



Supplementary Figure 3 – Behavioural data from the judgement bias task following acute treatment with phencyclidine.

Acute doses of PCP (0.0, 0.3, 1.0, 3.0 mg/kg) were administered by intraperitoneal injection to measure their effect on judgement bias. (A) PCP caused responses latencies to become slower across all three tones for the highest dose (3.0 mg/kg) only (significant session*tone interaction: $F_{4,49,67.41}=3.079$, $p=0.018$ and significant post-hoc tests: $p_s \leq 0.019$). (B) PCP did not alter percentage of positive responses. (C) The highest dose of PCP (3.0 mg/kg) caused rats to omit more responses for the midpoint and low reward tones (significant session*tone interaction: $F_{2,76,41.44}=4.101$; $p=0.014$ and significant post-hoc tests: $p_s \leq 0.002$). (D) 1.0 mg/kg PCP caused an increase in percentage of premature responses (main effect of session: $F_{3,45}=2.880$, $p=0.046$ and significant pairwise comparison against vehicle: $p=0.050$). Data represent mean \pm SEM. $n=16$, 40 min pre-treatment. *** $p < 0.001$, ** $p < 0.01$, * $p < 0.05$. HT - high reward tone; MT - midpoint tone; LT - low reward tone.