



# Pseudotumeurs et pièges en imagerie tumorale

## Session Thématique et Ateliers de Neuroradiologie

Delphine LECLERCQ

## Pièges diagnostiques

Pseudo-tumeurs

Caractérisation tumorale

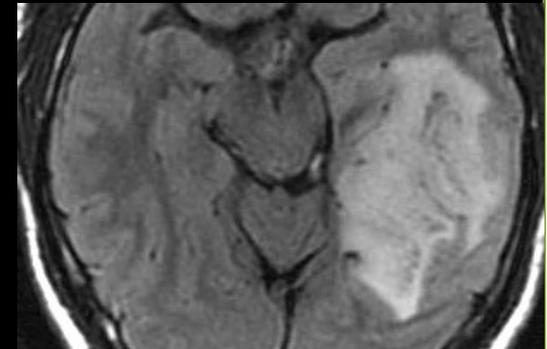
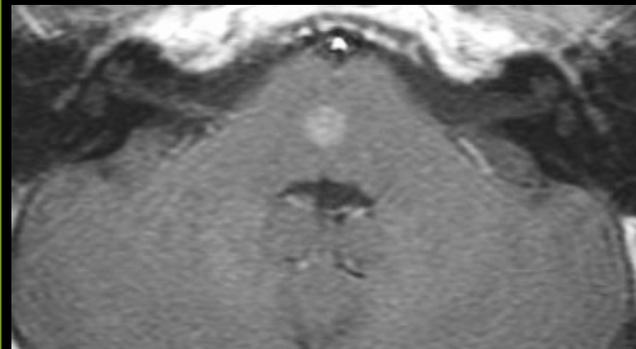
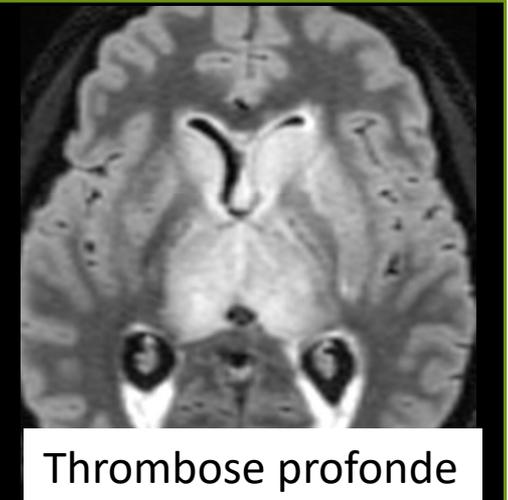
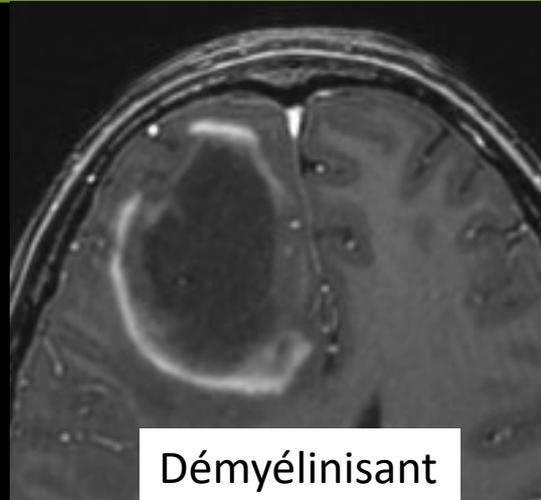
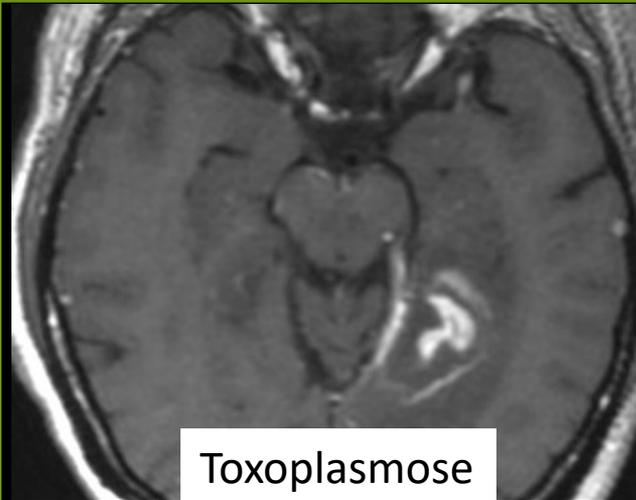
Dans la prise en  
charge = **urgence**

## Suivi

des lésions, post-op,  
chimio-radiothérapie

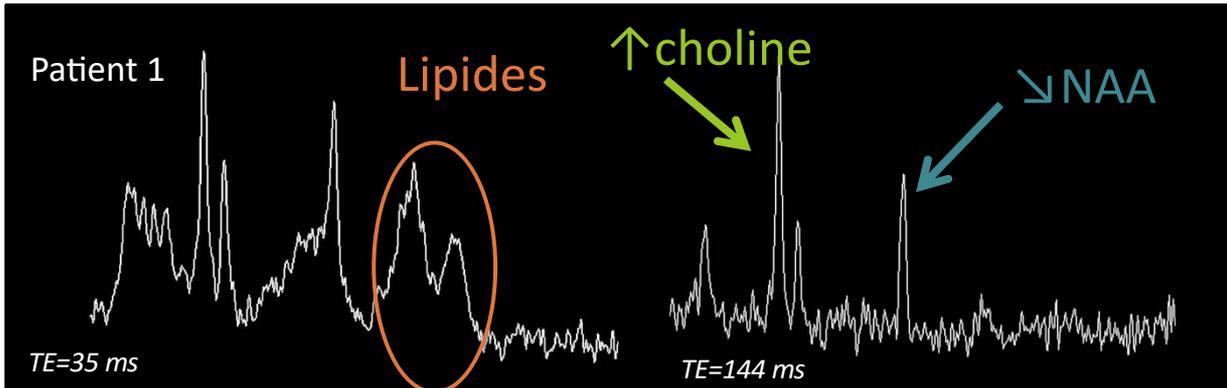
## Lésions à ne pas prendre pour une tumeur

## Pseudotumeurs



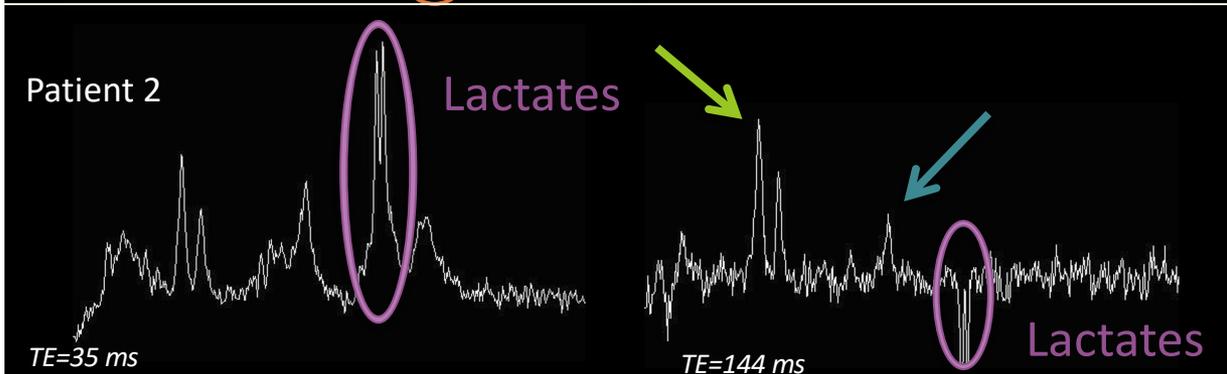
## Pseudo-tumeurs : apport de la multimodalité?

## Spectroscopie :

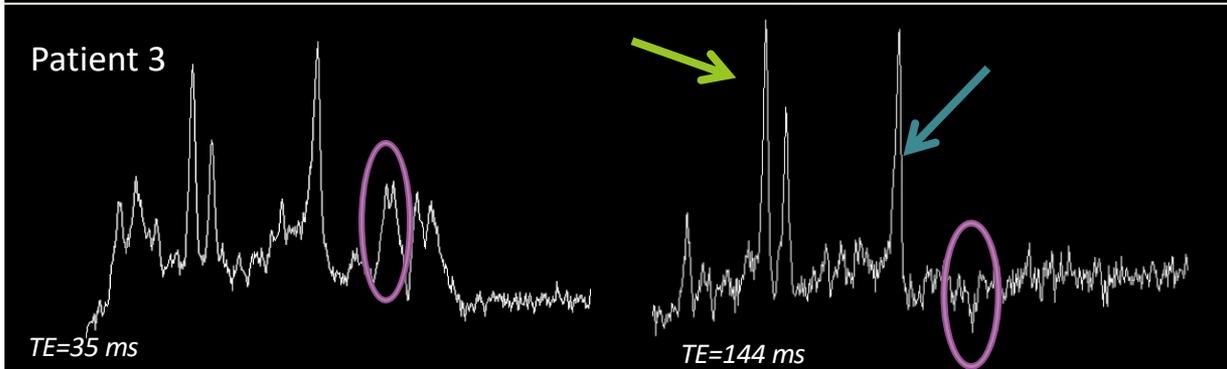


Non-spécifique!

= AVC



= démyélinisant

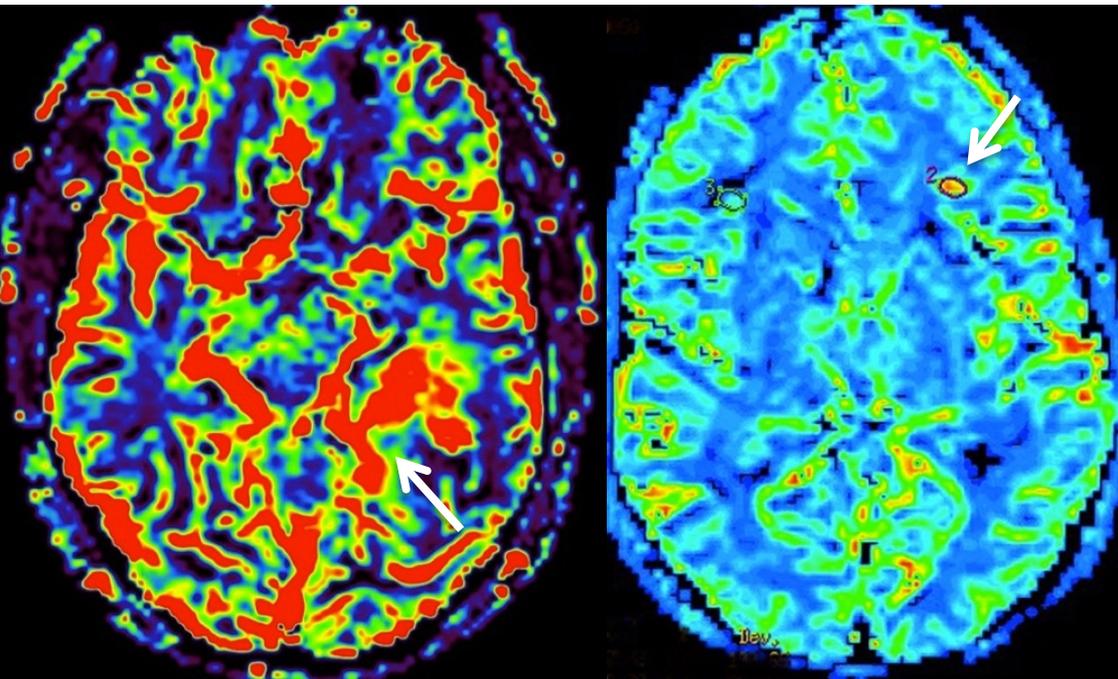


= AVC

## Pseudo-tumeurs : apport de la multimodalité?

Hyperperfusion  $\neq$  tumeur

Perfusion + gado (DSC)



