

Electric Tumor Treatment Field Therapy (for New Jersey Only)

Policy Number: CS146NJ.H
Effective Date: March 1, 2024

[➔ Instructions for Use](#)

Table of Contents	Page
Application	1
Coverage Rationale	1
Definitions	2
Applicable Codes	2
Description of Services	3
Clinical Evidence	4
U.S. Food and Drug Administration	10
References	10
Policy History/Revision Information	12
Instructions for Use	12

Related Policy
<ul style="list-style-type: none"> Clinical Trials (for New Jersey Only)

Application

This Medical Policy only applies to the state of New Jersey.

Coverage Rationale

The following is proven and medically necessary for treating newly diagnosed histologically-confirmed Supratentorial glioblastoma (GBM):

- The use of U.S. Food and Drug Administration (FDA) approved devices to generate electric tumor treatment fields (TTF) when used according to FDA labeled indications, contraindications, warnings and precautions, and when all of the following criteria are met:
 - Treatment with radiation therapy has been completed; **and**
 - Individual is receiving Temozolomide as the only cancer drug; **and**
 - Individual has a Karnofsky Performance Status (KPS) score of ≥ 60 or Eastern Cooperative Oncology Group (ECOG) Performance Status ≤ 2 ; **and**
 - Individual has been counselled that the device must be worn at least 18 hours daily.

When all of the above criteria are met for newly diagnosed GBM , an initial 3 months of electric TTF therapy will be approved.

The following is proven and medically necessary for treating radiologically confirmed recurrence of GBM in the Supratentorial region of the brain:

- The use of FDA approved devices to generate electric TTF after initial chemotherapy when used according to FDA labeled indications, contraindications, warnings and precautions and when all of the following criteria are met:
 - The device is used as the only treatment; **and**
 - Individual has a KPS score of ≥ 60 or ECOG Performance Status ≤ 2 ; **and**
 - Individual has been counselled that the device must be worn at least 18 hours daily.

When all of the above criteria are met for recurrent GBM, an initial 3 months of electric TTF therapy will be approved.

Subsequent approval(s) for continuation beyond the initial 3 months of electric TTF for treatment of histologically-confirmed Supratentorial glioblastoma (GBM) is based on:

- Magnetic resonance imaging (MRI) scan has been performed \leq 2 months prior to request and documents no evidence of disease progression; **and**
- Individual with newly diagnosed glioblastoma continues to receive Temozolomide as the only cancer drug or the device is used as the only treatment for an individual with recurrent GBM; **and**
- KPS score of \geq 60 or ECOG Performance Status \leq 2; **and**
- Documentation that the individual has been using the device at least 18 hours daily.

Due to insufficient evidence of efficacy, the use of devices to generate electric TTF is unproven and not medically necessary when the criteria above are not met and for all other indications including but not limited to the following:

- Treatment of tumors other than GBM.
- Use of electric TTF therapy with concurrent medical therapy (e.g., bevacizumab or chemotherapy) for treatment of recurrent GBM.

Computer software used for therapeutic radiology clinical treatment planning in conjunction with electric TTF therapy is unproven and not medically necessary due to insufficient evidence of efficacy.

Definitions

Eastern Cooperative Oncology Group (ECOG) Performance Status: A measurement of performance status that describes an individual's level of functioning. Individuals who have a worse performance status and limited functional capacity tend to have more difficulty tolerating rigorous cancer treatments. This scale may also be referred to as the WHO or Zubrod score.

- A score of zero indicates the individual is fully active and has no restrictions
- A score of one indicates that the individual is restricted in physically strenuous activity but is completely ambulatory and cares for self
- A score of two indicates the individual is ambulatory more than 50% of the time and is capable of all self-care but unable to carry out any work activities
- A score of three indicates that the individual is ambulatory 50 percent or less of the time and is capable of only limited self-care
- A score of four indicates bedbound (cannot perform any self-care and is totally confined to bed or chair)

(West and Jin, 2015; ECOG-ACRIN Cancer Research Group)

Karnofsky Performance Status (KPS): A standard way of measuring the ability of cancer patients to perform ordinary tasks. KPS scores range from 0 to 100; a higher score means a person is better able to carry out daily activities. For example, a KPS of 60 means a person requires occasional assistance but is able to care for most of their personal needs. KPS may be used to determine a patient's prognosis, to measure changes in a patient's ability to function, or to decide if a patient could be included in a clinical trial [National Cancer Institute (NCI), 2019; West and Jin, 2015].

Supratentorial: A term used to describe the upper portion of the brain comprised of the cerebrum, ventricles, choroid plexus, hypothalamus, pineal gland, pituitary gland, and optic nerve (NCI, 2019).

Temozolomide: An oral alkylating chemotherapy drug used in the treatment of some brain cancers. It is a first-line treatment for glioblastoma (NCI, 2019).

Applicable Codes

The following list(s) of procedure and/or diagnosis codes is provided for reference purposes only and may not be all inclusive. Listing of a code in this policy does not imply that the service described by the code is a covered or non-covered health service. Benefit coverage for health services is determined by federal, state or contractual requirements and applicable laws that may require coverage for a specific service. The inclusion of a code does not imply any right to reimbursement or guarantee claim payment. Other Policies and Coverage Determination Guidelines may apply.

CPT Code	Description
77299	Unlisted procedure, therapeutic radiology clinical treatment planning

CPT® is a registered trademark of the American Medical Association

HCPCS Code	Description
*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

Codes labeled with an asterisk (*) are not on the State of New Jersey Medicaid Fee Schedule and therefore may not be covered by the State of New Jersey Medicaid Program.

Description of Services

Electric tumor treatment field (TTF) therapy (also known as tumor-treating fields, TTFs, ETTFs) is based on the principle that low intensity, intermediate frequency electric fields (100 to 300 kHz) disrupt cell division and may destroy proliferating cells in brain tumors (Rulseh et al, 2012).

Glioblastoma multiforme (GBM) is the most prevalent and primary malignant brain tumor in adults. For patients with newly diagnosed glioblastoma the initial standard treatment consists of debulking surgery (when feasible), followed by radiation and chemotherapy (NCCN, 2023).

The Optune® Treatment Kit, formerly the NovoTTF-100A System, (Novocure) was approved by the FDA in April 2011, as a novel device to treat adults age 22 years or older with GBM that recurs or progresses after receiving chemotherapy and radiation therapy. The Optune Treatment Kit has also been approved by the FDA in combination with Temozolomide in adult patients with newly diagnosed, Supratentorial glioblastoma following maximal debulking surgery and completion of radiation therapy. For newly diagnosed GBM, Optune is not intended to be used as a substitute for standard treatments but rather as an adjunct therapy (Novocure, 2020).

