



Supplementary Materials

Facilitation of Dye-Based Quantitative Real-Time Polymerase Chain Reaction with Poly(ethylene glycol)-Engrafted Graphene Oxide

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Table S1. Oligonucleotides used in the study.

Primer		Amplicon size (bp)	Sequence (5'-3')
Influenza B (IFZ B)	F	391	CTC CAT AGA GGT TCT TCA TTT GGG T
Influenza B (IFZ B)	R	391	ACA CCT TCT GCG AAA GCT TCA ATA C
Influenza A (IFZ A)	F	193	TTC AGA CAA TGG AAC GTG TTA CCC
Influenza A (IFZ A)	R	193	AGA AGC TTT TTG CTC CAG CAT GAG

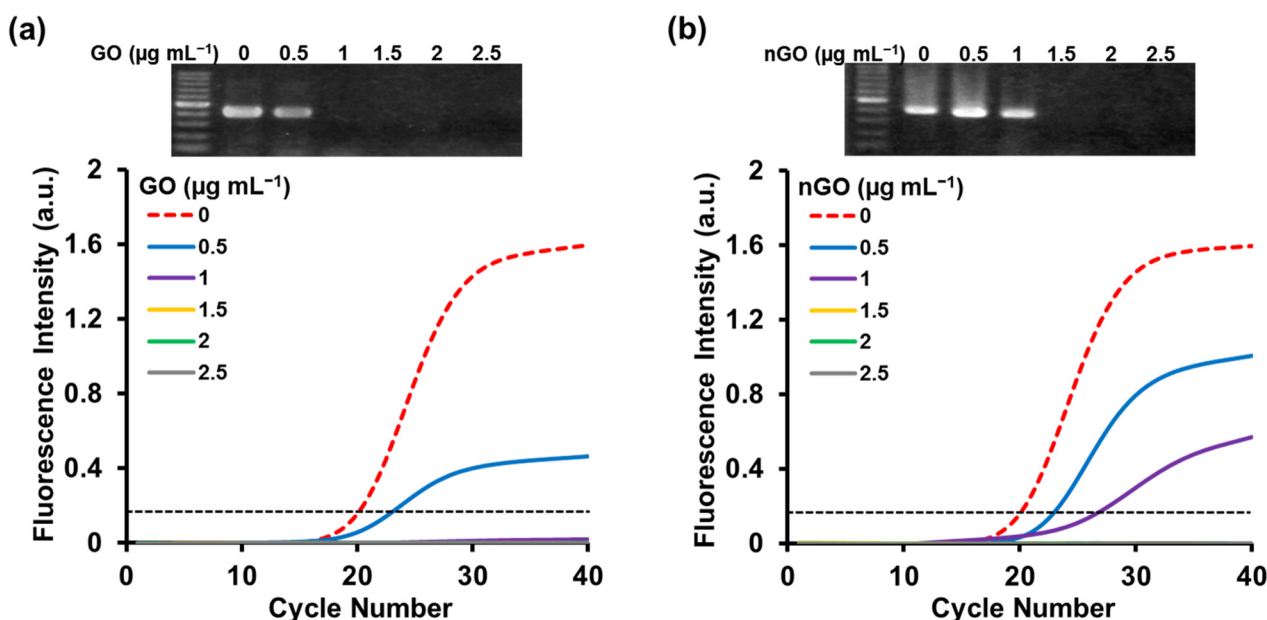


Figure S1. Real-time fluorescence monitoring of target RNA (IFZ B) amplification with GO and nGO. (a) Agarose gel electrophoretic analysis (top panel) and fluorescent amplification signals with various concentrations of GO (0, 0.5, 1, 1.5, 2, and 2.5 µg mL⁻¹). (b) Agarose gel electrophoretic analysis (top panel) and fluorescent amplification signals with various concentrations of nGO (0, 0.5, 1, 1.5, 2, and 2.5 µg mL⁻¹).

