Title: Tumor Heterogeneity Imaging (THI): Initial Experience in the Evaluation of Brain Gliomas

Name: Gloria J. Guzman Perez-Carrillo[1], Qing Wang[2], Yong Wang[2], Tammie Benzinger[2]
1. University of Arizona, USA; 2. Washington University in St. Louis, USA

Abstract (300 word limit)

Purpose:
Brain tumors are typically heterogeneous, and may contain different grades of tumor cells, different types of tumor cells, edema and/or abnormal vascular structures. Anatomical imaging alone can be limited in the evaluation of tumor heterogeneity, especially in those tumors that demonstrate little to no enhancement. While there are physiologic MR tools available in daily clinical practice such as perfusion or diffusion, we wanted to develop a more powerful, sensitive sequence for the characterization of tumor heterogeneity. We propose that Tumor Heterogeneity Imaging (THI)[1] can provide quantitative distributions of different grades of tumor cells and capillary blood perfusion within the tumor in a single clinical imaging scan with more accuracy than previously reported with traditional diffusion techniques.[2]
Materials and Methods:
11 adult patients with known or suspected brain gliomas that were non-enhancing or had substantial non-enhancing regions (>50%) underwent simultaneous 3,4-dihydroxy-6-[18F]fluoro-L- phenylalanine (18F-FDOPA)[3] PET/MRI prior to planned standard-of-care surgical resection and/or stereotactic biopsy. Of these, 7 patients also underwent THI, a new diffusion MRI protocol, microstructure modeling, and inverse computation technique. The THI maps were then compared to the 18F-DOPA and coordinate-guided biopsy or surgical resection results. Perfusion maps extracted from THI were calculated.[4] ADC cut-offs for tumor grade based on the THI data were then determined and tumor grade maps created. (See Figure 1).
Results:
Grade 4 tumor ADC cutoff was 0.3-0.5 10−3 mm2/s. Grade 3 was 0.5-0.8 10−3 mm2/s. Grade 1 and 2 was 0.8-1.5 10−3 mm2/s. Table 1 summarizes the subject's demographic characteristics and well as the correlation between 18F-FDOPA and THI maps. We found that in 7/7 patients (100%) THI maps correlated with tumor grade on pathological evaluation. Interestingly, 18F-FDOPA was negative on subject S7, whereas THI correctly identified not only the tumor, but the tumor grade at the region of stereotactic biopsy sample.
Conclusion:
This preliminary study demonstrated the capability of a new diffusion MRI method, THI, to noninvasively characterize the structural heterogeneity in brain tumors, including various grades of tumor cells and capillary blood perfusion within the tumors, consistent with pathology assessment on biopsy tissues. Although our preliminary data suggest THI is a promising multi-parametric imaging technique to accurately measure cellularity and tumor grade, larger studies will be needed before definitive conclusions can be made about the role of this technique. This preliminary study also suggested the unmet need to develop new generation of MRI technique that is capable to provide direct pathophysiological measures for tumor characterization.

Image

Figure1. A 66-year-old women (S3) with recurrent oligodendroglioma, WHO grade II was imaged using FLAIR, FDOPA PET, perfusion and THI imaging. Region of interest was label in magenta.

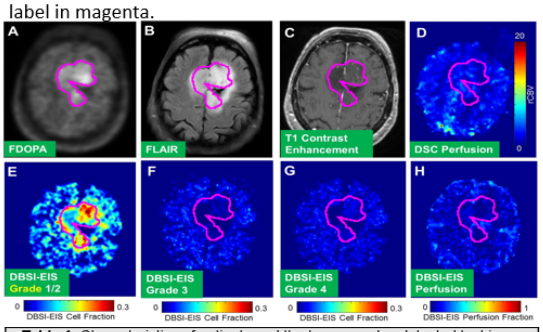
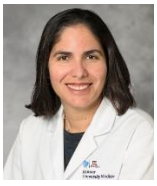


Table 1. Characteristics of patients and the tumor grades detected by biopsy and THI

Patient #	Age	Gender	FDOPA uptake	Tumor pathology	DBSI-EIS tumor grade
S3	66	Female	Yes	Oligoastrocytoma WHO grade II	Grade 1/2
S5	36	Female	Yes	Oligodendroglioma WHO grade II	Grade 1/2
S6	38	Male	Yes	Oligoastrocytoma WHO grade II	Grade 1/2
S7	30	Male	No	Diffuse astrocytoma WHO grade II	Grade 1/2
S8	33	Female	Yes	Oligodendroglioma WHO grade II	Grade 1/2
S9	48	Male	Yes	Oligodendroglioma WHO grade III	Grade 3
S11	62	Male	Yes	Glioblastoma WHO grade IV	Grade 4

References

1. Wang Y, Wang Q, Haldar JP, et al. Quantification of increased cellularity during inflammatory demyelination. Brain 2011; 134(Pt 12):3590-3601.
2. Sui Y, Xiong Y, Jiang J, et al. Differentiation of Low-and High-Grade Gliomas Using High b-Value Diffusion Imaging with a Non-Gaussian Diffusion Model. Am J Neuroradiol 2016; 37(9):1643-9.
3. Beuthien-Baumann B, Bredow J, Burchert W et al. 3-O-methyl-6-[18F] fluoro-L-DOPA and its evaluation in brain tumour imaging. Eur J Nucl Med Mol Imaging 2003; 30(7):1004-1008.
4. Le Bihan D, Breton E, Lallemand D, et al. MR imaging of intravoxel incoherent motions: application to diffusion and perfusion in neurologic disorders. Radiology 1986;161(2): 401-407.



Banner
University Medical Center
Tucson

Paste your Photograph Here

Dr. Guzman Perez-Carrillo is Assistant Professor of Radiology, Director of the Advanced Neuro-Imaging Initiative, Associate Residency Program Director for Research and Co-Chair of the University of Arizona Health Sciences LGBTQ+ Interest Group. She was born and raised in Puerto Rico. She completed her undergraduate studies at Johns Hopkins University in Baltimore, her medical degree at the University of Puerto Rico in San Juan, Puerto Rico, her radiology residency at West Virginia University in Morgantown, and her Neuroradiology Fellowship and Neuroradiology Research Fellowship at the Mallinckrodt Institute of Radiology at Washington University in St. Louis.

Dr. Guzman Perez-Carrillo also completed a Masters in Radiology at the University of Granada in Granada, Spain. Dr. Guzman has special research interests in advanced neuroimaging techniques, including functional imaging, diffusion spectrum based imaging, molecular imaging of tumor with F-DOPA in addition to outcomes and translational research of MRI in the field of Neuroradiology Imaging.

Email: gguzman@radiology.ariozona.edu

Notes/Comments: