Figure S1. *In vitro* HAT assays using recombinant histone H3 and either purified \( \text{ddHA} \text{GCN5b} \) or \( \text{ddHA} \text{GCN5b(E703G)} \). Proteins were purified from parasites cultured in the presence of 500 nM Shield for 48 hours. KAT reactions were analyzed by Western blotting with antibody recognizing acetylated H3. Parental strain was used as a negative control. Anti-HA was used to show that the same approximate amount of protein was used in each assay.