being the most common procedure[2,3]. Nevertheless, laparoscopic sleeve gastrectomy (LSG) has been proposed as a procedure capable of achieving the same goals, but with fewer complications[4].

A common complication in BS is the development of a postprandial hyperinsulinemic hypoglycemic state[5]. Hypoglycemia is defined as a glucose level below 70 mg/dL according to the American Diabetes Association guidelines[6]. The possible causes of hypoglycemia in patients who had undergone BS include late dumping syndrome, nesidioblastosis and, rarely, insulinoma[5,7]. Up to 40 cases of nesidioblastosis have been reported after RYGB, and only one case has been reported after sleeve gastrectomy [8,9]. To our knowledge, there are few reports of insulinoma after BS[10] but no reports after LSG. The purpose of these case reports is to inform clinicians that patients with neuroglycopenic symptoms during the fasting state could have hypoglycemia caused by insulinoma, which is not only due to late dumping syndrome.

A review of the medical literature for insulinoma and hypoglycemia after BS was performed in PubMed. We searched “insulinoma”, “hypoglycemia”, “sleeve gastrectomy”, “RYGB”, “glucagon-like peptide 1 (GLP-1)”, and “ghrelin” and a combination of the above terms including all dates up to October 2021. Herein, we present the case of a 43-year-old woman referred to the obesity clinic due to neuroglycopenic symptoms caused by an insulinoma 2 years after a sleeve gastrectomy.

CASE PRESENTATION

Chief complaints
A 43-year-old woman was referred to the obesity clinic due to neuroglycopenic symptoms caused by an insulinoma 2 years after a sleeve gastrectomy.

History of present illness
In March 2020, 2 years after LSG was performed, the patient developed neuroglycopenic symptoms including short-term memory loss, lingual nerve paresthesia, and nonspecific visual alterations predominantly during the morning in a fasting state. These symptoms were suppressed with food intake. Two months later, she visited a physician who documented fasting plasma glucose of 27 mg/dL, and in June 2020, the symptoms occurred more frequently, and she gained 14 kg. In the beginning, late dumping symptoms were suspected, but in September 2020, fasting plasma glucose of 30 mg/dL was docu-
mented, so she was hospitalized for the evaluation of hypoglycemia in a 72-h supervised fast test. She had baseline plasma glucose of 67 mg/dL, nonsuppressed insulin of 16.4 IU/mL, and C-peptide of 3.64 ng/mL. In the first hour after initiation, she developed Whipple’s triad symptoms, and her lab results detected plasma glucose of 38 mg/dL, insulin of 25.9 IU/mL, and C-peptide of 4.31 ng/mL. Thus, it was decided to stop the protocol and initiate 1000 mL of 20% glucose solution in 12 h.

**History of past illness**
In 2002, the patient was diagnosed with obesity and dyslipidemia (high triglycerides and cholesterol with low HDL) and treated with improvements in diet, physical activity, and statins without weight control. In 2016, a gastric balloon was placed, and although her body mass index (BMI) in 2018 was 34.4 kg/m², LSG was performed.

**Personal and family history**
The patient had no specific personal or family history.

**Physical examination**
After LSG, the patient weighed 74 kg, and her BMI was 32 kg/m². The physical examination showed no obvious cardiovascular or respiratory abnormalities. The abdomen was soft, and the only sign was the presence of postsurgery scars.

**Laboratory examinations**
Upon hospitalization prior to the surgery, the patient’s hemoglobin A1c level was 4.8% (normal range: < 5.7%). The C-peptide value was normal at 3.64 ng/mL (1.1-4.4 ng/mL), and insulin was mildly elevated at 16.40 µU/mL (3.21-16.30 µU/mL). Lipid levels indicated dyslipidemia with total cholesterol of 224 mg/dL and LDL-c of 142.8 mg/dL. Other biochemical parameters were normal and only an iron deficiency anemia was documented. Thyroid function was normal, with TSH 2.46 µU/mL (0.27-4.20 µU/mL), FT4 1.06 ng/dL (0.93-1.70 ng/dL), and cortisol level 15.04 ug/dL (3.70-19.40 µg/dL), all within the normal range.

