INTRODUCTION

Pain is the dominating symptom of chronic pancreatitis (CP) causing impairment of quality of life, decreased activity, unemployment and major healthcare costs. Opinions differ on the best strategy to treat CP-related pain. These differences are fuelled by the lack of a clear correlation between the severity of complaints and the presence and extent of morphological abnormalities, lack of a comprehensive pain assessment tools, deficiency in the knowledge about the natural course of CP and the presence of (changes in) both peripheral and central nervous system pain mechanisms.

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role in painful CP, and the severity of neuropathy is correlated with pain intensity. Neuropathic changes have been discovered at all levels of the nervous system (figure 1). Sensitisation of central nociceptive pathways is accelerated by acute disease flares and painful comorbid conditions. Hence, continued nociceptive input on central pathways may ultimately result in an autonomous and self-perpetuating pain state, which is independent of the peripheral nociceptive drive. When this state develops, it is unlikely that any local neuroablative procedure will be effective.

In support, generalised hyperalgesia in patients with painful CP, as objectively measured with quantitative sensory testing (QST), was associated with failure of thoracoscopic splanchnic denervation. The argument for invasive treatments of neuropathic pain is that removal of the source of nociceptive pain (eg, by pancreatic resection) would decrease the barrage from primary afferents and, over time this would reduce pain severity. This has some credence as neural blocks and local analgesics can be effective for peripheral nerve injury and painful polyneuropathy. However, while medical treatment aims to reduce the membrane potentials and firing thresholds of pathological neural circuits, there is evidence that neuroablative procedures (after initial alleviation of pain) can promote spontaneous firing of peripheral afferents due to disinhibition and upregulation of ion channels. This can lead to a cascade of central nervous system events followed by sensitisation and worsening of pain. Hence, procedures based on neural destruction may promote development of new neuropathic pain, formerly known as ‘deafferentation pain’. Taken together, these mechanisms may explain the reduced long-term efficacy of invasive treatments.

**What can surgeons learn from invasive treatments in other painful conditions?**

For lack of other options, surgeons have historically attempted to excise chronic pain. Although not studied after pancreas surgery, consequences after nerve damage are dependent on procedure rather than disease, and therefore, the experience from invasive treatment in other diseases is valid in this context. Destructive, irreversible neurolytic procedures have included cauterisation of the sciatic nerve for sciatica, amputations for limb pain, dorsal root entry zone rhizotomies and resection of neuromas for phantom limb pain. Such approaches have failed and the lack of controlled clinical data has meant that these techniques have become historical anecdotes. Despite this, the continuing support for pancreatic resection for chronic pain persists in the treatment of CP and has been given further impetus with popularisation of total pancreatectomy with islet auto transplantation (TPIAT), including resection even before the onset of pain.

Some surgical treatments for chronic pain can be considered neuroablative. A common example is microdiscectomy for the management of radicular pain. Although many patients with sciatica report pain relief in the first 3 months, most still had mild to moderate symptoms 5 years after surgery. Another...
example of this is revisional surgery after total knee arthroplasty, where 85% of revision patients have more intense pain and sensitisation compared with their baseline levels. More recently, neuromodulation has been developed as a less invasive treatment option, including spinal cord stimulation and targeted intrathecal pharmacotherapy (eg, morphine pump). While these are considered relatively safe and produce significant pain reduction, there is a lack of randomised placebo-controlled trials to support it.

**Flaws in studies of invasive treatments for painful chronic pancreatitis**

Treatment by endoscopy and/or surgery are commonly offered as part of a step-up approach, when lifestyle changes and drug treatment fails. Many observational studies or studies comparing different procedures have reported benefits from endoscopic and surgical treatment of CP pain. However, these symptomatic benefits are time dependent, and it is common for relapse to occur over time. This relapse may be due to progression of the neuropathy, development of further duct strictures and/or fibroinflammatory processes, as well as the loss of the placebo effect of intervention.

Studies related to neuroablative procedures are often biased by patient selection, have failed to control for differing aetiologies and definitions of CP have a short follow-up period, and/or use poor pain assessment methodology. For example, celiac plexus neurolysis improves analgesia and decreases opioid consumption in patients with pancreas cancer, but benefits are time limited (about 3 months). As the nerve damage may even worsen the neuropathic mechanisms as outlined above, it should be considered obsolete in patients with painful CP. TPIAT involves denervation and is becoming increasingly available. Although open studies have shown that both total pancreaticectomy (alone) and TPIAT can reduce pain, there are significant issues with risk, cost and consequences of such an invasive treatment. Some of the benefits may derive from better monitoring and control of diabetes and greater attention to nutritional management. Additionally, a critical follow-up study has showed that most TPIAT patients develop functional pain and other pain syndromes, and caution is advised to promote this treatment outside protocolled studies.

