Selection and characterization of ssDNA aptamer against furaneol

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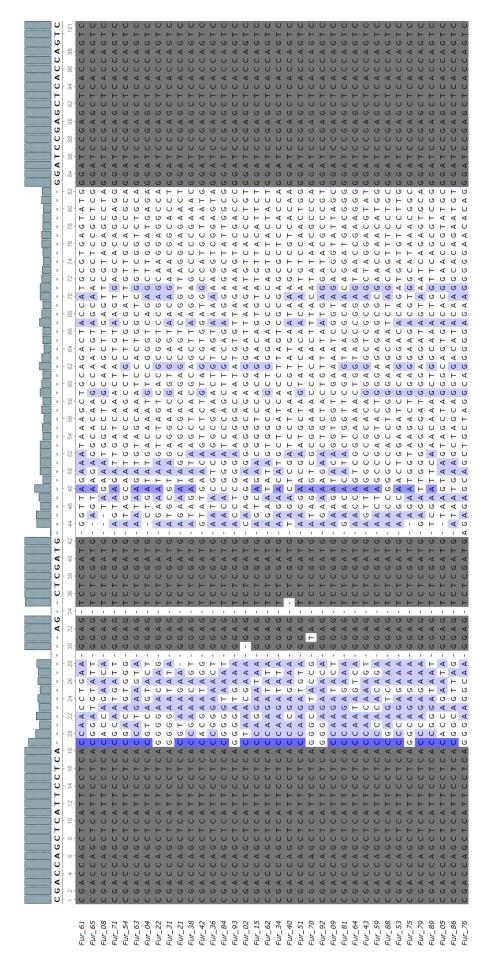


Figure S1. Alignment of the sequenced oligonucleotide pool. Constant regions of B1_bank library are highlighted in grey.

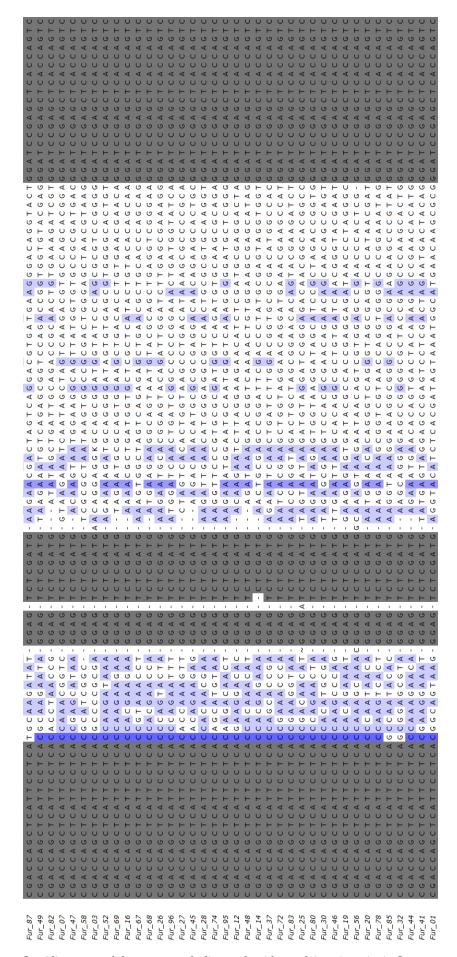


Figure S1. Alignment of the sequenced oligonucleotide pool (continuation). Constant regions of B1_bank library are highlighted in grey.

