Complex regional pain syndrome

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ABSTRACT
Complex regional pain syndrome (CRPS) previously known as reflex sympathetic dystrophy is a chronic neurological disorder involving the limbs characterized by disabling pain, swelling, vasomotor instability, sudomotor abnormality, and impairment of motor function. CRPS is not uncommon after hand surgery and may complicate post-operative care. There is no specific diagnostic test for CRPS and the diagnosis is based on history, clinical examination, and supportive laboratory findings. Recent modifications to diagnostic criteria have enabled clinicians to diagnose this disease more consistently. This review gives a synopsis of CRPS and discusses the diagnosis, pathophysiology, and treatment options based on the limited evidence in the literature.

KEY WORDS
Causalgia; complex regional pain syndrome; hand surgery; reflex sympathetic dystrophy; Sudeck atrophy; superficial radial nerve

INTRODUCTION
Complex regional pain syndrome (CRPS) is one of the most challenging chronic pain conditions of the limbs. There is little agreement with regards to the aetiology, symptoms, clinical presentation, diagnosis, or treatment of CRPS. In fact there is confusion regarding the terminology itself. Historically, CRPS has been described by a number of terms that include causalgia, Sudeck atrophy, reflex sympathetic dystrophy (RSD), algodystrophy, post-traumatic dystrophy and shoulder-hand syndrome. In order to bring some uniformity to this problem, the International Association for the Study of Pain (IASP) in 1994 introduced the term CRPS to describe a wide variety of post traumatic neuropathic pain conditions of the limbs.[1] The use of the term CRPS has also been questioned and perhaps another more appropriate term will be developed in the future.[2] In the meantime, CRPS is the term used routinely by pain specialists and neurologists and the use of traditional terminology like RSD and causalgia is dwindling. CRPS is most often associated with surgery of the distal upper extremity to complicate recovery, delay return to work, diminish health related quality of life, and increase the likelihood of poor outcomes and/or litigation.[3] This article presents a brief historical overview followed by a review of the definition, incidence, pathophysiology, current diagnostic criteria, and the basic approach to patients with CRPS following hand surgery.

HISTORICAL OVERVIEW
The earliest documented description of CRPS is probably Ambroise Pare’s report from the 16th century describing the persistent pain and contractures experienced by
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Table 1: IASP diagnostic criteria for CRPS* (1994)
The presence of an initiating noxious event, or a cause of immobilization†
Continuing pain, allodynia, or hyperalgesia in which the pain is disproportionate to any known inciting event
Evidence at some time of edema, changes in skin blood flow, or abnormal sudomotor activity in the region of pain (can be sign or symptom)
This diagnosis is excluded by the existence of other conditions that would otherwise account for the degree of pain and dysfunction
*If seen without “major nerve damage” diagnose CRPS I; if seen in the presence of “major nerve damage” diagnose CRPS II; †Not required for diagnosis; 5–10% of patients will not have this. Must meet criteria 2, 3, and 4 for diagnosis.

Table 2: Clinical diagnostic criteria for CRPS* – Budapest Criteria (2003)
Continuing pain, which is disproportionate to any inciting event
Must report at least one symptom in three of the four following categories
  - Sensory
  - Vasomotor
  - Sudomotor/ Oedema
  - Motor/ Trophic
Must display at least one sign at time of evaluation in two or more of the following categories
  - Sensory
  - Vasomotor
  - Sudomotor/ Oedema
  - Motor/ Trophic
There is no other diagnosis that better explains the signs and symptoms
* If seen without “major nerve damage” diagnose CRPS I; if seen in the presence of “major nerve damage” diagnose CRPS II.

