Chronic postsurgical pain: current evidence for prevention and management

1Department of Anaesthesiology and Critical Care, B. P. Koirala Institute of Health Sciences, Dharan, Nepal, 2Department of Anesthesiology, Siriraj Hospital, Mahidol University, Bangkok, Thailand

Parineeta Thapa1 and Pramote Euasobhon2

Chronic postsurgical pain (CPSP) is an unwanted adverse event in any operation. It leads to functional limitations and psychological trauma for patients, and leaves the operative team with feelings of failure and humiliation. Therefore, it is crucial that preventive strategies for CPSP are considered in high-risk operations. Various techniques have been implemented to reduce the risk with variable success. Identifying the risk factors for each patient and applying a timely preventive strategy may help patients avoid the distress of chronic pain. The preventive strategies include modification of the surgical technique, good pain control throughout the perioperative period, and preoperative psychological intervention focusing on the psychosocial and cognitive risk factors. Appropriate management of CPSP patients is also necessary to reduce their suffering. CPSP usually has a neuropathic pain component; therefore, the current recommendations are based on data on chronic neuropathic pain. Hence, voltage-dependent calcium channel antagonists, antidepressants, topical lidocaine and topical capsaicin are the main pharmacological treatments. Paracetamol, NSAIDs and weak opioids can be used according to symptom severity, but strong opioids should be used with great caution and are not recommended. Other drugs that may be helpful are ketamine, clonidine, and intravenous lidocaine infusion. For patients with failed pharmacological treatment, consideration should be given to pain interventions; examples include transcutaneous electrical nerve stimulation, botulinum toxin injections, pulsed radiofrequency, nerve blocks, nerve ablation, neuromodulation and surgical management. Physical therapy, cognitive behavioral therapy and lifestyle modifications are also useful for relieving the pain and distress experienced by CPSP patients. (Korean J Pain 2018; 31: 155-73)

Key Words: Chronic pain; Drug therapy; Incidence; Intractable pain; Neuropathic pain; Operative surgical procedure; Pain management; Physical therapy modalities; Postoperative pain; Prevention; Therapeutic methods.

INTRODUCTION

Pain lasting longer than the normal healing process after surgery is an unwanted adverse event in any operation, Chronic postsurgical pain (CPSP) was first defined in 1999 by Macrae and Davies [1], and later expanded by Macrae [2] in 2001, as “pain that develops after surgical intervention and lasts at least 2 months; other causes of pain...
have to be excluded, in particular, pain from a condition preceding the surgery”. An updated definition of CPSP, or persistent postsurgical pain (PPSP), was later proposed by Werner and Kongsgaard in 2014 [3]. The proposed definition was “pain persisting at least three months after surgery, that was not present before surgery, or that had different characteristics or increased intensity from preoperative pain, localized to the surgical site or a referred area, and other possible causes of the pain were excluded (e.g., cancer recurrence, infection)”. CPSP can represent a severe nuisance to patients, leading to functional limitation and psychological trauma, as well as a problem for the operative team in the form of feelings of frustration and disappointment.

We sought to explore the acute to chronic pain transition in postsurgical patients. To identify the incidence, risk factors, preventive and treatment strategies for CPSP, we searched the Ovid MEDLINE, EMBASE, and Cochrane databases between 1990 and 2017. The search utilized combinations of the following keywords: chronic postsurgical pain, persistent postsurgical pain, pain after surgery, chronic postoperative pain, incidence, risk factors, prevention, phantom limb pain, failed back surgery, post-laminectomy pain, post-thoracotomy pain and post-mastectomy pain. We limited our search to humans and English. Relevant articles were identified by the authors from the abstracts and the bibliographies.

### MAIN BODY

1. Incidence

A study by Fletcher et al. of surgical patients in Europe demonstrated that 11.8% of patients have moderate to severe pain, while 2.2% have severe pain (NRS ≥ 6), at 12 months after surgery [4]. Persistent pain can occur following various operations, ranging from simple and common ones (to illustrate, herniorrhaphy, caesarean section or dental extraction) to complicated surgeries (such as thoracotomy, radical mastectomy or hysterectomy).

The reported incidence of CPSP varies for different surgical procedures and in different studies, ranging from a low of 5% to a high of 85%. For example, studies have reported incidences ranging from 50%–85% following limb amputation, 11%–57% following mastectomy, 30%–55% after cardiac surgery, 5%–65% after thoracotomy, and 5%–63% following hernia repair [5]. One reason for this variability is the difference in the time reference considered by each researcher for labeling pain as CPSP (varying from 2 months to 1 year postoperatively).

Also, the amount of injury to the tissues or nerves and the degree of inflammation differ by operation type and procedure for the same surgery. For instance, according to the study by Fletcher, there is a reduced incidence of moderate to severe CPSP with laparoscopic cholecystectomy (8.8%) than with open cholecystectomy (28%). Fletcher’s study also found that orthopedic surgery is associated with an almost three-fold increased risk of moderate to severe CPSP, compared with all other procedures, at 12 months [4]. The sensitivity of patients to pain is also variable.

2. Risk factors

Besides the type and approach of surgery, various other risk factors have been attributed to CPSP. Some of them are patient factors (including female gender, being a young adult, genetic predisposition, and psychosocial factors), preexisting patient conditions (for example, pain present preoperatively, and any preexisting painful conditions in other parts of the body), and perioperative factors (for instance, duration and type of surgery, extent of nerve damage intraoperatively, and severity and duration of acute postoperative pain).

A recent study has also demonstrated that the severity of pain in CPSP is correlated with the mRNA expression of the signal transduction genes, representing the genetic influence on CPSP [6]. The most consistent feature associated with the occurrence of CPSP is the duration of the severe acute postoperative pain. Acute pain can lead to central sensitization, which reduces the mechanical threshold and exaggerates the response to noxious stimuli. The patient can thus present with both hyperalgesia and allodynia [7,8].

3. Prevention

Currently, there is no definitive way to prevent the occurrence of CPSP. Various techniques have been tried, by anesthesiologists and surgeons alike, to reduce the risks, but with variable success. Identifying the risk factors in each patient and applying a timely preventive strategy may help
patients to avoid the distress of chronic pain.

