Supporting Information

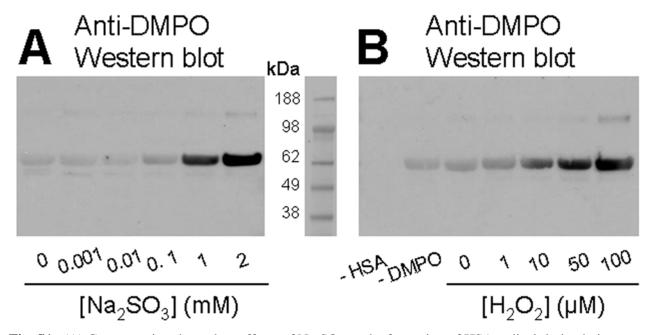


Fig. S1. (A) Concentration-dependent effects of Na₂SO₃ on the formation of HSA radical-derived nitrone adducts by Western blotting. Reactions including HSA (600 μM), H_2O_2 (100 μM), DMPO (1 mM) and sulfite as indicated were initiated with 1 μM EPO, and the mixtures were incubated for 1 h at 37 °C in 100 mM phosphate buffer (pH 7.4). (B) Concentration-dependent effects of hydrogen peroxide on the formation of HSA radical-derived nitrone adducts by Western blotting. Reactions including HSA (600 μM), Na₂SO₃ (2 mM), DMPO (1 mM) and H_2O_2 as indicated were initiated with 1 μM EPO, and the mixtures were incubated for 1 h at 37 °C in 100 mM phosphate buffer (pH 7.4). Each lane contained 3.8 μg of HSA.

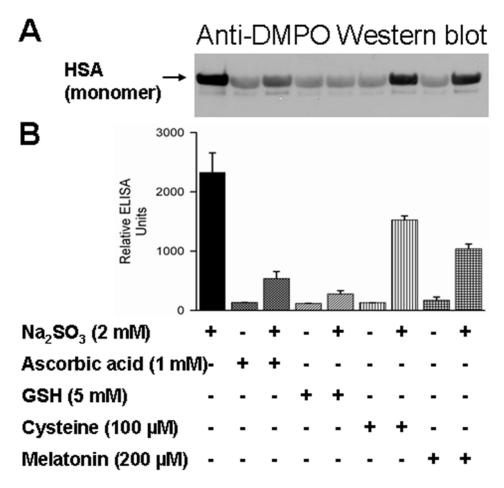


Fig. S2. Effect of inhibitors on the formation of DMPO-HSA-derived radical nitrone adducts. (A) Western blotting. (B) ELISA. Reaction mixtures containing HSA ($600 \, \mu M$), Na₂SO₃ (2 mM), DMPO (1 mM), and H₂O₂ ($100 \, \mu M$) with and without the indicated concentrations of ascorbic acid, GSH, cysteine, and melatonin were initiated by EPO (1 μM). ELISA data presented are the means \pm s.d. from three independent determinations using fresh preparations of all reaction components.