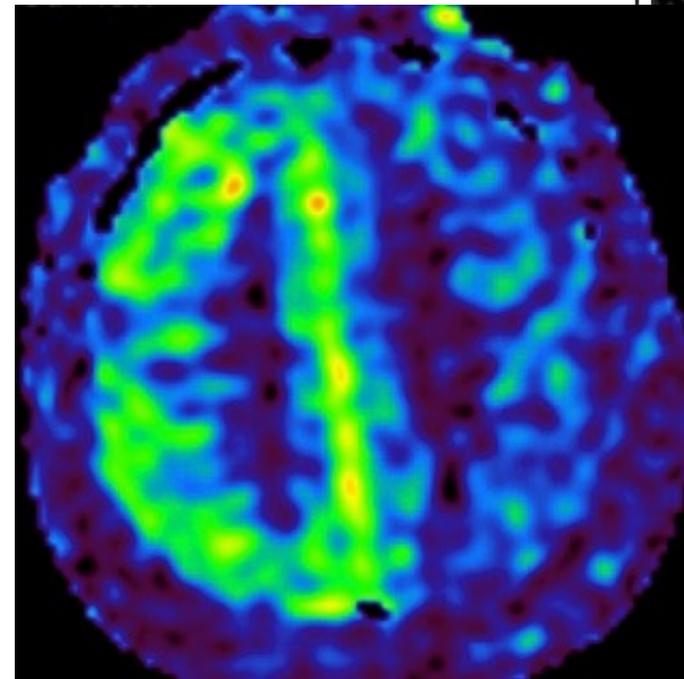
AVC

= perfusion de luxe

=&gt; 72h

Histiocytose

Perfusion ASL



Status epilepticus

= ASL

Diagnostic

Prise en charge

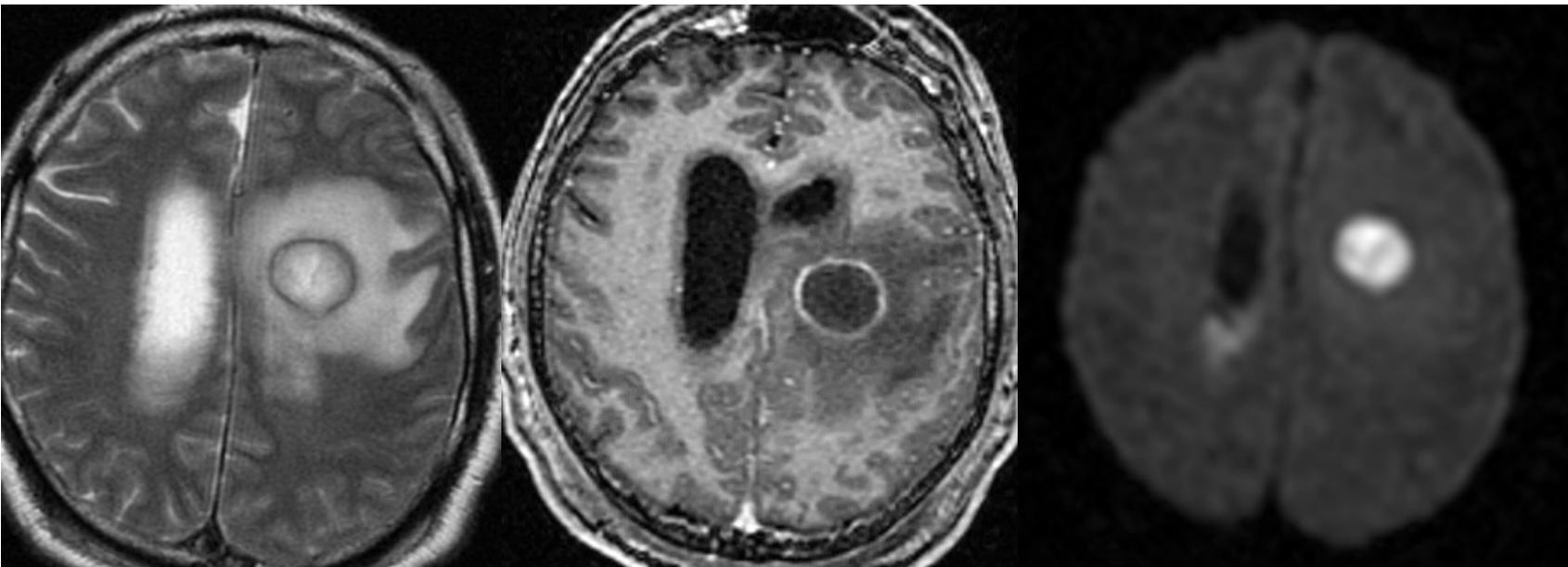
Suivi

Conclusion

## Diffusion DWI

Lésion nécrotique

K oesophage



Diagnostic

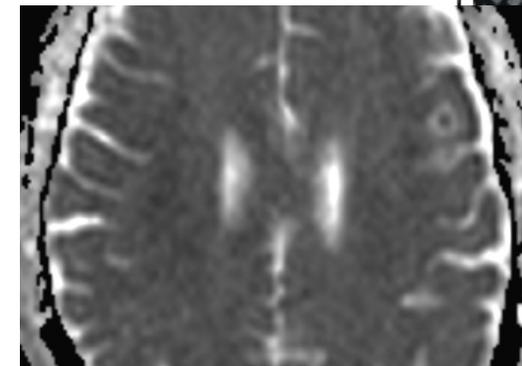
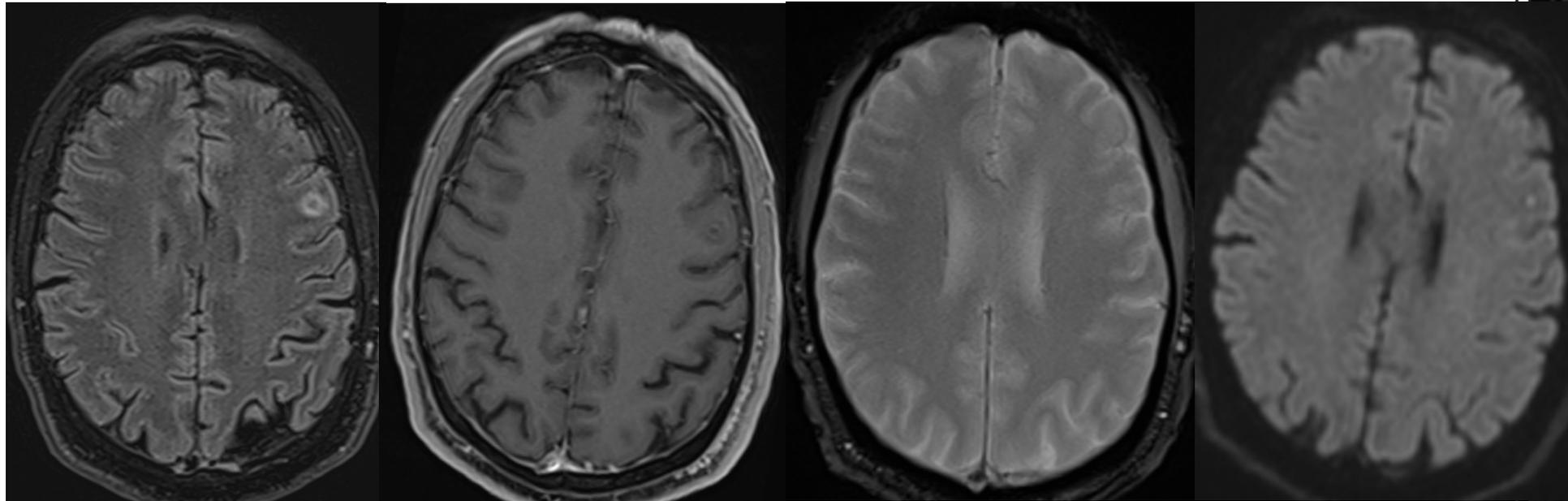
Prise en charge

Suivi

Conclusion

## Diffusion DWI

Lésion nécrotique



Diagnostic

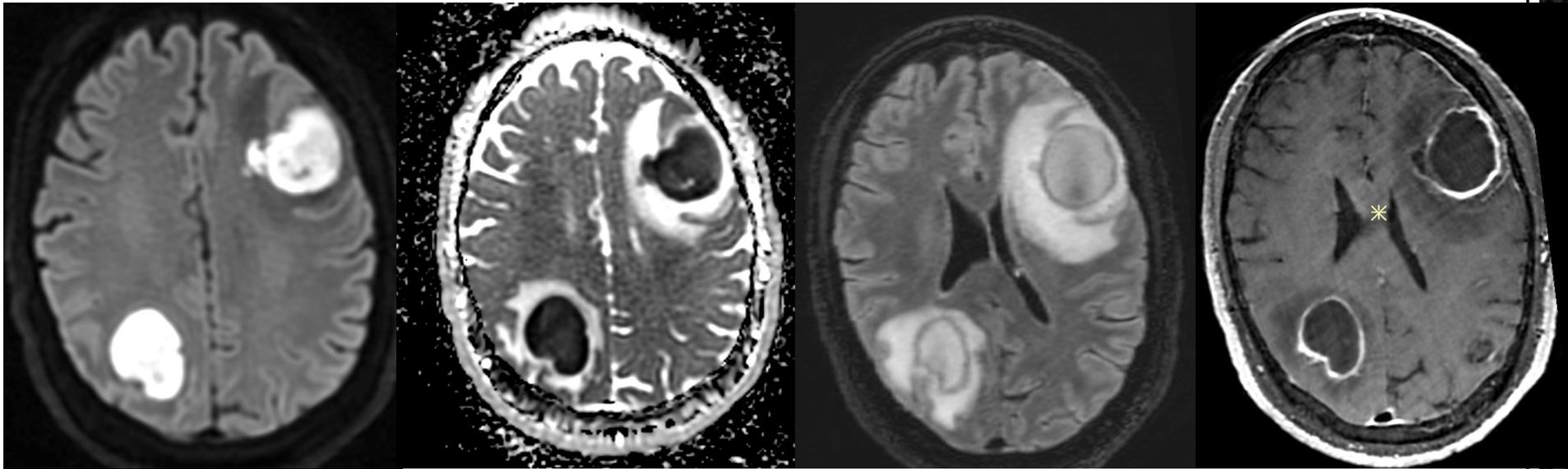
Prise en charge

Suivi

Conclusion

## Diffusion DWI

Lésion nécrotique

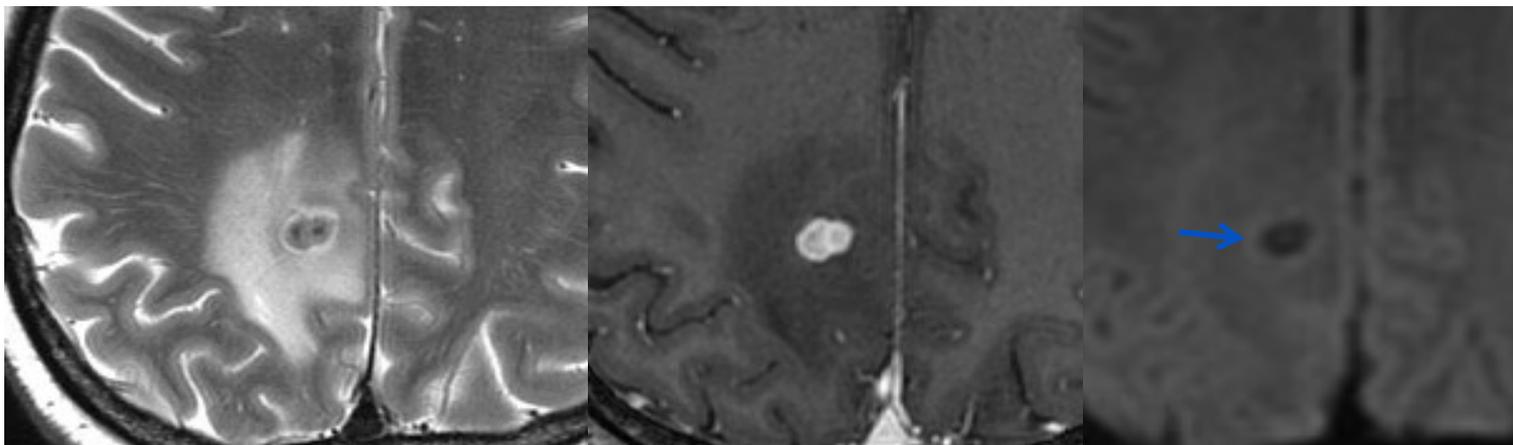


14 jours après...