1) Modification of surgical technique

One of the risk factors for CPSP is the extent of tissue damage during surgery and injury to the nerves during dissection or retraction. Nerves are at continuous risk of contusion, stretching, division or entrapment from insults like surgical retraction, diathermy, or compression with bones. Alfieri in his prospective study showed that a lack of identification of nerves (the ilioinguinal, iliohypogastric and genitofemoral nerves) is significantly correlated to the presence of chronic pain following herniorrhaphy, with the risk of the development of inguinal pain climbing with the number of nerves that are not detected [9].

(1) Minimally invasive surgery

Since there is less tissue trauma in minimally invasive surgery, less chronic pain is expected than in open procedures. However, results have not always been positive. A Cochrane study involving 41 published reports of eligible trials involving 7161 participants found that there was less persistent pain and numbness following laparoscopic repair [10]. Still, another meta-analysis by Karthikesalingam et al. did not find any significant difference in the incidence of chronic pain following the laparoscopic or open-mesh repair of a recurrent inguinal hernia. Nevertheless, only three trials were included in that study, and the meta-analysis defined CPSP as pain persisting for at least one year (instead of the generally more appropriate period of 3 months) after surgery [11]. Furthermore, a prospective randomized study on the incidence of chronic groin pain (CGP) and the impact on the quality of life 10 years after laparoscopic (transabdominal preperitoneal—TAPP) versus open (mesh) repair of a recurrent inguinal hernia, indicated that the resection, rather than the retraction, of a rib leads to reduced trauma to the intercostal nerve and thus decreases the incidence of CPSP. CPSP is also less following sternotomy than that following thoracotomy [15].

How ever, there are various other sources of pain; among those are the site of an internal mammary artery dissection, stainless steel suture, surgical scar, tissue destruction from surgery and inflammation, rib fracture, and intercostal nerve trauma. As well, the sources of chronic pain following a coronary artery bypass graft can be the upper or lower limb from where the vascular graft was harvested, or the site of the central venous catheter insertion [17].

During inguinal hernia repair, indiscriminate division of the subcutaneous tissue; excessive dissection of the ilioinguinal nerve; damage of the neural structures during stretching, cutting, suturing or cauterization; over-tightening of the inguinal ring; removal of the cremaster muscle fibers; and suturing of the edge of the internal oblique muscle have all been found to be associated with an increased occurrence of CPSP [18]. Another factor found to increase CPSP is the closure of the parietal peritoneum following an abdominal hysterectomy rather than the closure of the fascia, leaving the parietal peritoneum unsutured [13].

Post-mastectomy pain syndrome (PMPS), which pres-
ents with pain typically localized to the axilla, the medial upper arm, and/or the anterior chest wall on the affected side, occurs most probably due to damage to the intercostobrachial nerve, which can occur during axillary node dissection. Consequently, the risk of damage to the intercostobrachial nerve and the development of PMPS is as likely with a lumpectomy with an axillary dissection as with mastectomy. A sentinel lymph node biopsy during a lumpectomy or mastectomy can help prevent unnecessary axillary dissection, thus reducing the occurrence of CPSP [19]. Other sources of neuropathic pain following breast cancer surgery are damage to the medial and lateral pectoral, long thoracic, or thoracodorsal nerves.

A further surgical factor consistently related to CPSP is the duration of surgery. Operations lasting longer than three hours are found to be associated with an increased CPSP [20,21].

Despite there being insufficient evidence to recommend a definite surgical technique to eliminate the possibility of CPSP, surgeons can minimize the risk of CPSP by choosing a minimally invasive surgical technique, employing careful dissection to avoid injury to nerves, avoiding extensive surgery whenever possible, and/or minimizing the duration of surgery if possible.

2) Treatment of preoperative pain

The presence of pain before surgery and the severity and duration of acute postoperative pain are predictors of CPSP. Neuroplasticity (spinal sensitization) following trauma may transform an acute pain to chronic pain if not treated in a timely manner. This can be prevented by aggressive treatment of acute pain [18]. Therefore, good perioperative pain management is thought to prevent the occurrence of CPSP. In a study by Karanikolas et al, of 65 patients who underwent lower limb amputation, the incidence of phantom limb pain after 6 months decreased significantly more following the use of an optimized perioperative epidural analgesia, or if an intravenous patient-controlled analgesia was started 48 hours preoperatively and continued for 48 hours postoperatively, compared with patients who received conventional analgesia and general anesthesia [22].

3) Modification of anesthetic technique

Perioperative management, especially the anesthetic technique adopted, has a significant effect on the prevention of CPSP. Randomized controlled trials have demonstrated a positive effect of regional anesthesia on the prevention of CPSP following laparotomy, caesarean section, cardiac surgery, breast surgery, etc. Regional analgesia techniques, for instance, epidural anesthesia, wound infiltration and intercostal nerve blocks, have been studied [23]. Unfortunately, the results are not as consistent for the prevention of chronic pain as they are for the prevention of acute pain [24].

A Cochrane review has found epidural anesthesia and paravertebral blocks significantly decrease the incidence of CPSP at 6 months following thoracotomy and breast surgery, respectively. The review, however, did not comment on the benefits for other surgeries because of the lack of studies and the small sample size [25]. Another non-randomized study by Borghi et al, examined the preoperative percutaneous insertion of a peripheral nerve catheter for the postoperative infusion of local anesthetics following limb amputation. The infusion was continued for a median of 30 days. They found that the incidence of phantom limb pain at 12 months was 16%, much less than the background incidence quoted in other studies [26,27].

Intraarticular injections of local anesthetics during arthroplasty and other joint surgeries can be effective in achieving better postoperative pain control. Wound infiltration with local anesthetics following removal of an iliac crest bone graft led to lowered iliac bone chronic pain over 4 years of follow-up [18]. Still, the result with wound infiltration is not consistent with the prevention of CPSP [28].

The adoption of preventive analgesia (providing analgesia throughout the perioperative period, thereby blocking the noxious stimulus during this painful period) rather than preemptive analgesia (providing analgesia to block the noxious preoperative stimulus) has also shown benefit in preventing CPSP [21,29]. One study compared three analgesic techniques: thoracic epidural analgesia initiated preoperatively and intraoperatively (with patient-controlled epidural analgesia provided postoperatively in both instances); and intravenous patient-controlled analgesia with morphine, started postoperatively. It was found that CPSP was significantly reduced 6 months post-operatively by using the thoracic epidural analgesia ini-
tiated preoperatively [30]. In contrast, another study did not find a difference in the incidence of chronic phantom pain with an epidural analgesia throughout the perioperative period, compared with the use of a patient-controlled opioid analgesia throughout the perioperative period [27]. Consequently, good pain control throughout the perioperative period seems more important than the technique used to achieve the pain control.

