## 1 **5221.6200 LOW BACK PAIN.**

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Subp. 1. Diagnostic procedures for treatment of low back injury. A health care provider shall determine the nature of the condition before initiating treatment. [For text of items A to G, see Minnesota Rules] H. Diagnostic analgesic blocks or injections studies include facet joint injection, facet nerve injection, epidural differential spinal block, nerve block, and nerve root block. (1) These procedures are used to localize the source of pain before surgery and to diagnose conditions which fail to respond to initial nonsurgical management. (2) These injections are invasive and when done as diagnostic procedures only, are not indicated unless noninvasive procedures have failed to establish the diagnosis. (3) Selection of patients, choice of procedure, and localization of the level of injection should be determined by documented clinical findings indicating possible pathologic conditions and the source of pain symptoms. (4) These blocks and injections can also be used as therapeutic modalities and as such are subject to the parameters of subpart 5. [For text of items I and J, see Minnesota Rules] [For text of subparts 2 to 4, see Minnesota Rules] Subp. 5. Therapeutic injections Injection modalities. Injection modalities are indicated as set forth in items A to C. These diagnostic and therapeutic injections are invasive and when done as diagnostic procedures only, are not indicated unless noninvasive procedures have failed to establish the diagnosis. Selection of patients, choice of procedure, and localization of the level of injection should be determined by documented clinical findings indicating possible pathologic conditions and the source of pain symptoms. Use of injections can extend past the 12-week limit on passive treatment modalities so long as the maximum treatment for injections is not exceeded. Use of therapeutic injections beyond 12 months must be in accordance with subpart 7. A. Therapeutic injections, including injections of trigger points, facet joints, facet nerves, sacroiliac joints, sympathetic nerves, epidurals, nerve roots, and peripheral nerves. Therapeutic injections can only be given in conjunction with active treatment modalities directed to the same anatomical site. (1) Trigger point injections: (a) time for treatment response, within 30 minutes; (b) maximum treatment frequency, no more than four injection sites per patient

visit. Subsequent injections may occur once per week to any one site if a positive

34	response to the first injection at that site. If subsequent injections at that site fail to
35	demonstrate progressive improvement as specified in subpart 9, diminishing
36	control of symptoms or fail to facilitate objective functional gains, then trigger
37	point injections should be redirected to other areas or discontinued. No more than
38	three injections to different sites are reimbursable per patient visit; and
39	(c) maximum treatment, four injections to any one site visits in any 12-month
40	period.
41	(2) Sacroiliac joint injections:
42	(a) time for treatment response, within one week;
43	(b) maximum treatment frequency, no more than two injections per patient visit.
44	Subsequent injections may occur once every three months can repeat injection
45	two weeks after the previous injection if a positive response to the first injection.
46	If subsequent injections fail to demonstrate progressive improvement as specified
47	in subpart 9, injections should be discontinued at that joint Only two injections are
48	reimbursable per patient visit; and
49	(c) maximum treatment, two injections to any one site four injection visits in any
50	12-month period.
51	(3) Facet Intra-articular facet joint or nerve injections, may be considered for patients
52	with persistent symptoms that have not responded to six weeks of initial nonsurgical
53	treatment as described in subpart 2, item B, subitem (1):
54	(a) time for treatment response, within one two weeks;
55	(b) maximum treatment frequency, no more than three joint levels may be
56	injected, either unilaterally or bilaterally, per patient visit. Subsequent injections
57	may occur once every two weeks to any one site three months if a positive
58	response to the first injection. If subsequent injections <u>fail to</u> demonstrate
59	progressive improvement as specified in subpart 9, diminishing control of
60	symptoms or fail to facilitate objective functional gains, then injections should be
61	discontinued at that facet joint. No more than three injections to different sites are
62	reimbursable per patient visit; and
63	(c) maximum treatment, three injections to any on site visits in any 12-month
64	period.
65	(4) Nerve root blocks:
66	(a) time for treatment response, within one week;

67	(b) maximum treatment frequency, can repeat injection two weeks after the
68	previous injection if a positive response to the first injection. Only three injections
69	to different sites are reimbursable per patient visit; and
70	(c) maximum treatment, two injections to any one site.
71	Radiofrequency denervation injections of the facet joints, may be considered after a
72	positive response to a set of two diagnostic medial branch blocks as described in item
73	B, subitem (1):
74	(a) time for treatment response, within three weeks;
75	(b) maximum treatment frequency, no more than two facet joint levels, or three
76	medial branch nerves, may be injected, either unilaterally or bilaterally, per
77	patient visit. Subsequent injections may occur six months after the previous
78	injection if a positive response to the previous injection. Before a repeat injection
79	occurs, an additional confirmatory medial branch block must be performed, as
80	specified in item B, subitem (1), if the patient's pain presents differently than in
81	the initial evaluation; and
82	(c) maximum treatment, two injection visits in any 12-month period.
83	(5) Epidural injections:
84	(a) time for treatment response, within one two weeks;
85	(b) maximum treatment frequency, no more than one level may be injected, either
86	unilaterally or bilaterally, per patient visit. Subsequent injections may occur once
87	every two weeks if a positive response to the first injection. If subsequent
88	injections <u>fail to</u> demonstrate <u>progressive improvement as specified in subpart 9</u> ,
89	diminishing control of symptoms or fail to facilitate objective functional gains,
90	then injections should be discontinued at that level. Only one injection is
91	reimbursable per patient visit; and
92	(c) maximum treatment, three four injections visits in any 12-month period.
93	B. Permanent lytic or sclerosing injections, including radio frequency denervation of the facet
94	joints. These injections can only be given in conjunction with active treatment modalities
95	directed to the same anatomical site:
96	(1) time for treatment response, within one week;
97	(2) maximum treatment frequency, may repeat once for any; and
98	(3) maximum duration, two injections to any one site.

99	Diagnostic-only injections, including medial branch blocks and herve root blocks. These
100	injections may only be done as a diagnostic procedure and must not be used as an ongoing
101	therapeutic modality.
102 103 104 105	(1) Medial branch blocks, may be considered for patients with persistent symptoms that have not responded to six weeks of initial nonsurgical treatment as described in subpart 2, item B, subitem (1). These injections may be used to assess if a particular facet joint is the cause of symptoms and if the patient would benefit from other treatment modalities:
106	(a) time for treatment response, immediately or within one day;
107 108 109 110 111	(b) maximum treatment frequency, no more than two facet joint levels, or three medial branch nerves, either unilaterally or bilaterally, may be injected per patient visit. A confirmatory second injection to the same medial branch nerve may occur no sooner than one week after the initial injection if there is a positive response to the first injection; and
112 113	(c) maximum treatment, no more than two injections to any single medial branch nerve.
114 115	(2) Nerve root blocks, may be used to assess if a particular nerve root is the cause of symptoms and if the patient would benefit from other treatment modalities:
116	(a) time for treatment response, immediately or within one day;
117 118	(b) maximum treatment frequency, no more than one nerve root may be injected per patient visit; and
119 120	(c) maximum treatment, no more than one injection to any single nerve root.  Subsequent injections must be to an alternative nerve root.
121 122	C. Prolotherapy and botulinum toxin injections are not indicated in the treatment of low back problems and are not reimbursable.