



Corrigendum: A Novel RNA Editing Sensor Tool and a Specific Agonist Determine Neuronal Protein Expression of RNA-Edited Glycine Receptors and Identify a Genomic APOBEC1 Dimorphism as a New Genetic Risk Factor of Epilepsy

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Keywords: glycine receptors, epilepsy, temporal lobe, RNA editing, hippocampus, ligands

OPEN ACCESS

A Corrigendum on

Edited and reviewed by:

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Received: 26 March 2019 Accepted: 04 April 2019

Published: 24 April 2019 Citation:

Kankowski S, Förstera B, Winkelmann A, Knauff P, Wanker EE, You XA, Semtner M, Hetsch F and Meier JC (2019) Corrigendum: A Novel RNA Editing Sensor Tool and a Specific Agonist Determine Neuronal Protein Expression of RNA-Edited Glycine Receptors and Identify a Genomic APOBEC1 Dimorphism as a New Genetic Risk Factor of Epilepsy. Front. Mol. Neurosci. 12:103. doi: 10.3389/fnmol.2019.00103 A Novel RNA Editing Sensor Tool and a Specific Agonist Determine Neuronal Protein Expression of RNA-Edited Glycine Receptors and Identify a Genomic APOBEC1 Dimorphism as a New Genetic Risk Factor of Epilepsy

by Kankowski, S., Förstera, B., Winkelmann, A., Knauff, P., Wanker, E. E., You, X. A., et al. (2018). Front. Mol. Neurosci. 10:439. doi: 10.3389/fnmol.2017.00439

In the original article, there was an error. The incorrect oligonucleotide sequences were provided.

A correction has been made to the Methods, subsection PCR-RFLP Analysis of Human TLE Samples:

"Resected hippocampal tissue of human iTLE patients (Eichler et al., 2008) was analyzed with regard to *APOBEC1* gene dimorphism coding for 80M or 80I Apobec-1 protein variants. For this purpose, we developed a new PCR-based RFLP approach. Total RNA was isolated and reverse transcribed into cDNA as described earlier (Raltschev et al., 2016). Pre-amplification of Apobec-1 was performed using oligonucleotides 5'-CTTCAACCGGTGACCCCACTC-3' and 5'-TGCGTACAACATCATCCACAGAGG-3'. Then, 3.5 μ l of the pre-PCR were investigated in another PCR using oligonucleotides 5'-GAGTTTGACGTCTTCTATGACCC-3' and 5'-GTTGACAAAATTCCTCCAGCAG-3' to amplify a region spanning the 80M/I-coding position. This nested PCR amplification step yielded sufficient amount of DNA that was purified with Monarch[®] DNA Gel Extraction Kit (catalog no. L1020L, New England Biolabs GmbH) and digested using NlaIII restriction enzyme. NlaIII cuts at the <u>80M</u>-coding position (C<u>ATG</u>), and restriction fragments were separated using electrophoresis with 5% agarose gels to identify the

genotype of the iTLE patients. For control purpose, Apobec-1 80I- or 80M-coding vectors for transfection were processed in parallel. Ethidium bromide was used to stain DNA bands."

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The authors apologize for this error and state that this does not change the scientific conclusions of the article in any way. The original article has been updated.

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