**Imaging examinations**
Computed tomography (CT) demonstrated the presence of a focal asymmetric reinforcement area in the head of the pancreas (Figure 1A). Endoscopic ultrasound showed the presence of a tumoral lesion in the pancreas in close proximity to the main pancreatic duct and splenomesenteric confluence without evidence of invasion (Figure 1B and C).

**FINAL DIAGNOSIS**
The final diagnosis was insulinoma. This was confirmed by histology and immunohistochemistry of the tumor (Figure 2).

**TREATMENT**
After a surgery consultation, a laparoscopic insulinoma enucleation was performed without complications. No other tumors were identified in the upper abdomen.

**OUTCOME AND FOLLOW-UP**
Histopathological findings revealed a well-differentiated neuroendocrine grade 2 tumor with free edges. Immunohistochemical studies confirmed positive chromogranin and synaptophysin as well as a proliferative activity (Ki67) in 4% of neoplastic cells.

After surgery, the neuroglycopenic symptoms were relieved, and the patient had no hypoglycemic events. Her current treatment is diet and physical activity, targeting a BMI of 31.1 kg/m².

**DISCUSSION**
Since 2013, 468609 BSs have been performed worldwide[2]. LSG was initially introduced as a first-stage restrictive procedure to a more complex definitive one. At present, it is performed as a stand-alone BS [7]. Since 2008, the prevalence of the LSG procedure has increased from 5% to 37% worldwide[2], but in Mexico, it is performed only in 13% of patients, whereas LRYGB is performed in 85.8%, with a bypass/
sleeve ratio of 7:1. In our center, LSG accounts for 20% of total BSs (200 procedures since 2010).

LSG comprises vertical longitudinal resection of the greater gastric curve that includes the fundus, body, and antrum as well as the formation of a tubular conduit with a capacity of < 100 mL. Weight loss is achieved by restrictive and humoral effects[8,11].

Hypoglycemia is a well-documented complication after BS. Papamargaritis et al[12,13] recorded a study where 33% of patients experienced severe hypoglycemia a year after LSG due to late dumping symptoms, which usually occurs 1-3 h after a high-carbohydrate meal triggering a hyperinsulinemic response. Since 2005, up to 40 cases of nesidioblastosis after RYGB have been reported[8], and only one case after LSG was documented in 2019 by Kim et al[9]. While rare, insulinomas have been reported after BS. Mulla et al[10] described seven cases of insulinoma, one patient with pancreatic neuroendocrine tumor, and one patient with insulinoma and pancreatic neuroendocrine tumor after BS, 78% of whom were women. In these cases, hypoglycemia was more common in the fasting state.

The mechanism of the post-BS hyperinsulinemic hypoglycemic state and the changes in beta cell proliferation are not fully understood. In the LSG, the faster transit of undigested nutrients to the distal gastrointestinal tract due to rapid gastric emptying upregulates the production of GLP-1 secreted by enteroendocrine L cells in the distal intestine. This increase can normalize blood glucose and regulate insulin synthesis and proinsulin gene expression, as well as glucagon and somatostatin secretion[3].

GLP-1 has multiple beneficial effects on β cells, including an increase in their number by inhibiting apoptosis and enhancing neogenesis as well as promoting its proliferation. In a study carried out in 2016 by Xu et al[14], it was found that a chemically modified GLP-1 (mGLP-1) analog promotes the proliferation of pancreatic mouse β cells, upregulating the expression of cyclin E, CDK2, Bcl-2, Bax, and p21. The cyclin E-CDK2 complex plays an important role in the regulation of the G1 phase of the G1/S cell cycle, while p21 is a universal cyclin-dependent kinase (CKI) inhibitor. Meanwhile, the Bcl-2 and Bax genes, two important members of the Bcl-2 gene family, have opposite functions, inhibiting or promoting cell apoptosis, respectively[14].

