







## NERUODEGENERATIVE DISEASES









# What are Neurodegenerative Diseases?

- As the name suggests, in neurodegenerative diseases, there is progressive degeneration of neurons and nervous tissue.
- Some common examples are:
  - Alzheimer's Disease
  - Parkinson's Disease
  - Huntington's Disease





### Can you please explain Alzheimer's Disease?



### Etiology

- Increasing age
- Down's syndrome (as gene for beta amyloid present on chromosome 21)
- Deficiency of acetylcholine (neurotransmitter)

### Pathophysiology:

- Misfolding of beta amyloid proteins, leads to their extracellular accumulation, forming beta amyloid plaques.
- Tau protein is hyperphosphorylated and starts accumulating, forming Neurofibrillary tangles.
- Hippocampus and cerebral cortex also exhibit loss of neurons, presenting as cerebral atrophy.

### Clinical presentation:

- Progressive memory loss
- Decline in cognitive function

Imagine an old man forgetting everything, sometimes even his family members, and finds it difficult to carry out daily activities like putting on his shirt.





### Can you please explain its progression visually?



Progression of Alzheimer's Disease







Mild Alzheimer's Disease



Severe Alzheimer's Disease





### Can you please explain Parkinson's Disease?



### Etiology:

- Deficiency of dopamine
- Idiopathic
- Age-related

### Drugs:

- Too much of antipsychotics cause Parkinson's
- Too much of anti-Parkinson's drugs cause psychosis
- Parkinson's (dopamine deficiency) and psychosis (dopamine excess) are two ends of the spectrum.

### **Pathophysiology**

- Degeneration of dopaminergic neurons in substantia nigra
- Formation of Lewy bodies: alpha-synuclein containing eosinophilic inclusions in neurons of substantia nigra.

### Clinical presentation (mnemonic: RBC)

- Resting tremor (pin-rolling, coin-counting)
- Bradykinesia
- Cog-wheel rigidity
- Mask- like face
- Postural instability





### Can you please explain how to visually remember it?











### Can you please explain Huntington's Disease?



### **Etiology**

- Autosomal dominant
- CAG trinucleotide repeats in huntingtin gene.

### Pathophysiology

- Aggregates of huntingtin gene
- Atrophy of caudate nucleus

### Clinical presentation

- Chorea: involuntary rapid (jerky) movements
- Decline in cognitive function
- Psychiatric symptoms

