Evidence-based Guidelines for the Management of Exocrine Pancreatic Insufficiency After Pancreatic Surgery

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Objectives: To provide evidence-based recommendations for the management of exocrine pancreatic insufficiency (EPI) after pancreatic surgery.

Background: EPI is a common complication after pancreatic surgery but there is certain confusion about its frequency, optimal methods of diagnosis, and when and how to treat these patients.

Methods: Eighteen multidisciplinary reviewers performed a systematic review on 10 predefined questions following the GRADE methodology. Six external expert referees reviewed the retrieved information. Members from Spanish Association of Pancreatology were invited to suggest modifications to the drafts; all members were consulted for their comments. The final version of the guidelines was reviewed by the members of the study group. All authors declare that neither they nor their institutions at any time have received payment or support in kind for any aspect of the submitted work (including grants, data monitoring board, study design, manuscript preparation, statistical analysis, and so on).

Disclosure: All the authors declare that neither they nor their institutions at any time have received payment or support in kind for any aspect of the submitted work (including grants, data monitoring board, study design, manuscript preparation, statistical analysis, and so on). L. S. and E. de-M. contributed equally to this work.

Results: These guidelines analyze the definition of EPI after pancreatic surgery (one question), its frequency after specific techniques and underlying disease (four questions), its clinical consequences (one question), diagnosis (one question), and how to treat postsurgical EPI (two questions) and its impact on the quality of life (one question). Eleven statements answering those 10 questions were provided: one (9.1%) was rated as a strong recommendation according to GRADE, three (27.3%) as moderate and seven (63.6%) as weak. All statements had strong agreement.

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Conclusions: EPI is a frequent but under-recognized complication of pancreatic surgery. These guidelines provide evidence-based recommendations for the definition, diagnosis, and management of EPI after pancreatic surgery.

Keywords: pancreatic exocrine insufficiency, surgery, pancreas, pancreatic, diagnosis, treatment, guidelines


Exocrine pancreatic insufficiency (EPI) is a common complication after pancreatic surgery. Depending on the underlying disease, type of surgical procedure, extent of pancreatic resection, and anatomical reconstruction, EPI may vary in frequency and severity. Despite the large amount of information dealing with general post-operative complications, there is a lack of well-designed studies investigating EPI. This has led to a certain degree of confusion about the frequency of EPI after surgery, its optimal methods of diagnosis and when and how to treat these patients. The aim of these guidelines is to provide evidence-based recommendations for the diagnosis and treatment of EPI after pancreatic surgery.

METHODS

The Spanish Association of Pancreatology (AESPANC) led the initiative and chose two coordinators (E. de-M. and L. S.) who developed the methodology. Eighteen Spanish primary reviewers were chosen, based on their expertise in pancreatic surgery, clinical pancreatology or nutrition (nine surgeons, eight gastroenterologists, and one endocrinologist). A group of external expert referees, composed by three pancreatic surgeons and three gastroenterologists, were invited to participate in the project. These referees were selected among internationally renowned researchers in pancreatology. A draft of the questions to be addressed was proposed by the coordinators and discussed by the whole team (via e-mail) finally resulting in 10 questions.

The coordinators assigned each question to two or three primary reviewers based on their expertise. A working plan for the systematic review was provided, inspired by the IAP/APA evidence-based guidelines for the management of acute pancreatitis.1 All reviewers were asked to take a GRADE system tutorial (link on UpToDate: http://www.uptodate.com/home/grading-tutorial).

The systematic research for suitable articles was performed in the PubMed and Cochrane databases without language restriction. The authors were provided with a search algorithm for each question (see supplementary material 1, http://links.lww.com/SLA/B4). In addition, studies from the citations of the reviewed articles could also be included.

The inclusion criteria to select the articles were as follows: observational studies, clinical trials, and meta-analysis/systematic reviews relevant to the specific question. Studies published only as abstracts were excluded.

The primary reviewers were asked to write a report including:

1. A table with a structured summary of the included studies (authors, journal, date of publication, design, population, definition of outcome variable, results, and comments).
2. An evidence-based statement to the study question.
3. The strength of the recommendation (1 = strong, 2 = weak) and quality of evidence (A = high, B = moderate, C = low) according to the GRADE guidelines as adapted for “UpToDate” (Table 1).
4. Remarks: a brief (up to 750 words) commentary explaining current evidence to support the recommendation.

The external expert referees were asked to review the report of the primary reviewers; their task was to check that:

1. There was no relevant study missing.
2. Included studies met the eligible criteria.
3. There was no mistake in the report of the included studies.
4. The strength of recommendation was adequate according to the retrieved evidence.

With the retrieved information by primary reviewers and external expert referees, the coordinators wrote a first draft of the manuscript. This draft was reviewed by the whole team and afterwards shared electronically with the members of AESPANC. The members of AESPANC voted on a five-point Likert scale (A: “definitely yes”, B: “probably yes”, C: “no specific recommendation”, D: “probably no”, and E: “definitely no”) on the statements and their GRADE score. It was defined that “strong agreement” would require at least 70% of votes to be either “definitely yes” or “probably yes”. The members of AESPANC also had the possibility of making suggestions in open text for every question, aiming not to modify the statement but to include clinically relevant remarks.

With the feedback from AESPANC members, the coordinators wrote the second draft of the article that was shared again with the primary reviewers and with the external expert referees for suggestions and final approval.

RESULTS

Question 1

What Is the Definition of EPI After Pancreatic Surgery?

Statement. EPI after pancreatic surgery is defined as the condition in which the amount of secreted pancreatic enzymes is not enough to maintain a normal digestion because of modifications of gastrointestinal anatomy together with functional changes caused by underlying pancreatic disease, extent of pancreatic tissue removed, reduced postprandial stimulation, and asynchrony between gastric emptying of nutrients and pancreatic enzyme secretion.

Strength of the Recommendation and Quality of Evidence: 1C. Strong Agreement (A: 87.5%; B: 12.5%)

Remarks. There is no widely accepted consensus definition of EPI, and there are no studies aiming to validate different EPI definitions with outcome variables after pancreatic surgery. Published studies addressing EPI after surgery have different definitions according to the different pancreatic function test (PFT) used in each particular study. From a pragmatic point of view, EPI may be defined as the situation in which the disturbance of pancreatic function is associated with the inability of the pancreas to perform normal digestion.4 Thus, an abnormally high fecal fat excretion (FFE) (>7/g/day) or a Coefficient of Fat Absorption (CFA) <93% (equivalent to a FFE >7/g/day under a diet containing 100g of fat/day) is characteristically indicative of EPI in clinical practice5,6 and should be considered as a gold standard. EPI after surgery may be secondary to a reduced pancreatic secretion caused by the underlying pancreatic disease,3,5 extent of pancreatic resection,7 reduced postprandial stimulation,8 and gastrointestinal anatomical changes leading to an asynchrony between gastric emptying of nutrients and enzyme secretion.9

Question 2

What Is the Frequency of EPI in Patients With Acute Pancreatitis After Pancreatic Necrosectomy?

Statement. The frequency of EPI in patients with acute pancreatitis after necrosectomy is variable because of significant heterogeneity in the design and population of available studies addressing this issue. Pancreatic function tends to improve and consequently frequency of EPI diminishes over time after necrosectomy. About a quarter of patients with acute necrotizing pancreatitis present EPI after pancreatic necrosectomy.
Table 1. Grading Recommendations

<table>
<thead>
<tr>
<th>Grade of Recommendation</th>
<th>Clarity of Risk/Benefit</th>
<th>Quality of Supporting Evidence</th>
<th>Implications</th>
</tr>
</thead>
<tbody>
<tr>
<td>1A. Strong recommendation, high quality evidence</td>
<td>Benefits clearly outweigh risk and burdens, or vice versa</td>
<td>Consistent evidence from well performed randomized, controlled trials or overwhelming evidence of some other form. Further research is unlikely to change our confidence in the estimate of benefit and risk</td>
<td>Strong recommendations, can apply to most patients in most circumstances without reservation. Clinicians should follow a strong recommendation unless a clear and compelling rationale for an alternative approach is present</td>
</tr>
<tr>
<td>1B. Strong recommendation, moderate quality evidence</td>
<td>Benefits clearly outweigh risk and burdens, or vice versa</td>
<td>Evidence from randomized, controlled trials with important limitations (inconsistent results, methodologic flaws, indirect or imprecise), or very strong evidence of some other research design. Further research (if performed) is likely to have an impact on our confidence in the estimate of benefit and risk and may change the estimate</td>
<td>Strong recommendation and applies to most patients. Clinicians should follow a strong recommendation unless a clear and compelling rationale for an alternative approach is present</td>
</tr>
<tr>
<td>1C. Strong recommendation, low quality evidence</td>
<td>Benefits appear to outweigh risk and burdens, or vice versa</td>
<td>Evidence from observational studies, unsystematic clinical experience, or from randomized, controlled trials with serious flaws. Any estimate of effect is uncertain</td>
<td>Strong recommendation, and applies to most patients. Some of the evidence base supporting the recommendation is, however, of low quality</td>
</tr>
<tr>
<td>2A. Weak recommendation, high quality evidence</td>
<td>Benefits closely balanced with risks and burdens</td>
<td>Consistent evidence from well performed randomized, controlled trials or overwhelming evidence of some other form. Further research is unlikely to change our confidence in the estimate of benefit and risk</td>
<td>Weak recommendation, best action may differ depending on circumstances or patients or societal values</td>
</tr>
<tr>
<td>2B. Weak recommendation, moderate quality evidence</td>
<td>Benefits closely balanced with risks and burdens, some uncertainty in the estimates of benefits, risks, and burdens</td>
<td>Evidence from randomized, controlled trials with important limitations (inconsistent results, methodologic flaws, indirect or imprecise), or very strong evidence of some other research design. Further research (if performed) is likely to have an impact on our confidence in the estimate of benefit and risk and may change the estimate</td>
<td>Weak recommendation, alternative approaches likely to be better for some patients under some circumstances</td>
</tr>
<tr>
<td>2C. Weak recommendation, low quality evidence</td>
<td>Uncertainty in the estimates of benefits, risks, and burdens; benefits may be closely balanced with risks and burdens</td>
<td>Evidence from observational studies, unsystematic clinical experience, or from randomized, controlled trials with serious flaws. Any estimate of effect is uncertain</td>
<td>Very weak recommendation; other alternatives may be equally reasonable</td>
</tr>
</tbody>
</table>