**Sham-controlled studies are mandatory to confirm the effect of invasive treatments**

The well-known placebo effect has also been investigated in pharmacological pain management of CP. The effect size is about 20%, and this is similar to what has been found in studies with analgesics for other conditions, including painful gastrointestinal diseases. As no placebo/sham-controlled trials of invasive therapies for painful CP exist, previous studies may be subject to selection bias and the effects on pain per se are still unanswered (figure 2). Due to lack of studies in CP, it is worthwhile to consider the effect of sham-controlled surgical trials for other painful conditions. Examples include laparoscopic adhesiolysis for abdominal pain, arthroscopic meniscectomy for knee pain, subacromial decompression for shoulder pain, percutaneous coronary intervention for angina and vertebralplasty for painful vertebral compression fractures. Although these procedures are well established and the indication for pain has not been questioned (with results similar to surgery for CP (table 1)), all the studies demonstrated no difference in pain intensity between active treatment and sham. This was confirmed in meta-analyses of invasive procedures for painful conditions (other than CP), where improvements in the sham arm were seen in up to 75%. Another more recent meta-analysis confirmed that the effect of sham surgery on pain was about 40%, with studies reporting up to 60% pain improvement. More specifically for pancreatolgy, Cotton et al randomised 214 patients with sphincter of Oddi dysfunction to sphincterotomy or sham. There was reduction in pain in both groups, but the sham treatment performed significantly better. Later, the authors reported that the 47% of the sham-treated patients had >50% reduction in pain and 37% had no pain at all. This study has questioned the use of invasive treatments in patients with pancreatogenic pain. It is worth noting that the effect size of sham treatment in these studies is of a similar magnitude to that seen in studies investigating endoscopic and surgical treatment for painful CP. Unfortunately, the ethical challenges of conducting sham-controlled studies in surgery (open or laparoscopic) in CP are well recognised, but in contrast, it is possible to conduct sham-controlled endoscopic treatments.

**THE PROS OF INTERVENTIONAL TREATMENT OF PAIN IN CHRONIC PANCREATITIS**

In the recent international evidence-based (HaPanEU) guidelines, a multidisciplinary step-up approach for the treatment of pain in CP was recommended, although specific trials investigating this approach are yet to be performed. The first step is conservative therapy, including lifestyle management (eg, cessation of alcohol use and smoking), dietary advice and pain medication. The approach to pain management of the WHO is widely accepted for pain treatment in CP, although it has not been formally evaluated. If patients have persistent pain despite appropriate conservative measures, including optimisation of pain medication, subsequent interventional endoscopy or surgery is
recommended. While endoscopy is often used as the first approach in patients with solitary strictures and single pancreatic duct stones in the head of the pancreas, there are also indications for surgery as the first intervention, such as an inflammatory head mass in which cancer cannot be excluded. There is a large literature, both non-randomised and randomised, supporting the efficacy of interventional endoscopy and surgery in the treatment of symptomatic CP.\(^{13, 26, 61}\)

### Interventions in chronic pancreatitis: what we know

#### Medical therapy

Around 50% of patients with CP require pain medication on a daily basis, and the vast majority take opioids, despite the lack of evidence that they are effective for long-term pain management.\(^{62, 63}\) Further, the longer patients are treated with opioids, the greater is the risk of opioid dependence and side effects such as opioid-induced constipation, cognitive dysfunction and hyperalgesia.\(^{64, 65}\) As a result, there is acceptance that the lowest effective dose of opioids should be used.

