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> Supplemental Information for In Vitro Oxidations of Low-Density Lipoprotein and RAW 264.7 Cells with Lipophilic O(³P)-Precursors

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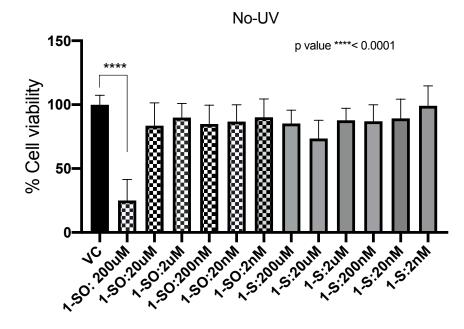


Figure S1: Cell viability when cells were treated with 1-SO and 1-S as a function of concentration without UV-A irradiation. Error bars – 95% CI. Cell viability calculated as $(Abs_{sample} - Abs_{blank})*100/(Abs_{VC}-Abs_{blank})$. N=10

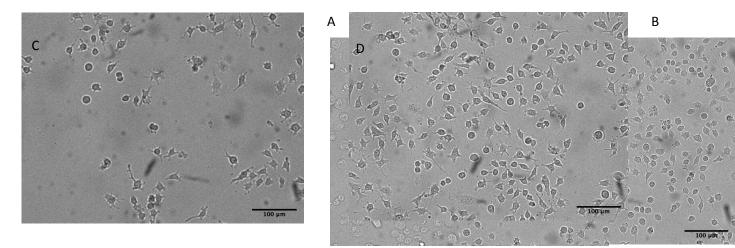


Figure S2. Visualization of RAW 264.7 cells pre- and post-UV treatment. Cells were plated on two 10X200 mm cell culture dishes (Cellstar by Grenier bio-one) in 5mL media (cell density 88,000 cells/mL) and incubated for 24 hours. One culture dish was irradiated in a Luzchem photoreactor with 14 UV-A LZC bulbs for 2 hours and a control dish was wrapped in foil and kept adjacent to the photoreactor. The dishes were then visualized using a Olympus BX60 Microscope and images were captured at 200X magnification. **A.** Cells prior to irradiation (plate 1). **B.** Cells prior to irradiation (plate 2) **C.** Cells after 2 hours irradiation (plate 1). **D.** Cells kept in dark for 2 hours (plate 2)

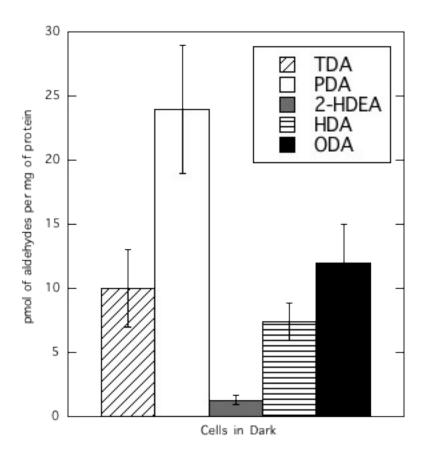
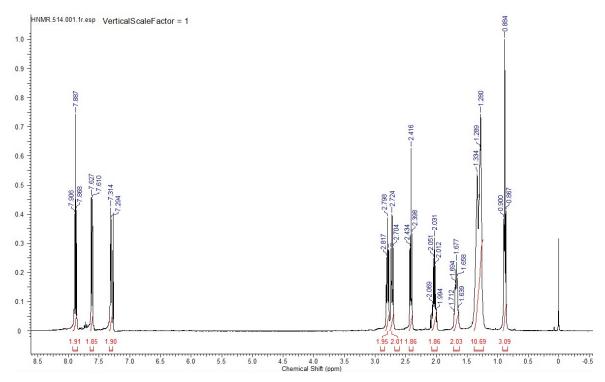
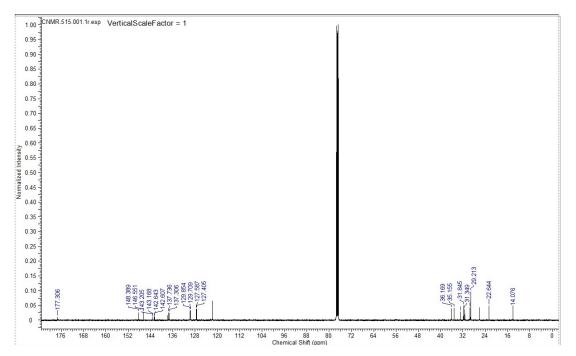