King Charles IX after a blood-letting procedure.[4] In 1766, Hunter described the pain syndrome after a joint injury.[5] Silas Weir Mitchell [Figure 1], the father of American neurology gave the first detailed description of CRPS in 1864.[6] Mitchell together with Morehouse and Keen noted the frequent occurrence of exaggerated presentation of pain in relation to the injury in veterans of the American civil war.[7] Mitchell coined the term causalgia from the Greek Kausis (fire) + Algos (pain). His graphic description remains the best depiction of the clinical presentation of CRPS even today.[2,8] Causalgia is characterized by exquisite burning pain that begins in the distribution of an injured peripheral nerve and then spreads beyond it. It is almost always associated with an injury to a major nerve trunk.[8]

In 1900, Sudeck described the clinical and radiological features of post-traumatic reflex atrophy of bone latter known as Sudeck atrophy.[9] He also proposed a post-traumatic pain syndrome associated with oedema and trophic changes. Rene Leriche in 1916 was the first to link the sympathetic nervous system to causalgia and reported pain relief in a patient after extensive periarterial nerve stripping.[10] In 1946, Evans introduced the term reflex sympathetic dystrophy (RSD) for this condition. Evans theorized that trauma that generated activity in afferents set up a reflex in the spinal cord which stimulated activity in sympathetic efferents, and in turn resulted in dystrophic changes in the periphery of the limb.[11] This theory gained acceptance because some patients had pain relief after a local anaesthetic or pharmacologic sympathetic ganglion blockade.[12] However, recent studies have failed to demonstrate a ‘reflex arc’ and suggest that the sympathetic nervous system may not be involved in every case and that dystrophy rarely occurs.[13] In 1986, Roberts used the term sympathetically mediated pain (SMP) to describe RSD, whereas Campbell and colleagues used the term sympathetically independent pain (SIP) for those patients with nerve injuries who did not respond to sympathetic blockage.[14] The use of differing terms with imprecise classifications and unclear pathogenesis led to confusion, misdiagnosis, and mistreatment of patients with this pathology.[15]

In 1993, IASP designated a Task Force to review the
nomenclature and develop diagnostic criteria. In 1994, at a consensus workshop in Florida, the Task Force developed the nomenclature, CRPS, and proposed the IASP diagnostic criteria [Table 1]. They further subdivided CRPS into types I and II. The clinical features of both types are identical and the only distinguishing feature is the presence of a peripheral nerve injury in type II. In essence, type I refers to RSD and type II refers to causalgia. An IASP consensus workshop held in Budapest in 2003 proposed modified clinical diagnostic criteria (Budapest criteria) to address the lack of specificity in the original IASP diagnostic criteria [Table 2].

DEFINITION

CRPS describes an array of painful conditions that are characterized by a continuing (spontaneous and/or evoked) limb pain that is seemingly disproportionate in time or degree to the usual course of any known trauma or other lesion. The pain is regional (not in a specific nerve territory or dermatome) and usually has a distal predominance of abnormal sensory, motor, sudomotor, vasomotor, and/or trophic findings [Table 3]. CRPS has been divided into three stages of progression based on the duration of symptoms. Although it is not necessary for each patient to develop all stages or for these stages to progress in a sequential fashion, recognizing the stage and the predominant complaint can help with management of patients.

Stage I (Acute stage: 0-3 months): It is characterized primarily by pain/sensory abnormalities (e.g. hyperalgesia, allodynia), signs of vasomotor dysfunction, and prominent edema and sudomotor disturbance [Figures 2-4].

Stage II (Dystrophic stage: 3-9 months): It is characterized by more marked pain/sensory dysfunction, continued evidence of vasomotor dysfunction, with development of significant motor/trophic changes [Figures 2-4].

Stage III (Atrophic stage: 9-18 months): It is characterized by decreased pain/sensory disturbance, continued vasomotor disturbance, and markedly increased motor/trophic changes.