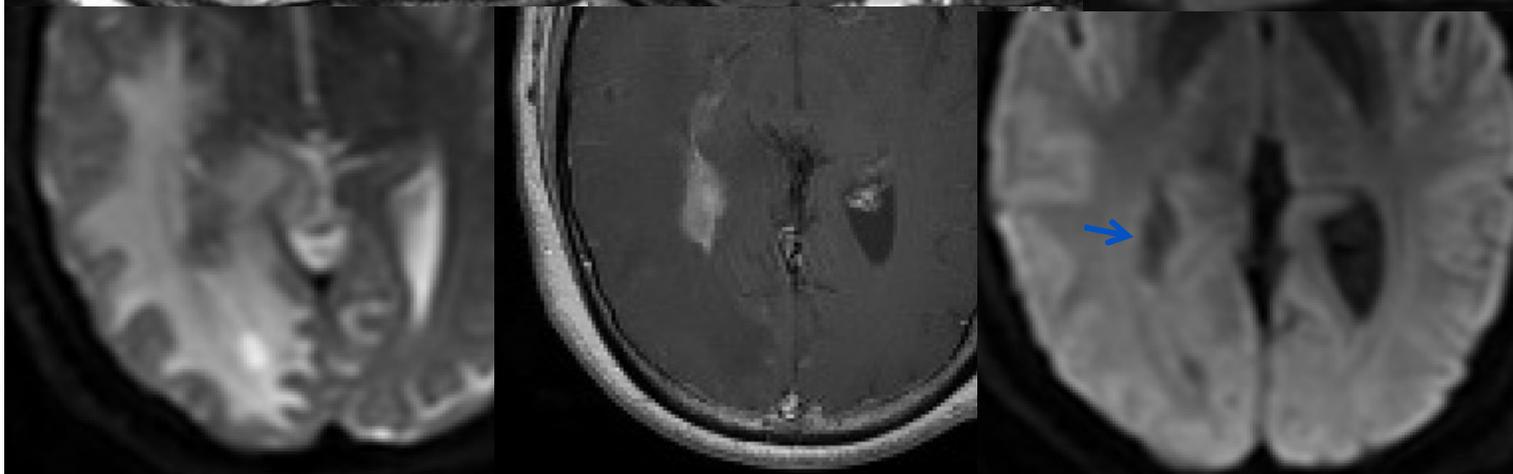
## hypoT2

Lésion solide

Pas de restriction de la diffusion  
= penser infection + inflammatoire



BK



Sarcoïdose

Hyposignal DWI

Diagnostic

Prise en charge

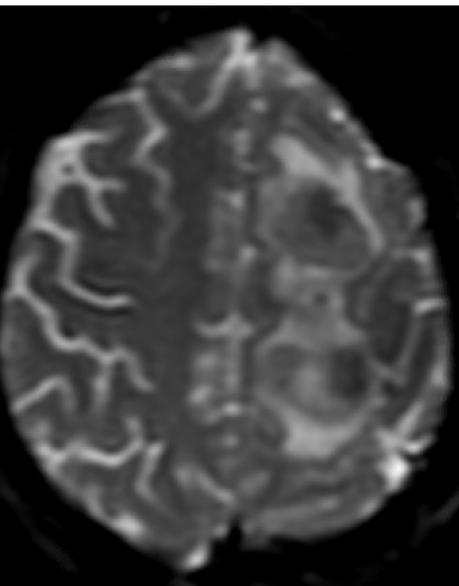
Suivi

Conclusion

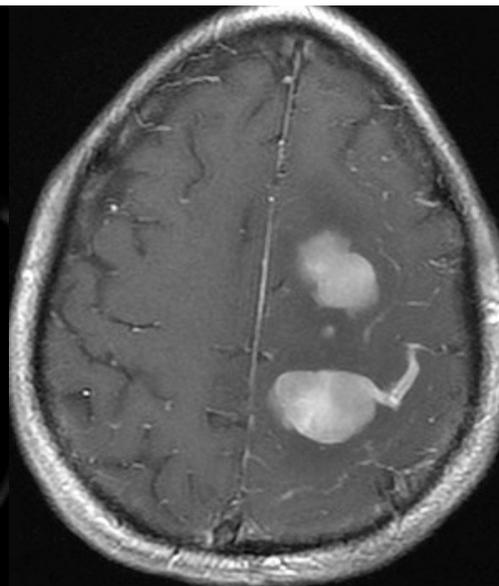
hypoT2

Lésion solide

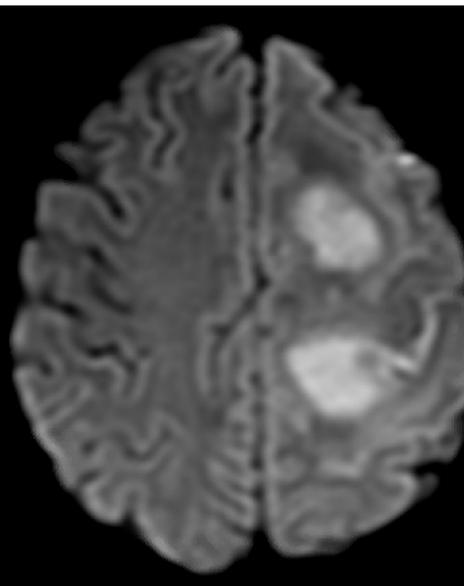
Tumeur hypoT2  
= habituellement hypercellulaires en DWI



