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gcm2 Promotes Glial Cell Differentiation and Is Required with *glial cells missing* for Macrophage Development in *Drosophila*Teresa B Alfonso ... Bradley W Jones¹ **Show more**<https://doi.org/10.1006/dbio.2002.0740>[Get rights and content](#)Under an Elsevier [user license](#)[open archive](#)

Abstract

glial cells missing (*gcm*) is the primary regulator of glial cell fate in *Drosophila*. In addition, *gcm* has a role in the differentiation of the plasmatocyte/macrophage lineage of hemocytes. Since mutation of *gcm* causes only a decrease in plasmatocyte numbers without changing their ability to convert into macrophages, *gcm* cannot be the sole determinant of plasmatocyte/macrophage differentiation. We have characterized a *gcm* homolog, *gcm2*. *gcm2* is expressed at low levels in glial cells and hemocyte precursors. We show that *gcm2* has redundant functions with *gcm* and has a minor role promoting glial cell differentiation. More significant, like *gcm*, mutation of *gcm2* leads to reduced plasmatocyte numbers. A deletion removing both genes has allowed us to clarify the role of these redundant genes in plasmatocyte development. Animals deficient for both *gcm* and *gcm2* fail to express the macrophage receptor Croquemort. Plasmatocytes are reduced in number, but still express the early marker Peroxidase. These Peroxidase-expressing hemocytes fail to migrate to their normal locations and do not complete their conversion into macrophages. Our results suggest that both *gcm* and *gcm2* are required together for the proliferation of plasmatocyte precursors, the expression of Croquemort protein, and the ability of plasmatocytes to convert into macrophages.



Keywords

glial cells missing; *gcm*; *gcm2*; glia; hemocyte; plasmatocyte; macrophage; blood cells; *Drosophila*[Recommended articles](#) [Citing articles \(115\)](#)






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