

EJEMPLO 7



ENFERMEDAD DE CREUTZFELDT-JACOB 129VV2

DR. IVÁN FERNÁNDEZ VEGA MD,PHD

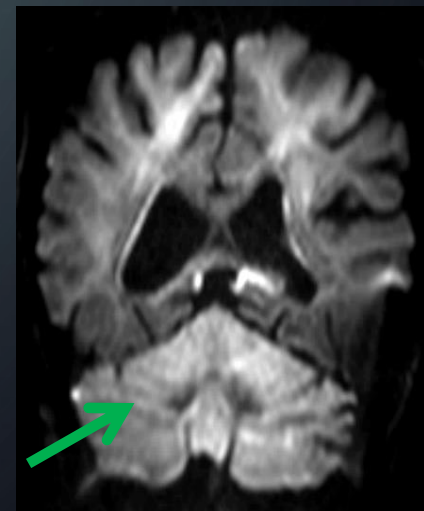
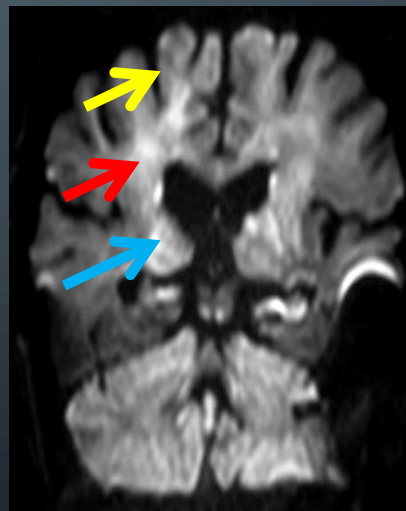
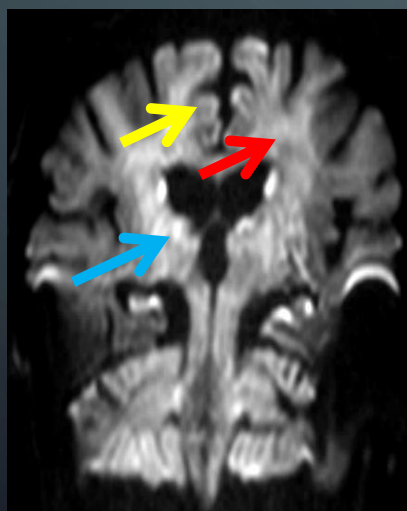
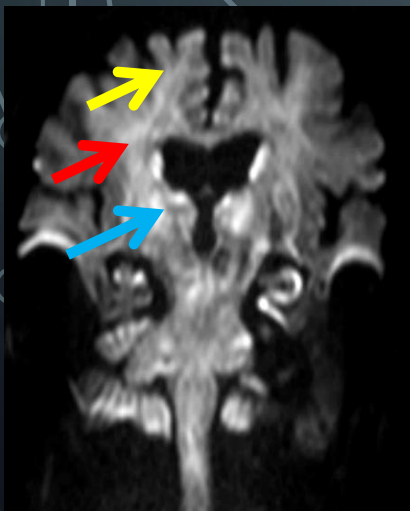
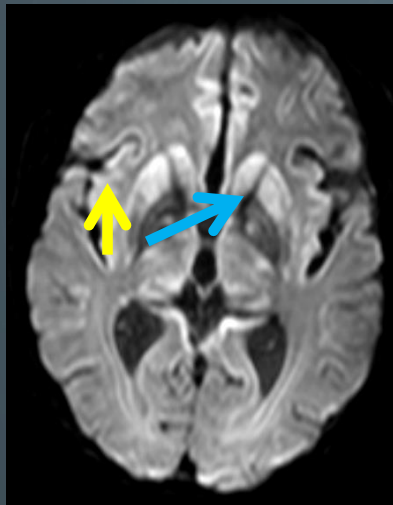
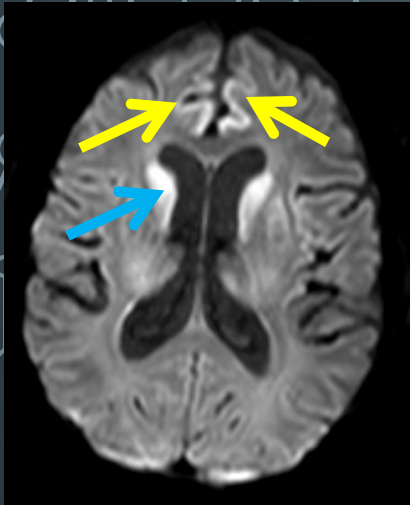
DEPARTAMENTO DE CIRUGÍA Y ESPECIALIDADES MÉDICO QUIRÚRGICAS