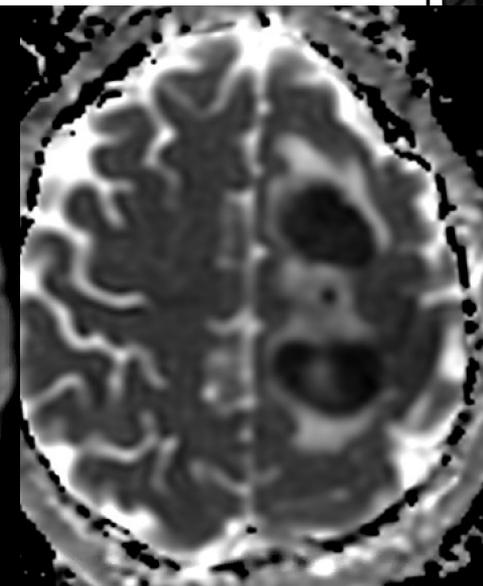
BO



Leucémie



Hypersignal DWI

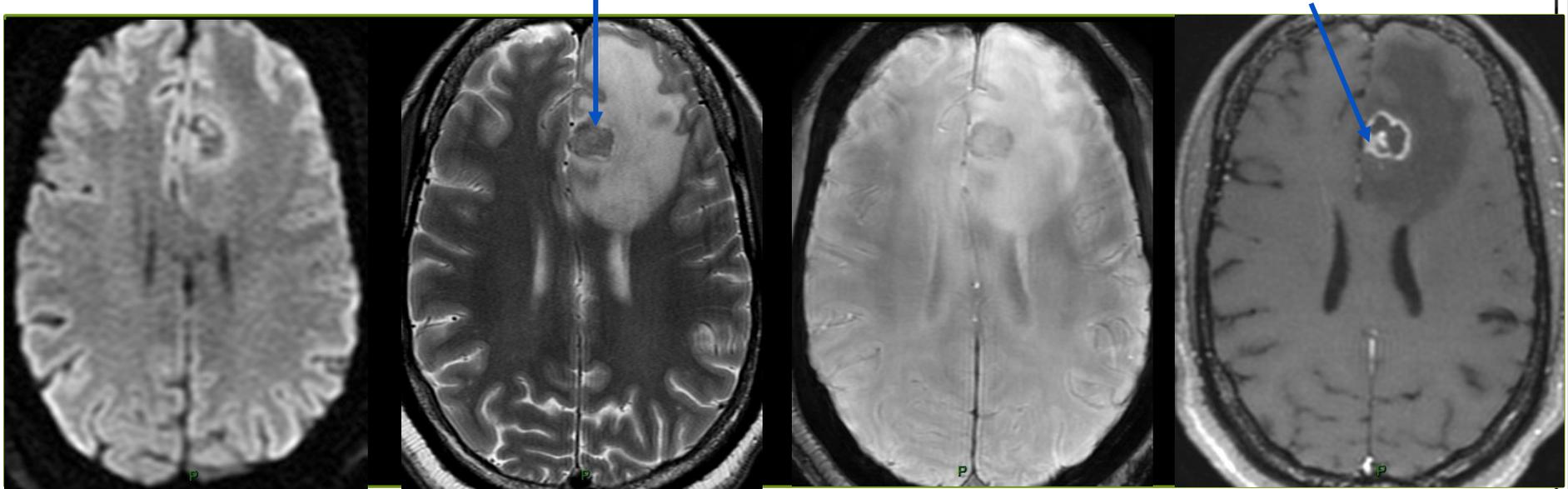


ADC bas  
= hypercellulaire

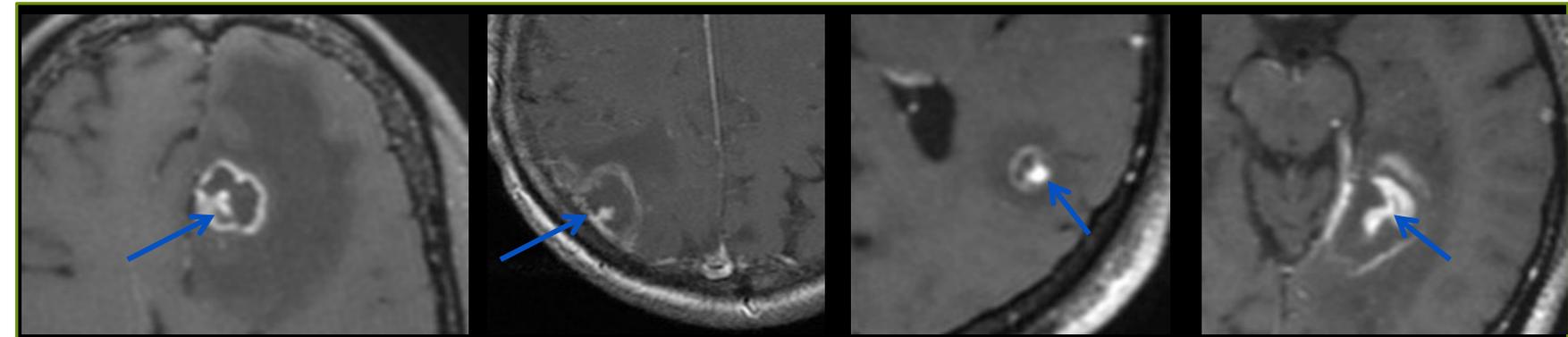
# Pseudotumeurs

HypoT2 = penser infection

Prise de contraste typique

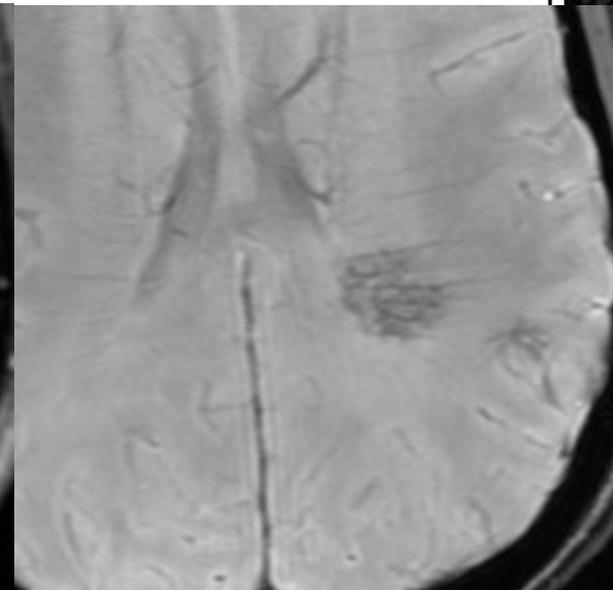
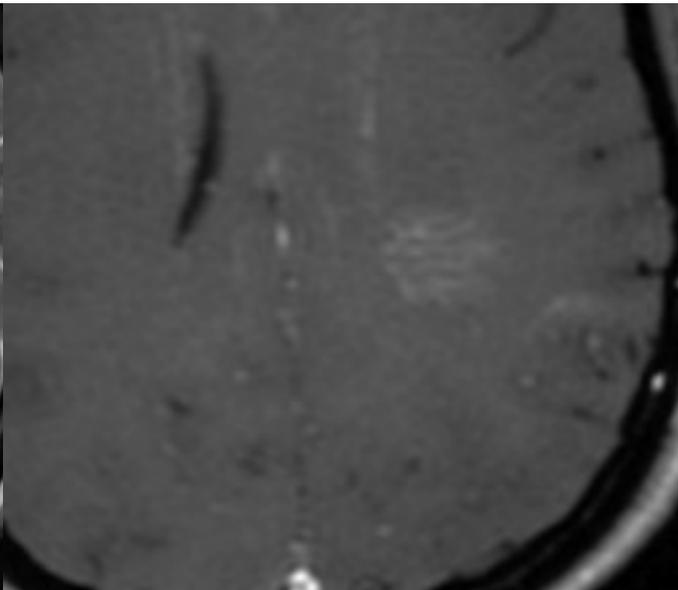
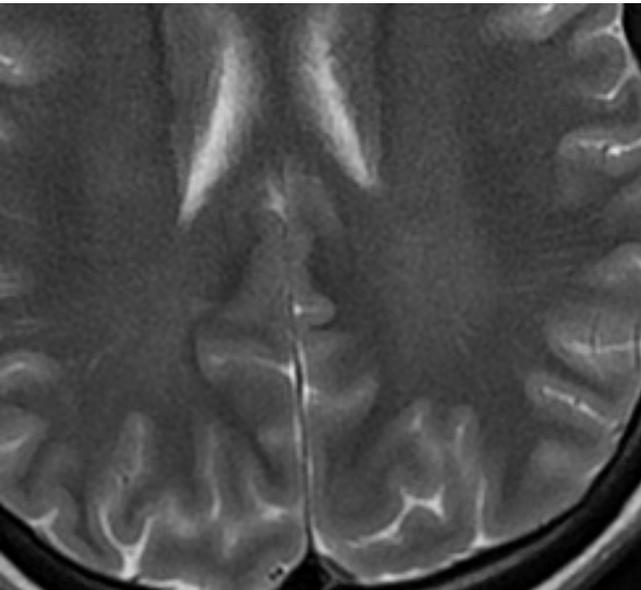


Eccentric target sign = toxoplasmose



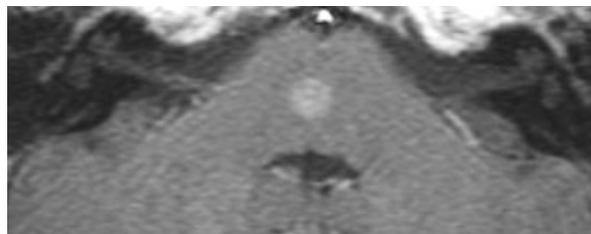
Pseudotumeurs

Aspects de pdc



Télangiectasie capillaire

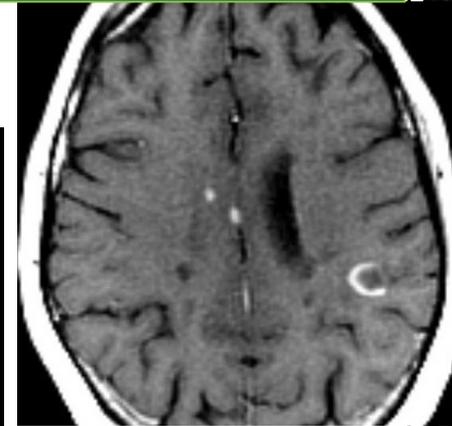
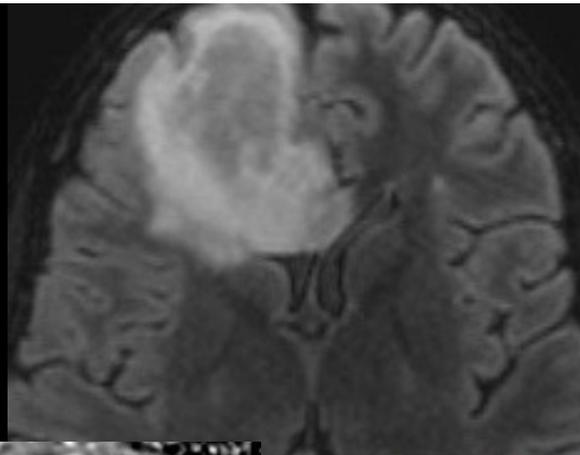
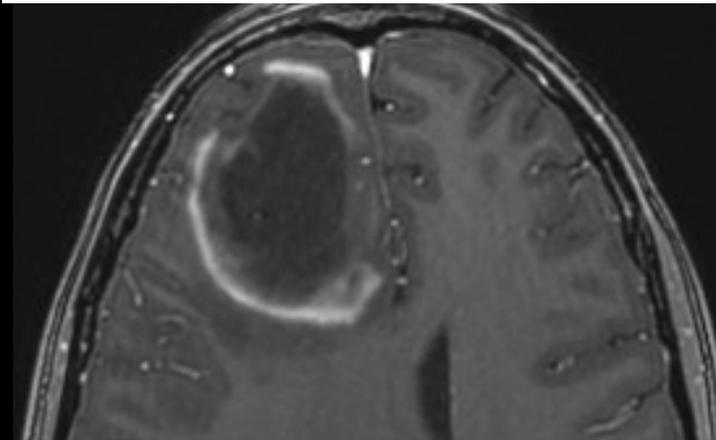
SWI +++



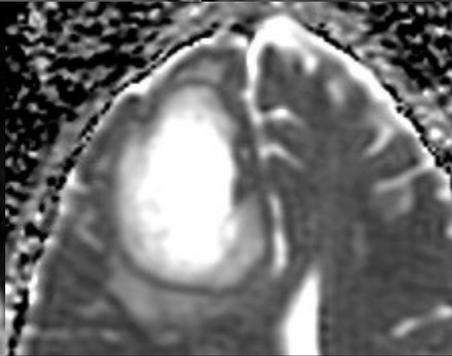
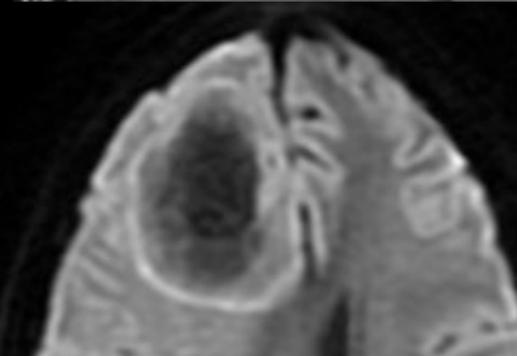
# Pseudotumeurs

