Proteinopathies

Amyloidopathy
- Alzheimers disease
- Cerebral beta amyloid angiopathy
- Retinal ganglion cell degeneration
- AL, AA and AH amyloidosis
- Hereditary cerebral hemorrhage with amyloidosis
- Familial amyloidotic neuropathy
- Corneal lactoferrin amyloidosis

Taupathies
- Progressive supranuclear palsy
- Corticobasal degeneration
- Primary age related taupathies
- Picks disease
- Age related astrogliopathy
- Argyrophilic grain disease
- Global glial taupathy

Syneuclopathies
- Parkinsons disease
- Dementia with lew body
- Pure autonomic failure
- REM sleep behaviour disorder
- Multiple system atrophy

Trinucleotide repeat protein disorder
- Hungtington dosease
- Dentatorubral-pallidoluysian atrophu
- Spinobulbar muscle atrophy
- Spino cerebellar ataxia
- Fragile X syndrome
- Baratela scott syndrome
- Myotonic dystrophy
- Fredreichs ataxia

TDP-43pathies
- Fronto temporal lobe degeneration
- Fronto temporal lobe degenration with neuron disease
- Hippocampal sclerosis
- Frontotemporal dementia with parkinsonism

Prion related proteinopathies
- Creutzfeldt-jakob disease
  - Iotragenic CJD
  - Variant CJD
  - Familial CJD
  - Sporadic CJD
- Gerstmann-straussler scheinker syndrome
- Fatal familial insomnia
- Kuru
- Familial spongiform encephalopathy
- Variable protease-sensitive prionopathy

Miscellaneous
- Familial british dementia
- Familial danish dementia
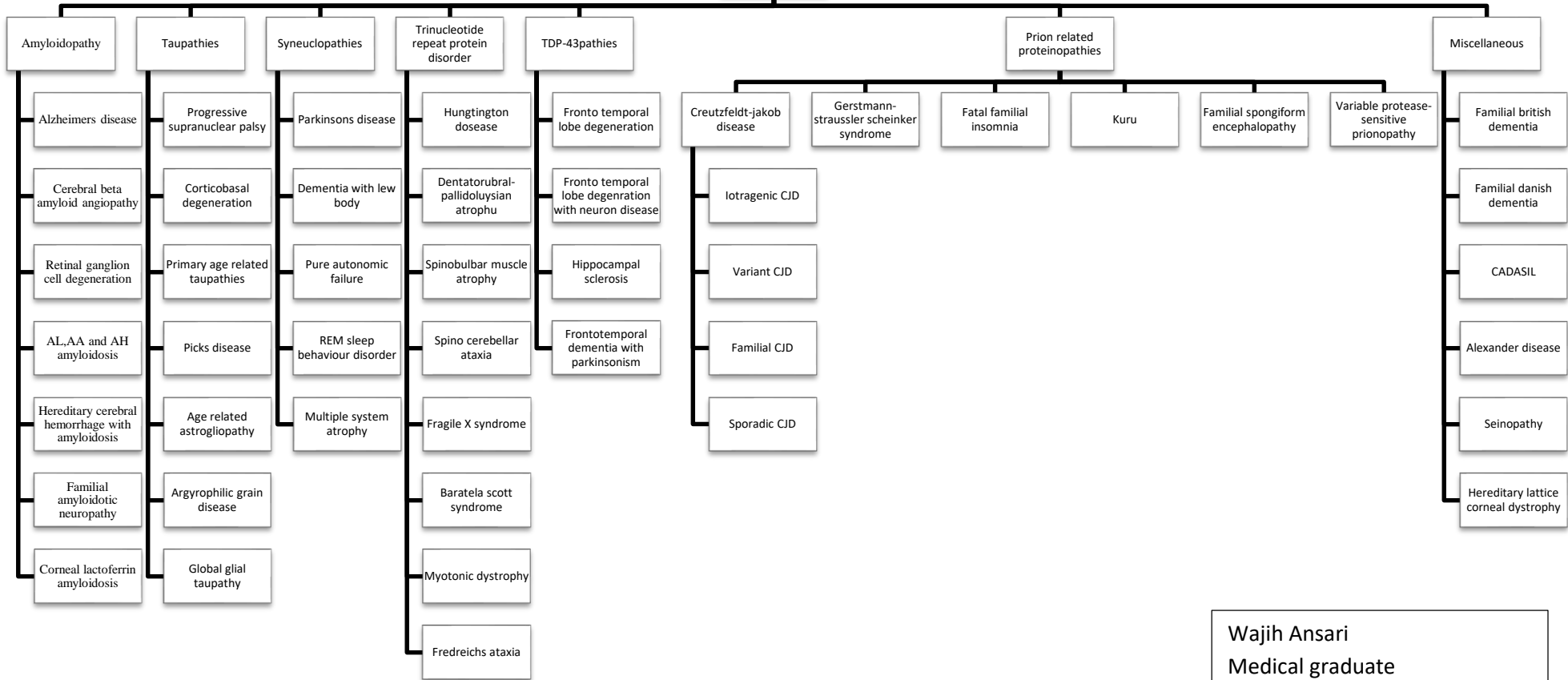- CADASIL
- Alexander disease
- Seinopathy
- Hereditary lattice corneal dystrophy

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