

1 **Clinical Practice Guideline:** **Quality Patient Management**

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3 **Date of Implementation:** **April 24, 2003**

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5 **Product:** **Specialty**

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8 Promoting quality patient management is an American Specialty Health – Specialty
9 (ASH) mandate. ASH requires that contracted practitioners adhere to reasonable practice
10 parameters. Diagnosis/evaluation and treatment are two significant parameters of clinical
11 decision-making. The practitioner must demonstrate a clinically appropriate approach to
12 his/her clinical decision-making process. This approach is dependent upon the clinical
13 knowledge and experience of the practitioner, his/her skill in clinical assessment,
14 deductive reasoning, and pattern recognition.

15
16 This document, in conjunction with Risk Factor Assessment as described in policy
17 Patient Safety – The Prevention, Recognition, and Management of Adverse Outcomes -
18 QM 7 - S, will assist the practitioner in understanding the level of assessment and
19 documentation that is appropriate and how this documentation demonstrates clinical
20 practices consistent with ASH-approved practice parameters and management of
21 expected clinical outcomes. Once received by ASH, the clinical data found in the
22 submitted documents serves as the basis for the clinical quality evaluator’s assessment of
23 the practitioner’s clinical decision-making and treatment utilization.

24
25 **Practitioner Involvement in the ASH Clinical Services Program**

26 With the exception of services covered under the practitioner’s applicable clinical
27 performance system, clinical information including current pertinent subjective and
28 objective clinical findings must be submitted to ASH for verification of medical necessity
29 of additional services. The practitioner must include adequate patient demographic
30 information to accurately identify the patient as a member; use of ASH forms is strongly
31 encouraged to ensure adequate information is submitted. In the event that the
32 documentation is either illegible or incomplete, Medical Necessity/Benefit
33 Administration (MNA) staff contacts the practitioner for clarification. The practitioner is
34 allowed the opportunity to provide the necessary information; failure to do so could result
35 in an administrative non-approval of the submitted treatment/services. Upon successful
36 administrative review, the documentation is sent to a licensed, credentialed, peer clinical
37 quality evaluator for verification of medical necessity.

38
39 Treating practitioners are expected to have ongoing communication with a referring
40 health care provider, where applicable, and co-management of the patient’s episode of
41 care between practitioners is expected. Factors that may affect the expected response of

1 the member are considered. Examples include: surgical procedures, member age, co-
2 morbidities, past medical history, response to previous treatment, mode of onset, severity,
3 and psychosocial and occupational factors. ASH does not set diagnosis-specific treatment
4 frequency or duration limitations. Each case is evaluated considering all pertinent clinical
5 evidence for that member's unique clinical situation. It is understood that similar case
6 presentations may be handled in similar fashion with reasonably consistent results. For a
7 given diagnosis the effect of variability in general health status (age, gender, past medical
8 history, psychosocial factors, and presence of co-morbid conditions) make the use of
9 diagnosis-specific treatment duration and frequency limits inherently untenable. If the
10 member has previously accessed a benefit managed by ASH, the results of previous case
11 evaluations are available to the clinical quality evaluator.

12
13 The practitioner is notified by fax of one of three potential outcomes of the evaluation of
14 the submitted treatment/services. These are: *Approval*, *Partial Approval*, or *Non-*
15 *approval* of the submitted treatment/services. The notification includes the name,
16 telephone number, and extension of the clinical quality evaluator who completed the
17 evaluation. Clinical quality evaluators are available by telephone to respond to any
18 questions or inquiries regarding the clinical services program or a specific issue related to
19 a case. In the event the fax transmission is unsuccessful, expedient alternatives for the
20 delivery of the response form are implemented.

21
22 *Approval*: ASH clinical quality evaluators have the responsibility to approve appropriate
23 care as medically necessary. The clinical quality evaluators assess the clinical data
24 supplied by the practitioner in order to determine whether the initiation or continuation of
25 care has been documented as medically necessary. The purpose of the practitioner's
26 initial assessment and subsequent assessments of the member is to estimate the treatment
27 plan/program needs of the member. The ASH practitioner is accountable to document the
28 medical necessity of all services submitted/provided. It is the responsibility of the peer
29 clinical quality evaluator to evaluate the documentation in accordance with their training,
30 understanding of practice parameters, and review criteria adopted by ASH.

