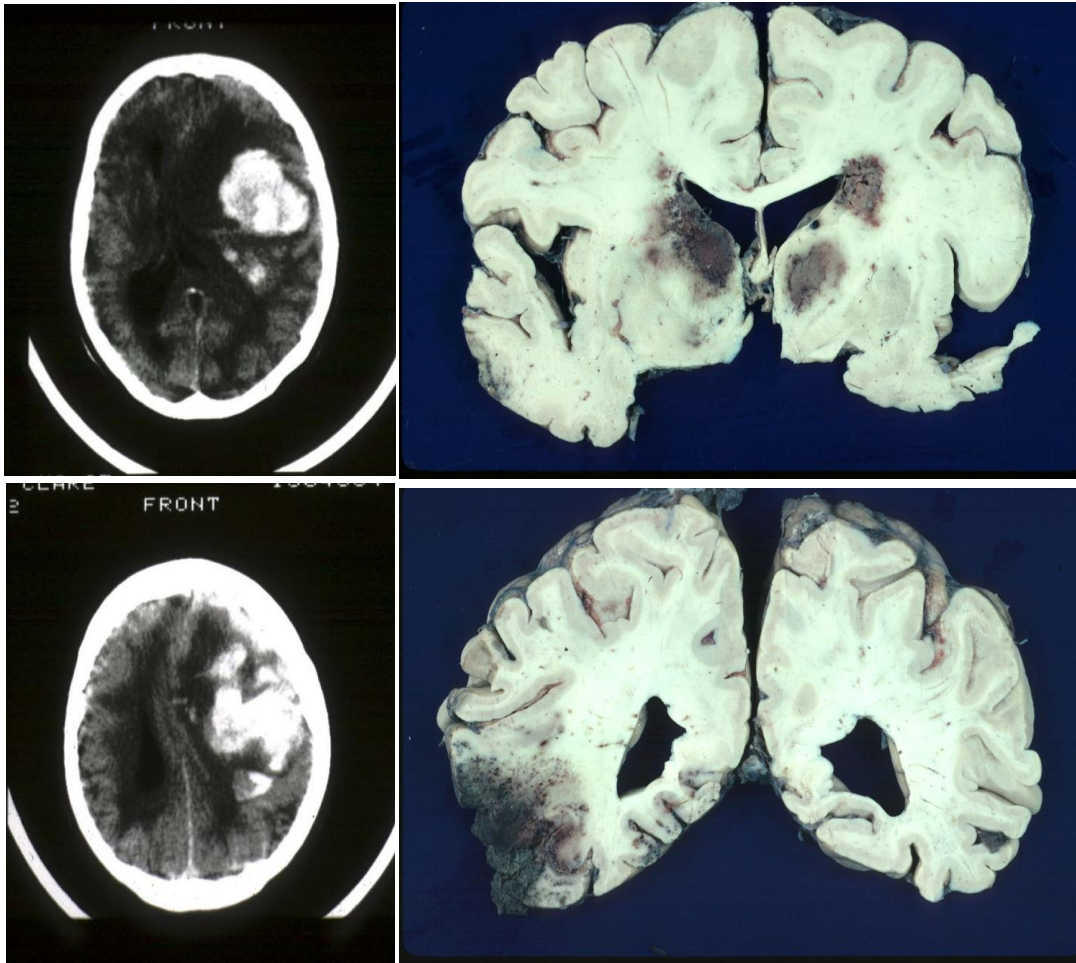


Cerebral Amyloid Angiopathies (CAA)

CAA related strokes

- **Usual vascular risks**
(↑ BP, lipids, HD risk)
- **Cognitive impairment**
- **Cerebral Haemorrhages**
- **Small vessel disease**
- **Microinfarcts**
- **WMLs**
- **Atrophy**
- **ε4 allele carriers**

