

Supporting Information

Development of electrochemical sensors for a humanized antibody, bevacizumab, employing anti-idiotypic aptamers

Madoka Nagata^{1, 2}, Jinhee Lee¹, Taro Saito², Kazunori Ikebukuro², and Koji Sode^{1*}

Table S1 DNA Sequences against bevacizumab.

Name	Sequence (5' > 3')	Length
A14#1	GCGGTTGGTGGTAGTTACGTTTCGC	24
BivA14#1	GCGGTTGGTGGTAGTTACGTTTCGCTTTTTTTTTTTTTTTTTTTT TTTTTTTTTTTTTTTTTTTTTTTGGCGGTTGGTGGTAGTTACGTTTC G	92

Table S2 Composition of artificial serum solution.

Artificial serum composition	pH 7.44
NaCl	123 mM
KCl	3.5 mM
CaCl ₂	2.5 mM
MgCl ₂	0.7 mM
NaH ₂ PO ₄	1.5 mM
HEPES	10 mM
Bovine serum albumin	533 μ M

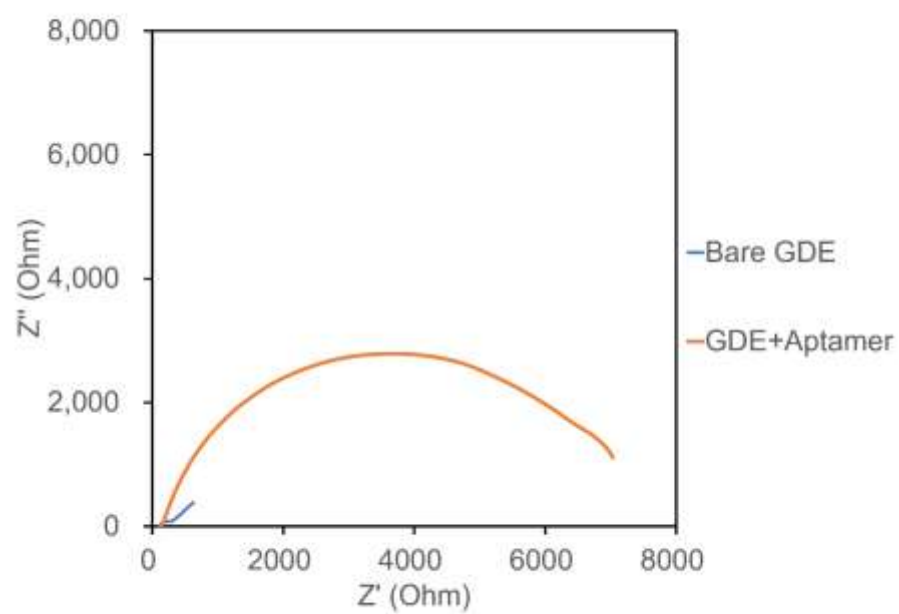


Figure S1 The Nyquist plots before and after the modification of aptamer

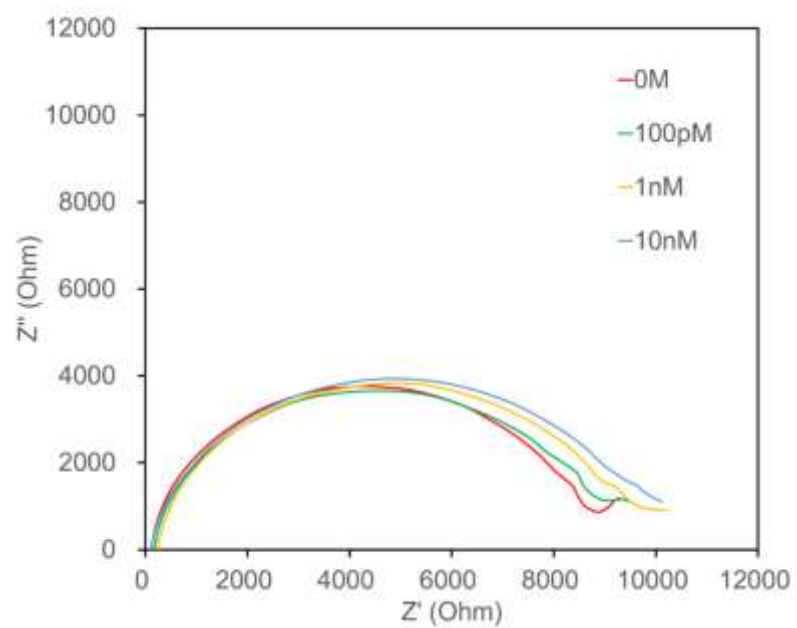


Figure S2 EIS monitoring of bevacizumab using anti-bevacizumab aptamer A14#1 immobilized electrode with the addition of human IgG kappa chain.

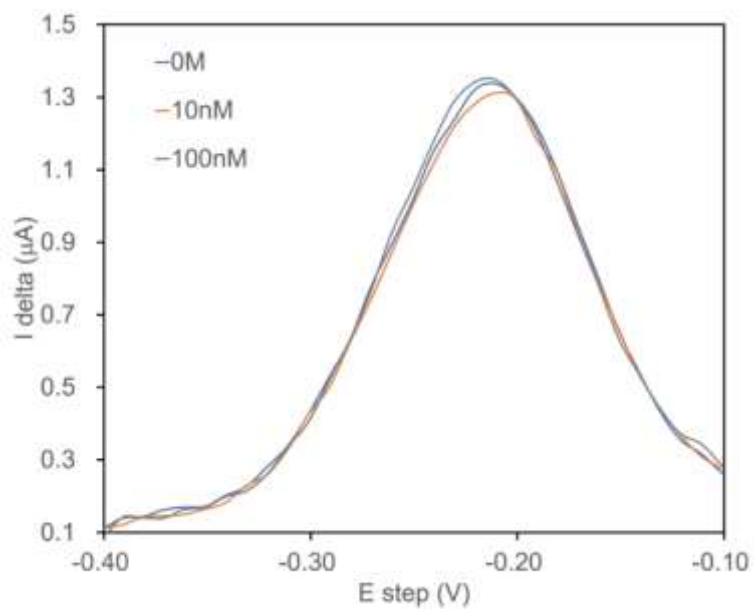


Figure S3 Square-wave voltammogram of PES-monovalent A14#1 aptamer immobilized sensor at 100Hz in PBS solution with the addition of several concentrations of bevacizumab. SWV scans were conducted as below conditions: -0.4 to -0.1 V, 25 mV amplitude.

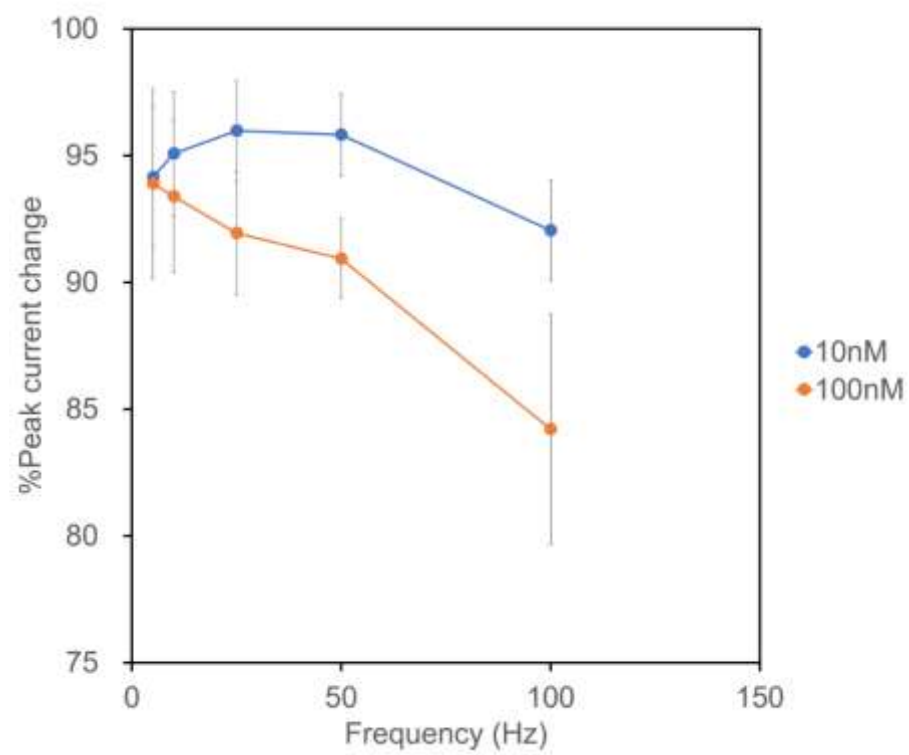


Figure S4 Peak current change dependencies on the frequency at each bevacizumab concentration. The error bars indicate the standard deviations (N=3).