ROS-Targeted Nanoprobes for Detection and Imaging of Cellular Senescence

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# ROS-Targeted Nanoprobe for Detection and Imaging of Cellular Senescence

## Background & Unmet Need
- Cellular senescence is a state of irreversible cell cycle arrest associated with aging, in which cells stop proliferating.
- Senescent cells are drug resistant and may secrete factors such as cytokines into surrounding tissues, causing low-grade inflammation.
- Senescence can be caused by cellular stress or damage, including mitochondrial dysfunction, oxidative stress, or DNA damage.
- Cells can also become senescent in response to chemotherapy and escape treatment, leading to future tumor recurrence.
- Senescence is currently imaged using beta-galactosidase (Xgal); however, this label is not senescence-specific and requires cell fixing and long incubation times.
- **Unmet Need:** Improved methods for detection and imaging of senescent cells.

## Technology Overview
- **The Technology:** A novel fluorogenic nanoprobe for labeling cellular senescence via detection of reactive oxygen species (ROS).
- ROS are known to play a role in progression and maintenance of cell senescence, and ROS levels are directly related to induction of cellular senescence.
- The inventors have created a novel nanoprobe, D3, which fluoresces in response to high levels of ROS, thereby labeling senescent cells.
- **PoC Data:** In tumor-bearing mice, D3 accumulated quickly and preferentially in tumors when administered intravenously.
- Fluorescent signal from D3 was specifically turned on in senescent tumors, which were induced via treatment of tumor-bearing mice with Palbociclib.
- The fluorescence signal from D3 in senescent tumors was 3-fold higher than that of non-senescent tumors.

## Inventors:
- Ching-Hsuan Tung
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## Patents:
- Provisional Filed

## Publications:
- N/A

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## Cornell Reference:
- D-10580
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**Technology Applications**

- Imaging nanoprobe to identify senescent tumors following chemotherapy
- Long-term study of disease progression and treatment response for senescence-associated conditions, including aging and fibrosis
- Real-time imaging of changes in cellular senescence
- Identification and isolation of senescent cells for further research

**Technology Advantages**

- D3 is remarkably stable in normal physiological conditions
- D3 does not require cells to be fixed or to undergo long incubation times
- Fluorescence intensity of D3 is dependent on ROS production level and corresponds to senescence progression, allowing for real-time imaging

**Supporting Data / Figures**

- Figure 1: D3 detects palbociclib-induced senescence both in vitro (Top) and in vivo (Bottom). DCF-DA is a marker for ROS production.

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