## MYELIN BASIC PROTEIN-PRIMED T CELLS INDUCE NEUROTROPHINS IN GLIAL CELLS VIA αVβ3 INTEGRIN

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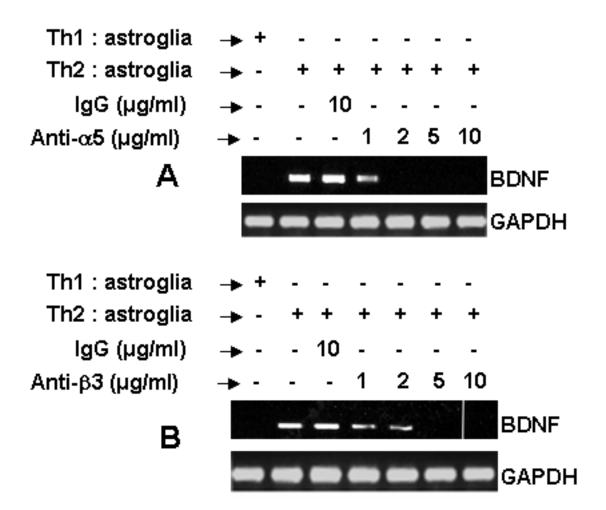
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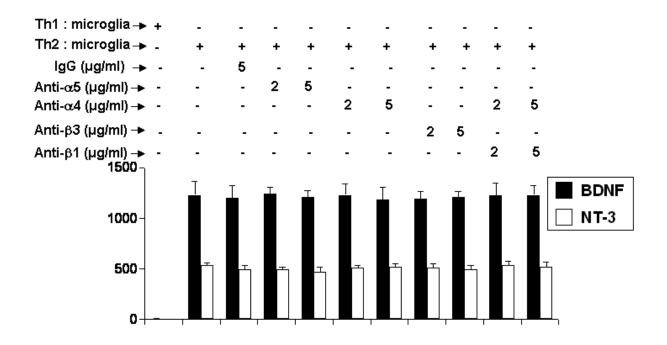
Supplementary Fig. 1. Role of  $\alpha V$  and  $\beta 3$  integrins in MBP-primed Th2 cell-induced expression of BDNF in mouse primary astroglia. (A) Th2 cells were treated with control IgG or different concentrations of antibodies against  $\alpha V$  (A) or  $\beta 3$  (B) as described above followed by addition to primary astroglia at a ratio of 0.5:1 T cell:glia. After 1 h of stimulation, T cells were removed followed by incubation of adherent astroglia in serum-free media. After 5 h of incubation (total), the expression of BDNF was analyzed by RT-PCR. Results represent three independent experiments.

Supplementary Fig. 2. Effect of neutralizing antibodies against  $\alpha V$ ,  $\alpha 4$ ,  $\beta 3$ , and  $\beta 1$  integrins on microglia. Primary microglia pre-incubated with control IgG and different concentrations of neutralizing antibodies against  $\alpha V$ ,  $\alpha 4$ ,  $\beta 3$ , or  $\beta 1$  integrins for 1h were washed three times to remove unbound antibodies. Then microglia were stimulated by gem-treated MBP-primed Th2 cells (0.5:1 T cell:glia). After 1 h of stimulation, T cells were removed followed by incubation of adherent microglia in serum-free media. After 24 h of incubation (total), supernatants were analyzed for BDNF and NT-3 by ELISA. Results represent mean  $\pm$  SD of three independent experiments.

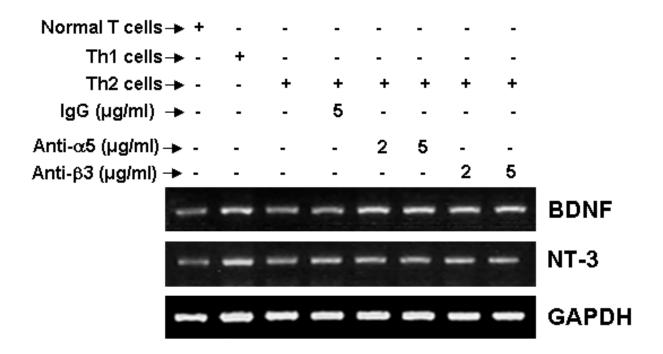
Supplementary Fig. 3. Effect of neutralizing antibodies against  $\alpha V$  and  $\beta 3$  integrins on the expression of BDNF in MBP-primed Th2 cells. Gem-treated MBP-primed Th2 cells were incubated with control IgG and different concentrations of neutralizing antibodies against either  $\alpha V$  or  $\beta 3$  for 6 h followed by analysis of BDNF mRNA by RT-PCR. Results represent three independent experiments.



## Supplementary Fig. 1



Supplementary Fig. 2



Supplementary Fig. 3