Original Article

5-Aminolevulinic acid fluorescence-guided endoscopic surgery for intraventricular tumors

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ABSTRACT

Background: In recent years, the efficacy of 5-aminolevulinic acid photodynamic diagnosis (5-ALA PDD) has been reported for various types of brain tumors, including malignant glioma. In addition, many reports have been published on the usefulness of neuroendoscopic surgery for intraventricular lesions. However, no systematic report is available on the combined use of 5-ALA PDD and neuroendoscopy for various intraventricular tumors.

Methods: We report 17 consecutive patients with intraventricular tumors. All patients received oral 5-ALA preoperatively and underwent endoscopic surgical treatment (resection or biopsy). We use a rigid endoscope with a built-in PDD system for intraoperative observation.

Results: Seven resections and 10 biopsies were performed. Histopathological diagnosis was confirmed in all 17 cases. Gross total resection was achieved in six of seven cases. The fluorescence positivity rates for each tumor were glioblastoma 100% (2/2), low-grade glioma 67% (2/3), subependymoma 0% (0/1), medulloblastoma 100% (1/1), pineoblastoma 0% (0/1), germ cell tumor 75% (3/4), diffuse large B-cell lymphoma 33% (1/3), and metastatic tumor 100% (2/2).

Conclusion: Our method has the potential to improve detection of residual tumors in blind spots and deep areas, as well as the accuracy and safety of biopsy procedures for intraventricular lesions that are difficult to view and treat under a microscope.

Keywords: 5-Aminolevulinic acid, Brain tumor, Intraventricular lesion, Neuroendoscope

INTRODUCTION

The efficacy of surgical treatment of malignant glioma following visualization with 5-aminolevulinic acid (5-ALA) has become widely known following the report by Stummer et al.[23,25] In recent years, the efficacy of 5-ALA photodynamic diagnosis (PDD) for various brain tumors as well as malignant glioma has been reported.[1,6,8,11,14-18,26,30] On the other hand, many reports have described the efficacy of neuroendoscopic surgery for intraventricular lesions.[5,7,28] Neuroendoscopic surgery can be used for removal and biopsy of deep-seated lesions as well as for performing endoscopic third ventriculostomy (ETV) simultaneously in cases of complicated hydrocephalus. Combined treatment with 5-ALA PDD and neuroendoscopy was first reported by Tamura et al. in 2007 for intraventricular glioma.[27] Since then, a number of reports have been
published on its usefulness.[4,20,21,26] However, no systematic report is available that describes this combined therapy for intraventricular tumors. In this study, we prospectively examined the usefulness of neuroendoscopic 5-ALA PDD for brain tumors located intraventricularly or paraventricularly.

MATERIALS AND METHODS

We included 17 consecutive patients with suspected malignant brain tumors located in the ventricles who underwent resection or biopsy at the Department of Neurosurgery, Kansai Medical University, between October 2014 and November 2021. All patients underwent neuroendoscopic surgery associated with 5-ALA PDD.

Surgical procedure

The patients were orally administered 20 mg/kg 5-ALA 3 h before induction of anesthesia. We used a special rigid endoscope (PDD endoscope system with D Light C system, KARL STORZ GmbH and Co., Tuttlingen, Germany) for intraoperative observation of 5-ALA fluorescence. The D Light C system has a 300 W Xe short-arc lamp and uses white light or blue light (380–420 nm) as excitation light. In addition, a dichroic long-pass filter is attached to the endoscope eyepiece for fluorescence observation. All fluorescence assessments were for in situ lesions in this study. The focus of the endoscope was set at a distance of 1 cm from the tip in the sheath before observing the lesion, which enabled the evaluation of fluorescence at the same distance from the lesion for all observations. The endoscope was used only at a tip angle of 0 degrees and the fluorescence was evaluated at the same angle by placing the lesion in the center of the field of view. For biopsies, the target and trajectory were determined with a neuronavigation system (Stealth Station or Stealth S7, Medtronic Navigation, Denver, CO.). A burr hole or small craniotomy was performed and a cortical incision of approximately 1 cm was performed. A transparent sheath (Neuropor Mini, Hakko Co., Japan) with a diameter of 6.8 mm was inserted into the target or ventricle and the neuroendoscope was inserted into the sheath to observe the fluorescence of the intraventricular lesion or the intraparenchymal lesion at the tip of the sheath. For resections, the navigation system was used and a small craniotomy was performed as the basic method, but in cases in which endoscopy was performed alone, the ViewSite Brain Access System (Vykor Medical, Inc., Boca Raton, FL, USA) was used, and forceps and suction tubes were used to remove the tumor in a piecemeal fashion. A brain retractor was used in cases in which the resection was performed using a combination of endoscopy and microscopy (OPMI Pentero, Carl Zeiss Surgical GmbH and Co., Oberkochen, Germany).