Refer to the [U.S. Food and Drug Administration \(FDA\)](#) section for additional information.

The Optune kit contains the portable electric field generator (Optune device), INE (insulated electrode) transducer arrays, power supply, and additional supplies. Prior to treatment, transducer arrays are placed on the individual's scalp according to the tumor's location, which are then covered by a lightweight white cap which resembles a bandage. The individual receives the noninvasive TTF treatment for at least 18 continuous hours per day for a minimum of 4 weeks. As the Optune device is portable, individuals are able to carry out every-day activities.

Treatment parameters are preset by the manufacturer such that there are no electrical output adjustments available to the individual being treated. The individual being treated, or caregiver must learn to change and recharge depleted device batteries and to connect to an external power supply overnight. In addition, the transducer arrays need to be replaced once to twice a week and the scalp re-shaved in order to maintain optimal contact.

The NovoTAL™ (transducer array layout) system is optional simulation software for use in clinical treatment planning with Optune therapy that may be leased from the manufacturer. Its purpose is to determine the optimal location of the transducer arrays based on the individual's most recent magnetic resonance imaging (MRI) scan, head size, and tumor location.

TTF technology is also being studied through ongoing clinical trials as a treatment for other solid tumors such as non-small cell lung cancer, brain metastasis, pancreatic cancer, ovarian cancer and mesothelioma.

Glioblastoma

The National Comprehensive Cancer Network (NCCN) Clinical Practice Guideline for Central Nervous System Cancers, anaplastic gliomas/glioblastoma, includes standard brain radiation therapy (RT) + concurrent temozolomide and adjuvant temozolomide + alternating electric field therapy for patients with good performance status (Karnofsky Performance Status (KPS) \geq 60) and either methylated or unmethylated/indeterminate MGMT promoter status, in whom maximal, safe resection was not feasible with the following footnote: "Alternating electric field therapy is only an option for patients with supratentorial disease" (category 1). For recurrence of GBM (GLIO-5), the guideline includes consideration of alternating electric treatment fields for glioblastoma after surgery, radiation and chemotherapy (category 2B). The guideline recommends that follow-up magnetic resonance imaging (MRI) of the brain be done 2 to 8 weeks after RT, then every 2–4 months for 3 years, then every 3–6 months indefinitely (NCCN,2023).

Regev et al. (2021) conducted a systematic review and meta-analysis of evaluating Tumor-Treating Fields (TTFields) mechanism of action, safety, and efficacy, for both newly diagnosed glioblastoma (ndGBM) and for recurrent GBM (rGBM). Twenty studies met the pre-defined inclusion criteria, including 1636 patients (542 ndGBM and 1094 rGBM), and 11 558 patients (6403 ndGBM and 5155 rGBM) analyzed for the clinical outcomes and safety endpoints, separately. This study demonstrated improved clinical efficacy and a good safety profile of TTFields. For ndGBM, pooled median overall survival (OS) and progression-free survival (PFS) were 21.7 (95%CI = 19.6-23.8) and 7.2 (95%CI = 6.1-8.2) months, respectively. For rGBM, pooled median OS and PFS were 10.3 (95%CI = 8.3-12.8) and 5.7 (95%CI = 2.8-10) months, respectively. Compliance of \geq 75% was associated with an improved OS and the predominant adverse events were dermatologic, with a pooled prevalence of 38.4% (95%CI = 32.3-44.9). Preclinical studies demonstrated TTFields' diverse molecular mechanism of action, its potential synergistic efficacy, and suggest possible benefits for certain populations. The findings showed that TTFields patients had significantly better emotional function and reported significantly lower incidence of side effects. Also, TTFields users had better emotional, physical, and cognitive functions than social and role functioning. The study showed TTFields frequently affected multiple aspects of the patients' daily lives; however, 70% would recommend TTFields to others, and 67% would reuse the device. This study supports the clinical benefit, safety and potential therapeutic synergism of TTFields for the treatment of GBM, alongside the standard-of-care treatment protocol (Mrugala 2014, Stupp 2017, Wong 2015 are included in this review).

A systematic review and a Bayesian network meta-analysis were performed by Chen et al. (2021) to compare and rank active therapies in recurrent glioblastoma (GBM). The authors obtained a treatment hierarchy using the surface under the cumulative ranking curve and mean ranks. A cluster analysis was conducted to aggregate the separated results of three outcomes. A total of 1,667 citations were identified, and 15 eligible articles with 17 treatments remained in the final network meta-analysis. Pairwise comparison showed no difference on the 6-month progression-free survival (6-m PFS) rate, objective response rate (ORR), and overall survival (OS). Among the reports, cediranib plus lomustine (CCNU) corresponded to the highest rates of grade 3-4 adverse events. Ranking and cluster analysis indicated that bevacizumab (BEV) plus CCNU and regorafenib had a higher efficacy on the ORR, 6-m PFS rate and OS, and that BEV monotherapy or BEV combined with active drug therapies was advantageous for the ORR and 6-m PFS rate. Additionally, tumor treatment fields (TTF) plus BEV showed a higher SUCRA value in OS. The authors concluded that according to ranking and cluster analysis, BEV plus CCNU and regorafenib are the primary recommendations for treatment. BEV monotherapy alone or combined with active drug therapies are recommended in patients with severe neurological symptoms. Limitations include inconsistency among researchers which may bring bias to the statistical results. In addition, the published results lag behind current therapy options. New therapies including neoadjuvant checkpoint inhibitor and laser interstitial thermotherapy have not been included. The findings of this study need to be validated by well-designed studies. Further investigation is needed before clinical usefulness of this procedure is proven.

Dono et al. (2021) performed a retrospective review of an institutional database with 530 patients with infiltrating gliomas to evaluate the survival effects of tumor treating fields (TTFields) in a cohort of patients with isocitrate dehydrogenase wild-type (IDH-WT) recurrent GBM (rGBM), and to investigate the possible clinical characteristics or genomic alterations that may predict responsiveness to TTFields. Patients with IDH-WT rGBM receiving TTFields at first recurrence were included. Tumors were evaluated by next-generation sequencing for mutations in 205 cancer-related genes. Post-progression survival (PPS) was examined using the log-rank test and multivariate Cox-regression analysis. A total of 149 rGBM patients were identified of which 29 (19%) were treated with TTFields. No difference in median PPS was observed between rGBM patients who received versus did not receive TTFields (13.9 versus 10.9 months, $p = 0.068$). However, within the TTFields-treated group ($n = 29$), PPS was improved in PTEN-mutant ($n = 14$) versus PTEN-WT ($n = 15$) rGBM, (22.2 versus 11.6 months, $p = 0.017$). Within the PTEN-

mutant group (n = 70, 47%), patients treated with TTFields (n = 14) had longer median PPS (22.2 versus 9.3 months, p = 0.005). No PPS benefit was observed in PTEN-WT patients receiving TTFields (n = 79, 53%). TTFields therapy conferred a PPS benefit in PTEN-mutant rGBM. The authors concluded that understanding the molecular mechanisms underpinning the differences in response to TTFields therapy could help elucidate the mechanism of action of TTFields and identify the rGBM patients most likely to benefit from this therapeutic option. This study is limited by its retrospective observations. In addition, a small sample size makes it difficult to decide whether these conclusions can be generalized to a larger population. Further research with randomized controlled trials is needed to validate these findings.