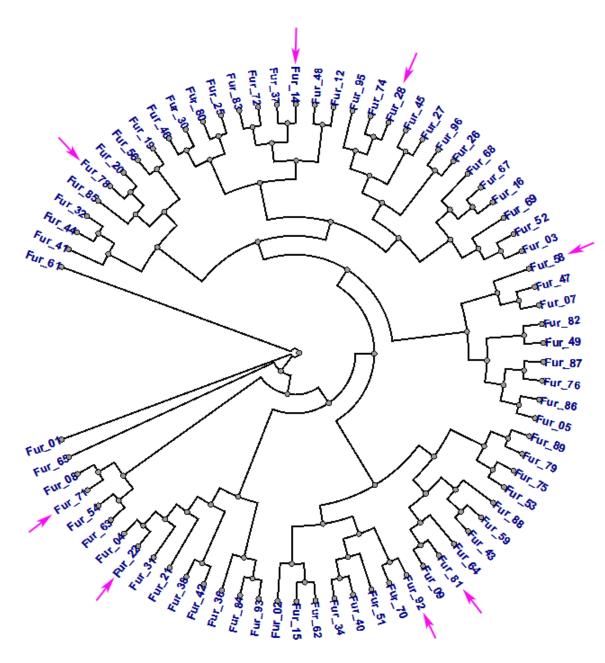


Figure S2. Phylogenetic tree of the sequenced pool of oligonucleotides.

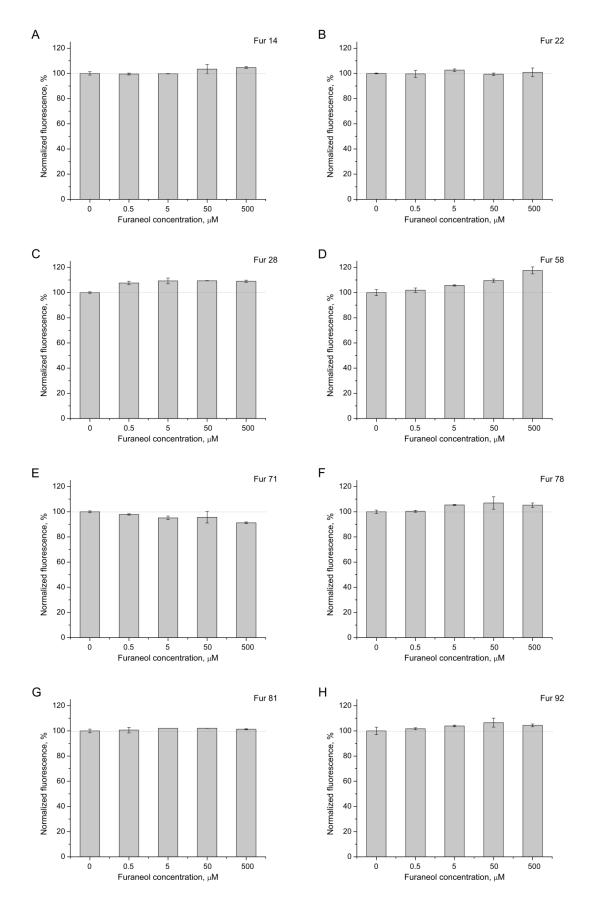


Figure S3. Results of elution-based binding assay for candidate aptamers.

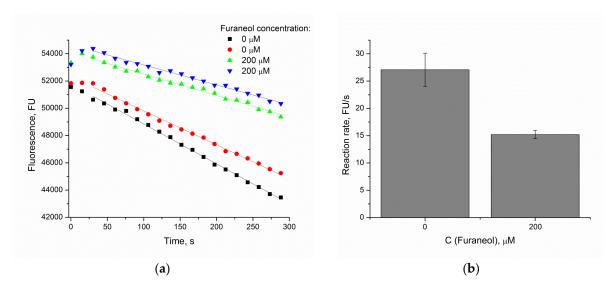


Figure S4. (a) Time dependence of fluorescent signal during digestion of Fur14 oligonucleotide by Exo I in the absence of furaneol and in the presence of 200 μ M furaneol; (b) Comparison of exonuclease reaction rate for Fur14 the absence of furaneol and in the presence of 200 μ M furaneol. Measurements were performed in duplicate, error bars were calculated as standard deviation. The difference between the bars proved to be statistically significant with p=0.05.