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## The homeostatic effects of the RE-1 Silencing Transcription Factor on cortical networks are altered under pro-epileptic conditions in the mouse

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### SUPPLEMENTARY MATERIALS





Bar plots of the mean (± SEM) membrane capacitance (*left*) and resting membrane potential (*right*) with superimposed individual experimental points. p>0.05, two-tailed unpaired Student's *t*-test ( $\Delta$ Cre n = 8, Cre n = 12).











Figure S3. REST knockout does not affect the kinetics of spontaneous and miniature IPSCs. (A) sIPSC kinetics. *Top:* Representative traces of sIPSCs recorded in ΔCre- (red) and Cre- (blue) transduced mice. Bottom: Bar plots of the mean ± SEM 80% decay, 10-90 rise and half-width of sIPSCs with superimposed individual experimental points. (B) mIPSC kinetics. Top: Representative traces of mIPSCs recorded in  $\Delta$ Cre- (red) and Cre- (blue) transduced mice. Bottom: Bar plots of the mean ± SEM 80% decay, 10-90 rise and half-width of mIPSCs with superimposed individual experimental points). p>0.05, two-tailed unpaired Student's t-test (sIPSCs and mIPSCs: n=8 and n=9 for  $\Delta$ Cre and Cre, respectively).

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**(A)** 



#### Figure S4. Increased excitation/inhibition (E/I) ratio in *Rest*-KO pyramidal neurons.

The E/I ratio was assessed by the analysis of the frequency of sEPSCs (*left*) and mEPSCs (*right*) of individual *Rest*-KO pyramidal neurons normalized over the average frequency of sIPSCs and mIPSCs, respectively. \* p<0.05, two-tailed unpaired Student's *t*-test (sPSCs: n=12; mPSCs: n=10 and n=9, for  $\Delta$ Cre and Cre, respectively).



# Figure S5. Administration of a tandem dose of PTZ in *Rest*-KO mice does not affect membrane capacitance and resting membrane potential in layer II/III pyramidal neurons of V1 cortex.

Bar plots of the mean (± SEM) membrane capacitance (*left*) and resting membrane potential (*right*) with superimposed individual experimental points. p>0.05, two-tailed unpaired Student's *t*-test (n=7 for both PTZ Cre and  $\Delta$ Cre).



## Figure S6. *Rest*-KO V1 pyramidal neurons display similarly increased intrinsic excitability and action potential firing after the administration of a tandem dose of PTZ.

(A) Number of elicited action potentials *versus* injected current in V1 pyramidal neurons from  $\Delta$ Cre- (red) and Cre- (blue) transduced *Rest*<sup>GTi</sup> mice. Points in the plot (means ± SEM) were fitted according to a logistic function. \* p<0.05 for genotype effect; two-way repeated measures ANOVA. (B-I) Bar plots showing rheobase (B), input resistance (C), rising slope (D), falling slope (E), AP peak (F), V threshold (G), width (H), and trough (I). Bar plots represent the means ± SEM with superimposed individual experimental points (n=6 for both  $\Delta$ Cre and Cre).



## Figure S7. Excitation/inhibition (E/I) ratio in *Rest*-KO pyramidal neurons after a tandem dose of PTZ.

The E/I ratio was assessed by the analysis of the frequency of sEPSCs (left) and mEPSCs (right) of individual *Rest*-KO pyramidal neurons normalized over the average frequency of sIPSCs and mIPSCs, respectively. p>0.05; two-tailed unpaired Student's *t*-test (sEPSCs and mEPSCs: n=7 for both PTZ Cre and  $\Delta$ Cre).