An increase in ghrelin levels has been observed a year after BS[15]. Ghrelin and the type 1a ghrelin receptor (GHS-R1A) are expressed in different types of neuroendocrine tumors. Recently, Wu et al[16]
found that GHS-R1a was found in 60% of insulinomas, suggesting that ghrelin may act through autocrine or paracrine pathways. The proliferative effects of ghrelin and its association with insulinoma have not been studied, although there is a clinical case report where a ghrelin-producing neuroendocrine tumor was transformed into an insulinoma[17].

The diagnosis of hypoglycemia after BS is challenging. The first step after identifying the presence of symptoms is to verify their relationship to hypoglycemia. A detailed clinical history must be performed to identify family or personal history of neuroendocrine tumors, if patients are taking any hypoglycemic medication such as sulfonylureas or if the symptoms are more common in fasting state, as in our case.

In a stepwise manner, biochemical analysis must be performed to rule out other causes[18]. Plasma glucose, insulin, C-peptide, proinsulin, beta-hydroxybutyrate, and cortisol levels should be measured. The development of provocative studies such as a 72-h fasting test is also recommended[10,18]. The goal is to determine whether beta-cell peptides are appropriately suppressed during hypoglycemia. If autonomous insulin secretion is identified, insulinoma should be suspected[10,18]. The next step is to determine the anatomical localization and to exclude other tumors. Multidetector contrast-enhanced imaging CT or dual phase helical CT with thin sections are the preferred initial imaging options. In patients in whom noninvasive radiologic techniques are negative or to improve the sensitivity for identifying an insulinoma, endoscopic ultrasound (EUS) must be performed. EUS has 80%-92% sensitivity for detecting tumors as small as 5 mm. Additionally, EUS-guided fine needle aspiration allows pathologic confirmation in 57% of patients. If the techniques mentioned above fail to detect the tumor, selective arteriography and intra-arterial calcium stimulation tests with hepatic venous sampling can be performed. They should be used only as a last resort because they are invasive techniques[5,10]. In our case, we performed CT and EUS that allowed us to identify insulinoma.

Finally, histopathologic and immunohistochemical confirmation is necessary to classify the type of tumor and to determine the patient’s follow-up[19].

The definitive treatment for insulinoma comprises complete surgical resection. However, there are other treatment options such as octreotide or EUS-guided alcohol tumor ablation, radiofrequency ablation (RFA), or embolization[20]. There is superior short-term recovery, shorter length of stay, decreased hemorrhage, and improved cosmesis when performing minimally invasive pancreatic resection compared to open pancreatic surgery[10]. However, the technique used depends on the size, extension, localization, and type of lesion. Atypical resection, including enucleation and partial or
middle pancreatectomy, has the advantage of pancreatic parenchyma preservation, thereby reducing the risk of late exocrine and/or endocrine insufficiency [20]. As in the case of our patient, when the lesion was small, benign, solitary, and superficial and when the pancreatic duct was not involved, the best surgical approach was laparoscopic enucleation [21]. It is important to note that positive resection margins are not associated with increased recurrence rates [10].

CONCLUSION

This is the first case of insulinoma after sleeve gastrectomy. Although this is a very rare case, clinicians must be aware of it, especially if the patient has hypoglycemic symptoms during the fasting state.

FOOTNOTES

Author contributions: Lobaton-Ginsberg M participated in the conception and design of the report and wrote the paper; Sotelo-González MP made substantial contributions to the acquisition, analysis, and interpretation of the patient data and helped write the paper; Júarez-Aguilar FG performed the histopathological and immunohistochemical report; Ramírez-Rentería C and Ferreira-Hermosillo A were involved in the coordination and design of the report and the revision of the manuscript; all authors read and approved the final manuscript.

