## **Supporting information**

## Synthesis and Characterization of a Fluorinated S-Nitrosothiol as the Nitric

## **Oxide Donor for Fluoropolymer-Based Biomedical Device Applications**

Yang Zhou,<sup>a</sup> Qi Zhang,<sup>a</sup> Jianfeng Wu,<sup>b</sup> Chuanwu Xi<sup>b</sup> and Mark E. Meyerhoff\*<sup>a</sup>

 <sup>a</sup> Department of Chemistry, University of Michigan, Ann Arbor, MI 48109, USA
<sup>b</sup> Department of Environmental Health Sciences, University of Michigan School of Public Health, Ann Arbor, MI 48109, USA



**Fig. 1S** UV-Vis spectra for the photolysis of SNAP (250.0  $\mu$ M) at 23 °C in a mixture of PBS and DMSO (50:50, v/v). Inset: Plot of absorbance at 342 nm vs. time for this reaction and solid line is fit to first order rate equation line (rate constant  $k_{obs} = (1.50 \pm 0.03) \times 10^{-2} \text{ s}^{-1}$  and  $t_{1/2} \sim 46 \text{ s}$ ).



**Fig. 2S** Plot of absorbance at 342 nm vs. time for the thermal decomposition of SNAP (206.0  $\mu$ M) at 37 °C in a mixture of PBS and DMSO (50:50, v/v), (rate constant  $k_{obs} = (3.45 \pm 0.02) \times 10^{-5} \mu$ M Min<sup>-1</sup>).



Fig. 3S NO flux for the PDVF tubings swelled in C<sub>2</sub>F<sub>5</sub>-SNAP (400.5 mg/mL in THF) under physiological conditions (PBS, 10 mM, pH 7.4, with 100  $\mu$ M EDTA at 37 °C in the dark), and the total loading of C<sub>2</sub>F<sub>5</sub>-SNAP in PVDF tubing was estimated to be ~ 32.3 nmol/mg.





(b)

Fig. 4S (a) Calibration curve for the concentration of  $C_2F_5$ -NAP obtained in HPLC-MS and (b) Calibration curve for the concentration of  $C_2F_5$ -NAP disulfide obtained in HPLC-MS.



Fig. 5S The corresponding black image for Fig. 9d

## NMR spectra