31
32 *Partial Approval*: Occurs when only a portion of the submitted treatment/services is
33 initially approved. The partial approval may refer to a decrease in treatment frequency,
34 treatment duration, DME/supply/appliance, or type of services submitted. This decision
35 may be due to the practitioner's documentation of findings that are inconsistent with the
36 clinical conclusion, and/or treatment dosage (frequency/duration) is not supported by the
37 underlying diagnostic or clinical features. In many cases, the clinical documentation
38 supplied only provides sufficient information to establish the need to initiate a trial of
39 care. In these circumstances, the clinical quality evaluator will provide a partial approval
40 of the practitioner's submitted treatment/services. Additional submitted
41 treatments/services may be reviewed after evaluating the progress of the initial

1 treatment/services. These procedures allow the member to receive appropriate care but
2 take into account the variable responses that a member may have to the clinical
3 intervention. If the practitioner disagrees with the partial approval to the submitted
4 treatment/services, he/she may contact the clinical quality evaluator listed on their
5 response form to discuss the case, submit additional documentation utilizing the reopen
6 process, or submit additional documentation to appeal the decision.

7
8 *Non-approval:* Occurs when none of the submitted treatment/services are approved. The
9 most common causes for a non-approval of treatment/services are
10 administrative/contractual (e.g., ineligibility, reached plan benefit limits). It is appropriate
11 not to approve ongoing treatment/services if the member's condition is no longer
12 responding favorably to the treatment/services being rendered by the treating practitioner.

13 14 **Additional Care**

15 Approval of an additional course of treatment/services requires submission of additional
16 information, including patient response and updated clinical findings. In those cases
17 where an additional course of treatment/services is submitted, the decision to approve
18 additional treatment/services will be based on the following criteria:

- 19 • The member has made clinically significant progress under the initial treatment
20 plan/program. Clinically significant progress may be noted on a reliable and valid
21 outcome tool. Determining that progress is clinically significant requires
22 correlation with the overall clinical presentation, including updated subjective and
23 objective examination findings.
- 24 • Additional clinically significant progress can be reasonably expected by
25 continued treatment.
- 26 • The member has not reached maximum therapeutic benefit (MTB) or maximum
27 medical improvement (MMI).
- 28 • There is no indication that immediate care/evaluation is required by other health
29 care professionals.

30
31 Any exacerbation or flare-up of the condition that contributes to the need for additional
32 treatment/services must be documented.

33 34 **Supportive Care**

35 Supportive care is defined as treatment/services for members having reached MTB and
36 for whom periodic trials for therapeutic withdrawal fail to sustain previous therapeutic
37 gains that would otherwise progressively deteriorate. Supportive care follows appropriate
38 application of passive and active care, including lifestyle modification.

39
40 It is appropriate to approve additional treatment/services if the need for supportive care is
41 demonstrated. Supportive care may be inappropriate when it interferes with other

1 appropriate primary care or when the risk of supportive care (e.g., practitioner
2 dependence, somatization, illness behavior, or secondary gain) outweighs its benefits.

3
4 Ancillary diagnostic procedures should be selected based on clinical history and
5 examination findings that suggest the necessity to rule out underlying pathology or to
6 confirm a diagnosis that cannot be verified through less invasive methods.

- 7 • Information is expected to directly impact the treatment/services and course of
8 care
- 9 • Benefit of the procedure outweighs the risk to the member's health (short and
10 long term)
- 11 • Procedure is sensitive and specific for the condition being evaluated (e.g., an
12 appropriate procedure is utilized to evaluate for pathology).

13 14 **Clinical Decision-Making Process**

15 The goals of the clinical decision making process, which occur at both the practitioner-
16 member interface and the interface between practitioner and ASH, are to review for
17 approval, as appropriate, those clinical treatments/services necessary to return the
18 member to pre-clinical/pre-morbid health status or stabilize a chronic condition.

19
20 The clinical information the clinical quality evaluator expects to see when evaluating the
21 documentation in support of the medical necessity of submitted treatment/services may
22 include but is not limited to:

- 23 • History:
 - 24 ○ Past and familial history
 - 25 ○ Chief complaint
 - 26 ▪ Onset/Duration
 - 27 □ Type/Mechanism
 - 28 □ Insidious/Spontaneous
 - 29 ▪ Initial date of onset/surgery
 - 30 ▪ Stage/Nature/Cause(s)
 - 31 □ Acute, sub-acute, chronic
 - 32 □ Initial occurrence, exacerbation, chronic recurrent
 - 33 ▪ Severity of pain/functional limitation
 - 34 ▪ Frequency of pain/functional limitation
 - 35 ○ Other co-morbidity and medical or surgical management
- 36
37 • Physical Examination/Evaluation [commensurate with the nature and severity of
38 the presenting complaint(s) and scope of the practitioner of services]:
 - 39 ○ General review of systems
 - 40 ○ Orthopedic assessment
 - 41 ○ Neurological assessment

- 1 ○ Biomechanical assessment
- 2 ○ Functional outcome measure
- 3 ○ Nutritional assessment
- 4 ○ Psychosocial/Lifestyle
- 5 ○ Specialty/situation –specific evaluation (traditional oriental medicine, ADLs,
- 6 disability/impairment rating, etc.)