INCIDENCE

CRPS is an uncommon disease with a prevalence of <2% in most retrospective series. A study from Netherlands reported an incidence of 26.2 cases per 100,000 person years, whereas a study from the United States estimated the incidence at 5.5 cases per 100,000 person years. A higher incidence of CRPS is reported in patients between the ages of 40-49 and in women.

| Diagnostic category | Symptom | Sign |
|---------------------|---------|------|
| Sensory             | Continuous burning pain in the distal part of the affected extremity | Stimulus-evoked pains include mechanical and thermal allodynia and/or hyperalgesia, and deep somatic allodynia (pain due to touching the joints and movement of joints). |
|                     | Pain is disproportionate in intensity to the inciting event and usually increases when the extremity is in a dependent position. | Evidence of temperature asymmetry |
|                     | Sensory abnormalities are most pronounced distally, and have no consistent spatial relationship to individual nerve territories or to the site of the inciting lesion. | Evidence of skin color changes/ asymmetry |
| Vasomotor           | Reports of temperature asymmetry | Evidence of edema |
|                     | Reports of skin color changes/ asymmetry | Evidence of sweating changes and/or sweating asymmetry |
| Sudomotor/Edema     | Reports of edema | Evidence of temperature asymmetry |
|                     | Reports of sweating changes and/or sweating asymmetry | Evidence of skin color changes/ asymmetry |
| Motor/Trophic       | Reports of decreased range of motion | Evidence of decreased range of motion |
|                     | Reports of motor dysfunction | Evidence of motor dysfunction |
|                     | Weakness | Weakness |
|                     | Tremor | Tremor |
|                     | Dystonia | Dystonia |
|                     | Coordination deficits | Coordination deficits |
|                     | Disturbed body perception of the affected extremity | Evidence of trophic changes |
|                     | Reports of trophic changes | Hair |
|                     | Hair | Nail |
|                     | Nail | Skin |
|                     | Skin | Osteoporosis |
The upper extremity is affected twice as commonly as the lower limb, and a fracture is the most common trigger (46%). In 10-26% of patients with CRPS, no precipitating factors can be found. No correlation to diabetes, smoking or alcohol has been found. The incidence of recurrent CRPS type I is 1.8% per year, and 50% are spontaneous recurrences.

The exact incidence and prevalence of CRPS after hand surgery is unknown. Rates of the occurrence of CRPS range from 4.5% to 40% after fasciectomy for Dupuytren contracture, 2% to 5% after carpal tunnel surgery, and 22% to 39% after distal radius fractures. A higher incidence of CRPS has been reported in highly comminuted, intra-articular distal radius fractures, fractures with associated ulnar styloid injuries, and in fractures treated with closed reduction and casting (versus percutaneous pinning). An elevated intra-cast pressure as a result of a tight cast or extreme positions was also a common risk factor for development of CRPS.

PATHOPHYSIOLOGY

Most patients (90%) with CRPS have an initiating noxious event (trauma/ischemia/nerve compression) in the clinical history. The reason why only some patients develop CRPS is unclear. There is also no single pathophysiological...
mechanism that can explain the diversity and the heterogeneity of the symptoms (oedema, central nervous system symptoms, joint involvement, etc). It is now accepted that multiple mechanisms are involved and the presentation depends on the relative contribution of each mechanism. The pathophysiologic mechanisms that are believed to contribute to CRPS include alterations in cutaneous innervation (lower density of small fibres—C and Aδ), central and peripheral sensitization (increased excitability of nociceptive neurons in the spinal cord and local tissues as a result of persistent noxious input from tissue damage/nerve injury mediated by neuropeptides like substance P and bradykinin), altered function of the sympathetic nervous system; lower levels of circulating catecholamines; increased levels of local and systemic inflammatory cytokines (TNF-α, interleukin-1, -2, and -6), lower systemic levels of anti-inflammatory cytokines (interleukin-10), genetic factors (HLA-b62 and HLA-DQ8 alleles), and psychologic factors (anxiety, anger, and depression). The most prominent mechanism appears to be the inflammatory process because all the classic signs of inflammation (oedema, redness, hyperthermia, and impaired function) are conspicuous in the early stages of CRPS.