FACULTAD DE MEDICINA

UNIVERSIDAD DE OVIEDO

ESTUDIO RADIOLÓGICO

RESONANCIA NUCLEAR MAGNÉTICA



ESTUDIO MACROSCÓPICO

MACROSCOPÍA

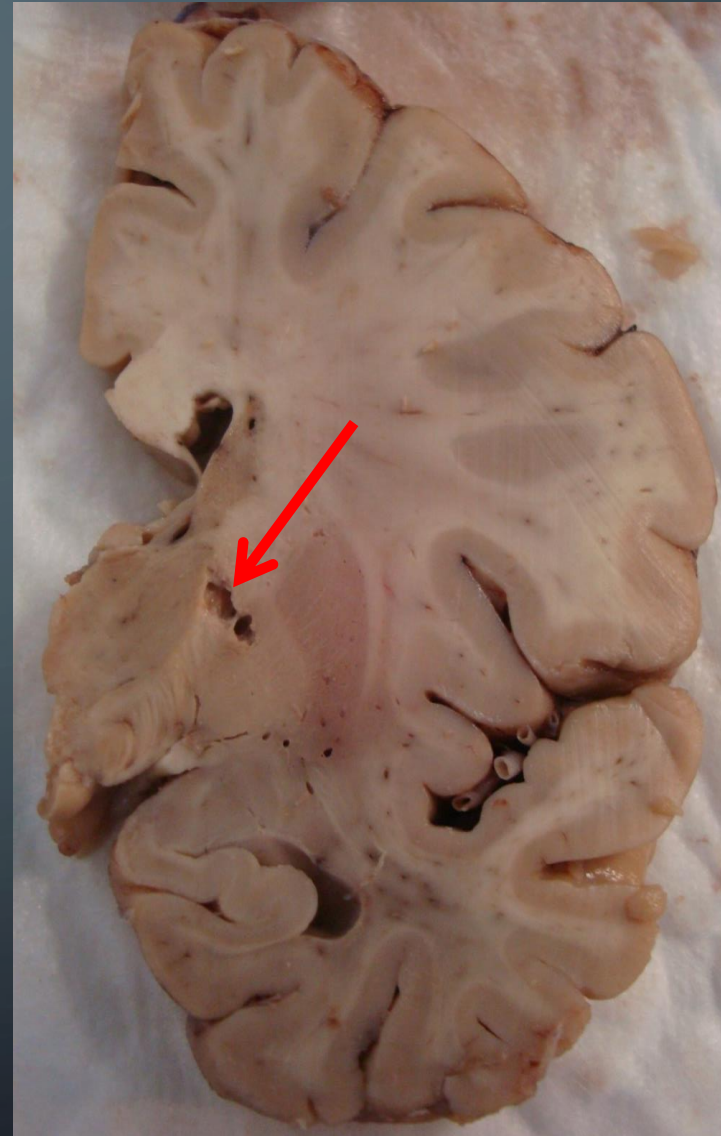
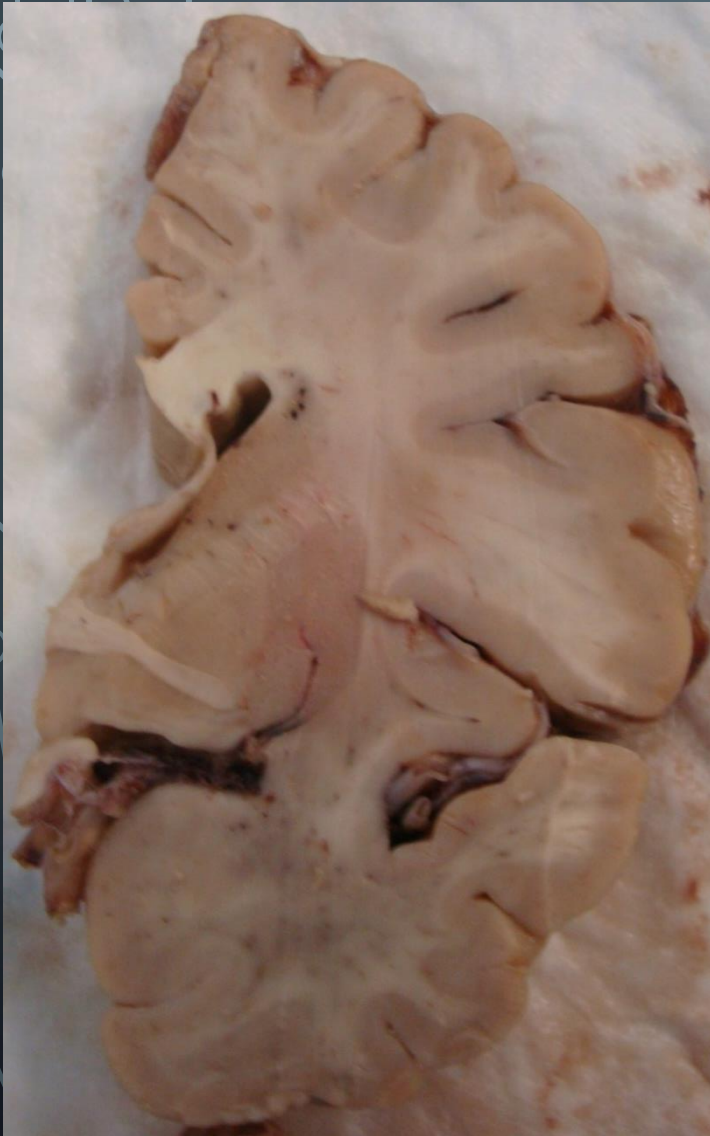


PESO completo:1400g

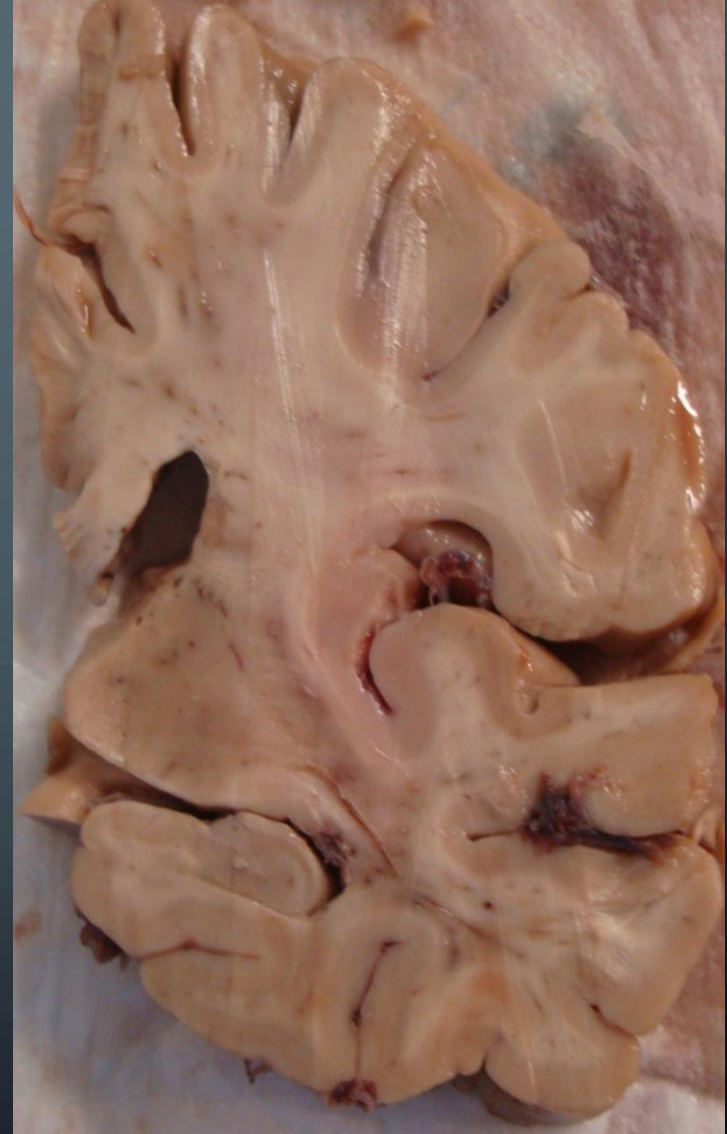
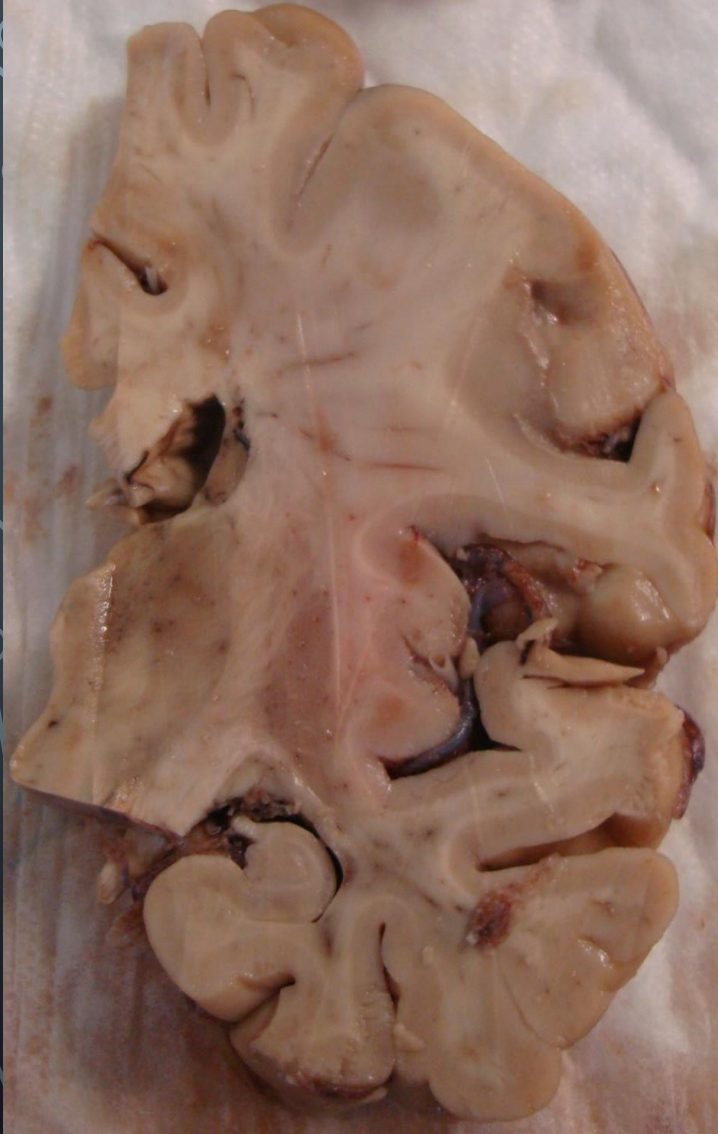
MACROSCOPÍA



