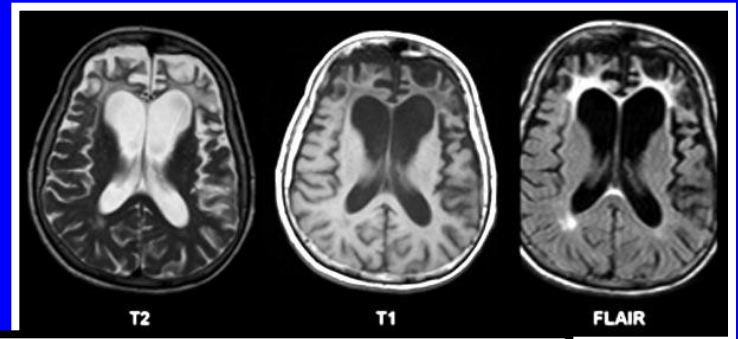
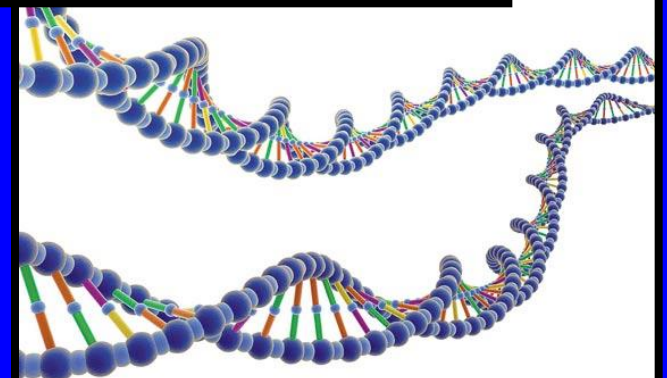
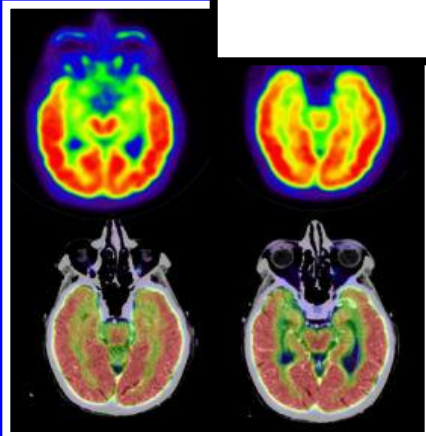


NEUROIMAGENES EN ENFERMEDADES NEURODEGENERATIVAS



**DIAGNOSTICO
MAIPU**



OBJETIVOS

- Formar al neurologista en la relación entre las demencias y la neuroimagen.
- Neuropatología de las demencias.
- Neuroimagen en las demencias.
- Neuroimagen en la práctica clínica.
- Neuroimagen en la investigación.

Journal of Neurology (2020) 267:3429–3435
<https://doi.org/10.1007/s00415-020-10040-0>

NEUROLOGICAL UPDATE

Neurological update: neuroimaging in dementia

Timothy Rittman¹ 

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Neuroimaging in Dementias

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Semin Neurol 2019;39:188–199.

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DEMENCIAS

- PRIMARIAS:
 - a. Alzheimer (50-60%)
 - b. Cuerpos de Lewy
 - c. DFT
 - d. Mixta
 - e. Parkinson, PSP, Huntington
- SECUNDARIAS:
 - a. Vascular
 - b. CJD
 - c. HPN
 - d. VIH
- DEMENCIAS RAPIDAMENTE PROGRESIVAS

Enfermedad de Alzheimer

- Lóbulo temporal mesial, cíngulo posterior y precuneo (áreas de comienzo)
- Extensión hacia corteza temporal postero-superior.
- Aproximadamente 10/15% de los casos con DCL evolucionan a EA.
- Valoración morfológica a través de RM.
- Biomarcadores (beta amiloide, TAU hiperfosforilada)
- Evaluación metabólica (PET: FDG/amiloide/tau).
- Evaluación clínica: fundamental.

PROTOCOLOS DE IMAGENES

- **RESONANCIA MAGNETICA:**
(ej:espectroscopia, RM funcional, RM-f en estado de reposo, conectividad cerebral).
- **S.P.E.C.T.**
- **P.E.T. (Beta-amiloide y Proteína Tau)**

RM Protocolo cognitivo

- Agregar a las secuencias convencionales:

A. Adquisición volumétrica T1

B. Adquisición volumétrica SWI
(susceptibilidad magnética)

C) Coronal T2 Alta Resolución (*Perpendiculares al eje mayor del Hipocampo*)

TABLA 3. ESCALA DE EVALUACIÓN VISUAL DE ATROFIA DEL LÓBULO TEMPORAL MEDIAL (MTA-SCALE). VER FIGURA 1

PUNTAJE	Ancho de fisura coroidea	Ancho del cuerno temporal	Altura del hipocampo
0	N	N	N
1	↑	N	N
2	↑ ↑	↑	↓
3	↑ ↑ ↑	↑ ↑	↓ ↓
4	↑ ↑ ↑	↑ ↑ ↑	↓ ↓ ↓

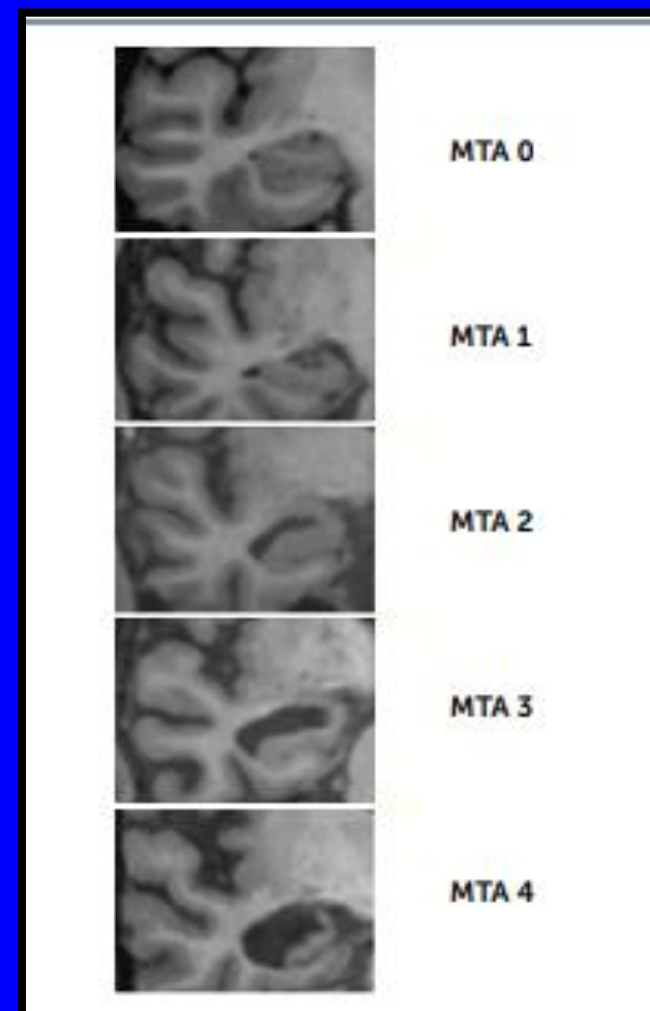


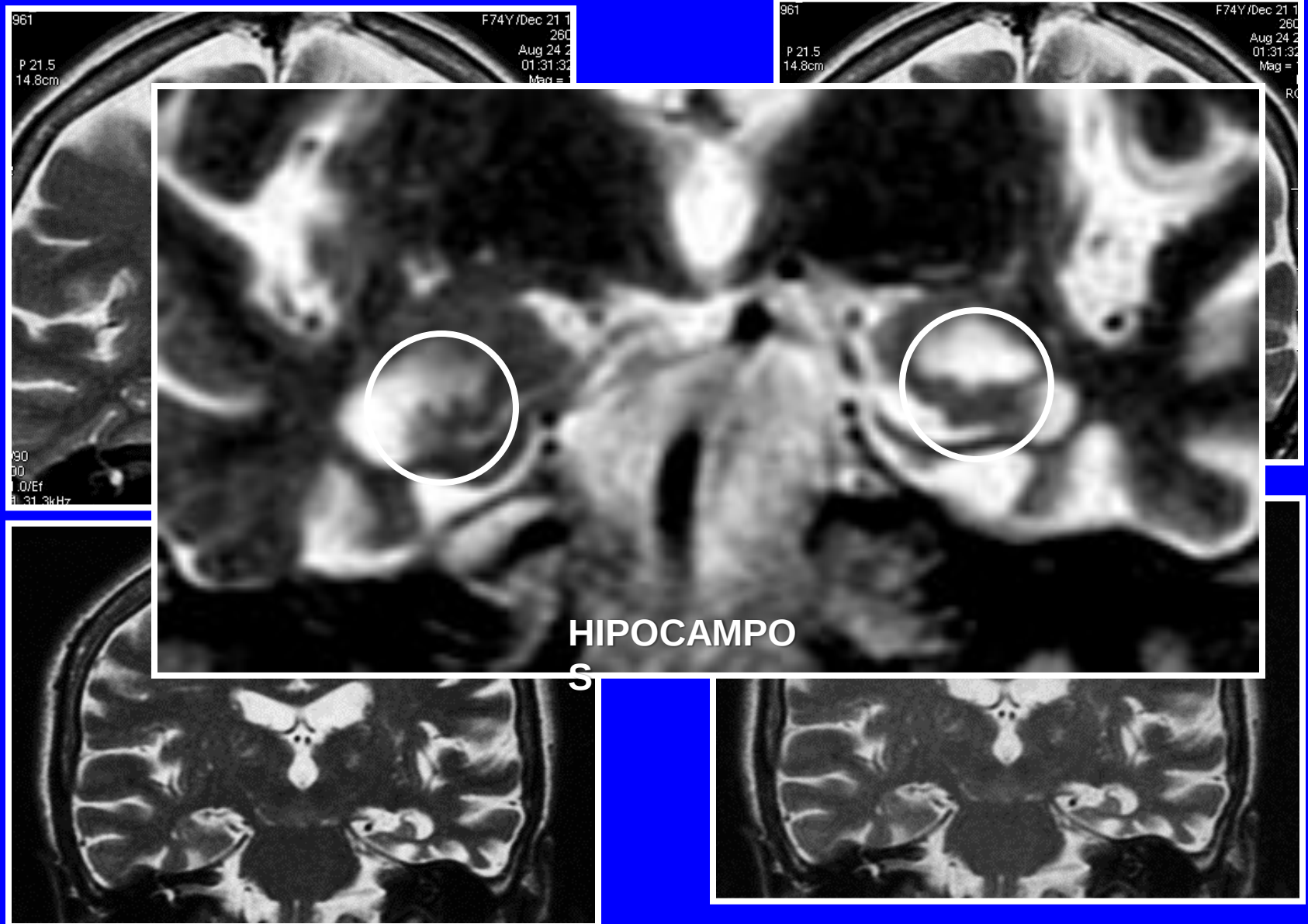
TABLA 5. ESCALA VISUAL DE ATROFIA POSTERIOR/PARIETAL A PARTIR DE MR MULTIPLANAR

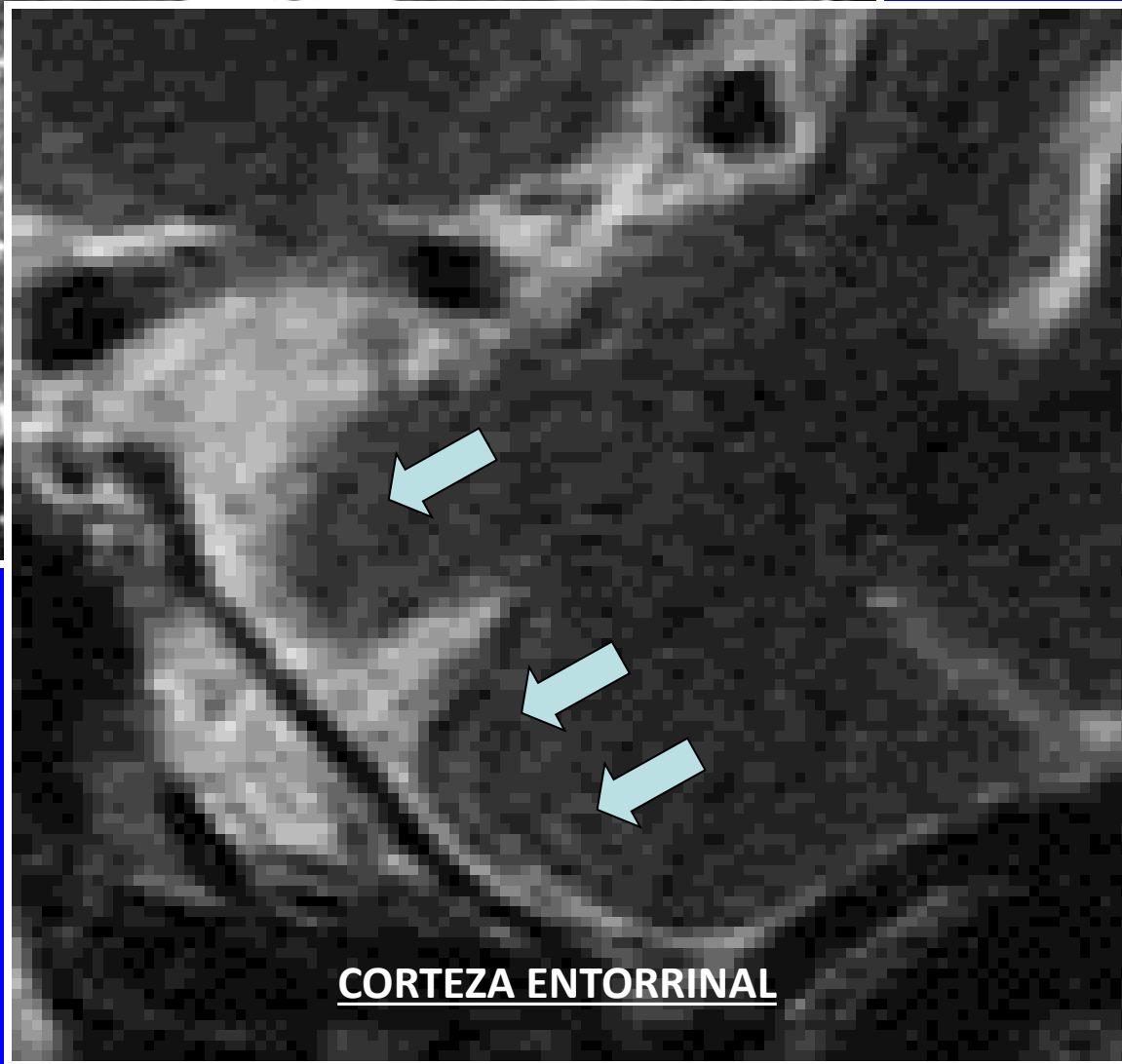
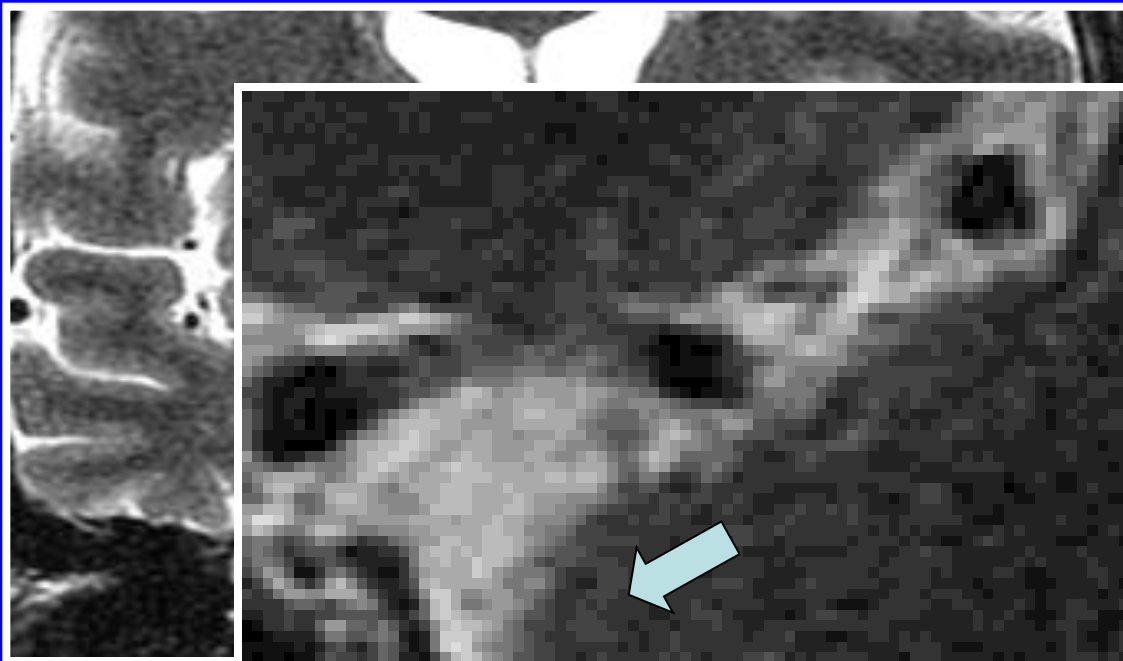
	SAGITAL	AXIAL	CORONAL
PUNTAJE	Ancho del surco cingulado posterior y el surco parieto-occipital, atrofia del precúneo	Amplitud del surco cingulado posterior y dilatación surcal en lóbulos parietales	Amplitud del surco cingulado posterior y dilatación surcal en lóbulos parietales
0	Normal	Normal	Normal
1	↑	↑	↑
2	↑↑	↑↑	↑↑
3	↑↑↑	↑↑↑	↑↑↑