Postoperative evaluation

All patients underwent a computed tomography scan within 24 h postoperatively to assess the possibility of postoperative bleeding. For resections, magnetic resonance imaging (MRI) examinations were performed within 72 h postoperatively, and T1-weighted images (noncontrast and contrast) were used to evaluate the removal rate, which was classified as gross total resection (GTR) and partial resection. Histopathological diagnosis was performed for each patient and postoperative complications were evaluated.

Ethics committee approval and statement regarding patient consent

The protocol for this study was approved by the Ethics Committee of Kansai Medical University (No. 2018037). The requirement for written patient consent was waived because data were deidentified.

Statistical analysis

All statistical analyses were performed using JMP version 14.2 (SAS Institute, Cary, NC, USA). Fisher’s exact test was performed to compare the fluorescence evaluation between the following two groups: presence or absence of MRI contrast effect, intraventricular lesion or paraventricular lesion, and certain entity of tumor or not. A significant difference was considered to be \( P < 0.05 \).

RESULTS

The histopathological diagnosis was obtained in all 17 patients. Tumors were diagnosed as glioblastoma (GBM) in two cases, low-grade gliomas (LGGs) in three cases, subependymoma in one case, medulloblastoma in one case, pineoblastoma in one case, germ cell tumor in four cases, diffuse large B-cell lymphoma (DLBCL) in three cases, and metastatic tumor in two cases. Ten patients underwent biopsy and seven patients underwent resection. Of the seven resections, four cases were performed with endoscopy alone and GTR was achieved in 86% (6/7) of the cases. ETV was performed simultaneously with surgery in six cases (three biopsies and three resections). The clinical and histological characteristics of the patients, the details of the surgical procedures, and the fluorescent diagnoses are summarized in Table 1. No cases of postoperative worsening of neurological symptoms occurred, but postoperative hemorrhage was observed in patient No. 11 (metastatic brain tumor), which was consistent with the excision cavity. Patient No. 10 (pilocytic astrocytoma) showed postoperative leakage of cerebrospinal fluid. No adverse events related to 5-ALA administration occurred.
Fluorescence properties of tumors

The fluorescence positivity rates by tumor were GBM 100% (2/2), LGGs 67% (2/3), subependymoma 0% (0/1), medulloblastoma 100% (1/1), pineoblastoma 0% (0/1), germ cell tumor 75% (3/4), DLBCL 33% (1/3), and metastatic tumor 100% (2/2).

Relationship between 5-ALA fluorescence positivity and tumor characteristics

We performed statistical analysis of the relationship between 5-ALA positivity and MRI contrast enhancement (one patient was not given contrast medium due to renal dysfunction), intraventricular or paraventricular localization, and the histology of each tumor. No statistically significant correlations were found for any items, but relationships between 5-ALA positivity and MRI contrast enhancement (71%; 10/14), intraventricular localization (82%; 9/11), and histology of germ cell tumor (75%; 3/4) may exist [Table 2].

Representative cases

Case No. 4: A biopsy was performed for a contrast-enhancing lesion located in the left caudate nucleus. We planned a contralateral transventricular approach. The lesion was approached from the right anterior horn and reached the contralateral anterior horn after performing a septostomy. 5-ALA PDD showed diffuse fluorescence at the wall of the ventricle presumably overlying the tumor, and we performed a biopsy of the same lesion and diagnosed it as DLBCL [Figure 1].

Case No. 10: A patient who had undergone a transventricular biopsy 10 years earlier at another hospital underwent resection of a pilocytic astrocytoma located in the third
ventricle. A right anterior horn approach was performed through a small craniotomy and a 1.5 cm cortical incision. Because we were able to use the previous surgical corridor, we reached the right lateral ventricle directly with the endoscope without a sheath. PDD performed in the ventricles showed strong fluorescence in lesions consistent with tumors. Removal was performed in a piecemeal fashion using forceps and sometimes suction tubes, and hemostasis was confirmed under direct observation. The postoperative course was uneventful and GTR was confirmed by postoperative contrast-enhanced MRI [Figure 2].

