

Supporting Information

Sulfated *Undaria pinnatifida* polysaccharides inhibit kidney stone formation through crystalline modulation and relieving cellular oxidative damage and inflammation

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Supplementary Figures

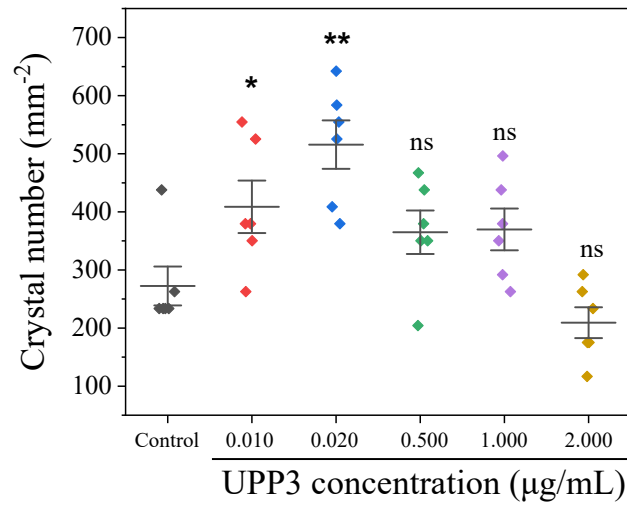


Figure S1. Quantitative statistical diagram of calcium oxalate crystal number, Compared with control group, **P<0.05, **P<0.01.

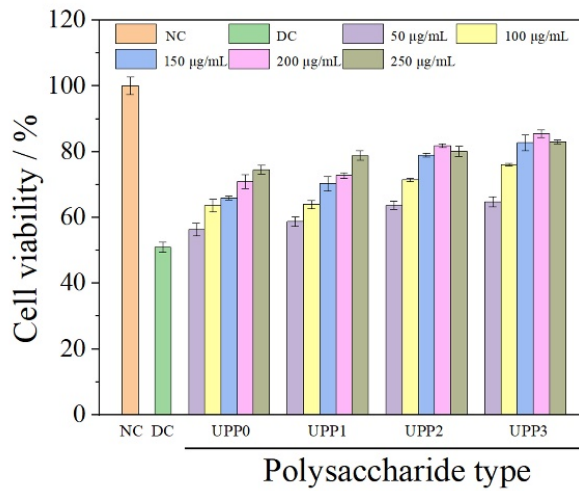


Figure S2. Detection of cell viability before and after UPPS protection of NRK-52E cells. NC: Normal control group, DC: Damaged control group; polysaccharide concentration: 50 and 200 µg/mL; sodium oxalate concentration: 2 mM; protection time: 24h.