TABLA 6. ESCALA DE EVALUACIÓN VISUAL DE HIPERINTENSIDADES DE SUSTANCIA BLANCA (FAZEKAS, 20)

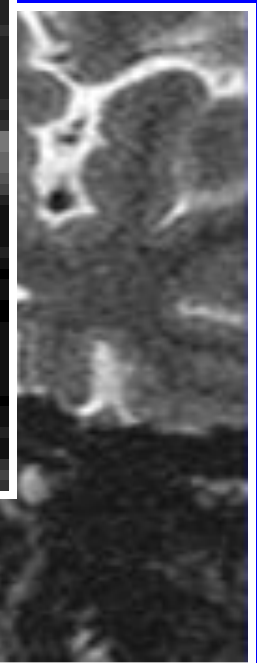
PUNTAJE	Característica
0	Ninguna lesión o una lesión única punteada
1	Lesiones punteadas múltiples
2	Comienzo de confluencia de lesiones (puentes)
3	Lesiones grandes confluentes

Deterioro cognitivo leve tipo amnésico

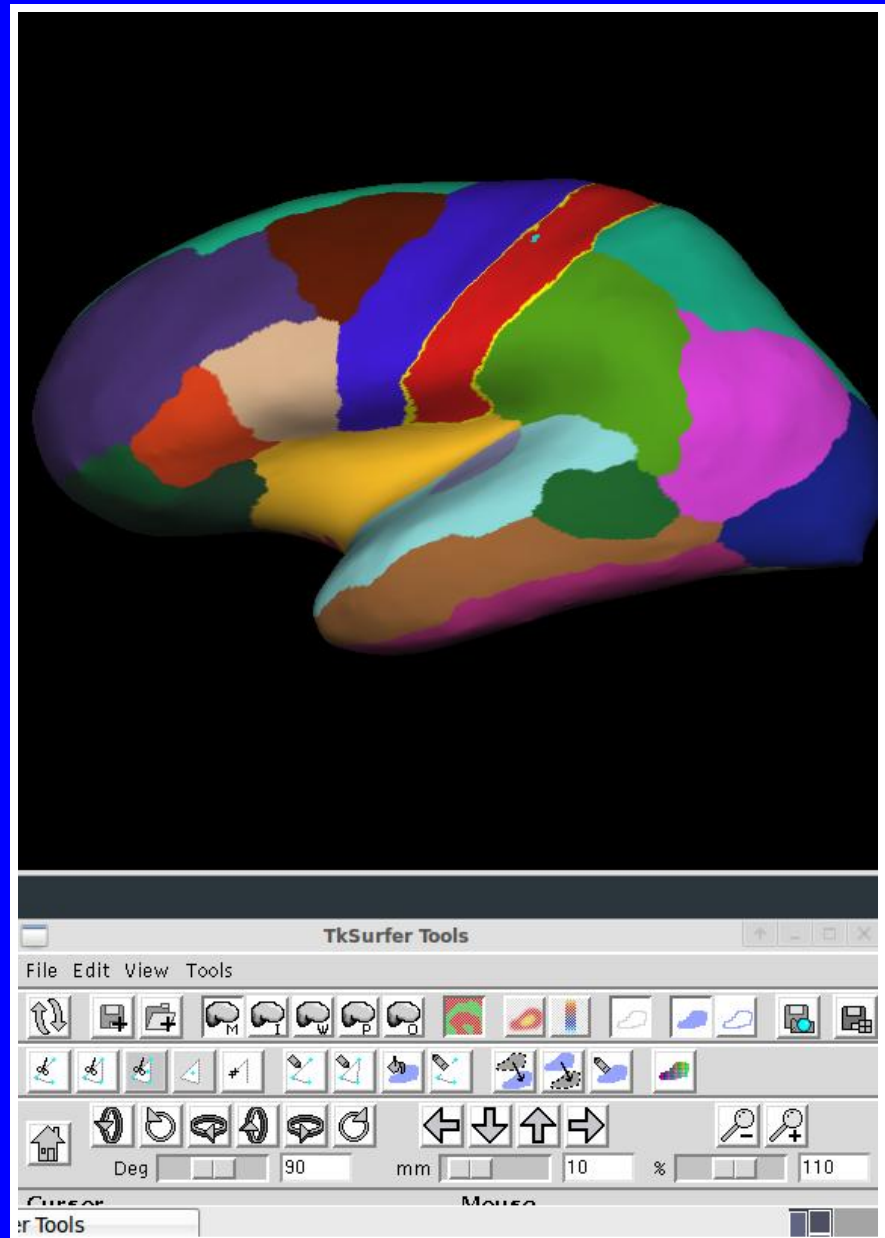
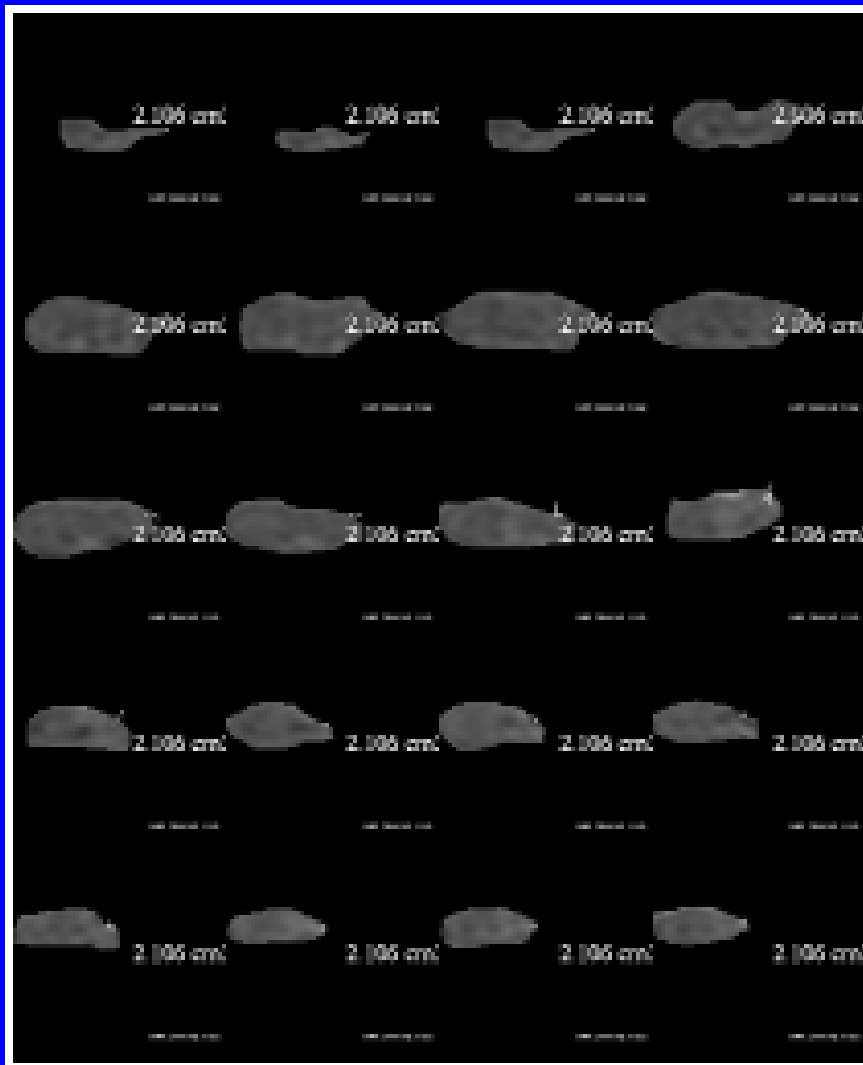




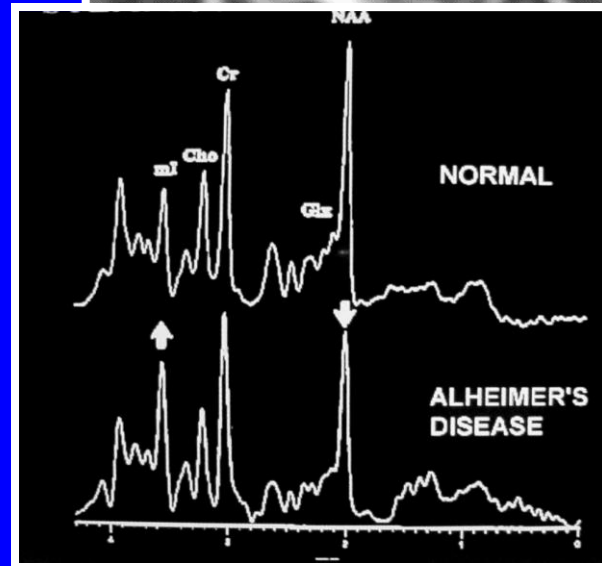
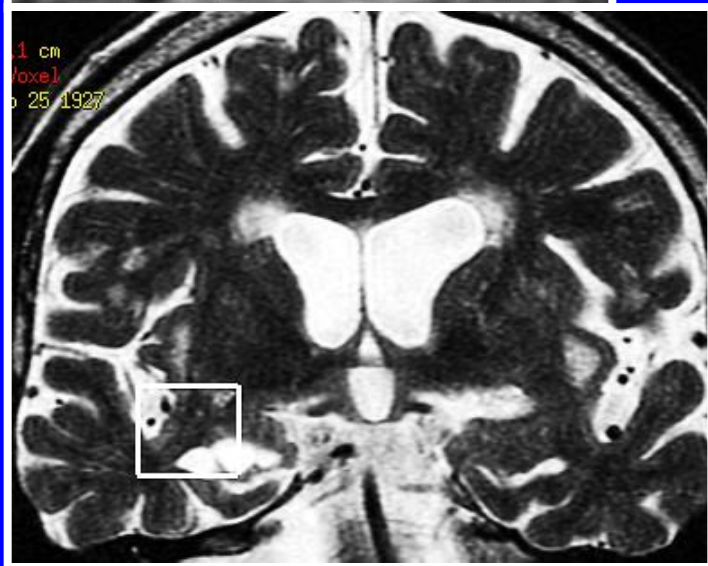
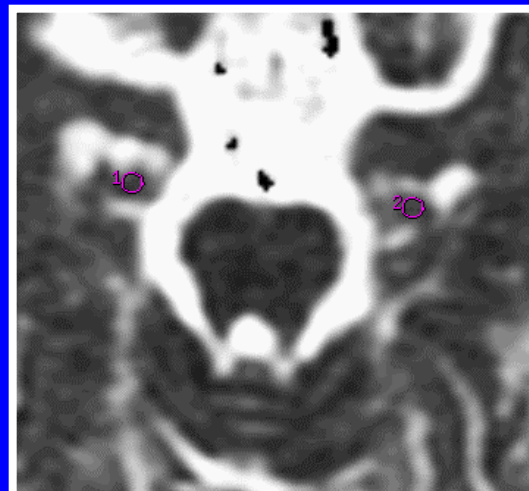
CORTEZA ENTORRINAL



Método semi-automático

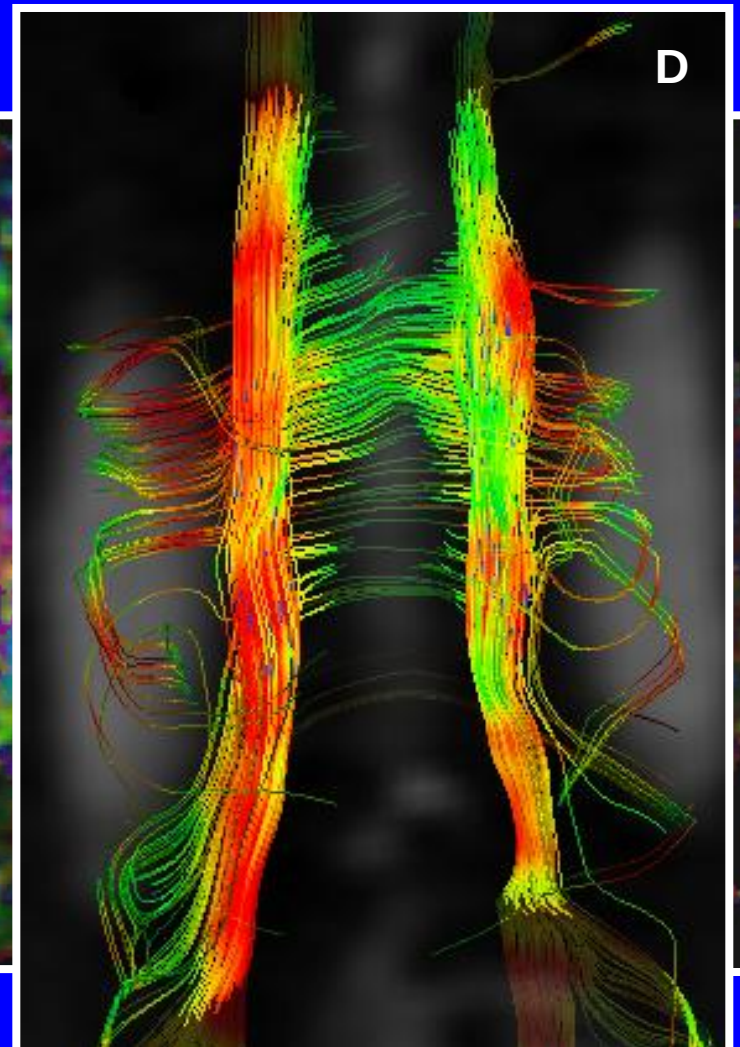
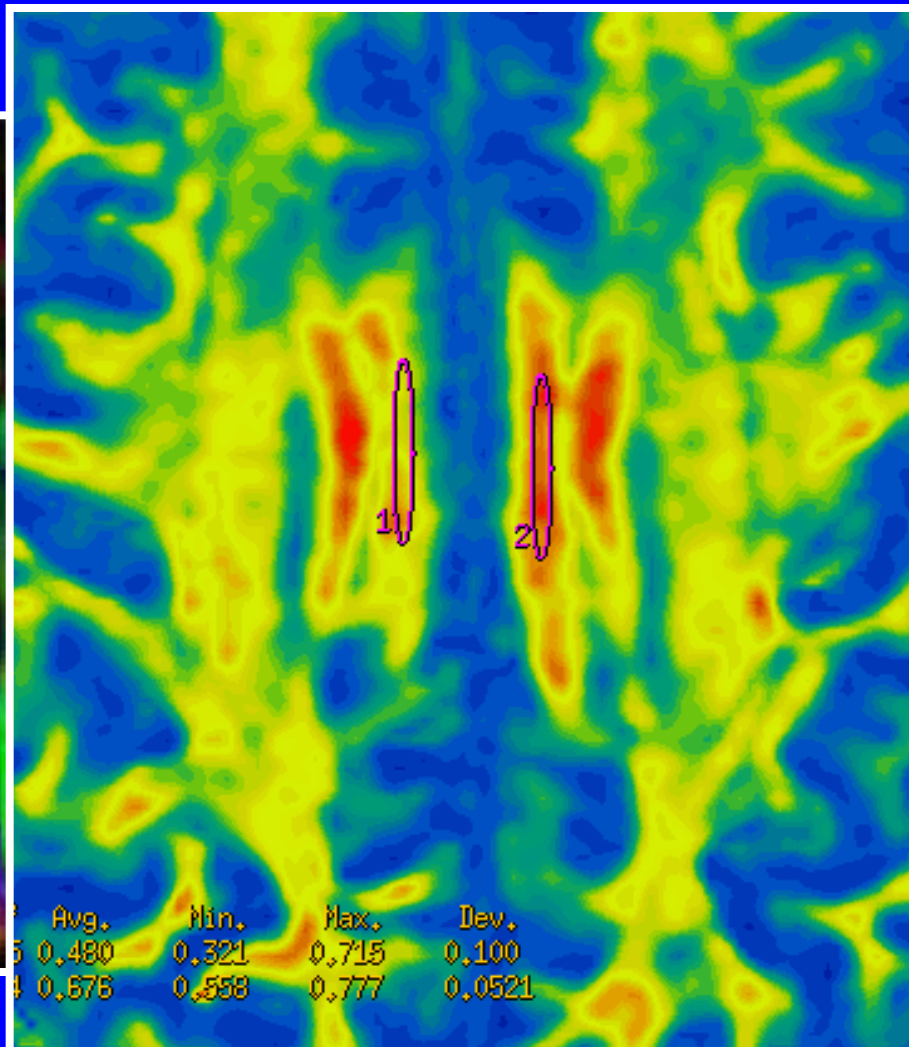


