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Supporting Information

A Thermodynamic Perspective on Potential G-Quadruplex Structures as Silencer Elements in the MYC Promoter

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Author Contributions

J.J. Formal analysis: Equal; Investigation: Lead; Methodology: Equal; Validation: Equal; Writing - Original Draft: Equal; Writing - Review & Editing: Equal

K.W. Conceptualization: Lead; Formal analysis: Equal; Funding acquisition: Lead; Methodology: Equal; Resources: Lead; Supervision: Lead; Validation: Equal; Writing - Original Draft: Equal; Writing - Review & Editing: Equal.

Supporting Information

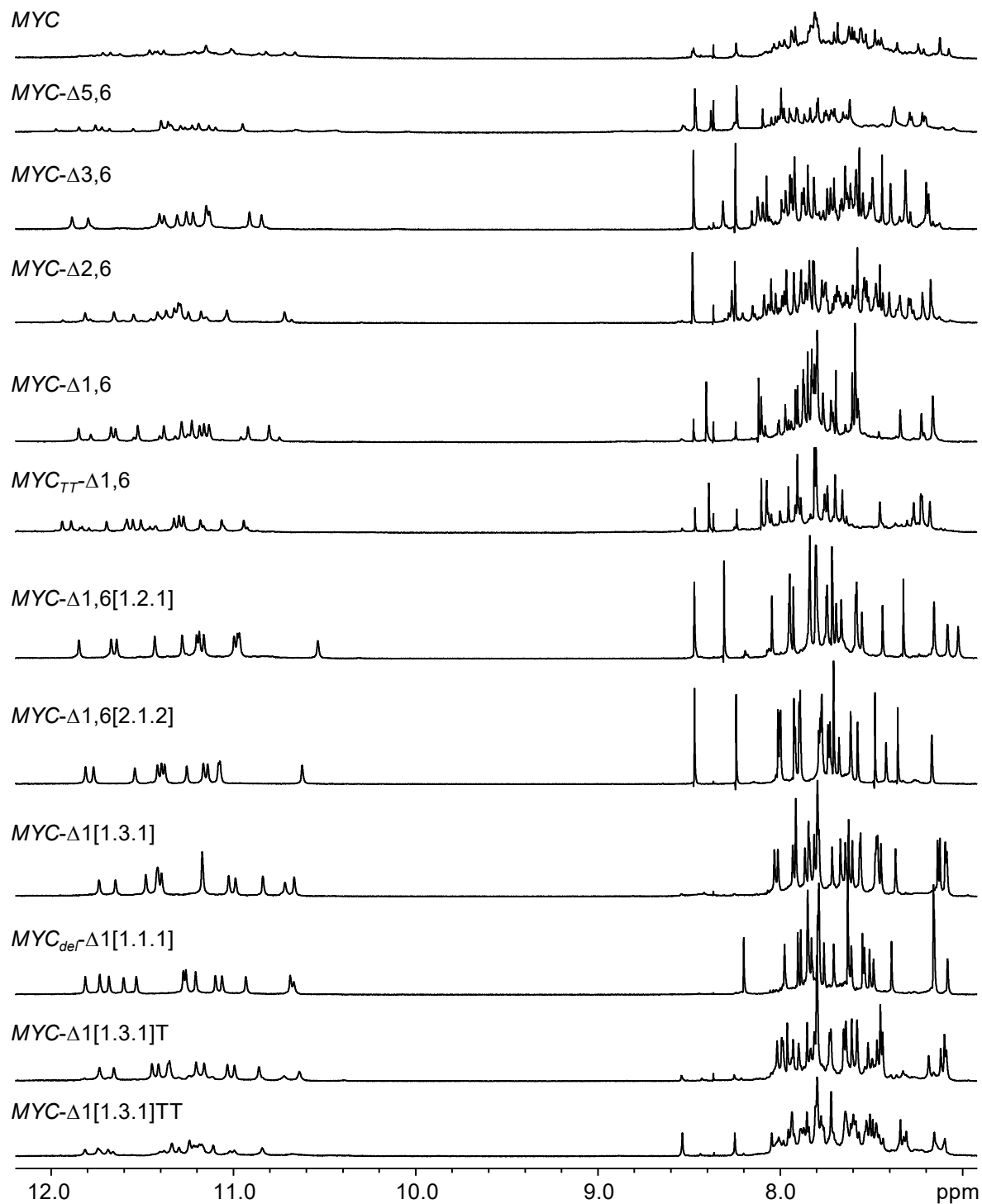


Figure S1. Imino and aromatic proton NMR spectral region of *MYC* sequence variants (0.2 mM) at 298 K in 10 mM potassium phosphate buffer, pH 7.

Gel electrophoresis

For non-denaturing polyacrylamide gel electrophoresis, annealed oligonucleotide samples (200 pmol in 10 μ L per lane) were loaded onto a 15% polyacrylamide gel (acrylamide:bis-acrylamide 19:1). Separation was performed at room temperature in TBE buffer supplemented with 10mM KCl using a voltage of 80 V. Bands were visualized by staining with a 50 μ M thiazole orange solution.