Patients with long-standing CP can develop changes in the central nervous system leading to pain that is less dependent on peripheral driving factors (nociception from local inflammation of the pancreas).\(^{66}\) To treat this ‘central sensitisation’ and reduce the effective opioid dose, neuropathic pain medication (eg, gabapentin or amitriptyline) is widely used in clinical practice in conjunction with other analgesics and interventional treatment in CP. Only one trial investigated this strategy in the short term (see table 1), and long-term efficacy is unknown.\(^{67}\) What is important to note, especially in the absence of compelling evidence, is that all analgesic strategies are adjunctive because they do not specifically target the drivers of painful CP. All patients requiring opioid analgesia for CP should be discussed at a multidisciplinary meeting to determine whether they should be referred for interventional endoscopy or surgery.\(^{1}\) A large proportion of these patients will have morphological abnormalities such as a ductal obstruction with upstream dilatation due to ductal strictures, ductal stones, inflammatory pancreatic (head) mass and/or pseudocysts.\(^{32, 68, 69}\) Although the correlation between pain and the presence and extent of morphological abnormalities is highly variable, several RCTs have confirmed pain relief attributable to the interventions in the majority of these patients, see table 1.\(^{1, 2}\)

#### Interventional endoscopy

There are a number of morphological abnormalities that can be treated by interventional endoscopy on the premise that the intervention reduces pancreatic duct obstruction and hypertension. These interventions include pancreatic and biliary stricture dilation/stenting, pancreatic duct stone removal and drainage of pseudocysts.\(^{70}\) Several observational cohort studies have reported a long-term pain relief in substantial proportions of patients following interventional endoscopy. Two studies of 57 and 1018 patients reported pain relief in >60% of the patients after 1 and 5 years, respectively.\(^{13, 52}\) In patients with large intraductal stones (>5 mm), ESWL and subsequent endoscopic stone extraction can be effective.\(^{70}\) There is one RCT in which ESWL with and without subsequent endoscopic stone extraction was compared in 55 patients. After 2 years, the pain relief was comparable between both groups (53% vs 62%).\(^{72}\) See table 1 for a structured overview of studies mentioned above.

### Surgery

Surgery is effective both in patients with a ductal obstruction and an inflammatory head mass. The most commonly performed procedure is ductal drainage by a lateral pancreaticojejunostomy, in which the pancreatic duct is opened along its entire length and anastomosed (side to side) to a Roux-Y jejunal limb.\(^{73}\) An inflammatory head mass can be resected using pancreaticoduodenectomy or a duodenum preserving pancreatic head resection such as the Berne, or Beger procedures.\(^{74, 75}\) Ductal drainage and duodenum preserving (partial) head resection are combined in a Frey procedure.\(^{76}\) Surgery is reported to have a good long-term effect although approximately 10% of patients will not respond, and this risk is higher in patients who have had a large number of endoscopic procedures prior to surgery.\(^{77}\)

Two RCTs have compared endoscopic and surgical management in patients with painful obstructive CP.\(^{13, 64}\) Both studies showed superiority of surgical over endoscopic management. The study of Dite et al randomised 72 patients with CP and reported, after 5 years of follow-up, 34% complete and 52% with partial relief after surgery, versus 15% who had complete pain absence and 46% partial relief after endoscopy. However, endoscopy in this study was not optimal since no ESWL was applied.\(^{61}\) The study by Cahen et al randomised 39 patients with painful obstructive CP and reported, after 2 years of...
of follow-up, 75% pain relief after surgery versus 32% after endoscopy, which included ESWL (p=0.007). In long-term follow-up, these results remained stable after >6 years, with 80% pain relief in the surgery group versus 38% pain in the endoscopy group. Complete pain relief was seen in 53% of patients after surgery versus 25% after endoscopy. It was also noted that 47% (9/16) of the endoscopic-treated patients required delayed surgery, and complete pain relief was found in only 2/9 (22%) patients with delayed surgery compared with the 8/15 (53%) who had primary surgery. Several RCTs that compared different surgical strategies for CP have reported long-term pain relief of around 80% after 7–15 years follow-up. The most recent international RCT compared duodenum-preserving pancreatic head resection with pancreatoduodenectomy in 250 patients with CP and an inflammatory pancreatic head mass. Two years after surgery, pain scores and the mean quality of life significantly improved in both groups, without significant differences.

An evolution in the surgical intervention for CP is TPIAT, first reported in 1977. As conventional surgery and endoscopic treatment may not be feasible or successful in up to 50% of patients, TPIAT is based on the concept of complete removal of the source of pain while maintaining islet cell function to avoid or mitigate the risk of diabetes by infusing the patient’s own islet cells into the portal circulation to engraft within the liver. Application of TPIAT is growing worldwide and it might be considered appropriate for pediatric patients, those without pre-existing diabetes, those with hereditary pancreatitis and those with small duct disease for whom decompression is not possible. A single-centre observational study has reported on 215/742 TPIAT patients with at least a 10-year follow-up. Complete and sustained pain relief was present in 82% at 10 years and 90% at 15 years, with either full or partial islet graft function in nearly 50% of patients.