MACROSCOPÍA



MACROSCOPÍA



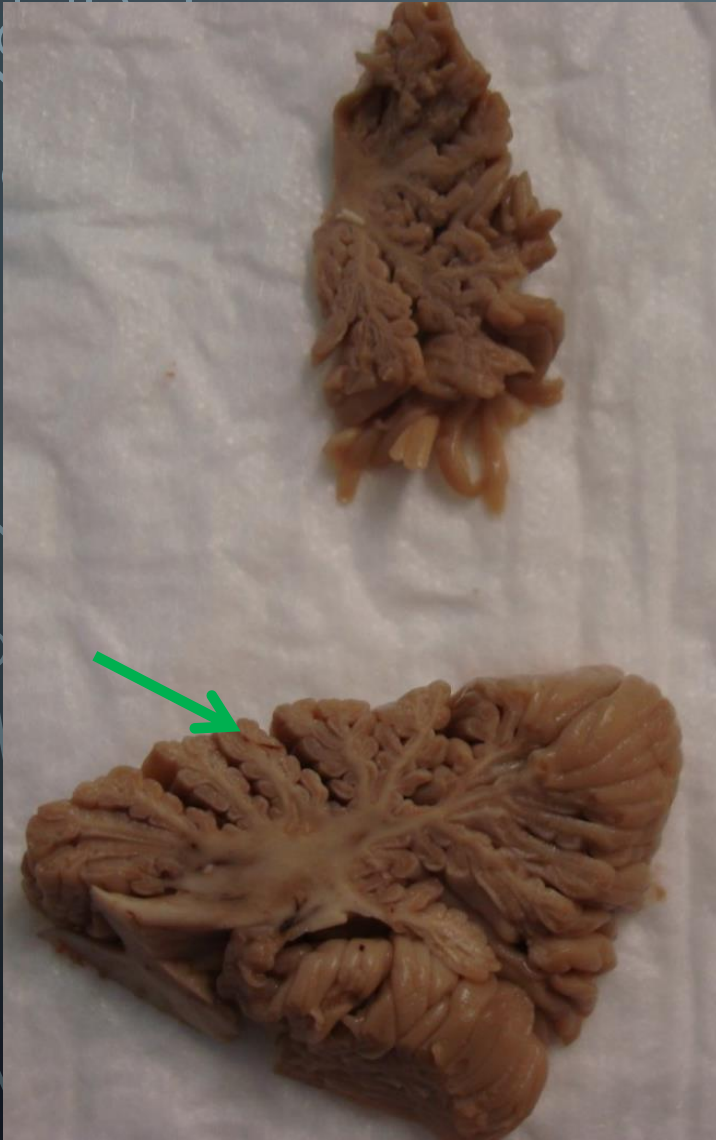
MACROSCOPÍA



MACROSCOPÍA



MACROSCOPÍA

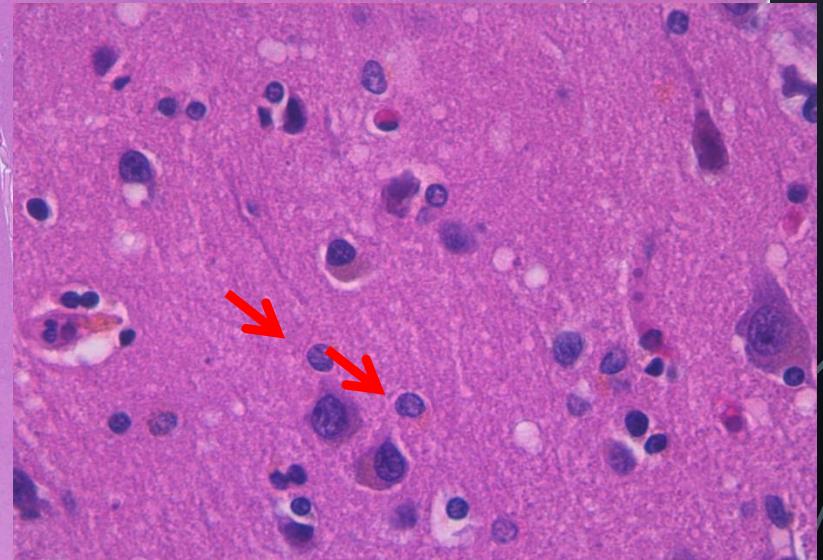
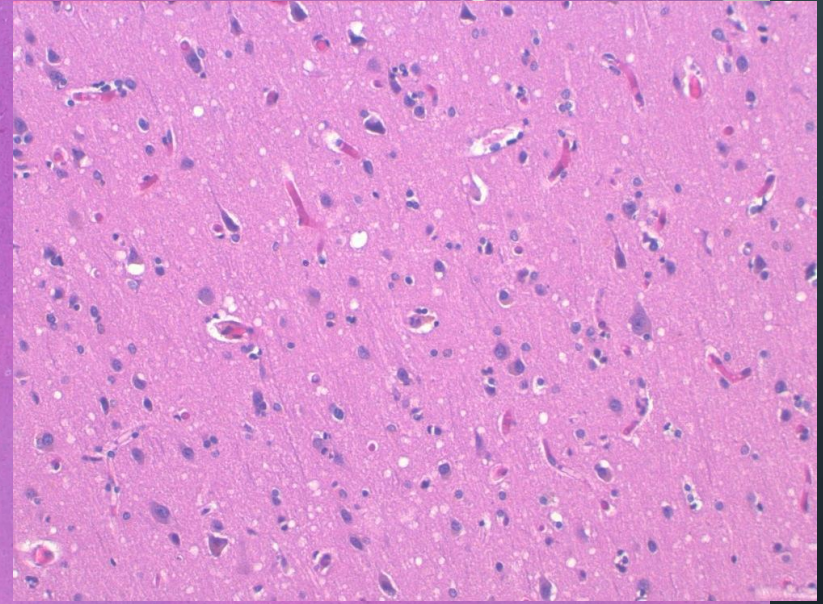
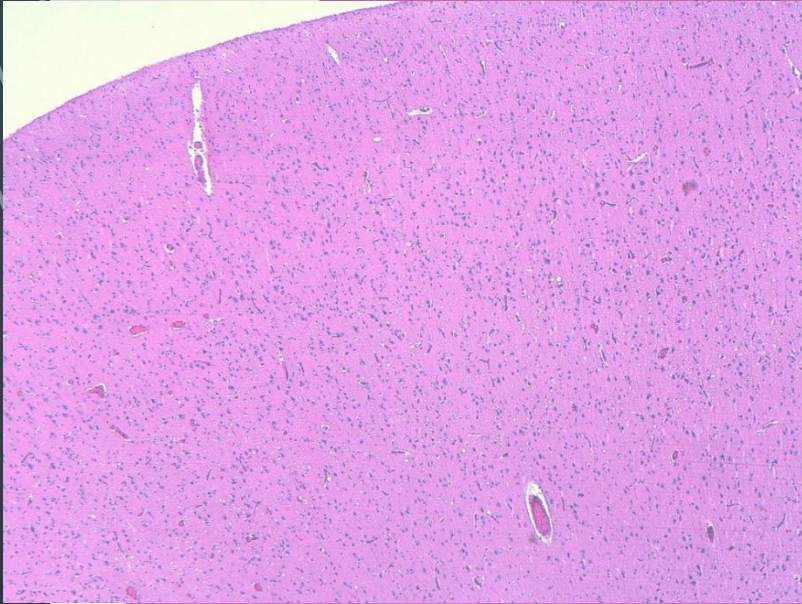


1 INTENSA ATROFIA CEREBRAL

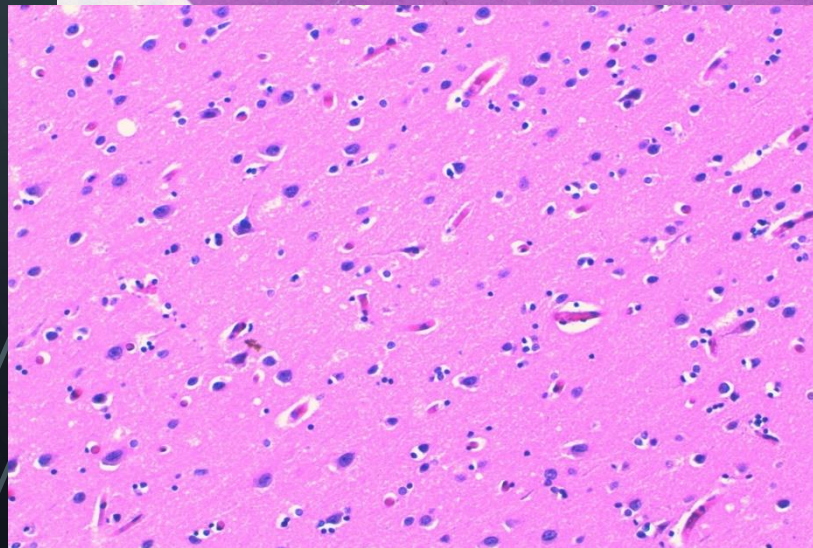
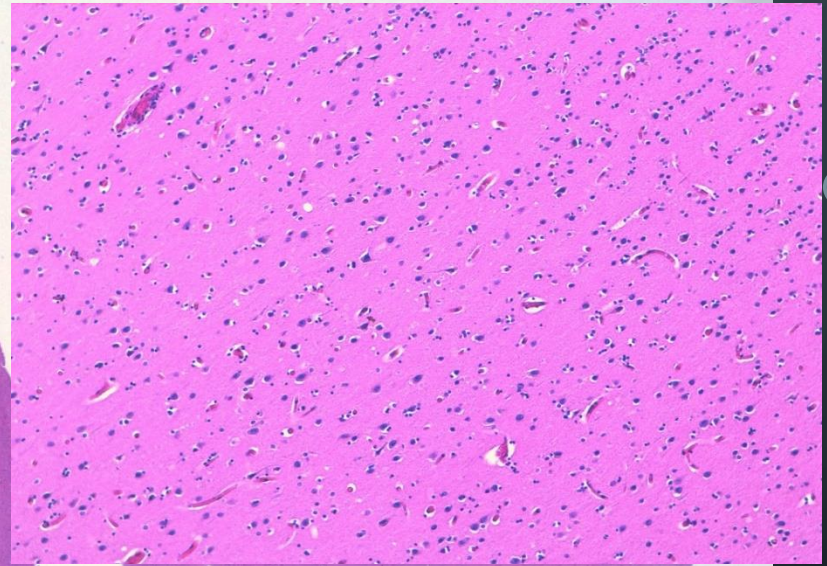
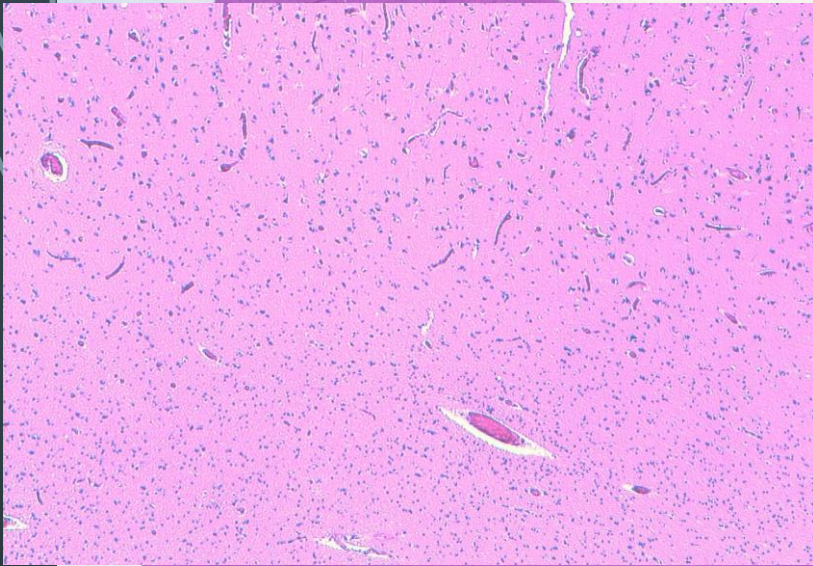
2 INFARTO LACUNAR DE 1CM EN EL NÚCLEO PÁLIDO Y PARTE DE LA CÁPSULA BLANCA INTERNA DERECHA