## Aspects de pdc

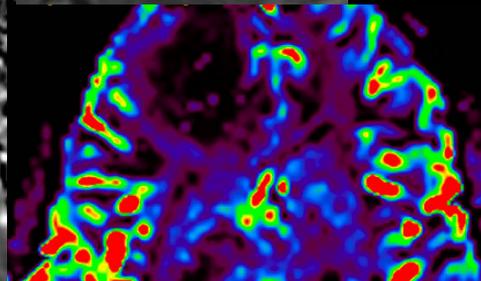
Anneau ouvert = démyélinisation



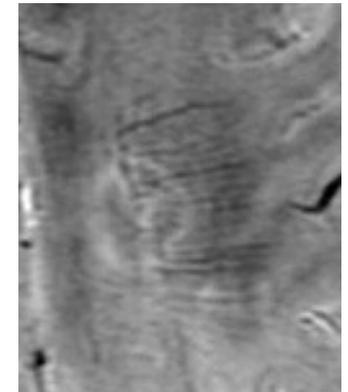
MS



restriction diffusion

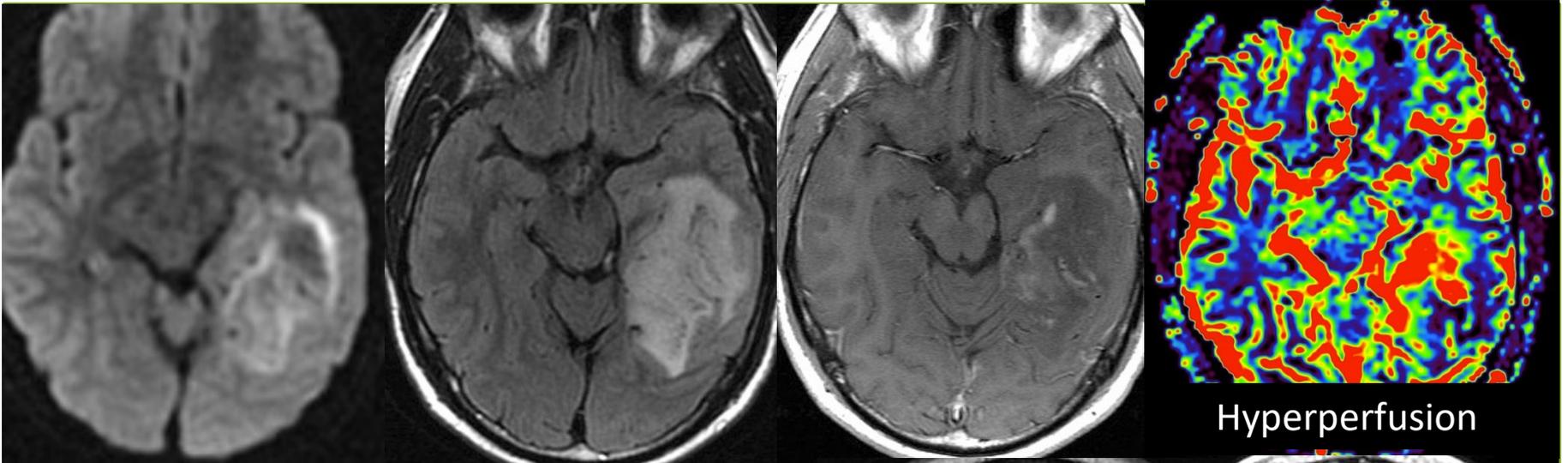


Pas hyperperfusé  
≠ glioblastome

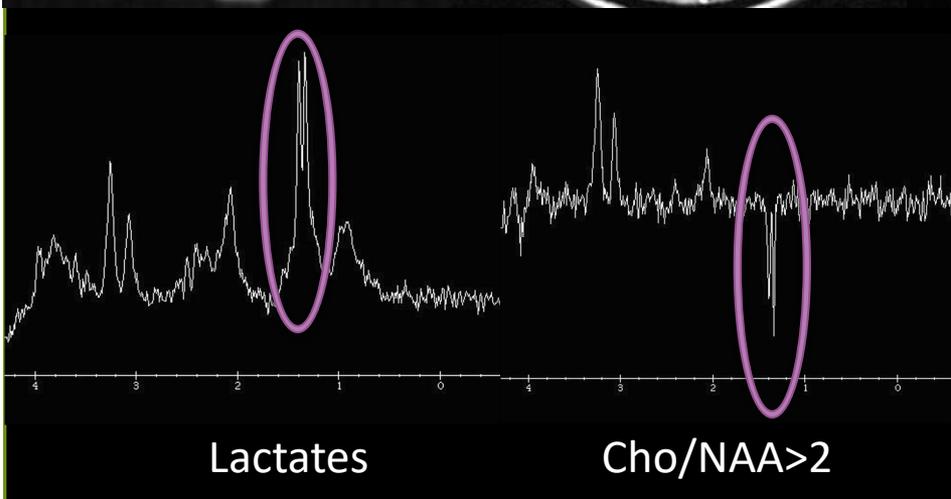


SWI

# Pseudotumeurs

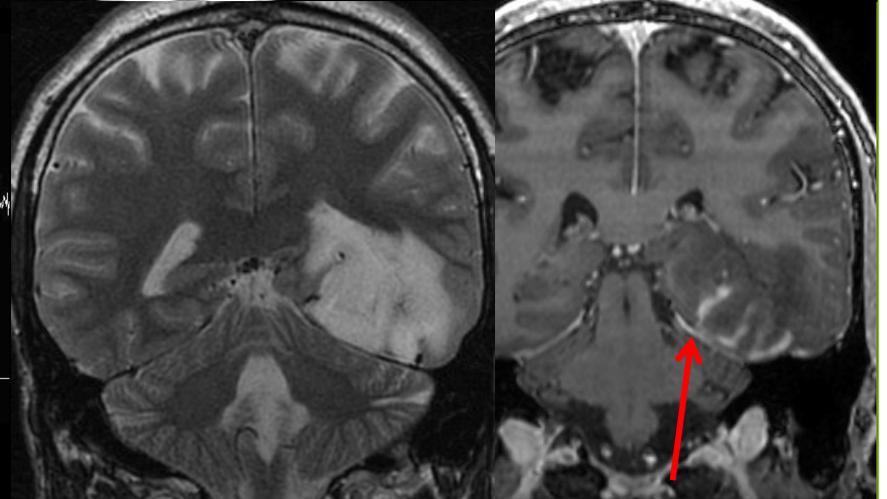


Hyperperfusion



Lactates

Cho/NAA > 2



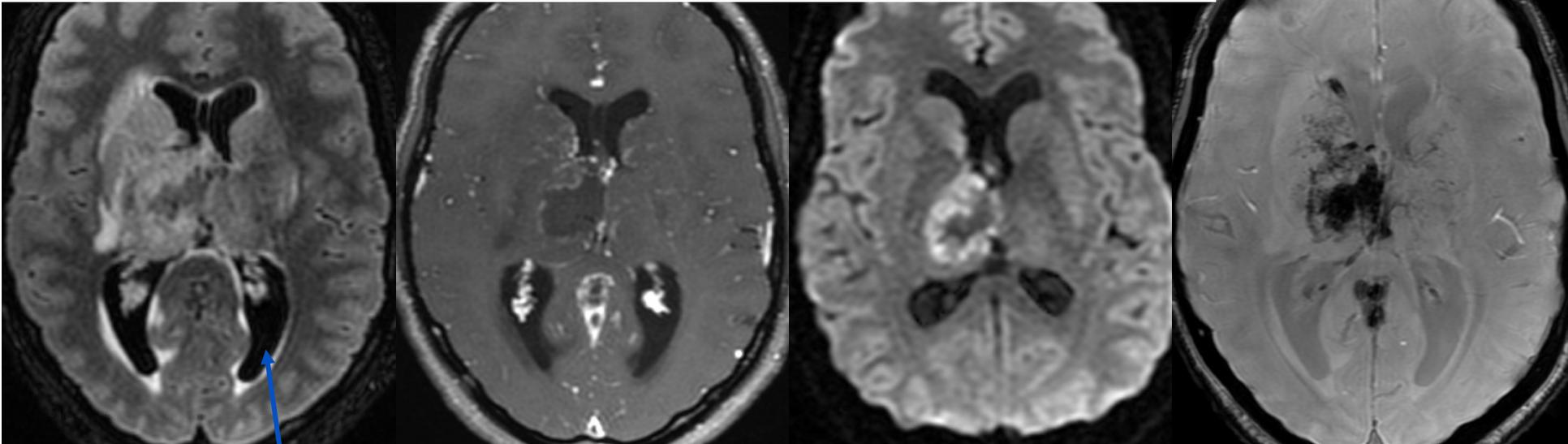
PDC gyriforme

## Pseudotumeurs : topographie

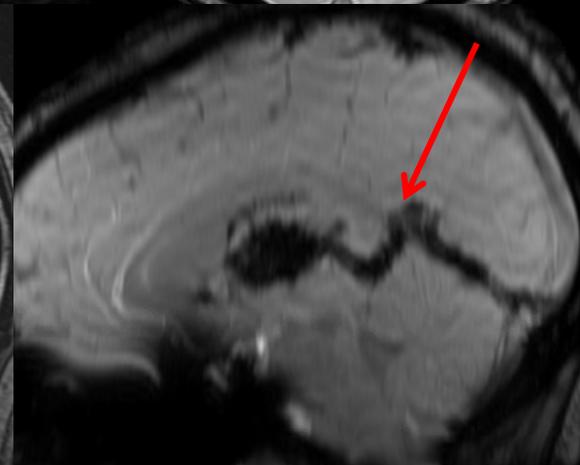
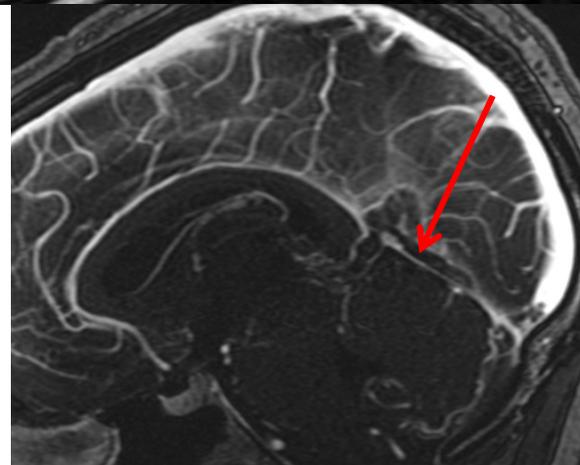
## Thrombose veineuse profonde

