ERRATUM



Erratum to: Prenatal nicotine exposure induces HPA axis-hypersensitivity in offspring rats via the intrauterine programming of up-regulation of hippocampal GAD67

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The authors of the above article would like to apologise for the mistakes which are present in Fig. 7b and in Fig. 7

legend. In Fig. 7b, the immunofluorescence pictures should be corrected as below. In Fig. 7 legend, the "(nt -358 to -77)" should be corrected as "(nt -1019 to -689)".

A corrected version of this figure and figure legend are as below:

The online version of the original article can be found under doi:10.1007/s00204-017-1996-8.



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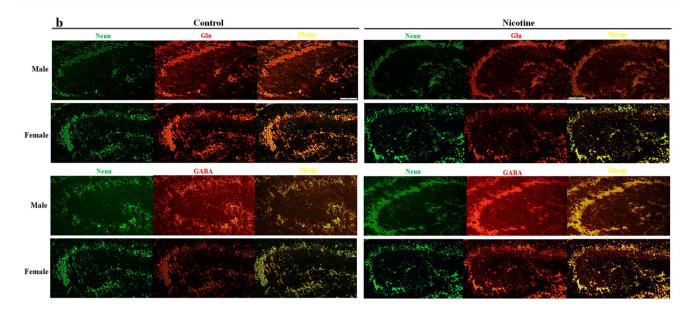


Fig. 7 Effects of prenatal nicotine exposure on the function of the hippocampus of fetal rats. Pregnant rats were subcutaneously administered 2.0 mg/kg day of nicotine from gestational day (GD) 9 to GD20, and then the fetal rats were extracted. The number of pregnant rats in each group was set to 15 (a litter size of 8 to 14 was considered qualified). The fetal hippocampus tissues were collected, and samples of each gender collected from each littermate were combined. The mRNA expression levels of glutamic acid decarboxylase 67 (GAD67), $\alpha 4$ and $\beta 2$ subtype of nicotinic acetylcholine receptor ($\alpha 4\beta 2nAChR$) and DNA methyltransferase 1 (Dnmt1) were detected by RT-qPCR (a,

d, n=15 per gender per group). Hippocampal neurons were detected by immunofluorescence staining, while the representative confocal laser-scanning microscopic images were double-stained for Glu (red, glutamatergic neuronal marker) or GAD67 (red, GABAergic neuronal marker) and NeuN (green, neuronal nuclei marker) (\mathbf{b} , n=4 per gender per group). Quantitative analysis for Glu-positive and GABA-positive cells in immunofluorescence was calculated (\mathbf{c}). The methylation status of GAD67 promotor (nt -1019 to -689) was detected by bisulfite-sequencing PCR method (\mathbf{d} , n=4 per gender per group). Mean \pm SD, *P < 0.05 compared with control