**DISCUSSION**

**Evaluation of 5-ALA fluorescence in intraventricular lesions**

Intraventricular areas are rather difficult to reach during microscopic surgery and the possibility of endoscopic surgery has been discussed.[5,7] ETV can be performed simultaneously with tumor removal and biopsy, and its usefulness has already been established. Difficulty in identifying the tumor has been reported, making surgery difficult.[10,12,29] Fluorescence-guided endoscopic surgery for tumor identification has been

![Figure 1](image1.png)

**Figure 1:** The patient in Case 4 underwent tumor biopsy for a basal ganglia DLBCL through the contralateral anterior horn. Diffuse fluorescence was observed in the lateral ventricle under blue light. Red dots indicate the site of observation. Arrowheads indicate the site of positive fluorescence.

![Figure 2](image2.png)

**Figure 2:** The patient in Case 10 underwent tumor resection for a third ventricle pilocytic astrocytoma through the right anterior horn. (a and b) MRI shows the presurgical view and postsurgical state of gross total removal. (c and d) Observation in the third ventricle by white light and blue light. Strong fluorescence was observed at the site of the tumor lesion. (e) Removal in a piecemeal manner was performed with forceps. (f) Gross total removal was achieved. Red dots indicate the site of observation. Arrowheads indicate the site of positive fluorescence.
reported using indocyanine green and fluorescein sodium in addition to 5-ALA. Indocyanine green and fluorescein sodium are particularly useful from an economic viewpoint and in periventricular structures because of their ability to enable visualization of not only tumors but also the vascular architecture. However, these fluorescent molecules also show disruption of the blood–brain barrier in a way that is fundamentally different from 5-ALA, which shows the fluorescence of protoporphyrin IX, a metabolic product of 5-ALA by tumor cells. 5-ALA may be more useful for discriminating false positives with regard to inflammatory and edematous lesions, tumor margins, and disseminated disease.

**Indications for 5-ALA**

Recently, fluorescence positivity of 5-ALA has been reported in various types of tumors as well as malignant gliomas. Although off-label use of 5-ALA is expanding, its routine use is being questioned because it may not be useful in all tumors. Stummer et al. reported that it is useful for supratentorial and contrast-enhanced lesions in pediatric brain tumors. Other conditions that may benefit from the use of 5-ALA, such as other tumor types and tumors with surrounding invasion, have also been reported. Results from this study suggested that contrast-enhanced lesions, intraventricular lesions, and germ cell tumors may be conditions that benefit from the use of 5-ALA. Further prospective studies may help to determine the precise indications.

**Advantages of our procedure**

Since Tamura et al. reported the first 5-ALA PDD-guided endoscopic surgery for intraventricular glioma biopsy in 2007 as a technical note, scattered case reports have been published on this technique. Rapp et al. reported nine cases and Choo et al. reported 20 cases, but these were reports of intraparenchymal tumors. Takeda et al. reported only two cases limited to intraventricular tumors. We analyzed 17 cases of various types of tumors in the ventricles and this is the first systematic report to our knowledge. In the biopsy procedure, we were able to obtain a sufficient amount of specimen for histological diagnosis without complications. The advantages of biopsy with 5-ALA PDD are that the core of the lesion can be visualized accurately and the number of samplings can be reduced, thus reducing the risk of damaging critical structures. In particular, in this study, intraventricular localization tended to be positively correlated with 5-ALA fluorescence positivity. Although reports have suggested that no clear difference exists in the accuracy and safety between simple endoscopic and stereotactic biopsy for intraventricular lesions, the use of fluorescent diagnosis may increase the accuracy and safety of endoscopic biopsy. In addition, even when microscopy is used mainly during tumor removal, the combined use of neuroendoscopic 5-ALA PDD may allow observation of blind spots in deep lesions and residual tumors in areas with insufficient fluorescence excitation, thereby maximizing tumor removal.