8 **Outcome Expectations Considered in Case Evaluation**

9 Within the context of the expected natural progression of the condition and considering
 10 member compliance, submitted treatment/services are evaluated to see if they are
 11 expected and likely to:

- 12 • Increase rate or quality of tissue repair;
- 13 • Accelerate return to functional status or stabilize functional capacities;
- 14 • Decrease time to reach pre-clinical status, if clinically appropriate;
- 15 • Substantially decrease or resolve pain and/or other symptoms;
- 16 • Decrease or prevent adverse sequelae or complications;
- 17 • Reduce or eliminate risk of relapse or recurrence.

18
 19 Ancillary diagnostic procedures should be selected based on clinical history and
 20 examination findings that suggest the necessity to rule out underlying pathology or to
 21 confirm a diagnosis that cannot be verified through less invasive methods.

- 22 • Information derived from the procedures is expected to directly impact the
 23 treatment/services and course of care
- 24 • The benefit of the procedure should outweigh the risk to the member's health
 25 (short and long term)
- 26 • The procedure must be sensitive and specific for the condition being evaluated
 27 (e.g., an appropriate procedure is utilized to evaluate for pathology).

29 **Principles of Practitioner Clinical Services**

30 The first principle of clinical services is to facilitate the early return to activity with
 31 associated reduction of symptoms, decrease of impairments, and the restoration of
 32 function. A second principle is that care should provide for improvement/recovery more
 33 efficiently than if no care had been delivered (improve upon the expected natural
 34 progression of the condition). The third principle is that chronicity should be prevented
 35 whenever possible. Psychosocial warning signs and/or over-dependence on the
 36 practitioner should be evaluated and monitored as appropriate. A fourth principle is that
 37 repeated use or reliance on acute care measures alone may foster chronicity, practitioner
 38 dependence by the member, and over-utilization of the practitioner's services. In
 39 addition, the use of passive modalities that have redundant physiological effect should
 40 not be employed.

1 The level of the patient's compliance with the recommended treatment regimen can affect
2 the outcome of passive or active care.

3 Passive Care: Treatment/care that is rendered to the patient by the practitioner.

4 Active Care: Treatment/care performed by the patient (e.g., therapeutic exercise
5 program or lifestyle modification).

6
7 The planning for therapeutically necessary care can be divided into four phases, each
8 having distinct objectives.

9
10 Phase 1: Acute Intervention

- 11 A. To promote anatomical rest
- 12 B. To diminish severity of acute symptoms
- 13 C. To reduce inflammation
- 14 D. To alleviate pain

15
16 Phase 2: Remobilization/Functional Improvement

- 17 A. To increase the range of pain-free motion/ADL function
- 18 B. To minimize deconditioning

19
20 Phase 3: Rehabilitation

- 21 A. To restore strength and endurance
- 22 B. To increase physical work capacity

23
24 Phase 4: Lifestyle Adaptations

- 25 A. To modify social and recreational activity, if necessary
- 26 B. To diminish work environmental risk factors
- 27 C. To adapt psychological factors affecting or altered by the disorder.

28
29 The practitioner should keep in mind that these phases will overlap and there may not be
30 clear cut delineation among them. It is beneficial to proceed to the remobilization phase
31 (if warranted) as rapidly as possible and to minimize dependency upon passive forms of
32 treatment/care.

33
34 In general, the initiation of care is warranted if there are no contraindications to
35 prescribed care, there is reasonable evidence to suggest the efficacy of the prescribed
36 intervention, and the intervention is within the scope of services permitted by State or
37 Federal law. The treatment submission for a disorder is typically structured in time-
38 limited increments (e.g., 30-day increments with follow-up reporting every 30 days).
39 However, when the practitioner discovers that a member is non-responsive to the applied
40 interventions within a two (2) week interval, re-evaluation and treatment modification
41 should be implemented and documented.

1 Successful management of the member's treatment plan/program involves the effective
2 exchange of clinical information between the contracted practitioner and the clinical
3 quality evaluator. By following these practice parameters and review criteria as decision-
4 assist tools, the contracted practitioner will effectively interact within the ASH clinical
5 services evaluation system.