Evidence shows that the duration of acute pain influences the development of CPSP. A one-year follow-up study showed that the sum of the postoperative visual analog scores (VAS) during the first week after a laparoscopic cholecystectomy was a better predictor of the development of CPSP than the maximum reported VAS [31]. Hence, aggressive postoperative pain management might reduce the chance of developing CPSP.

The use of multimodal analgesia during the perioperative period has been proven to be better for acute postoperative pain management. Drugs acting by various mechanisms can more effectively manage pain by modulating pain signals at various points of the pain pathway than by using a single drug.

Multimodal pain regimens might include combinations of gabapentin, NSAIDs, acetaminophen, and regional anesthesia with the conventional analgesia technique [32]. Very few studies have been done to evaluate the effects of multimodal analgesia on CPSP. Fassoulaki et al. randomized 50 patients undergoing breast cancer surgery to receive gabapentin, a eutectic mixture of local anesthetic cream and ropivacaine wound infiltration, or three placebos. They found that pain and analgesic consumption was significantly less 3 months after the surgery in the multimodal analgesia group than the control [33].

4) Pharmacological treatment

Various pharmacologic agents have been tried to prevent CPSP, with some agents showing promising results. Some of the reference studies are listed in Table 1.

(1) Gabapentin and pregabalin

Most, if not all, cases of CPSP include neuropathic pain. Since the anticonvulsants gabapentin and pregabalin are the preferred agents for neuropathic pain, they have been tried for CPSP. A systematic review and meta-analysis by Clarke et al., which included 8 studies on gabapentin and 4 studies on pregabalin for use in the prevention of CPSP, found that 6 of the gabapentin trials showed a moderate to large reduction in the occurrence of CPSP (pooled odds ratio [OR] 0.52; 95% confidence interval [CI], 0.27 to 0.98; \( P = 0.04 \)), while a large reduction was found in 2 pregabalin trials (pooled OR 0.09; 95% CI, 0.02 to 0.79; \( P = 0.007 \)) [29]. Nevertheless, the dosage regimen used differed in the various studies, ranging from a single preoperative dose (of 600 or 1200 mg of gabapentin) to perioperative use (starting with 300 or 1200 mg of gabapentin preoperatively, and continuing for 8 to 10 days postoperatively).

Similarly, pregabalin (150 or 300 mg) was used preoperatively and then continued postoperatively in some studies, for only 2 more doses, but in others, for up to 2 weeks. Another systematic review and meta-analysis was conducted by Mishriky et al. to assess the analgesic efficacy of perioperative pregabalin. They found that pregabalin significantly reduced the incidence of pain at 6 months (4% vs 15%) and 12 months (9% vs 20%; RR [95% CI] = 0.31 [0.10, 0.92], \( I^2 = 15\% \)) and 0.47 [0.23, 0.97], \( I^2 = 0\% \), respectively) [34].

This promising result with gabapentin and pregabalin is now considered to be due to the preventive analgesic effect provided by gabapentinoids [35]. Despite that, a recent review of 18 randomized controlled trials (RCTs) with published and unpublished studies demonstrated that pregabalin could not reduce the incidence of CPSP at 3 months with a moderate quality of evidence [36].

(2) Antidepressants

Tricyclic antidepressants and serotonin-norepinephrine reuptake inhibitors are commonly used for chronic pain patients, both due to their efficacy in reducing neuropathic pain and the common association of depression with chronic pain. Nonetheless, studies of their use in the prevention of CPSP are too heterogeneous to allow a definite conclusion to be made.

