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Supporting information for article:

Crystal structure of a GCN5-related *N*-acetyltransferase from *Lactobacillus curiae*

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Figure S1 Representative electron density. $(2mF_{obs}-DF_{calc})\alpha_{calc}$ electron density map contoured at 1σ for the region local to the catalytically relevant residue Y97.

A

В





Figure S2 Size exclusion chromatogram of LcGNAT. Size exclusion was performed on a Superdex S75 16/60 column (GE Healthcare) pre-equilibrated in 50 mM Tris pH 8.0, 100 mM NaCl. The loaded sample (volume approx. 3ml) had been concentrated to 6 mg/ml; **B.** The exclusion volume (V_e) of *Lc*GNAT (orange), interpreted in the light of column calibration standards (blue), suggests a monomeric form of the protein (MM= 20.25 kDa; calculated from sequence data). For each protein, exclusion volume and molecular mass are quoted in brackets.



Figure S3 Topological differences in GNAT enzymes. (A) Canonical topology of the GNAT superfamily (B) and *Lc*GNAT topology representative of the subfamily of polyamine acetyltransferases. In the latter, β -strand β 0 is not present. Instead, an extra C-terminal β -strand β 7 (dark blue) forms an antiparallel interaction with β -strand β 1.