Supporting Information

Design and properties of ligand-conjugated guanine oligonucleotides for recovery of mutated G-quadruplexes

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Content

Analytical reverse-phased (RP) HPLC profiles and MALDI-TOF mass spectra for the ligandconjugated guanine tract oligonucleotides **PySG3** (S2), **G3PyS** (S3), **PyLG3** (S4), **G3PyL** (S5), **PEPyG3** (S6), **G3PEPy** (S7), **BPEAG3** (S8), **G3BPEA** (S9), **PerG3** (S10) and **G3Per** (S11). CD spectra of native VEGF G4 (S12). PAGE image of replication assay for native VEGF G4 (S13). $PySG_3 = 5'-XpGpGpGpTpT-3'$



RP-HPLC profile



MALDI-TOF MS



 $G_3PyS = 5$ '-TpTpGpGpGpX-3'



RP-HPLC profile



MALDI-TOF MS



PyLG₃ = 5'-**X**pGpGpGpTpT-3'





MALDI-TOF MS



 $G_3PyL = 5$ '-TpTpGpGpGpX-3'





MALDI-TOF MS



 $PEPyG_3 = 5'-XpGpGpGpTpT-3'$









 $G_3PEPy = 5$ '-TpTpGpGpGpX-3'









 $BPEAG_3 = 5'-XpGpGpGpTpT-3'$



RP-HPLC profile







 $G_3BPEA = 5$ '-TpTpGpGpGpX-3'



RP-HPLC profile









 $PerG_3 = 5'-XpGpGpGpTpT-3'$



RP-HPLC profile







 $G_3Per = 5$ '-TpTpGpGpGp**X**-3'











CD spectra of 10 μ M VEGF native G4. All the experiments were performed in a buffer consisting of 10 mM Tris- HCl (pH 7.5), 8 mM MgCl₂, and 50 mM KCl [1].

Time (min)	0	0.5	1 1	10	30	
Full-length product	-	-	-		18	
Stalled product —	_	-	-	-		
Primer —		•				

Denaturing-PAGE images of replication products from a native VEGF containing template in a buffer consisting of 10 mM Tris-HCl (pH 7.5), 8 mM MgCl₂, and 10 mM KCl with 1 μ M primer, 1 μ M template, 250 μ M dNTPs, and 1 μ M KF exo- at 37 °C [1].

Reference:

[1] Takahashi, S.; Kim, K. T.; Podbevsek, P.; Plavec, J.; Kim, B. H.; Sugimoto, N., Recovery of the Formation and Function of Oxidized G-Quadruplexes by a Pyrene-Modified Guanine Tract. *J. Am. Chem. Soc.* **2018**, 140, (17), 5774-5783.