Ram et al. (2021) performed a subgroup analysis of the multicenter, phase 3, EF-14 randomized clinical trial (Identifier: NCT00916409) subgroup analysis of patients with newly diagnosed glioblastoma to evaluate the safety and efficacy of tumor treating fields (TTFields) in elderly patients. All 134 patients who are ≥ 65 years of age were included (TTFields/TMZ combination, n = 89; TMZ monotherapy, n = 45; 2:1 ratio of randomization). Progression-free survival (PFS) and overall survival (OS) were analyzed using Kaplan-Meier methodology ($\alpha = 0.05$). Health-related quality-of-life (HRQoL) was assessed using the European Organization for Research and Treatment of Cancer (EORTC) quality-of-life questionnaire QLQ-C30 supplemented with the brain tumor module (QLQ-BN20). Adverse events (AEs) were evaluated using Common Terminology Criteria for AEs (CTCAE) v4.0. The PFS was 6.5 months in patients randomized to the treatment group with TTFields/TMZ combination *versus* 3.9 months in patients treated with TMZ monotherapy (HR, 0.47; 95% CI, 0.30-0.74; p = 0.0236). The OS was 17.4 months in patients treated with TTFields/TMZ combination *versus* 13.7 months in patients treated with TMZ monotherapy (HR, 0.51; 95% CI, 0.33-0.77; p = 0.0204). Annual survival rates with TTFields/TMZ *versus* TMZ monotherapy were 39% (95% CI, 29-50%) *versus* 27% (95% CI, 15-41%; p = 0.072) at 2 years, 19% (95% CI, 11-29%) *versus* 11% (95% CI, 4-23%; p = 0.135) at 3 years, and 15% (95% CI, 7-25%) *versus* 0% at 5 years, respectively. There were no differences between groups in the pre-selected items of HRQoL assessment. Grade ≥ 3 systemic AEs were 46% in the TTFields/TMZ group *versus* 40% in the TMZ monotherapy group, without statistically significant difference between the two groups. The only TTFields-related AEs were reversible scalp skin reactions, with grades 1-2 and grade 3 skin reactions reported by 51% and 2% of patients, respectively. The authors concluded that combining TTFields with maintenance TMZ improved PFS and OS in elderly patients with ndGBM in the phase 3 EF-14 clinical trial, without increases in systemic toxicity or negatively affecting patient HRQoL. TTFields-related skin AEs were low-grade and manageable. Limitations of this analysis include the small sample sizes of both treatment groups due to the limited number of patients enrolled in the EF 14 trial who were 65 years of age or older. There is also a lack of available molecular data in $\sim 20\%$ of patients; 4% of TTFields plus TMZ-treated patients had an IDH mutation compared with 0% of the TMZ-treated patients; and only 1 patient had a 1p19q codeletion (TTFields plus TMZ group). Long-term evaluations of the results and prospective randomized studies are still needed.

Jin et al. (2020) conducted a systematic review and meta-analysis to compare the efficacy and safety of treatments based on the Stupp protocol for adult patients with newly diagnosed glioblastoma and to determine the optimal treatment option for patients with different O-6-methylguanine-DNA methyltransferase (MGMT) promoter methylation statuses. The author estimated hazard ratios (HRs) for overall survival (OS) and odds ratios (ORs) for adverse events of grade 3 or higher (AEs ≥ 3). Twenty-one randomized controlled trials including 6478 patients treated with 21 different treatment strategies were included. Results of the pooled HRs indicated tumor-treating fields (TTF) combined with the Stupp protocol resulted in the most favorable overall survival for patients with and without MGMT promoter methylation. Subgroup analyses by the two MGMT promoter statuses indicated that lomustine-temozolomide plus radiotherapy or TTF combination therapy was associated with the best overall survival for patients with methylated MGMT promoter [HR, 1.03; 95% credible interval (CI), 0.54-1.97], and standard cilengitide combination therapy or TTF combination treatment was linked with the best overall survival for patients with unmethylated MGMT promoter (HR, 1.05; 95% CI, 0.67-1.64). Regarding AEs ≥ 3 , there were no significant differences in pooled ORs. However, Bayesian ranking profiles that verified intensive cilengitide combination therapy and TTF combination therapy have a similar possibility to cause the least toxicity. Results indicated that TTF combination therapy was associated with increased survival, regardless of the MGMT promoter methylation status, and a relatively tolerated safety profile compared with other combination treatments. The optimal treatment option for glioblastoma patients with different MGMT promoter methylation statuses was different. These findings are helpful in establishing a standard of care and plan for adults patients with newly diagnosed glioblastoma (Stupp 2017 and Stupp 2014 are included in this review).

Marenco-Hillebrand et al. (2020) conducted a systematic review to describe the current status and advances in the survival of patients with glioblastoma by analyzing median overall survival through time and between treatment modalities. Full-text glioblastoma papers with human subjects ≥ 18 years old and n ≥ 25 were included for evaluation. The central tendency of median overall survival (MOS) was 13.5 months (2.3-29.6) and cumulative 5-year survival was 5.8% (0.01%-29.1%), with a significant difference in survival between studies that predate versus postdate the implementation of temozolomide and

radiation, [12.5 (2.3-28) vs 15.6 (3.8-29.6) months, $p < 0.001$]. Within clinical trials, the highest MOS involved tumor treating fields (TTF) with 20.7 (range 20.5–20.9) months. According to the authors, therapies such as TTF provide a means of prolonging the survival of glioblastoma patients.