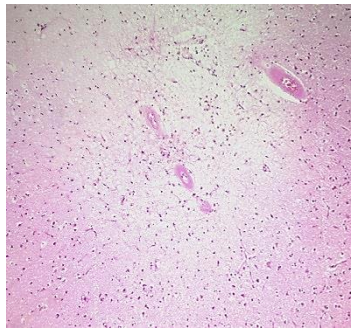
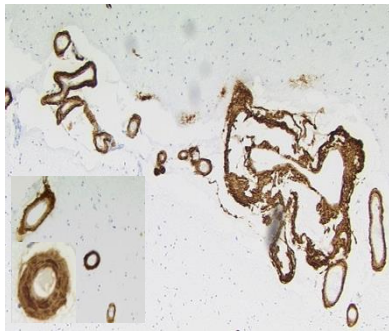
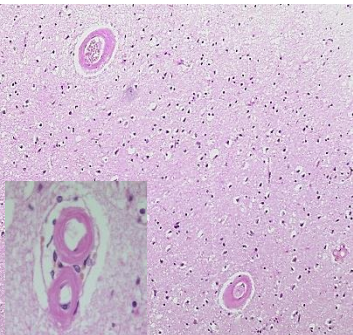
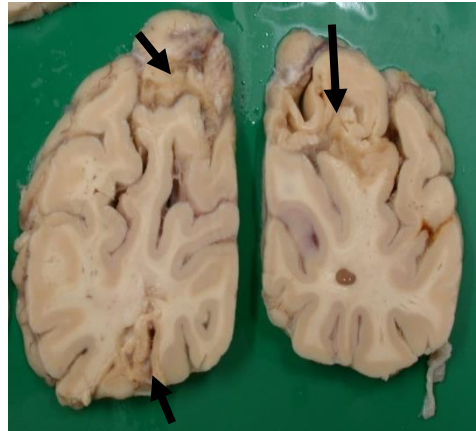


Typical case of 92 year old woman:

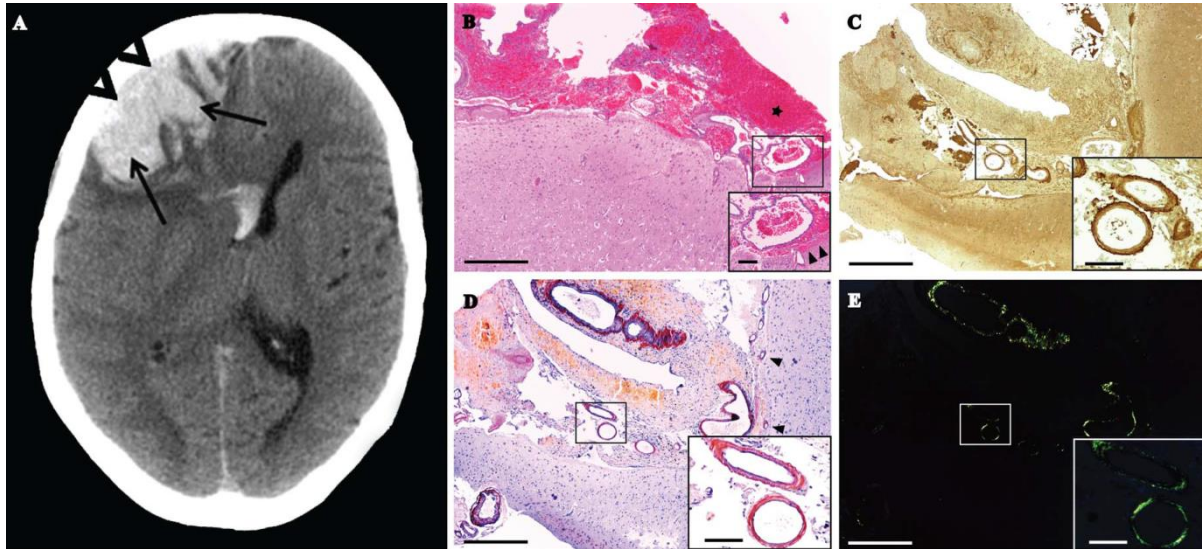
Main neuropathological findings



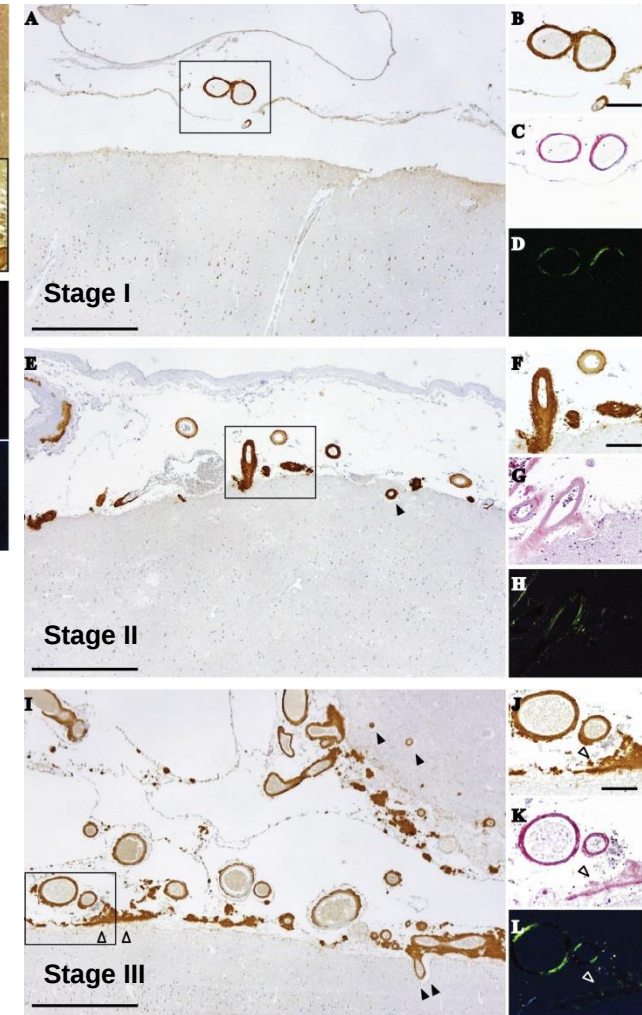
- Confusion, bilateral weakness, TIAs
- CT- infarction in the rt posterior parieto-occipital region
- Few macro- cortical infarcts
- Lobar haemorrhage
- Numerous microinfarcts
- Severe CAA
- Other findings:
 - Sparse diffuse senile plaques.
 - Braak stage II for NFT
 - Moderate density of AGD grains in limbic cortex.
 - No α -synuclein pathology



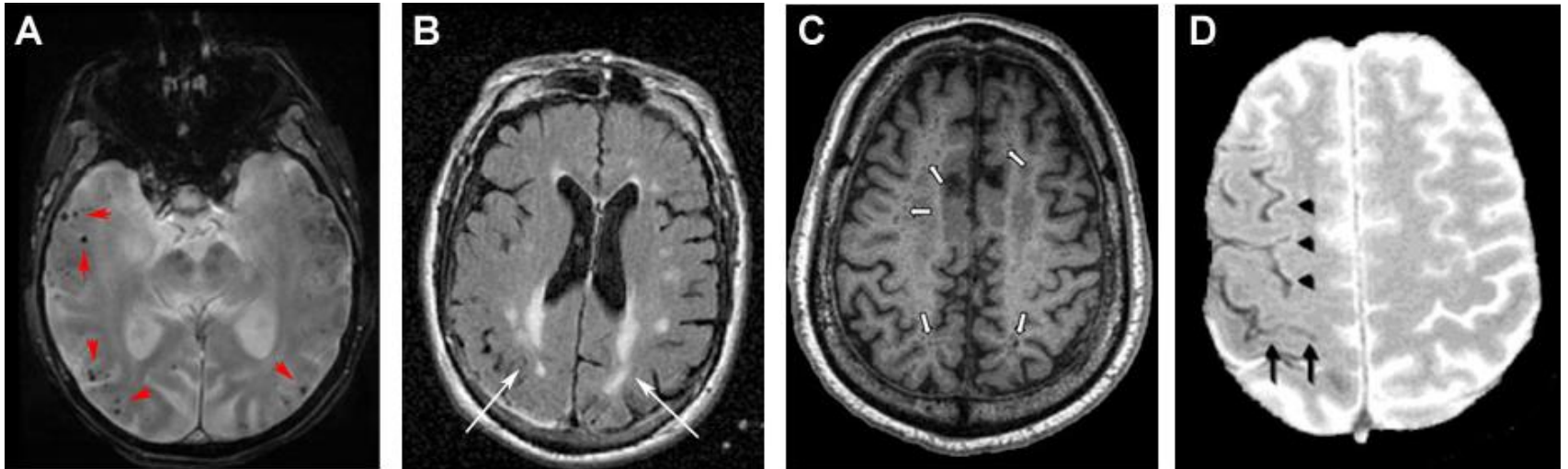
CNS patterns of Transthyretin-related amyloidosis in Familial Amyloid polyneuropathy



- Onset 35 ± 9 yrs, disease duration 8 ± 4 yrs
- CNS TTR noted 3 yrs onset of peripheral neuropathy
- Pattern: meninges and its vessels and progresses to the meningocortical arteries and subpial parenchyma
- Subpial TTR amyloid associated with astrogliosis
- No cortical microbleeds, superficial siderosis or A β IR



Radiological signature of CAA



A, lobar microbleeds on GRE images; B posterior dominant WMH; C, dilated PVS on WM T1-WI; D, multiple areas of superficial siderosis