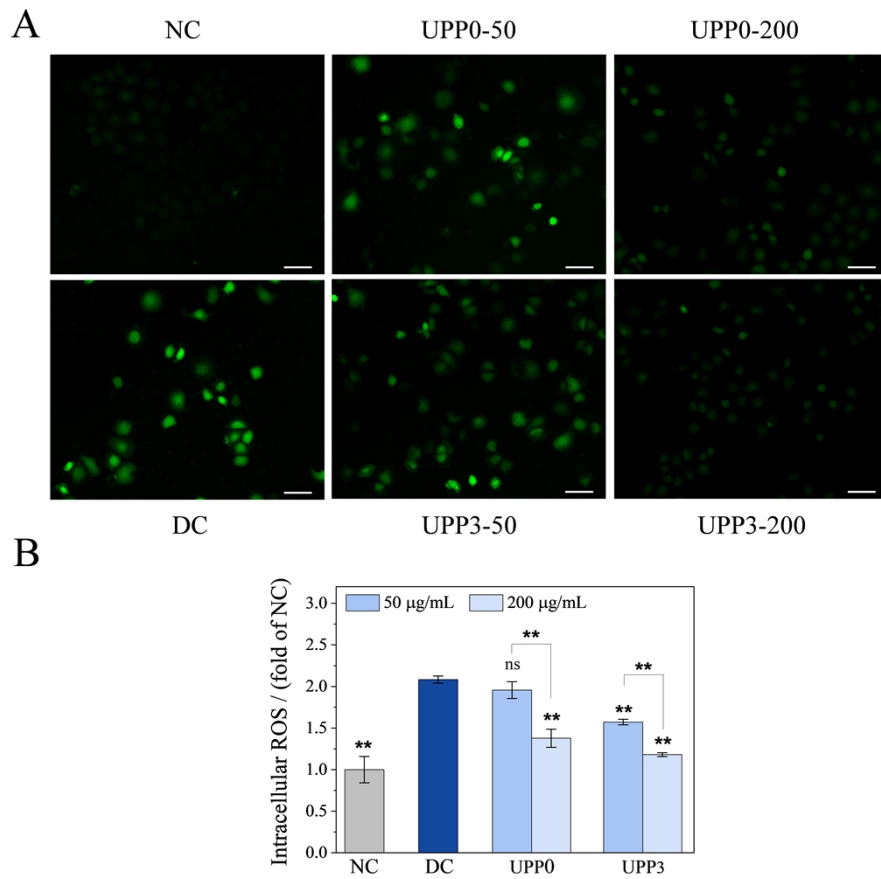


Figure S3. The ROS level of NRK-52E cells was detected by DCFH-DA staining, scale bar: 50 μm . NC: Normal control group, DC: Damaged control group; polysaccharide concentration: 50 and 200 $\mu\text{g}/\text{mL}$; sodium oxalate concentration: 2 mM; protection time: 24h; Compared with DC group, $**P < 0.01$.

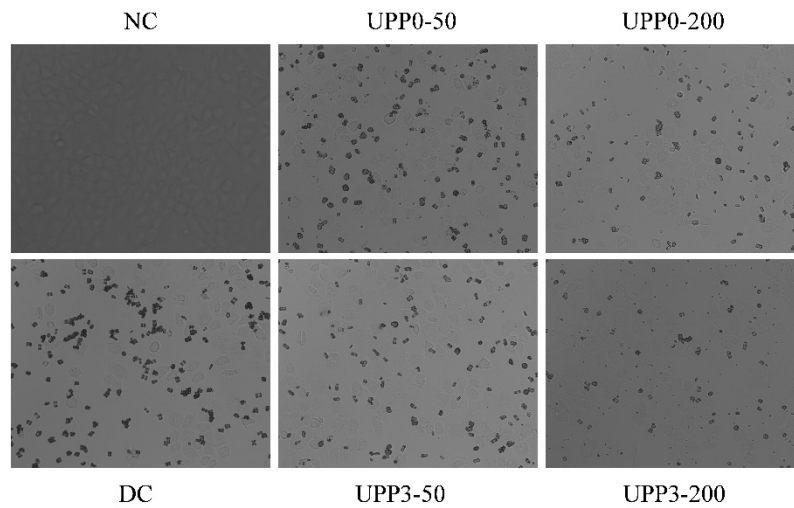


Figure S4. The morphology of NRK-52E cells before and after UPPS protection was observed, scale bar: 50 μm . NC: Normal control group, DC: Damaged control group; polysaccharide concentration: 50 and 200 $\mu\text{g}/\text{mL}$; sodium oxalate concentration: 2 mM; protection time: 24h.

