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Notice of Independent Review Decision

DATE OF REVIEW: MARCH 9, 2010

IRO CASE #:

DESCRIPTION OF THE SERVICE OR SERVICES IN DISPUTE

Medical necessity of proposed 6 sessions of psychotherapy (90806)

A DESCRIPTION OF THE QUALIFICATIONS FOR EACH PHYSICIAN OR OTHER HEALTH CARE PROVIDER WHO REVIEWED THE DECISION

This case was reviewed by a clinician with a Ph.D. in clinical Psychology and who is licensed in the State of Texas. The reviewer specializes in general psychology and behavioral pain management and is engaged in full time practice.

REVIEW OUTCOME

Upon independent review the reviewer finds that the previous adverse determination/adverse determinations should be:

- Upheld (Agree)
- Overturned (Disagree)
- Partially Overturned (Agree in part/Disagree in part)

Primary Diagnosis	Service being Denied	Billing Modifier	Type of Review	Units	Date(s) of Service	Amount Billed	Date of Injury	DWC Claim#	IRO Decision
307.89	90806		Prosp	6					Upheld

INFORMATION PROVIDED TO THE IRO FOR REVIEW

TDI-HWCN-Request for an IRO-17 pages

Respondent records- a total of 54 pages of records received to include but not limited to: TDI letter 2.16.10; Group letters 12.7.09, 1.4.10, 2.10.10, 2.12.10, 2.17.10; Request for an IRO forms; report 1.4.10, 2.12.10; Healthcare records 8.25.09-12.24.09; Reviews report 12.7.09

Requestor records- a total of 82 pages of records received to include but not limited to:
TDI letter 2.16.10;A- Healthcare records 7.6.04-1.19.10; FCE 2.6.06

PATIENT CLINICAL HISTORY [SUMMARY]:

Records indicate that the claimant is a female who was injured at work on xx/xx/xx. At the time, she was lifting heavy boxes when she felt severe pain in her back. She was taken off work, and appears to remain in an off-work status.

Claimant has received the following diagnostics and treatments to date: X-rays, MRI, lumbar fusion x2, TENS unit, physical therapy, massage therapy, injections, psychological testing, chronic pain programs x 2, and medications management. Office notes and psychological eval give diagnoses as: failed surgery syndrome, lumbar facet syndrome, depression, and sleep disturbance. Medications that have been prescribed include Hydrocodone, Soma, Ultram, Elavil, and Amitriptyline. Patient has treatment plan to wean off Hydrocodone and Soma, but this has yet to be accomplished.

Patient was referred for psychological eval on 09/01/09, and patient was interviewed and evaluated by Healthcare Systems in order to make psychological treatment recommendations. As a result, patient was diagnosed with 307.89 chronic pain disorder, 311.0 depressive disorder, and 300.0 anxiety disorder. Request is for individual therapy 1x6 overall goals of: "increasing coping with negative emotions, stress relief, management of pain, and expression of emotions." Long term goals are listed as: stabilization of depressed/anxious/irritable mood, independent utilization of stress management skills, increase confidence in managing pain flare-ups, and compliance with medication reduction plan.

ANALYSIS AND EXPLANATION OF THE DECISION INCLUDE CLINICAL BASIS, FINDINGS AND CONCLUSIONS USED TO SUPPORT THE DECISION. IF THERE WAS ANY DIVERGENCE FROM DWC'S POLICIES/GUIDELINES OR THE NETWORK'S TREATMENT GUIDELINES, THEN INDICATE BELOW WITH EXPLANATION.

Patient has been advanced through the stepped-care approach recommended by ODG, and having completed a pain program, should be at, or close to, MMI. Although severe clinical depression can require medication, and/or therapy follow-up visits, there is no rationale presented as to why patient would benefit at this time from more of the same intervention when her pain continues to be reported as 7/10 on average, and she continues to have elevated BDI and BAI scores. There is also no information regarding the CPMP outcomes to determine whether she failed these programs, was partially successful, etc., and no explanation of whether or not previous failed attempts could be modified somehow to provide for better intervention in the future. As such, request cannot be considered medically appropriate at this time.

ODG Work Loss Data, 2010, Texas

Psychological evaluations: Recommended. Psychological evaluations are generally accepted, well-established diagnostic procedures not only with selected use in pain problems, but also with more widespread use in subacute and chronic pain populations. Diagnostic evaluations should distinguish between conditions that are preexisting, aggravated by the current injury or work related. Psychosocial evaluations should determine if further psychosocial interventions are indicated. The interpretations of the evaluation should provide clinicians with a better understanding of the patient in their social environment, thus allowing for more effective rehabilitation. ([Main-BMJ, 2002](#)) ([Colorado, 2002](#)) ([Gatchel, 1995](#)) ([Gatchel, 1999](#)) ([Gatchel, 2004](#)) ([Gatchel, 2005](#))

Aetna Clinical Policy Bulletins. Chronic Pain Programs Number 0237. Reviewed: May 5, 2006.

