

Early Detection of Parkinson's Disease using Noninvasive Biomarkers

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Background & Unmet Need

- Parkinson's Disease (PD) is the second most common neurodegenerative disease, affecting 10 million people worldwide
- The presence of Lewy bodies, which are made up by aggregated α-synuclein protein deposits, are a hallmark of PD
- Emerging diagnostics for PD measure levels of αsynuclein in spinal fluid, which is collected from invasive lumbar punctures
- Lumbar punctures can be painful and put patients at risk for spinal fluid leakage, prolonged headaches, back pain, and bleeding
- It is currently difficult to accurately measure pathological α-synuclein aggregates in living patients using non-invasive methods
- Unmet Need: Noninvasive biomarkers for early assessment of Parkinson's disease

Technology Overview

- The Technology: A method for early detection of Parkinson's Disease using an imaging modality to measure a novel biomarker
- **The Discovery:** In a novel mouse model of PD, the inventor has discovered a new biomarker that is visually apparent using a readily available imaging modality
- The emergence of this biomarker temporally coincides with onset and progression of disease as well as Lewy body deposition, and could be used a biomarker for PD detection
- **PoC Data**: In a mouse model of PD, diseased mice had significantly more expression of the biomarker (p<0.001) than control mice starting at 2 months old
- The expression of the biomarker increases over time in diseased mice, matching PD disease progression

- Inventor: Ching-Hwa Sung
- Patents: Provisional Filed

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Technology Applications

- Early, noninvasive detection of Parkinson's Disease
- · Method of monitoring PD progression over time
- Method of assessing treatment efficacy during clinical trials
- Diagnosis of other neurodegenerative diseases involving aggregated α-synuclein deposition, such as Lewy body dementia

Technology Advantages

- Noninvasive, unlike current diagnostics that utilize spinal taps or biopsies
- Measures actual levels of pathological α-synuclein inclusions, rather than amplifying the quantity, enabling more accurate assessment of the disease
- Increased expression of the biomarker correlates to disease progression, allowing for better assessment of the state of the disease or treatment efficacy
- More cost effective than methods requiring sampling and assays for protein or nucleic acid levels

Weill Cornell Medicine



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