**DIAGNOSTIC CRITERIA**

CRPS is a clinical diagnosis based on patient symptoms and signs elicited on physical examination. The IASP diagnostic criteria [Table 1] have very good sensitivity (0.99), but poor specificity (0.41), potentially leading to over diagnosis. The Budapest clinical diagnostic criteria retain the sensitivity of the IASP criteria, but improve the specificity (0.68) [Table 2]. The IASP Budapest consensus group also retained the division of CRPS into types I and II subtypes depending on the absence or presence of evidence of peripheral nerve injury, respectively. In addition, a third diagnostic subtype called CRPS-NOS was recommended. This is because approximately 15% of the patients previously diagnosed with CRPS (based on 1994 IASP criteria-Table 1) would not fully meet the new clinical diagnostic criteria [Table 2]. Patients who have fewer than three symptom or two sign categories, or who were not showing a sign at the time of the examination, but had exhibited this previously, and whose signs and symptoms were felt to be best explained by CRPS would receive a diagnosis of CRPS-NOS. The common sensory, vasomotor, sudomotor, motor, and trophic changes associated with CRPS have been tabulated in Table 3.

The IASP Budapest consensus group also felt that a higher level of specificity was desirable in the research context. The Budapest research diagnostic criteria for CRPS are therefore slightly more stringent and require the patient to have at least two positive sign categories and all four positive symptom categories (versus at least two positive sign categories and three positive symptom categories for clinical diagnosis). This improves the specificity of the Budapest research diagnostic criteria to 0.79. Harden has also suggested the use of measurement techniques and laboratory testing to further improve the specificity of the research diagnostic criteria. Some of these techniques include the use of 100 mm visual analogue scale for quantifying pain, measuring temperature alldynia using a Peltier type device, mechanical alldynia using von Frey testing, Laser Doppler for vasomotor tone, and volumetry for oedema.

**APPROACH TO A PATIENT WITH CRPS FOLLOWING HAND SURGERY**

**History**

In patients with a suspected diagnosis of CRPS, history of the period prior to surgery should be obtained to determine any pre-existing conditions, past traumatic injuries, or pain issues. Pre-existing subclinical problems can be exacerbated in the perioperative period. One should ascertain the time of onset of symptoms relative to surgery. The symptoms of CRPS (pain, numbness, swelling, stiffness, etc) are often non-specific and frequently seen in most post-operative patients. It is therefore important to evaluate the response of patients to standard treatment of these symptoms. Patients with CRPS do not respond to narcotics, are irritable, do not cooperate with therapy, resist returning to work, and often adopt a protective posture to guard the affected extremity.

**Physical examination**

The affected extremity should be exposed to allow a full examination from the neck downwards. The examination should be carried out at rest, during activity, and during ambulation. The three goals of physical examination are (1) comparison of the affected extremity against the unaffected limb and the pre-operative examination; (2) to determine evidence of sensory, vasomotor, sudomotor/oedema, and motor/trophic signs diagnostic of CRPS [Table 3]; and (3) identify any possible nerve injuries. The commonly reported nerve injuries associated with CRPS include the superficial branch of the radial nerve near the radial styloid, the palmar cutaneous branch of
the median nerve near the wrist crease, and the dorsal branch of the ulnar nerve close to the ulnar styloid. The extremity examination should also include assessment of skin integrity, range of motion, joint stability, motor power, neurologic, and vascular functions. Table 4 lists the definitions of the commonly used terms used in describing the signs and symptoms associated with CRPS.[31]

**Investigations**

CRPS is a clinical diagnosis, but one of the criteria for diagnosis is the absence of any other diagnosis to explain the findings. The main purpose of investigations in CRPS is therefore to rule out other conditions. The ESR, C-reactive protein, complete blood count, and serum auto-antibodies results are useful in ruling out infections and rheumatologic conditions.[32] Electrodiagnostic studies may be indicated to rule out specific neuropathic conditions, such as peripheral neuropathy, entrapment neuropathies, or nerve injury. Radiographic studies including magnetic resonance imaging are often necessary to rule out bone or soft tissue pathology as the source of pain.[12,19]

