Tumor treating fields is a newly epoch making therapeutic method for malignant brain tumors

Abstract

Glioblastoma (GBM) is a malignant brain tumor and extremely difficult to treat with no specific curative treatment. Tumor treating fields (TTFIELDS/TTF) are a new cancer treatment modality for patients with GBM, which uses alternating electrical fields. This review introduces the principle of TTF and reports on recent GBM treatments and prospects.

Keywords: tumor treating fields, glioblastoma, alternating electric fields

Introduction

Glioblastoma (GBM) possesses a lethal prognosis. Despite surgery, radiotherapy, and chemotherapy, the median survival time for a patient with GBM is only 15.6months. TTF have a very different mechanism from conventional treatment methods in which malignant tumor cells placed in a specific electric field environment are broken down and lead to apoptosis. Several multicenter studies to treat GBM using TTF have been performed so far and have shown superior results.

Discussion

Basics and theory of TTF

To date, many studies have shown the influence of the electric field on the body. Kirson et al. explained the basic principle of TTF in 2004. TTF create an alternating electric field in the brain that inhibits cell division, thereby causing cells to trigger apoptosis. TTF are thought to be effective in two different phases during cell division. First, when microtubules are polymerized and mitotic spindles are formed (from the early to middle stages of mitosis), an electric field inhibits polymerization and cell division, leading to apoptosis. Second, the cells take an hourglass-like form in the late stage of mitosis and during cytokinesis, and a slight electric field change occurs in the cells. At this time, when the electric field is applied, the component is pushed out toward the constricted part of the form, destroying the structure of the cell and leading to apoptosis. TTF utilize the effect obtained by providing an electric field to the cells at the time of cytokinesis from mitosis in a very brief period during cell division and inhibits cell division. As a matter of course, this therapy has no effect on cells that are not dividing. Furthermore, it has the highest therapeutic effect when the axes of division and electric field are parallel, but when it is at a right angle, the effect becomes the lowest. Therefore, it does not mean that all cells are affected by the electric field and cause apoptosis.

Clinical trial for GBM using TTF

Stupp et al. reported EF-11 in 2012 as the first multicenter randomized clinical trial conducted using TTF for GBM. Patients with recurrent GBM were randomized to receive a treatment with TTF alone (n=120) or chemotherapy (n=117). The median overall survival (OS) after randomization was 6.6 versus 6.0months. The median progression free survival (PFS) rate was 2.2 and 2.0months. No significant difference was observed among them. However, significantly fewer systematic side effects were observed with TTF alone. Gastrointestinal disorders, blood system disorders, and infections, which are typical adverse events of chemotherapy, were significant low in the TTF alone. Contact dermatitis of the scalp, which is an adverse event related to TTF, was found in 16% of patients and could be treated only by a steroid ointment application. This result showed that toxicity and quality of life clearly favorable TTF.

Stupp et al. reported EF-14 as a phase 3 randomized clinical trial of TTF for newly diagnosed GBM in 2015. After completion of chemoradiotherapy, patients with newly diagnosed GBM were randomized for treatment with TTF plus temozolomide maintenance therapy (n=466) or temozolomide maintenance therapy alone (n=229). The median OS was 19.6 months in the TTF plus temozolomide group and 16.6 months in the temozolomide alone group, and the median PFS was 7.1 and 4.0months, respectively. These result showed that TTF significantly prolongs PFS and OS in patients with newly diagnosed GBM.

Prospects

It is expected that TTF become a new cancer treatment modality for patients with GBM. Because of its mechanism of action, TTF therapy alone only effects cells in the division phase, but it has been reported that enhance a cell proliferation inhibitory effect in combination with chemotherapy and affect cell self-repair function. Currently, clinical trials are being planned for GBM as well as for lung, pancreatic, breast, and ovarian cancers.
Conclusion

In recent years, TTF have gained attention as a new treatment for GBM. Many multicenter clinical trials have shown that TTF prolong the prognosis without weakening Activity of Daily Living.

Conflict of interest

The authors declare that there are no conflicts of interest regarding to the publication of this paper.

References


