

Advanced Imaging for Biopsy Guidance in Primary Brain Tumors

Nelson Moussazadeh¹, Apostolos J. Tsiouris², Rohan Ramakrishna³

1. Neurological Surgery, NewYork-Presbyterian/Weill Cornell Medical Center 2. Radiology, Division of Neuroradiology, Weill Cornell Medical College, New York Presbyterian Hospital, New York 3. Department of Neurological Surgery, Weill Cornell Medical College

✉ **Corresponding author:** Nelson Moussazadeh, nem9026@nyp.org

Disclosures can be found in Additional Information at the end of the article

Abstract

Accurate glioma sampling is required for diagnosis and establishing eligibility for relevant clinical trials. MR-based perfusion and spectroscopy sequences supplement conventional MR in noninvasively predicting the areas of highest tumor grade for biopsy. We report the case of a patient with gliomatosis cerebri and multifocal patchy enhancement in whom the combination of advanced and conventional imaging attributes successfully guided a diagnostic biopsy.

Categories: Neurosurgery, Oncology, Radiology

Keywords: magnetic resonance imaging, glioma surgery, perfusion imaging, gliomatosis cerebri, magnetic resonance spectroscopy, image-guided therapy

Introduction

Brain tumors are imaged with magnetic resonance imaging (MRI), given its exquisite anatomic definition and diagnostic versatility. Many pathologies can be suggested on the basis of features seen on standard T1- and T2-weighted sequences alone, for example, diffuse intrinsic pontine glioma or juvenile pilocytic astrocytoma. Diffusion-weighted imaging demonstrates hypercellular and ischemic areas, with restricted Brownian motion correlating with a higher grade in gliomas [1]. Contrast enhancement indicates areas of high-grade disease with blood-brain barrier porosity. In focally enhancing and/or diffusion-restricting astrocytomas, a biopsy is targeted towards these areas as the highest histologic grade, in turn, guides treatment.

Supplemental MR perfusion imaging aids in tumor diagnosis, adding three to four minutes to the scanning time. In primary brain tumors, dynamic susceptibility contrast-enhanced relative cerebral blood volume (DSC rCBV) and dynamic contrast-enhanced plasma volume (DCE V_p) have been used as tumor neovascularity surrogates. MR spectroscopy adds five to 10 minutes and noninvasively quantitates brain metabolites. N-acetyl aspartate (NAA; a neuron-specific molecule), choline (Cho; a cell membrane component reflective of cellularity), creatine (a metabolic surrogate), and lactate/lipid quantification (indicative of necrosis) improve upon standard diffusion and contrast mapping in predicting areas of low versus high (WHO II vs. III) histologic grade in astrocytoma [2-3]. In some cases, for example, where decreased choline differentiates contrast-enhancing radiation necrosis from the active tumor, spectroscopy can spare patients invasive sampling [4].

Case Presentation

History and examination

How to cite this article

Moussazadeh N, Tsiouris A J, Ramakrishna R (February 22, 2016) Advanced Imaging for Biopsy Guidance in Primary Brain Tumors. Cureus 8(2): e504. DOI 10.7759/cureus.504

Received 11/07/2015

Review began 11/11/2015

Review ended 02/15/2016

Published 02/22/2016

© Copyright 2016

Moussazadeh et al. This is an open access article distributed under the terms of the Creative Commons Attribution License CC-BY 3.0., which permits unrestricted use, distribution, and reproduction in any medium, provided the original author and source are credited.

A 57-year-old right-handed woman with hypertension and tobacco use presented with progressive word-finding difficulty. Neurologic examination was notable for mild anomic dysphasia, truncal ataxia upon tandem gait testing, and right-sided dysrhythmokinesia. She also reported nighttime headaches and memory loss. She was immunocompetent, afebrile, and had no clinical evidence of active infection or primary malignancy.

Imaging attributes

Non-contrast head CT revealed bilateral poorly defined intra-axial cerebral hemispheric lesions with vasogenic edema and scattered internal microcalcifications.

Gadolinium-enhanced brain MRI demonstrated predominately non-enhancing mass-like T2-hyperintense infiltrating lesions involving the left superior temporal gyrus, extending through the posterior left temporal and parietal lobes across the splenium of the corpus callosum into the right peritrial white matter, with additional noncontiguous signal abnormality in the left thalamus, corona radiata, and cortical regions in the left parietal and occipital lobes. In addition to the posterior left superior temporal gyrus, which was deemed inoperable, three other areas were diffusion-restricting: the medial left parietal lobe, left corona radiata and splenium of the corpus callosum, and the right peritrial white matter.