**Strength of the Recommendation and Quality of Evidence: 1C Strong Agreement (A: 45%; B: 55%)**

Remarks. There is a great heterogeneity in the design of studies addressing EPI after pancreatic necrosectomy. Some studies used FFE to assess pancreatic function and define EPI as FFE > 7 g/24 h. Gupta et al. reported increased FFE in six out of 21 patients (28.6%) at least 6 months after necrosectomy. Sabater et al. compared exocrine pancreatic function in patients with severe biliary AP with and without necrosectomy. Pancreatic function was assessed by FFE, fecal chymotrypsin and secretin-cerulein test (SCT). Twelve months after AP, seven out of 12 patients with necrosectomy (58.3%) had an abnormal PFT, with steatorrhea in three patients (25%). Reddy et al. reported increased FFE in eight out of 10 (80%) patients with necrosectomy, but no patient had symptoms of steatorrhea or EPI. Tsilotos et al. and Bavare et al. defined EPI with FFE, but it was only performed in patients with significant changes in bowel habit; thus, the prevalence of EPI could be underestimated.

Angelini et al. reported EPI (evaluated with SCT) in eight out of 20 patients with necrosectomy (40%) at 12 to 36 months and in 6.6% at 36 to 48 months after the onset of disease. Seligson et al. detected EPI in 7/10 (70%) patients with Lundh Test.

Other studies are hampered by important biases: the presence of acute and chronic pancreatitis or the inclusion of nonoperated patients. Finally, in some studies, EPI was reported on the basis of the need for pancreatic enzymes or clinical symptoms of steatorrhea, with figures between 23% and 25%, respectively. In this regard, it is noteworthy to highlight the results of the PANTHER trial from the Dutch Pancreatitis Study group, in which the minimal invasive step-up approach was significantly associated with a lower need for pancreatic enzymes than in primary open necrosectomy (7 vs 33%).

**Question 3**

**What Is the Frequency of EPI in Patients with Chronic Pancreatitis After Pancreatic Surgery?**

Statement. The incidence of EPI in patients with chronic pancreatitis after derivative surgery or hybrid procedures is the...
following: (i) after Partington-Rochelle procedure there are clinical steatorrhea and/or other clinical symptoms in 0 to 32% of patients and altered PFT in 80%; (ii) after Frey procedure there are clinical steatorrhea and/or other clinical symptoms in 33% of patients and altered PFT in 86%; (iii) after duodenal preserving pancreatic head resection (DPHPR) there are clinical steatorrhea and/or other clinical symptoms in 26 to 34% of patients and altered PFT in more than 80% of patients.

The incidence of EPI after pancreatectoduodenectomy (PD) for chronic pancreatitis is high, within the range of 35 to 100%, most of the studies showing >60%. The incidence of EPI after distal pancreatectomy (DP) for chronic pancreatitis seems to be lower, ranging from 27.5 to 63%.

As there is a high prevalence of EPI in chronic pancreatitis patients, and few studies evaluate EPI before pancreatic surgery, the specific contribution of the surgical procedure to EPI is difficult to quantify.