ESTUDIO MICROSCÓPICO

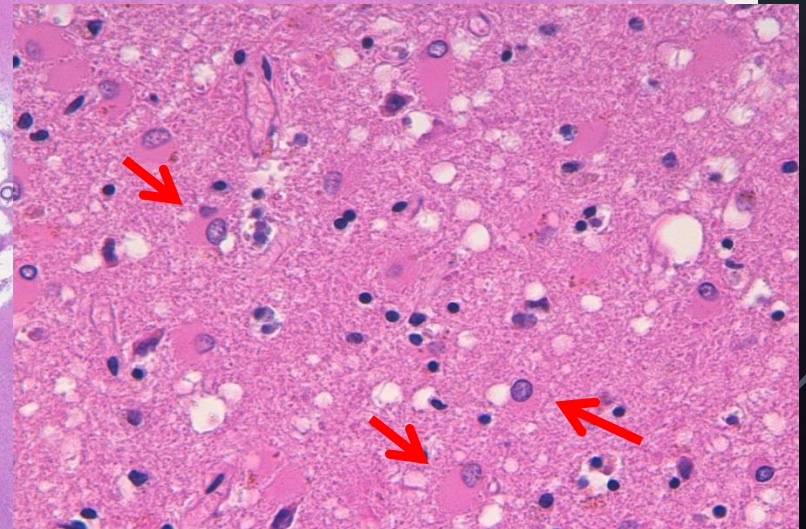
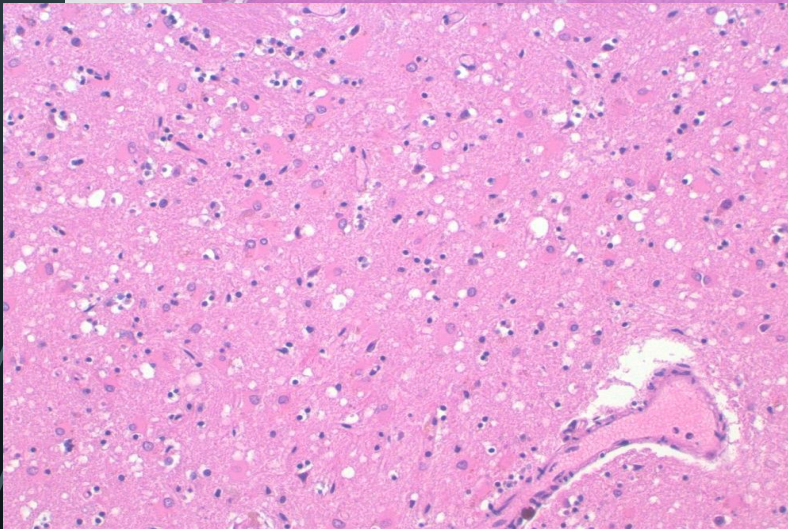
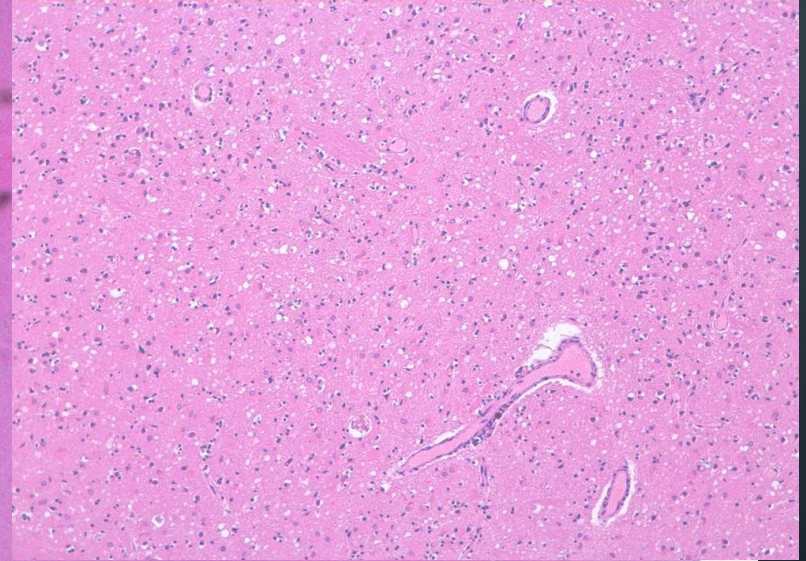
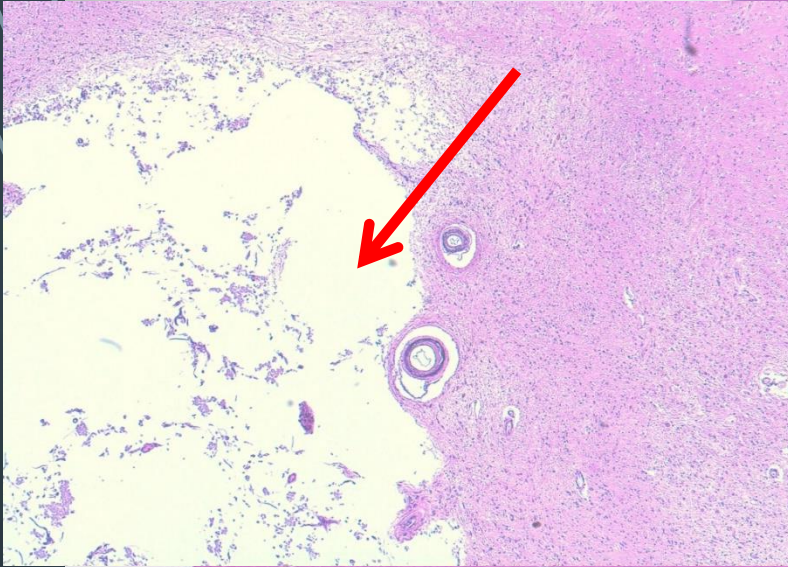
MICROSCOPÍA: CIRCUNVOLUCIÓN FRONTAL MEDIA



