

## **Supporting Information**

### **Sirtuin 6 (SIRT6) regulates redox homeostasis and downstream signaling events in human articular chondrocytes**

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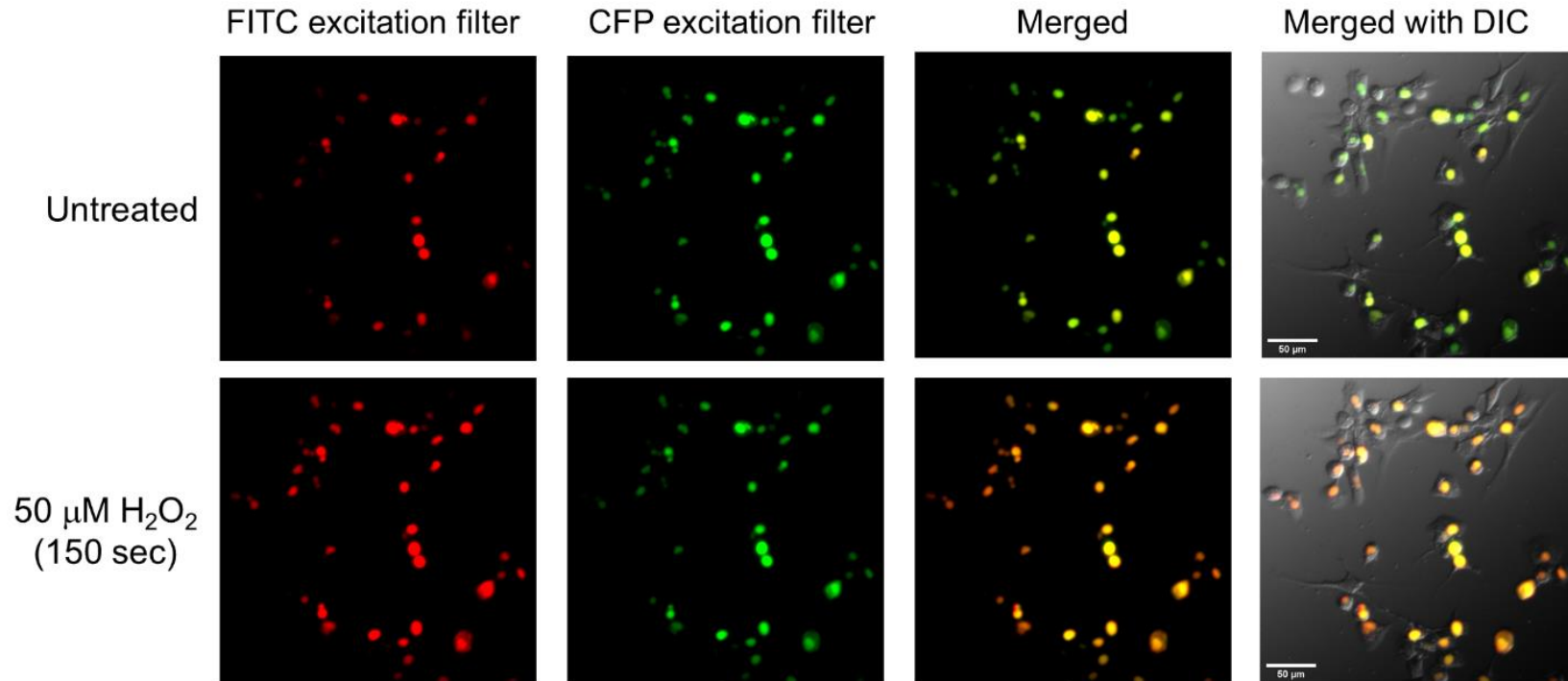
#### **Material Included**

Supplementary Figure 1. Validation of the NLS-HyPer-DAAO measurement of H<sub>2</sub>O<sub>2</sub>.

Supplementary Figure 2. Identification of nuclear ROI using CellProfiler software for subsequent measurement of fluorescence intensity and determination of H<sub>2</sub>O<sub>2</sub> generation.

Supplementary Figure 3. Expression and nuclear localization of NLS-HyPer-DAAO.