Aetna considers a screening examination medically necessary for members who are being considered for admission into a chronic pain program.

1. Outpatient Pain Management Programs

Aetna considers outpatient multidisciplinary pain management programs medically necessary when all of the following criteria are met:

- * Referral for entry has been made by the primary care physician/attending physician; and
- * Member has experienced chronic non-malignant pain (not cancer pain) for 6 months or more; and
- * The cause of the member's pain is unknown or attributable to a physical cause, i.e., not purely psychogenic in origin; and
- * Member has failed conventional methods of treatment; and
- * The member has undergone a mental health evaluation, and any primary psychiatric conditions have been treated, where indicated; and
- * Member's work or lifestyle has been significantly impaired due to chronic pain; and
- * If a surgical procedure or acute medical treatment is indicated, it has been performed prior to entry into the pain program.

Aetna considers entry into an outpatient multidisciplinary chronic pain program not medically necessary for members with any of the following contraindications:

- * The member is unable to understand and carry out instructions; or
- * The member exhibits aggressive and/or violent behavior; or
- * The member exhibits imminently suicidal tendencies; or
- * The member has unrealistic expectations of what can be accomplished from the program (i.e., member expects an immediate cure); or
- * The member is medically unstable (e.g., due to uncontrollable high blood pressure, unstable congestive heart failure, or other medical conditions); or
- * Member has previously failed an adequate multidisciplinary (e.g., Commission on Accreditation of Rehabilitation Facilities (CARF) accredited) chronic pain management program.

Pain is considered chronic if it results from a chronic pathological process, has recurred periodically over months or years, or persists longer than expected after an illness or injury. Typically, pain is considered chronic if it has persisted for 6 months or more.

Modality-oriented pain clinics and single disciplinary pain clinics are considered not medically necessary and inappropriate for comprehensive treatment of members with chronic pain.

Note: Dependence or addiction to narcotics or other controlled substances is frequently part of the presentation of a member with chronic pain. Issues surrounding addiction, detoxification must be considered and evaluated prior to enrollment of a member into a pain management program.



FRP's: Recommended, although research is still ongoing as to how to most appropriately screen for inclusion in these programs. Functional restoration programs (FRPs), a type of treatment included in the category of interdisciplinary pain programs (see [Chronic pain programs](#)), were originally developed by Mayer and Gatchel. FRPs were designed to use a medically directed, interdisciplinary pain management approach geared specifically to patients with chronic disabling occupational musculoskeletal disorders. These programs emphasize the importance of function over the elimination of pain. FRPs incorporate components of exercise progression with disability management and psychosocial intervention. Long-term evidence suggests that the benefit of these programs diminishes over time, but still remains positive when compared to cohorts that did not receive an intensive program. ([Bendix, 1998](#)) A Cochrane review suggests that there is strong evidence that intensive multidisciplinary rehabilitation with functional restoration reduces pain and improves function of patients with low back pain. The evidence is contradictory when evaluating the programs in terms of vocational outcomes. ([Guzman 2001](#)) It must be noted that all studies used for the Cochrane review excluded individuals with extensive radiculopathy, and several of the studies excluded patients who were receiving a pension, limiting the generalizability of the above results. Studies published after the Cochrane review also indicate that intensive programs show greater effectiveness, in particular in terms of return to work, than less intensive treatment. ([Airaksinen, 2006](#)) There appears to be little scientific evidence for the effectiveness of multidisciplinary biopsychosocial rehabilitation compared with other rehabilitation facilities for neck and shoulder pain, as opposed to low back pain and generalized pain syndromes. ([Karjalainen, 2003](#)) Treatment is not suggested

for longer than 2 weeks without evidence of demonstrated efficacy as documented by subjective and objective gains. For general information see [Chronic pain programs](#).

Biopsychosocial model of chronic pain; ODG Pain section, December, 2009

See [Chronic pain programs](#) (functional restoration programs), which are recommended where there is access to programs with proven successful outcomes, for patients with conditions that put them at risk of [delayed recovery](#), including the detailed "Criteria for use of multidisciplinary pain management programs" highlighted in blue. *Definition:* The biopsychosocial model, first proposed by George Engel, MD, acknowledges the important interplay between the biological, psychological, and social systems in illness. While disease is defined as the objective effect of pathology, illness includes the patient's perception of lack of health. An exclusively biomedical focus on objective pathology and disease is of limited usefulness in conditions like chronic pain. A focus on the patient's illness, which includes his or her psychological reactions and social function, may lead to more effective involvement in treatment, with diminished disability, improved function, and diminished co-morbidity. The model focuses on disease and illness, with illness being viewed as an interaction of biological (physiological), psychological and social factors. Disease is defined as the objective event that involves the actual pathology. Pain is experience as a unique experience, and a range of psychological and socioeconomic factors can modulate physical pathology to affect symptoms and subsequent disability. The model is utilized in interdisciplinary pain clinics as patients with chronic pain are at increased risk for emotional disorders, maladaptive cognitions, functional deficits, nociceptive dysregulation, and physical deconditioning. See also [Psychosocial adjunctive methods](#) in the Mental Illness & Stress Chapter

