## Supplementary File 1

Based on the results, the final protocol is as following:

- 1. Rinse screen-printed substrates with Isopropyl Alcohol/Milli-Q water.
- 2. Dry using a N2 stream.
- 3. Immerse screen-printed substrates in ODT (10mM in ethanol) solution at room temperature (RT) in dark overnight.
- 4. Rinse electrode array with ethanol
- 5. Dry using a N2 stream.
- 6. Deposit each antibody (CTx, PINP 100 μg/mL in PBS) on corresponding working electrodes & incubate at 37 °C in a humidity-controlled environment for 2 hrs.
- 7. Rinse electrode array with PBS solution (10 mM pH 7.4)
- 8. Dry gently using a N2 stream.
- 9. Deposit Blocker Casein 1% (in PBS) on working electrodes & incubate at 37 °C in a humiditycontrolled environment for 1 hr.
- 10. Rinse electrode array with PBS Tween-20 solution (0.05% Tween 20 in 10mM PBS, pH 7.4)
- 11. Rinse electrode array with PBS solution (10 mM pH 7.4)
- 12. Dry gently using a N2 stream.

The screen-printed electrode arrays are characterized after each step using Cyclic Voltammetry/EIS electrochemical measurements.