## Pencil lead as a material for microfluidic 3D-electrode assemblies

Emilia Witkowska Nery \*, Magdalena Kundys-Siedlecka, Yoshitaka Furuya and Martin Jönsson-Niedziółka \*

Institute of Physical Chemistry, Polish Academy of Sciences, Kasprzaka 44/52, 01-224 Warsaw, Poland; mkundys@ichf.edu.pl (M.K.-S.); yfuruya@iis.u-tokyo.ac.jp (Y.F.)

\* Correspondence: ewitkowskanery@ichf.edu.pl (E.W.N.); martinj@ichf.edu.pl (M.J.-N.); Tel.: +48-22-343-3306 (E.W.N. & M.J.-N.)



**SI Figure 1.** Cyclic voltammetry studies of oxidation of FcDM performed for different flow rates (0-1000  $\mu$ L min<sup>-1</sup>). Electrode cell consisted of a 3D working electrode formed from 16 graphite posts, steel counter electrode and an Ag/AgCl reference. Scan rate 100 mV s<sup>-1</sup>.



SI Figure 2. Limiting current calculated for 3D working electrodes formed from 4 to 16 graphite posts.



**SI Figure 3.** Concentration profile of the reduced form of the redox probe along a cut-line through the second row of electrodes for different flow rates: black 10  $\mu$ l min<sup>-1</sup>, red 25  $\mu$ l min<sup>-1</sup>, green 100  $\mu$ l min<sup>-1</sup>. The profile is taken at the turning point in the CV (fig. 3b) at 0.6 V.



**SI Figure 4.** Concentration profile of the reduced form of the Redox probe around the electrodes for different flowrates: A) 10  $\mu$ l min<sup>-1</sup>, B) 25  $\mu$ l min<sup>-1</sup>, C) 100  $\mu$ l min<sup>-1</sup>. The image is saved at the turning point in the CV (fig. 3B) at 0.6 V. The flow is from the left to the right side.



**SI Figure 5.** Cyclic voltammetry of FcDM performed using a pencil graphite electrode: prepared by cycling in FcDM, oxidized by cycling in NaOH and oxidized by applying constant potential in PBS.



**SI Figure 6.** Square wave voltammetry of dopamine solutions recorded using the array device, corresponding to the calibration curve presented in the manuscript in Fig. 7