Chronic pain/functional restoration programs (2009)

Recommended where there is access to programs with proven successful outcomes (i.e., decreased pain and medication use, improved function and return to work, decreased utilization of the health care system), for patients with conditions that have resulted in delayed recovery. There should be evidence that a complete diagnostic assessment has been made, with a detailed treatment plan of how to address physiologic, psychological and sociologic components that are considered components of the patient's pain. Patients should show evidence of motivation to improve and return to work, and meet the patient selection criteria outlined below. While these programs are recommended (see criteria below), the research remains ongoing as to (1) what is considered the "gold-standard" content for treatment; (2) the group of patients that benefit most from this treatment; (3) the ideal timing of when to initiate treatment; (4) the intensity necessary for effective treatment; and (5) cost-effectiveness. It has been suggested that interdisciplinary/multidisciplinary care models for treatment of chronic pain may be the most effective way to treat this condition. ([Flor, 1992](#)) ([Gallagher, 1999](#)) ([Guzman, 2001](#)) ([Gross, 2005](#)) ([Sullivan, 2005](#)) ([Dysvik, 2005](#)) ([Airaksinen, 2006](#)) ([Schonstein, 2003](#)) ([Sanders, 2005](#)) ([Patrick, 2004](#)) ([Buchner, 2006](#)) These treatment modalities are based on the biopsychosocial model, one that views pain and disability in terms of the interaction between physiological, psychological and social factors. ([Gatchel, 2005](#)) See [Biopsychosocial model of chronic pain](#).

Types of programs: There is no one universal definition of what comprises interdisciplinary/multidisciplinary treatment. These pain rehabilitation programs (as described below) combine multiple treatments, and at the least, include psychological care along with physical and/or occupational therapy (including an active exercise component as opposed to passive modalities). The most commonly referenced programs have been defined in the following general ways ([Stanos, 2006](#)):

(1) **Multidisciplinary programs:** Involves one or two specialists directing the services of a number of team members, with these specialists often having independent goals. These programs can be further subdivided into four levels of pain programs:

- (a) Multidisciplinary pain centers (generally associated with academic centers and include research as part of their focus)
- (b) Multidisciplinary pain clinics

(c) Pain clinics

(d) Modality-oriented clinics

(2) **Interdisciplinary pain programs:** Involves a team approach that is outcome focused and coordinated and offers goal-oriented interdisciplinary services. Communication on a minimum of a weekly basis is emphasized. The most intensive of these programs is referred to as a Functional Restoration Program, with a major emphasis on maximizing function versus minimizing pain. See [Functional restoration programs](#).

Types of treatment: Components suggested for interdisciplinary care include the following services delivered in an integrated fashion: (a) physical treatment; (b) medical care and supervision; (c) psychological and behavioral care; (d) psychosocial care; (e) vocational rehabilitation and training; and (f) education.

Outcomes measured: Studies have generally evaluated variables such as pain relief, function and return to work. More recent research has begun to investigate the role of comorbid psychiatric and substance abuse problems in relation to treatment with pain programs. Recent literature has begun to suggest that an outcome of chronic pain programs may be to “demedicalize” treatment of a patient, and encourage them to take a more active role in their recovery. These studies use outcomes such as use of the medical care system post-treatment. The role of the increasing use of opioids and other medications (using data collected over the past decade) on outcomes of functional restoration is in the early stages, and it is not clear how changes in medication management have affected outcomes, if at all. (See [Opioids for chronic pain](#).)

Outcomes (in terms of body parts)

Neck and Shoulder: There are limited studies about the efficacy of chronic pain programs for neck, shoulder, or upper extremity musculoskeletal disorders. ([Karjalainen, 2003](#)) This may be because rates of cervical claims are only 20-25% of the rates of lumbar claims. In addition, little is known as to chronicity of outcomes. Researchers using PRIDE Program (Progressive Rehabilitation Institute of Dallas for Ergonomics) data compared a cohort of patients with cervical spine disorders to those with lumbar spine disorders from 1990-1995 and found that they had similar outcomes. Cervical patients were statistically less likely to have undergone pre-rehabilitative surgery. ([Wright, 1999](#))