A study by Amr et al., comparing the effects of gabapentin and venlafaxine on post-mastectomy pain found that venlafaxine extended-release (37.5 mg/d) significantly reduced chronic pain at 6 months compared to gabapentin (300 mg/d); both started on the night before surgery and continued till 30 days postoperatively [37]. Wong et al. tried
|Author, year of publication| Intervention| Sample size| Type of surgery| Observed outcomes| Time of observation| Results |
|---------------------------|-------------|-------------|----------------|-------------------|-------------------|---------|
|Gabapentinoids             |             |             |                |                   |                   |         |
|Fassoulaki et al., 2002    | Mexiletine vs gabapentin vs placebo, for 10 days | 75 | Breast cancer surgery | Incidence and severity of chronic pain | 3 months | No differences in overall incidence of chronic pain |
|Fassoulaki et al., 2005    | Gabapentin + ropivacaine infiltration + topical local anesthetics cream vs placebo | 50 | Breast cancer surgery | Incidence and severity of chronic pain | 3 and 6 months | Gabapentin and local anesthetic reduced incidence of chronic pain at 3 months, but not at 6 months |
|Nikolajsen et al., 2006    | Gabapentin vs placebo for 30 days | 46 | Limb amputation | Incidence intensity of phantom pain | 3 and 6 months | No effect on incidence of phantom limb pain |
|Brogly et al., 2008        | Gabapentin vs placebo | 50 | Thyroidectomy | Chronic neuropathic pain | 6 months | Lower risk with gabapentin |
|Clarke et al., 2009        | Gabapentin vs placebo | 126 | Total hip arthroplasty | Incidence and severity of chronic pain | 6 months | Gabapentin did not decrease risk of CPSP |
|Sen et al., 2009           | Gabapentin vs placebo | 60 | Inguinal herniorrhaphy | Incidence and severity of chronic pain | 3 and 6 months | Numerical rating scale scores lower in the gabapentin group at 3 and 6 months after surgery |
|Moore et al., 2011         | Gabapentin vs placebo | 46 | Cesarean delivery | Incidence and severity of chronic pain | 3 months | Gabapentin did not decrease risk of CPSP |
|Kinney et al., 2012        | Gabapentin vs placebo | 120 | Thoracotomy | Incidence of CPSP | 3 months | Gabapentin did not change incidence of chronic post-thoracotomy pain |
|Acin et al., 2009          | Pregabalin vs placebo | 140 | Mesh hernia repair | Persistent pain | 6 and 12 months | Reduced incidence of pain with pregabalin |
|Burke et al., 2010         | Pregabalin vs placebo | 38 | Lumbar discectomy | Pain score | 3 months | Less pain intensity and better functional outcomes in pregabalin group |
|Buvanendran et al., 2010   | Pregabalin vs placebo | 240 | Total knee arthroplasty | Neuropathic pain | 6 months | Pregabalin-treated at lower risk than placebo-treated at 3 months, but not at 6 months |
|Kim et al., 2010           | Pregabalin vs placebo | 99 | Thyroidectomy | Chronic postsurgical pain | 3 months | No differences between the groups |
|Pesonen et al., 2011       | Pregabalin vs placebo | 64 | Cardiac surgery | Incidence of CPSP | 3 months | Pregabalin-treated at lower risk than placebo-treated |
|Gianesello et al., 2012    | Pregabalin vs placebo | 60 | Spinal surgery | EuroQOL questionnaire | 3 and 12 months | At 3 months, subjective qualification of overall quality of life was better in the pregabalin group, but no differences at 1 year |
|Fassoulaki et al., 2012    | Pregabalin vs placebo | 80 | Abdominal hysterectomy or myomectomy | Presence of pain and analgesic needs due to surgery | 3 months | No differences in outcomes of the groups |
| Author, year of publication | Intervention | Sample size | Type of surgery | Observed outcomes | Time of observation | Results |
|----------------------------|--------------|-------------|----------------|-------------------|--------------------|---------|
| Choi et al., 2013           | Pregabalin vs pregabalin + dexamethasone vs placebo | 108 | Lumbar spinal surgery | Pain intensity and daily activity performance | 3 and 6 months | No significant differences in back and leg pain VAS scores of the groups over a 6-month period |
| Joshi et al., 2013          | Pregabalin vs placebo | 40 | Coronary artery bypass | Pain score | 3 months | No differences in CPSP of the groups |
| Khurana et al., 2014        | Pregabalin vs gabapentin vs placebo | 90 | Spinal surgery | Static and dynamic pain and functional outcomes | 3 months | Less pain intensity and improved functional outcomes with pregabalin |
| Bruulotte et al., 2015      | Pregabalin vs placebo | 114 | Thoracotomy | Incidence of persistent post-thoracotomy pain | 3 months | Pregabalin did not reduce the incidence of post-thoracotomy pain |
| Matsutani et al., 2015      | Pregabalin vs Loxoprofen | 68 | Thoracotomy | The pain scores, sleep interference and the incidence of neuropathic pain | 12 weeks | Pain scores, sleep interference and incidence of neuropathic pain were significantly lower for the pregabalin group |
| Antidepressants             |              |             |                |                   |                    |         |
| Amr et al., 2010            | Venlafaxine vs gabapentin vs placebo | 150 | Mastectomy | Pain intensity | 6 months | Venlafaxine superior to gabapentin for pain, with movement at 6 months |
| Ho et al., 2010             | Duloxetine vs placebo | 50 | Knee replacement | Pain intensity | 3 and 6 months | No significant differences noted throughout the duration of follow-up |
| Chocron, et al., 2013       | Escitalopram vs placebo | 368 | CABG | Quality of life (36-item) and Short Form (SF-36) | 3 and 6 months | Pain score was better overall in the escitalopram group for the preoperatively-depressed subset |
| Ketamine                   |              |             |                |                   |                    |         |
| De Kock et al., 2001        | Different doses of IV or epidural ketamine vs placebo | 36 | Rectal cancer surgery | Wound mechanical hyperalgesia and residual pain | 6 and 12 months | Significantly less residual pain until the sixth postoperative month with IV ketamine (0.5 mg/kg bolus, followed by 0.25 mg/kg/hr) |
| Hayes et al., 2004          | Ketamine vs placebo | 45 | Limb amputation | Incidence of phantom pain and stump pain | 6 months | No differences in phantom and stump pain of the groups |
| Katz et al., 2004           | Ketamine vs placebo | 74 | Prostatectomy | Incidence and intensity of pain | 6 months | No differences between the groups |
| Suzuki et al., 2006         | Ketamine vs placebo | 49 | Thoracotomy | Pain and abnormal sensation on the wound | 3 and 6 months | Lower pain score and analgesic requirement for the ketamine group at 3 months |
| Wilson et al., 2008         | Epidural bupivacaine ± epidural ketamine infusion vs. placebo | 53 | Lower limb amputation | Incidence of phantom and stump pain | 12 months | Ketamine had no significant impact on the incidence of stump or phantom pain |
| Author, year of publication | Intervention | Sample size | Type of surgery | Observed outcomes | Time of observation | Results |
|-----------------------------|--------------|-------------|-----------------|-------------------|---------------------|---------|
| Crousier et al., 2008       | Ketamine vs placebo | 36 | Mastectomy       | Incidence and characteristics of PMPS | 3 months | No significant differences in the incidence of chronic pain, but a tendency to a decrease of hyperalgesia near the scar with ketamine |
| Sveticic et al., 2008       | Ketamine vs placebo | 352 | Orthopedic surgery | Incidence of CPSP | 3 and 6 months | No differences in chronic pain of the groups |
| Dualé et al., 2009          | Ketamine vs placebo | 80 | Thoracotomy      | Incidence of CPSP | 4 months | No differences in incidence of CPSP of the groups |
| Dullenkopf et al., 2009     | Intraoperative single-dose of low- or moderate-dose of ketamine vs placebo | 120 | General and orthopedic surgery | Pain score | 3 months | No differences between the groups |
| Remerand et al., 2009       | Ketamine vs placebo | 154 | Total hip arthroplasty | Pain score | 3 and 6 months | Ketamine decreased the proportion of patients with persistent pain at rest in the operated hip |
| Ryu et al., 2011            | Epidural ketamine vs placebo | 133 | Thoracotomy      | Incidence of CPSP | 3 months | No differences in incidence of CPSP |
| Mendola et al., 2012        | S-ketamine vs placebo | 66 | Thoracotomy      | Incidence of CPSP | 3 and 6 months | No differences in incidence of CPSP of the groups |
| Joseph et al., 2012         | Ketamine vs placebo | 60 | Thoracotomy      | Incidence of CPSP | 3 months | No differences in incidence of CPSP |
| Bilgen et al., 2012         | Ketamine vs placebo | 140 | Cesarean delivery | Incidence of CPSP | 6 and 12 months | No reduction in risk of CPSP with ketamine |
| Suppa et al., 2012          | S-ketamine vs placebo | 56 | Cesarean delivery | Pain at scar area | 3 years | No differences between the groups |
| Miscellaneous               | EMLA cream vs inactive placebo cream perioperatively | 45 | Breast cancer surgery | Incidence and intensity of CPSP | 3 months | Significantly reduced incidence of total chronic pain in EMLA group |
|                            | celecoxib vs placebo | 200 | Anterior cruciate ligament reconstruction | Anterior knee pain and CRPS | 6 months | Fewer patellofemoral complication, including anterior knee pain and CRPS following celecoxib |
|                            | Intravenous lidocaine vs placebo | 36 | Breast cancer surgery | CPSP and secondary hyperalgesia | 3 months | Significantly lower incidence of CPSP and area of secondary hyperalgesia with lidocaine |
to review the use of antidepressants for the prevention of CPSP but examined only three trials, using venlafaxine, duloxetine, and escitalopram, with a positive outcome seen only with venlafaxine [38].

