GBM, unlike melanoma and lung cancers, does not attract robust T cell immune responses, and, so far, immunotherapies have had little success against it. GBM is considered "immunologically cold" or unreactive likely due to tumor elaborated immune suppressive factors.

Findings from the study indicate FGL2 (fibrinogen-like protein 2), which is known for suppressing the immune system, is highly expressed in GBM. Investigators showed that inactivating or "knocking out" FGL2 from the tumor cells can eliminate tumor progression in mice with intact immune systems. Understanding this type of expression is key to discovering causes of GBM progression.

First author Jun Yan, Ph.D., a research scientist, showed the FGL2 present in tumor cells controls a specialized group of dendritic cells which activates T cells. More specifically, FGL2 secreted from tumor cells prevents the differentiation of a special subpopulation of CD103 dendritic cells that are essential for triggering the activation of the tumor killing T cells. The study also showed these dendritic cells must find a way to the tumor microenvironment in the central nervous center (CNS) in order to activate the T cells.

"This study is important because it shows that the immune system must interact in the CNS and in the tumor to be effective. Previously, this interaction was only thought to be necessary in specialized immune organs such as the lymph nodes," said Heimberger. "It also shows a new mechanism of immune suppression that hasn't been described before, and it further supports how important FGL2 is to this disease."

The team also analyzed human GBM from The Cancer Genome Atlas and found that lower levels of FGL2 protein expression coupled with high levels of GM-CSF or IFN, DC differentiation inducer or T cell activator, are associated with longer survival of GBM patients.
Li and Heimberger are actively working on therapeutic strategies to target FGL2.


Provided by University of Texas M. D. Anderson Cancer Center

*This document is subject to copyright. Apart from any fair dealing for the purpose of private study or research, no part may be reproduced without the written permission. The content is provided for information purposes only.*