Multidisciplinary back training: (involvement of psychologists, physiotherapists, occupational therapists, and/or medical specialists). The training program is partly based on physical training and partly on behavioral cognitive training. Physical training is performed according to the “graded activity” principle. The main goal is to restore daily function. A recent review of randomized controlled studies of at least a year’s duration found that this treatment modality produced a positive effect on work participation and possibly on quality of life. There was no long-term effect on experienced pain or functional status (this result may be secondary to the instrument used for outcome measure). Intensity of training had no substantial influence on the effectiveness of the treatment. ([van Geen, 2007](#)) ([Bendix, 1997](#)) ([Bendix, 1998](#)) ([Bendix2, 1998](#)) ([Bendix, 2000](#)) ([Frost, 1998](#)) ([Harkapaa, 1990](#)) ([Skouen, 2002](#)) ([Mellin, 1990](#)) ([Haldorsen, 2002](#))

Intensive multidisciplinary rehabilitation of chronic low back pain: The most recent Cochrane study was withdrawn from the Cochrane (3/06) as the last literature search was performed in 1998. Studies selected included a physical dimension treatment and at least one other treatment dimension (psychological, social, or occupational). Back schools were not included unless they included the above criteria. There was strong evidence that intensive multidisciplinary biopsychosocial rehabilitation with functional restoration improved function when compared to inpatient or outpatient nonmultidisciplinary rehabilitation. Intensive (> 100 hours), daily interdisciplinary rehabilitation was moderately superior to noninterdisciplinary rehabilitation or usual care for short- and long-term functional status (standardized mean differences, -0.40 to -0.90 at 3 to 4 months, and -0.56 to -1.07 at 60 months). There was moderate evidence of pain reduction. There was contradictory evidence regarding vocational outcome. Less intensive programs did

not show improvements in pain, function, or vocational outcomes. It was suggested that patients should not be referred to multidisciplinary biopsychosocial rehabilitation without knowing the actual content of the program. ([Guzman, 2001](#)) ([Guzman-Cochrane, 2002](#)) ([van Geen, 2007](#)) ([Bendix, 1997](#)) ([Bendix, 1998](#)) ([Bendix2, 1998](#)) ([Bendix, 2000](#)) ([Frost, 1998](#)) ([Harkapaa, 1990](#)) ([Skouen, 2002](#)) ([Mellin, 1990](#)) ([Haldorsen, 2002](#))

Multidisciplinary biopsychosocial rehabilitation for subacute low back pain among working age adults: The programs described had to include a physical component plus either a psychological, social and/or vocational intervention. There was moderate evidence of positive effectiveness for multidisciplinary rehabilitation for subacute low back pain and that a workplace visit increases effectiveness. The trials included had methodological shortcomings, and further research was suggested. ([Karjalainen, 2003](#))

Role of opioid use: See [Chronic pain programs, opioids](#).

Role of comorbid psych illness: Comorbid conditions, including psychopathology, should be recognized as they can affect the course of chronic pain treatment. In a recent analysis, patients with panic disorder, antisocial personality disorder and dependent personality disorder were > 2 times more likely to not complete an interdisciplinary program. Personality disorders in particular appear to hamper the ability to successfully complete treatment. Patients diagnosed with post-traumatic stress disorder were 4.2 times more likely to have additional surgeries to the original site of injury. ([Dersh, 2007](#)) The prevalence of depression and anxiety in patients with chronic pain is similar. Cohort studies indicate that the added morbidity of depression and anxiety with chronic pain is more strongly associated with severe pain and greater disability. ([Poleshuck, 2009](#)) ([Bair, 2008](#))

Predictors of success and failure: As noted, one of the criticisms of interdisciplinary/multidisciplinary rehabilitation programs is the lack of an appropriate screening tool to help to determine who will most benefit from this treatment. Retrospective research has examined decreased rates of completion of functional restoration programs, and there is ongoing research to evaluate screening tools prior to entry. ([Gatchel, 2006](#)) There is need for research in terms of necessity and/or effectiveness of counseling for patients considered to be “at-risk” for post-discharge problems. ([Proctor, 2004](#)) The following variables have been found to be negative predictors of efficacy of treatment with the programs as well as negative predictors of completion of the programs: (1) a negative relationship with the employer/supervisor; (2) poor work adjustment and satisfaction; (3) a negative outlook about future employment; (4) high levels of psychosocial distress (higher pretreatment levels of depression, pain and disability); (5) involvement in financial disability disputes; (6) greater rates of smoking; (7) increased duration of pre-referral disability time; (8) higher prevalence of opioid use; and (9) elevated pre-treatment levels of pain. ([Linton, 2001](#)) ([Bendix, 1998](#)) ([McGeary, 2006](#)) ([McGeary, 2004](#)) ([Gatchel2, 2005](#)) ([Dersh, 2007](#))

Role of duration of disability: There is little research as to the success of return to work with functional restoration programs in long-term disabled patients (> 24 months).