Based on the available evidence, both surgery and endoscopy are durable (and not time dependent as suggested by the con group), although surgery provides the superior outcome (table 1). There are some patients who will benefit from an endoscopy-first approach, including those with (single) pancreatic duct stones or strictures confined to the head of the pancreas in whom complete ductal clearance can be achieved. While this is a reasonable approach, there is a risk of delay in referral for surgery when the outcome of endoscopy proves unsatisfactory. An increasing body of evidence suggests that earlier surgical intervention leads to better pain control.

After the onset of pain, a (long) delay before invasive treatment increases the risk of opioid dependence and may lead to sensitisation of central pain pathways and changes in pain perception, which increases the chance of treatment failure. On this basis, the CON group actually favour earlier intervention in painful, obstructive CP.

## Interventions in chronic pancreatitis: what we do not know

### Timing of intervention

In current clinical guidelines, there is inconsistency regarding the timing and whether endoscopy or surgery should be offered as the first intervention for patients with painful obstructive CP. There is increasing evidence that suggests that earlier surgery leads to better pain relief rather than the unproven step-up approach, in which medication and then endoscopy are offered first.

It is known that delay in intervention, prolonged and repeated episodes of pain and opioid dependency are associated with the development of neuropathic pain. To resolve this longstanding debate about the timing of intervention, the Dutch Pancreatitis Study Group conducted the ESCAPE trial. Patients with painful obstructive CP and a dilated pancreatic duct in a very early phase of treatment, with a maximum of 2 months of opioid use and no prior interventions, were eligible. These patients were randomised to either surgical treatment within 6 weeks or an optimised current step-up approach including endoscopic treatment. Primary outcome was pain as assessed by the Ibzik pain score, a validated pain score specific for CP and widely used. The preliminary results of the ESCAPE trial were presented at United European Gastroenterology Week 2018 but final results are still awaited.

### Sham-controlled trials: good in theory, problematic in practice

Although all endoscopic and surgical intervention trials in CP have shown pain relief in a clinically relevant proportion of patients, none of them used sham interventions. Trials with a sham intervention as comparator can estimate the placebo effect of an intervention, something of particular interest with studies of pain treatment. There is indeed evidence that the placebo effect can account for up to 65% of the treatment effect for invasive procedures, especially when treating pain. Sham-controlled studies, however, are subject to debate for both ethical and practical reasons.

A sham intervention might be considered as dishonest in clinical practice, whereas it is a legitimate aspect of research design. There is an inherent difficulty in incorporating a blinded sham-control arm in surgical procedures, although this can be done more easily in endoscopic studies. It is also difficult for clinicians to deny treatment that they consider effective and when not convinced that equipoise exists, even in the context of an RCT, such as trial requires long-term follow-up, and many would consider it unethical to deny potentially effective treatment for years. Lastly, the effect size of endoscopic and surgical treatment is such that it seems highly improbable that this could be accounted for by a sham intervention alone. Is it ethical to withhold patients from other treatment options for such a prolonged period, even if these other options have been shown to be successful without sham comparison?

## FACTORS THAT INFLUENCE THE OUTCOME FROM INVASIVE TREATMENT: RECOMMENDATIONS FOR FUTURE TRIALS

Advances in the treatment of patients with painful CP will require an improved study design that takes into account the many risk factors that influence outcome (table 2). Studies have shown that demographic variables and imaging features are important in the outcomes from surgery. Alcohol and smoking also influences outcomes, and alcohol abstinence can improve the effect of surgery. Compared with hereditary or idiopathic CP, patients with toxic aetiologies have improved outcomes from resectional or decompressive procedures, but worse after TPIAT. This implies that the aetiology of CP is a more important factor than any morphological changes of the pancreas, and future trials should more carefully consider the effect of smoking and drinking continuance/cessation. Other factors including depression, widespread pain, expectations, catastrophizing and sleep deprivation may also be important although less well studied in CP.
Pain evolution has a significant impact on the outcomes from invasive treatment. Observational studies of surgical and endoscopic therapy have shown improved outcomes when performed during the early phase of CP, especially prior to the onset of narcotic dependence and central sensitisation. The previously mentioned ESCAPE trial may provide data on this factor.

