

Supporting Information

Development of bioluminescent enzyme immunoassay for glycocholic acid based on
single-chain variable fragment-nanoluciferase fusion protein

Yunping Mu^a, Huajian Cai^a, Xindan Zhang^b, Mingxia Lin^a, Junhao Zeng^a, Yiyun
Wang^a, Ziming Wu^a, Suqing Zhao^a, Xiping Cui^{a, *}

^a *Department of Biomedical Engineering, School of Biomedical and Pharmaceutical
Sciences, Guangdong University of Technology, Guangzhou 510006, People's
Republic of China*

^b *Guangzhou Quality Testing & Inspection Institute, Guangzhou 511447, People's
Republic of China*

*Corresponding Author:

Dr. Xiping Cui

E-mail address: xipcui@gdut.edu.cn

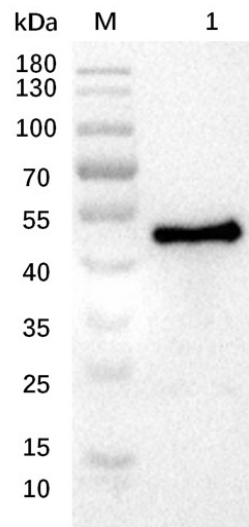


Figure S1 Analysis of scFv-Nluc fusion protein

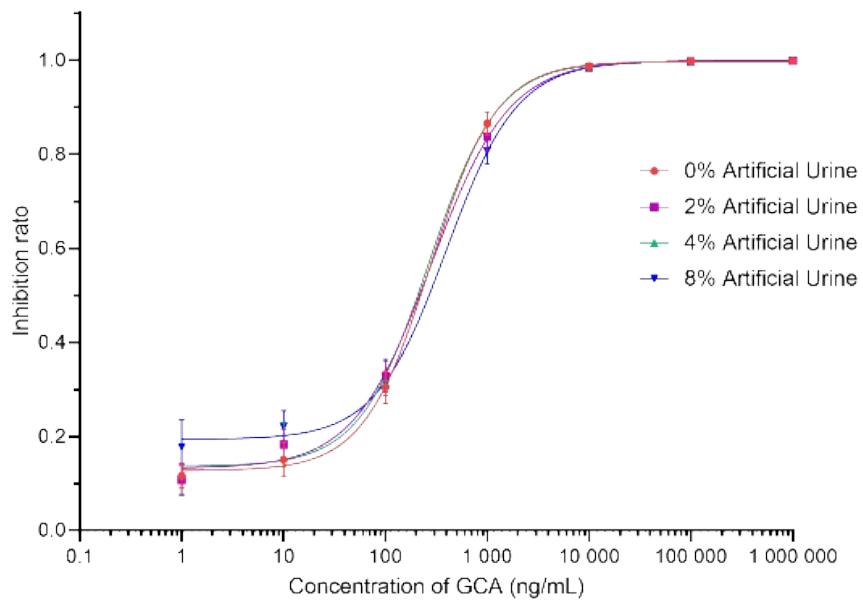


Figure S2 BLEIA competition curves of GCA prepared at various artificial urine concentrations

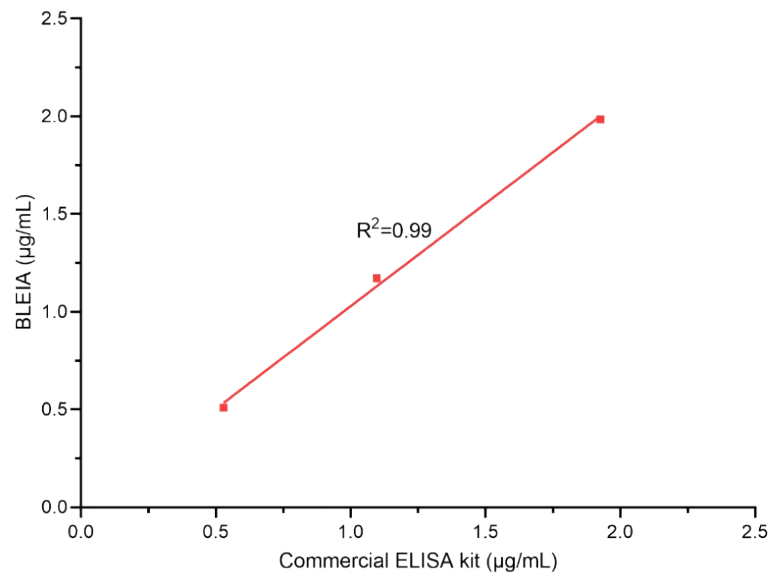


Figure S3 Correlation between BLEIA and Commercial ELISA kit results for the artificial urine sample

Table S1 Primer sequence information

Name	Sequence (5'-3')
SNF1	CC <u>G</u> AATTCGGTGGCCCAGGCGGCCCTGACTCAG
SNR1	ACCGCCGCTGCCGCCACCACCCTGGCCGGCCTGG
SNF2	GGTGGTGGCAGCGGCGGTATGGTCTTCACACTCGAA
SNR2	CCGCTC <u>G</u> AGCGCCAGAATGCGTTCGCA

Underline indicates *EcoR* I and *Xho* I restriction enzyme sites