A few tests have been described that are complementary to the clinical diagnostic criteria and are useful in demonstrating objective abnormalities characteristic of CRPS.[32] They include:

- **Plain radiography**
  - If they are normal at baseline, serial examination can reveal significant change over time and aid in confirmation of diagnosis [Figure 4]. Periarticular osteopaenia and patchy osteoporosis can be seen in plain radiographs as early as 2 weeks after onset of CRPS.[12,19,32]

- **Bone scans**
  - Three phase (blood pool, blood phase, scan phase) bone scanning of the affected extremity using technetium Tc 99m-labelled bisphosphonates is a highly sensitive (but relatively nonspecific) test that may detect osseous changes earlier than plain radiographs. Classic findings include increased periarticular uptake in the third phase (scan phase) and evidence of vasomotor instability and abnormal patterns of flow distribution in the first and second phase (blood pool and blood phase). Band-shaped increased radionuclide accumulation in the metacarpophalangeal and interphalangeal joints of the affected extremity during the scan phase is characteristic for CRPS. Multiple studies have found that the sensitivity of bone scans decreases and the specificity increases with disease duration.[12,26,32]

- **Bone densitometry**
  - They will show a lowered bone mineral density and bone mineral content in affected limb of patients with CRPS. This can be used to monitor treatment efficacy as these indices often improve in patients undergoing treatment.[12]

- **Thermography**
  - An infrared thermometer (accuracy of ±0.1 °C) is used to measure several symmetrical points on the affected and contralateral extremity, making comparisons between the two extremities. A difference of 0.5 °C is considered mildly asymmetrical, and a difference of 1.0 °C is considered significant.[12]

- **Sweat testing**
  - Subjective sweat testing can be performed by applying an indicator-starch powder to the limbs. The indicator changes colour when the limb sweats. Quantitative

| Term               | Meaning                                                                 |
|--------------------|-------------------------------------------------------------------------|
| Pain               | An unpleasant sensory and emotional experience associated with actual or potential tissue damage |
| Noxious stimuli    | A stimulus that is damaging or threatens damage to normal tissues       |
| Nociception        | The neural process of encoding noxious stimuli.                         |
| Analgesia          | Absence of pain in response to stimulation which would normally be painful. |
| Hyperalgesia       | Increased pain from a stimulus that normally provokes pain.            |
| Hypoalgesia        | Diminished pain in response to a normally painful stimulus             |
| Allodynia          | Pain due to a stimulus that does not normally provoke pain.            |
| Paresthesia        | An abnormal sensation, whether spontaneous or evoked.                  |
| Dysesthesia        | An unpleasant abnormal sensation, whether spontaneous or evoked.       |
| Hyperesthesia      | Increased sensitivity to stimulation (includes both allosthesia and hyperalgesia) |
| Hypoesthesia       | Decreased sensitivity to stimulation                                    |

**Table 4: Common term used to describe symptoms and signs in CRPS**
sweat production can be measured using the resting sweat output and quantitative sudomotor axon reflex test (provocative test that determines the sweat output in response to an iontophoresis cholinergic challenge, such as acetylcholine or methacholine). Both of these tests have been shown to correlate with clinical signs of CRPS.\textsuperscript{[3,12]}

**Sympathetic blocks**
Sympathetically maintained pain (SMP/ RSD) may be differentiated from sympathetically independent pain (SIP) by an intravenous injection of phentolamine (\(\alpha_1\)- and \(\alpha_2\)-adrenergic receptor blocker). In patients with SMP, the injection results in transient pain relief. Another option is to perform a sympathetic block using a local anaesthetic. A cervicothoracic block (stellate ganglion/ upper thoracic) is used for upper extremity symptoms and a lumbar paravertebral block is used for lower extremity symptoms. In patients who respond during the block, pain is relieved, whereas motor function is retained.\textsuperscript{[3]}

**Treatment**
The aim of treatment of CRPS is pain control followed by recovery of limb function.\textsuperscript{[26]} Although a number of treatment options have been reported in the literature, these are empirical at best and no reliable protocol is available for use in all patients. Early recognition and prompt initiation of treatment improve patient outcomes. Even suspected cases of CRPS should be referred to a pain specialist early. A multidisciplinary approach to treatment with the hand surgeon as the lead with inputs from the hand therapist and anaesthesiologist (pain clinic) is required for optimum outcomes. The inclusion of psychologist or the workmen’s compensation coordinator may be beneficial in certain circumstances. The initial treating surgeon should be involved throughout the process and one must resist the temptation to dump patients to pain clinics. Comorbidities such as depression, sleep disturbance, anxiety, and loss of body image should be treated concurrently.\textsuperscript{[19]} The modalities for treatment of CRPS include the following.

**Physical therapy**
This is the first line treatment for CRPS and has been shown to be effective in multiple controlled studies.\textsuperscript{[12]} The initial treatment is immobilization and splinting of the extremity. Elevation, massage, gentle range of motion, and isometric strengthening exercises are gradually incorporated into the treatment with provision of adequate analgesia. As the patient improves, therapy becomes more aggressive. However, therapy must not exacerbate the pain. If it leads to escalation of pain, it must be given up.\textsuperscript{[26]} Contrast baths (alternating heat and cold), transcutaneous electrical nerve stimulation (TENS), H-wave therapy, and a stress loading program of traction and compression exercises have also been shown to be effective in CRPS.\textsuperscript{[3]}

**Pharmacologic therapy**
The selection of the drug is determined by the severity of the pain. Traditional NSAID’s (ibuprofen) or COX-2 inhibitors (celecoxib) are useful in the acute stage. Metamizol and controlled release opioids (hydrocodone or oxycodone) can be considered in patients with more severe pain to allow participation in therapy.\textsuperscript{[26]} Other drugs for which data from controlled clinical trials have established efficacy in CRPS include gabapentin, Ca-modulating drugs (nifidipine, amlodipine, calcitonin, and bisphosphonates), and free radical scavengers.\textsuperscript{[33]} Bisphosphonates (alendronate, pamidronate, and clodronate) are the only class of drugs that have survived placebo controlled trials and shown a statistically significant pain reduction in patients with CRPS.\textsuperscript{[32,34]} Randomized controlled trials from the Netherlands have

| Drug        | Initial dose | Dosage (mg) | Maintenance dose | Side-Effects                                           |
|-------------|--------------|-------------|-----------------|-------------------------------------------------------|
| Oxycodone   | 10 mg tid    | 10-80 mg tid| Tapering dose   | Nausea, dry mouth, hives, hallucination               |
| Gabapentin  | 100 mg tid   | 300-600 mg tid| Tapering dose   | Ataxia, dizziness, somnolence, purpura, fatigue       |
| Nifedipine  | 10 mg tid    | 10-30 mg tid|                 | Headache, postural hypotension                         |
| Alendronate | 40 mg OD     | 40 mg OD    |                 | Nausea, dizziness, heartburn, jaw numbness             |
| Amtriptiline| 10 mg ON     | 25-75 mg ON |                 | Drowsiness                                             |
| Prednisone  | 30 mg tid    | Tapering dose|                 | Adrenal suppression, avascular necrosis                |
| Phenytoin   | 50 mg OD     | 50-100 mg tid|                 | Ataxia, liver damage, convulsion                       |
| Pregabalin  | 50 mg tid    | 50-200 mg tid|                 | Dizziness, somnolence, peripheral oedema               |
| Fluoxetine  | 20 mg OD     | 20-80 mg OD |                 | Headache, postural hypotension                         |
shown promising results with the use of free radical scavengers [topical application of 50% dimethylsulfoxide (DMSO) and oral N-acetylcysteine(NAC)] in the treatment of CRPS.[25] High dose vitamin C (500 mg/day) has been shown to have a prophylactic role in diminishing the incidence of CRPS in patients with distal radius fractures.[26] There is also substantial anecdotal evidence for the efficacy of antidepressants (amitriptyline, nortriptyline, and doxepine), anticonvulsants (dilantin), opioids, topical capsaicin, and lignocaine transdermal in the treatment of CRPS.[29] The dosage and major complications of use of the common pharmacologic agents has been summarized in Table 5.

**Regional anaesthesia techniques**

These are useful in patients with moderate to severe pain that do not respond to physical and pharmacologic therapy. Two types of regional anaesthesia techniques are available, namely a sympathetic nerve block and a combined somatic and sympathetic nerve block. Sympathetic nerve blocks are chosen when the patient has marked improvement after a diagnostic sympathetic block. A somatic plus sympathetic block is used in patients who do not respond to the diagnostic sympathetic block.
Physical therapy should be initiated immediately after the block. A series of daily or every alternate day blocks using local anaesthetic agents is usually required for 1-3 weeks. Blocks are considered successful if the patient can participate in physical therapy and progresses in rehabilitation. If severe pain persists beyond 3 weeks, long-term pain-relieving options may need to be considered.[12]

Neuromodulation
It involves modulation of central pain pathways by delivery of an electrical current or chemical application to the central neural axis. Neuromodulation techniques include spinal cord and peripheral nerve stimulation, TENS, and intrathecal injection of baclofen, clonidine, or opioids. These techniques are invasive and should be reserved for patients in whom other measures have failed.[12]

Surgery
The benefit of surgical (and chemical sympathectomy) in CRPS has not been confirmed by controlled studies. In addition these procedures are associated with significant complications and therefore are not currently recommended in the treatment of CRPS.[3,7] However, surgery has a role in management of nerve related nociceptive foci. The most common neural diagnoses contributing to CRPS are neuroma, neuroma-in-continuity, and secondary compression neuropathies.[3,7]

One needs to address the nerve, the underlying tissue bed, and the overlying skin/scar. The nerve should be explored by an extensible incision, freed of all adhesions, and evaluated using appropriate magnification. A complete release of the involved nerve should be done in patients with secondary compression neuropathies. Internal neurolysis should be avoided. In patients with a nerve transection or a neuroma-in-continuity, a tension free repair should be done and nerve grafts used, if required. Sural nerve grafts are preferred to medial or lateral antebrachial cutaneous nerves to avoid creating another nociceptive focus in the same extremity. Local fat, muscle or fascial flaps can be used to resurface a scarred bed. The nerve repair site can be wrapped using a vein graft or an artificial conduit. The adhesions between the skin and nerve can be managed by a Z-plasty, local transposition flaps, or a distant flap. It is important to secure meticulous haemostasis and avoid the use of constrictive postoperative dressings. The aim is to initiate physical therapy immediately after surgery and appropriate continuous sympathetic blocks and/or drugs should be used for pain relief.[3,7]

An algorithm for the treatment of CRPS is depicted in Figure 5.[12]

CONCLUSIONS
The diagnosis of CRPS should be made cautiously. In patients who develop CRPS after surgery of the upper limb, one must look for an underlying nerve related nociceptive focus and treat it early before it gets to the chronic phase. Although outcomes for patients with CRPS are very difficult to predict, an early diagnosis and treatment increase the likelihood of a successful outcome. Mild cases respond to physical therapy, whereas moderate cases may require adjuvant analgesics, such as gabapentin and/or an anti-depressant medication. Patients with severe pain and/or sympathetic dysfunction require regional anaesthetic blockade to participate in physical therapy. A small percentage of patients with refractory, chronic pain will require long-term multidisciplinary treatment, including physical therapy, psychological support, and pain-relieving measures.

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