

**State of New Mexico - Taxation and Revenue Department**  
**CANCER CLINICAL TRIAL TAX CREDIT CLAIM FORM**

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**Who May Claim This Credit:** For tax years beginning on or after January 1, 2012, but before January 1, 2016, a taxpayer who files an individual New Mexico income tax return, who is not a dependent of another taxpayer, who is an oncologist that is a physician licensed pursuant to the Medical Practice Act (Section 61-6-1 NMSA 1978) and whose practice is located in rural New Mexico may claim a tax credit of \$1,000 for each patient participating in a cancer clinical trial under the physician's supervision during the tax year, but not to exceed \$4,000 for all cancer clinical trials conducted by that physician. The credit may only be claimed for the tax year in which the physician participates as an investigator in a clinical trial. The credit may not be carried forward to another year, or refunded. This credit can only be claimed against personal income tax owed by the licensed physician. A husband and wife who file separate returns for a tax year in which they could have filed a joint return may each only claim one-half of the tax credit that would have been allowed on a joint return.

Only a qualified licensed physician may claim the credit. If the physician belongs to a business association in which one or more members qualifies for a cancer clinical trial tax credit, the credit is to be equally apportioned between the eligible physicians conducting, supervising or participating in the cancer clinical trial for which the credit is allowed. If not apportioned equally, provide an explanation in the space provided in Part III, Section 2. The total cancer clinical trial tax credit allowed for all the members of a partnership or business association shall not exceed the amount of credit that could have been claimed by one qualified physician.

When claiming the cancer clinical trial tax credit, this form must accompany the personal income tax or fiduciary income tax return to which the taxpayer wishes to apply the credit and mailed to the address on the tax return. For assistance call 505-827-1746.

**Part I - Qualified physician or practice**

Name of the qualified physician or the name of the practice	SSN	FEIN	New Mexico CRS ID Number
Physical address of clinic where the clinical trial took place	City, state and ZIP code		Medical License Number (MLN)
Mailing address, if different than the physical address	City, state and ZIP code		Expiration Date of MLN
Name of contact	Phone number	E-mail address	

**Part II - Total credit amount allowed**

1. Last day of the tax year for this claim (Format for the date is mm/dd/yyyy)	1.
2. Enter the number of patients who participated in a qualified cancer clinical trial under the claimant's supervision during the tax year. ....	2.
3. Multiply line 2 times \$1,000, but do not enter more than \$4,000. This is the amount of tax credit that maybe claimed. ....	3. \$

**Part III - Owners, members or partners, if the cancer clinical trial is performed within a partnership or business association**

**Section 1.** If the cancer clinical trial is performed by a partnership or business association in which one or more members qualify because they are eligible physicians conducting, supervising or participating in the cancer clinical trial for which the credit is allowed, complete the following for each member, partner or owner who is eligible to claim the credit. If additional space is needed, continue the list on a separate page.

	<u>Name</u>	<u>SSN</u>	<u>MLN</u>	<u>Expires</u>	<u>Owner's Share of the Credit</u>
a.					\$
b.					\$
c.					\$
d.					\$

**Section 2.** If the credit is not evenly distributed to each member, partner or owner, include an explanation in the space below. If additional space is needed, continue the explanation on a separate page.

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**Part IV - Qualifying the cancer clinical trial**

The cancer clinical trial must meet all requirements in Section I, below, to qualify. Check all boxes that apply. In Section II, you must provide the name of the organization and contact information for the entity approving the cancer clinical trial. Enter that information in Section II below.

**Section I**

Check all that apply:

- ☐ The cancer clinical trial was conducted for the purposes of:
- a) the prevention of or the prevention of reoccurrence of cancer, or
  - b) the early detection or treatment of cancer for which no equally or more effective standard cancer treatment exists.
- ☐ The clinical trial is not designed exclusively to test toxicity or disease pathophysiology and has a therapeutic intent.
- ☐ The clinical trial is provided in this state as part of a scientific study of a new therapy or intervention and is for the prevention of, prevention of reoccurrence, early detection, treatment or palliation of cancer in humans and in which the scientific study includes all of the following:
- 1) specific goals;
  - 2) a rationale and background for the study;
  - 3) criteria for patient selection;
  - 4) specific direction for administering the therapy or intervention and for monitoring patients;
  - 5) a definition of quantitative measures for determining treatment response;
  - 6) methods for documenting and treating adverse reactions; and
  - 7) a reasonable expectation that the treatment will be at least as efficacious as standard cancer treatment.
- ☐ The clinical trial is being conducted with approval of at least one of the following:
- 1) one of the federal national institutes of health;
  - 2) a federal national institutes of health cooperative group or center;
  - 3) the United States Department of Defense;
  - 4) the Federal Food and Drug Administration in the form of an investigational new drug application;
  - 5) the United States Department of Veterans Affairs; or
  - 6) a qualified research entity that meets the criteria established by the federal national institutes of health for grant eligibility;
- ☐ The clinical trial is considered part of a cancer clinical trial;
- ☐ The clinical trial has been reviewed and approved by an institutional review board that has an active federal-wide assurance of protection for human subjects; and
- ☐ The clinical trial in which the personnel conducting the clinical trial are working within their scope of practice, experience and training and are capable of providing the clinical trial because of their experience, training and volume of patients treated to maintain their expertise.