Kim et al. (2020) reported on Korean newly diagnosed GBM patients who participated in the EF-14 trial. Thirty-nine participants of the EF-14 trial were enrolled at 8 sites in South Korea. Patients (24 TTF/TMZ; 14 TMZ alone) received: TTF (200 kHz) for > 18 h/day; TMZ at 120-150 mg for 5 days per a 28 day cycle. Safety and efficacy were assessed. Patient baseline characteristics were balanced in the 2 arms and the mean age was 52.1 years, 66.7% were male with a mean KPS of 90. Safety incidence was comparable between the 2 arms. In the TTF/TMZ arm, 30% suffered from skin irritation versus 52% in the entire study population. No TTF-related serious adverse events were reported. The median progression-free survival (PFS) in the TTF/TMZ arm was 6.2 months (95% CI 4.2-12.2) versus 4.2 (95% CI 1.9-11.2) with TMZ alone ($p = 0.67$). Median overall survival was 27.2 months (95% CI 21-NA) with TTF/TMZ versus 15.2 months (95% CI 7.5-24.1; HR 0.27, $p = 0.01$) with TMZ alone. The authors concluded that median overall survival and 1- and 2-year survival rates were higher with TTF/TMZ and similar to the entire EF-14 population. According to the authors, these results demonstrate the efficacy and safety of TTF in this population of newly diagnosed glioblastoma patients.

A retrospective analysis was performed by Shi et al. (2020) with unsolicited, post-marketing surveillance data from tumor treating fields (TTF) treated patients with high-grade glioma (October 2011-February 2019) using Medical Dictionary for Regulatory Activities (MedDRA) v21.1 preferred terms, stratified by region [US, EMEA (Europe, Middle East, Africa), Japan], diagnosis (ndGBM, rGBM, anaplastic astrocytoma/oligodendroglioma, other brain tumors), and age [< 18 (pediatric), 18-64 (adults), ≥ 65 (elderly); years of age]. The aim of this analysis was to assess the safety of TTF in real-world, clinical practice settings, using global post-marketing surveillance data from a large patient cohort. Of 11,029 patients, 53% were diagnosed with ndGBM and 39% were diagnosed with rGBM at any line of disease recurrence. Most were adults (73%), 26% were elderly, and the male-to-female ratio was $\sim 2:1$ (close to published ratios of typical GBM populations). The most commonly reported TTF-related adverse event (AE) was array-associated skin reaction, occurring in patients with ndGBM (38%), rGBM (29%), anaplastic astrocytoma/oligodendroglioma (38%), and other brain tumors (31%); as well as 37% of pediatric, 34% of adult, and 36% of elderly patients. Most skin AEs were mild/moderate and manageable. Other TTF-related AEs in patients with ndGBM/rGBM included under-array heat sensation (warmth; 11%, 10%, respectively) and electric sensation (tingling; 11%, 9%, respectively), and headache (7%, 6%, respectively). The authors concluded that this TTF safety surveillance analysis in $> 11,000$ patients revealed no new safety concerns, with a favorable safety profile comparable with published TTF/GBM trials. The safety profile remained consistent among subgroups, suggesting feasibility in multiple populations, including elderly patients. Limitations include the retrospective and observational design. Safety data were collated only from TTF-treated patients that reported AEs, therefore incidence of the overall cohort and sub-groups is likely overestimated. In addition, no efficacy, survival, or standardized QoL assessment data were included. This analysis did not translate research data into guidelines to improve patient care, and there is no evidence that this analysis will affect patient management. A Hayes Health Technology (HT) Assessment report published December 27, 2019, of tumor treating fields (Optune) for treatment of glioblastoma provides a Hayes rating of C for the use of tumor treating fields (TTF) as monotherapy in adult patients (22 years of age and older) with recurrent glioblastoma (GBM) following surgery and radiotherapy. A Hayes rating of C is also provided for TTF treatment with concomitant temozolomide (TMZ) in adult patients (22 years of age and older) with newly diagnosed GBM following surgery and radiation therapy with concomitant chemotherapy. A small, low-quality body of evidence suggests that TTF therapy results in overall survival (OS) and progression-free survival (PFS) at least equivalent to chemotherapy in patients with recurrent GBM. A small, low-quality body of evidence suggests that TTF plus TMZ increases OS and PFS in patients with newly diagnosed GBM compared with TMZ alone. HT annual review performed on December 13, 2021, reveals no changes in current ratings of C.

Toms et al. (2019) analyzed compliance data from Tumor treating fields (TTF)/temozolomide (TMZ) patients in a subgroup analysis of the phase 3 EF-14 trial (Stupp et al., 2017) to correlate TTF compliance with progression free survival (PFS) and overall survival (OS) and identify potential lower boundary for compliance with improved clinical outcomes. Compliance was assessed by usage data from the NovoTTF-100A device and calculated as percentage per month of TTF delivery. TTF/TMZ patients were segregated into subgroups by percent monthly compliance. A Cox proportional hazard model controlled for sex, extent of resection, MGMT methylation status, age, region, and performance status was used to investigate the effect of compliance on PFS and OS. A threshold value of 50% compliance with TTF/TMZ improved PFS and OS versus TMZ alone with improved outcome as compliance increased. At compliance $> 90\%$, median survival was 24.9 months (28.7 months from diagnosis) and 5-year survival rate was 29.3%. The authors concluded that a compliance threshold of 50%

with TTFields/TMZ correlated with significantly improved OS and PFS versus TMZ alone. Patients with compliance > 90% showed extended median and 5-year survival rates.

Magouliotis et al. (2018) performed a systematic review on the literature for patients with glioblastoma treated with tumor-treating fields (TTFields) plus radio chemotherapy or conventional radio chemotherapy alone, to compare the efficacy and safety of the two methods. Six studies met the inclusion criteria incorporating 1806 patients for the qualitative analysis and 1769 for the quantitative analysis. This study reveals increased median overall survival at 1 year and 2 years and median progression-free survival along with progression-free survival at 6 months for the patients treated with TTFields. Survival at 3 years was comparable between the two groups. TTFields were associated with fewer adverse events compared to chemotherapy along with similar incidence of skin irritation. The authors indicated that this review suggests that TTFields are a safe and efficient novel treatment modality. Stupp et al. (2017) reported final outcomes from the randomized, open-label trial of 695 patients with glioblastoma whose tumor was resected or biopsied and had completed concomitant radiochemotherapy (median time from diagnosis to randomization, 3.8 months) and Optune therapy. Of the 695 randomized patients [median age, 56 years; IQR, 48-63; 473 men (68%)], 637 (92%) completed the trial. Median progression-free survival from randomization was 6.7 months in the TTFields-temozolomide group and 4.0 months in the temozolomide-alone group (HR, 0.63; 95% CI, 0.52-0.76; $p < .001$). Median overall survival was 20.9 months in the TTFields-temozolomide group vs 16.0 months in the temozolomide-alone group (HR, 0.63; 95% CI, 0.53-0.76; $p < .001$). Systemic adverse event frequency was 48% in the TTFields-temozolomide group and 44% in the temozolomide-alone group. Mild to moderate skin toxicity underneath the transducer arrays occurred in 52% of patients who received TTFields-temozolomide vs no patients who received temozolomide alone. In the final analysis of this randomized clinical trial of patients with glioblastoma who had received standard radiochemotherapy, the addition of TTFields to maintenance temozolomide chemotherapy vs maintenance temozolomide alone, resulted in statistically significant improvement in progression-free survival and overall survival. These results are consistent with the previous interim analysis.