SWI +++thrombus

HS Bithalamique et NGC + hypertrophie + nécrose ischémohémorragique



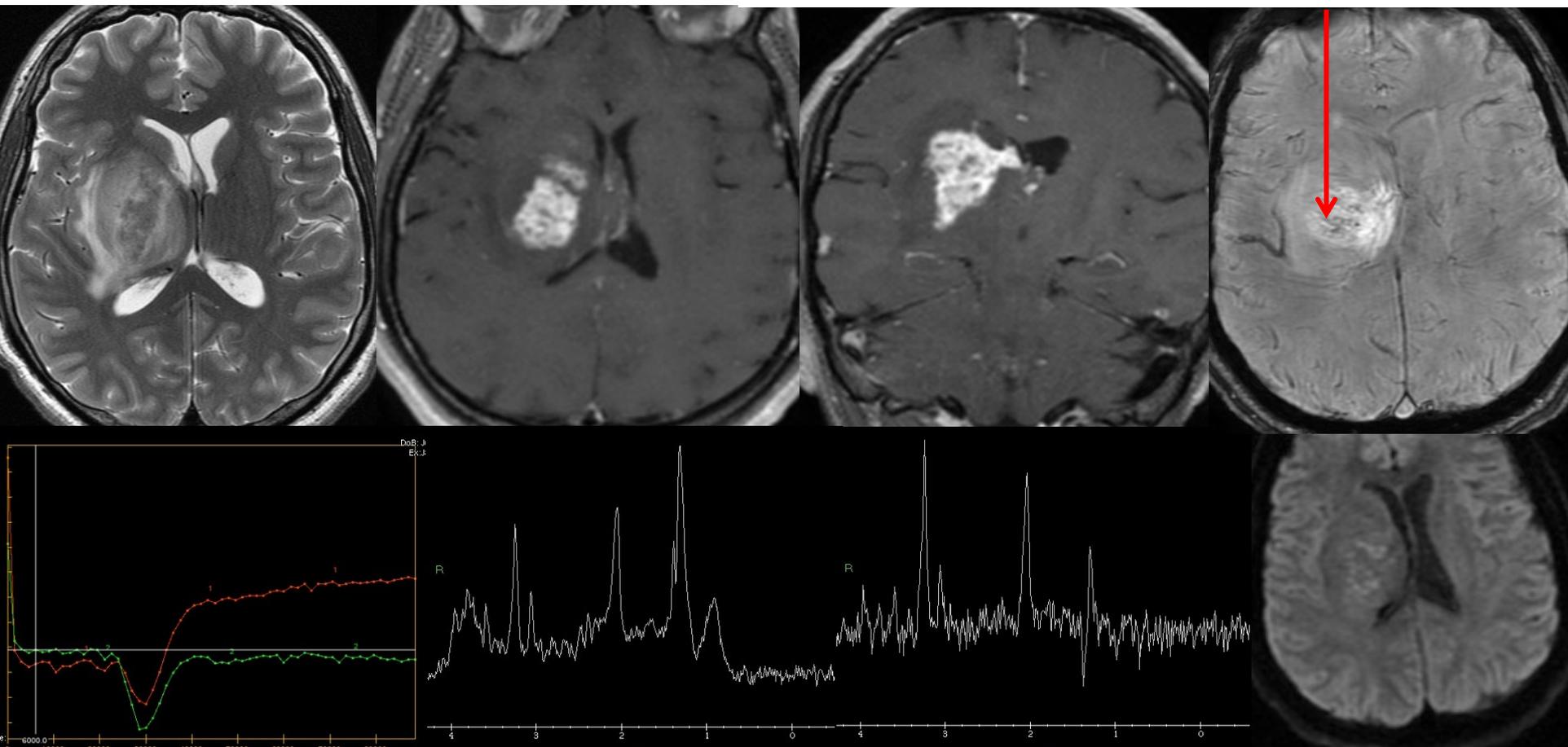
Hydrocéphalie  
= obstruction Monro



## Pseudotumeurs : topographie

## Neuro-Behçet

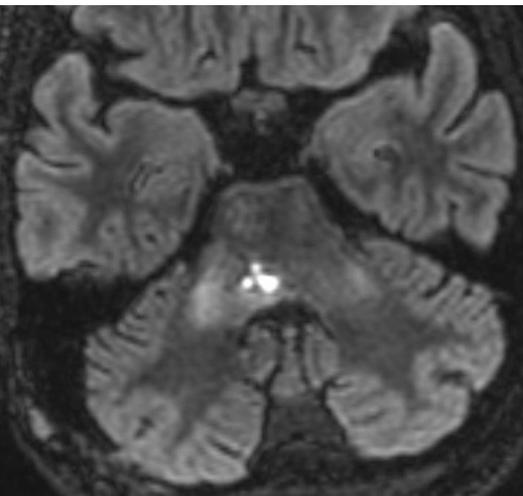
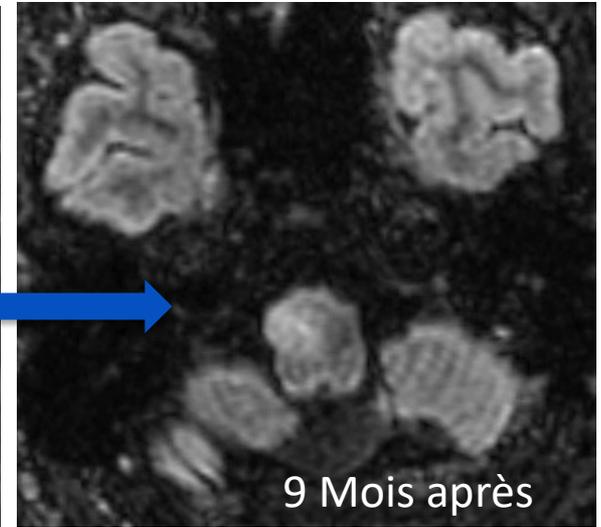
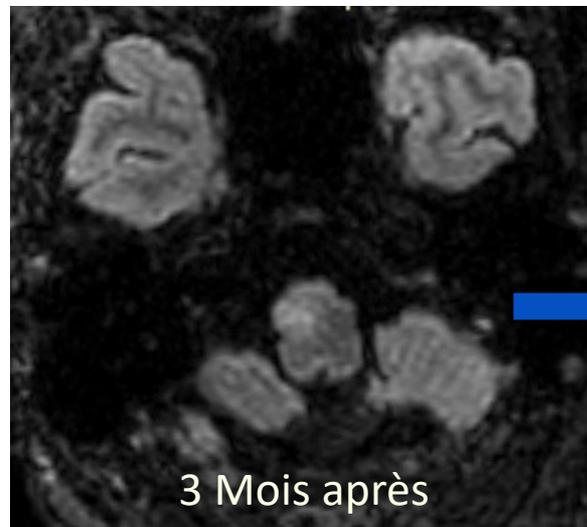
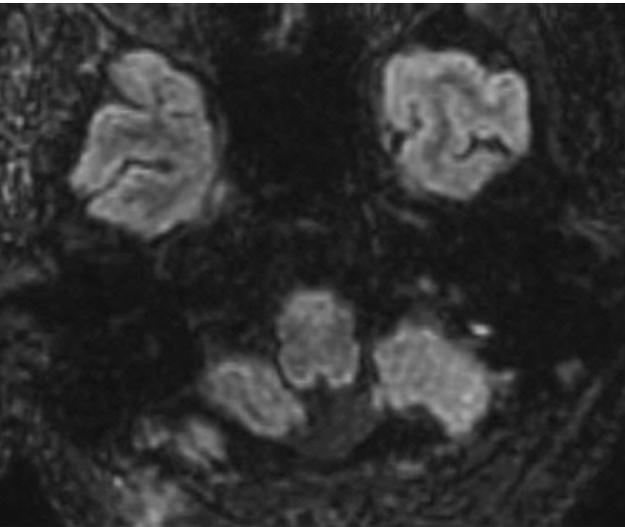
SWI = hémorragies, veines dilatées, thromboses



Capsule interne ++ NGC + thalamus

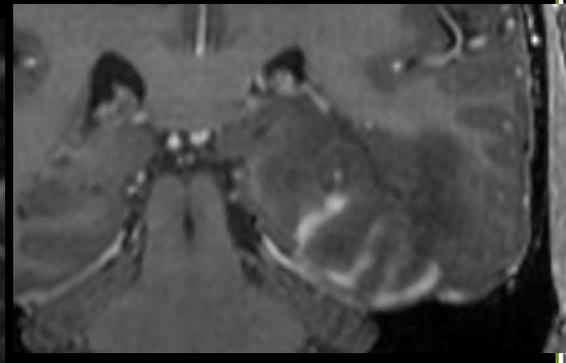
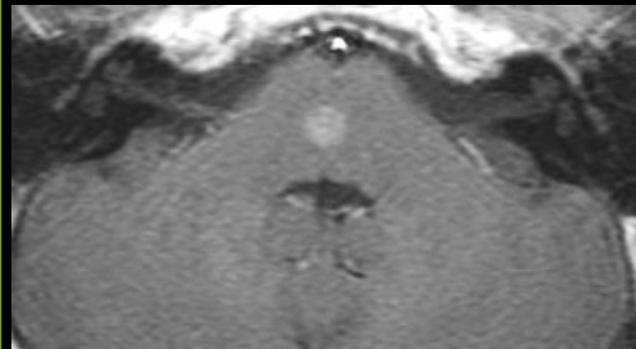
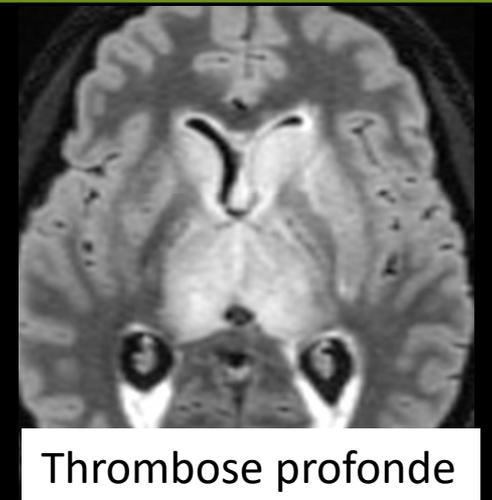
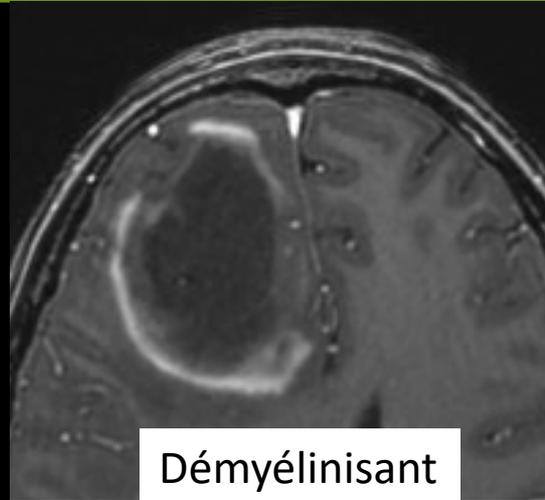
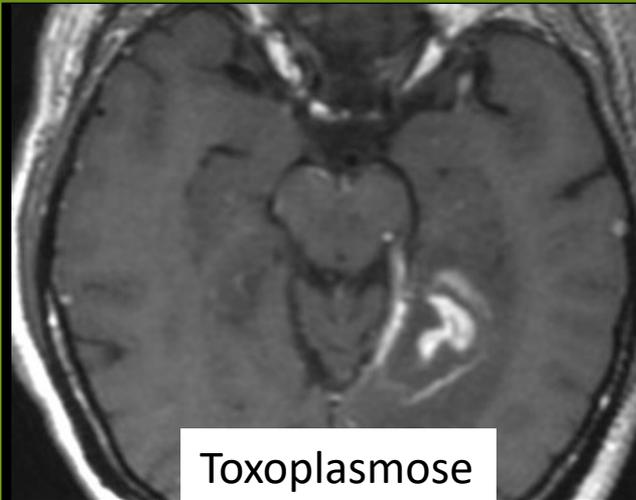
## Pseudotumeurs : topographie

## Dégénérescence hypertrophique olive bulbaire



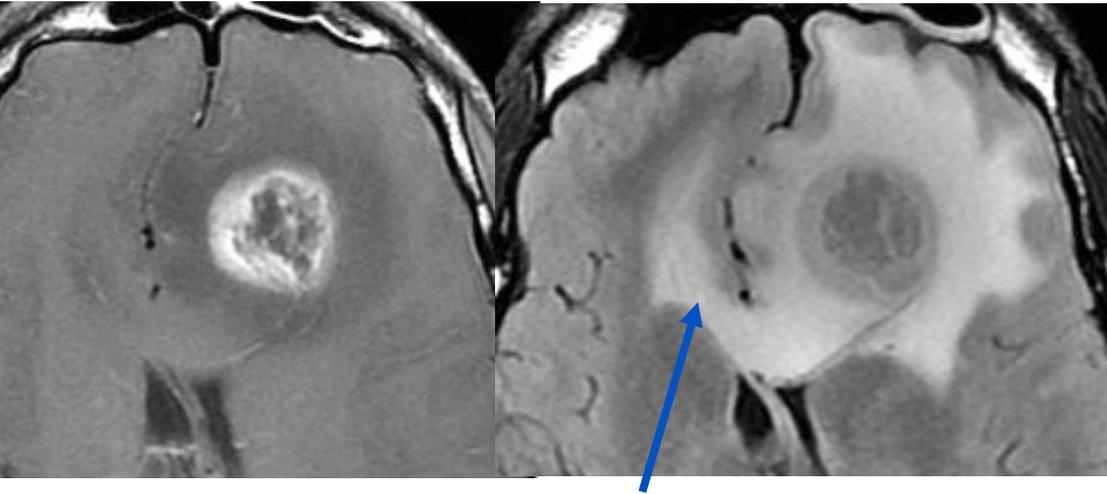
Dégénérescence trans-synaptique  
= interruption triangle Guillain-Mollaret  
→ hypertrophie transitoire olive  
bulbaire  
DDx= Gliome bulbaire hypertrophique

# Pseudotumeurs



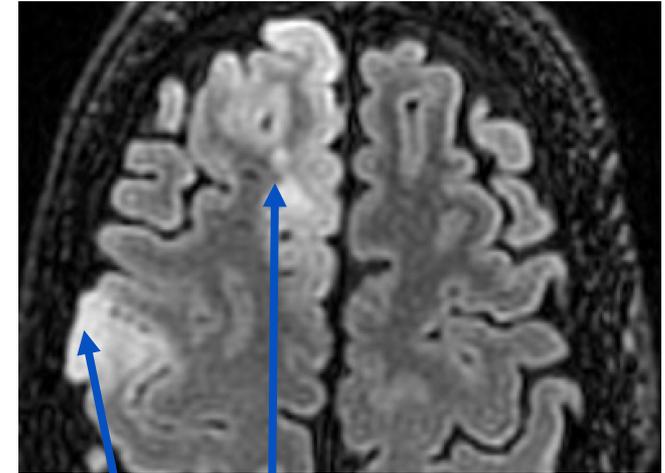
## Gliome ou métastase?