Overall, the limited data on the use of antidepressants presently precludes their use for the prevention of CPSP.

(3) NMDA antagonists

The benefits of the perioperative use of a subanesthetic dose of ketamine for the prevention of various types of CPSP, including phantom limb pain, has been demonstrated in different studies [39]. Significant reduction in pain for up to 6 months postoperatively has been seen with intravenous ketamine in patients undergoing colon resection [18]. A systematic review and meta-analysis was done by McNicol et al. [40] to evaluate the effectiveness of ketamine in reducing the prevalence and severity of PPSP, and to assess the safety associated with its use.

Although their meta-analysis of combined routes of ketamine use did not show any significant difference from a placebo, the analysis of the exclusively intravenous route showed a statistically significant reduction in the risk at 3 and 6 months ($P = 0.01$ and $P = 0.04$, respectively), with a risk reduction of 25% and 30%, respectively. They recommended intravenous bolus doses in the range of 0.2–0.75 mg/kg, and infusions of 2–7 mcg/kg/min for the prevention of CPSP. They did not find any statistically significant difference in the incidence of side effects (hallucinations, nightmares, excessive sedation, nausea and vomiting) except for visual disturbances (in particular, nystagmus and diplopia) with the use of ketamine, as opposed to the concern of many clinicians regarding the use of ketamine.

The use of other NMDA antagonists, one example being memantine, has not yet been established, but the pain relief does not seem to persist for long enough to prevent CPSP [21,39].

(4) Clonidine

Alpha-2 agonists such as clonidine are now well recognized for their acute analgesic effect and are frequently used perioperatively. A study of subarachnoid clonidine (300 mg) and bupivacaine, compared with bupivacaine alone, for colon surgery found that the incidence of chronic pain after 6 and 12 months was significantly less [18].

Even though there are currently few studies on the use of these agents in preventing CPSP, it has been suggested that their anti-inflammatory and anti-sensitizing effects warrants the further investigation of these drugs for the prevention of chronic pain [27].

(5) Lidocaine

Besides the use of local anesthetics for perineural injections, the use of intravenous lidocaine is increasing for the management of chronic pain, as well as perioperatively for the reduction of acute postoperative pain. Local anesthetics interrupt the sensory information to the spinal cord and thus reduce sensitization.

An intraoperative, intravenous, lidocaine infusion has been found to be effective for the prevention of CPSP after breast cancer surgery in a study conducted by Grigoras et al. Their study used a 1.5 mg/kg bolus of intravenous lidocaine before induction of general anesthesia, which was then followed by lidocaine infusion at 1.5 mg/kg/hour: the control group used an equal volume of saline. They found a significant reduction in the incidence and severity of CPSP with lidocaine at 3 months postoperatively ($P = 0.031$) [41].

Perioperative EMLA (eutectic mixture of local anesthetic) cream has also been found to reduce the incidence of chronic pain after a mastectomy [28].

However, a Cochrane review did not find many studies to further analyze the effects of lidocaine. Consequently, more research is required before its use in the prevention of CPSP can be recommended [42].

(6) NSAIDs and acetaminophen

Though NSAIDs have a beneficial effect on acute pain, are opioid sparing, and are believed to reduce secondary hyperalgesia and central sensitization, their effects on the prevention of CPSP has not been demonstrated in any study. The use of ibuprofen for the prevention of chronic pain following hip replacement surgery or a mastectomy, and the use of parecoxib during augmentation mammoplasty, have not shown any significant reduction in the incidence of CPSP [32,43]. As the data presently available is very limited and restricted to only a few agents, it is not possible to draw any conclusions about the use of
NSAIDs for CPSP.

No RCT has been conducted to show the effect of acetaminophen in preventing CPSP, though its use is now an integral part of multimodal perioperative pain management. Its role in CPSP prevention has not yet been identified.

A study by Reuben et al. of 200 patients undergoing anterior cruciate ligament surgery who received acetaminophen (1 gram) and either celecoxib or a placebo for 1–2 hours preoperatively, along with intraarticular analgesics, found that more patients in the control group developed patellofemoral complications, which included anterior knee pain and complex regional pain syndrome, among others, 6 months after the surgery [44].

(7) Steroids

Since development of chronic pain involves neuroinflammation and as steroids have an anti-inflammatory effect, steroids might be a promising agent for CPSP prevention. Although its effect on acute pain management has already been established, the data available for chronic pain is limited to date.

A Cochrane review found only 3 studies of steroids for chronic pain, too few and too heterogeneous for a meta-analysis. One of the studies found no difference in CPSP when 40 mg of dexamethasone was given before a total hip arthroplasty, yet another study found a significant difference for the prevalence of hyperesthesia, but not pain, with a single dose of methylprednisolone (125 mg) given prior to augmentation mammoplasty. The third trial using a stress dose of intravenous hydrocortisone (a loading dose plus four days infusion) after cardiac surgery found a significant positive impact on chronic pain and chronic stress symptoms [42,45].

By contrast, a recent prospective, randomized, 1-year follow-up study of the use of 16 mg intravenous dexmethasone after lumbar discectomy found a significantly higher pain score for the dexmethasone group, though there was no difference in the patients’ ability to work, disability, or self-reported health [46].

Thus, the use of steroids for the prevention of CPSP cannot be recommended with the data currently available.

(8) Opioids

Opioids are the analgesics of choice for intraoperative and postoperative analgesia for moderate to severe pain. Since severe postoperative pain is a CPSP risk factor, opioids may help in preventing CPSP. Unfortunately, strong opioids are also associated with opioid–induced hyperalgesia.