MR perfusion (MRP), spectroscopy (MRS), and post-contrast T1 imaging findings are presented in Figure 1.

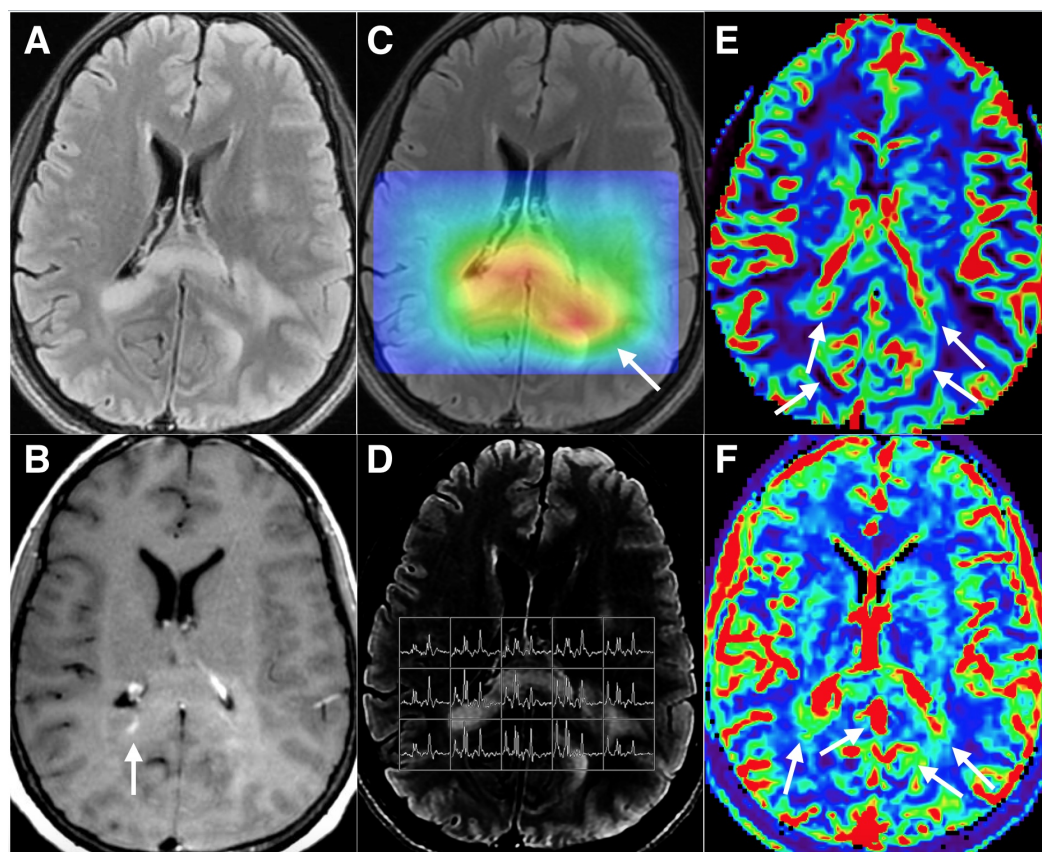


FIGURE 1: Advanced MR Characteristics

A) Axial T2W-FLAIR demonstrates infiltrative disease in the right peritrial white matter, the bilateral forceps major, and deep left parietal lobe. B) Post-contrast T1W demonstrates a focus of enhancement in the right forceps major (arrow). C) Elevated choline (Cho):N-acetyl aspartate (NAA) ratios correspond well to the areas of T2 signal abnormality and represent neoplastic infiltration, best demonstrated on a color "heat map" generated from the multivoxel MR

spectroscopy (GE AW READY View). D) Compared to the normal spectra (far right lateral voxels) with matched Cho and creatine (Cr) levels and high NAA levels, the infiltrated brain regions have a reversal of this pattern with elevated Cho and decreased NAA. E) The bilateral forceps major and splenium demonstrate multifocally increased cerebral blood volume (rCBV) on dynamic susceptibility contrast-enhanced (DSC) MRP (arrows). F) These same areas demonstrate increased plasma volume (V_p) on dynamic contrast-enhanced (DCE) MRP (arrows). The left forceps major was chosen as the sampling site, despite not being avidly contrast-enhancing, as it was the area of highest Cho:NAA ratio, rCBV, and V_p .