Studies supporting programs for patients with long-term disability: Long-term disabled patients (at least 18 months) vs. short-term disabled (4 to 8 months) were evaluated using Pride data (1990-1993). No control was given for patients that did not undergo a program. During the time studied program dropouts averaged 8% to 12%. (It does appear that at the time of this study, participants in the program were detoxified from opioids prior to beginning.) The long-term disabled group was more likely to have undergone spinal surgery, with this likelihood increasing with time. Return to work was statistically different between the short-term disabled (93%) and the long-term disabled-18 months (80%). The long-term disabled-24 months group had a 75% return to work. Long-term disabled-18 month patients were statistically more likely to visit new health providers than short-term disabled patients (34% and 25% respectively). Work retention at one year in groups up to 24 months duration of disability was 80%. This dropped to 66% in the group that had

been disabled for > 24 months. The percentage of recurrent lost time injury claims increased from around 1% in the groups disabled for < 35 months to 8.3% in the groups disabled for > 36 months. A main criterion for success appeared to be the decision of the patient to actively participate in the program rehabilitation goals.

([Jordan, 1998](#))

Studies suggesting limited results in patients with long-term disability: While early studies have suggested that time out-of-work is a predictor of success for occupational outcomes, these studies have flaws when an attempt is made to apply them to chronic pain programs. ([Gallagher, 1989](#)) ([Beals, 1972](#)) ([Krause, 1994](#)) Washington State studied the role of duration of work injury on outcome using a statistical model that allowed for a comparison of patients that participated in a multidisciplinary pain program (using data from 1991-1993) vs. those that were evaluated and not treated. This was not an actual study of time of disability, but of duration of injury (mean years from injury to evaluation of 2.6 years for the treated group and 4.0 years for the evaluated only group). The original statistical analysis allowed for a patient to be included in a “treated group” for those individuals that both completed and did not complete the program. Data was collected from 10 sites. Each of the centers was CARF approved and included Pysch/behavioral treatment, vocation counseling and physical therapy. A sub-study evaluated a comparison of patients that were treatment completers vs. those that did not participate (78.6%, N=963). No information was given in terms of surgical procedures or medications. The primary outcome was time loss status of subjects 2 years after they had undergone the index pain center evaluation. In the 2001 study, if chronicity of duration of injury was controlled for, there was no significant benefit produced in terms of patients that were receiving time-loss benefits at 2-years post treatment between the two groups. Approximately 60% of both groups were not receiving benefits at the two-year period. As noted, the “treated patient” was only guaranteed to have started a program. A repeat analysis of only the patients who completed the study did not significantly change the results of the study. In a 2004 survey follow-up no significant difference was found between treated and untreated groups, although the treated group had better response. The survey response was 50%, and the treatment responders were more likely to be disabled at the time of the survey. The authors suggest that the results indicated early intervention was a key to response of the programs, and that modest goals (improvement, not cure) be introduced. ([Robinson, 2004](#)) ([Robinson, 2001](#)) [The authors also concluded that there was no evidence that pain center treatment affects either disability status or clinical status of injured workers.]

Timing of use: Intervention as early as 3 to 6 months post-injury may be recommended depending on identification of patients that may benefit from a multidisciplinary approach (from programs with documented positive outcomes). See [Chronic pain programs, early intervention](#).

Role of post-treatment care (as an outcome): Three variables are usually examined; (1) New surgery at the involved anatomic site or area; (2) Percentage of patients seeking care from a new provider; (3) Number of visits to the new provider over and above visits with the health-care professional overseeing treatment. It is suggested that a “new provider” is more likely to reorder diagnostic tests, provide invasive procedures, and start long-term analgesics. In a study to determine the relationship between post-treatment healthcare-seeking behaviors and poorer outcomes (using prospectively analyzed PRIDE data on patients with work-related musculoskeletal injuries), patients were compared that accessed healthcare with a new provider following functional restoration program completion (approximately 25%) to those that did not. The former group was significantly more likely to have an attorney involved with their case (22.7% vs. 17.1%, respectively), and to have had pre-rehabilitation surgery (20.7% vs. 12.1%, respectively). Return to work was higher in the group that did not access a new provider (90% vs. 77.6% in the group that did access). The group that did not access new providers also was more likely to be working at one year (88% vs. 62.2% in the group that accessed new providers). It should be noted that 18% of the patients that entered the program

dropped out or were asked to leave. The authors suggested monitoring of additional access of healthcare over and above that suggested at the end of the program, with intervention if needed. (Proctor, 2004)

See also [Chronic pain programs, intensity](#); [Chronic pain programs, opioids](#); [Functional restoration programs](#); & [Chronic pain programs, early intervention](#).

Criteria for the general use of multidisciplinary pain management programs:

Outpatient pain rehabilitation programs may be considered medically necessary in the following circumstances:

- (1) The patient has a chronic pain syndrome, with evidence of loss of function that persists beyond three months and has evidence of three or more of the following:
 - (a) Excessive dependence on health-care providers, spouse, or family;
 - (b) Secondary physical deconditioning due to disuse and/or fear-avoidance of physical activity due to pain;
 - (c) Withdrawal from social activities or normal contact with others, including work, recreation, or other social contacts;
 - (d) Failure to restore preinjury function after a period of disability such that the physical capacity is insufficient to pursue work, family, or recreational needs;
 - (e) Development of psychosocial sequelae that limits function or recovery after the initial incident, including anxiety, fear-avoidance, depression, sleep disorders, or nonorganic illness behaviors (with a reasonable probability to respond to treatment intervention);
 - (f) The diagnosis is not primarily a personality disorder or psychological condition without a physical component;
 - (g) There is evidence of continued use of prescription pain medications (particularly those that may result in tolerance, dependence or abuse) without evidence of improvement in pain or function.
- (2) Previous methods of treating chronic pain have been unsuccessful and there is an absence of other options likely to result in significant clinical improvement.
- (3) An adequate and thorough multidisciplinary evaluation has been made. This should include pertinent validated diagnostic testing that addresses the following:
 - (a) A physical exam that rules out conditions that require treatment prior to initiating the program. All diagnostic procedures necessary to rule out treatable pathology, including imaging studies and invasive injections (used for diagnosis), should be completed prior to considering a patient a candidate for a program. The exception is diagnostic procedures that were repeatedly requested and not authorized. Although the primary emphasis is on the work-related injury, underlying non-work related pathology that contributes to pain and decreased function may need to be addressed and treated by a primary care physician prior to or coincident to starting treatment;
 - (b) Evidence of a screening evaluation should be provided when addiction is present or strongly suspected;
 - (c) Psychological testing using a validated instrument to identify pertinent areas that need to be addressed in the program (including but not limited to mood disorder, sleep disorder, relationship dysfunction, distorted beliefs about pain and disability, coping skills and/or locus of control regarding pain and medical care) or diagnoses that would better be addressed using other treatment should be performed;
 - (d) An evaluation of social and vocational issues that require assessment.
- (4) If a goal of treatment is to prevent or avoid controversial or optional surgery, a trial of 10 visits may be implemented to assess whether surgery may be avoided.
- (5) If a primary reason for treatment in the program is addressing possible substance use issues, an evaluation with an addiction clinician may be indicated upon entering the program to establish the most appropriate treatment approach (pain program vs. substance dependence program). This must address evaluation of drug abuse or diversion (and prescribing drugs in a non-therapeutic manner). In this particular case, once drug abuse or diversion issues are addressed, a 10-day trial may help to establish a diagnosis, and determine if the patient is not better suited for treatment in a substance dependence program. Addiction consultation can be incorporated into a pain program. If there is indication that substance dependence may be a problem, there should be evidence that the program has the capability to address this type of pathology prior to approval.
- (6) Once the evaluation is completed, a treatment plan should be presented with specifics for treatment of identified problems, and outcomes that will be followed.

(7) There should be documentation that the patient has motivation to change, and is willing to change their medication regimen (including decreasing or actually weaning substances known for dependence). There should also be some documentation that the patient is aware that successful treatment may change compensation and/or other secondary gains. In questionable cases, an opportunity for a brief treatment trial may improve assessment of patient motivation and/or willingness to decrease habituating medications.

(8) Negative predictors of success (as outlined above) should be identified, and if present, the pre-program goals should indicate how these will be addressed.

(9) If a program is planned for a patient that has been continuously disabled for greater than 24 months, the outcomes for the necessity of use should be clearly identified, as there is conflicting evidence that chronic pain programs provide return-to-work beyond this period. These other desirable types of outcomes include decreasing post-treatment care including medications, injections and surgery.

(10) Treatment is not suggested for longer than 2 weeks without evidence of compliance and significant demonstrated efficacy as documented by subjective and objective gains. (Note: Patients may get worse before they get better. For example, objective gains may be moving joints that are stiff from lack of use, resulting in increased subjective pain.) However, it is also not suggested that a continuous course of treatment be interrupted at two weeks solely to document these gains, if there are preliminary indications that they are being made on a concurrent basis.

(11) Integrative summary reports that include treatment goals, compliance, progress assessment with objective measures and stage of treatment, must be made available upon request at least on a bi-weekly basis during the course of the treatment program.

(12) Total treatment duration should generally not exceed 20 full-day (160 hours) sessions (or the equivalent in part-day sessions if required by part-time work, transportation, childcare, or comorbidities). (Sanders, 2005) Treatment duration in excess of 160 hours requires a clear rationale for the specified extension and reasonable goals to be achieved. Longer durations require individualized care plans explaining why improvements cannot be achieved without an extension as well as evidence of documented improved outcomes from the facility (particularly in terms of the specific outcomes that are to be addressed).

