1	Supporting information
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4	Aptasensing of ciprofloxacin residue using graphene oxide modified by gold nanoparticle
5	and branched polyethyleneimine
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43 Fig. S2. A, B. FE-SEM with MAP images of GO-PEI-AuNPs/AuE surface with different magnifications.





- Fig. S3, A, B. FE-SEM with MAP images of Apt/GO-PEI-AuNPs/AuE with various magnifications.











79 Fig. S5. (A) CVs of GO-PEI-AuNPs/AuE in the potential range of -1.2 to +1.2V and sweep rate of 100



81 Histograms of peak current versus number of cyclic.











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Fig. S6. A) CVs of GO-PEI-AuNPs/AuE after attachment of aptamer with various volume of aptamer (5, 10, and 15μL) B) Histogram of peak current versus aptamer volume. C) CVs of Apt/GO-PEI-AuNPs/AuE in different incubation time of aptamer (4, 6, 8, 12, and 24 h), D) Histogram of peak current based on incubation time of aptamer on the surface of GO-PEI AuNPs/AuE. E) CVs of MCH/Apt/GO-PEI-AuNPs/AuE in diverse time (10, 20, 30, 40, 50 and 60 min) after CFX incubation, F) Histogram of peak current versus incubation time of CFX on the modified AuE surface.











Fig. S8. (A) DPVs of MCH/Apt/GO-PEI-AuNPs/AuE established aptasensor in the existence of some
interfering species (serine, valine, cysteine, methionine, and glutamine) in the solution of 0.01 M of
K₃Fe(CN)₆ (0.1M KCl). (B) Histogram of peak current versus various kind of interfering agent.







144Fig. S9. (A, C) The CVs of Apt/EDC-NHS/GO-PEI-AuNPs/AuE suggested probe to study the inter and145intra-day stability in 0.01 M of $K_3Fe(CN)_6$ (0.1M KCl) solution. (B, D) Histogram of peak current based146ondifferentstoragetimeviaCVtechnique.(SD-2.06, n=3).