**Section II**

Enter the name and contact information for the organization approving the cancer clinical trial. Include the contact's name, phone number and e-mail address.

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NOTE: Failure to attach this fully completed form to your New Mexico return will result in denial of the credit.

Under penalty of perjury I declare that I have examined this claim, and to the best of my knowledge and belief, it is true, correct and complete.

Signature of claimant \_\_\_\_\_ Date \_\_\_\_\_

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Instructions

**The purpose** of the cancer clinical trial tax credit is to encourage physicians to participate as clinical trial investigators by performing cancer clinical trials of new cancer treatments in New Mexico and making cancer clinical trials more readily available to cancer patients in New Mexico.

**To complete the form**

**Part I.** Enter the information for the qualified physician, or if a partnership or business association, enter the information for the partnership or business association, in which the cancer clinical trials were conducted. You must provide the physical address of the clinic, to show where the cancer clinical trials were conducted. Enter the qualified physician's medical license number and expiration date if the applicant is a physician. If a partnership or business association, leave these boxes blank and enter the medical license number and expiration date of each owner, member or partner in Part III, Section I.

**Part II.** Complete this section to compute the total credit amount allowed during the tax year. On line 1, enter the last day of the tax year in which the cancer clinical trial was performed for this claim. The format to be used for the date is mm/dd/yyyy. On line 2, enter the number of patients who participated in a qualified cancer clinical trial under the claimant's supervision during the tax year of the claim. On line 3, multiply line 2 times \$1,000, but not more than \$4,000 and enter the amount of tax credit that may be claimed.

**Part III, Section 1.** This section is used to identify the owners, members or partners eligible to claim the credit if the cancer clinical trial is performed within a partnership or business association in which one or more members qualifies. For each owner, member or partner, enter their name, social security number, medical license number, the date their medical license expires and the owner, member or partners share of the total credit allowed on line 3, Part II of this claim form.

**Part III, Section 2.** If the credit is not evenly distributed to each member, owner or partner listed in Section 1, enter the reason in Section 2.

**Part IV.** Complete Sections 1 and 2 affirming that the cancer clinical trial qualifies for the cancer clinical trial tax credit. You must be able to answer yes to all of the questions listed, and to provide a name and contact information for the organization that approved the cancer clinical trial.

**Sign and date** the claim form affirming that the information provided is correct.

**Important Definitions**

*"Cancer clinical trial"* means a clinical trial:

- conducted for the purposes of the prevention of or the prevention of reoccurrence of cancer or the early detection or treatment of cancer for which no equally or more effective standard cancer treatment exists;
- that is not designed exclusively to test toxicity or disease

- pathophysiology and has a therapeutic intent;
- that is provided in this state as part of a scientific study of a new therapy or intervention and is for the prevention of, prevention of reoccurrence, early detection, treatment or palliation of cancer in humans and in which the scientific study includes all of the following:
  - 1) specific goals;
  - 2) a rationale and background for the study;
  - 3) criteria for patient selection;
  - 4) specific direction for administering the therapy or intervention and for monitoring patients;
  - 5) a definition of quantitative measures for determining treatment response;
  - 6) methods for documenting and treating adverse reactions; and
  - 7) a reasonable expectation that the treatment will be at least as efficacious as standard cancer treatment;
- that is being conducted with approval of at least one of the following:
  - 1) one of the federal national institutes of health;
  - 2) a federal national institutes of health cooperative group or center;
  - 3) the United States Department of Defense;
  - 4) the Federal Food and Drug Administration in the form of an investigational new drug application;
  - 5) the United States Department of Veterans Affairs; or
  - 6) a qualified research entity that meets the criteria established by the federal national institutes of health for grant eligibility;
- that is considered part of a cancer clinical trial;
- that has been reviewed and approved by an institutional review board that has an active federal-wide assurance of protection for human subjects; and
- in which the personnel conducting the clinical trial are working within their scope of practice, experience and training and are capable of providing the clinical trial because of their experience, training and volume of patients treated to maintain their expertise.

*"Rural New Mexico"* means a class B county in which no municipality has a population of 60,000 or more according to the most recent federal decennial census and includes the municipalities within that county. This includes areas within New Mexico that are outside of Bernalillo, DeBaca, Dona Ana, Los Alamos, Sandoval, San Juan, and Santa Fe Counties.