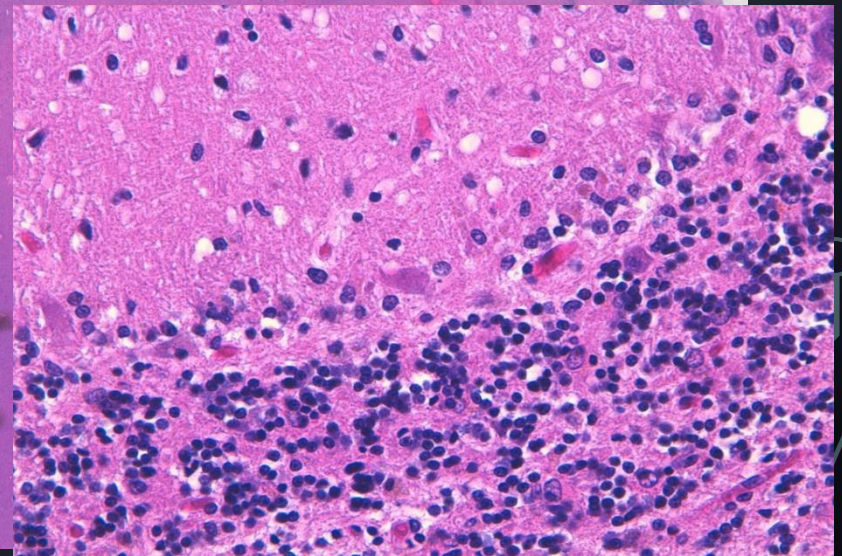
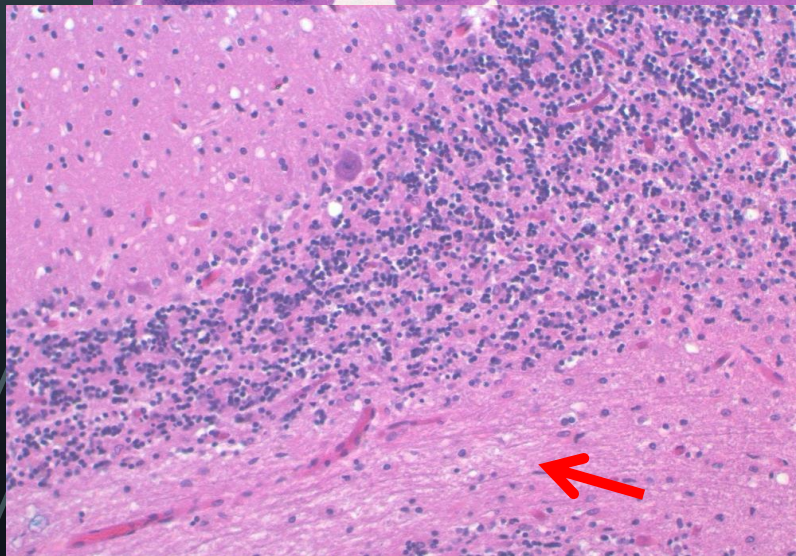
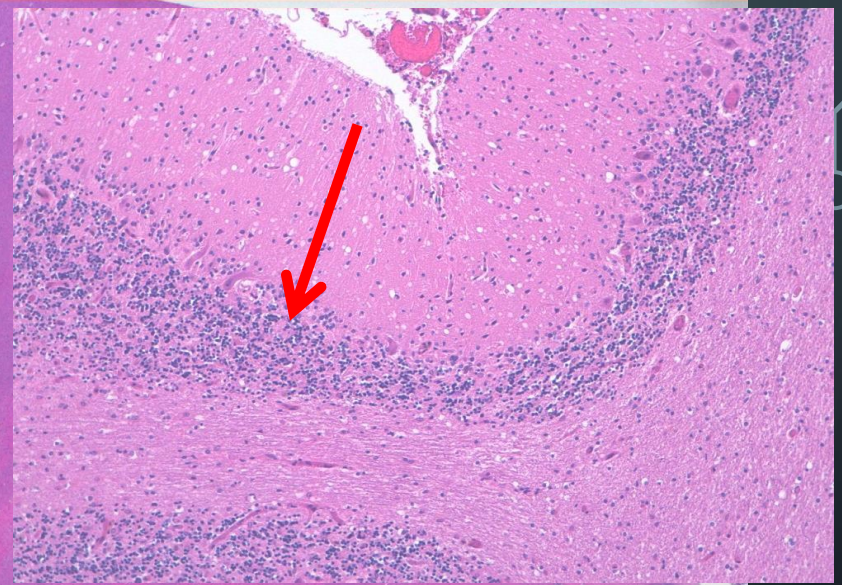
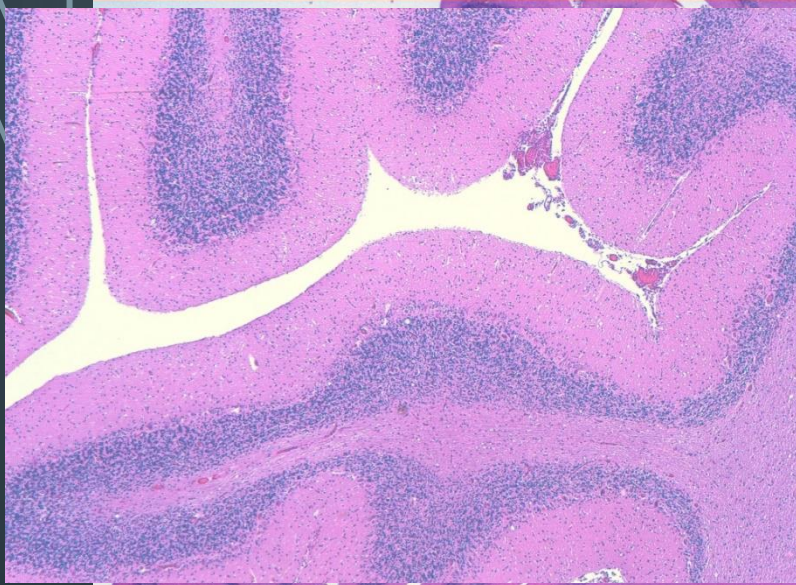
MICROSCOPÍA: CIRCUNVOLUCIÓN TEMPORAL MEDIA Y SUPERIOR