Informed patient consent was obtained for this patient's treatment. No reference to the patient's identity was disclosed in the figure displayed in this paper.

Discussion

This patient's gliomatosis cerebri is unresectable, and the primary concern is establishing an accurate diagnosis via a minimally-invasive biopsy. Anatomic MR establishes this as an intra-axial tumor, most likely a glioma, given its confinement to and expansion of normal sulcal boundaries and diffuse infiltration of multiple lobes. Extension through multiple vascular territories and discontinuous gray and white matter involvement exclude ischemia and demyelination while viral encephalitis can be excluded clinically. By definition, this constitutes Type I gliomatosis cerebri as it diffusely involves ≥ 3 cerebral lobes [5]. Importantly, cell type and histologic grade guide gliomatosis therapy. While microcalcifications suggest oligodendroglial lineage, this and the tumor grade cannot be definitively assessed noninvasively. In combination with an ability to rule out atypical primary CNS lymphoma and cryptogenic metastasis, the goal of a biopsy is to safely sample the area of the highest predicted grade.

Standard contrast-enhanced T1- and diffusion-weighted sequences are typically adequate for biopsy planning. In this patient, where tissue adequacy for molecular profiling is critical, the only diffusion-restricting and contrast-enhancing area, which was in the right peritrial white matter, was small and in an area where a needle biopsy might return a non-diagnostic sample. However, elevated perfusion V_p and spectroscopic choline corroborated that area as the most likely to harbor high-grade features.

Additional advanced imaging modalities include CT/MR-fused positron-emission tomography (PET), a noninvasive adjunct examination for brain lesion diagnosis. Lacking the resolution of anatomic or perfusion MRI, PET may be useful in distinguishing radiation necrosis from an active tumor or biopsy planning in multifocal metastatic cases with unknown primary and sufficient spatial separation. While standard FDG-PET may be falsely negative and nonspecific for brain tumors, other amino acid tracers, including ^{11}C -MET, have been shown to be more sensitive and specific in gliomatosis [6]. Diffusion tensor tractography may be informative in cases where normal eloquent fibers are suspected of traversing a lesion requiring biopsy.

This patient underwent navigation-assisted open biopsy with awake cortical language mapping to minimize the risk of postoperative dysphasia, given the proximity to Wernicke's area, and local sampling of the posterior left temporal lesion, which revealed anaplastic oligodendroglioma (WHO Grade III). Open biopsy was selected in this case over stereotactic biopsy given the small and deep target volume ($< 0.5 \text{ cc}^3$) and the patient's desire for thorough tissue sampling and genetic sequence analysis for potential future clinical trial eligibility. Fluorescence in situ hybridization demonstrated 1p/19q codeletion (favorable), and immunohistochemistry was negative for IDH1 R132H mutation (wild-type is a negative prognostic) [7-8]. On the third postoperative day, the patient was discharged home with stable

mild anomic dysphasia. At her outpatient visit three weeks later, she remained complication-free. However, breakthrough partial seizures required anticonvulsant titration, and she is planned for adjuvant whole brain radiation and procarbazine, lomustine, and vincristine therapy on the basis of RTOG 9402 and EORTC 26951 [9-10].

Conclusions

Advanced MR-based imaging techniques may aid in glioma surgical planning. In gliomatosis cerebri, which is characterized by infiltration of ≥ 3 brain lobes, a stereotactic biopsy (or open biopsy, as clinically indicated) should be directed at the areas of predicted highest grade for treatment stratification. Lesional enhancement signifies breakdown of the blood-brain barrier, which is seen in tumors, but can also be seen in active demyelination, infection, and radiation necrosis, among other causes. Diffusion restriction is present in regions of tumor hypercellularity as well as infarction or purulence. The degree of Cho elevation and NAA depression on MR spectroscopy as well as plasma volume on DCE MRP, correlate with glioma grade.