(13) At the conclusion and subsequently, neither re-enrollment in repetition of the same or similar rehabilitation program (e.g. work hardening, work conditioning, out-patient medical rehabilitation) is medically warranted for the same condition or injury (with possible exception for a medically necessary organized detox program). Prior to entry into a program the evaluation should clearly indicate the necessity for the type of program required, and providers should determine upfront which program their patients would benefit more from. A chronic pain program should not be considered a “stepping stone” after less intensive programs, but prior participation in a work conditioning or work hardening program does not preclude an opportunity for entering a chronic pain program if otherwise indicated.

(14) Suggestions for treatment post-program should be well documented and provided to the referral physician. The patient may require time-limited, less intensive post-treatment with the program itself. Defined goals for these interventions and planned duration should be specified.

(15) Post-treatment medication management is particularly important. Patients that have been identified as having substance abuse issues generally require some sort of continued addiction follow-up to avoid relapse.

Inpatient pain rehabilitation programs: These programs typically consist of more intensive functional rehabilitation and medical care than their outpatient counterparts. They may be appropriate for patients who: (1) don't have the minimal functional capacity to participate effectively in an outpatient program; (2) have medical conditions that require more intensive oversight; (3) are receiving large amounts of medications necessitating medication weaning or detoxification; or (4) have complex medical or psychological diagnosis that benefit from more intensive observation and/or additional consultation during the rehabilitation process. (Keel,

1998) ([Kool, 2005](#)) ([Buchner, 2006](#)) ([Kool, 2007](#)) As with outpatient pain rehabilitation programs, the most effective programs combine intensive, daily biopsychosocial rehabilitation with a functional restoration approach. If a primary focus is drug treatment, the initial evaluation should attempt to identify the most appropriate treatment plan (a drug treatment /detoxification approach vs. a multidisciplinary/interdisciplinary treatment program). See [Chronic pain programs, opioids](#); [Functional restoration programs](#).

Psychological Evaluations: Recommended based upon a clinical impression of psychological condition that impacts recovery, participation in rehabilitation, or prior to specified interventions (e.g., lumbar spine fusion, spinal cord stimulator, implantable drug-delivery systems). ([Dolevs, 2003](#)) Psychological evaluations are generally accepted, well-established diagnostic procedures not only with selected use in pain problems, but also with more widespread use in subacute and chronic pain populations. Diagnostic evaluations should distinguish between conditions that are preexisting, aggravated by the current injury or work related. Psychosocial evaluations should determine if further psychosocial interventions are indicated. The interpretations of the evaluation should provide clinicians with a better understanding of the patient in their social environment, thus allowing for more effective rehabilitation. ([Main-BMJ, 2002](#)) ([Colorado, 2002](#)) ([Gatchel, 1995](#)) ([Gatchel, 1999](#)) ([Gatchel, 2004](#)) ([Gatchel, 2005](#)) For the evaluation and prediction of patients who have a high likelihood of developing chronic pain, a study of patients who were administered a standard battery psychological assessment test found that there is a psychosocial disability variable that is associated with those injured workers who are likely to develop chronic disability problems. ([Gatchel, 1999](#)) Childhood abuse and other past traumatic events were also found to be predictors of chronic pain patients. ([Goldberg, 1999](#)) Another trial found that it appears to be feasible to identify patients with high levels of risk of chronic pain and to subsequently lower the risk for work disability by administering a cognitive-behavioral intervention focusing on psychological aspects of the pain problem. ([Linton, 2002](#)) Other studies and reviews support these theories. ([Perez, 2001](#)) ([Pulliam, 2001](#)) ([Severeijns, 2001](#)) ([Sommer, 1998](#)) In a large RCT the benefits of improved depression care (antidepressant medications and/or psychotherapy) extended beyond reduced depressive symptoms and included decreased pain as well as improved functional status. ([Lin-JAMA, 2003](#)) See "[Psychological Tests Commonly Used in the Assessment of Chronic Pain Patients](#)" from the Colorado Division of Workers' Compensation, which describes and evaluates the following 26 tests: (1) BHI 2nd ed - Battery for Health Improvement, (2) MBHI - Millon Behavioral Health Inventory [has been superseded by the MBMD following, which should be administered instead], (3) MBMD - Millon Behavioral Medical Diagnostic, (4) PAB - Pain Assessment Battery, (5) MCMI-111 - Millon Clinical Multiaxial Inventory, (6) MMPI-2 - Minnesota Inventory, (7) PAI - Personality Assessment Inventory, (8) BBHI 2 - Brief Battery for Health Improvement, (9) MPI - Multidimensional Pain Inventory, (10) P-3 - Pain Patient Profile, (11) Pain Presentation Inventory, (12) PRIME-MD - Primary Care Evaluation for Mental Disorders, (13) PHQ - Patient Health Questionnaire, (14) SF 36, (15) SIP - Sickness Impact Profile, (16) BSI - Brief Symptom Inventory, (17) BSI 18 - Brief Symptom Inventory, (18) SCL-90 - Symptom Checklist, (19) BDI-II - Beck Depression Inventory, (20) CES-D - Center for Epidemiological Studies Depression Scale, (21) PDS - Post Traumatic Stress Diagnostic Scale, (22) Zung Depression Inventory, (23) MPQ - McGill Pain Questionnaire, (24) MPQ-SF - McGill Pain Questionnaire Short Form, (25) MPQ - Oswestry Disability Questionnaire, (26) Visual Analogue Pain Scale - VAS. ([Bruns, 2001](#)) Chronic pain may harm the brain, based on using functional magnetic resonance imaging (fMRI), whereby investigators found individuals with chronic back pain (CBP) had alterations in the functional connectivity of their cortical regions - areas of the brain that are unrelated to pain - compared with healthy controls. Conditions such as depression, anxiety, sleep disturbances, and decision-making difficulties, which affect the quality of life of chronic pain patients as much as the pain itself, may be directly related to altered brain function as a result of chronic pain. ([Baliki, 2008](#)) See also [Comorbid psychiatric disorders](#). See also the [Stress/Mental Chapter](#)

