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Common pathophysiology underpinning Parkinson's disease and other neurodegenerative diseases Sheila Pirooznia, Nikhil Panicker ,Liana Rosenthal, 2025-09-30 A wide range of neurodegenerative disorders affect the central nervous system causing changes in neuronal circuitry loss in neuronal connections and eventual neuronal death The high prevalence of neurodegenerative diseases like Parkinson's disease Alzheimer's disease and other degenerative brain

disorders in the aging population poses a significant burden on global healthcare systems Given their highly complex nature etiologic research on many of these neurodegenerative diseases are often conducted in isolation However elucidating conserved mechanistic underpinnings of neurodegenerative diseases could be utilized to develop pan neurodegenerative treatments interventions Is it possible to tease apart the similarities and differences between different neurodegenerative diseases at the molecular cellular and organismal levels Could there be common pathophysiological pathways that lead to synaptic loss and neuronal death This research topic intends to address these questions broadly and encourages the submission of research findings that could help understand the interplay between different neurodegenerative diseases and uncover novel therapeutic opportunities for neuroprotection during aging Some examples include Proteotoxicity Clearance of misfolded proteins and proteostasis endoplasmic reticulum stress and the unfolded protein response e q proteasomes lysosomes autophagy Mitochondrial function Mitochondrial mechanisms in Parkinson's Disease Huntington's Disease and Amyotrophic Lateral Sclerosis ALS ataxias and other diseases RNA transcription and processing Synuclein Parkinson's Trinucleotide repeat expansions Huntington s ataxias hexanucleotide repeat expansion familial FTD and ALS TDP43 Protein translation deficits Defects in mRNA localization mRNA sequestration ribosome biogenesis effects of mutant tRNA synthetases Inflammation Involvement of microglial astrocytic activation and the innate immune system Adaptive immune responses to neurodegeneration Prion like spread of pathological proteins Pathology stemming from distinct strains of tau and alpha synuclein TDP43 the gut brain axis Connectomics Identification of neural cell populations brain regions neural circuits and or large scale networks connectome that are vulnerable during brain aging and contribute to neurodegeneration Genomics and epigenomics Identification of genetic and epigenetic contributions to neurodegenerative syndrome and the overlapping phenotypic presentations of individuals with the same genetic mutations Further identification of genetic and epigenetic mechanisms that are associated with motor cognitive decline Neurogenesis or adaptive cell stress response pathways molecular cellular synaptic and neural circuitry mechanisms underlying brain plasticity Develop and characterize novel animal models of neuropathology Human cell reprogramming approaches iPSCs 3D or organoid culture approaches to study molecular physiological and systems cell biology Identification of biomarkers that could distinguish different neurodegenerative diseases Clinical correlates of neuroanatomical changes Brain iron accumulation The relationship between clinical symptoms and brain iron accumulation and comparisons between localization and amount of brain iron accumulation between different neurodegenerative diseases Non motor symptoms Sleep and circadian disturbances Smell Evidence-Based Imaging in Pediatrics L. Santiago Medina, Kimberly E. Applegate, C. Craig anosmia cognitive changes Blackmore, 2010-03-10 This practical book is ideal for readers who want to rapidly determine the appropriate imaging for pediatric patients The text provides a concise and accessible summary of the literature on how and when to use imaging studies Chapters address the essentials such as cost effectiveness and are written in collaboration by renowned specialists in

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