In a secondary analysis of the Stupp et al. (2017) trial, Taphoorn et al. (2018) examined the association of TTFields therapy with progression-free survival and HRQoL among patients with glioblastoma. Of the 695 patients in the study, 639 (91.9%) completed the baseline HRQoL questionnaire. Of these patients, 437 (68.4%) were men; mean (SD) age, 54.8 (11.5) years. Health-related quality of life did not differ significantly between treatment arms except for itchy skin. Deterioration-free survival was significantly longer with TTFields for global health (4.8 vs 3.3 months; $p < .01$); physical (5.1 vs 3.7 months; $p < .01$) and emotional functioning (5.3 vs 3.9 months; $p < .01$); pain (5.6 vs 3.6 months; $p < .01$); and leg weakness (5.6 vs 3.9 months; $p < .01$), likely related to improved progression-free survival. Time to deterioration, reflecting the influence of treatment, did not differ significantly except for itchy skin (TTFields worse; 8.2 vs 14.4 months; $p < .001$) and pain (TTFields improved; 13.4 vs 12.1 months; $p < .01$). Role, social, and physical functioning were not affected by TTFields. The addition of TTFields to standard treatment with temozolomide for patients with glioblastoma results in improved survival without a negative influence on HRQoL except for more itchy skin, an expected consequence from the transducer arrays.

In a multinational, open-label, randomized phase III trial (EF-14 trial), Stupp et al. (2015) compared Optune in combination with temozolomide to temozolomide alone in 700 patients age 18 and over with newly diagnosed GBM. The interim report revealed that in the intent-to-treat population, patients treated with TTFields plus temozolomide showed a statistically significant increase in progression free survival (PFS), the primary endpoint, compared to temozolomide alone (median PFS 7.1 months versus 4.0 months, hazard ratio = 0.62, $p = 0.0013$). In the per-protocol population, patients treated with TTFields plus temozolomide demonstrated a statistically significant increase in OS, a powered secondary endpoint, compared to temozolomide alone (median OS 20.5 months versus 15.6 months, hazard ratio = 0.64, $p = 0.0042$). In the intent-to-treat population, the median OS was 19.6 months versus 16.6 months, respectively, hazard ratio = 0.74 ($p = 0.0329$). The two-year survival rate was 50 percent greater with TTFields plus temozolomide versus temozolomide alone: 43 percent versus 29 percent. The trial's independent data monitoring committee concluded that the study met its endpoints at its pre-specified interim analysis of the first 315 patients with 18 months or more of follow-up. The committee recommended that the trial be terminated early for success and that all control patients be offered TTFields therapy even prior to progression. In addition, the authors reported that the trial showed Optune could be safely combined with temozolomide. There was no significant increase in systemic toxicities from Optune reported in combination with temozolomide versus temozolomide alone. The most common adverse reaction from Optune treatment was mild to moderate skin irritation, which according to the authors was easily managed, reversible and did not result in treatment discontinuation. This clinical trial has some important limitations. Patient enrollment occurred only after the end of radiochemotherapy, leading to some variation in the delivery of standard treatment of temozolomide and radiotherapy. Patients who had progressed early during radiochemotherapy were not eligible for randomization, thus excluding patients with very poor prognoses. There is likely reporting bias for second-line therapies after tumor progression because in

the TTFields plus temozolomide group, TTFields were to be continued, and thus, more detailed treatment information has been tracked for this group.

Wong et al, (2015) conducted a retrospective chart review from a single institution on patients treated with NovoTTF-100A and bevacizumab between November 2011 and December 2013. The patients were segregated into two cohorts: (i) those treated with NovoTTF-100A and bevacizumab only and (ii) those treated with NovoTTF-100A, bevacizumab and TCCC. Response to treatment was measured according to the Response Assessment in Neuro-Oncology criteria. Progression-free survival (PFS) and OS were measured from the time of application of these treatments to death or last follow up. The cohort treated with NovoTTF-100A, bevacizumab, and TCCC (n = 3) did not differ significantly from the rest of the cohort treated with NovoTTF-100A and bevacizumab only (n = 34). Potential reasons for this include baseline clinical characteristics and dexamethasone use. The authors note limitations with this review to be the number of patients treated with NovoTTF-100A, bevacizumab, and TCCC is small and therefore they cannot recommend this combination as standard clinical practice. However, they commented that the findings in their patients are notable and it can serve as a basis for future clinical trials. Second, it is unclear what the relative contribution of immunosuppression in the periphery versus the tumor microenvironment has on treatment resistance in recurrent glioblastomas. Therefore, they conclude that combination treatment, rather than single-agent monotherapy, will more likely affect meaningful clinical results.

A treatment-based analysis of data from the pivotal phase III trial of the NovoTTF-100A System™ versus best physician's choice (BPC) chemotherapy was conducted by Kanner et al. (2014) in patients with recurrent glioblastoma multiforme (GBM), with particular focus on efficacy in patients using NovoTTF therapy as intended. Median overall survival (OS) was compared for recurrent GBM patients receiving at least one full cycle of treatment with NovoTTF-100A System (monotherapy) or BPC chemotherapy [modified intention-to-treat (mITT) population]. The relationship between NovoTTF-100A System compliance and OS was evaluated in the ITT population. Kaplan-Meier analyses examined treatment-related differences in OS for various patient subgroups. Median OS was significantly higher in patients receiving ≥ 1 course of NovoTTF therapy versus BPC [7.7 v 5.9 months; hazard ratio, 0.69; 95% confidence interval (CI), 0.52-0.91; p = .0093]. Median OS was also significantly higher in patients receiving NovoTTF therapy with a maximal monthly compliance rate $\geq 75\%$ (≥ 18 hours daily) versus those with a $< 75\%$ compliance rate (7.7 v 4.5 months; p = .042), and Kaplan-Meier analysis demonstrated a significant trend for improved median OS with higher compliance (p = .039). Additional post hoc analysis showed significantly higher median OS with NovoTTF therapy than with BPC for patients with prior low-grade glioma, tumor size ≥ 18 cm², Karnofsky performance status ≥ 80 , and those who had previously failed bevacizumab therapy. This contrasts with the equivalent efficacy reported previously based on analysis of all randomized ITT subjects, including many who did not receive a full cycle of treatment. The authors summarized that results from the present study suggest that when used as intended, NovoTTF Therapy provides efficacy superior to that of chemotherapy in a heterogeneous population of patients with recurrent GBM. Post hoc analyses identified subgroups of patients who may be particularly good candidates for NovoTTF Therapy, pending further confirmatory studies.