Figure S3. Amount of oxidized lipid product arising from RAW 264.7 cells after irradiation without any additional compounds.

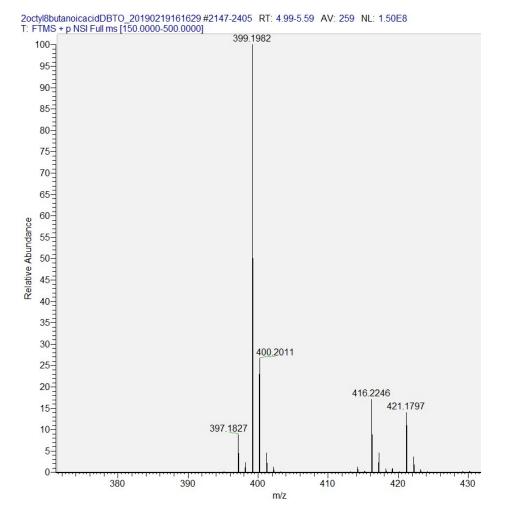
HNMR for 1-SO



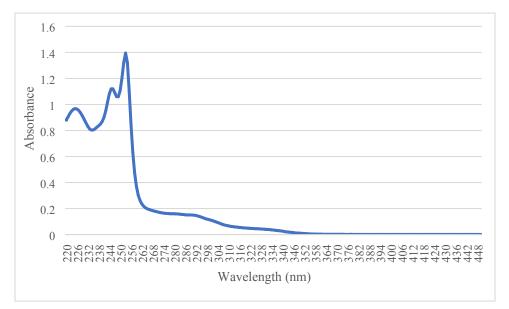
CNMR for 1-SO



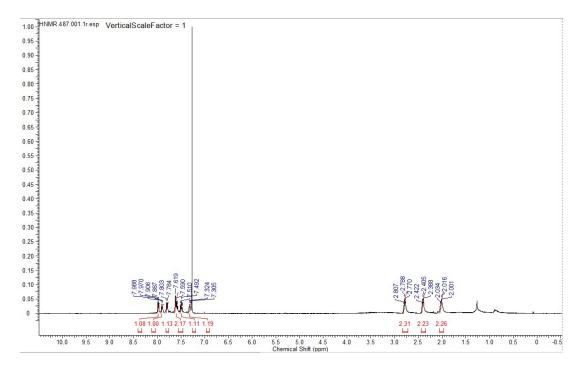
HRMS for 1-SO



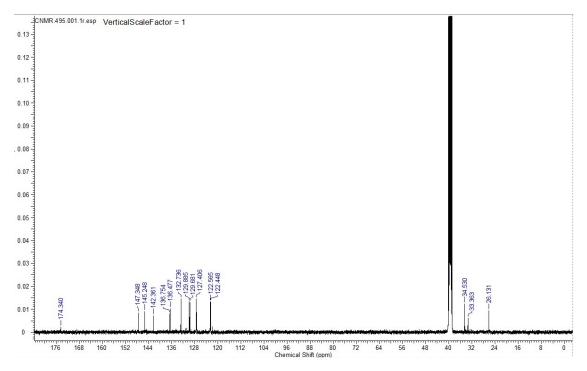




HNMR for 2-SO



CNMR for 2-SO



HRMS for 2-SO

