

TFields In Germany in routine clinical care

Submission date 27/07/2017	Recruitment status No longer recruiting	<input checked="" type="checkbox"/> Prospectively registered <input type="checkbox"/> Protocol
Registration date 08/08/2017	Overall study status Completed	<input type="checkbox"/> Statistical analysis plan <input type="checkbox"/> Results
Last Edited 21/02/2023	Condition category Cancer	<input type="checkbox"/> Individual participant data <input type="checkbox"/> Record updated in last year

Plain English summary of protocol

Background and study aims

Glioblastoma (GBM) is the most common malignant primary tumor (cancer) of the brain. The current standard of care for patients with newly diagnosed GBM consists of maximal surgical resection (surgically removing the cancer), radiotherapy together with chemotherapy using temozolomide (TMZ), followed by maintenance TMZ for six months. This treatment scheme was shown to extend median survival from 12.1 to 14.6 months compared to surgery and radiotherapy alone. This survival was essentially unchanged since 2005 despite numerous other studies. Although immense efforts have been made over the years with different treatment strategies, the survival of patients with newly diagnosed GBM remained very poor until recently. Tumor-treating fields (TFields) are low-intensity, intermediate frequency, alternating electric fields delivered continuously through adhesive patches, called transducer arrays, to the area of the brain where the GBM tumor is located and help slow down or stop glioblastoma cancer cells from dividing. These transducer arrays are applied to the scalp and are connected to the wearable and portable device. The aim of this study is to collect real life data on the use of tumor-treating fields (TFields) in patients with newly diagnosed GBM in routine clinical care in Germany.

Who can participate?

Patients aged 18 and older who have GBM

What does the study involve?

This study is a chart review study. Participants who receive TFields treatment have data collected about their care for 24 months. Data is collected about their health and any serious adverse events that occurred during their care. Participants complete a quality of life questionnaire and questionnaire about TFields at baseline and during their follow up treatment.

What are the possible benefits and risks of participating?

There are no direct benefits or risks with participating

Where is the study run from?

This study is being run from 5 hospitals in Germany

When is the study starting and how long is it expected to run for?

July 2017 to August 2022

Who is funding the study?
Novocure GmbH (Germany)

Who is the main contact?

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2. Prof. Dr Oliver Bähr

Contact information

Type(s)

Public

Contact name

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Additional identifiers

EudraCT/CTIS number

IRAS number

ClinicalTrials.gov number

NCT03258021

Secondary identifying numbers

05.07.2017

Study information

Scientific Title

TTFields In GERMANY in Routine clinical care (TIGER): the use of TTFields for newly diagnosed GBM patients in Germany in routine clinical care

Acronym

TIGER

Study objectives

The purpose of this post-authorisation medical device study is to obtain real life data on the use of tumor-treating fields (TTFields) in patients with newly diagnosed GBM in routine clinical care in Germany.

Ethics approval required

Old ethics approval format

Ethics approval(s)

Ethikkommission des FB Medizin der J.W. Goethe-Universität, 23/08/2017, ref: 280/17

Study design

Multicentre prospective non-interventional observational post-authorisation medical device study

Primary study design

Observational

Secondary study design

Cohort study

Study setting(s)

Other

Study type(s)

Treatment

Participant information sheet

Not available in web format, please use contact details to request a participant information sheet.

Health condition(s) or problem(s) studied

Glioblastoma

Interventions

This is non-interventional study evaluating use of tumor-treating fields (TTFields) in newly diagnosed glioblastoma (GBM) patients in routine clinical care. TTFields help slow down or stop

glioblastoma cancer cells from dividing by disrupting dividing mechanism of cancer cells leading to apoptosis. TTFields are low-intensity, intermediate frequency, alternating electric fields delivered continuously through adhesive patches, called transducer arrays, to the area of the brain where the GBM tumor is located. These transducer arrays are applied to the scalp and are connected to the wearable and portable device. TTFields are approved for the treatment of newly diagnosed and recurrent GBM.