Strength of the Recommendation and Quality of Evidence: 1C Strong Agreement (A: 52.5%; B: 45%; C: 2.5%)

Remarks. The studies addressing EPI in patients with chronic pancreatitis after derivative surgery or hybrid procedures can be divided into five groups: articles comparing Partington-Rochelle versus PD22–25; articles comparing Frey versus PD26–28; articles comparing DPHPR versus PD20–36; articles comparing DPHPR versus Frey procedure37,38 and miscellaneous retrospective series.39–41 According to these studies, (i) after Partington-Rochelle procedure there are clinical steatorrhea and/or other clinical symptoms in 0 to 32% of patients and altered PFT in 80%; (ii) after Frey procedure there are clinical steatorrhea and/or other clinical symptoms in 33% of patients and altered PFT in 86%; and (iii) after duodenal preserving pancreatic head resection (DPPHR) there are clinical steatorrhea and/or other clinical symptoms in 26 to 34% of patients and altered PFT in more than 80% of patients.

Regarding the frequency of EPI in patients with chronic pancreatitis after resectional procedures (PD, DP), four prospective RCT30,31,33 (two of them with long-term follow-up), two prospective non randomized studies,26,27 two meta-analyses,2,25 and 10 retrospective studies,22–24,27,28,42–45 were included. The incidence of EPI after PD operation ranged from 35 to 100%. However, some concerns can be raised as to the quality of these findings. First, EPI was not the primary outcome in the majority of these studies, which were mainly designed to compare different surgical techniques. Furthermore, definition of EPI was not homogeneous and it seems that clinical definition (expressed by questionnaire or need for biliary and pancreatic secretions and are therefore difficult to be absorbed by the intestine; and (iv) the resection of duodenum which avoids the release of cholecystokinin57,58 and consequently there is a reduction of postprandial hormonal pancreatic stimulation.

Regarding DP, the incidence of EPI ranged from 27.5 to 63%. As mentioned before, the definition of EPI and the scarce preoperative assessment of pancreatic function can be considered strong biases.

Question 4

What Is the Frequency of EPI in Patients With Pancreatic Tumors After Resection (PD, DP)?

Statement. The incidence of EPI after PD for pancreatic tumors is high, especially in patients undergoing PD caused by malignancy, with a range of 64 and 100% The incidence of EPI after DP is lower than after PD, within a range of 0 to 42%.

Strength of the Recommendation and Quality of Evidence: 1C Strong Agreement (A: 67.5%; B: 32.5%)

Remarks. Information regarding the incidence of EPI in patients with pancreatic tumors after resection is limited and there is a lack of well-designed studies. Most available studies are retrospective and cross-sectional, limited by small sample size and single-institution designs. They also include a heterogeneous patient population with malignant and benign diseases. They include different types of surgery: PD, DP, and atypical resections. In addition, most of the reports include patients with and without chronic pancreatitis, and have used different methods to assess the pancreatic exocrine function. As mentioned before, malnutrition after PD has a complex pathophysiology that involves other factors besides the removal of pancreatic tissue (see questions 1 and 3).

Twenty-two studies have described the impairment of exocrine function after pancreatic head resection;4–6,49–57 14 studies included only patients who underwent PD,52,54,56–62,64–67 seven studies included both PD and DP4,49,50,55,55,63 (central or total pancreatectomy in three of them),50,53,55 and one study included PD and total pancreatectomy.51 Fourteen studies included a heterogeneous patient population with malignant and benign disease.6,50–54,56–60,62–64 five studies included only patients with malignant disease,4,5,49,61,66 and one study covered only benign tumors.55 There was also one meta-analysis.85

Among the studies, seven different methods for the assessment of EPI were applied (Table 2). As seen in this table, depending on the method used to evaluate exocrine pancreatic function, results vary considerably.

