

S4 Fig. Improved signal ratio of edited Nano gene over Firefly gene upon transcription by P+L MeV polymerase with (a) raw data and (b) improved dynamic range. Data were obtained using dual-luciferase minigenome with Firefly and edited Nano gene separated by canonical N-P IGRs in the presence of wt, N, P and L or a truncated inactive L ( $L^{ko}$ ) as a negative control to measure the translation background from the minigenome transcribed by the T7 polymerase. The level of the background RLU signal is shown by the grey zone in (a)