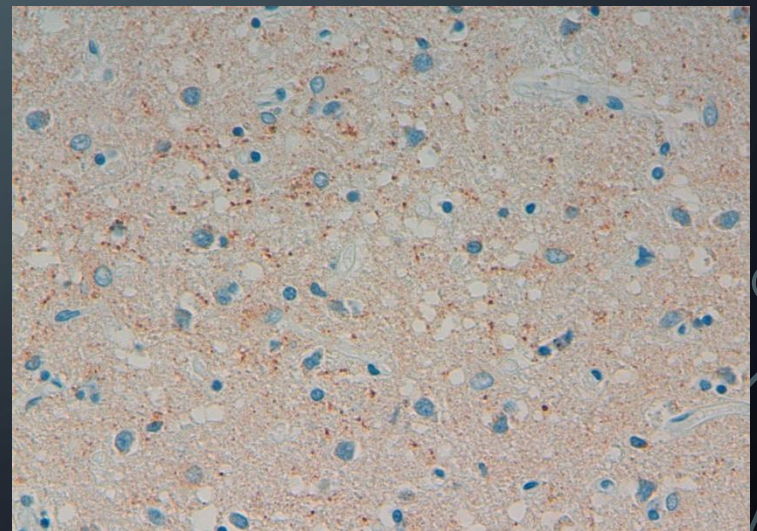
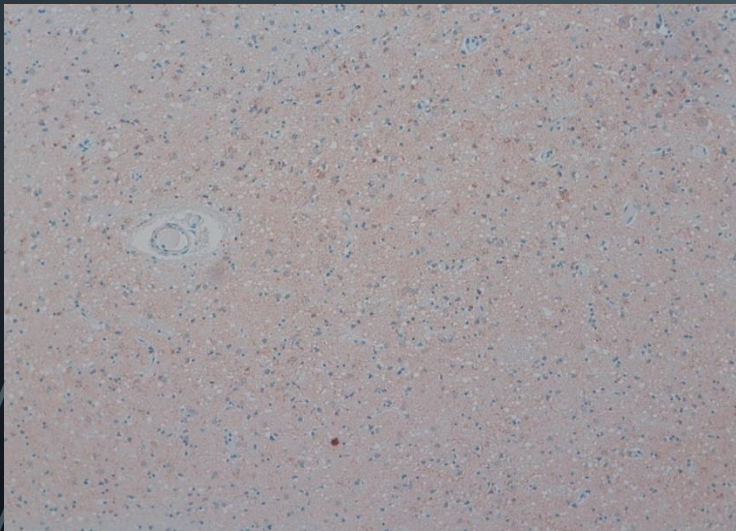
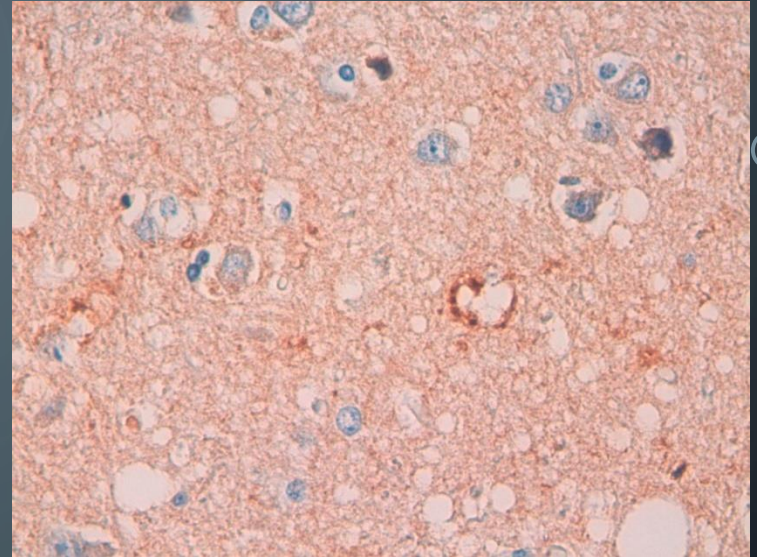
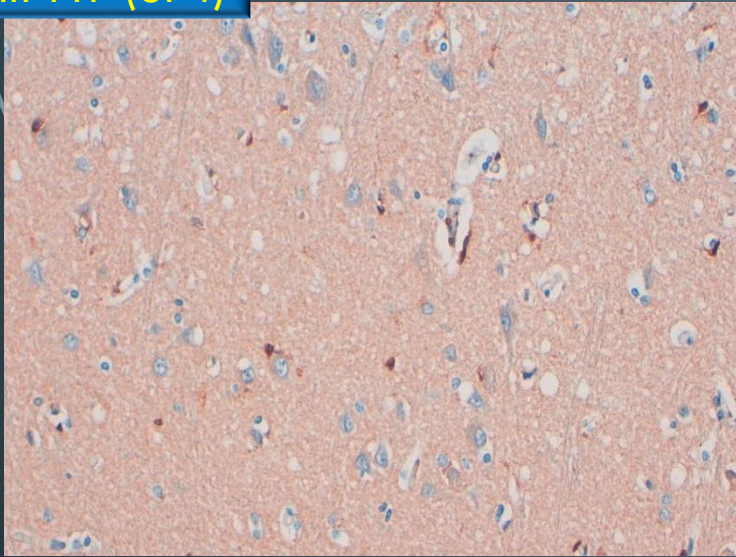
MICROSCOPÍA: NÚCLEO PÁLIDO