Eligible participants are enrolled in the study after signing informed consent to use their data and process it centrally for research purposes. Participants do not experience any tests or procedures that are not part of routine clinical care. For patients refusing TTFields treatment at baseline only baseline data is collected. Patient who initiate TTFields Therapy are followed-up during the first 24 months after study enrolment, LPI has a max. of 18 months follow-up.

Data obtained from patient files: Only data which can be obtained from routine clinical care files of the patients is recorded. Data of each routine clinical care visit during the first 24 months after study enrolment is reported in the eCRF. Data is derived from clinical records and findings, radiological assessments (e.g. contrast enhanced MRI), observations or other sources (e.g. hospital records, clinical and office charts, electronic patient records, laboratory notes, recorded data from automated device). Serious Adverse Events (SAEs) are documented during follow-up visits by the treating investigator and subsequently centrally assessed by an independent Clinical Event Committee.

Participants also complete quality of life questionnaires and a study-specific questionnaire on TTFields at baseline and during follow-up.

Intervention Type

Other

Primary outcome measure

1. Time to death of any cause (overall survival [OS]) from diagnosis is measured using the patient data from date of enrollment until the date of death from any cause
2. Number of TTFields treatment-related (serious adverse events) SAEs standardised to one year of FU is measured using the collection of SAEs during the follow-up period
3. Number of SAEs after start of TTFields treatment is measured using the collection of SAEs at the follow-up period
4. Time of usage (compliance) of TTFields treatment over time is measured using the treatment compliance report at the follow-up period
5. Time to first progression of GBM (progression-free survival [PFS]), defined within radiological and/or clinical/neurological assessment during routine clinical care in patients who started TTFields treatment at baseline from date of enrollment until the date of first progression of GBM
6. Changes in quality of life after start of TTFields treatment compared to baseline in patients who started TTFields treatment at baseline is measured using the QoL questionnaires at months two and four after start of TTFields treatment
7. Patients' reason(s) for refusing TTFields is measured using a questionnaire at baseline

Secondary outcome measures

There are no secondary outcome measures

Overall study start date

05/07/2017

Completion date

31/08/2022

Eligibility

Key inclusion criteria

1. Newly diagnosed histologically confirmed GBM (WHO-Grade IV)
2. Patient within first 3 cycles of first-line tumor-specific maintenance chemotherapy
3. ≥ 18 years of age
4. Clinical indication for TTFields treatment
5. Given informed consent for use and processing of data

Participant type(s)

Patient

Age group

Adult

Lower age limit

18 Years

Sex

Both

Target number of participants

About 1000

Total final enrolment

709

Key exclusion criteria

1. Present or planned pregnancy
2. Significant additional neurological disease (e.g. significantly increased intracerebral pressure (ICP) with a significant midline shift of the brain)
3. Active implanted medical device (e.g. deep brain stimulator)
4. Documented allergy to conductive hydrogel
5. Skull defect (e.g. missing bone with no replacement, bullet fragments in the skull)

Date of first enrolment

31/08/2017

Date of final enrolment

30/11/2019

Locations

Countries of recruitment

Germany

Study participating centre
Clinical Centre Aschaffenburg
Aschaffenburg
Germany
63739

Sponsor information

Organisation
Novocure GmbH

Sponsor details
Elektrastraße 6
Munich
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81925

Sponsor type
Industry

ROR
<https://ror.org/04pspdc11>

Funder(s)

Funder type
Industry

Funder Name
Novocure GmbH

Results and Publications

Publication and dissemination plan
Planned publication in a high-impact peer-reviewed journal.

Intention to publish date
30/04/2022

Individual participant data (IPD) sharing plan

The datasets generated during and/or analysed during the current study are not expected to be made available due to data protection.

IPD sharing plan summary

Not expected to be made available