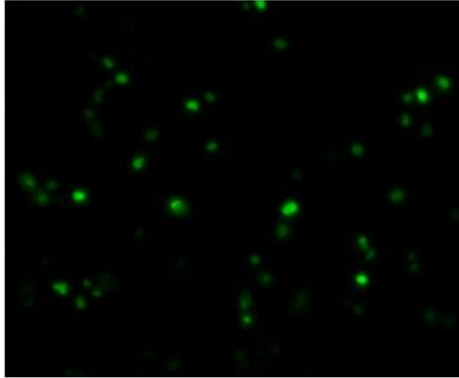
**Supplementary Figure 1.**



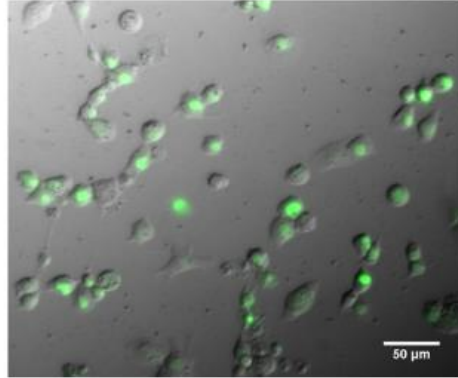
**Supplementary Figure 1. Validation of the NLS-HyPer-DAAO measurement of H<sub>2</sub>O<sub>2</sub>.** Human chondrocytes were transduced with NLS-HyPer-DAAO for 48 hours and then treated with 50 μM H<sub>2</sub>O<sub>2</sub>. An increase in fluorescence emission intensity with FITC excitation filter (red), which is most notable on the merged image, demonstrates that the HyPer probe is oxidized in response to exogenously added H<sub>2</sub>O<sub>2</sub>.

**Supplementary Figure 2.**

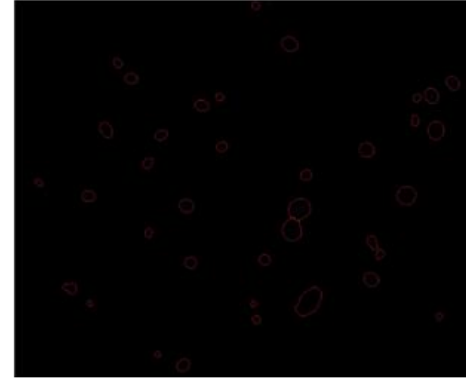
Nuclear fluorescence  
signal (CFP)



Nuclear fluorescence  
signal merged with DIC

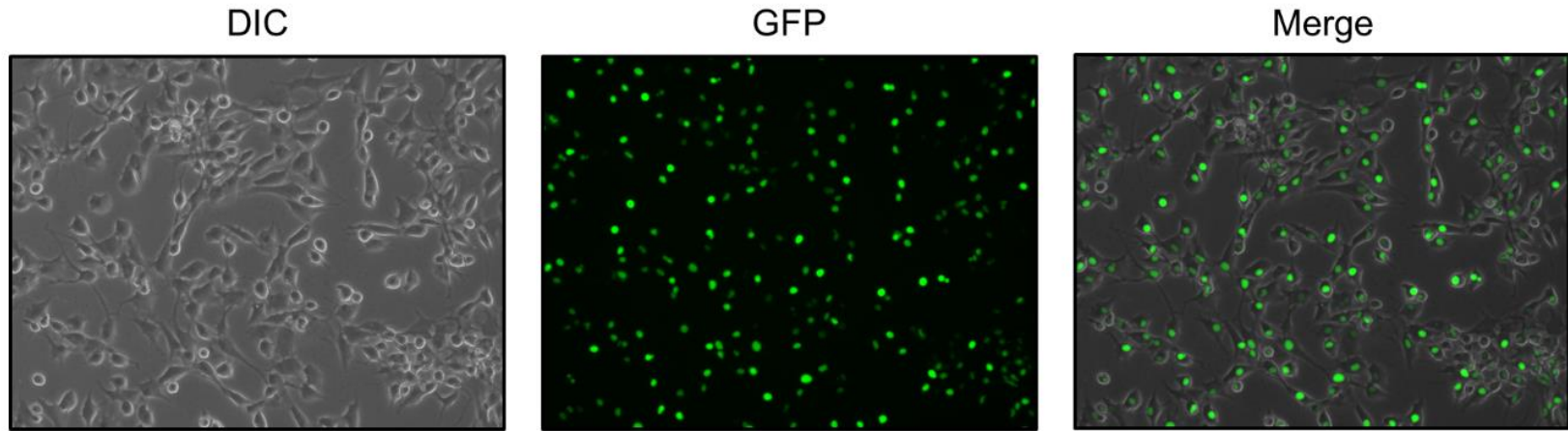


Identified nuclear  
ROI



**Supplementary Figure 2. Identification of nuclear ROI using CellProfiler software for subsequent measurement of fluorescence intensity and determination of H<sub>2</sub>O<sub>2</sub> generation.**

**Supplementary Figure 3**



**Supplementary Figure 3. Expression and nuclear localization of NLS-HyPer-DAAO.** Human chondrocytes were transduced with the adenoviral vector encoding NLS-HyPer-DAAO for 48 hours and nuclear fluorescence was detected using an EVOS m5000 imaging system at 20x magnification.