EPI rates varied widely from 24 to 100%. When only considering patients who underwent PD for malignant disease, EPI was present in 64 to 100%,5,49,61. Five studies9,55,58,63,64 have evaluated the preoperative and postoperative exocrine function. In the study by Sikkens et al,49 EPI was present in 44.8% at the time of diagnosis of pancreatic cancer increasing to 89% at the end of follow up. However, follow up was limited to 6 months, the long-term course was not evaluated, and the study covered two types of surgery (DP and PD). In
the series by Falconi et al\textsuperscript{55} including 51 PD for benign tumors with normal preoperative pancreatic exocrine function, EPI was observed in 33\% at the end of follow up. Matsumoto and Traverso\textsuperscript{58} reported a preoperative EPI rate of 33\% (68\% in pancreatic adenocarcinoma, and 46\% in malignant vs 21\% in benign disease), increasing to 73\% after 1 year. In the study of Sato et al\textsuperscript{63} the frequency of EPI increased from 44\% in the preoperative period to 81\% after pancreatic resection, but follow up was limited to only 2 months. One study\textsuperscript{66} suggested that postoperative impairment of pancreatic exocrine function was transient and reversible. EPI was present in 46\% preoperatively, rose to 75\% at the short-term (within 2 months), and then decreased to 33\% after 12 months, but this observation was based on data from only nine patients. Furthermore, the study included a heterogeneous group of patients with malignant and benign diseases.

Regarding DP, the incidence of EPI varied from 0 to 42\% depending on the method used to assess pancreatic exocrine function. Similar biases can be observed in the studies evaluating this procedure as in PD and in fact most of the studies include both PD and DP\textsuperscript{4,49,50,53,55,63} One study\textsuperscript{66} showed that most patients who underwent DP for benign or malignant pancreatic disease did not experience permanent postoperative EPI: all patients had normal exocrine function after DP or extended DP at 24 months after surgery and in the few cases where lower values were observed at 3 and 12 months after DP, the effect was transient. In the study by Falconi et al\textsuperscript{55} including 50 left pancreatectomies for benign tumors with normal preoperative fecal chymotrypsin levels, 18\% presented EPI at the end of follow up. Sato et al\textsuperscript{63} studied 12 patients who underwent DP for benign or malignant tumors of the pancreas and did not observe a significant decline in exocrine function after DP. Finally the meta-analysis by Xu et al\textsuperscript{68} showed an EPI rate of 10.8\% in DP.

Table 2: Variability in Exocrine Pancreatic Insufficiency After Pancreatoduodenectomy According to the Different Methods Used for Measuring EPI

<table>
<thead>
<tr>
<th>Method</th>
<th>EPI %</th>
<th>References</th>
</tr>
</thead>
<tbody>
<tr>
<td>Coefficient of fat absorption</td>
<td>55</td>
<td>4</td>
</tr>
<tr>
<td>Fecal fat excretion</td>
<td>87.5, 94</td>
<td>51, 62</td>
</tr>
<tr>
<td>13C-labelled mixed triglyceride breath test</td>
<td>64, 62.3, 51</td>
<td>6, 54, 67</td>
</tr>
<tr>
<td>Urinary PABA excretion rate</td>
<td>33, 75</td>
<td>64, 65</td>
</tr>
<tr>
<td>Fecal elastase 1</td>
<td>91, 59, 87.5</td>
<td>5, 49–51, 57, 50, 74.5, 100, 94.5, 97.5, 100</td>
</tr>
<tr>
<td>Fecal chymotrypsin levels</td>
<td>24, 33</td>
<td>53, 55</td>
</tr>
<tr>
<td>Clinical steatorrhea</td>
<td>52.8, 52.4, 42, 64.5</td>
<td>52, 56, 59, 66</td>
</tr>
</tbody>
</table>

EPI indicates exocrine pancreatic insufficiency; PABA, para-aminobenzoic acid.

Question 5

What Is the Frequency of EPI in Patients With Central Pancreatectomy? Statement

Central pancreatectomy is a conservative resectional procedure that is associated with low rates of EPI, approximately 10\%.

Strength of the Recommendation and Quality of Evidence: 1C Strong Agreement (A: 77.5\%; B: 17.5\%; C: 2.5\%; D: 2.5\%)

Remarks. Studies addressing EPI in central pancreatectomy (CeP) have two important shortcomings: (i) most studies do not report PFT in patients with CeP and most reports of EPI are based on clinical suspicion of steatorrhea and/or need for enzymes. Furthermore, the only two prospective studies\textsuperscript{70,71} did not perform PFT on patients with CeP.

Two studies reported FFE after CeP in patients with benign/low grade pancreatic tumors resulting in only one among 28 (3.6\%) patients with EPI\textsuperscript{72,73} In seven studies other PFT were performed after CeP\textsuperscript{53,55,74–78} reporting a range of EPI between 0\%\textsuperscript{74,76,77} and 21\%.\textsuperscript{78} Studies reporting clinical EPI (steatorrhea and/or weight loss and/or need for enzymes) describe a range between 0\%\textsuperscript{74,76,77} and 43\%.\textsuperscript{71} In a systematic review published in 2013, which included 21 studies, EPI (diagnosed either clinically or by means of diverse PFT) was noted in 9.9\% of the patients.\textsuperscript{69}

Question 6

What Are the Clinical Consequences of EPI?

