HEALTH PLAN POLICY

Policy Title: Prior Authorization for National Cancer Institute Clinical Trials  Policy Number: MUM03
Revision: C

Department: Medical Management  Sub-Department: Utilization Management

Applies to Product Lines:
- Medicaid
- Children’s Health Insurance Plan
- USFHP
- Health Insurance Exchange
- Medicare
- Commercial Insured
- Non Insured Business

Origination/Effective Date: 08/26/2014
Reviewed Date(s): 03/04/2016, 06/01/2017, 09/20/2018

SCOPE:

To assist health plan members in participating in National Cancer Institute (NCI) sponsored Phase I, Phase II or Phase III protocols and to ensure participation is reviewed and authorized in accordance with TRICARE policy.

DEFINITIONS AND ACRONYMS:

- **Clinical trials** - Clinical trials are research studies that involve people and test new ways to prevent, detect, diagnose, or treat cancer and other diseases.

- **National Cancer Institute (NCI)**

- **Phase I** - These trials are conducted to evaluate safety of chemical or biologic agents or other types of interventions (e.g., radiation therapy technique).

- **Phase II** – These trials test the effectiveness of interventions in people who have specific type of cancer or related cancers.

- **Phase III** – These trials compare effectiveness of new intervention, or new use of existing intervention, with the current standard of care for a particular type of cancer.

POLICY:

Members who are eligible for clinical trials will be identified to the health plan by the participating physician. The participating physician will provide the following to the health plan:

- The member’s name and last four digits of sponsor’s social security number.
- Request for authorization for services that specifies the clinical trial on the facility’s letter head.
- Certification that the patient meets all entry criteria for the said protocol.
- Copy of NCI clinical trial protocol information.
- Proof of NCI Sponsorship.

Once received, the request for authorization will be subject to the authorization review process to include verification of clinical trial by utilizing NCI’s comprehensive cancer data base, physician data query.
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The health plan will assign a utilization management registered nurse to work closely with the member, physician, and facility throughout the member’s participation in the trial. The member will also be assigned to the case management team.

If the request meets TRICARE policy guidelines for approval, the authorization will be entered in the system and the requesting physician will be notified.

If the request does not meet TRICARE policy guidelines it will be sent to the chief medical officer or designee for the review and determination.

REFERENCES:

- National Cancer Institute: www.cancer.gov/cancertopics/factsheet/clinicaltrials
- TRICARE policy manual 6010.57-M, February 1, 2008, Chapter 7, Section 24.1

RELATED DOCUMENTS:

None

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REVISION HISTORY:

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<td>New</td>
<td>08/26/2014</td>
<td>Initial Release</td>
<td>Board of Directors</td>
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<tr>
<td>A</td>
<td>03/04/2016</td>
<td>Yearly review – updated to current template. Updated Definitions and Acronyms.</td>
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<tr>
<td>B</td>
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