Supporting Information

Quench-release-based Fluorescent Immunosensor for the Rapid Detection of Tumor Necrosis Factor-alpha

Haimei Li¹, Xinyu Li¹, Limei Chen¹, Baowei Li¹, Hang Dong², Hongying Liu¹, Xueying Yang¹, Hiroshi Ueda^{3,4*} and Jinhua Dong ^{1,3,4*}

¹Key Laboratory for Biological Medicine in Shandong Universities, Weifang Key Laboratory for Antibody Medicine, School of Life Science and Technology, Weifang Medical University, Weifang, 261053, China.

²School of Basic Medical Sciences, Peking University, Beijing 100191, China.

³World Research Hub Initiative, Institute of Innovative Research, Tokyo Institute of Technology, Yokohama 226-8503, Japan.

⁴Laboratory for Chemistry and Life Science, Institute of Innovative Research, Tokyo Institute of Technology, Yokohama, 226-8503 Japan

*Authors to whom correspondence should be addressed;

E-mail: dongjh@wfmc.edu.cn (J.D.); ueda@res.titech.ac.jp (H.U.)

Figure S1. The amino acid sequence of UQ1H-Fab. V_H and V_L sequences are shown

in magenta and blue, respectively. Tryptophan residues in V_H and V_L are shown in bold.

Cysteine in cys-tag is underlined.

UQ1H-Fab V_H-C_H1

MAQIEVNCSNETGEVQLVESGGGLVQPGRSLRLSCAASGFTFDDYAMHWVRQAPGKGLEWVS AITWNSGHIDYADSVEGRFTISRDNAKNSLYLQMNSLRAEDTAVYYCAKVSYLSTASSLDYWGQ GTLVTVSSASTKGPSVFPLAPSSKSTSGGTAALGCLVKDYFPEPVTVSWNSGALTSGVHTFPAVL QSSGLYSLSSVVTVPSSSLGTQTYICNVNHKPSNTKVDKKVEPKSASAAHHHHHHGAAEQKLIS EEDLNGAA

UQ1H-Fab Light Chain

MDIQMTQSPSSLSASVGDRVTITCRASQGIRNYLAWYQQKPGKAPKLLIYAASTLQSGVPSRFSG SGSGTDFTLTISSLQPEDVATYYCQRYNRAPYTFGQGTKLEIKRADAAPSVFIFPPSDEQLKSGTA SVVCLLNNFYPREAKVQWKVDNALQSGNSQESVTEQDSKDSTYSLSSTLTLSKADYEKHKVYAC EVTHQGLSSPVTKSFNRGEGGGSDYKDDDDK

Figure S2. Absorption spectra of QB1-TMR, QB2-TMR, QB1-ATTO and QB2-ATTO. For double labeled Q-body with TAMRA, a peak at 520 nm, for that with ATTO520, a peak at around 490 nm were also observed, which are derived from H-dimer, respectively.