Additional Information

Disclosures

Human subjects: Consent was obtained by all participants in this study. **Conflicts of interest:** In compliance with the ICMJE uniform disclosure form, all authors declare the following:

Payment/services info: All authors have declared that no financial support was received from any organization for the submitted work. **Financial relationships:** All authors have declared

that they have no financial relationships at present or within the previous three years with any organizations that might have an interest in the submitted work. **Other relationships:** All

authors have declared that there are no other relationships or activities that could appear to have influenced the submitted work.

References

1. Bai X, Zhang Y, Liu Y, Han T, Liu L: Grading of supratentorial astrocytic tumors by using the difference of ADC value. *Neuroradiology*. 2011, 53:533-39. [10.1007/s00234-011-0846-2](https://doi.org/10.1007/s00234-011-0846-2)
2. Kitis O, Altay H, Calli C, Yuntun N, Akalin T, Yurtseven T: Minimum apparent diffusion coefficients in the evaluation of brain tumors. *Eur J Radiol*. 2005, 55:393-400. [10.1016/j.ejrad.2005.02.004](https://doi.org/10.1016/j.ejrad.2005.02.004)
3. Sibtain NA, Howe FA, Saunders DE: The clinical value of proton magnetic resonance spectroscopy in adult brain tumours. *Clin Radiol*. 2007, 62:109-19. [10.1016/j.crad.2006.09.012](https://doi.org/10.1016/j.crad.2006.09.012)
4. Hollingworth W, Medina LS, Lenkinski RE, Shibata DK, Bernal B, Zurakowski D, Comstock B, Jarvik JG: A systematic literature review of magnetic resonance spectroscopy for the characterization of brain tumors. *AJNR Am J Neuroradiol*. 2006, 27:1404-11.
5. Chen S, Tanaka S, Giannini C, Morris J, Yan ES, Buckner J, Lachance DH, Parney IF: Gliomatosis cerebri: clinical characteristics, management, and outcomes. *J Neurooncol*. 2013, 112:267-75. [10.1007/s11060-013-1058-x](https://doi.org/10.1007/s11060-013-1058-x)
6. Sato N, Inoue T, Tomiyoshi K, Aoki J, Oriuchi N, Takahashi A, Otani T, Kurihara H, Sasaki T, Endo K: Gliomatosis cerebri evaluated by 18Falpha-methyl tyrosine positron-emission tomography. *Neuroradiology*. 2003, 45:700-707. [10.1007/s00234-003-1057-2](https://doi.org/10.1007/s00234-003-1057-2)
7. Taillibert S, Chodkiewicz C, Laigle-Donadey F, Napolitano M, Cartalat-Carel S, Sanson M: Gliomatosis cerebri: a review of 296 cases from the ANOCEF database and the literature. *J Neurooncol*. 2006, 76:201-205. [10.1007/s11060-005-5263-0](https://doi.org/10.1007/s11060-005-5263-0)
8. Desestret V, Ciccarino P, Ducray F, Crinière E, Boisselier B, Labussière M, Polivka M, Idhah A, Kaloshi G, von Deimling A, Hoang-Xuan K, Delattre JY, Mokhtari K, Sanson M: Prognostic stratification of gliomatosis cerebri by IDH1 R132H and INA expression. *J Neurooncol*. 2011, 105:219-24. [10.1007/s11060-011-0587-4](https://doi.org/10.1007/s11060-011-0587-4)
9. Cairncross G, Wang M, Shaw E, Jenkins R, Brachman D, Buckner J, Fink K, Souhami L,

- Laperriere N, Curran W, Mehta M: Phase III trial of chemoradiotherapy for anaplastic oligodendroglioma: long-term results of RTOG 9402. *J Clin Oncol.* 2013, 31:337-43. [10.1200/JCO.2012.43.2674](https://doi.org/10.1200/JCO.2012.43.2674)
10. van den Bent MJ, Brandes AA, Taphoorn MJ, Kros JM, Kouwenhoven MC, Delattre JY, Bernsen HJ, Frenay M, Tijssen CC, Grisold W, Sipos L, Enting RH, French PJ, Dinjens WN, Vecht CJ, Allgeier A, Lacombe D, Gorlia T, Hoang-Xuan K: Adjuvant procarbazine, lomustine, and vincristine chemotherapy in newly diagnosed anaplastic oligodendroglioma: long-term follow-up of EORTC brain tumor group study 26951. *J Clin Oncol.* 2013, 31:344-50. [10.1200/JCO.2012.43.2229](https://doi.org/10.1200/JCO.2012.43.2229)