Statement. EPI after pancreatic surgery may be subclinical or associated with symptoms secondary to the presence of undigested food in the intestinal lumen (fatty diarrhea, flatulence, and dyspeptic symptoms) and/or those associated with the loss of nutrients (weight loss, fat-soluble vitamin deficit).

Strength of the Recommendation and Quality of Evidence: 1C Strong Agreement (A: 77.5\%; B: 17.5\%; C: 2.5\%; D: 2.5\%)

Remarks. EPI after pancreatic surgery is associated with abnormal total energy absorption because of decreased digestion of fat, proteins, and carbohydrates.\textsuperscript{90} EPI may be subclinical or associated with two kinds of symptoms: those associated with the presence of undigested food within the intestinal lumen (fatty diarrhea, flatulence, dyspeptic symptoms)\textsuperscript{91} and those associated with the loss of nutrients (mainly weight loss and fat-soluble vitamin deficit). The pancreas is involved in the digestion of proteins, carbohydrates, fat, and other nutrients, but pancreatic lipase is so essential for fat absorption that most of the clinical consequences of EPI are related to fat malabsorption. The extent of malabsorption depends on the original disease process and the type and extent of surgical resection.\textsuperscript{92} The main clinical manifestation of fat malabsorption is steatorrhea typically reported as an increase in bowel movements, particularly after fatty meals, with loose, greasy, foul-smelling voluminous stools.\textsuperscript{47,93} Steatorrhea, however, may be not present or present because of another cause. Postprandial abdominal pain and abdominal bloating may also be associated with EPI.\textsuperscript{47}

In patients with untreated EPI, potential additional complications such as weight loss, poor wound healing, vitamin deficiencies, osteomalacia, osteoporosis, and low-trauma fractures, electrolyte imbalance, increased adverse effects of oncological treatments, and lethargy can theoretically appear. One study compared pancreatic enzymes and placebo after surgery for chronic pancreatitis; four out of five patients receiving pancreatic enzymes gained weight but none of those six patients receiving placebo did.\textsuperscript{90} Apart from weight loss, there are no specific studies demonstrating a different nutritional status in patients with or without EPI after pancreatic surgery.\textsuperscript{4,49}

Question 7

What Is the Optimal Method for the Diagnosis of EPI After Pancreatic Surgery?

Statement. PFT are of limited clinical value after pancreatic surgery as the prevalence of EPI is high and PFT are either difficult to

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perform or have poor predictive values. In cases when objective evidence for EPI is needed, FFE may be considered as the gold standard. Human elastase-1 (FE1) is easy to perform, has a high sensitivity to detect steatorrhea but its specificity seems lower. The 13C-MTG might be an alternative method but further studies are needed. The absence of clinical symptoms of steatorrhea is an inaccurate method to rule out the existence of EPI.

**Strength of the Recommendation and Quality of Evidence: 2B Strong Agreement (A: 35%; B: 40%; C: 22.5%; E: 2.5%)**

Remarks. Currently FFE/CFA may be considered as a gold standard for EPI (see question 1). Unfortunately, this technique is cumbersome to perform: it requires a specific diet with a given amount of fat per day and stools from 3 days must be collected and processed. For these reasons, it would be very useful to have simpler PFTs like FE-1 and/or 13C-MTG but few studies have tried to validate them for the diagnosis of steatorrhea by means of FFE or CFA after pancreatic surgery.  

Halloran et al studied 40 operated patients for pancreatic cancer (37 PD and only three left pancreatectomies) by FE-1 and CFA. A comparison of FE-1 using a cut-off point of 200 mcg/g for EPI against CFA showed a diagnostic accuracy of 70%, with a sensitivity of 91%, a specificity of 35%, a positive predictive value of 70%, and a negative predictive value of 71% for FE-1. There was no clear association between CFA and FE-1 levels. Overall, this study suggests the limited accuracy of FE-1 to diagnose EPI after pancreatic surgery. In another study Benini et al studied 40 operated patients (37 pylorus preserving PD, one Whipple procedure, and two total pancreatectomies) and 42 nonoperated patients with pancreatic diseases, and evaluated EPI by FE-1 compared with FFE. Sensitivity and specificity of FE-1 in operated patients to detect steatorrhea were as follows: 100% and 83.3% for FE-1<200 mcg/g; 100% and 100% for FE-1<100 mcg/g and 61.8%, respectively, and 100% for FE-1<15 mcg/g. The cut-off for FE-1 in the diagnosis of EPI was considerably higher in operated compared with nonoperated patients. Another conclusion of this study is that the relationship between both tests is not linear but logarithmic. The rate of increase of 24 hours fecal fat output with decreasing FE-1 levels is not constant but depends on FE-1 values, with rates much higher when FE-1 values are low. The information regarding the correlation between FE-1 and FFE in left pancreatectomy is lacking. Nakamura et al investigated the usefulness of 13C-MTG compared with FE-1 concentration, but they used clinical steatorrhea as a gold standard. According to their results, the 13C-MTG might be more useful than the FE-1 for the diagnosis of EPI after pancreatic surgery because of its higher accuracy, which could be explained by the fact that fecal water content influences the fecal enzyme concentration, resulting in falsely decreased FE-1 levels. The advantages and disadvantages of the different available PFT are shown in Table 3.