Wong et al. (2014) analyzed the characteristics of responders and non-responders in both cohorts of the phase III trial which compared NovoTTF-100A Best Physician's Choice (BPC) chemotherapy for recurrent glioblastoma to determine the characteristics of response and potential predictive factors. Their analysis showed that a significantly higher proportion of NovoTTF-100A responders, five of 14 (36%), had prior low-grade histologies while none of seven (0%) BPC responder had this type of histological characteristics, suggesting that secondary glioblastoma may be more responsive to NovoTTF-100A treatment. Because primary and secondary glioblastomas have different genetic alterations, notably *EGFR* and *MDM2* amplifications together with *p16* deletion in primary glioblastomas and mutation, *IDH1* mutation and *PDGFR* amplification in secondary glioblastomas, the distinct genetic makeup in these two subtypes of glioblastomas could make secondary glioblastomas more susceptible to NovoTTF-100A treatment. Secondary glioblastomas and low dexamethasone usage are associated with a higher proportion of NovoTTF-100A responders but not BPC chemotherapy responders. The authors surmise that during treatment with NovoTTF-100A, this slower rate of tumor progression might allow enough time for the efficacy of TTFields to emerge because it may take multiple mitotic cycles to reduce the number of tumor cells and the size of the glioblastoma. The authors recommend that future clinical trials on the NovoTTF-100A device must include stratification of potential predictive factors of response that include both genetic and epigenetic determinants.

Mrugala et al. (2014) evaluated data collected from all adult patients with recurrent GBM who began commercial Novocure TTF therapy through the Patient Registry Dataset (PRiDe), which is a post-marketing registry of all recurrent GBM patients who received NovoTTF therapy in a real-world, clinical practice setting in the United States between 2011 and 2013. Data from 457 recurrent GBM patients who received Novocure TTF therapy in 91 US cancer centers were analyzed. More patients in PRiDe than the EF-11 trial received Novocure TTF therapy for first recurrence (33% v 9%) and had received prior bevacizumab therapy

(55.1% v 19%). Median OS was significantly longer with Novocure TTF therapy in clinical practice (PRiDe data set) than in the EF-11 trial (9.6 v 6.6 months; HR, 0.66; 95% CI, 0.05 to 0.86, p = .0003). One- and 2-year OS rates were more than double for Novocure TTF therapy patients in PRiDe than in the EF-11 trial (1-year: 44% v 20%; 2-year: 30% v 9%). First and second versus third and subsequent recurrences, high Karnofsky performance status (KPS), and no prior bevacizumab use were favorable prognostic factors. No unexpected adverse events were detected in PRiDe. As in the EF-11 trial, the most frequent adverse events were mild to moderate skin reactions associated with application of the Novocure TTF therapy transducer arrays. The authors concluded that results from PRiDe, together with those previously reported in the EF-11 trial, indicate that Novocure TTF therapy offers clinical benefit to patients with recurrent GBM, has high patient tolerability and favorable safety profile in the real-world, clinical practice setting.

Tumors Other Than Glioblastoma

There is a lack of published evidence from randomized controlled trials examining the long-term safety and effectiveness of TTF as a treatment for tumors other than GBM, including non-small cell lung, brain metastasis, pancreatic cancer, ovarian cancer and mesothelioma.

Ceresoli et al. (2019) conducted a prospective, single-arm, phase 2 trial (STELLAR study) with the aim to test the activity of tumor treating fields (TTFields) delivered to the thorax in combination with systemic chemotherapy for the front-line treatment of patients with unresectable malignant pleural mesothelioma. Patients were at least 18 years old, had an Eastern Cooperative Oncology Group performance status of 0-1, and at least one measurable or evaluable lesion according to modified Response Evaluation Criteria in Solid Tumors for mesothelioma. Patients received continuous TTFields at a frequency of 150 kHz to the thorax and concomitant chemotherapy with intravenous pemetrexed (500 mg/m² on day 1) plus intravenous platinum (either cisplatin 75 mg/m² on day 1 or carboplatin area under the curve 5 on day 1) every 21 days for up to six cycles. Patients not progressing after completion of chemotherapy received TTFields as maintenance treatment until progression, patient or physician decision, or unacceptable toxic effects. The primary endpoint of the trial was overall survival. Survival analyses were done in the intention-to-treat population, and safety analyses were done in all patients who received at least 1 day of TTFields treatment. A total of 80 patients were enrolled in the study. Median follow-up was 12.5 months. Median overall survival was 18.2 months (95% CI 12.1-25.8). The most common grade 3 or worse adverse events were anaemia [nine (11%) patients], neutropenia [seven (9%)], and thrombocytopenia [four (5%)]. Skin reaction was the only adverse event associated with TTFields and was reported as grade 1-2 in 53 (66%) patients, and as grade 3 in four (5%) patients. No treatment-related deaths were observed. According to the authors, the trial showed encouraging overall survival results, with no increase in systemic toxicity. TTFields (150 kHz) delivered to the thorax concomitant with pemetrexed and platinum was an active and safe combination for front-line treatment of unresectable malignant pleural mesothelioma. The lack of a comparison group limits the conclusions that can be drawn from the study. The authors indicated that further investigation in a randomized trial is warranted.

NovoTAL™ Simulation System

There is limited published clinical evidence related to the NovoTAL™ simulation system, and insufficient data to support improved long-term health outcomes with its use. This includes a small case series (Connelly et al., 2016), human head model (Wenger et al., 2016), and a user group survey (Chaudry et al., 2015). A framework for the use of NovoTAL in treatment planning has been proposed by Trusheim et al. (2016).

Clinical Practice Guidelines

American Society of Clinical Oncology (ASCO) and the Society for NeuroOncology (SNO)

ASCO and SNO published (2022) published a clinical guideline for therapy for diffuse astrocytic and oligodendroglial tumors in adults. This guideline recommends Alternating electric field therapy may be added to adjuvant TMZ in people with newly diagnosed supratentorial glioblastoma, IDH-wildtype, CNS WHO grade 4 who have completed chemoradiation therapy (Type: evidence-based, benefits outweigh harms; Evidence quality: moderate; Strength of recommendation: weak).

