



Supp. Fig. 10: Inhibition of AMPA or NMDA signalling results in decreased neuronal function. **A** Quantification of soma which show at least 1 characteristically neuronal calcium event ($n \geq 3$). Both genotype ($F_{2,19}=97.44$; $P < 0.0001$; $n \geq 3$ /group) and drug ($F_{2,19}=100.5$; $P < 0.05$; $n \geq 3$ /group) had significant effects on the percentage of active cells. There was also a significant interaction between the effect of genotype and drug ($F_{4,19}=10.16$; $P = 0.0001$; $n \geq 3$ /group) on the percentage of active cells. Data sets were analysed by two-way ANOVA with post hoc comparisons using Dunnett's multiple comparisons test comparing to control samples. Stars represent Dunnett-corrected post hoc tests. All data presented as means \pm SEM * $P < 0.05$; **** $P < 0.0001$ vs. untreated. **B** Example of an array-wide spike detection rate (ASDR) plot from 1q21.1 deletion cultures after 100 days of differentiation. **C** Example of an array-wide spike detection rate (ASDR) plot from 1q21.1 deletion cultures after 100 days of differentiation. The culture was incubated with AP5 immediately before recording. **D** Example of an array-wide spike detection rate (ASDR) plot from 1q21.1 deletion cultures after 100 days of differentiation. The culture was incubated with CNQX immediately before recording. **E** Representative raster plot of neuronal activity exhibited by control and duplication-derived neurons at early (D60) neurodevelopmental stage. **F** The average number of spikes recorded per electrode at D60 of differentiation in control and duplication-derived neurons, there was no significant difference between the spike counts in both models.