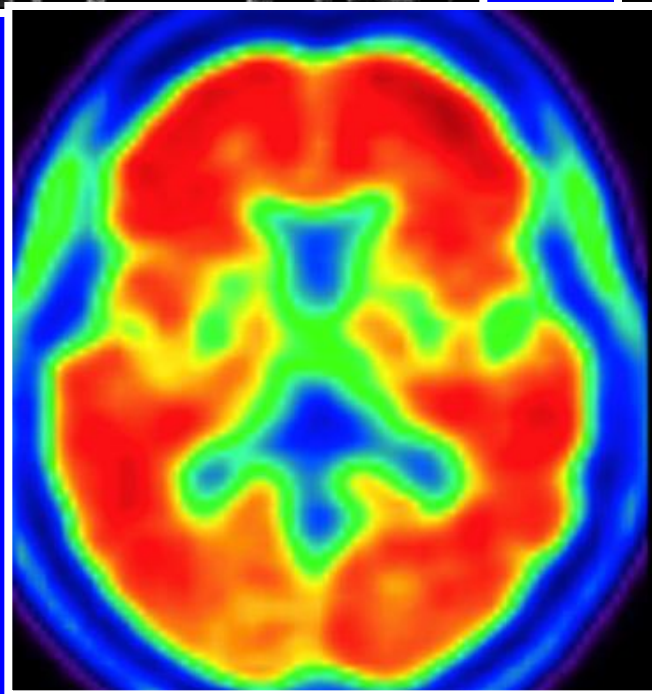
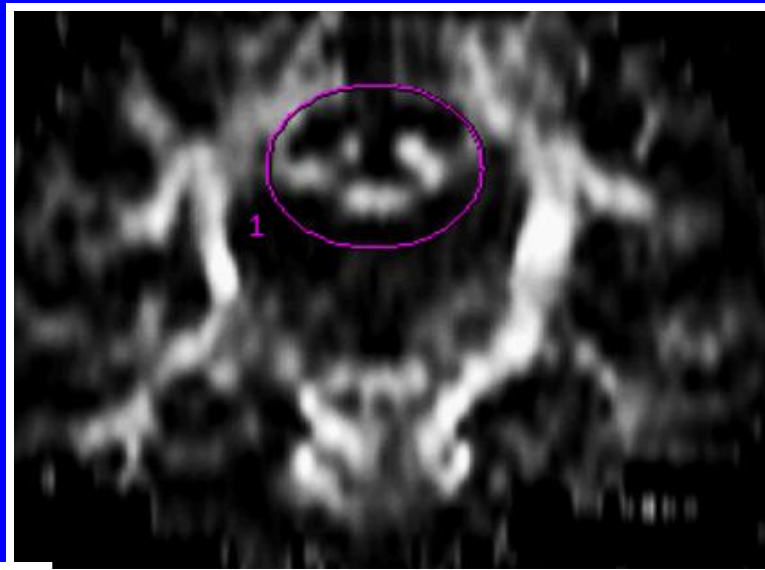
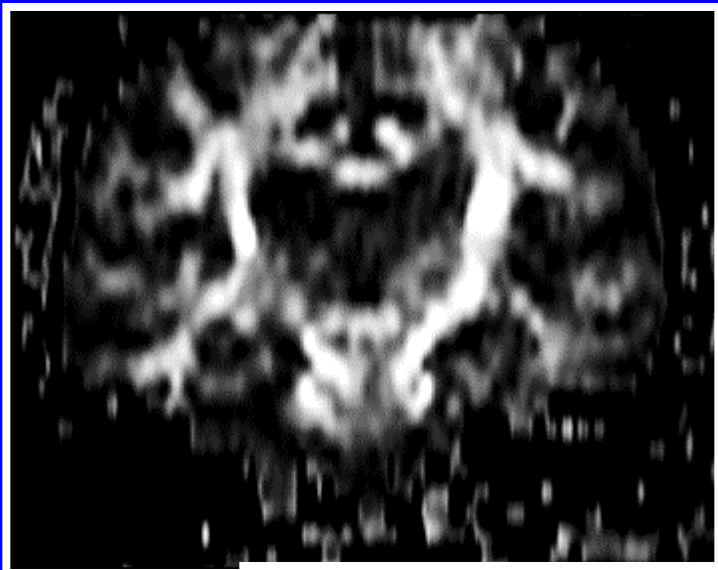
DWI y ADC



ESPECTROSCOPIA

DTI (Tensor de Difusión)





C-Pib-PET (b-AMILLOIDE)

Enf Alzheimer:

Estructural:

Atrofia Lob. Temporal mesial (CA2 y CA3 del hipocampo), cíngulo posterior y lóbulo parietal pósteromedial.

Molecular:

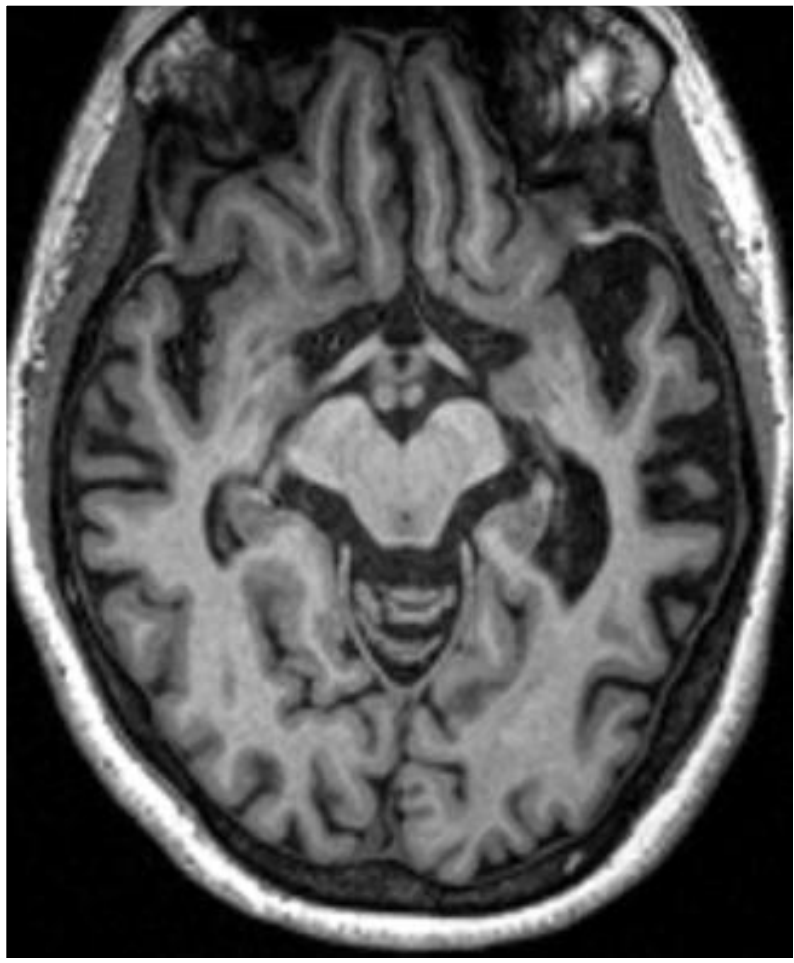
SPECT (Perfusión disminuida) **y PET** (Menor captación de glucosa).

Ambos en lóbulo temporal mesial (hipocampo)
C-PiB Florbetapir (captación por placas de amiloide)

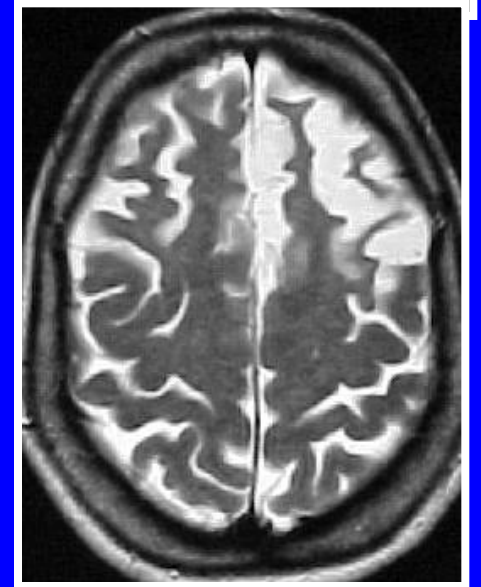
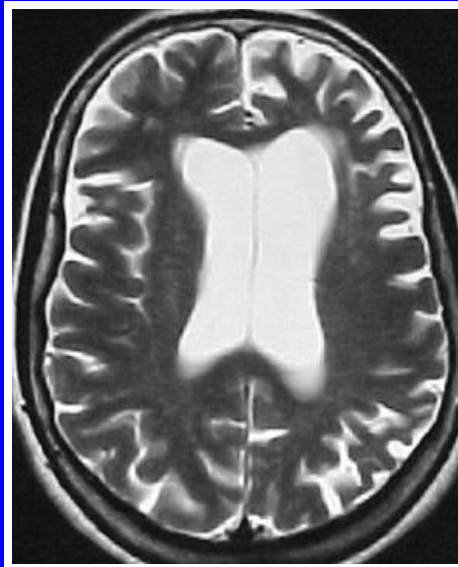
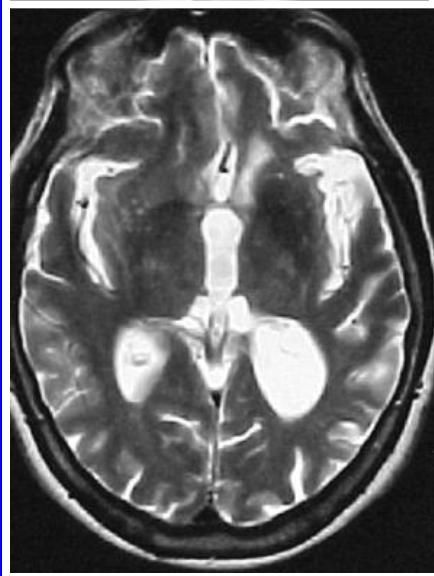
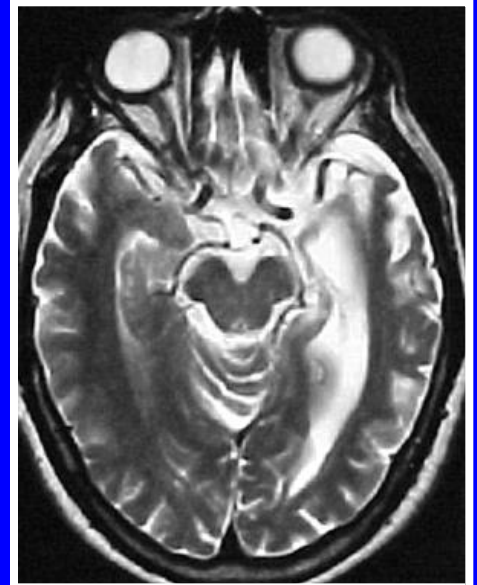
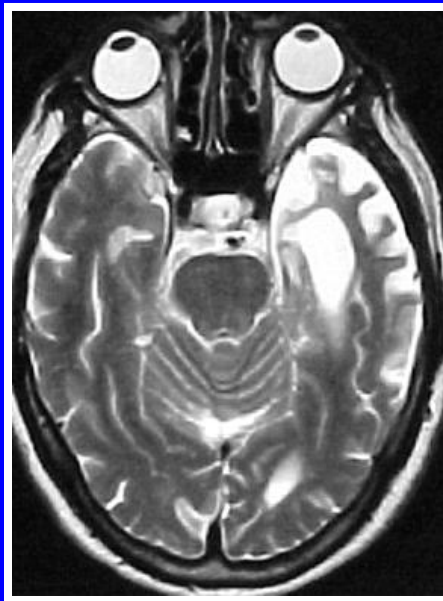
Demencia fronto-temporal

- CLINICA:
- A. Variante conductual (bvFTD)
- B. Afasia progresiva primaria (PPA)
- b.1) Afasia progresiva NO fluente
- b.2) APP semántica
- b.3) Afasia logopénica