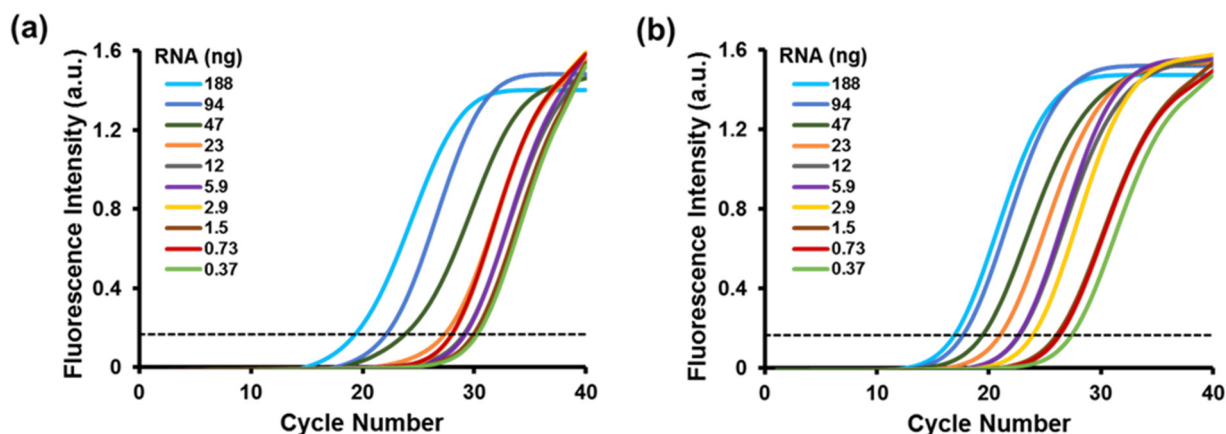


Figure S2. Real-time fluorescence monitoring of qPCR-amplified products with serially diluted (188 ng to 0.37 ng) target (IFZ B) RNA (a) in the absence of PEG-nGO and (b) in the presence of PEG-nGO.

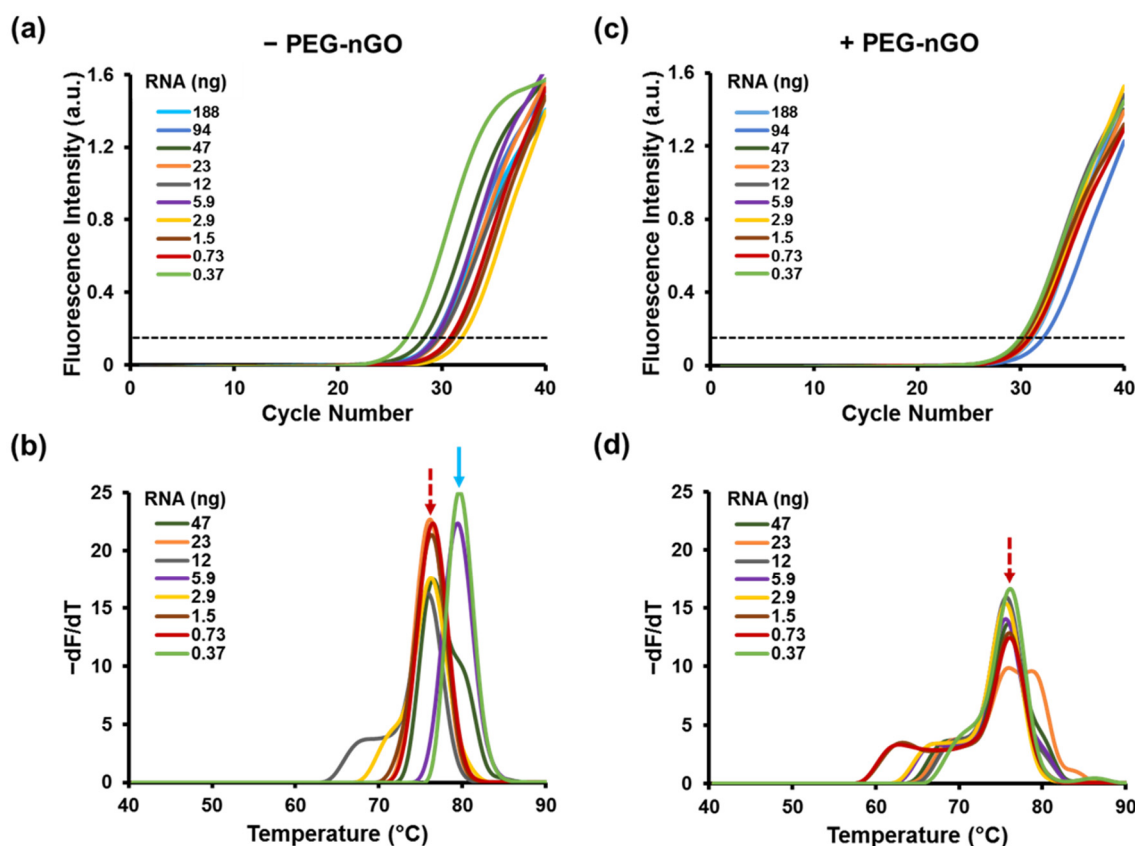


Figure S3. Analysis of qPCR-amplified products with non-target (IFZ A) RNA in the presence or absence of PEG-nGO. (a) Real-time fluorescence monitoring of qPCR-amplified products with serially diluted (188 ng to 0.37 ng) non-target RNA in the absence of PEG-nGO. (b) Melting curve analysis of qPCR-amplified products with serially diluted (47 ng to 0.37 ng) non-target RNA in the absence of PEG-nGO. (c) Real-time fluorescence monitoring of qPCR-amplified products with serially diluted (188 ng to 0.37 ng) non-target RNA in the presence of PEG-nGO. (d) Melting curve analysis of qPCR-amplified products with serially diluted (47 ng to 0.37 ng) non-target RNA in the presence of PEG-nGO. Red dotted arrow and cyan arrow represents nonspecific and specific amplicon dsDNA peak, respectively.