Œdème vasogénique ne franchit pas le corps calleux



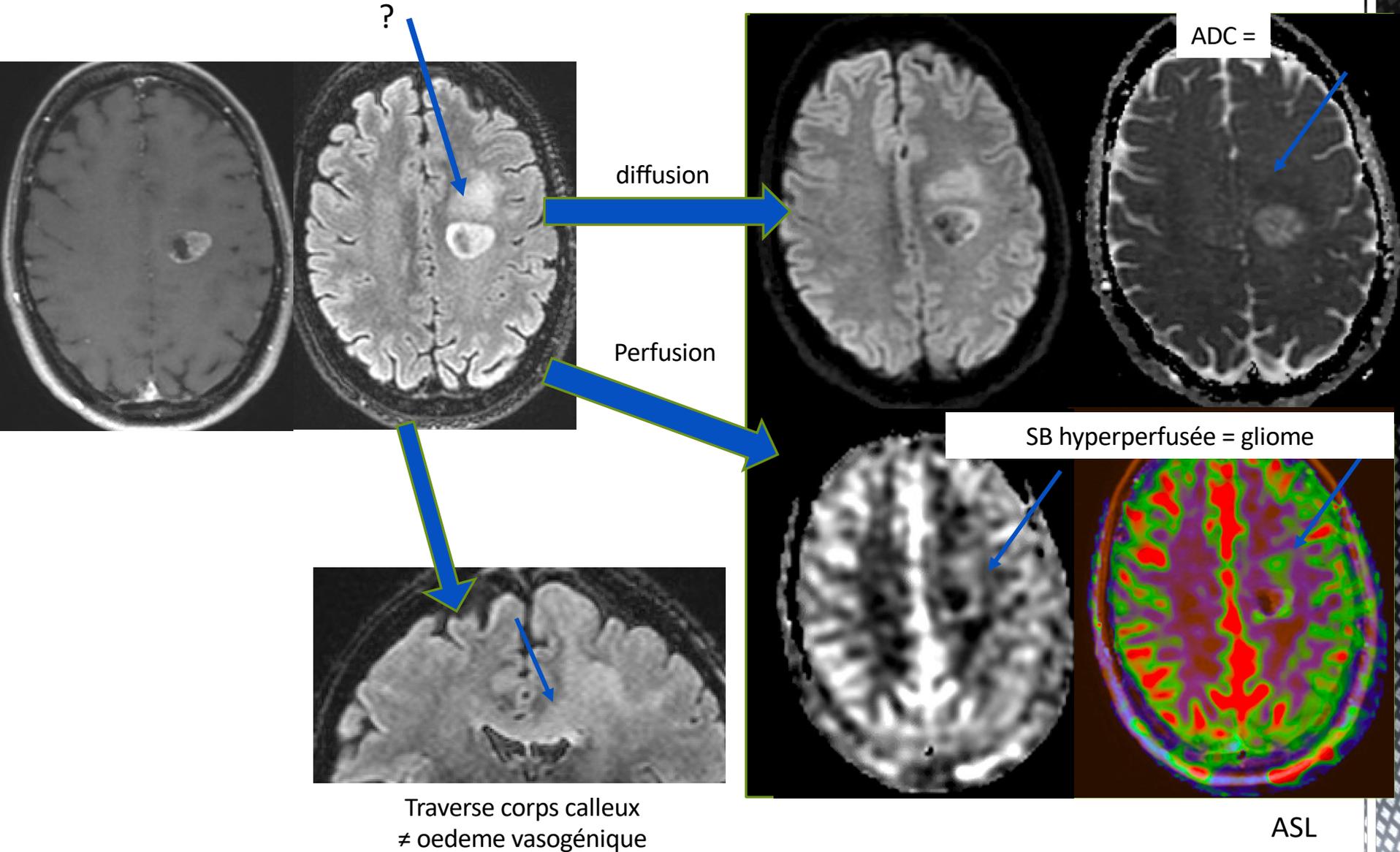
**= Gliome**

Œdème vasogénique n'infiltré pas le cortex



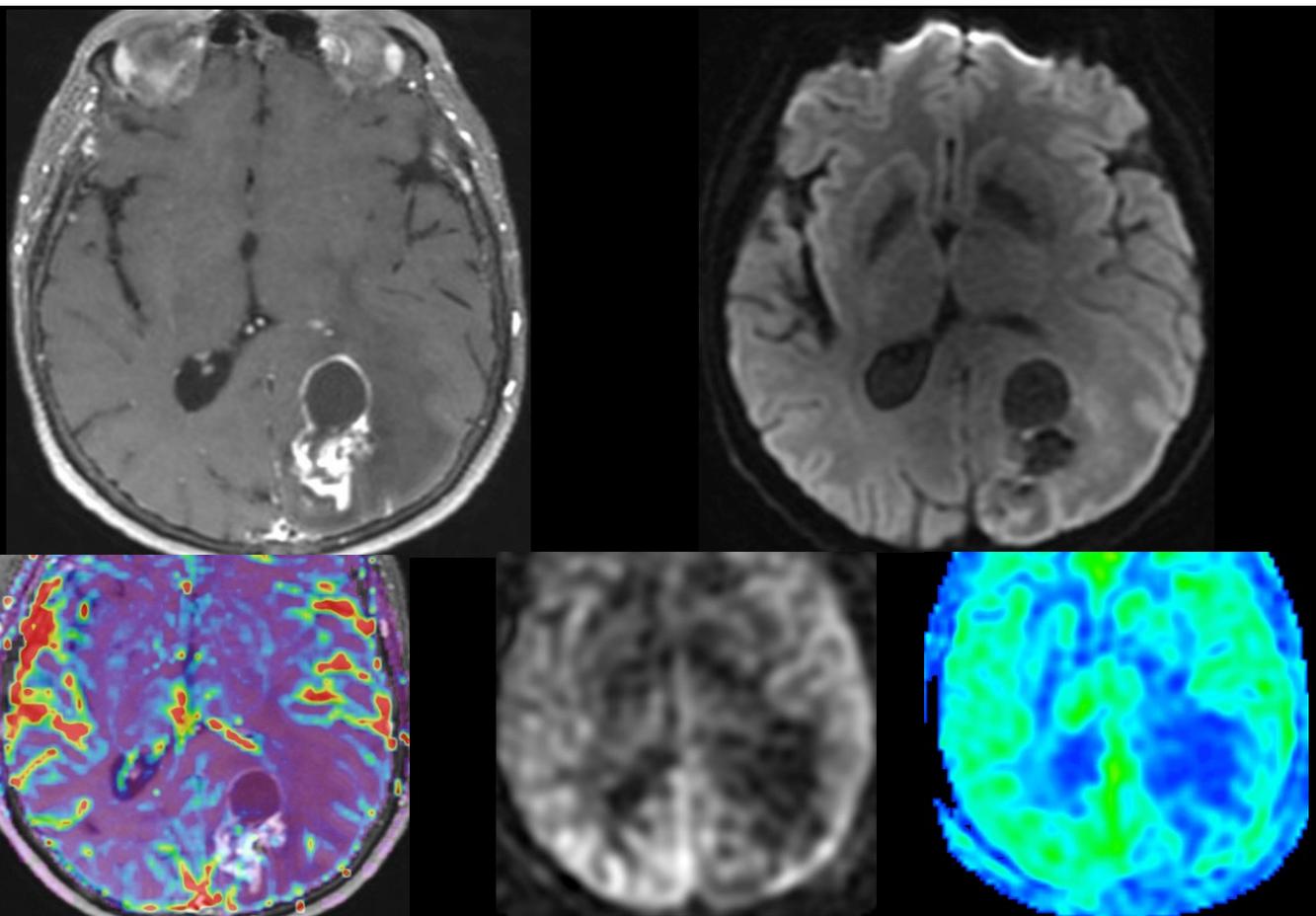
Infiltration corticale

# Œdème vasogénique ou infiltration ?



## Gliome ou métastase?

Coregistration +++ (cortex, vx)



Morphologie compatible  
Avec gliome

Pas d'hyperperfusion

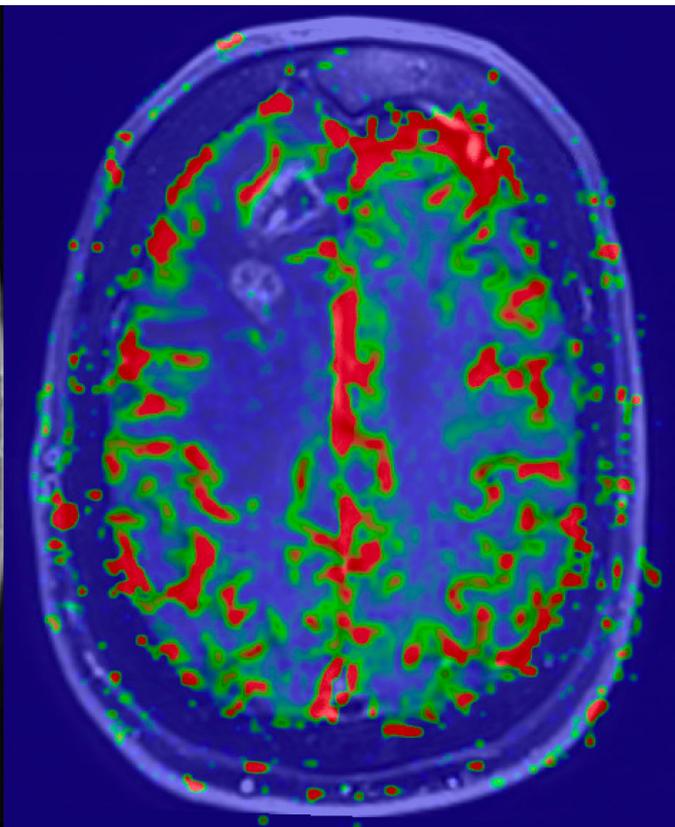
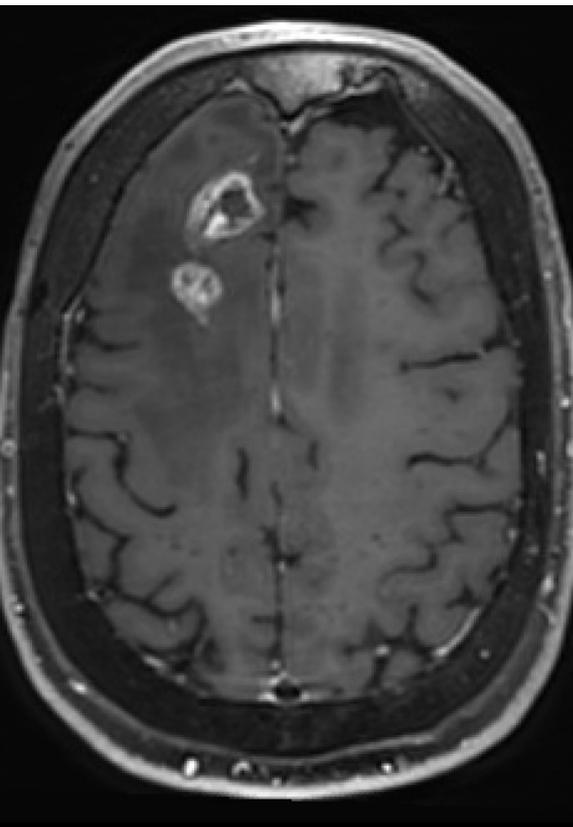
**Métastase**