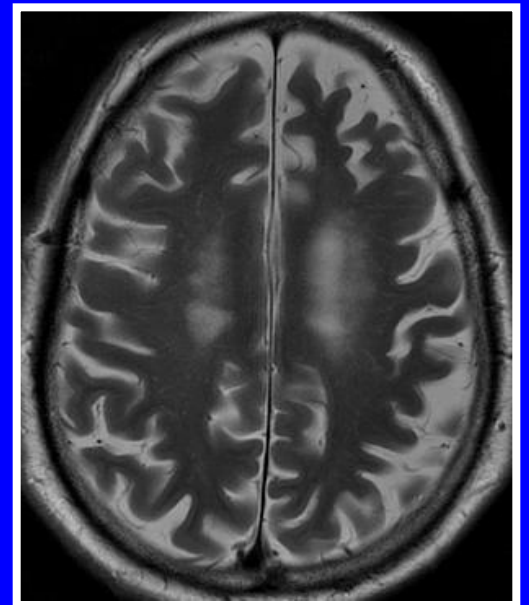
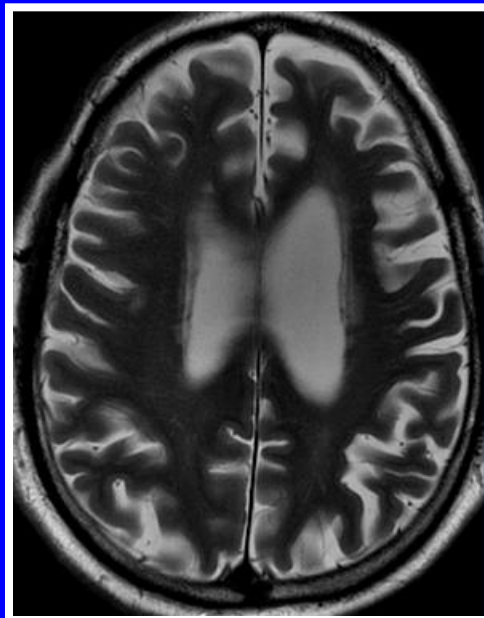
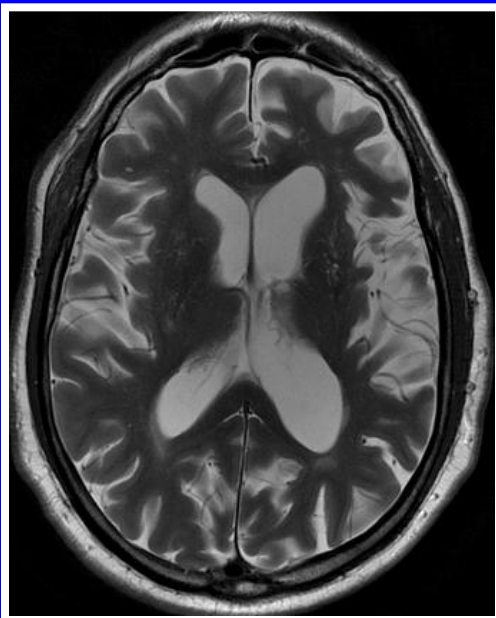
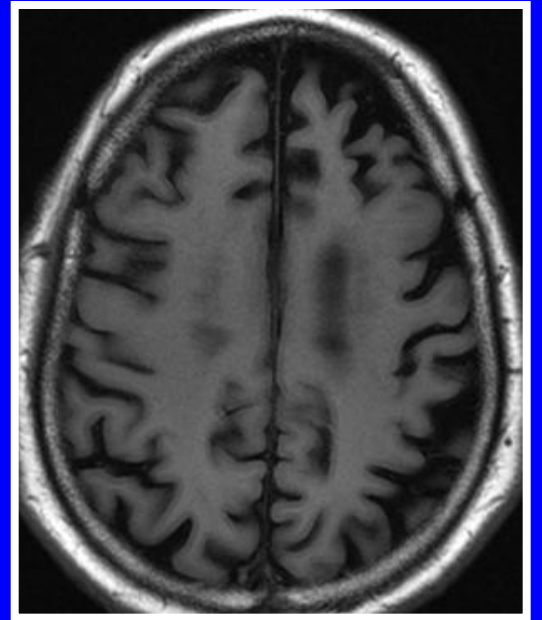
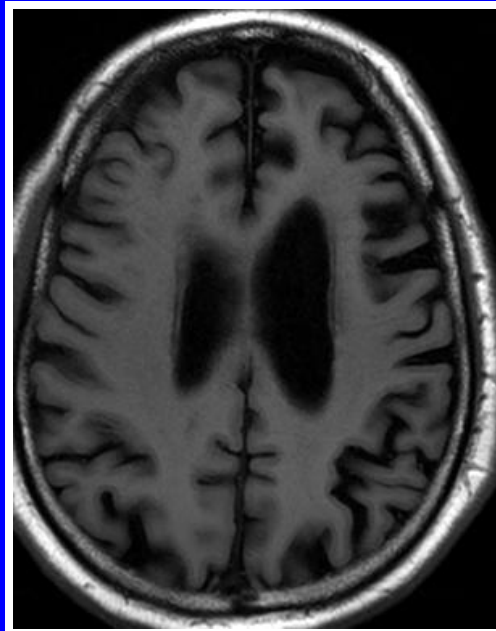
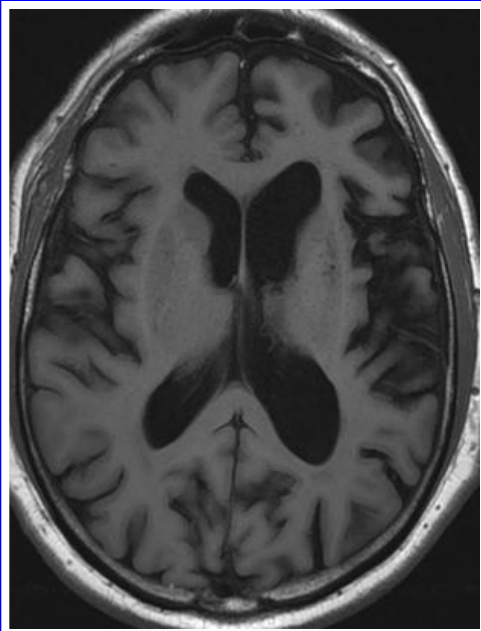
AFASIA PROGRESIVA PRIMARIA:

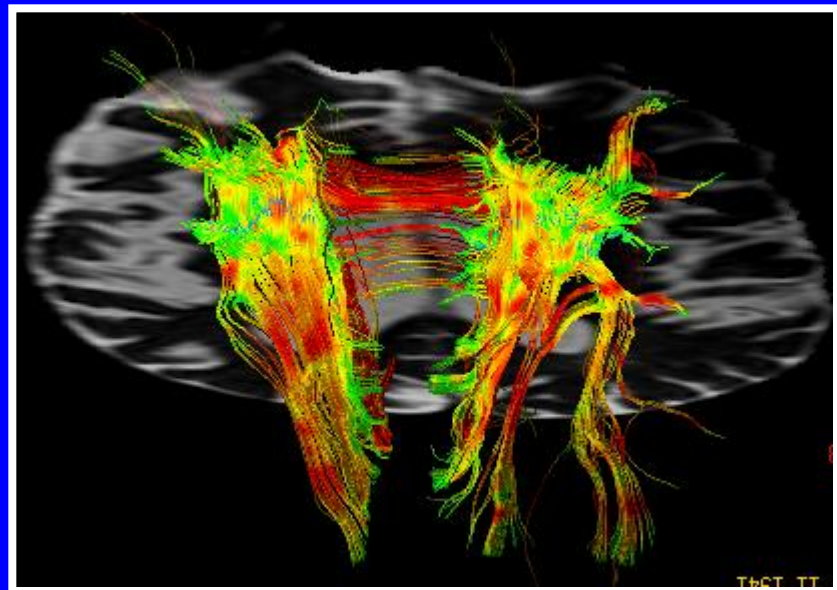
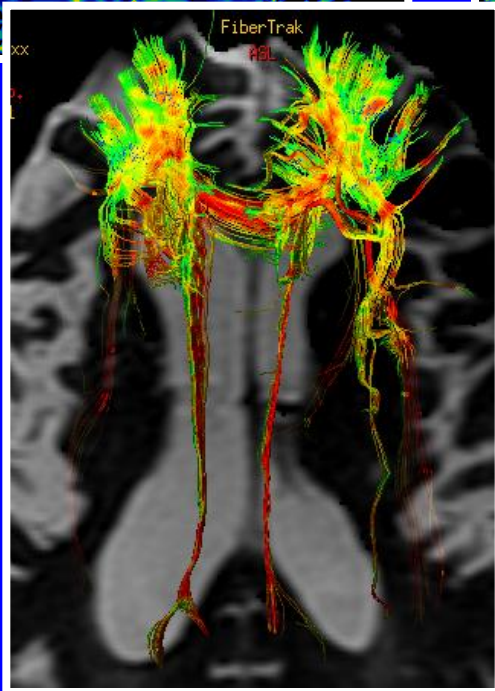
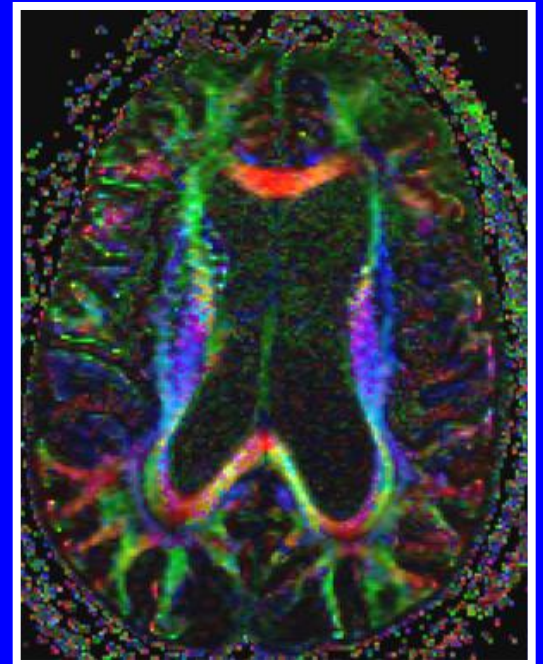
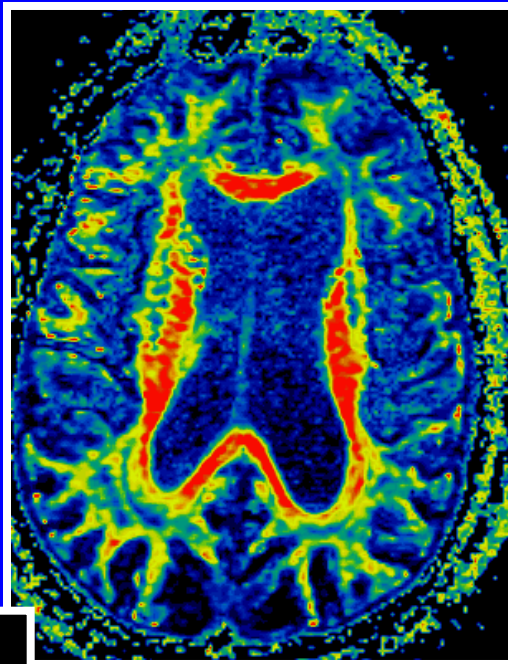
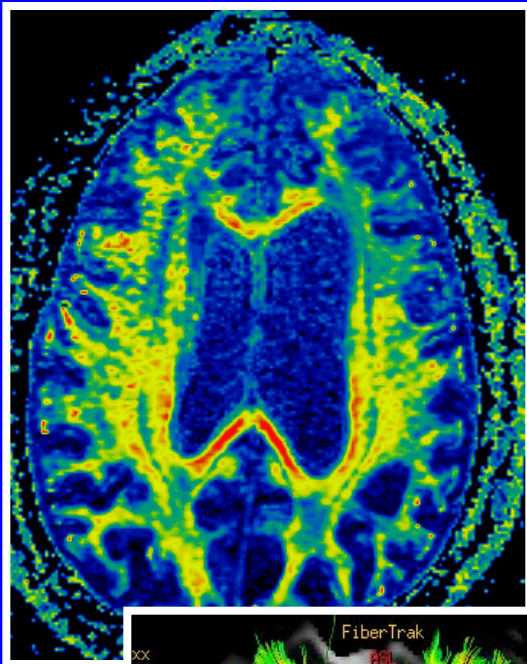


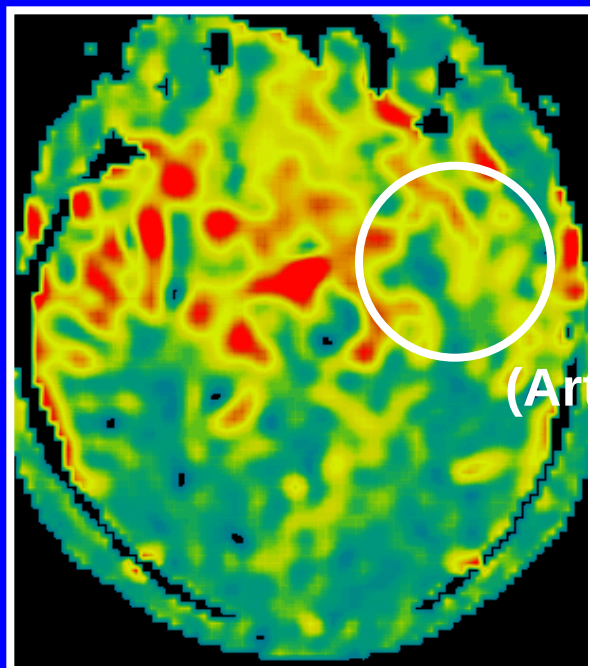
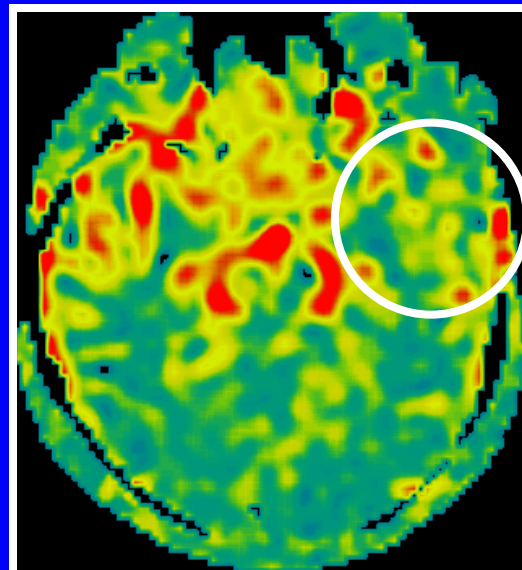
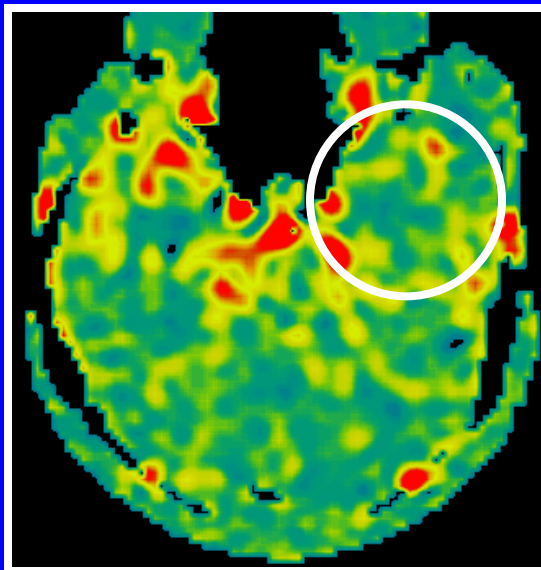
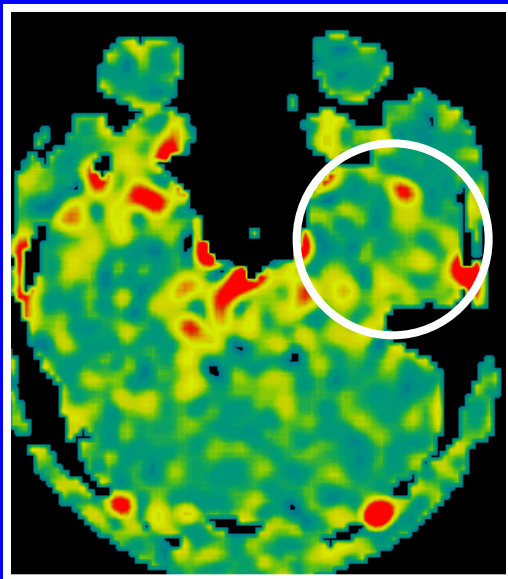
- **APP No Fluente:** (tau)
- **APP SEMANTICA :** (TDP-43)
- **LOGOPÉNICA:** (Alzheimer)



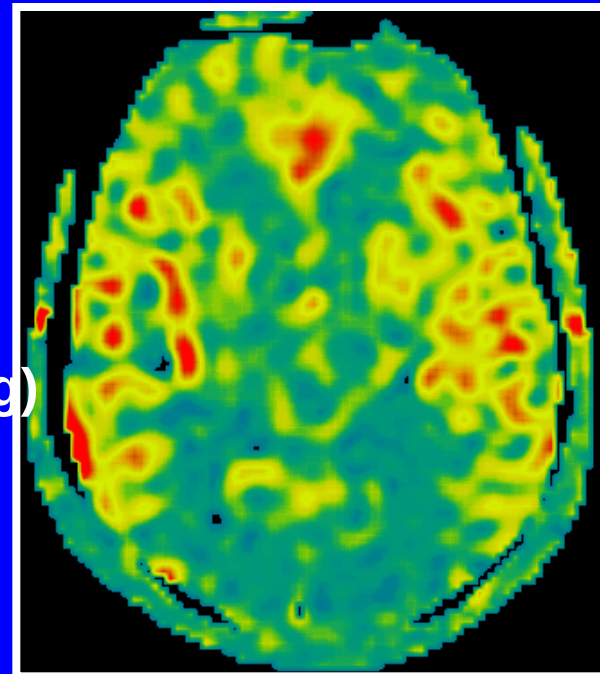
AFASIA PROGRESIVA PRIMARIA

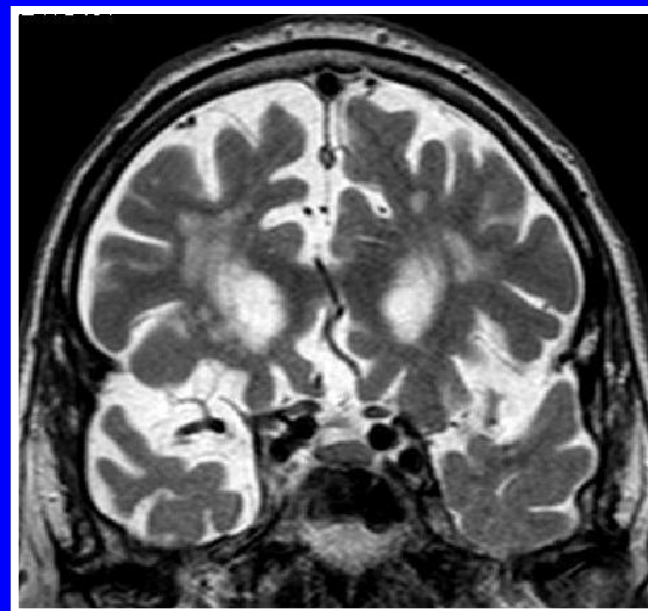
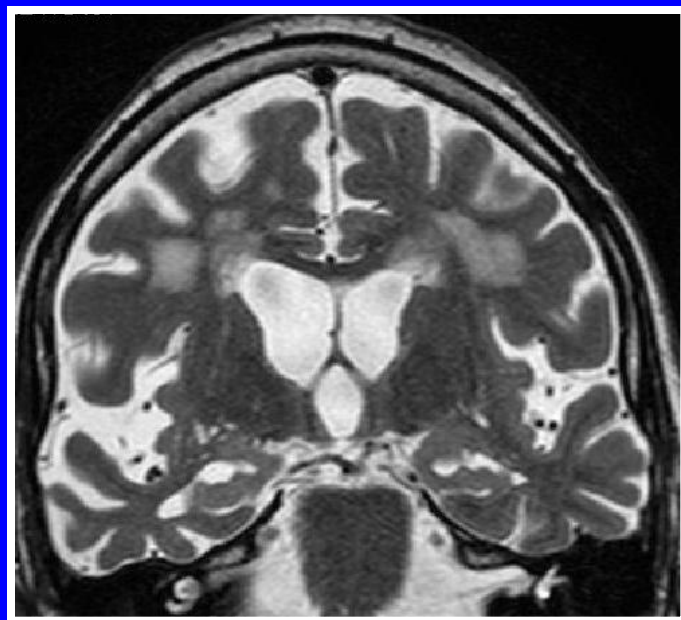
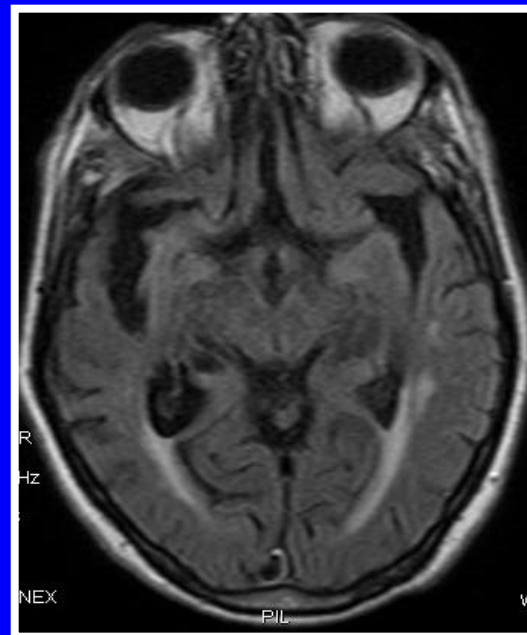
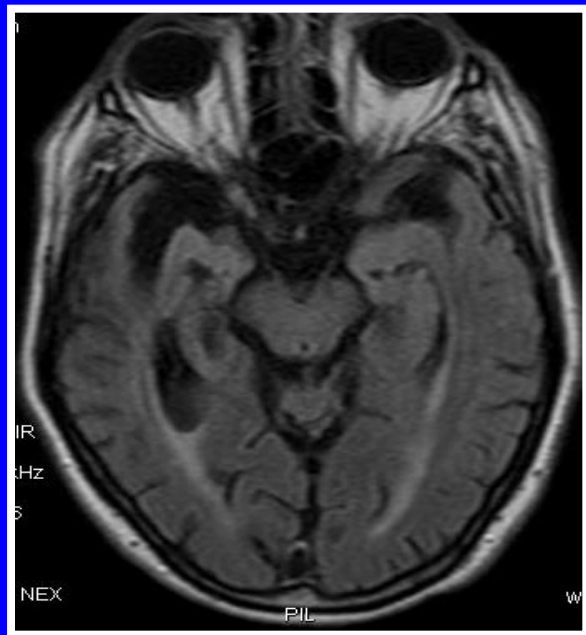




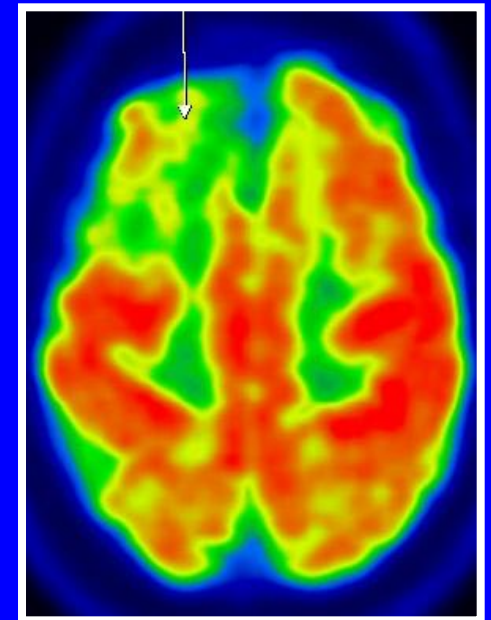
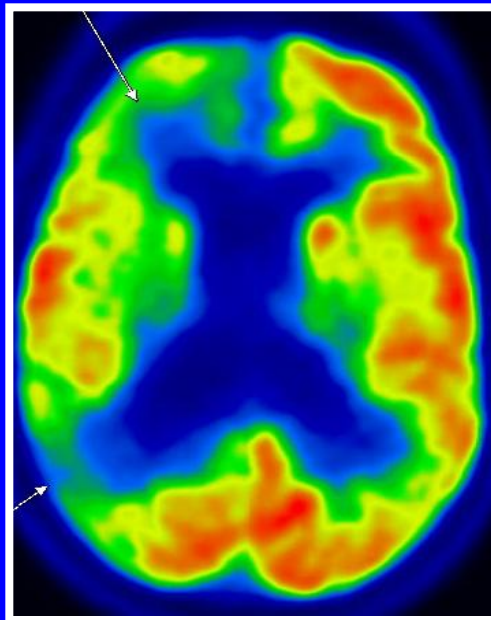
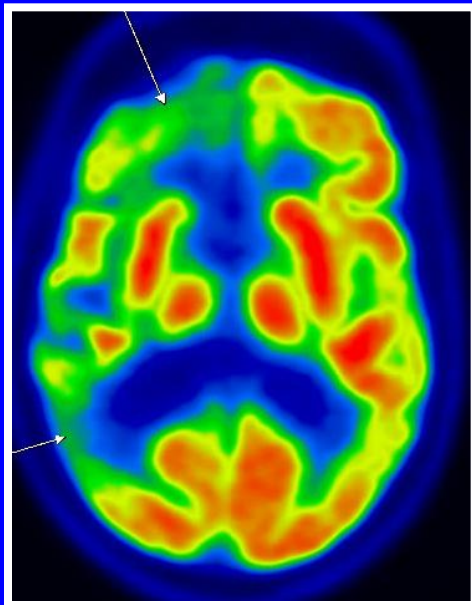
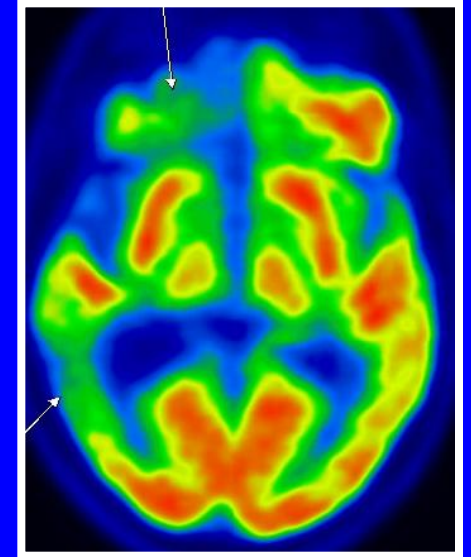
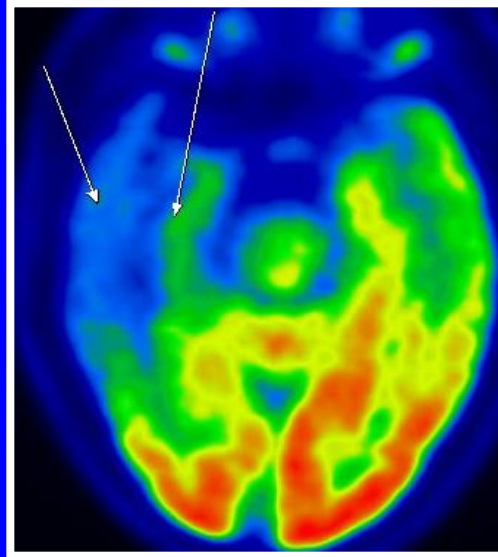
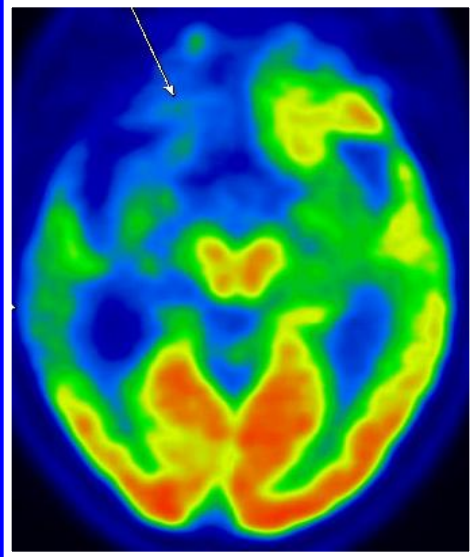


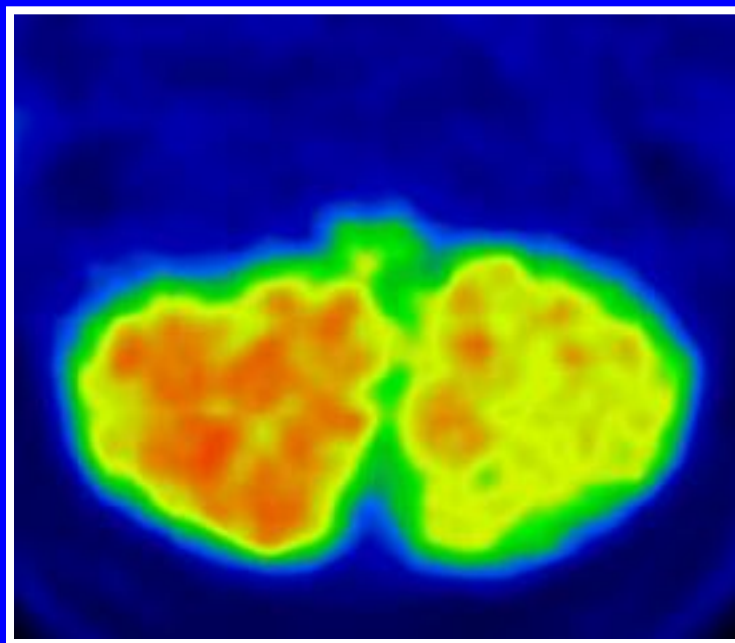
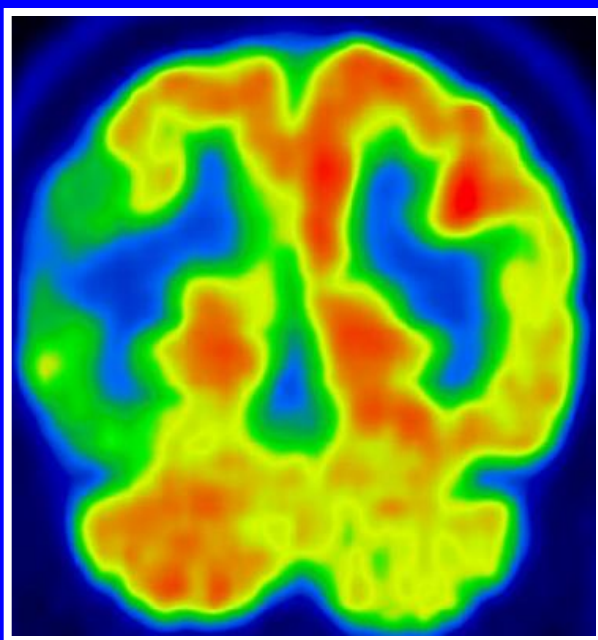
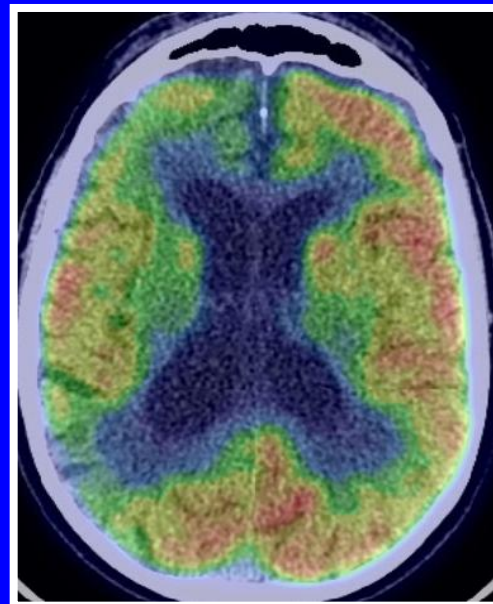
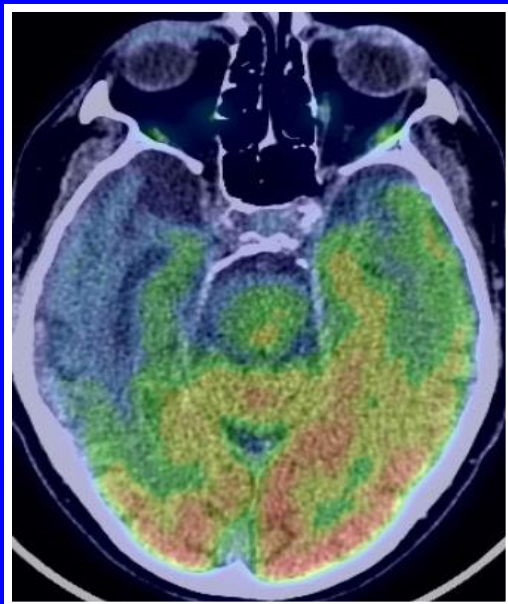
ASL
(Arterial Spin-labeling)

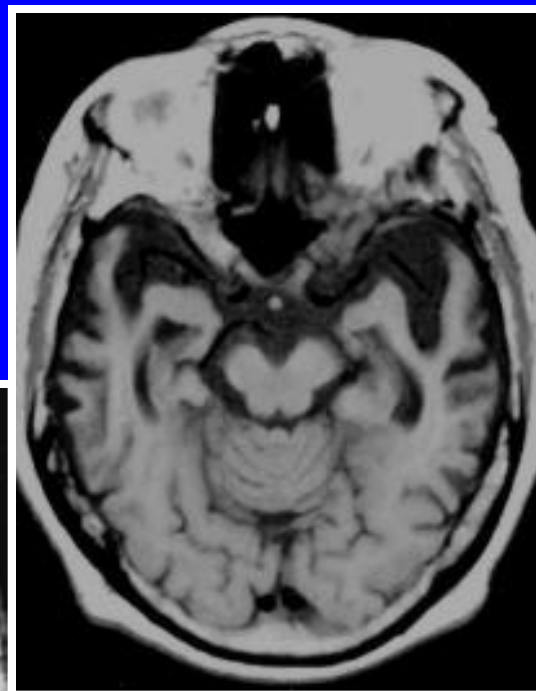
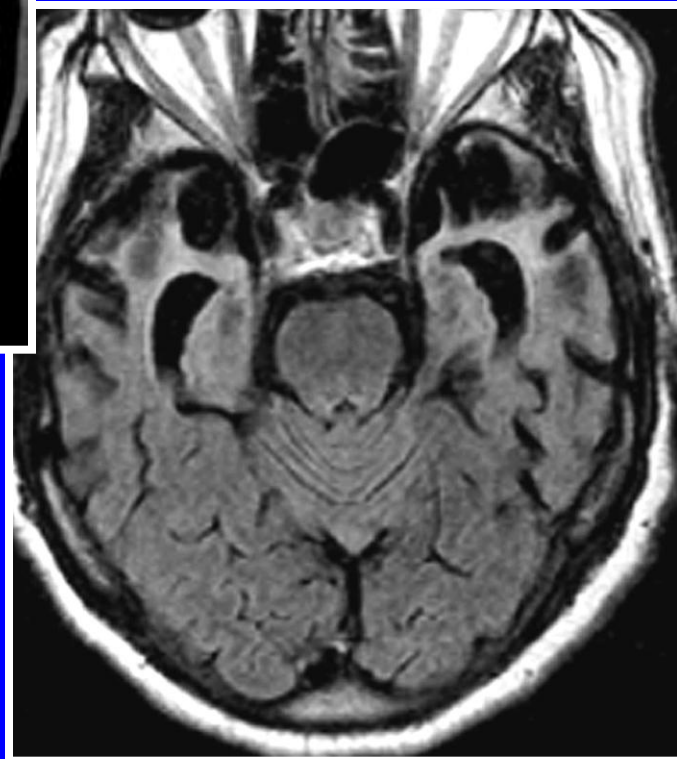
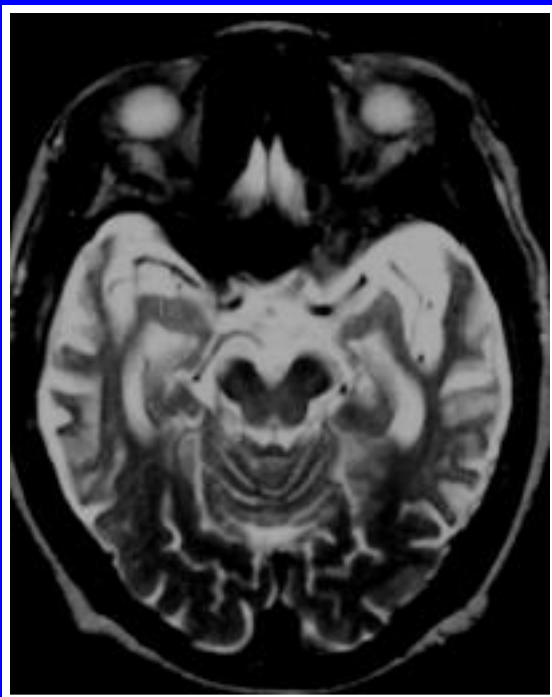




PET-FDG







A.P. SEMÁNTICA

DFT:

Estructural:

Atrofia predominante en regiones frontal y temporal anterior.

Molecular:

PET-FDG y SPECT (Disminuidos en flujo y captación de glucosa en similares regiones).

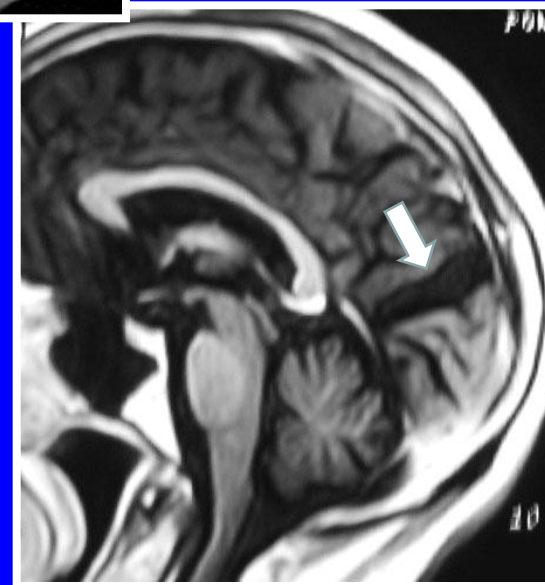
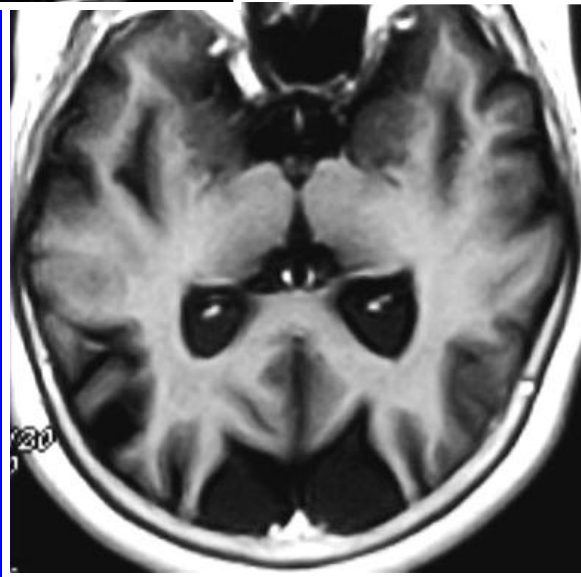
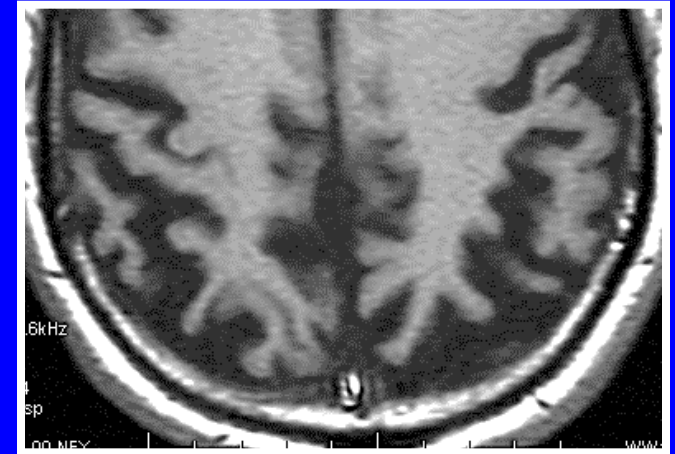
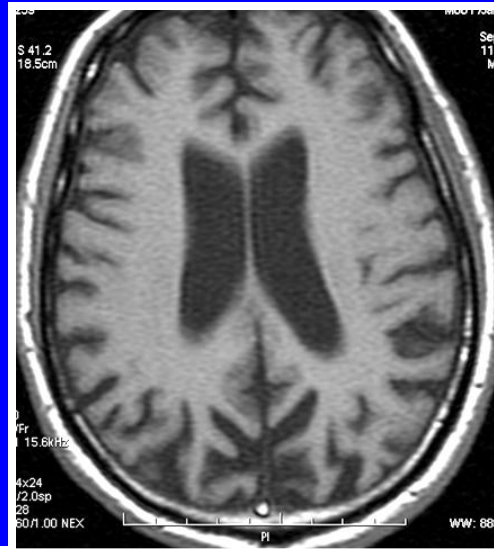
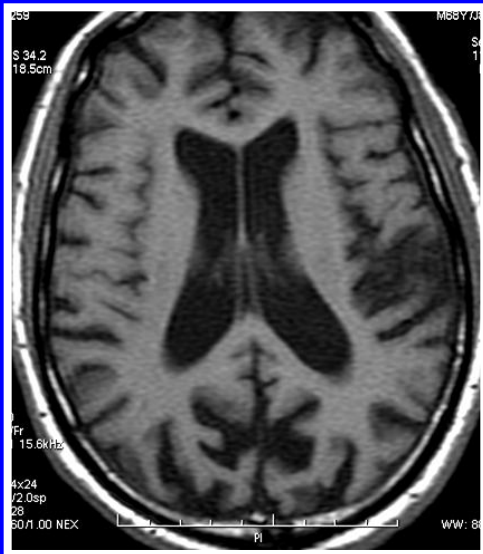
CUERPOS DE LEWY (DLB)

- Leve atrofia cortical difusa (**predomina a nivel insular y frontotemporal**, como asi tambien en regiones parieto-occipitales: *Fibras de asociación*).
- Atrofia Putaminal
- *Atrofia mesencefàlica*



DD con EP.

Demencia por Cuerpos de Lewy



DLB:

Estructural:

Atrofia lob frontal inferior, corteza visual, insula, región diencéfalo-mesencefálica, caudado, putámen e hipocampo anterior (CA1)

Molecular:

SPECT (Perf disminuida en putámen, caudado y corteza visual), PET-FDG (Menor captación en corteza visual)

DEMENCIA VASCULAR

1) Macroangiopática:

(30% de los casos: demencia)

2) Microangiopática:

Leucoaraiosis

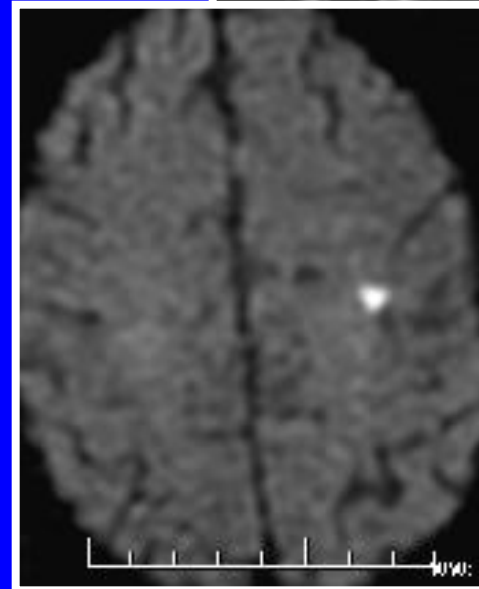
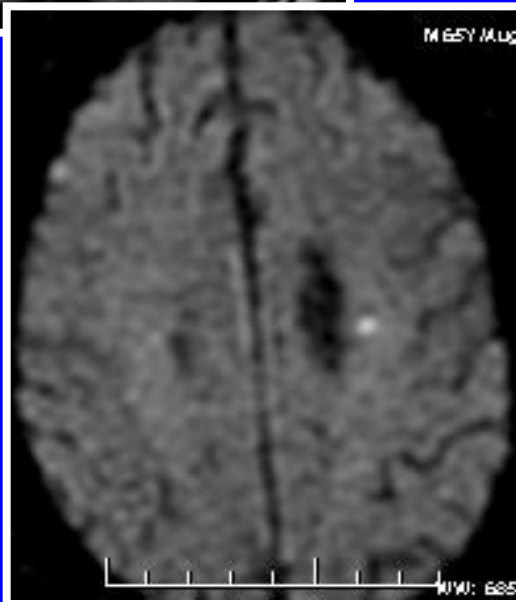
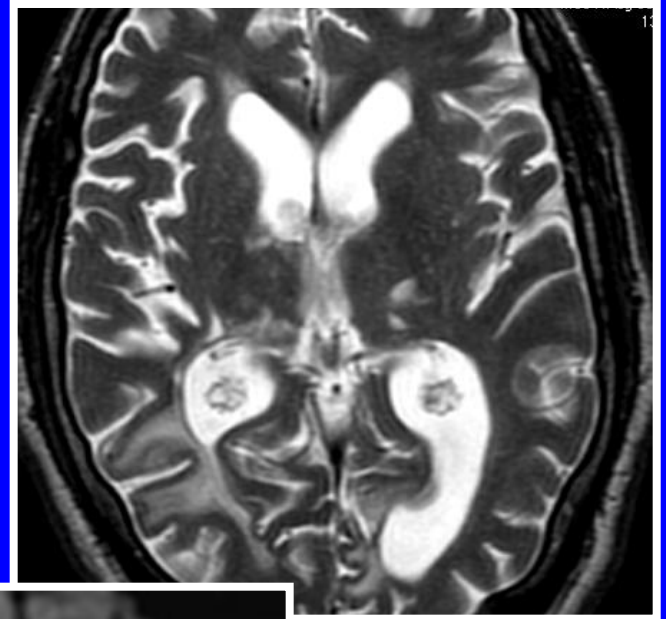
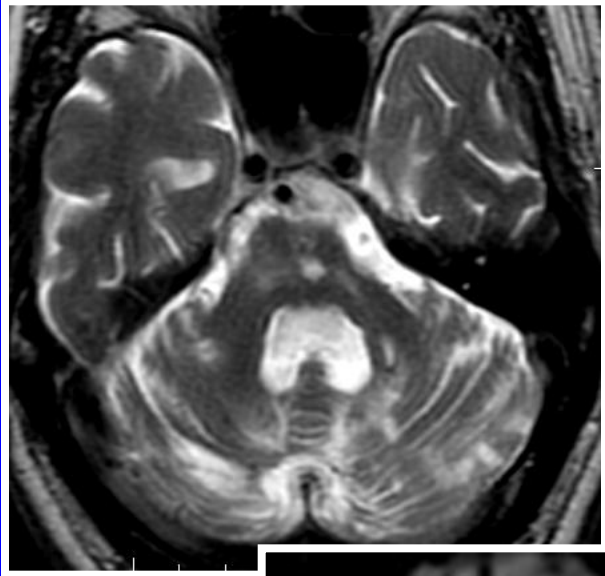
Infartos lacunares

3) Microhemorragias:

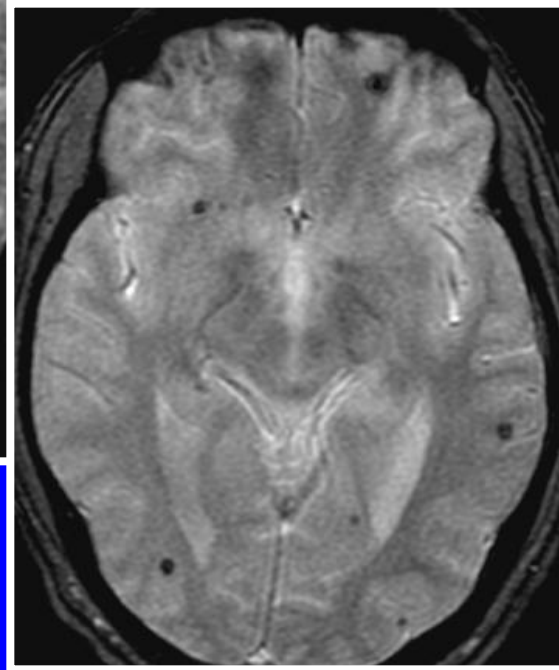
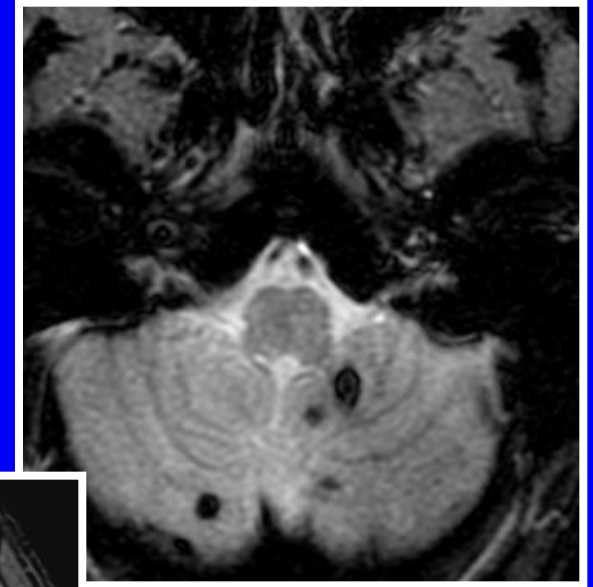
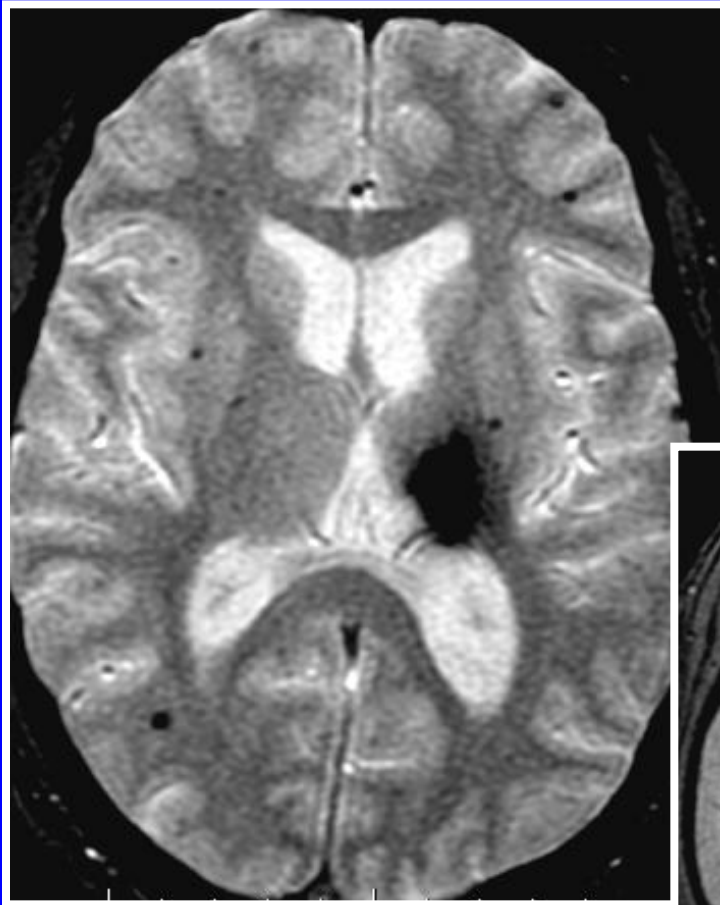
65% de casos de DV.

18% cursan con aumento de b-amiloide y
de ubicación cortico-subcortical fronto-
parietal

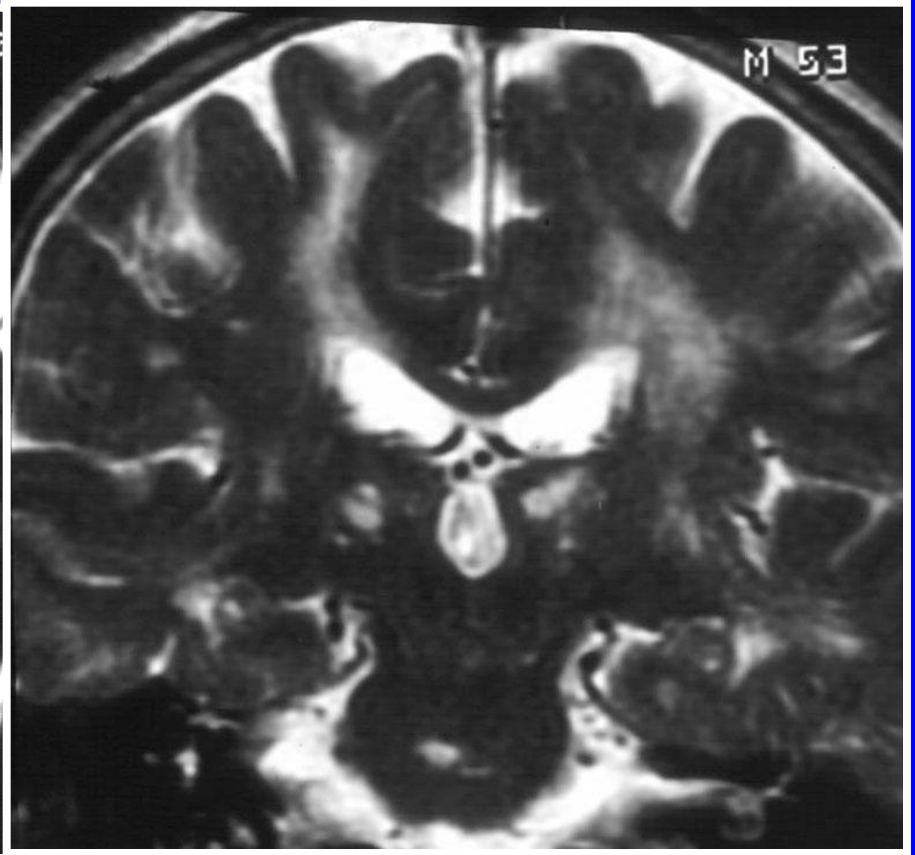
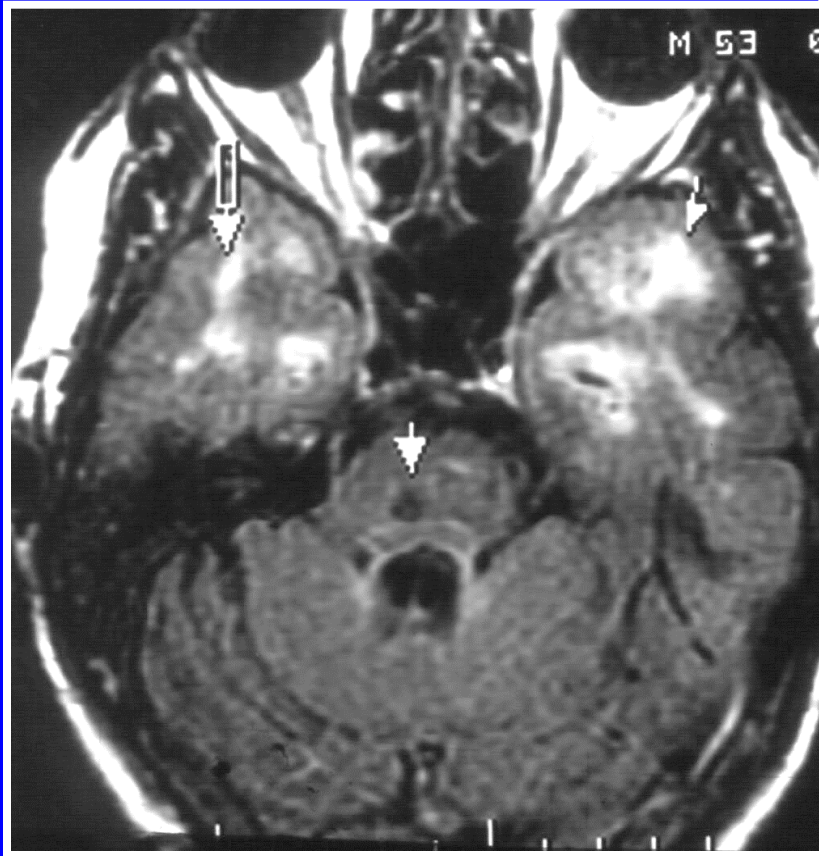
Demencia Multi-infarto



Micro/Macrosangrados crónicos en HTA



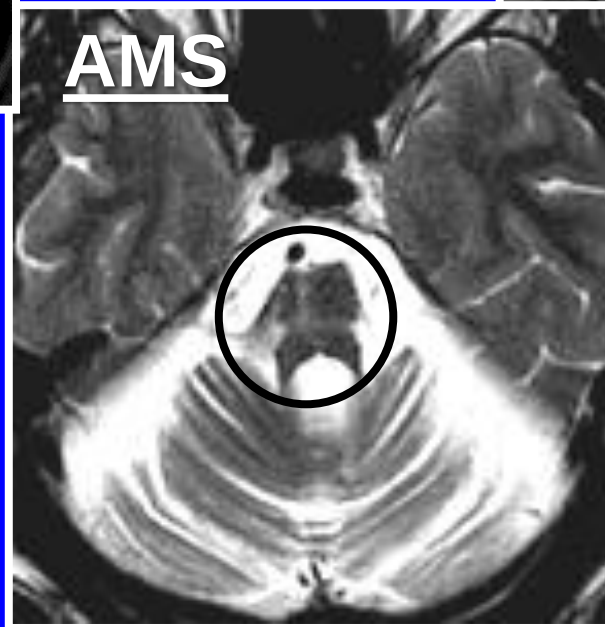
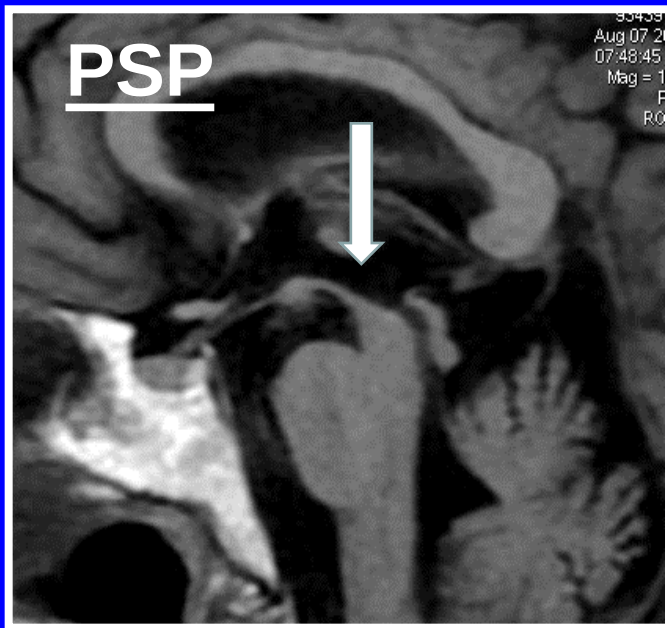
CADASIL



SINDROMES PARKINSONIANOS

ATÍPICOS

- **Parálisis Supranuclear Progresiva (PSP)**
- **Atrofia multisistémica (AMS)**
- **Degeneración córtico-basal (DCB)**
- **Demencia cuerpos de Lewy (DCL)**

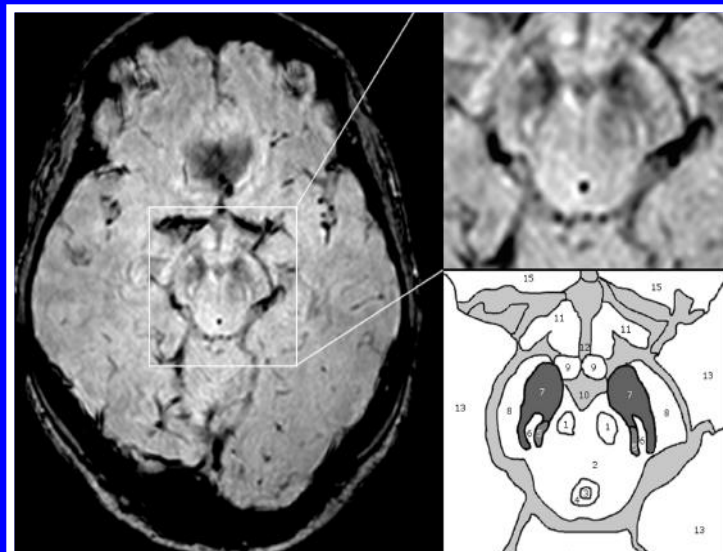


*De Diego S, Meli F., Diagnóstico Maipú,
Buenos Aires, Argentina.
AJNR, Noviembre 8, 2010. Case of the week*

The 'Swallow Tail' Appearance of the Healthy Nigrosome – A New Accurate Test of Parkinson's Disease: A Case-Control and Retrospective Cross-Sectional MRI Study at 3T

Stefan T. Schwarz^{1*}, Mohammed Afzal², Paul S. Morgan³, Nin Bajaj⁴, Penny A. Gowland⁵, Dorothee P. Auer¹

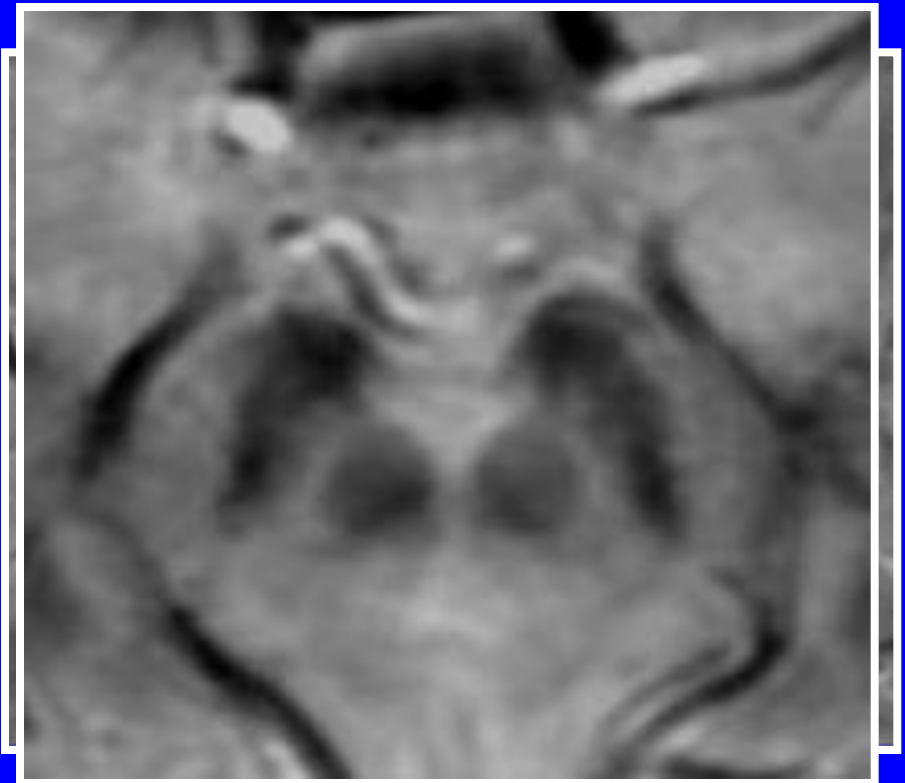
¹ Radiological Sciences, Division of Clinical Neurosciences, School of Medicine, University of Nottingham, Nottingham, United Kingdom, ² Department of Medicine, Nottingham University Hospitals NHS Trust, Nottingham, United Kingdom, ³ Medical Physics, Nottingham University Hospitals NHS Trust, Nottingham, United Kingdom, ⁴ Department of Neurology, Nottingham University Hospitals NHS Trust, Nottingham, United Kingdom, ⁵ Sir Peter Mansfield Magnetic Resonance Centre, School of Physics and Astronomy, Nottingham, United Kingdom



NIGROSOMA 1



NORMAL



ENF PARKINSON

DEMENCIAS RAPIDAMENTE PROGRESIVAS (RPD)

- **Curso subagudo**
- **Declinacion progresiva menor a 2 años**
- Trastornos neurodegenerativos
- Enfermedades autoinmunes
- Enfermedades infecciosas
- Enfermedades metabólicas

DEMENCIAS RAPIDAMENTE PROGRESIVAS (RPD)

- Enf por priones: 62%
- Otras enf. ND: 15%
- Autoinmunes: 8%
- Infecciones: 4%
- Otras: 2% (tox, metab, cancer, vascular)
- Indeterminadas: 4% (leucoencefalopatias)
- *(UCSF, Memory and aging center), Sep 2012)*

Neurology® Clinical Practice

Diagnosis and treatment of rapidly progressive dementias

Ross W. Paterson, MRCP*

Leonel T. Takada, MD*

Michael D. Geschwind, MD, PhD

DEMENCIAS RAPIDAMENTE PROGRESIVAS (RPD)

- Regla “mnemotécnica clínica”:
 - **V** : Vascular
 - **I** : Infecciones
 - **T** : Tóxicas-metabólicas
 - **A**: Autoinmunes
 - **M**: Mts/neoplasias
 - **I**: Iatrogenia/errores cgts del metabolismo
 - **N**: Neurodegenerativas
 - **S**: Enfermedades sistémicas/ status

DEMENCIAS RAPIDAMENTE PROGRESIVAS

1) PRIONICAS: CJD.

2) NO PRIONICAS:

- Encefalitis límbica :

- Autoinmune.
- Paraneoplásica
- Infecciosa (Herpes)

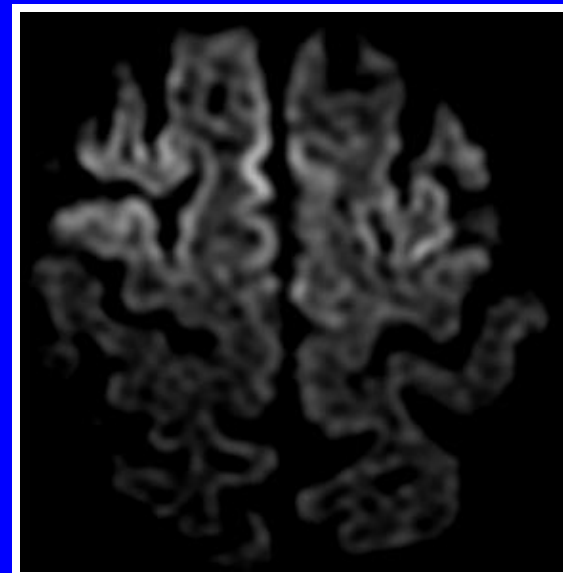
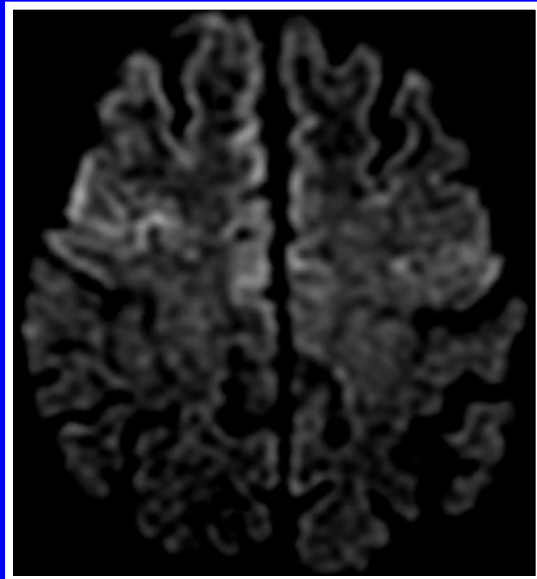
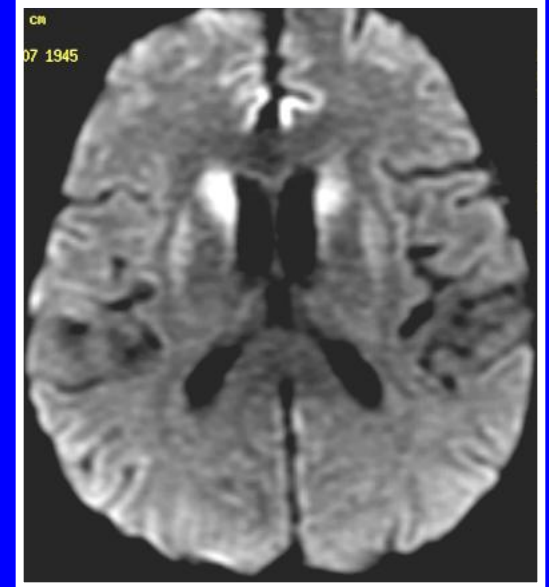
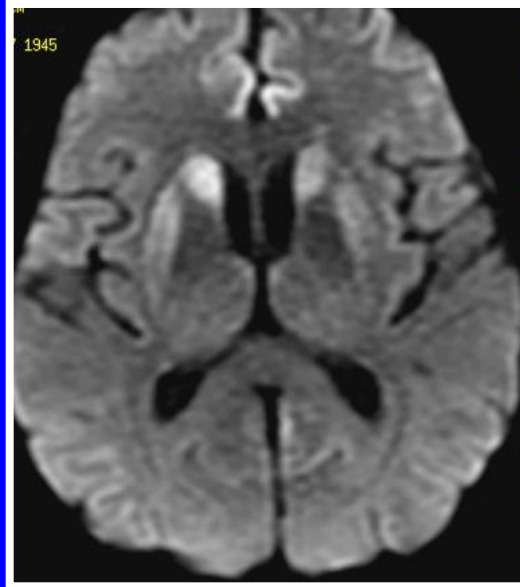
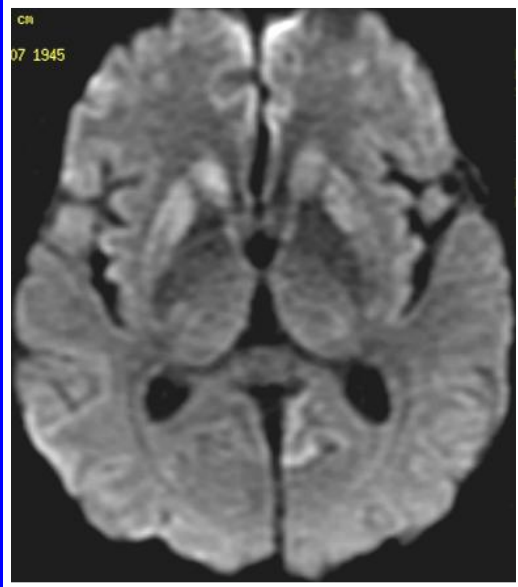
- Encefalitis anti NMDA (*N-metil-D-aspartato*)

- Encefalitis anti AMPA-R (*alfa-amino-3-hidroxi-5-metil-4ácido isoxazolepropionic*)

- Encefalitis anti canales de K

- Encefalopatía respondedora a corticoides o encefalopatía de Hashimoto.

Enfermedad de Creutzfeldt-Jakob (CJD)



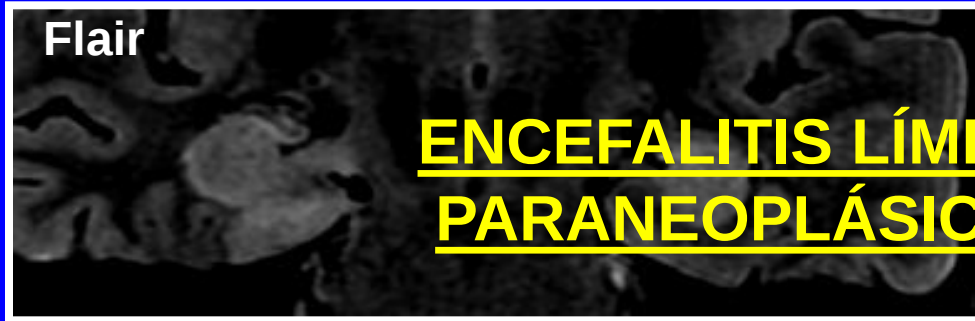
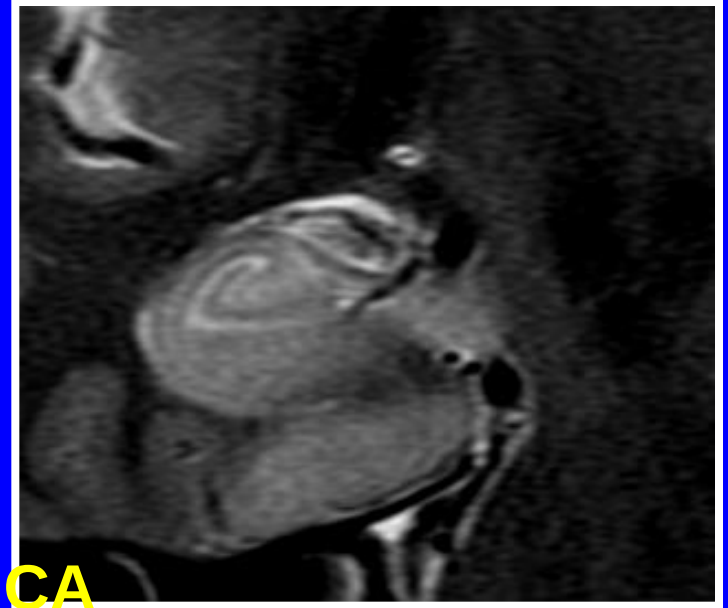
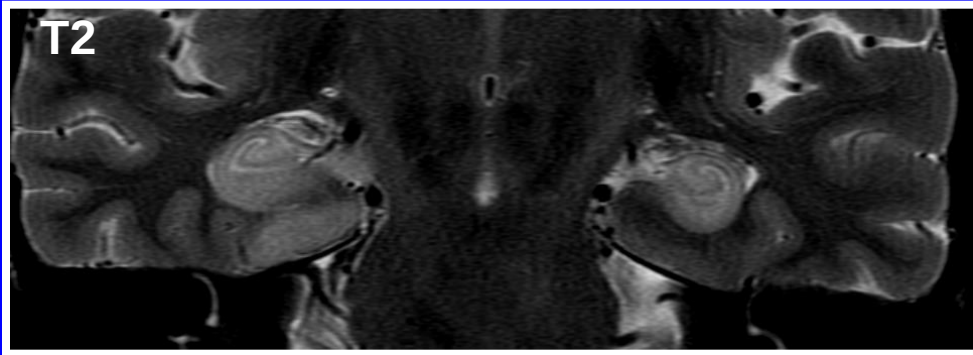
Características encontradas en RM con secuencias DWI y FLAIR

1. **HIPER DWI** en sCJD **mayor que en FLAIR.**
2. **HIPER en FLAIR mayor que DWI en DRPnp.**
3. **HIPER DWI subcortical en sCJD con ADC descendido (siempre).**
4. **Compromiso límbico NO se encontró en sCJD. Frecuente en DRP.**

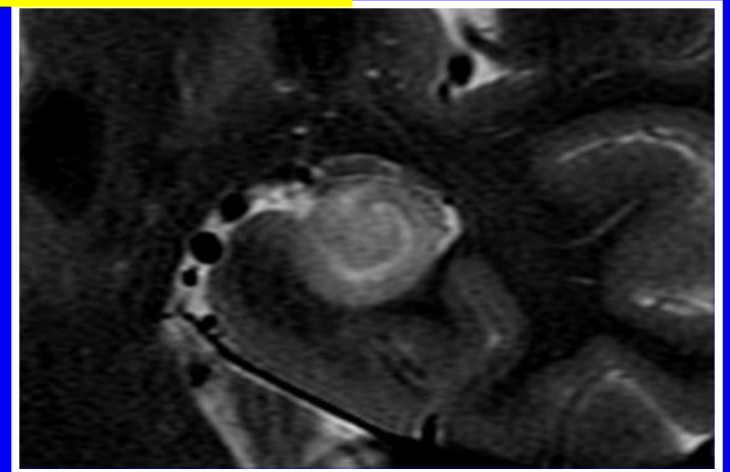
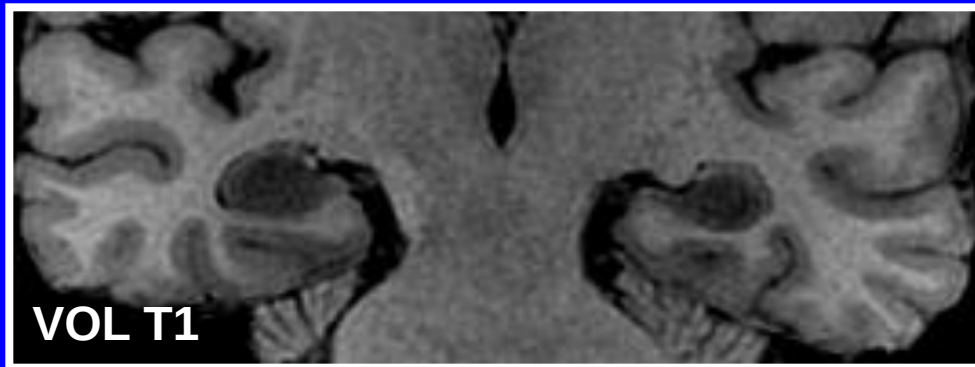
CASO 1

PACIENTE FEMENINA, 42 AÑOS DE EDAD

- CRISIS CONVULSIVA TCG
- POSTERIORMENTE COMIENZA CON
MARCADA ALTERACION EN LA MEMORIA
ANTEROGRADA.
- REPITE EN FORMA AISLADA CRISIS
PARCIALES COMPLEJAS
SECUNDARIAMENTE GENERALIZADAS.



ENCEFALITIS LÍMBICA
PARANEOPLÁSICA (Ca. Mama)



OTRAS DEMENCIAS

- LMP
- LEUCODISTROFIAS
- ENFERMEDADES MITOCONDRIALES
- HIDROCEFALIA NORMOTENSIVA
- LINFOMA

CONCLUSIONES:

- Las imágenes disponibles (RM, PET, SPECT) podrían llegar a ser indispensables en el diagnóstico y tratamiento de estas enfermedades, especialmente cuando haya tratamientos definitivos.
- Hipoflujo/hipometabolismo (SPECT/PET) simétrico parieto-temporal bilateral (precuneo/cingulo post)  ALZHEIMER.
- Compromiso anterior y asimétrico  DFT.
- Distribución más posterior (Lób. Occipital)  DLB.

MUCHAS GRACIAS