**Limitations and future**

In some cases in our study, the ventricular wall was fluorescence positive despite the absence of an obvious tumor. We did not perform histopathological examination of those fluorescence-positive areas. Reporting on GBM, Hayashi et al. described the disruption of ependymal cells and the presence of tumor cells in fluorescence-positive areas of the ventricular wall. On the other hand, Moon et al. reported the presence of tumor cells in the fluorescence-positive ventricular wall in 5/11 cases. This issue cannot be avoided when performing 5-ALA PDD-guided endoscopic surgery for intraventricular tumors and pathological studies will have to be conducted in the future. In addition, the fluorescence intensity was not quantitatively evaluated in this study and quantitative evaluation such as spectroscopy is necessary in the future.

**CONCLUSION**

We report the results of 5-ALA PDD combined with neuroendoscopic surgery for various types of intraventricular tumors. We suggest that our procedure for intraventricular lesions may improve the detection of blind spots and deep-seated residual tumors, as well as the accuracy and safety of biopsies.

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**Declaration of patient consent**

Institutional Review Board (IRB) permission obtained for the study.

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**Conflicts of interest**

There are no conflicts of interest.

**REFERENCES**

1. Barbagallo GM, Certo F, Heiss K, Albanese V. 5-ALA fluorescence-assisted surgery in pediatric brain tumors: Report of three cases and review of the literature. Br J Neurosurg
2. Birski M, Furtak J, Krystkiewicz K, Birksa J, Zielsinska K, Sokal P, et al. Endoscopic versus stereotactic biopsies of intracranial lesions involving the ventricles. Neurosurg Rev 2021;44:1721-7.

3. Boschi A, Della Puppa A. 5-ALA fluorescence on tumors different from malignant gliomas. Review of the literature and our experience. J Neurosurg Sci 2019;63:661-9.

4. Choo J, Takeuchi K, Nagata Y, Ohka F, Kishida Y, Watanabe T, et al. Neuroendoscopic cylinder surgery and 5-aminolevulinic acid photodynamic diagnosis of deep-seated intracranial lesions. World Neurosurg 2018;116:e33-41.

5. Chowdhry SA, Cohen AR. Intraventricular neuroendoscopy: Complication avoidance and management. World Neurosurg 2013;79 Suppl 2:S15.e1-10.

6. Cornelius JR, Kamp MA, Tortora A, Knipps J, Krause-Molle Z, Beer T, et al. Surgery of small anterior skull base meningiomas by endoscopic 5-aminolevulinic acid fluorescence guidance: First clinical experience. World Neurosurg 2019;122:e890-5.

7. Du B, Shan AJ, Peng YP, Wang J, Peng KW, Zhong XL, et al. A new modified neuroendoscope technology to remove severe intraventricular haematoma. Brain Inj 2018;32:1142-8.

8. Eljamel MS, Leese G, Moseley H. Intraoperative optical identification of pituitary adenomas. J Neurooncol 2009;92:417-21.

9. Fiorindi A, Boaro A, Del Moro G, Longatti P. Fluorescein-guided neuroendoscopy for intraventricular lesions: A case series. Oper Neurosurg (Hagerstown) 2017;13:173-81.

10. Fiorindi A, Longatti P. A restricted neuroendoscopic approach for pathological diagnosis of intraventricular and paraventricular tumours. Acta Neurochir (Wien) 2008;150:1235-9.

11. García LM, Artero JM, Zamorano MB, Tejero IG. Fluorescence-guided resection with 5-aminolevulinic Acid of subependymomas of the fourth ventricle: Report of 2 cases: Technical case report. Neurosurgery 2015;11 Suppl 2:E364-71.

12. Diem LA, Rauhman BA, de Lima TO, Padrosa HA, Souweidane MM. Neuroendoscopic biopsy of brain lesions: Accuracy and complications. J Neurosurg 2015;122:34-9.

13. Hayashi Y, Nakada M, Tanaka S, Uchiyama N, Hayashi Y, Kita D, et al. Implication of 5-aminolevulinic acid fluorescence of the ventricular wall for postoperative communicating hydrocephalus associated with cerebrospinal fluid dissemination in patients with glioblastoma multiforme: A report of 7 cases. J Neurosurg 2010;112:1015-9.

14. Inoue T, Endo T, Nagamatsu K, Watanabe M, Tominaga T, et al. Intraventricular 5-aminolevulinic acid fluorescence-guided resection of intramedullary ependymoma: Report of 9 cases. Neurosurgery 2013;72:159-68.

