



## Correction to: Matrine suppresses expression of adhesion molecules and chemokines as a mechanism underlying its therapeutic effect in CNS autoimmunity

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**Correction to: Immunologic Research (2013) 56:189–196**  
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In the originally published version of this article, the authors have noticed mistakes during figure preparation in the H&E-stained image in the MAT-M group of Fig. 2 and immunohistochemistry images of ICAM-1 in the MAT-treated groups of Fig. 3A. The correct Figs. 2 and 3A are displayed in the next page.

The authors confirm that all of the results and conclusions of the article remain unchanged. The authors sincerely apologize for this mistake.

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The original article can be found online at <https://doi.org/10.1007/s12026-013-8393-z>

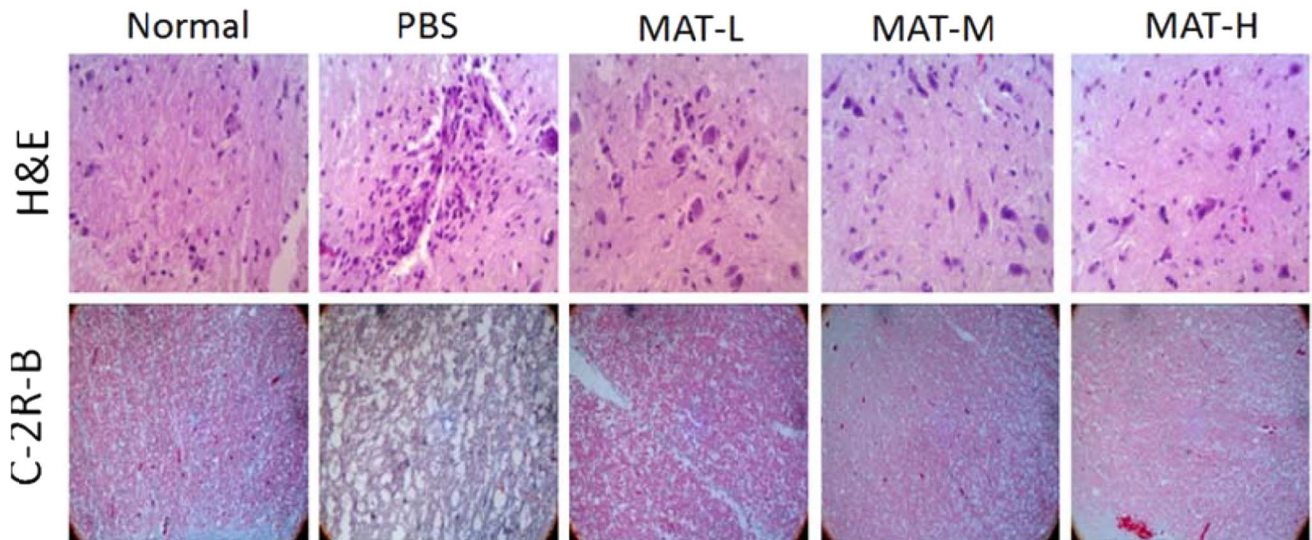
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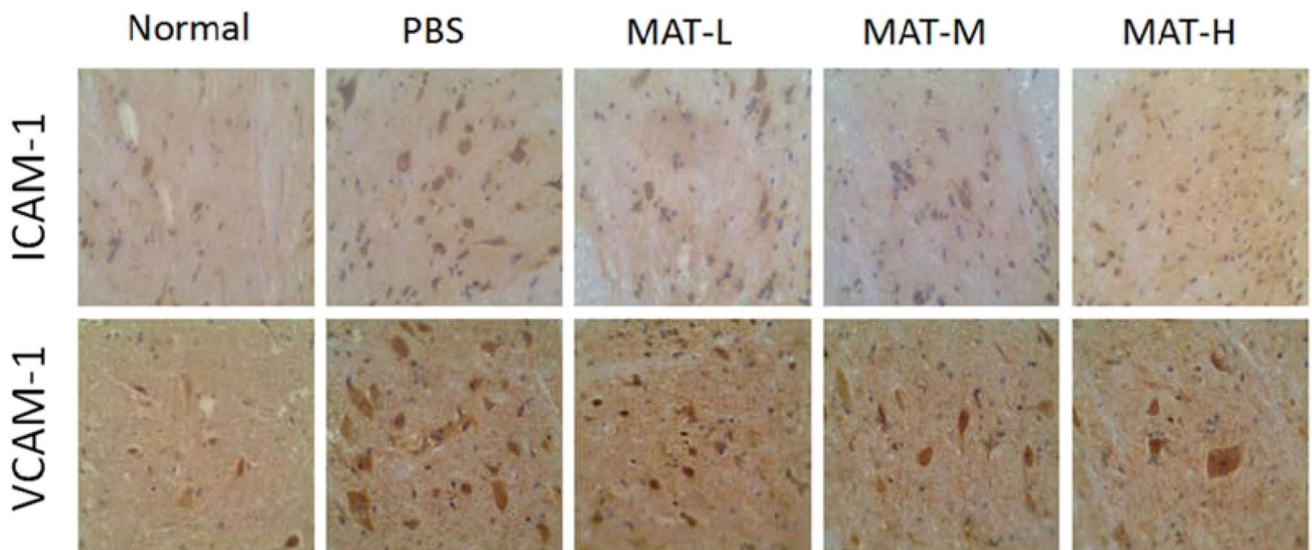
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**Fig. 2** CNS inflammatory infiltration and demyelination. On day 17 p.i., lumbar enlargement of the spinal cord was harvested after extensive perfusion and assayed by H&E and C-2R-B staining. The spinal cords from non-immunized animals serve as a normal control. The

PBS-injected group exhibited severe cellular infiltration and demyelination, while these pathological changes were significantly decreased in three MAT-treated groups. Magnification:  $\times 40$ . Quantitative analysis and statistics were summarized in Table 1 ( $n = 10$  each group)



**Fig. 3** Immunohistochemistry of ICAM-1 and VCAM-1. At day 17 p.i., the spinal cords were harvested from MAT-treated and untreated EAE rats, as well as from normal rats, and stained with anti-ICAM-1 and VCAM-1 antibodies (A). Magnification:  $\times 40$

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