Bulky cations greatly increase the turnover of a native hammerhead ribozyme

Shu-ichi Nakano, Hirofumi Yamashita, Kazuya Tanabe, and Naoki Sugimoto

Department of Nanobiochemistry, Faculty of Frontiers of Innovative Research in Science and Technology (FIRST), Konan University, 7-1-20, Minatojima-minamimachi, Chuo-ku, Kobe, 650-0047, Japan

Frontier Institute for Biomolecular Engineering Research (FIBER), Konan University, 7-1-20, Minatojimaminamimachi, Chuo-ku, Kobe, 650-0047, Japan

Electronic supporting information

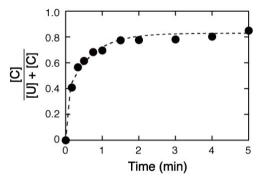
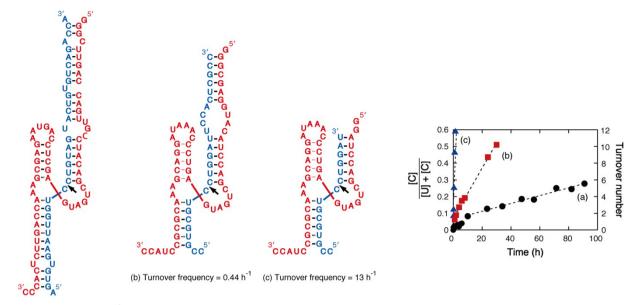


Fig. S1 The kinetic trace of the ribozyme-catalyzed substrate cleavage under the single-turnover condition conducted with 10 mM MgCl₂ at 37°C. The relative proportions of the populations that cleave fast and slowly are about 1:1.



(a) Turnover frequency = 0.050 h^{-1}

Fig. S2 The ribozyme-substrate complexes formed by HH10 and short ribozyme constructs derived from *Schistosoma mansoni*. Kinetic traces and the turnover frequencies at 10 mM MgCl₂ at 37°C are also presented.

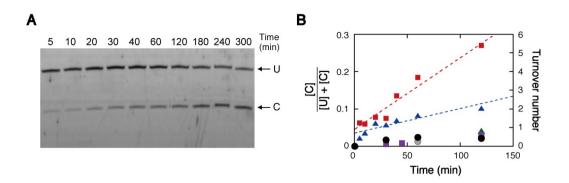


Fig. S3 (A) Polyacrylamide gel electrophoresis of the ribozyme-catalyzed substrate cleavage under multiple-turnover conditions in the presence of 100 mM TPeA. The upper and lower bands are uncleaved substrates and cleaved fragments, respectively. (B) Kinetic traces for substrate cleavage under multiple-turnover conditions in the absence (black circles) and presence of TMA (gray circles), TEA (green triangles), TPrA (purple squares), TBA (blue triangles), or TPeA (red squares) at 100 mM.