15. Kamp MA, Grosser P, Felsberg J, Slattery PJ, Steiger HJ, Reifenberger G, et al. 5-aminolevulinic acid (5-ALA)-induced fluorescence in intracerebral metastases: A retrospective study. Acta Neurochir (Wien) 2012;154:223-8.

16. Marbacher S, Klinger E, Schwyzer L, Fischer I, Nevzati E, Diepers M, et al. Use of fluorescence to guide resection or biopsy of primary brain tumors and brain metastases. Neurosurg Focus 2014;36:E10.

17. Micko A, Rapoport BI, Youngerman BE, Fong RP, Kosty J, Brunswick A, et al. Limited utility of 5-ALA optical fluorescence in endoscopic endonasal skull base surgery: A multicenter retrospective study. J Neurosurg 2020;30:1-7.

18. Millesi M, Kiesel B, Mischkulnig M, Martinez-Moreno M, Wührer A, Wolfsberger S, et al. Analysis of the surgical benefits of 5-ALA-induced fluorescence in intracranial meningiomas: Experience in 204 meningiomas. J Neurosurg 2016;125:1408-19.

19. Moon JH, Kim SH, Shim JK, Roh TH, Sung KS, Lee JH, et al. Histopathological implications of ventricle wall 5-aminolevulinic acid-induced fluorescence in the absence of tumor involvement on magnetic resonance images. Oncol Rep 2016;36:837-44.

20. Rapp M, Kamp M, Steiger HJ, Sabel M. Endoscopic-assisted visualization of 5-aminolevulinic acid-induced fluorescence in malignant glioma surgery: A technical note. World Neurosurg 2014;82:e277-9.

21. Ritz R, Feigl GC, Schuhmann MU, Ehrhardt A, Danz S, Noell S, et al. Use of 5-ALA fluorescence guided endoscopic biopsy of a deep-seated primary malignant brain tumor. J Neurosurg 2011;114:1410-3.

22. Roth J, Constantini S. 5ALA in pediatric brain tumors is not routinely beneficial. Childs Nerv Syst 2017;33:787-92.

23. Stummer W, Pichlmueier U, Meinl T, Westler OD, Zanella F, Reulen HJ, et al. Fluorescence-guided surgery with 5-aminolevulinic acid for resection of malignant glioma: A randomised controlled multicentre phase III trial. Lancet Oncol 2006;7:392-401.

24. Stummer W, Rodrigues F, Schucht P, Preuss M, Wiewrodt D, Nestler U, et al. Predicting the “usefulness” of 5-ALA-derived tumor fluorescence for fluorescence-guided resections in pediatric brain tumors: A European survey. Acta Neurochir (Wien) 2014;156:2315-24.

25. Stummer W, Stocker S, Wagner S, Stepp H, Fritsch C, Goetz C, et al. Intraoperative detection of malignant gliomas by 5-aminolevulinic acid-induced porphyrin fluorescence. Neurosurgery 1998;42:518-25.

26. Takeda J, Nonaka M, Li Y, Komori Y, Kamei T, Iwata R, et al. 5-ALA fluorescence-guided endoscopic surgery for mixed germ cell tumors. J Neurooncol 2017;134:119-24.

27. Tamura Y, Kuroiwa T, Kamitani Y, Miki Y, Miyatake S, Tsuji M. Endoscopic identification and biopsy sampling of an intraventricular malignant glioma using a 5-aminolevulinic acid-induced protoporphyrin IX fluorescence imaging system. Technical note. J Neurosurg 2007;106:507-10.

28. Tokairin K, Kazumata K, Gotoh S, Sugiyama T, Kobayashi H. Neuroendoscope-assisted aneurysm trapping for ruptured intraventricular aneurysms in moyamoya disease patients. World Neurosurg 2020;141:278-83.

29. Tsuzuki S, Aihara Y, Eguchi S, Amano K, Kawamata T, Hashiba T, et al. Application of indocyanine green (ICG) fluorescence system lymphoma. Photochem Photobiol 2015;91:1452-7.

30. Yamanaka T, Ishikawa E, Miki S, Sakamoto N, Zaboronok A, Nishikawa H, et al. Limitation and complications of 5-aminolevulinic acid photodynamic diagnosis of deep-seated intracranial lesions: Experience in 204 meningiomas. J Neurosurg 2016;125:1408-19.