Remifentanil, which is a short-acting, strong opioid, and a popular component of balanced anesthesia, has been found to increase CPSP when used intraoperatively. However, total intravenous anesthesia with propofol and remifentanil has been more closely associated with reduced chronic post–thoracotomy pain than inhalation anesthesia with sevoflurane [28].

Phantom pain has also been found to be similar when either PCA with fentanyl or an epidural infusion with bupivacaine was started 48 hours preoperatively and continued for 48 hours postoperatively [22].

Thus, good pain control with opioids is important for CPSP prevention, despite their known hyperalgesia risk.

(9) Other pharmacological agents

Limited studies are available for other drugs, including memantine, dextromethorphan, mexiletine, and nitrous oxide, with variable effects shown on CPSP [42]. Based on the available data, the use of these drugs for the prevention of CPSP cannot be recommended.

5) Psychological intervention

The association of psychological factors with chronic pain has been well documented. Hinrichs–Rocker did a systematic review on the psychological predictors and correlates for CPSP, and found that depression, psychological vulnerability, stress and late return to work showed a probable correlation with CPSP [47]. Pain catastrophizing is sometimes found to be related to decreased CPSP, which can be due to early medical help being sought [48]. Patients having negative beliefs about opioids have a higher CPSP risk [32].

Adequate preoperative counselling regarding the surgery and expected outcomes can alleviate stress and help prevent CPSP. Identifying psychologically vulnerable patients and early intervention pre– as well as post–operatively may help prevent the development of chronic pain.
4. Management of CPSP

The management of CPSP depends on the proper identification of the etiology and type of pain via a thorough history-taking and physical examination. Pain existing from the preoperative period, postoperative complications (notable among these are infections), or the recurrence of the primary disease should be ruled out before labeling it as CPSP. More than half of CPSP patients have neuropathic pain, the remainder having nociceptive (somatic or visceral) pain. A patient may have different components of pain, and these must be identified for effective management.

During the preoperative and early postoperative period, it is very important to provide patient education and counselling about the chances of developing CPSP. Similarly, should chronic pain develop, patients must be counselled about their prognosis, the management plan, and their rehabilitation. Patients should also be counselled about their self-management strategies and their return to normal functioning.

1) Pharmacotherapy

RCTs of drugs for the management of CPSP are limited, and most of the recommendations have been extrapolated from data for other types of chronic pain, especially neuropathic pain. Similar to other neuropathic pain conditions, anticonvulsants (gabapentin and pregabalin), tricyclic antidepressants (amitriptyline and nortriptyline), serotonin-norepinephrine reuptake inhibitors (duloxetine and venlafaxine), topical lidocaine or topical capsaicin form the first line of treatment for most patients (Table 2).

Paracetamol, NSAIDs, and weak opioids (tramadol and codeine) can be used according to symptom severity, but strong opioids should be used with great caution, weighing the risks and benefits. Other drugs that may be helpful are ketamine, muscle relaxants, clonidine and intravenous lidocaine infusion.

Some studies have found low-concentration capsaicin to be useful for post-mastectomy pain [49]. In contrast, a recent Cochrane review of low concentration (< 1%) capsaicin cream was not found useful for neuropathic pain [50]; a high concentration was also not useful for persistent pain following inguinal herniorrhaphy, though it was useful for postherpetic neuralgia, painful diabetic neuropathy and HIV neuropathy [51]. The number of studies was too limited to form any recommendations.

A 5% lidocaine patch was found effective for neuropathic pain associated with allodynia following cancer surgery [52]. A conclusion from a meeting of 44 pain specialists from 17 countries, and based on a retrospective analysis of case reports, concluded that CPSP associated with localized, superficial pain and allodynia showed a positive response to 5% lidocaine plaster [53]. An observational study of patients with posttraumatic and postsurgical localized neuropathic pain also showed a significant reduction in pain and a decrease in painful areas following treatment with lidocaine-mediated plaster [54,55]. A randomized, double-blind, placebo-controlled, cross-over trial among 21 male patient having unilateral severe persistent inguinal pain following herniorrhaphy, showed no significant improvement with a 14-day application of a 5% lidocaine patch [56].

A nitroglycerine transdermal patch has been found useful for chronic post-thoracotomy pain [57]. A topical 5% amitriptyline cream has also been tried, but it was not found to be effective compared with 5% lidocaine cream or a placebo [58]. A pilot study of topical 1% amitriptyline, 0.5% ketamine or their combination for the treatment of neuropathic pain, including CPSP, found the combination cream to be effective after 7 days’ application, whereas they were ineffective when applied individually [59]. The subsequent open-label, prospective study with a 2% amitriptyline/1% ketamine combination cream applied for 6–12 months showed significant pain relief, with long-term patient satisfaction and minimal side effects [60].

2) Pain interventions

For patients who do not improve with pharmacotherapy, various interventions have been tried, ranging from nerve blocks to nerve ablation and neuromodulation (Table 2).

