

Phase II And Phase III Cancer Clinical Trials

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1.0 DESCRIPTION

The Department of Defense (DoD) Cancer Prevention and Treatment Clinical Trials Demonstration was conducted from 1996 through March 2008 to improve access to promising new cancer therapies, assist in meeting the National Cancer Institute's (NCI) clinical trial goals, and to assist in the formulation of conclusions regarding the safety and efficacy of emerging therapies in the prevention and treatment of cancer. This Demonstration included Phase II and Phase III protocols sponsored by the NCI for the prevention, screening, early detection, and treatment of all types of cancer (see the TRICARE Operations Manual (TOM), [Chapter 18, Section 2](#)). The Demonstration is to end on March 31, 2008 and applicable coverage guidance has been incorporated into this policy. A new Interagency Agreement between DoD and the NCI has been entered into which is effective April 1, 2008.

2.0 POLICY

2.1 Cancer clinical trial participation is authorized for those TRICARE-eligible patients selected to participate in NCI-sponsored Phase II and Phase III studies for the prevention, screening, early detection, and treatment of cancer. TRICARE will cost-share all medical care and testing required to determine eligibility for an NCI-sponsored trial, including the evaluation for eligibility at the institution conducting the NCI-sponsored study. TRICARE will cost-share all medical care required as a result of participation in NCI sponsored studies. This includes purchasing and administering all approved chemotherapy agents (except for NCI-funded investigational drugs), all inpatient and outpatient care, including diagnostic and laboratory services not otherwise reimbursed under an NCI grant program if the following conditions are met:

- 2.1.1** The provider seeking treatment for a TRICARE-eligible beneficiary in an NCI approved protocol has obtained preauthorization for the proposed treatment before initial evaluation; and
- 2.1.2** Such treatments are NCI sponsored Phase II or Phase III protocols; and
- 2.1.3** The patient continues to meet entry criteria for said protocol; and
- 2.1.4** The institutional and individual providers are TRICARE-authorized providers.

3.0 POLICY CONSIDERATIONS

3.1 Referral by Attending Physician

The attending physician, Primary Care Manager (PCM), or oncologist shall determine the eligible patient's needs and consult with the TRICARE contractor's cancer clinical trials case manager/NCI to determine which, if any, Phase II or Phase III, NCI-sponsored studies are appropriate for the patient.

3.2 Identification of Eligible NCI-sponsored Clinical Trials

3.2.1 NCI sponsorship of clinical trials occurs through the Cancer Therapy Evaluation Program (CTEP), Cooperative Group Studies, NCI Grants or Cancer Center Studies. Evidence of NCI sponsorship in one of these categories will be that it is identified in the NCI comprehensive database, Physicians's Data Query (PDQ), or NCI supplements to that database; formal notification of approval from The Clinical Protocol Review and Monitoring Committee; or verification from the NCI project officer; or through protocols co-sponsored by the NCI and other Federal Agencies.

3.2.2 Unlike the NCI-sponsored protocols for CTEP, Cooperative Group Studies, or NCI Grants, protocols for Cancer Center Studies are not individually reviewed by the NCI. Instead, the NCI designates specific institutions as meeting NCI criteria for clinical and comprehensive cancer centers. Cancer center protocols receive approval through an NCI approved institutional peer review and quality control system at the institution. Protocols which have been through this process receive formal notification of approval from The Clinical Protocol Review and Monitoring Committee and, therefore, are considered NCI sponsored, but may not appear in the PDQ. A provider who is seeking to enter a patient into a Cancer Center Study must provide evidence of NCI sponsorship by forwarding the formal notification of approval from this specific committee. Formal notification of approval by the Clinical Protocol Review and Monitoring Committee will be required for approval of treatment in Cancer Center Studies which are not otherwise sponsored through the CTEP program, NCI cooperative groups, or NCI grants.

3.2.3 Certain protocols listed in the PDQ may not be clearly identified in terms of NCI sponsorship. Clinical trials conducted as part of an NCI grant, or those identified with a "V" number, must be verified for NCI sponsorship with the NCI project officer. Physicians who are holders of the grant at the institution must provide written clarification that the proposed treatment is a protocol under their NCI grant. The grant title and number must be specified.