To sum-up we need more studies to validate the use of FE-1 (which was associated with a poor correlation with FFE in two studies and poor accuracy for the diagnosis of steatorrhea in one of them) and 13C-MTG in surgical patients. In this scenario, the diagnosis of EPI may be assumed in patients with symptoms suggesting malabsorption. On the other hand, the absence of clinical symptoms of steatorrhea is not an accurate method to exclude the existence of EPI, and therefore PFT may play a role in the diagnosis of EPI in asymptomatic patients.

**Question 8**

**When Should EPI Be Treated? Statement**

Pancreatic enzyme replacement therapy should start once EPI is diagnosed or when there is a high clinical suspicion of EPI.

**Strength of the Recommendation and Quality of Evidence: 2B Strong Agreement (A: 72.5%; B: 25%; C: 2.5%)**

Remarks. There is a paucity of high quality trials specifically designed to assess when to treat EPI in patients with previous pancreatic surgery. Most recommendations come from expert opinion or guidelines from medical societies. The incidence of EPI associated with different surgical techniques, its clinical consequences, and diagnosis have been addressed in specific questions in this review. As a summary, deterioration of pancreatic function frequently occurs after pancreatic surgery; this condition is associated with relevant consequences. In patients with pancreatic surgery and EPI, pancreatic enzyme replacement therapy improves the CFA, the coefficient of nitrogen absorption, and reduces flatulence, diarrhea, and abdominal pain and therefore EPI should be treated as soon as it is diagnosed. However, the task of establishing the diagnosis of EPI in patients with previous pancreatic surgery does not have a straightforward approach. To overcome this limitation in patients with a high clinical suspicion of EPI, its diagnosis may be accepted after an empiric therapeutic trial showing that symptoms, nutritional markers or body weight improve after pancreatic enzyme replacement therapy.

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**TABLE 3. Advantages and Disadvantages of the Main Different Available Pancreatic Function Tests**

<table>
<thead>
<tr>
<th>Pancreatic Function Test</th>
<th>Advantages</th>
<th>Disadvantages</th>
</tr>
</thead>
<tbody>
<tr>
<td>Fecal fat excretion/ Coefficient of Fat Absorption</td>
<td>Clinically relevant</td>
<td>Very cumbersome and difficult to perform</td>
</tr>
<tr>
<td>Fecal elastase-1</td>
<td>It detects other causes of maldigestion Useful for monitoring response to treatment</td>
<td>Not widely available</td>
</tr>
<tr>
<td>13C-labeled mixed triglyceride breath test</td>
<td>Theoretically it detects other causes of maldigestion Probably useful for monitoring response to treatment</td>
<td>Expensive Scarcely available</td>
</tr>
</tbody>
</table>

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Question 9

How Should EPI be Treated and How Should Follow-up Be Performed?

Statement. EPI after pancreatic surgery should be treated with pancreatic enzyme replacement therapy with pancreatin in form of enteric-coated minimicrospheres. Enzyme doses of 72,000–75,000 Ph.U. of lipase with main meals and 36,000–50,000 Ph.U. with snacks have shown to be effective in terms of improvement in fat digestion.

Strength of the Recommendation and Quality of Evidence: 1A. Strong Agreement (A: 70%; B: 27.5%; D: 2.5%)

Statement B. Follow up should be based on symptoms and nutritional evaluation, including body weight and routine nutritional parameters in blood.