Nerve blocks, neuromaxial blocks and sympathectomies have been tried. Successful treatment of sternotomy-induced neuralgia has been reported, using repeated bupivacaine blocks, phenol blocks or alcohol blocks [49]. Pain relief was seen with epidural injections for lumbar or cervical post-surgery syndrome [61], but no significant difference was seen when caudal or cervical epidural local anesthetic
| Author                | Methods                                | Intervention                          | Participants                                                                 | Outcomes                                                                 | Author’s conclusions                                                                 |
|-----------------------|----------------------------------------|---------------------------------------|------------------------------------------------------------------------------|---------------------------------------------------------------------------|-------------------------------------------------------------------------------------|
| **Topical agents**    |                                        |                                       |                                                                              |                                                                           |                                                                                      |
| Fleming et al., 2009  | Retrospective study                    | 5% Lidocaine patch                    | 26 patients: persistent post-surgical neuropathic pain 24 patients: post herpetic neuralgia, 18 patients: cancer related neuropathic pain | Potent analgesic efficacy in 35% of patients with postsurgical pain       | Supports trial of 5% lidocaine patch for cancer patients with neuropathic syndromes associated with allodynia |
| Nicolaou et al., 2011 | Retrospective study and expert consensus | 5% Lidocaine patch                    | 58 patients with post-operative/post-traumatic neuropathic cutaneous pain     |                                                                           | Positive response for CPSP associated with localized superficial pain and allodynia       |
| Correa-Illanes et al., 2012 | Prospective observational study       | 5% Lidocaine patch                    | 19 patients with traumatic injuries                                           | NRS: 3-point reduction in 79% of patients, 50% reduction in 57.9% of patients, Painful area: reduction by median 1 cm², 50% reduction in 94.7% of patients, Functional improvement after treatment in 14/19 patients (73.7%) | Lidocaine medicated patch effectively treated traumatic injuries of peripheral nerves which presented with chronic localized neuropathic pain, reducing both pain intensity and the size of the painful area |
| Hans et al., 2009     | Prospective observational study        | 5% Lidocaine patch                    | 40 patients with chronic neuropathic pain after surgical or non-surgical trauma | Significant improvement in pain intensity, neuropathic pain scores         | Lidocaine 5% patches seem to be an effective treatment for post-surgical and post-traumatic pain |
| Bischoff et al., 2013 | RCT                                    | 5% Lidocaine patch                    | 21 patients with severe, unilateral, persistent inguinal postherniorrhaphy pain | No differences in "summed pain intensity differences" following treatment between lidocaine and placebo patch treatments in all patients | Lidocaine patch treatment did not reduce combined resting and dynamic pain ratings, compared with placebo in patients with severe, persistent inguinal postherniorrhaphy pain |
| Glantz et al., 2004   | Prospective observational study        | Transdermal Nitroglycerine (NTG) with oral etodolac | 30 patients with moderate to severe pain 1.5 years after thoracotomy         | Significant reduction in VAS scores, breakthrough pain intensity and sleep efficiency | NTG added to etodolac appears to be effective for the treatment of chronic post-thoracotomy pain, with minimal side effects |
| Ho et al., 2008       | Randomized controlled cross-over study | 5% amitriptyline cream vs 5% lidocaine cream vs placebo | 35 patients with postsurgical neuropathic pain, postherpetic neuralgia, or diabetic neuropathy with allodynia or hyperalgesia | Reduction in pain intensity observed with topical lidocaine (p <0.05). No significant change in pain intensity with topical amitriptyline or placebo | Topical lidocaine reduced pain intensity, but the clinical improvement was minimal, and the topical 5% amitriptyline was not effective |
| Lynch et al., 2003    | 2-day randomized, double blind, lacebo-controlled, 4-way cross-over trial | Topical 1% amitriptyline vs 0.5% ketamine vs combination amitriptyline 1%/ketamine 0.5% vs placebo | 20 patients with chronic neuropathic pain | A lack of effect for all treatments in the 2-day double blind placebo controlled trial, followed by analgesia in an open label trial in a subgroup of subjects who chose to use the combination cream for 7 days |                                                                                      |
| Author               | Methods                          | Intervention                                      | Participants                          | Outcomes                                                                 | Author’s conclusions                                                                 |
|----------------------|----------------------------------|---------------------------------------------------|---------------------------------------|---------------------------------------------------------------------------|---------------------------------------------------------------------------------------|
| Lynch et al., 2005   | Open-label prospective trial     | Topical combination cream of amitriptyline 2%/ketamine 1% | 28 patients with refractory, moderate to severe peripheral neuropathic pain | Treatment was associated with long-term reduction (6–12 months) in perceived pain, moderate to complete satisfaction, and was well-tolerated in the treatment of neuropathic pain | Topical 2% amitriptyline/1% ketamine, given over 6–12 months, is associated with long-term perceived analgesic effectiveness in treatment of neuropathic pain |
| Revel et al., 1996   | RCT, multicenter study           | Forceful injection: 125 mg of prednisolone acetate with 40 mL of normal saline vs 125 mg of prednisolone | 60 post-lumbar laminectomy patients with chronic low back pain | The proportion of patients relieved of sciatica was 45% in the forceful injection group, compared to 19% in the control group, with significant difference at 6 months | Positive short-term and negative long-term pain relief |
| Meadeb et al., 2001  | RCT                              | Forceful injection of 20 mL of normal saline with/without 125 mg of epidural prednisolone vs 125 mg of epidural prednisolone | 47 post-lumbar laminectomy syndrome patients | The VAS scores improved steadily in the forceful injection group, producing a nearly significant difference on day 120 compared to the baseline. Epidural glucocorticoids used alone induced only short-lived pain relief | Negative short-term and long-term pain relief |
| Manchikanti et al., 2010 | RCT                              | Caudal epidural with 0.5% lidocaine (LA) vs caudal epidural with 0.5% lidocaine with 6 mg betamethasone | 140 patients with chronic function-limiting low back pain with or without lower extremity pain of at least 6 months duration post-surgery | Combined pain relief (≥ 50%) and disability reduction recorded for 53% of the patients in the LA group, vs 59% in the LA and steroid group, with no significant differences noted with or without steroids over a period of one year | In post-surgery patients who have chronic function-limiting low back and/or lower extremity pain, and who receive caudal epidural injections, either with or without steroids, may provide significant pain relief in 70%–75% of patients |
| Manchikanti et al., 2012 | RCT                              | Cervical interlaminar epidural with 0.5% lidocaine vs epidural with 0.5% lidocaine + 6 mg betamethasone | 120 patients with cervical post-surgery syndrome | Cervical epidural injections with LA, with or without steroids, were effective in 67% of patients overall, and 87% in the group without steroids, and 72% in the group with steroids | Patients with continued pain following previous cervical surgery may be treated with cervical interlaminar epidural injections with or without steroids |
| Stolker et al., 1994 | Prospective observational study  | Posterior thoracic percutaneous partial rhizotomy (PPR) | 45 patients with irradiating pain in the thoracic region with temporary positive response to an intercostal blockade with lidocaine | Excellent long-term effect (average 24 months) in 48.8% good result in 36.6% poor result in 14.6% | When conservative treatment fails, thoracic PPR may prove an effective and safe treatment for chronic segmental thoracic pain |
injections alone were used, compared with combined epidural steroid and local anesthetic injections [62,63].

Entrapment of the superficial nerves of the abdominal wall, or trauma to nerves during surgery, can be a source of chronic abdominal pain following any abdominal surgery. The injection of local anesthetic agents, with or without steroids, in the transverse abdominis plane, rectus sheath or tender point can produce pain relief in such patients [64-66]. An axillary brachial plexus block with patient-controlled analgesia was found to be useful for the treatment of complex regional pain syndrome I, developed after surgical release in case of carpel tunnel syndrome [67].