European Association of Neuro-Oncology (EANO) and the European Society for Medical Oncology (ESMO)

A clinical practice guideline published (2014) by EANO and ESMO recommends that at recurrence of GBM, nitrosourea regimens, TMA, and bevacizumab are options for pharmacotherapy; when available, recruitment into appropriate clinical trials should be considered. New approaches, including suicide gene therapy, immunotherapy, or TTF should only be administered in the context of clinical trials.

National Comprehensive Cancer Network (NCCN)

NCCN clinical practice guidelines for Central Nervous System Cancers (2022) state that based on results of the open-label phase III EF-14 clinical trial, concurrent treatment with adjuvant TMZ and alternating electric fields is FDA approved and recommended for newly diagnosed glioblastoma patients 70 years of age or younger who have good PS. It should also be considered a reasonable treatment option for patients older than 70 years of age with good PS and newly diagnosed glioblastoma who are treated with standard focal brain radiation and concurrent daily TMZ (p. MS-16). Alternating electric field therapy is also FDA approved for treating recurrent glioblastoma based on the safety results of this medical device from the EF-11 clinical trial. Due to a lack of clear efficacy data, the NCCN panel is divided about recommending it (p. MS-17). However, management of recurrent tumors depends on the extent of disease and patient condition. At the time of publication (June 2022), a panel discussion update was in progress. Currently, NCCN Category 1 recommendations for patients aged 70 years and younger with a good PS, regardless of the tumor's MGMT methylation status, include standard brain RT plus concurrent and adjuvant TMZ with or without alternating electric field therapy (p. MS-18). Category 1 treatment recommendations for patients older than 70 years of age with newly diagnosed glioblastoma, a good PS, and MGMT promoter methylated tumors include hypo-fractionated brain RT plus concurrent and adjuvant TMZ or standard brain RT plus concurrent and adjuvant TMZ and alternating electric field therapy. For those patients older than 70 years with newly diagnosed glioblastoma, a good PS, and with MGMT unmethylated or indeterminate tumors, hypo-fractionated brain radiation with concurrent and adjuvant TMZ is preferred, but standard brain RT plus concurrent and adjuvant TMZ and alternating electric field therapy is also a reasonable option for those elderly patients who want to be treated as aggressively as possible (p. MS-19).

U.S. Food and Drug Administration (FDA)

This section is to be used for informational purposes only. FDA approval alone is not a basis for coverage.

The Optune Treatment Kit, formerly the NovoTTF-100A System, (Novocure) was approved by the FDA in April 2011, as a novel device to treat adults age 22 years or older with glioblastoma (GBM) that recurs or progresses after receiving chemotherapy and radiation therapy. The Optune is categorized by the FDA as a stimulator, low electric field, tumor treatment; refer to the following website for the initial Premarket Approval information:

<https://www.accessdata.fda.gov/scripts/cdrh/cfdocs/cfpma/pma.cfm?id=P100034>. (Accessed June 5, 2023)

A supplemental FDA premarket approval was received in October 2015 for Optune with Temozolomide in adults with newly diagnosed, Supratentorial glioblastoma following maximal debulking surgery and completion of radiation therapy together with concomitant standard of care chemotherapy. Refer to the following website for more information:

<https://www.accessdata.fda.gov/scripts/cdrh/cfdocs/cfpma/pma.cfm?id=P100034S013>. (Accessed June 5, 2023)

The FDA has approved a humanitarian device exemption (HDE) application for the NovoTTF™-100L System for mesothelioma. Refer to the following website for more information: https://www.accessdata.fda.gov/cdrh_docs/pdf18/H180002B.pdf. (Accessed June 5, 2023)

Refer to the following website for additional information on supplemental FDA approvals for the Optune using product code NZK: <http://www.accessdata.fda.gov/scripts/cdrh/cfdocs/cfpma/pma.cfm>. (Accessed June 5, 2023)

NovoTAL simulation software is not regulated by the FDA.

References

American Brain Tumor Association (ABTA). Glioblastoma (GBM). 2016.

Ceresoli GL, Aerts JG, Dziadziuszko R, et al. Tumour treating fields in combination with pemetrexed and cisplatin or carboplatin as first-line treatment for unresectable malignant pleural mesothelioma (STELLAR): a multicentre, single-arm phase 2 trial. *Lancet Oncol*. 2019 Dec;20(12):1702-1709.

Chaudhry A, Benson L, Varshaver M, et al. NovoTTF™-100A System (Tumor Treating Fields) transducer array layout planning for glioblastoma: a NovoTAL™ System user study. *World J Surg Oncol*. 2015;13:316.

Chen W, Wang Y, Zhao B, et al. Optimal therapies for recurrent glioblastoma: a Bayesian Network meta-analysis. *Front Oncol*. 2021 Mar 29;11:641878.

Connelly J, Hormigo A, Mohile N, et al. Planning TTFIELDS treatment using the NovoTAL system-clinical case series beyond the use of MRI contrast enhancement. *BMC Cancer*. 2016 Nov 4;16(1):842.

Dono A, Mitra S, Shah M, et al. PTEN mutations predict benefit from tumor treating fields (TTFIELDS) therapy in patients with recurrent glioblastoma. *J Neurooncol*. 2021;153(1):153-160.

ECOG-ACRIN Cancer Research Group; ECOG Performance Status. 2020. Available at: <http://ecog-acrin.org/resources/ecog-performance-status>. Accessed June 5, 2023.

ECRI Institute. Custom Product Brief. Optune treatment kit (Novocure, Ltd.) for treating newly diagnosed glioblastoma. January 2019.

ECRI Institute. Custom Product Brief. Optune treatment kit (Novocure, Ltd.) for treating recurrent glioblastoma. January 2019.

ECRI Institute. Custom Product Brief. Optune treatment kit (Novocure, Ltd.) for treating newly diagnosed glioblastoma in patients aged 65 years or older. March 2019.

Hayes, Inc. Health Technology Assessment. Tumor treating fields (Optune) for treatment of glioblastoma. Lansdale, PA: Hayes, Inc.; December 2019, updated January 5, 2023.

Jin L, Guo S, Zhang X, Mo Y, et al. Optimal treatment strategy for adult patients with newly diagnosed glioblastoma: a systematic review and network meta-analysis. *Neurosurg Rev*. 2020 Oct 10.

Kanner AA, Wong ET, Villano JL, et al. Post hoc analyses of intention-to-treat population in phase III comparison of NovoTTF-100A™ system versus best physician's choice chemotherapy. *Semin Oncol*. 2014;41(suppl 6):S25-S34.