MICROSCOPÍA: CEREBELO

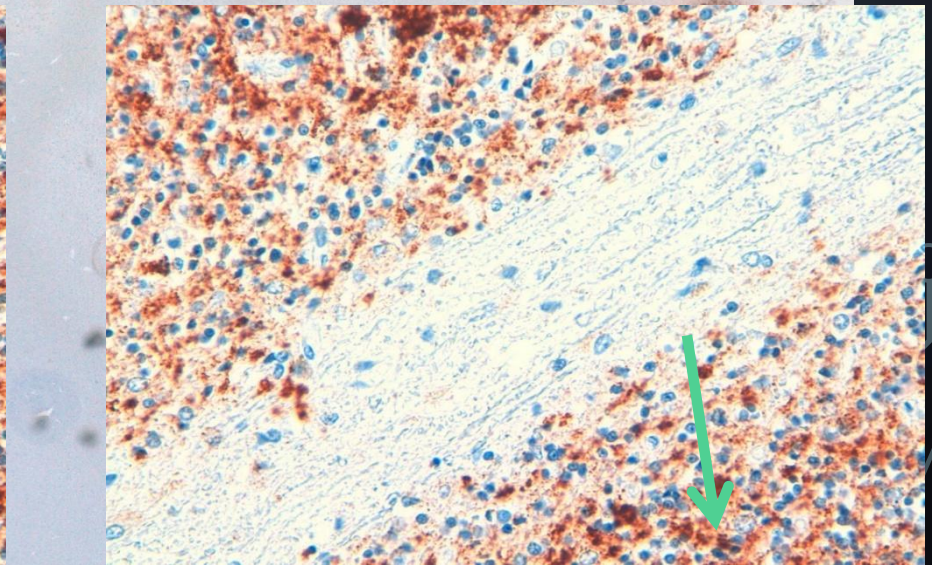
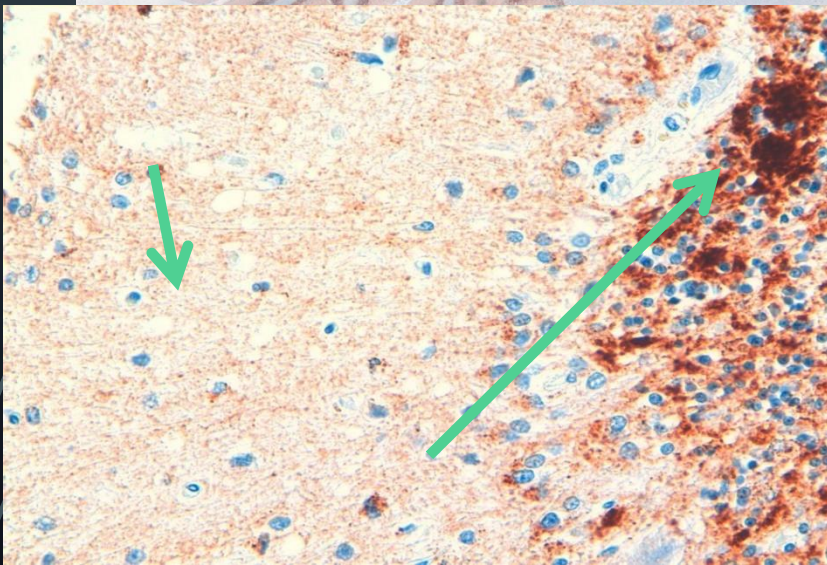
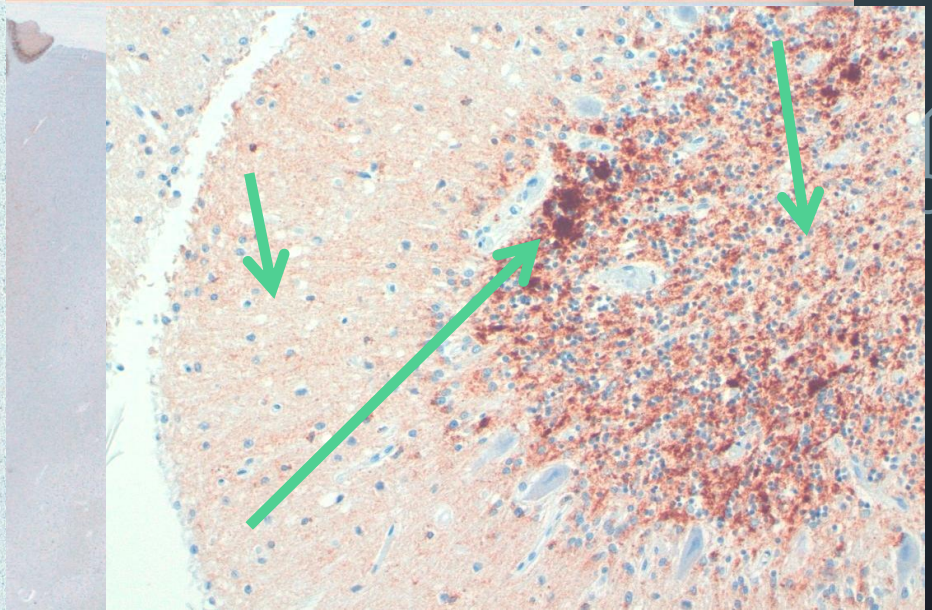
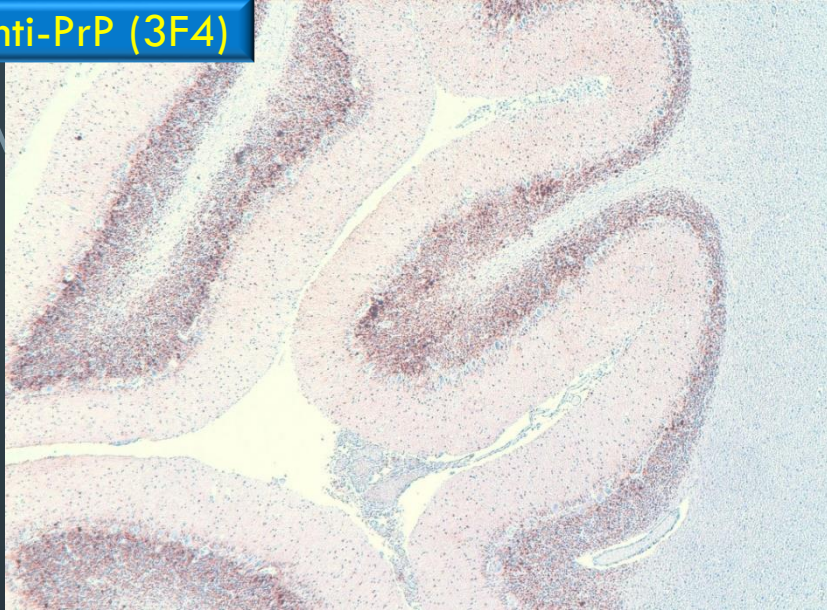


Anti-PrP (3F4)



MICROSCOPIA: INMUNOHISTOQUÍMICA

Anti-PrP (3F4)



1

CEREBRO

CAMBIO ESPONGIFORME: Degeneración pan cortical microvacuolar (principalmente en los lóbulos frontal, temporal y parietal). Cambio espongiforme microvacuolar difuso afectando al estriado predominantemente.

MODERADA PÉRDIDA NEURONA Y ATROFIA DE REMANENTES

INTENSA GLIOSIS REACTIVA: cortical y subcortical

2

TRONCO EL ENCÉFALO: sin alteraciones destacables

3

CEREBELO: CAMBIOS ATRÓFICOS SEVEROS

CAMBIO ESPONGIFORME: degeneración microvacuolar de la capa molecular

Atrofia y pérdida de las células de Purkinje

Intensa gliosis reactiva en la sustancia blanca.

HALLAZGOS INMUNOHISTOQUÍMICAS

1

CEREBRO

Depósitos extracelulares de PrPsc: Tipo sináptico y perivacuolar.

2

TRONCO EL ENCÉFALO: no estudiado. No alteraciones destacables.

3

CEREBELO:

Depósitos extracelulares de PrPsc: Tipo sináptico, granular y en placa.

No se observaron depósitos para las principales proteínas implicadas en neurodegeneración (Tau, alfa-synuclein, TDP43, amyloid beta).

***ENCÉFALO DE 1400g CON:
ENFERMEDAD DE CREUTZFELDT-JACOB 129VV2***

