Electronic Supplementary Information

Remarkable increase in luminol electrochemiluminescence

by sequential electroreduction and electrooxidation

Xiaoyun Liu,^{a,b} Wenjing Qi,^{a,c} Wenyue Gao,^{a,c} Zhongyuan Liu,^a Wei Zhang,^a Ying Gao^b and Guobao Xu*,^a

^a State Key Laboratory of Electroanalytical Chemistry, Changchun Institute of Applied Chemistry, Chinese Academy of Sciences, Changchun, Jilin 130022, China. Fax: +86 431 85262747; Tel: +86 431 85262747; E-mail: guobaoxu@ciac.ac.cn

^b School of Chemistry and Environmental Engineering, Changchun University of Science and Technology, Changchun 130022, PR China.

University of the Chinese Academy of Sciences, Chinese Academy of Sciences, No. 19A
Yuquanlu, Beijing 100049, China;

*Corresponding author: Prof. G. B. Xu

Experimental Section

Chemicals and Reagent. Luminol was obtained from Aldrich and used without any further purification. A 1.0×10^{-2} M stock solution of luminol was prepared by dissolving luminol in 0.2 M NaOH. Luminol working solutions were prepared by diluting the stock solution. Supporting electrolyte was 0.2 M phosphate buffer solution (PBS). Other chemicals were analytical-reagent grade and used as received. Doubly distilled water was used throughout the experiments.

Apparatus. Cyclic voltammetric measurements were carried out using a model CHI 660C electrochemical working station (Shanghai CHI Instruments Co., China). ECL was performed using a BPCL ultra-weak luminescence analyzer. All experiments were carried out with a homemade three-electrode electrochemical cell system. The glassy carbon electrode (GCE), platinum wire, and Ag/AgCl (saturated

with KCl) were used as working, counter, and reference electrodes, respectively. All measurements were performed at room temperature.