sociation with malignant brain tumor was analyzed. The frequency of stroke in 287 patients with primary glioblastoma and 217 patients with metastatic brain tumor was also analyzed.

RESULTS: Twenty one (4.1%) patients with ischemic stroke and 26 (5.1%) patients with hemorrhagic stroke patients had malignant brain tumor, and most tumors were either malignant glioma or metastatic brain tumor. A medical history of cranial irradiation was seen in 66.7% of patients with ischemic stroke, and 80% of hemorrhagic stroke occurred within the tumor before starting the treatments. Either ischemic or hemorrhagic stroke occurred in 9.1% of patients with glioblastoma and 4.1% of patients with metastatic brain tumor, and the number of ischemic and hemorrhagic were almost the same. In patients with glioblastoma, nearly half of the stroke cases were associated with bevacizumab. Half of the cases of bevacizumab-related stroke were asymptomatic, while asymptomatic cases were seen in 21.4% for non-bevacizumab cases.

DISCUSSION: Stroke is not an uncommon complication in patients with malignant brain tumor but only a restricted number of cases are preventable. Including the cases of bevacizumab-related stroke, which is often asymptomatic, accurate diagnosis and the second prevention would be important.

COT-08
ANALYSIS OF PROGNOSIS OF BIOPSY/PARTIAL RESECTION CASES OF MALIGNANT GLIOMA
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INTRODUCTION: Malignant glioma is the most common and aggressive primary brain tumor and requires multimodality treatment. Regarding surgical treatment, it is desirable to achieve maximum resection while considering function preservation. There is consensus that the survival prognosis is prolonged in gross or subtotal resection. However, there are cases in which biopsy or partial resection is performed due to the spread of lesions at the time of onset, underlying diseases, and social background. The purpose of this study was to retrospectively analyze the cases of malignant glioma at our university and to find out the factors related to the prognosis of cases in which removal was insufficient.

TARGET: 55 cases of malignant glioma treated at our university since 2013 who underwent biopsy or partial resection.

METHOD: Overall/progression-free survival period is the end point, and parameters are age, bevacizumab use, pathological diagnosis, photodynamic diagnosis use at operation, immunotherapy, ventricular invasion, contralateral invasion, sex, preoperative Performance Status (PS), postsurgical PS, left or right, navigation use, steroid use, anticoagulant drug type, radiation, IDH mutation, 1p19q co-deletion, MGMT methylation, TERT mutation, p53 mutation, biopsy or partial resection. After narrowing down the evaluation items by univariate analysis(Logrank test), multivariate analysis(Cox proportional hazard model) was performed.

RESULT: The univariate analysis was significant in 5 items including bevacizumab use, radiation therapy, levantacetaxan use, postoperative PS70 or higher, and partial resection instead of biopsy. Multivariate analysis detected two statistically significant differences, bevacizumab use and post-operative PS70 and above. There was no difference in the timing of bevacizumab use.

CONSIDERATIONS: In poorly resection cases, the weight of postoperative treatment is high, so continuity of treatment and selection of postoperative treatment are important, and maintenance of ADL and use of bevacizumab are significant among them.

COT-11
ADMINISTRATION OF BEVACIZUMAB FOR PATIENTS WHO FAILED TO COMPLETE STUPP REGIMEN AFTER GliOBlastoma SURGERY
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Stupp regimen is widely used as the standard treatment after glioblastoma surgery, but in some cases treatment must be discontinuated for various reasons. We experienced Bevacizumab in two patients who were unable to continue treatment in the Stupp regimen, and report our experience with literature review. First patient is a man in his 60s. Resection of glioblastoma of the left cerebral hemisphere was performed, and postoperatively right hemiparesis and aphasia remained. Irradiation and administration of Temozolomide were performed, but Temozolomide was unable to continue because of side effects. After system management, Bevacizumab was administered, and resection of residual tumor and peripheral edema were observed, and the patient began to speak. After 12 cycles of administration, the tumor regrew, and he died. Second patient is a woman in her 80s. Cricotomaty was performed for hemiorthic infarction of the left cerebral hemisphere, postoperatively, aphasia, right hemiparesis remained, bedridden, and was unable to eat. Four months after initial surgery, a tumor was found in left parietal lobe and was resected. The pathological diagnosis was glioblastoma. For the treatment of recurrence, the patient was unable to be transferred for radiochemotherapy, so the patient was treated with Temozolomide and Bevacizumab. The patient’s condition became better, eat by herself, and could play in rehabilitation facility on the wheelchair. After 12 cycles of bevacizumab, the tumor subsequently enlarged, and died. Although the effect is limited, there are some cases in which Bevacizumab administration could maintain patient’s condition by controlling tumor growth for a certain period of time. From the experience of these patients, it seems that even in patients in postoperative poor Karnofsky Performance Status (KPS)and elderly people, Bevacizumab administration would be an option before transitioning to end-of-life care.

COT-12
THE ROLE OF CLINICAL RESEARCH PROFESSIONAL IN THE REGISTRATION STUDY OF PEDIATRIC SOLID TUMOR IN JAPAN CHILDREN’S CANCER GROUP
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A larger scale prospective registration study for pediatric solid tumor has been conducted nationwide in Japan since 2011 in Japan’s Children’s Cancer Group (JCCG). In the study, the clinical data and surgical specimen are collected into the National Center for Child Health. Kyoto University Hospital has been participated in the study since IRB approval in 2011. We reviewed 115 patients registered to the study and assessed the role of clinical research professional in the registration study. Fifty-one patients with pediatric brain tumors participated in this study from 2011 to 2020. There were 17 intracranial germ cell tumors, 9 medulloblastomas, 14 gliomas and ependymomas in 5 diffuse midline gliomas, 9 pilocytic astrocytoma, and 2 other types of tumor. Forty surgical specimens were collected for central review. The status of clinical data entry was complete in 33 patients. The registrations and sending of clinical data and specimens have remarkably increased without exceptions since a clinical research professional supported the study in 2018. The study collecting and analyzing pathological diagnosis, molecular diagnosis, treatment, and clinical information in patients with pediatric brain tumor are important to realize the current status. The clinical research professional plays an important role to register patients and to send the specimens and clinical data into the study.