Botulinum toxin injections in the painful areas of chronic post-thoracotomy pain have been reported to provide significant pain relief for patients unresponsive to oral therapies [68]. Positive results have also been seen with botulinum injections in the area of abdominal wall pain following ventral hernia repair, and for trigeminal neuropathic pain following dental implants [69,70].

Phenol injections or radiofrequency ablation of stump neuroma and dorsal root ganglia have been found useful for stump and phantom pain [71,72]. If other options fail, lesioning the dorsal root entry zone (DREZ) and motor cord stimulation have been found useful [73]. Radiofrequency percutaneous partial rhizotomy has been found to be useful in the management of chronic thoracic segmental pain, including post-thoracotomy and post-mastectomy pain [74]. A retrospective study comparing the effects of pulsed radiofrequency (PRF) of the dorsal root ganglia (DRG) with motor cord stimulation, in terms of success in pain relief and the duration of pain relief, found the PRF of DRG to be superior to PRF of the ICN [75]. Radiofrequency neurolysis appears to be significantly more effective than local nerve infiltrations [76]. Pulsed radiofrequency treatment to the ilioinguinal and genitofemoral nerves and nerve roots have resulted in complete relief lasting in 6 months [77]. Transcutaneous electrical nerve stimulation [78].

Spinal cord stimulation has provided promising results and is commonly used for phantom pain [79]. Recently, spinal cord stimulation has provided promising results in the management of phantom limb syndrome [80]. It has been used for the management of chronic pain [81].

It has been used for the management of chronic pain [82]. RCT: Randomized controlled trial.

| Author, Year | Methods | Intervention | Participants | Outcomes | Author’s conclusions |
|-------------|---------|--------------|--------------|----------|---------------------|
| Cohen et al., 2006 | Retrospective study | Pharmacotherapy vs pulsed radiofrequency (PRF) of intercostal nerve (ICN) vs PRF of dorsal root ganglion (DRG) | 49 patients with chronic post-thoracotomy pain (CPTP) | At 6-week follow-up: > = 50% pain relief in 61.5% in PRF of DRG group vs. 27.3% in the medical management (MM) group vs 21.4% in the ICN group (P = 0.12). At 3-month follow-up: 53.8% in the DRG group continued to report > = 50% pain relief vs. 19.9% in the MM and 6.1% in the ICN groups, respectively (P = 0.02) | PRF of the DRG was a superior treatment to pharmacotherapy and PRF of the ICN in patients with CPTP |
| Kastler et al., 2012 | Retrospective study | Radiofrequency ablation vs local infiltration | 42 patients with chronic inguinal pain refractory to specific medications | Maximum early pain relief did not statistically differ (77% in the RFN group and 81.5% in the injection group). Mean duration of pain relief was statistically significant (P = 0.005) in the RF group (12.5 months) compared to the infiltration group (1.6 months) | Radiofrequency neurolysis appears to be significantly more effective than local nerve infiltrations |

Table 2. Continued
post-thoracotomy pain, but the evidence is presently limited to case studies [80]. Its use has also been found effective for the management of neuropathic arm or leg pain following cervical or spinal surgeries, and in complex regional pain syndrome type I [81].

3) Surgical management

For patients with post-mastectomy pain syndrome, surgical resection of the neuroma and allowing the cut-ending of the nerve to retract deep into the intercostal muscles has been found to be helpful [82]. Since a neuroma can reform, relocation of the nerve to a protected site and helping regrowth via a nerve graft can be a better option [19]. Excision of a painful intercostal neuroma and implantation of the proximal end of the nerve into the latissimus dorsi muscle has been found to be effective for neuromas developed following thoracic and upper abdominal surgeries [83].

An autologous fat graft to the dermo-hypodermal junction at the painful scar area has been shown to be effective [84]. A scar excision can also help relieve pain following surgery. In the case of patients with chronic neuropathic pain following an inguinal herniorrhaphy who are not relieved by conservative measures, some pain relief can be provided by revision surgery: a triple neurectomy of the ilioinguinal, iliohypogastric and genitofemoral nerves; or removal of the fixation material and the

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**Fig. 1.** Step-by-step approach to the prevention of chronic postsurgical pain.
Though surgery is an option for pain management in CPSP when other options have failed, we must consider the possibility of the recurrence of CPSP following surgery. Preventive measures, including effective perioperative pain management and counselling, should be instituted on time.

4) Lifestyle modification

Physical therapies, including massage, physiotherapy and acupuncture, have been tried in the management of CPSP. These modalities can reduce pain, but only temporarily. Lifestyle modifications in the form of rest and activity limitation are not advised as these can lead to further complications, especially poor functional outcomes.

5) Psychological interventions

Operant conditioning and cognitive behavioral therapy (CBT) have been found to be useful for the management of chronic pain. In operant conditioning, the clinician emphasizes the modification of responses to maladaptive behaviors, and the modification of behaviors that consist of overt expressions of pain, distress and suffering in response to chronic pain. By comparison, CBT focuses on improving physical and emotional functioning despite the pain, rather than attempting to eliminate the pain. It provides positive reinforcement to wellness behavior, physical fitness and cognitive reframing so that the patients can become desensitized to the persistent pain and can function better. It usually combines stress management, problem solving, goal setting, relaxation, and the pacing of activities [86,87].

In a randomized study comparing the effectiveness of lumbar fusion with posterior transpedicular screw fixation, and cognitive intervention and exercises among patients with CPSP after previous surgery for disc herniation, the surgery and the combination of cognitive intervention and exercise showed similar levels of effectiveness in managing the pain [88]. Aggressive exercise with CBT has been found to be useful for various back pain conditions, including those following surgery [89].

CONCLUSIONS

CPSP is a common but overlooked complication of surgery which can cause functional limitation and psychological distress to patients. Though we are still unable to completely prevent this problem, with a proper knowledge of the condition, we can easily identify the risk factors in a patient undergoing surgery and institute appropriate preventive measures.

These range from modification of the surgical, anesthetic and analgesic techniques to psychological counselling. Any event of CPSP should be identified in a timely manner, and proper management—consisting of pharmacotherapy, appropriate pain interventions, surgery and/or psychological managements by a multidisciplinary team—can improve the pain as well as the physical and social functionality of the patient (Fig. 1).

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CONFLICTS OF INTEREST

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