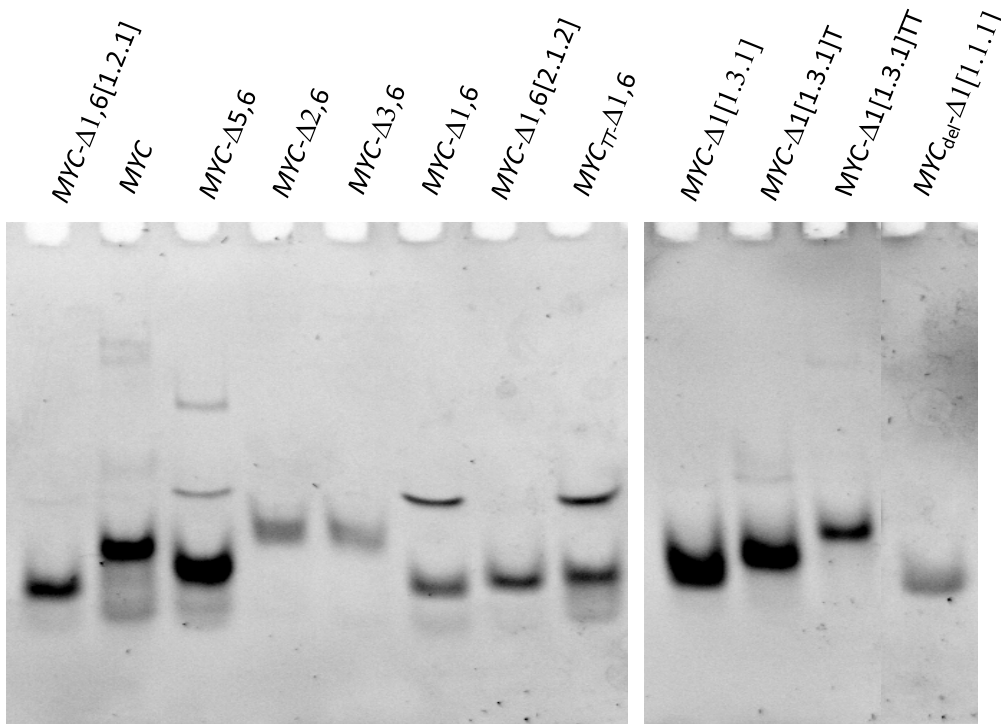


Figure S2. Non-denaturing polyacrylamide gel electrophoresis of *MYC* sequences after annealing in 10 mM potassium phosphate buffer, pH 7.

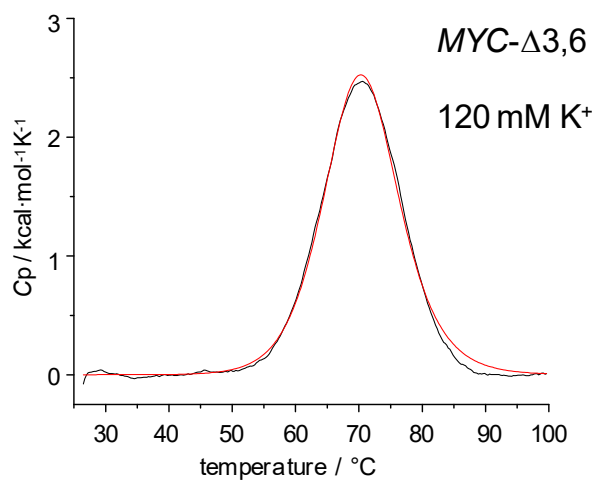


Figure S3. DSC thermogram for melting of *MYC*- Δ 3,6 in 20 mM potassium phosphate, 100 mM KCl, pH 7. Fitted curve based on a non-two-state model with $\Delta H_{\text{cal}}^{\circ} \neq \Delta H_{\text{vH}}^{\circ}$ is shown in red.

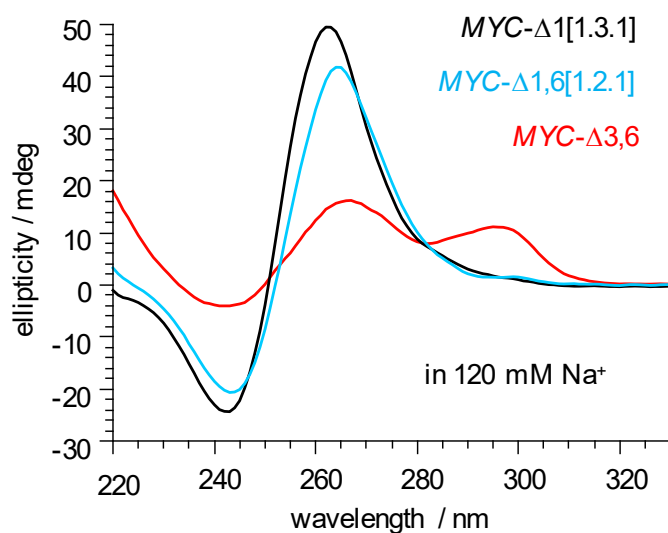


Figure S4. CD spectra of *MYC* sequence variants (5 μ M) at 293 K in 20 mM sodium phosphate, 100 mM NaCl, pH 7.