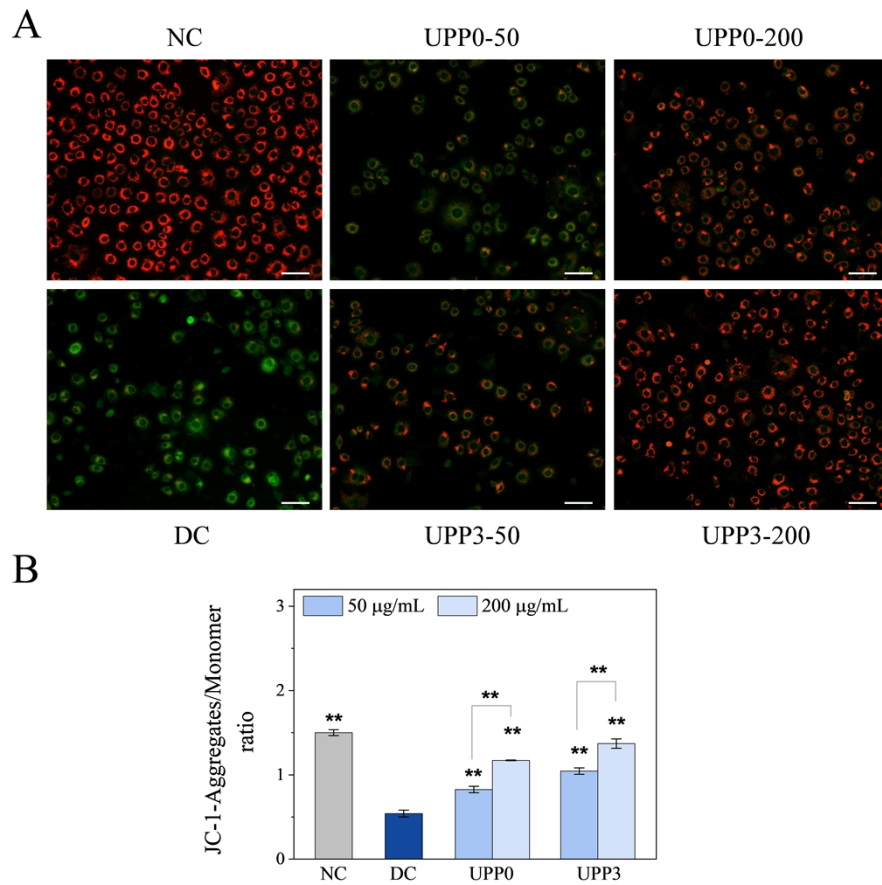


Figure S5. The mitochondrial membrane potential of NRK-52E cells before and after UPPS protection was detected by JC-1 staining, scale bar: 50 μ m. NC: Normal control group, DC: Damaged control group; polysaccharide concentration: 50 and 200 μ g/mL; sodium oxalate concentration: 2 mM; protection time: 24h; Compared with DC group, ** $P < 0.01$.

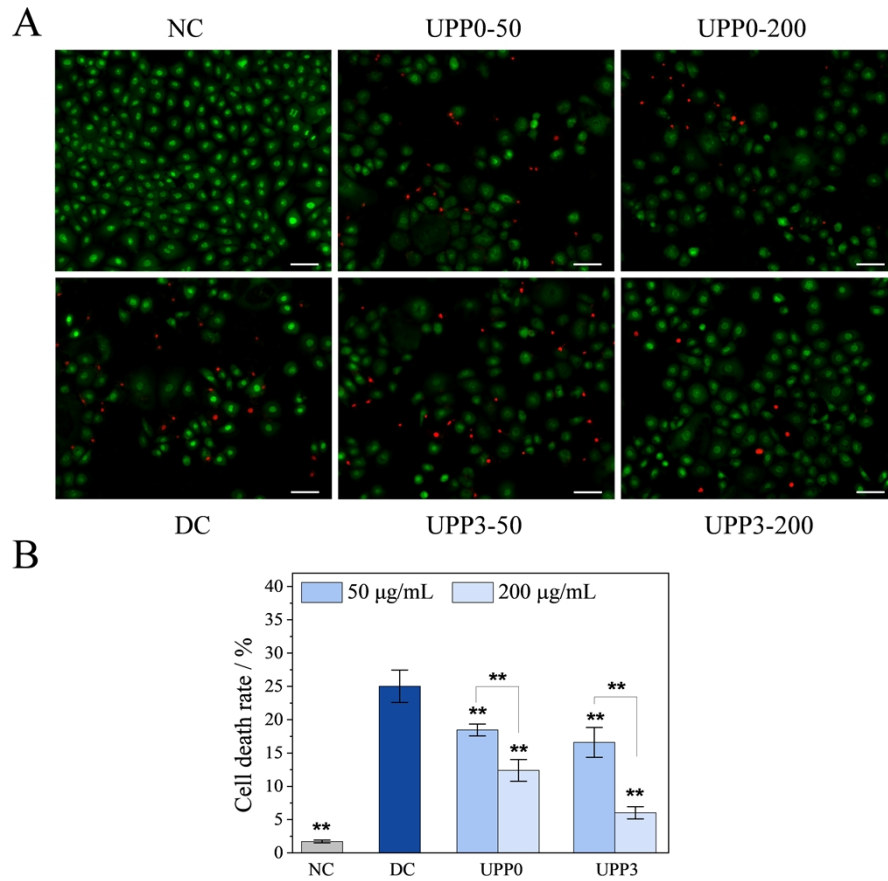


Figure S6. AO/EB double staining was used to detect the death of NRK-52E cells before and after UPPS protection, scale bar: 50 μm . NC: Normal control group, DC: Damaged control group; polysaccharide concentration: 50 and 200 $\mu\text{g}/\text{mL}$; sodium oxalate concentration: 2 mM; protection time: 24h; Compared with DC group, $^{**}P < 0.01$.