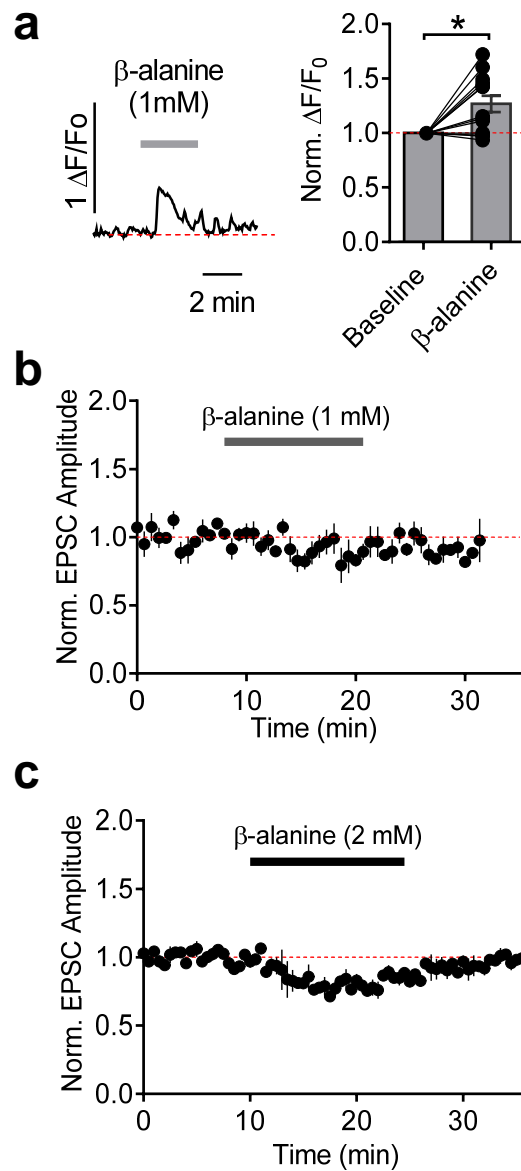
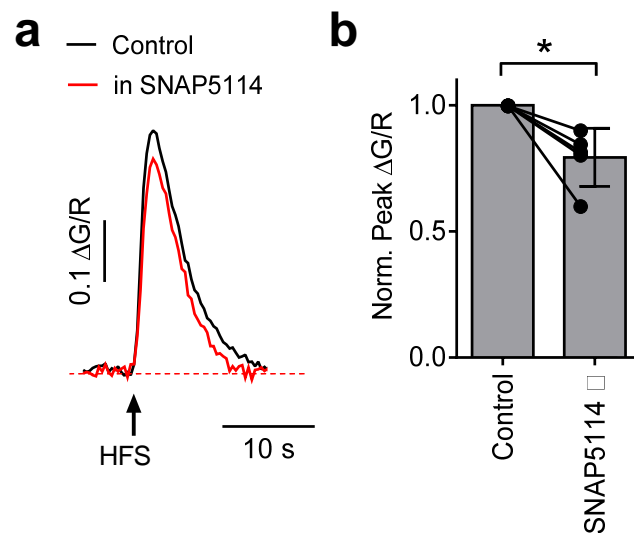


Supplementary Figure 1. Activation of metabotropic glutamate receptors induces Ca²⁺ increase in the presence of GAT-3 inhibitor. *Top:* representative trace showing change in astrocytic Ca²⁺ level following application of t-ACPD (50 μM) in the presence of SNAP5114. *Bottom:* Summary graph (n = 14 cells/2 animals); individual recordings represented by dots, means displayed by bars, error bars represent SEMs; * p = 0.003 (paired t-test compared to baseline).



Supplementary Figure 2. β -alanine action on astrocytic Ca^{2+} responses and EPSCs. **(a)** β -alanine (1 mM) induces Ca^{2+} response in astrocytes ($n = 13$ cells/2 animals). Note that Ca^{2+} increase is smaller than that induced by $30 \mu\text{M}$ GABA (c.f. Fig. 4e). **(b)** Normalized EPSC amplitudes in response to application of 1 mM β -alanine ($n = 6$ cells/2 animals). **(c)** Normalized EPSC amplitudes in response to application of 2 mM β -alanine ($n = 3$ cells/2 animals). Error bars represent SEMs, * $p = 0.004$ (paired t-test compared to baseline).



Supplementary Figure 3. Astrocytic Ca^{2+} responses to a one second, 100 Hz tetanus are reduced in the presence of the GAT-3 inhibitor SNAP5114 (100 μM). **(a)** Mean Ca^{2+} indicator fluorescence response ($n = 5$ cells/5 animals) to 100 Hz Schaffer collateral stimulation in control conditions (*black*) and in SNAP5114 (*red*). **(b)** Peak Ca^{2+} response amplitude for individual recordings (*dots*) and mean of five cells (*bars*). Error bars represent SEMs; * $p = 0.016$ (paired t-test).