# Author's Accepted Manuscript

Multimodal MRI Evaluation of Primary Brain Tumors

Daniel Treister, Sara Kingston, Kristina E Hoque MS MD PhD, Meng Law MD, Mark S. Shiroishi MD



www.elsevier.de/endend

PII: S0093-7754(14)00142-0

DOI: http://dx.doi.org/10.1053/j.seminoncol.2014.06.006

Reference: YSONC51721

To appear in: Semin Oncol

Cite this article as: Daniel Treister, Sara Kingston, Kristina E Hoque MS MD PhD, Meng Law MD, Mark S. Shiroishi MD, Multimodal MRI Evaluation of Primary Brain Tumors, *Semin Oncol*, http://dx.doi.org/10.1053/j.seminoncol.2014.06.006

This is a PDF file of an unedited manuscript that has been accepted for publication. As a service to our customers we are providing this early version of the manuscript. The manuscript will undergo copyediting, typesetting, and review of the resulting galley proof before it is published in its final citable form. Please note that during the production process errors may be discovered which could affect the content, and all legal disclaimers that apply to the journal pertain.

# **ACCEPTED MANUSCRIPT**

### **Multimodal MRI Evaluation of Primary Brain Tumors**

**Daniel Treister** 

Sara Kingston

Kristina E Hoque, MS MD PhD

Meng Law, MD

Mark S. Shiroishi, MD

From the Department of Radiology, Division of Neuroradiology, Keck School of Medicine, University of Southern California.

#### Introduction

Gliomas comprise 80% of primary brain neoplasms, with glioblastoma multiforme being the most commonly diagnosed glioma.<sup>1</sup> The annual incidence is 5.26 per 100,000, or 17,000 newly diagnosed cases per year in the United States.<sup>2</sup> The incidence increases with age, peaking between the 6<sup>th</sup> and 8<sup>th</sup> decades.<sup>2</sup> Gliomas are more common among Caucasians and occur more commonly in men.<sup>3</sup> They can be associated with certain rare hereditary syndromes including Cowden, Turcot, Li-Fraumeni, neurofibromatosis type 1 and type 2, tuberous sclerosis, and familial schwannomatosis.<sup>4, 5</sup> Known risk factors include a history of ionizing radiation, family history of glioma, and certain genetic susceptibility variants that are weakly associated with glioma.<sup>6, 7</sup> Preventative measures have not been shown to decrease the risk of later development.<sup>3</sup> In addition, screening tests are unwarranted since early diagnosis and treatment have not been shown to improve outcome.<sup>3</sup>

The malignant phenotype of GBM that lends to poor prognostic outcomes is due to its invasive and infiltrative abilities, which have been correlated with the degree of

## Download English Version:

# https://daneshyari.com/en/article/2161979

Download Persian Version:

https://daneshyari.com/article/2161979

<u>Daneshyari.com</u>