**Medical Coverage Policy** | Tumor-Treating Fields Therapy for Glioblastoma



**EFFECTIVE DATE:** 01 | 20 | 2014 **POLICY LAST UPDATED:** 08 | 16 | 2016

#### **OVERVIEW**

Glioblastoma multiforme (GBM) is the most common and deadly malignant brain tumor. It has a very poor prognosis and is associated with low quality of life during the course of treatment. Tumor treating fields (TTF) therapy is a new, noninvasive technology that is intended to treat glioblastoma using electric fields.

#### PRIOR AUTHORIZATION

Not applicable

## POLICY STATEMENT

#### **BlueCHiP for Medicare and Commercial Products**

Tumor Treating Fields Therapy for Glioblastoma as an alternative to standard chemotherapy or as an adjunct to standard maintenance therapy following initial treatment with surgery and/or radiotherapy is considered not medically necessary as there is insufficient peer-reviewed scientific literature that demonstrates that the procedure/service is effective.

#### MEDICAL CRITERIA

None

#### BACKGROUND

Glioblastomas, also known as glioblastoma multiforme, are the most common form of malignant primary brain tumor in adults, and they comprise approximately 15% of all brain and central nervous system tumors and more than 50% of all tumors that arise from glial cells. The peak incidence for GBM occurs between the ages of 45 and 70 years. GBMs are grade IV astrocytomas, the most deadly type of glial cell tumor, and are often resistant to standard chemotherapy. According to the National Comprehensive Cancer Network, GBM is the "deadliest brain tumor with only a third of patients surviving for 1 year and less than 5% living beyond 5 years."

The primary treatment for GBM is debulking surgery to remove as much of the tumor as possible. At that time, some patients may undergo implantation of the tumor cavity with a carmustine (BCNU)-impregnated wafer. Depending on the patient's physical condition, adjuvant radiation therapy, chemotherapy (typically temozolomide), or a combination of the two are sometimes given. After adjuvant therapy, some patients may undergo maintenance therapy with temozolomide. In patients with disease that recurs after these initial therapies, additional debulking surgery may be used if recurrence is localized. Treatment options for recurrent disease include various forms of systemic medications such as bevacizumab, bevacizumab plus chemotherapy (e.g., irinotecan, BCNU/CCNU, temozolomide), temozolomide, nitrosourea, PCV (procarbazine, CCNU, and vincristine), cyclophosphamide, and platinum-based agents. Response rates in recurrent disease are less than 10%, and progression-free survival rates at 6 months are less than 20%.

Tumor treating fields therapy is a new, noninvasive technology that is intended to treat GBM on an outpatient basis using electrical fields. (3-5) TTF therapy exposes cancer cells to alternating electric fields of low-intensity and intermediate frequency, which are purported to both selectively inhibit tumor growth and reduce tumor angiogenesis. Tumor treating fields are proposed to inhibit rapidly dividing tumor cells by two mechanisms, arrest of cell proliferation and destruction of cells while undergoing division.

The NovoTTF-100A<sup>™</sup> System (Novocure Ltd., Haifa, Israel) has been approved by the U.S. Food and Drug Administration (FDA) to deliver TTF therapy. TTF therapy via the NovoTTF-100A<sup>™</sup> System is delivered by a battery-powered, portable device that generates the fields via disposable electrodes that are noninvasively attached to the patient's shaved scalp over the site of the tumor. The device is used by the patient at home on a continuous basis (20–24 hours per day) for the duration of treatment, which can last for several months. Patients can carry the device in a backpack or shoulder pack while carrying out activities of daily living.

Relevant outcomes are overall survival, disease-specific survival, quality of life, and treatment-related morbidity. The single randomized controlled trial (RCT) reported that patients who received TTF therapy plus temozolomide have longer progression-free survival (3.1 months) and overall survival (4.9 months) than patients who received temozolomide alone. The trial had methodologic limitations, including the lack of placebo control, differential dropout between groups, and the possibility of adherence bias for outcomes reported with per protocol analysis. Further corroboration of these results are needed in high-quality RCTs. The evidence is insufficient to determine the effects of the technology on health outcomes and is considered not medically necessary.

### COVERAGE

Benefits vary between groups/contracts. Please refer to the appropriate Evidence of Coverage or Subscriber Agreement for applicable services not medically necessary coverage.

#### CODING

#### BlueCHiP for Medicare and Commercial Products

The following codes are not medically necessary:

- A4555 Electrode/transducer for use with electrical stimulation device used for cancer treatment, replacement only
- E0766 Electrical stimulation device used for cancer treatment, includes all accessories, any type

# RELATED POLICIES

None

#### **PUBLI SHED**

Provider Update, October 2016 Provider Update, April 2015 Provider Update, March 2014

#### REFERENCES

1. Novo TTF-100A System

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