3.2.4 Requests for treatment in clinical trials overseas must be verified as to NCI sponsorship with the NCI project officer.

3.2.5 Protocols that are co-sponsored by the NCI and other Federal Agencies must be verified by the NCI project officer.

3.2.6 Some NCI-sponsored clinical trials are designated as multiple-phased trials (e.g., Phase I/II). Multi-phase NCI-sponsored clinical trials are eligible for TRICARE coverage as long as the beneficiary is a participant in a trial phase that would normally be covered in a single-phase trial.

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3.3 The DoD has no authority regarding the NCI protocol eligibility for the sponsored study. Therefore, if a patient does not meet the protocol eligibility criteria for enrollment, appeal rights do not apply.

3.4 Retroactive authorizations can be authorized in accordance with the provisions outlined in [32 CFR 199.4\(g\)\(19\)](#).

3.5 Claims will be paid from the applicable underwritten Contract Line Item Number (CLIN) and submitted through normal TRICARE Encounter Data (TED) system processing as required in the TRICARE Systems Manual (TSM) with the applicable coding for cancer clinical trials with enrollment effective on or after April 1, 2008.

3.6 Normal TRICARE eligibility, reimbursement, co-payments, cost-shares, deductibles, TRICARE for Life (TFL), and double coverage rules apply.

3.7 The contractor shall:

3.7.1 Provide a registered nurse to serve as case manager for inquiries and actions pertinent to the cancer clinical trials benefit.

3.7.2 Ensure the provider has submitted a letter on the facility's letterhead:

3.7.2.1 Provide the patient's name and the last four digits of the sponsor's Social Security Number (SSN); and

3.7.2.2 Certify the protocol is an NCI-sponsored study and providing the title and phase of the protocol and the NCI number of the protocol and/or other appropriate evidence of NCI sponsorship; and

3.7.2.3 Certify the patient meets all entry criteria for said protocol; and

3.7.2.4 Certify notification will be provided to the contractor's cancer clinical trials benefit case manager of the patient's registration date when treatment actually begins; and

3.7.2.5 Certify notification will be provided to the contractor's cancer clinical trials benefit case manager if the patient becomes ineligible for the study prior to the treatment.

3.7.3 Utilize the NCI's Comprehensive Cancer Database known as the Physician's Data Query (PDQ), to assist in determining whether a particular study meets the requirements of the cancer clinical trials benefit and whether the patient is eligible for a particular protocol. For those studies that are not listed on the PDQ, the contractor will work with NCI staff to verify NCI sponsorship and phase of the study.

3.8 The contractor may at its discretion establish a dedicated toll-free telephone number to receive inquiries from both patients and providers regarding the cancer clinical trials benefit. If a dedicated toll-free telephone number is established, the phone shall be staffed seven hours a day during normal business hours in the contractor's time zone where the inquiries are received. In the absence of a dedicated toll-free number for cancer clinical trials benefit inquiries, contractors shall

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use their primary toll-free telephone inquiry system (see the TOM, Chapter 11, Section 7 and Chapter 20, Section 4).

The contractor may at its discretion establish a dedicated mailing address where cancer clinical trials benefit inquiries and claims shall be sent for expedited response and/or claims adjudication. In the absence of a dedicated mailing address for cancer clinical trials benefit inquiries and claims, contractors shall use their primary address(es) for written correspondence and claims (see the TOM, [Chapter 11, Sections 5 and 6](#), and [Chapter 20, Section 4](#)).

3.9 The Cancer Clinical Trials Demonstration project rules in the TOM [Chapter 18, Section 2](#), will continue to apply to those TRICARE beneficiaries who began participation in Cancer Clinical Trials Demonstration before termination of the Demonstration. Such rules will continue to apply until the beneficiary is discharged from the clinical trial.

4.0 EXCLUSIONS

4.1 Care rendered in the National Institutes of Health Clinical Center.

4.2 Costs associated with non-treatment research activities related to clinical trials.

4.3 Phase I clinical trials (including Phase I arm of multi-phase clinical trials).

5.0 EFFECTIVE DATE

April 1, 2008.

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