Kim CY, Paek SH, Nam DH, et al. Tumor treating fields plus temozolomide for newly diagnosed glioblastoma: a sub-group analysis of Korean patients in the EF-14 phase 3 trial. *J Neurooncol*. 2020 Feb;146(3):399-406.

Marenco-Hillebrand L, Wijesekera O, Suarez-Meade P, et al. Trends in glioblastoma: outcomes over time and type of intervention: a systematic evidence based analysis. *J Neurooncol*. 2020 Apr;147(2):297-307.

Magouliotis DE, Asproдини EK, Svokos KA, et al. Tumor-treating fields as a fourth treating modality for glioblastoma: a meta-analysis. *Acta Neurochir (Wien)*. 2018 Jun;160(6):1167-1174.

Mohile NA, Messersmith H, Gatson NT, et al. Therapy for Diffuse Astrocytic and Oligodendroglial Tumors in Adults: ASCO-SNO Guideline. *J Clin Oncol*. 2022 Feb 1;40(4):403-426.

Mrugala MM, Engelhard HH, Dinh Tran D, et al. Clinical practice experience with NovoTTF-100A system for glioblastoma: the patient registry dataset (PRiDe). *Semin Oncol*. 2014 Oct;41 Suppl 6:S4-S13.

National Cancer Institute (NCI) Dictionary of Cancer Terms. Available at: <https://www.cancer.gov/publications/dictionaries/cancer-terms>. Accessed May 22, 2023.

National Comprehensive Cancer Network (nccn). NCCN Clinical practice guidelines in oncology (NCCN Guidelines®). Central nervous system cancers. Version 1.2023 - March 24, 2023.

Onken J, Goerling U, Heinrich M, et al. Patient reported outcome (PRO) among high-grade glioma patients receiving TTFIELDS treatment: a two center observational study. *Front Neurol*. 2019 Oct 1;10:1026.

Ram Z, Kim CY, Hottinger AF, et al. Efficacy and safety of tumor treating fields (TTFIELDS) in elderly patients with newly diagnosed glioblastoma: subgroup analysis of the phase 3 EF-14 clinical trial. *Front Oncol*. 2021 Sep 27;11:671972.

Regev O, Merkin V, Blumenthal DT, et al. Tumor-treating fields for the treatment of glioblastoma: a systematic review and meta-analysis. *Neurooncol Pract*. 2021 Apr 20;8(4):426-440.

Shi W, Blumenthal DT, Oberheim Bush NA, et al. Global post-marketing safety surveillance of Tumor Treating Fields (TTFIELDS) in patients with high-grade glioma in clinical practice. *J Neurooncol*. 2020 Jul;148(3):489-500.

Stupp R, Taillibert S, Kanner A, et al. Effect of tumor-treating fields plus maintenance temozolomide vs maintenance temozolomide alone on survival in patients with glioblastoma: a randomized clinical trial. *JAMA* 2017 Dec 19;318(23):2306-2316.

Stupp R, Taillibert S, Kanner AA, et al. Maintenance therapy with tumor-treating fields plus temozolomide vs temozolomide alone for glioblastoma. *JAMA*. 2015;314(23):2535-2543.

Taphoorn MJB, Dirven L, Kanner AA, et al. Influence of treatment with tumor-treating fields on health-related quality of life of patients with newly diagnosed glioblastoma: a secondary analysis of a randomized clinical trial. *JAMA Oncol.* 2018 Apr 1;4(4):495-504.

Toms SA, Kim CY, Nicholas G, et al. Increased compliance with tumor treating fields therapy is prognostic for improved survival in the treatment of glioblastoma: a subgroup analysis of the EF-14 phase III trial. *J Neurooncol.* 2019 Jan;141(2):467-473.

Trusheim J, Dunbar E, Battiste J, et al. A state-of-the-art review and guidelines for tumor treating fields treatment planning and patient follow-up in glioblastoma. *CNS Oncol.* 2017 Jan;6(1):29-43.

Wenger C, Salvador R, Basser PJ, et al. Improving tumor treating fields treatment efficacy in patients with glioblastoma using personalized array layouts. *Int J Radiat Oncol Biol Phys.* 2016 Apr 1;94(5):1137-43.

West HJ, Jin JO. *JAMA Oncology Patient Page.* Performance status in patients with cancer. *JAMA Oncol.* 2015 Oct;1(7):998.

Wong ET, Lok E, Swanson KD, et al. Response assessment of NovoTTF-100A versus best physician's choice chemotherapy in recurrent glioblastoma. *Cancer Med.* 2014;3(3):592-602.

Wong ET, Lok E, Swanson KD. Clinical benefit in recurrent glioblastoma from adjuvant NovoTTF-100A and TCCC after temozolomide and bevacizumab failure: a preliminary observation. *Cancer Med.* 2015 Mar; 4(3): 383–391.

Policy History/Revision Information

Date	Summary of Changes
03/01/2024	<p>Coverage Rationale</p> <ul style="list-style-type: none"> Replaced language indicating “subsequent approval(s) for continuation of electric TTF for treatment of histologically-confirmed supratentorial glioblastoma (GBM) is based on [the listed criteria]” with “subsequent approval(s) for continuation <i>beyond the initial 3 months</i> of electric TTF for treatment of histologically-confirmed Supratentorial glioblastoma (GBM) is based on [the listed criteria] <p>Applicable Codes</p> <ul style="list-style-type: none"> Removed notation indicating CPT code 77299 is not on the New Jersey Medicaid Fee Schedule and therefore may not be covered by the state of New Jersey Medicaid Program <p>Supporting Information</p> <ul style="list-style-type: none"> Updated <i>Clinical Evidence</i> and <i>References</i> sections to reflect the most current information Archived previous policy version CS146NJ.G

Instructions for Use

This Medical Policy provides assistance in interpreting UnitedHealthcare standard benefit plans. When deciding coverage, the federal, state or contractual requirements for benefit plan coverage must be referenced as the terms of the federal, state or contractual requirements for benefit plan coverage may differ from the standard benefit plan. In the event of a conflict, the federal, state or contractual requirements for benefit plan coverage govern. Before using this policy, please check the federal, state or contractual requirements for benefit plan coverage. UnitedHealthcare reserves the right to modify its Policies and Guidelines as necessary. This Medical Policy is provided for informational purposes. It does not constitute medical advice.

UnitedHealthcare may also use tools developed by third parties, such as the InterQual[®] criteria, to assist us in administering health benefits. The UnitedHealthcare Medical Policies are intended to be used in connection with the independent professional medical judgment of a qualified health care provider and do not constitute the practice of medicine or medical advice.