Psychological treatment: Recommended for appropriately identified patients during treatment for chronic pain. Psychological intervention for chronic pain includes setting goals, determining appropriateness of treatment, conceptualizing a patient's pain beliefs and coping styles, assessing psychological and cognitive function, and addressing co-morbid mood disorders (such as depression, anxiety, panic disorder, and posttraumatic stress disorder). Cognitive behavioral therapy and self-regulatory treatments have been found to be particularly effective. Psychological treatment incorporated into pain treatment has been found

to have a positive short-term effect on pain interference and long-term effect on return to work. The following “stepped-care” approach to pain management that involves psychological intervention has been suggested:

Step 1: Identify and address specific concerns about pain and enhance interventions that emphasize self-management. The role of the psychologist at this point includes education and training of pain care providers in how to screen for patients that may need early psychological intervention.

Step 2: Identify patients who continue to experience pain and disability *after the usual time of recovery*. At this point a consultation with a psychologist allows for screening, assessment of goals, and further treatment options, including brief individual or group therapy.

Step 3: Pain is sustained in spite of continued therapy (including the above psychological care). Intensive care may be required from mental health professions allowing for a multidisciplinary treatment approach. See also [Multi-disciplinary pain programs](#). See also [ODG Cognitive Behavioral Therapy \(CBT\) Guidelines](#) for low back problems. ([Otis, 2006](#)) ([Townsend, 2006](#)) ([Kerns, 2005](#)) ([Flor, 1992](#)) ([Morley, 1999](#)) ([Ostelo, 2005](#))

A DESCRIPTION AND THE SOURCE OF THE SCREENING CRITERIA OR OTHER CLINICAL BASIS USED TO MAKE THE DECISION:

- ACOEM- AMERICAN COLLEGE OF OCCUPATIONAL & ENVIRONMENTAL MEDICINE UM KNOWLEDGEBASE
- AHCPR- AGENCY FOR HEALTHCARE RESEARCH & QUALITY GUIDELINES
- XX DWC- DIVISION OF WORKERS COMPENSATION POLICIES OR GUIDELINES
- EUROPEAN GUIDELINES FOR MANAGEMENT OF CHRONIC LOW BACK PAIN
- INTERQUAL CRITERIA
- XX MEDICAL JUDGEMENT, CLINICAL EXPERIENCE AND EXPERTISE IN ACCORDANCE WITH ACCEPTED MEDICAL STANDARDS
- MERCY CENTER CONSENSUS CONFERENCE GUIDELINES
- MILLIMAN CARE GUIDELINES
- XX ODG- OFFICIAL DISABILITY GUIDELINES & TREATMENT GUIDELINES
- PRESSLEY REED, THE MEDICAL DISABILITY ADVISOR
- TEXAS GUIDELINES FOR CHIROPRACTIC QUALITY ASSURANCE & PRACTICE PARAMETERS
- TEXAS TACADA GUIDELINES
- TMF SCREENING CRITERIA MANUAL
- PEER REVIEWED NATIONALLY ACCEPTED MEDICAL LITERATURE (PROVIDE A DESCRIPTION)
- OTHER EVIDENCE BASED, SCIENTIFICALLY VALID, OUTCOME FOCUSED GUIDELINES (PROVIDE A DESCRIPTION)