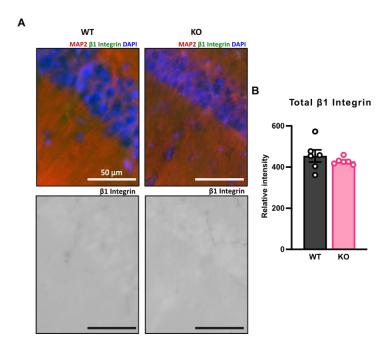
The Serine/threonine kinase NDR2 regulates integrin signalling, synapse formation and synaptic plasticity in the hippocampus. Supplementary figures.

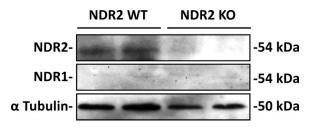
Miguel del Angel¹, Atsuhiro Tsutiya¹, Hussam Hayani², Deniz Madencioglu¹, Emre Kul¹, Gürsel Caliskan^{1,3}, Yunus Emre Demiray¹, Alexander Dityatev^{2,3,4}, Oliver Stork^{1,3,5,6}

- 1- Department of Genetics & Molecular Neurobiology, Institute of Biology, Otto-Von-Guericke University, Magdeburg, Germany.
- 2-Molecular Neuroplasticity Group, German Center for Neurodegenerative Diseases, 39120 Magdeburg, Germany.
- 3-Center for Behavioural Brain Sciences, Magdeburg, Germany.
- 4-Medical Faculty, Otto-Von-Guericke University, Magdeburg, Germany.
- 5-Center for Intervention and Research on adaptive and maladaptive brain Circuits underlying mental health (C-I-R-C), Jena-Magdeburg-Halle, Germany.
- 6- German Center for Mental Health (DZPG), Site Jena-Magdeburg-Halle, Germany.



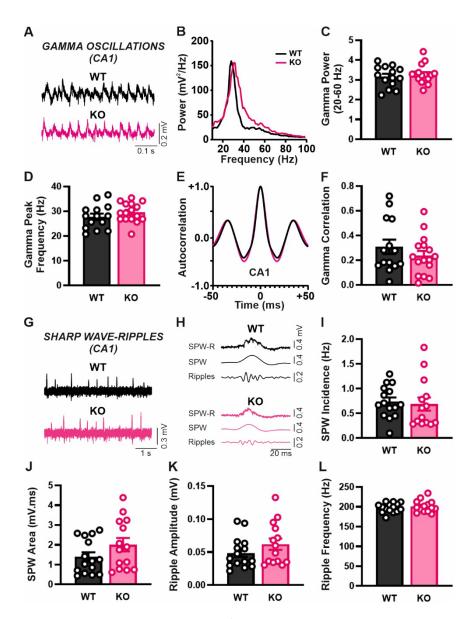
Supplementary Figure 1. Ndr2 deficient mice do not show alterations in total $\beta1$ integrin in the CA1

- (A) Representative image of dorsal hippocampus sections for WT and Ndr2 KO mice stained against total $\beta 1$ Integrin.
- (B) Quantification in the CA1 neurons shows no changes in total β 1 Integrin labelling *in vivo*, n= 6 mice each. All data are presented as mean \pm SEM. Student's t-test. *p<0.05; ****, p<0.0001.



Supplementary Figure 2. Ndr2 deficient mice do not show increase in NDR1 protein expression

Representative WB of dorsal hippocampus samples of 3-month-old Ndr2 wildtype (WT) & Knockout (KO) mice against NDR1 and NDR2 n=4 (WT); 4 (KO).



Supplementary Figure 3. Ndr2 deficient mice do not show any alterations in hippocampal network oscillations *ex vivo*.

- (A) Representative carbachol-induced gamma oscillation traces from the CA1 pyramidal layer of wildtype (WT) and knockout (KO) mice, n=14 slices from 3 mice (WT), 16 slices from 4 mice (KO).
- (B) Representative power spectra of CA1 gamma oscillations of WT and KO mice illustrating no changes between WT and KO mice.
- (C) Summary data illustrating no changes in the integrated gamma power (20-60 Hz) (T_{28} =0.6149, p=0.5436) between WT & KO mice.
- (D) Gamma peak frequency from the CA1 pyramidal neurons showing no difference between the WT and KO mice (T_{28} =1.115, p=0.2745).
- (E) Representative correlograms of the gamma peak frequency.
- (F) Summary data illustrating no changes in gamma correlation in the CA1 region of KO mice (T_{28} =1.073, p=0.2925).

Representative sharp wave-ripple (SPW-R) traces of WT and KO mice, n=15 slices from 4 mice (WT), 14 slices from 4 mice(KO).

- (G) Low pass-filtered SPW component of WT and KO mice, n=15 slices from 4 mice (WT), 14 slices from 4 mice(KO).
- (H) Band pass-filtered band-pass component of WT and KO mice, n=15 slices from 4 mice (WT), 14 slices from 4 mice(KO).
- (I) Summary data of Band pass-filtered band-pass component shows no significant alterations in SPW incidence (T_{27} =0.3204, p=0.7511) n=15 slices from 4 mice (WT), 14 slices from 4 mice(KO).
- (J) SPW area (T₂₇=1.537, p=0.1360)
- (K) Ripple amplitude (T_{27} =1.276, p=0.2128).
- (L) Ripple frequency (T_{27} =0.6739, p=0.5061).

Data are presented as mean ± SEM. Panels A, C, D, F, I, J, K and L: Student's two-tailed T-test.