Informed consent statement: Informed written consent was obtained from the patient for the publication of this report and any accompanying images.

Conflict-of-interest statement: The authors declare that they have no conflict of interest to disclose.

CARE Checklist (2016) statement: The authors have read the CARE Checklist (2016), and the manuscript was prepared and revised according to the CARE Checklist (2016).

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

Country/Territory of origin: Mexico

ORCID number: Miry Lobaton-Ginsberg 0000-0001-8227-087X; Pilar Sotelo-González 0000-0002-4132-5093; Claudia Ramírez-Rentería 0000-0003-3025-8339; Fany Gabriela Júarez-Aguilar 0000-0002-6204-3677; Aldo Ferreira-Hermosillo 0000-0002-5119-9856.

S-Editor: Chen YL
L-Editor: Wang TQ
P-Editor: Chen YL

REFERENCES


ectopic insulinoma: a rare case.

Liu J

Okabayashi T

gastric bypass.

Heterogeneity of proliferative markers in pancreatic

Patti ME

Sheehan A

10.3978/j.issn.2078-6891.2014.113

gastrinoma to insulinoma: a case report.

Chauhan A

Fasting ghrelin levels after gastric bypass and vertical sleeve gastrectomy: An analytic cohort study.

Navarro García MI

e0171601 [PMID: 32142259]

Wu HY

Li NS, Song YL, Bai CM, Wang Q, Zhao YP, Xiao Y, Yu S, Li M, Chen YJ. Plasma levels of acylated ghrelin in

Panousopoulos SG, Memos N, Menenakos E, Zografos G, Leandros E, Albanopoulos K. Laparoscopic sleeve
24185751 DOI: 10.1007/s00464-013-3275-y]

Service GJ, Thompson GB, Service FJ, Andrews JC, Collazo-Clavell ML, Lloyd RV. Hyperinsulinemic hypoglycemia
10.1056/NEJMoa043690]

Endocrinol (Oxf) 2019; 91: 906-908 [PMID: 31465534 DOI: 10.1111/cen.14083]

Koukoulis G, Sioka E, Zachari A, Zacharoulis D, Tzovaras G. Dumping symptoms and incidence of hypoglycaemia after
provocation test at 6 and 12 months after laparoscopic sleeve gastrectomy. Obes Surg 2012; 22: 1600-1606 [PMID:
22773085 DOI: 10.1007/s11695-012-0711-3]


e0171601 [PMID: 28152036 DOI: 10.1371/journal.pone.0171601]

Navarro García MI, González-Costea Martínez R, Torregrosa Pérez N, Romera Barba E, Periago MJ, Vázquez Rojas JL.

Wu HY, Li NS, Song YL, Bai CM, Wang Q, Zhao YP, Xiao Y, Yu S, Li M, Chen YJ. Plasma levels of acylated ghrelin in
patients with insulinoma and expression of ghrelin and its receptor in insulinomas. Endocrine 2020; 68: 448-457 [PMID:
32124259 DOI: 10.1007/s12020-020-02233-4]

Chauhan A, Ramírez RA, Stevens MA, Burns LA, Woltering EA. Transition of a pancreatic neuroendocrine tumor from

Sheehan A, Patti ME. Hypoglycemia After Upper Gastrointestinal Surgery: Clinical Approach to Assessment, Diagnosis,

Heterogeneity of proliferative markers in pancreatic β-cells of patients with severe hypoglycemia following Roux-en-Y

Okabayashi T, Shimah, Y, Sumiyoshi T, Kozuki A, Itos O, Ogawa Y, Kobayashi M, Hanazaki K. Diagnosis and management

Liu J, Zhang CW, Hong DF, Wu J, Yang HG, Chen Y, Zhao DJ, Zhang YH. Laparoscopic resection of retroperitoneal