Pain pattern is another important factor to consider. CP patients with constant pain pattern often have a worse quality of life, more lost work days, increased number of hospitalisations, higher pain intensity and more night pain. Constant pain was also associated with higher levels of exocrine insufficiency and malnutrition. Finally, constant pain is a predictor of failed surgical treatment for painful CP.

On the other hand, intermittent pain associated with recurrent acute pancreatitis is a positive predictor in patients having TPIAT. Correspondingly, TPIAT is more effective in the paediatric CP patients, which might be explained by an immature pain system that is less likely to undergo the irreversible neuroplastic changes observed in adult patients. The effect of pain pattern is supported by studies in patients with other painful neuropathic conditions, where a constant pain pattern was also associated with a worse outcome from treatment.

Opioid use is associated with more hospitalisations, and is a predictor of failure from surgical treatment. Opioids can interfere with pain processing, and in some instances, contribute to hyperalgesia. In an individual patient, it is difficult to determine whether opioids worsen the response to treatment. Patients with opioid dependence are generally those with more severe pain and lower quality of life, and as such are more likely to be refractory to treatment.

Subjective pain assessment is highly variable and the instruments are poorly validated. This is the primary indication for the treatment of patients with CP and the most important endpoint of trials. The ‘IMMPACT guidelines’ recommend that patient factors, study design, study site factors and outcome measurements are all used in pain evaluation, and patient-reported characteristics should also be used in assessing the response to invasive treatments.

As most invasive treatment studies have used only simple pain assessment instruments, it is mandatory to use better validated and more comprehensive measures in future trials.

Objective pain assessment includes QST and can be combined with subjective pain assessment, neurophysiological and imaging methods. Visceral QST has been used in the evaluation of many pain conditions, but the invasiveness has limited its use in clinical practice. Visceral QST has also been used in CP to explore different pain mechanisms. A bedside solution that assesses somatic pain sensitivity as a proxy for the changes in the central nervous system caused by CP has recently been developed and validated.

This has been shown to reflect disease severity and to predict the outcome of medical and surgical treatment for painful CP. However, further studies, including normative reference data and additional validation in CP patients without pain, are required before this approach can be used in clinical practice.

**CONCLUSION**

Pain is the predominant and most incapacitating symptom of CP, and current guidelines recommend treatment by a step-up approach comprising an initial pain medication, followed by interventions including endoscopy and surgery. However, as highlighted in this debate article, the conceptual basis for these interventions (ie, reducing pressure in pancreatic duct, removing ductal stones and resecting inflammatory masses to remove local nociceptive source) does not take into account the advances in our understanding of pain mechanisms, and in particular, the importance of neuropathy and neuroplastic changes in the central nervous system. Furthermore, there is a paucity of data from prospective, randomised and adequately controlled clinical trials as basis for the current interventional treatment recommendations. Although showing that interventions improve pain in CP, most randomised trials compare different types of interventions or the timing of interventions, rather than questioning the indication for interventions to treat pain in CP, an assumption that may indeed be challenged, as described by the ‘con-authors’ in this debate. Furthermore, many trials do not take into account that CP is heterogeneous in many factors such as aetiology, stage, duration and pre-treatments, and thus different patients may require different treatment approaches, making a point for the personalisation of algorithms for the treatment of pain in CP. To make progress towards a better understanding of the benefits of invasive treatments in CP, it is necessary to conduct studies with appropriate control arms wherever possible.
Additionally, further trials should take a more comprehensive and validated approach to pain assessment, taking into account the many factors that influence pain and the efficacy of treatment, especially the development of central sensitisation. Future RCTs should involve all required disciplines such as pain specialists, pancreatologists, interventional endoscopists and pancreatic surgeons to ensure an appropriate design and a wide-spread acceptance of the results.

Overall, this debate shows that there is an urgent need for well-designed, prospective RCTs in this field of research. Gut hopes that this debate article has contributed to summarise the current evidence and the controversies in the treatment of pain in CP, and will foster future research.

Correction notice This article has been corrected on August 8, 2021 by guest. Protected by copyright. http://gut.bmj.com/ Gut: first published as 10.1136/gutjnl-2019-318742 on 25 May 2019. Downloaded from http://gut.bmj.com/ on August 8, 2021 by guest. Protected by copyright.
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Endoscopic treatment of chronic pancreatitis:


