Early experience with hand continuous passive motion

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S Morris, A Freiberg, L Harris. Early experience with hand continuous passive motion. Can J Plast Surg 1993;1(1):19-23. Clinical experience with continuous passive motion (CPM) has increased in recent years. However, little information is available in the literature of objective evaluation of the results obtained using this treatment modality. The purpose of this study was to review both the indications for hand CPM at the Toronto Western Hospital, Toronto, Ontario, and the course and the outcome of the patients treated. Between 1984 and 1989 the Mobilimb H1 CPM was used on 43 patients for a mean period of 39±6 days (mean total hours of CPM 741±84 h). Indications for hand CPM included hand trauma, capsulectomy and tenolysis and other hand conditions. Mean follow-up was 32±3 months. Overall, hand CPM was well tolerated and highly effective in relieving hand pain and increasing active and passive range of motion. Compliance was excellent.

Key Words: Continuous passive motion (CPM), Hand surgery, Indications, Outcome, Patient compliance

Expérience préliminaire avec mouvement passif continu de la main

RÉSUMÉ: L'expérience clinique portant sur le mouvement passif continu (MPC) n'a cessé de s'étendre au cours des dernières années. La littérature renferme cependant peu de renseignements sur l'évaluation objective des résultats obtenus à l'aide de cette modalité thérapeutique. Le but de la présente étude a été de passer en revue tant les indications du MPC de la main au Toronto Western Hospital de Toronto, en Ontario, que l'évolution et l'issue du traitement chez les patients. Entre 1984 et 1989, le MPC Mobilimb H1 a été utilisé chez 43 patients durant une période moyenne de 396 jours (nombre d'heures totales moyen de MPC 7414 heures). Les indications de cette modalité thérapeutique incluaient: traumatisme de la main, capsulectomie, ténolyse et autres problèmes de la main. La période de suivi moyenne a été de 323 mois.a'De façon globale, le MPC de la main a été bien toléré et très efficace pour le soulagement de la douleur et l'augmentation de l'amplitude du mouvement actif et passif. L'observance au traitement a été excellente.

ince the pioneering laboratory research by Salter (1-5) on the effects of continuous passive motion on the healing and regeneration of articular cartilage, the clinical applications of continuous passive motion (CPM) have broadened to include the knee, hip, elbow, shoulder and temporomandibular joints (6-11). Recently, interest has steadily increased in the use of hand CPM (Figure 1). Therapists and surgeons have been reluctant to employ this modality and there is a paucity of information in the literature reporting results obtained using hand CPM. Several authors have reported on experience with isolated indications for hand CPM (12-17). No long term studies are available, however, to document the course and outcome of patients treated with hand CPM. This review studies the indications, course and outcome in patients treated with hand CPM used as an adjunct to hand surgery. The majority of these cases represent salvage situations in which hand CPM was used in the presence of devastating

hand injuries. A number of the reconstructive cases would not have been attempted had hand CPM not been available.

MATERIALS AND METHODS

All patients who were placed on hand CPM at Toronto Western Hospital between 1984 and 1989, and who were available for follow-up examination and interview, were included in the study. Of a total of 43 patients, 36 were available for review, while seven were lost to follow-up. These 36 patients were interviewed and a standard questionnaire was administered. As well, all inpatient, outpatient and therapy charts were reviewed. Hand dominance, injured hand, type of injury, age, sex, type of surgery, duration and method of CPM use and overall outcome were assessed. Each patient was asked to complete an assessment of pain and outcome according to a visual analogue scale. This test was administered by a person other than the patient's surgeon or therapist. The patient was asked to mark a cross on a 10 cm line to indicate their assessment of overall outcome, function, pain and satisfaction (0 = low or unfavourable; 10 = high or favourable). This test was selected to determine in an unbiased manner the patient's subjective impression of the CPM device. In all 43

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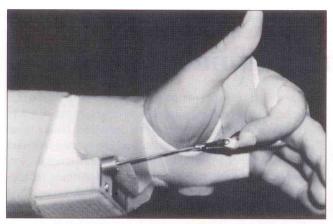


Figure 1) The Mobilimb H1 continuous passive motion (CPM) unit, in situ

patients the Mobilimb CPM unit was used (H1 or H2 models, Toronto Medical Corporation, Pickering, Ontario). All surgical procedures were performed or directly supervised by the senior author (AF).

Postoperatively, the hand CPM unit was applied within the first 24 h, often in the operating room or recovery room. The patients were instructed on application, removal and care of the CPM device prior to discharge from hospital. All patients were advised to use the hand CPM device 23 h per day. The Mobilimb CPM unit was used for periods of 11 to 140 days (39 \pm 6 days; mean \pm SEM), and was removed when the hand active range of motion (ROM) matched that achieved on the CPM unit. If the ROM was not maintained during the first week post-CPM, hand CPM was restarted. Frequent follow-up was arranged when possible in the hand surgery clinic and hand therapy unit. A hand therapist supervised all aspects of hand CPM use. ROM results are herein reported according to the Strickland formula for proximal (PIP) and distal interphalangeal (DIP) joint motion and as total active motion (TAM) (18). The Mobilimb CPM unit was chosen for use in these patients because it is durable, lightweight and well accepted by patients.

Statistical analysis of the data was performed with a Macintosh SE microcomputer. Paired and unpaired t tests (two-tailed) were used to compare two means. Multiple comparisons of means were performed using one-way analysis of variance (ANOVA) or two-way ANOVA. The level of probability accepted as significant was P<0.05. Variation of means is reported as standard error unless otherwise specified.

RESULTS

Indications for the use of hand CPM were divided into three groups. Group A: trauma (total nine patients, 16 digits) severe lacerations (n=3), crush injuries (n=4), fractures (n=l) and joint injuries (1); Group B: reconstructive (total 22 patients, 26 digits) metacarpal phalangeal (MCP) arthroplasty (n=5), extensor tenolysis and proximal interphalangeal (PIP) joint capsulectomy (n=17); and Group C: miscellaneous (total five patients, nine digits) reflex sympathetic dystrophy

TABLE 1: Duration of hand CPM unit use

	Acute Group A	Elective Group B	Total
Duration (days) (range)	56±15	33±5	39±6
	(14 to 140)	(11 to 96)	(11 to 140)
Time/day	18±1	19±1	19±1
(range)	(10 to 23)	(8 to 24)	(8 to 24)
Total hours (range)	1008±222	627±78	741±84
	(196 to 2160)	(152 to 1440)	(152 to 2160)

Values are presented as means ± standard error

TABLE 2: Removal of CPM machine by patient during therapy period

Removed/day	Acute Group A	Elective Group B	Total
0 to 2	5	7	12
3 to 5	4	11	15
>5	0	4	4

No significant difference in splint removal was detected between group A (acute post injury) and group B (elective postoperative)

TABLE 3: Reasons cited by patients for the removal of hand CPM device

Reason	Number of patients
Prevents ADL	13
Pain/fingernail problems	12
Sleep disruption	10
Cumbersome	7
Unable to wear at work	2
Cost (batteries)	2
Machine breakdown	2
Swelling	1

ADL Activities of daily living

(n=2), intra-articular sepsis (n=1), tenosynovectomy (n=1) and Dupytren's contracture (1).

In group A, patients were started on CPM because the complexity of injuries would otherwise prohibit early active physiotherapy. Group B patients were, in general, started on CPM in order to maintain the ROM achieved in the operating room. Group C patients had unusually resistant hand surgical problems which did not respond to conventional therapy. Since group C is so diverse it will not be statistically assessed as an individual group and omitted in ROM outcome studies.

Patients were seen either in the emergency room acutely or were referred to the Hand Surgery Clinic for evaluation. The mean age at surgery was 38±2 years. There were 22 males and 14 females. Patients were treated with hand CPM for a mean duration of 39±6 days and most were compliant (mean daily duration of CPM therapy was 19±1 h, Table 1). The mean total duration of CPM treatment was 741±84 h (range 152 to 2160 h). The mean follow-up was 32±3 months (range seven to 72 months). Patients were off work for a mean of 190±53 days (range zero to 1071). Five patients (8%) were unable to return to their former employment. Nine of the patients (23%) had suffered injuries at work and were followed by the Workman's Compensation Board.

TABLE 4: Patient satisfaction with hand CPM using physician administered visual analogue type questionnaire

	Acute (%)	Elective (%)	Total (%)
Outcome	71±6	66±6	68±4
Function	70±6	67±5	68±4
Pain on CPM	7±2	23±5	18±4
Satisfaction with CPM	79±9	71±6	73±5

No significant difference was detected between the acute post injury (group A) and elective postoperative (group B) groups (ANOVA)

TABLE 5: Range of motion (ROM) outcome (Strickland formula) for patients treated with hand CPM in group A (acute post injury) and group B (elective reconstruction)

	Acute Group A	Elective Group B	Total
Number of patients	9	22	
Number of digits	16	26	42
DIP, PIP range preoperatively (degrees)	N/A	28±4	28 <u>±</u> 4
DIP, PIP range postoperatively (degrees)	63±7	67±5	66±7

There was a significant ($P \le 0.05$) improvement in group B patients postoperatively compared to preoperative ROM (paired t test)

The CPM device was removed zero to two times per day in 12 patients, three to five times per day in 15 patients and greater than five times per day in four patients (Table 2). Reasons cited by the patient for removal of the CPM unit included: fingernail pain and/or deformity, sleep disruption, awkwardness, prevention of activities of daily living and inability to work (Table 3). No wound breakdowns were observed. No significant difference between overall outcome or ROM was observed between the compliant and noncompliant patients. Overall, there was excellent compliance and a high level of satisfaction among patients with the CPM machine. In order to assess the patients' subjective acceptance of the hand CPM, a visual analogue scale was used. In general, the patients reported that CPM of the hand reduced pain, increased active and passive range of motion and improved overall outcome (Table 4).

In group A, the final ROM (Strickland formula) was 63±7 (Table 5). In group B, the mean ROM was 28±4 preoperatively and 67±5 postoperatively. In group A, the final TAM was 164±29 (Table 6). The mean TAM was 112±14 preoperatively in group B and 186±22 postoperatively. Overall, results were judged to be excellent in 29%, good in 39%, fair in 26% and failure in 6% of cases (Table 7).

DISCUSSION

Traditional management of injured or inflamed joints, bones or tendons included prolonged periods of immobilization. Salter demonstrated that CPM increased periosteal chondrogenesis, augmented extrasynovial tendon healing and improved outcome in intra-articular sepsis. The proposed

TABLE 6: Total active motion (TAM) of patients treated with hand CPM in group A (acute post injury) and group B (elective reconstruction)

	Acute Group A	Elective Group B	Total
Number of digits	16	26	42
TAM preoperatively	N/A	112±14	112±14
TAM postoperatively	164±29	186±22	167±25

There was a significant (P<0.05) improvement in group B patients postoperatively compared to preoperative ROM (paired t test)

TABLE 7: Subjective assessment of overall outcome by patient and surgeon (in parentheses) in patients treated with hand CPM in group A (acute post injury) and group B (elective reconstruction)

Outcome	Group A Acute	Group B Elective	Total	Percentage
Excellent	4 (4)	5 (5)	9 (9)	29
Good	4 (1)	11 (11)	15 (12)	39
Fair	1 (3)	4 (4)	5 (7)	26
Poor	0	0	0	0
Failure	0	2 (2)	2 (2)	6

No significant difference in outcome between outcome in the two groups or between the assessment by patient and surgeon was detected (two-way ANOVA)

mechanism for these beneficial effects includes the prevention of synovial adhesions, increased diffusion of nutrients via the increased synovial fluid uptake, stimulation of living chondrocytes, and clearance of lysozymal enzymes and purulent exudate from the synovial cavity (1-5). A wide variety of applications have been evaluated. The most impressive clinical results have emerged using CPM for postoperative rehabilitation after intra-articular fractures (6), knee ligament repairs (7), osteotomies of the hip or knee (19,20) and elbow (8,9) and temporomandibular joint surgery (10,11).

Ketchum (14) initially reported experience with the electrically driven hand splint in 1972. This device was developed to extend passively the fingers for durations of 6 to 8 h per day in patients who had suffered extensor tendon injuries which had been surgically repaired. The use of this creative prototype was later evaluated in 426 stiff joints in 142 patients (15) and found to improve significantly the TAM and total passive motion.

Clearly, the indications for the use of hand CPM are evolving. A summary of the available literature is presented in Table 8. Prosser et al (21) studied the use of CPM in 22 patients who had suffered hand injuries including fractures, joint and crush injuries, and septic arthritis. The authors found that CPM treatment was generally effective; however the beneficial effects plateaued (ROM) approximately seven days after commencing. It was concluded that CPM can be a useful adjunct to hand therapy in a wide variety of hand conditions with restricted ROM. Other authors have recommended hand CPM for the following: patients in whom pain and edema persists longer than expected post injury or post-

TABLE 8: Indications for hand CPM

	Author (reference)
Acute	, and the first transfer to the first transfer transfer to the first transfer t
Burns	Covey, 1987 (12)
	Kealey, 1987 (24)
Intra-articular sepsis, fractures,	Prosser, 1987 (21)
crush injuries	
Flexor tendon repairs	Gelberman, 1990 (16)
Extensor tendon repairs	Bunker, 1990 (17)
Joint injuries	Ketchum, 1972 (15)
out injuries	Prosser, 1989 (21)
Reconstructive	
Tenolysis and capsulotomy	Cannon, 1989 (22)
	Scirven, 1987 (25)
Rheumatoid and implant surgery	Merritt, 1987 (32)
Reflex sympathetic dystrophy	Gelberman and
	Dimick, 1987 (33)

operatively; patients in whom passive range of motion is not full preoperatively; patients in whom an extensive tenolysis is required; and in fearful patients (22).

A recent survey of therapists at 100 burn centres revealed that 44% of respondents (response rate 54%) used hand CPM in burn patients (23). The hand CPM units used by these therapists included the Kintec, Stryker, Richards, Chattanooga, Toronto Medical Corporation Mobilimb, Therakinetics, Danniger and Invacare models. Covey et al (12) reported the use of hand CPM (Sutter 5000) in 10 burn patients and suggested that indications for the use of hand CPM include extensive burn involvement of multiple kinetic areas, patients who are unable to participate actively in rehabilitation and in patients in whom excessive pain or anxiety precludes normal hand therapy (12,24).

Scirven et al (25) reviewed the use of hand CPM in 120 patients for a variety of indications including joint release, tendon repair, tenolysis, Dupytren's contracture release and PIP or MCP joint replacement. They used three CPM machines, (Sutter, Zimmer and Richards) and found a dramatic relief of postoperative pain and edema. Evidence had accumulated in experimental and clinical studies that indicates that intermittent passive motion is beneficial to functional outcome of repaired flexor tendons (18,26-30). Gelberman (16) found that the use of hand CPM following flexor tendon repairs resulted in significantly improved ROM compared to traditional early passive motion treatment. A subsequent uncontrolled study of hand CPM use in Verdan zone II flexor tendon injuries produced good or excellent results in 86% of patients (17). McCarthy et al (31) found that chicken feet treated with continuous passive motion as an adjunct to tenolysis had an overall reduction in the ROM compared to animals which were treated with restricted activity.

Several reports of hand CPM use in rheumatoid arthritis hand reconstruction have emerged (32,33). To date, this work is preliminary and inconclusive; however, it appears that hand CPM may aid in the reduction of joint motion after arthroplasty and in rehabilitation following synovectomy and soft tissue reconstruction.

In the present review we describe the use of hand CPM in acute and elective reconstructive settings. It appears from the subjective evaluations that the CPM was well accepted by patients, principally because it relieved pain. Various regimens for use of hand CPM have been described. An early application of the machine in the initial 24 h postoperatively is important to improve patient's acceptance. In the present series, it is noteworthy that there did not appear to be a significant difference in outcome between the patients who were very compliant (used CPM machines 23 h per day or longer) and those who were less compliant and used their machines for shorter durations. The patient's progress and state of wound healing are the most important determinants of the duration of CPM required. Each patient's care should be individualized based on close hand therapy follow-up. If the patient's pain and swelling persist, then CPM may be continued for several weeks. In crush injuries and other salvage situations, CPM may be continued for several months, often at the request of the patient. When the range of motion off CPM equals that on CPM, and when postoperative pain and edema are resolving, use of the machine may be discontinued.

Since Ketchum's earliest description of the electric hand splint and the developmental work done by Toronto Medical Corporation, a number of models of hand CPM have become available. Cannon et al (22) reported use of the Sutter model hand CPM and has found that the ability to adjust the force, speed and arc of motion are important features. The problem with all hand CPM units available, however, is that multiple joints will be affected by a single deforming force leaving the more damaged joints to move the least. The Sutter CPM 5000 unit has been used for capsulectomy of the PIP joint (28) flexor tendon repairs (29) and for hand burns (12). In a review of the hand therapy units in 100 burn units, the Sutter was the most commonly used for burn rehabilitation (23). Bunker has noted difficulty in obtaining full extension of the PIP joint, adequate mobilization of the DIP joint and optimal adhesion of the CPM unit to the fingertips in flexor tendon repairs (17). Despite this, excellent results were obtained.

In this review, hand CPM appears to provide excellent pain relief and improved results in salvage situations. The disadvantages include the requirement for active ROM exercises by a trained hand therapist, and the cost of the unit. The theoretical problem with hand CPM today, which has not been answered by any of the existing units, includes the deficiency of tendon excursion through the tendon sheaths of the hand. As well, in the three joint digital skeletal system, one joint may not move through a complete range of movement despite maximal motion of the machine. Inadequate tendon excursion in the hand with the CPM machine pushes and does not pull the flexor tendon, and hence the tendon may become adherent to the surrounding sheath lining. Bunker (17), for example, found the relative lack of movement at the DIP joint to be a problem. This problem could be addressed by adding an electrical nerve stimulation device which could stimulate the profundus muscle synchronously with flexion excursion of the hand CPM device. Other important unanswered questions persist. What are the best indications for use of hand CPM? What is the optimal duration and frequency of CPM use? What are the requirements for active hand therapy when hand CPM is used?

CONCLUSIONS

In certain salvage situations, CPM of the hand appears to be effective in providing a painless range of motion in otherwise stiff and painful digits. CPM of the hand does not appear to provide adequate excursion of the flexor tendons and

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hence should be combined with active hand therapy. The patient must be intermittently monitored while on hand CPM to avoid the prolonged lack of progression at a certain joint which may be ameliorated by splint adjustment. (Therefore, close hand therapy follow-up will both provide active ROM exercises and splint surveillance). High compliance was a consistent feature of hand CPM. This may be due to the fact that the device actually relieved pain in the vast majority of patients. For a variety of difficult hand surgical problems good or excellent results were obtained in 68% of patients treated with postoperative hand CPM.

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