## **Electronic Supplementary Information (ESI)**

## Versatile multi-functionalization of protein nanofibrils for biosensor applications

L. Sasso,\*<sup>*abc*</sup> S. Suei,<sup>*bd*</sup> L. Domigan,<sup>*abce*</sup> J. Healy,<sup>*bc*</sup> V. Nock,<sup>*abf*</sup> M. A. K. Williams<sup>*adg*</sup> and J. A. Gerrard\*<sup>*abch*</sup>

<sup>a</sup> MacDiarmid Institute for Advanced Materials and Nanotechnology, Wellington 6140, New Zealand.

<sup>b</sup> Biomolecular Interaction Centre, University of Canterbury, Christchurch 8140, New Zealand.

<sup>c</sup> School of Biological Sciences, University of Canterbury, Christchurch 8140, New Zealand.

<sup>d</sup> Institute of Fundamental Sciences, Massey University, Palmerston North 4442, New Zealand.

<sup>e</sup> Department of Biomedical Engineering, Tufts University, Medford 02155, MA, USA.

<sup>*f*</sup> Department of Electrical and Computer Engineering, University of Canterbury, 8140, New Zealand.

<sup>g</sup> Riddet Institute, Massey University, Palmerston North 4442, New Zealand.

<sup>h</sup> Callaghan Innovation Research Ltd, Lower Hutt 5040, New Zealand.

S1. Cyclic voltammetry characterization of biosensor platform	2	
S2. Biosensor response to various glucose concentrations	3	
S3. Electrode roughness due to protein nanofibril modifications	4	



S1. Cyclic voltammetry characterization of biosensor platform

**Figure S1.** Comparison of glucose responses with same concentration (500 mM) for different enzyme immobilization methods: adsorption onto a bare gold electrode (Au/GOx), GOx-functionalized WPNFs deposition (Au/WPNF-GOx), dual GOx- and SH-functionalized WPNFs deposition (Au/WPNF-GOx-SH), and a bare electrode with no enzyme immobilization (Au), where the signal is solely caused by the ferrocene mediator. Potential sweep rates are 100 mV/s for all curves, using an Au counter and Ag reference electrode.





**Figure S2.** Comparison of biosensor current peak response for different concentrations of glucose. Inset shows the linear response for glucose concentrations smaller than 2 mM. Potential sweep rates are 100 mV/s for all curves, using an Au counter and Ag reference electrode. The limit of detection was calculated to be 0.11 mM (3 sigma method).



S3. Electrode roughness due to protein nanofibril modifications

**Figure S3.** Comparison of electrode roughness due to WPNF modifications, from AFM imaging: A) bare Au electrode, B) electrode after WPNFs deposition, and C) electrode after SH-functionalized WPNFs deposition. All samples were rinsed with milliQ water before AFM measurements.