Strength of the Recommendation and Quality of Evidence: 2C. Strong Agreement (A: 70%; B: 27.5%; D: 2.5%)

Remarks. Treatment of EPI after any pancreatic surgical procedure should be based on oral pancreatic enzyme replacement therapy. Enzyme doses of 72,000–75,000 Ph.U. of lipase with main meals and 36,000–50,000 Ph.U. with snacks have shown to be effective in terms of improvement in fat digestion in RCTs. Only two double-blind RCTs evaluating pancreatic enzyme replacement therapy for EPI in patients after pancreatic surgery have been reported. An open-label long-term follow-up study was reported, with the patients from the double-blind study previously published by Whitcomb et al. In these two latter studies, results of operated patients are reported together with nonoperated patients with chronic pancreatitis, but the study from Seiler et al. only addresses operated patients, which also includes data from open-label pancreatic enzyme replacement therapy administration for 1 year.

Compared with placebo, pancreatic enzyme replacement therapy with pancreatin in form of enteric-coated minimicrospheres is associated with a significant improvement of fat (CFA) and protein digestion (coefficient of nitrogen absorption) in patients after pancreatic resection for chronic pancreatitis or pancreatic cancer. In addition, pancreatic enzyme replacement therapy is associated with a significant improvement of fat (CFA) and protein digestion (coefficient of nitrogen absorption) in patients with pancreatic cancer. In addition, pancreatic enzyme replacement therapy is associated with a significant weight gain and reduced stool frequency.

No study has been published which specifically focused on dietary advice for patients after pancreatic surgery, it seems reasonable that a normal healthy diet should be generally recommended if tolerated.

No study has been found to answer the question about the follow up of EPI in patients after pancreatic surgery. In our opinion, follow up should be based on symptoms and nutritional evaluation, including body weight and some routine nutritional parameters in blood (eg, albumin, fat-soluble vitamins). Frequency of visits should be defined depending on the clinical and nutritional status of patients. Once the therapy has been optimized and the clinical and nutritional evaluation is normal, further follow up should probably be on-demand.

Question 10

What Is the Quality of Life in Operated Patients With EPI?

Statement. Exocrine pancreatic insufficiency is a relevant prognostic factor related to impaired quality of life in patients who undergo pancreatic surgery.

Strength of the Recommendation and Quality of Evidence: 1B Strong Agreement (A: 72.5%; B: 25%; C: 2.5%)

Remarks. Quality of life deteriorates in patients who develop EPI after pancreatic resection. In addition, the development of postoperative EPI is a relevant prognostic factor, significantly affecting the postoperative quality of life. Patients undergoing surgery caused by pancreatic cancer with EPI score lower on quality of life and functional scores. Long-term follow up survivors are generally satisfied with their quality of life, but bowel function, steatorrhea, need for treatment of diarrhea, or need for pancreatic enzyme replacement therapy and food intolerance may impair quality of life. Patients with benign pancreatic tumors had higher quality of life values at all time points compared with patients with pancreatitis and cancer; however, it is interesting to point out that quality of life in this group did not reach normal values for the healthy population even late after surgery, although these patients underwent curative therapy and did not suffer per se a chronic pancreatic disease. This is probably because of a higher rate of postoperative EPI. Total pancreatectomy (TP) has a deep influence on short- and long-term changes in the quality of life and EPI appears to be an important factor because it especially affects symptom scales. In acute pancreatitis, patients with long-term survival after surgical treatment for infected pancreatic necrosis have a quality of life comparable with that of the normal population. In patients with chronic pancreatitis requiring surgery, quality of life improves significantly both in the short and long term. Although the number of patients with exocrine insufficiency is very high in CP, in some studies such complication does not seem to have relevance in the overall reported quality of life. Finally, in operated patients, regardless of the disease requiring pancreatic resection, the type of reconstruction technique does not affect their well being, and also among different types of pancreatic head resection, the majority of functional and symptom scales revealed a better quality of life and less steatorrhea in duodenum-preserving pancreatic head resection.

CONCLUSIONS

Pancreatic surgery is still a great challenge as it is frequently associated with immediate surgical complications and long-term sequelae. EPI is a frequent but under-recognized and under-treated complication of pancreatic surgery. The lack of awareness and information regarding the frequency, diagnostic methods, and recommended therapy prompted the Spanish Association of Pancreatology to design the present systematic review. EPI is commonly observed after pancreatic surgery, it is clinically relevant and affects quality of life; thus, it should be investigated, treated, and followed-up appropriately. The most important limitation of the literature and the origin of much of the confusion on this topic is that the diagnosis of EPI depends on the definition and the method used for measuring EPI and there is a lack of studies trying to validate PFT with a proper gold standard in operated patients. Therefore, further research is needed to look for better and simpler diagnostic tools.

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REFERENCES