DSC rCBV

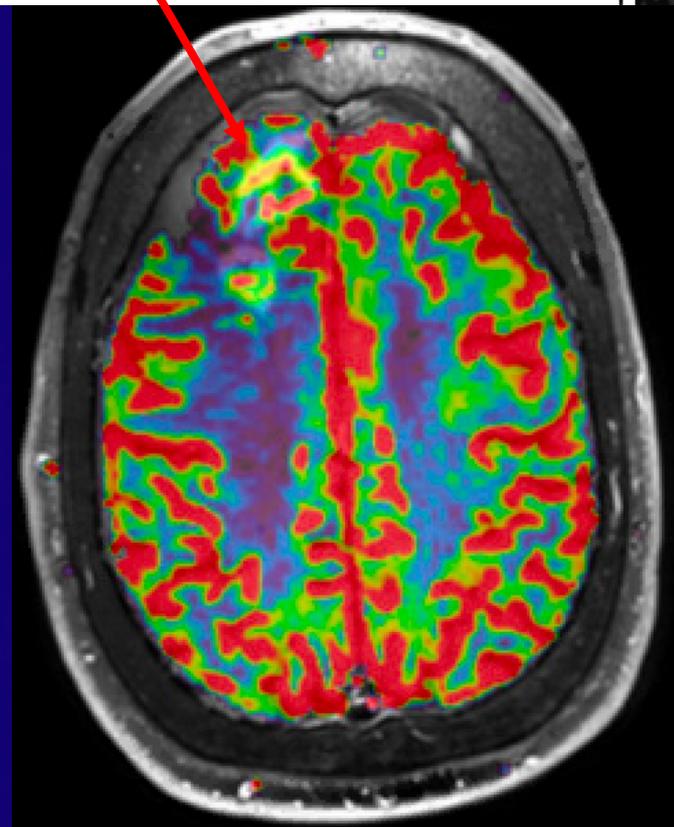
ASL

ASL CBF

# DSC perfusion : leakage



Coregistration = rCBV  
Sans correction leakage

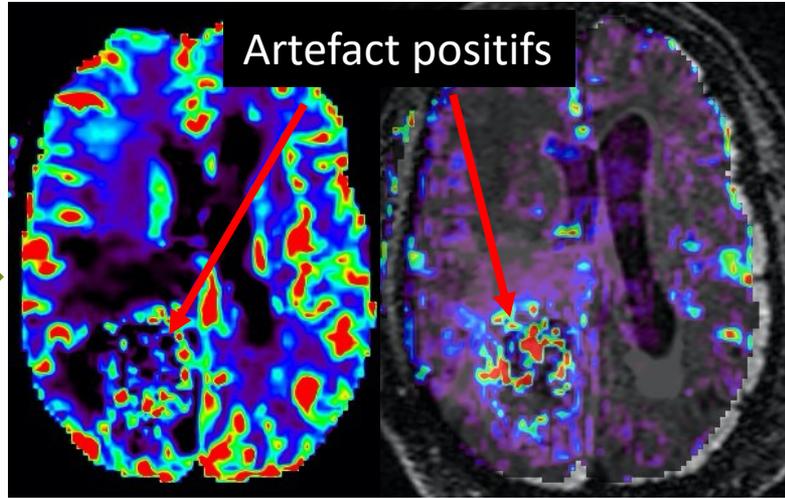


Coregistration + rCBV  
Avec correction leakage

# DSC perfusion : lésion hémorragique



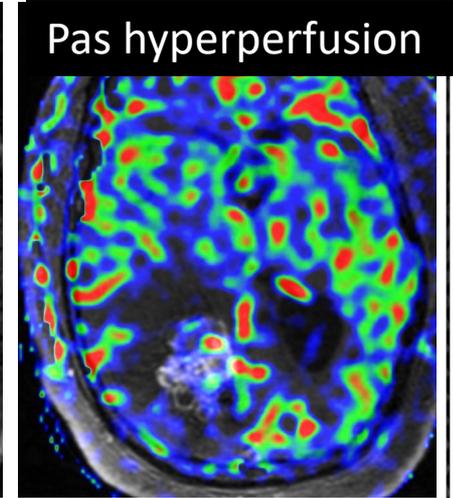
Native DSC image



Artefact positifs

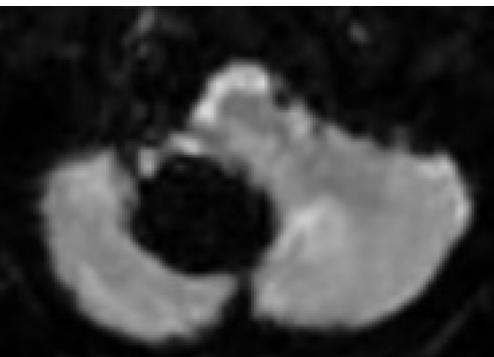
rCBV

K2

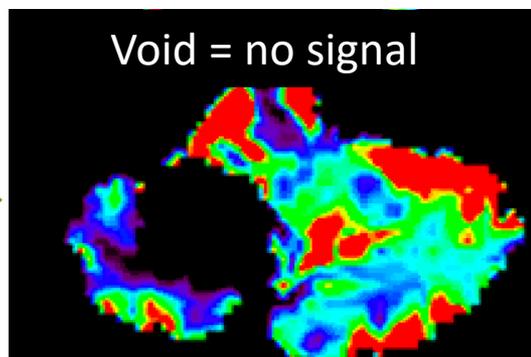


Pas hyperperfusion

ASL

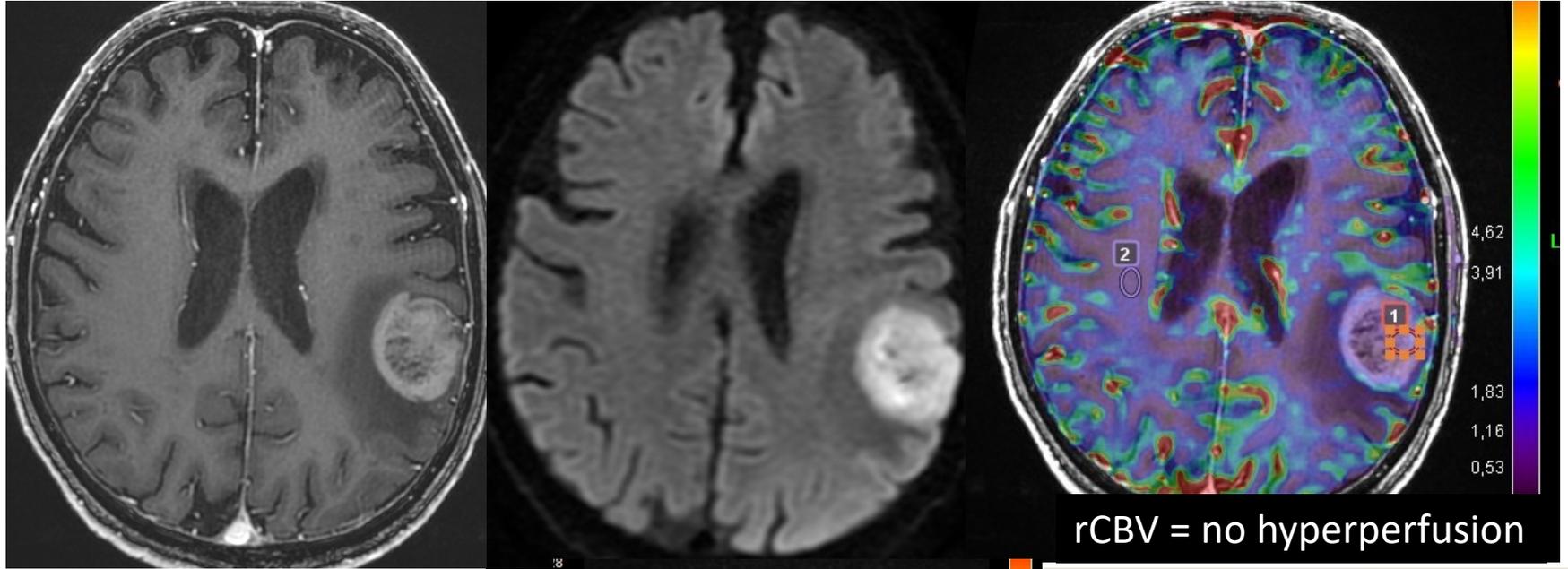


Native DSC image



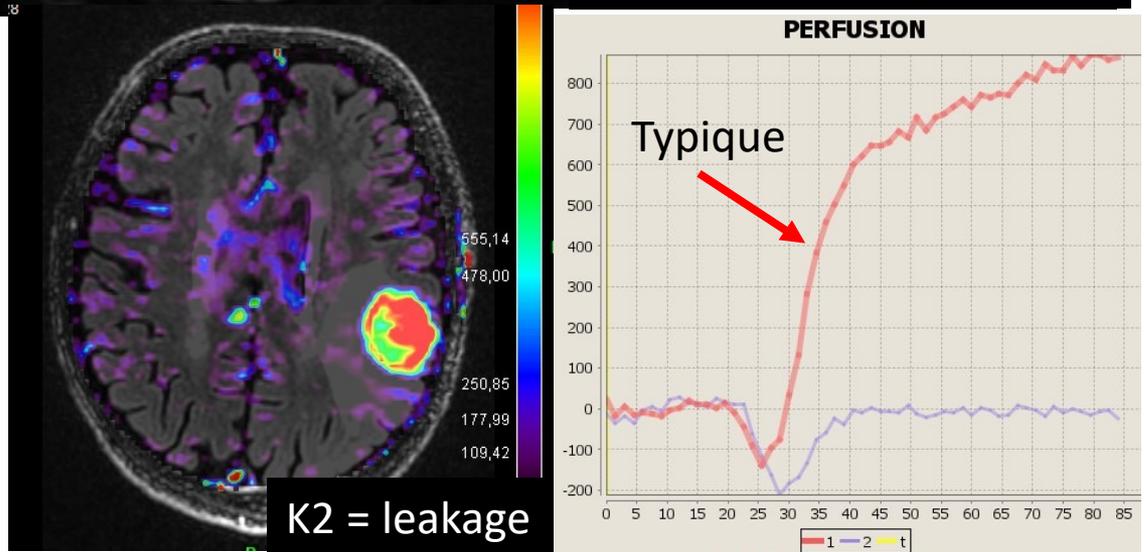
Void = no signal

rCBV

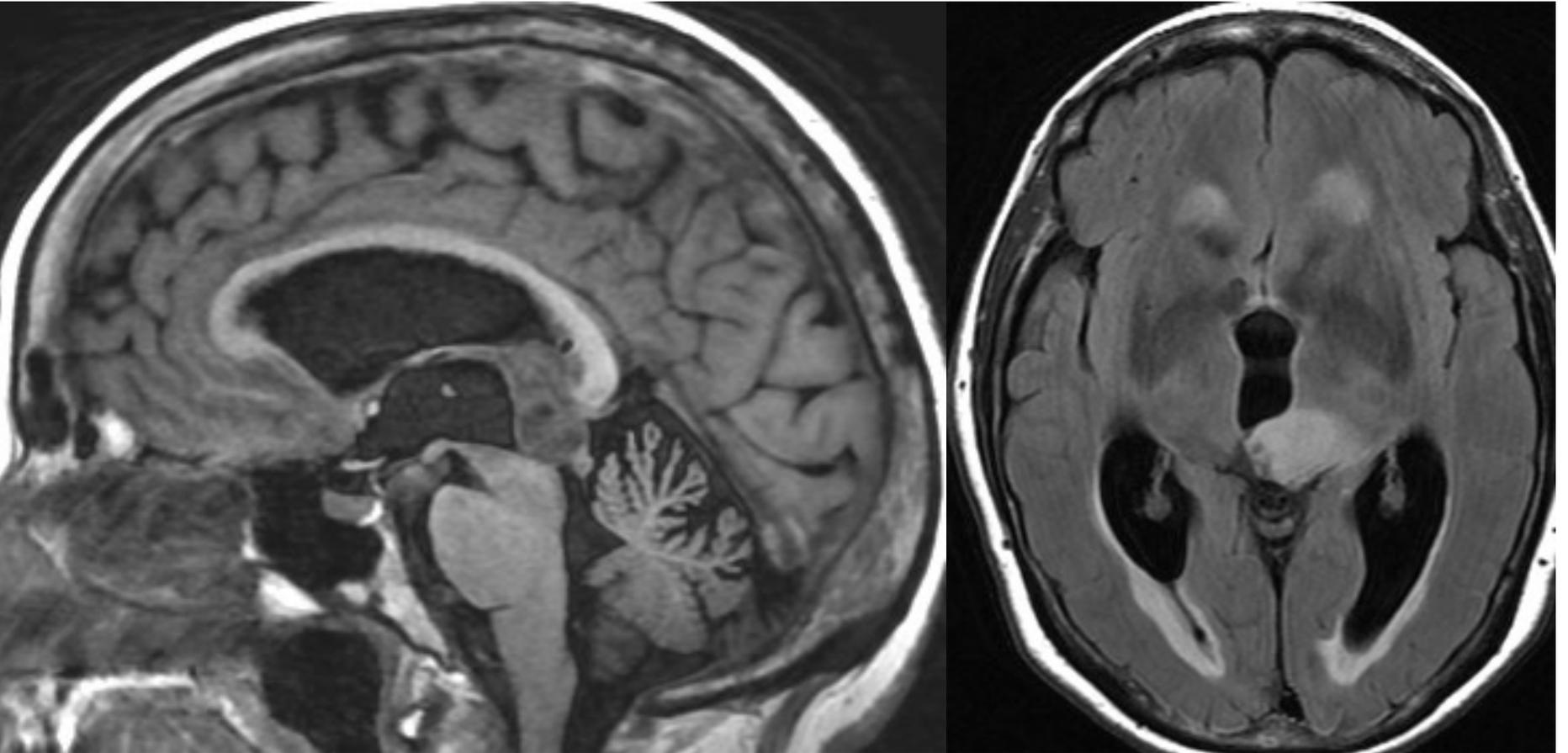


Lymphome nécrotique  
= corticoïdes  
3 jours avant MRI

Perfusion aide+++



## Identifier l'urgence = Hydrocéphalie



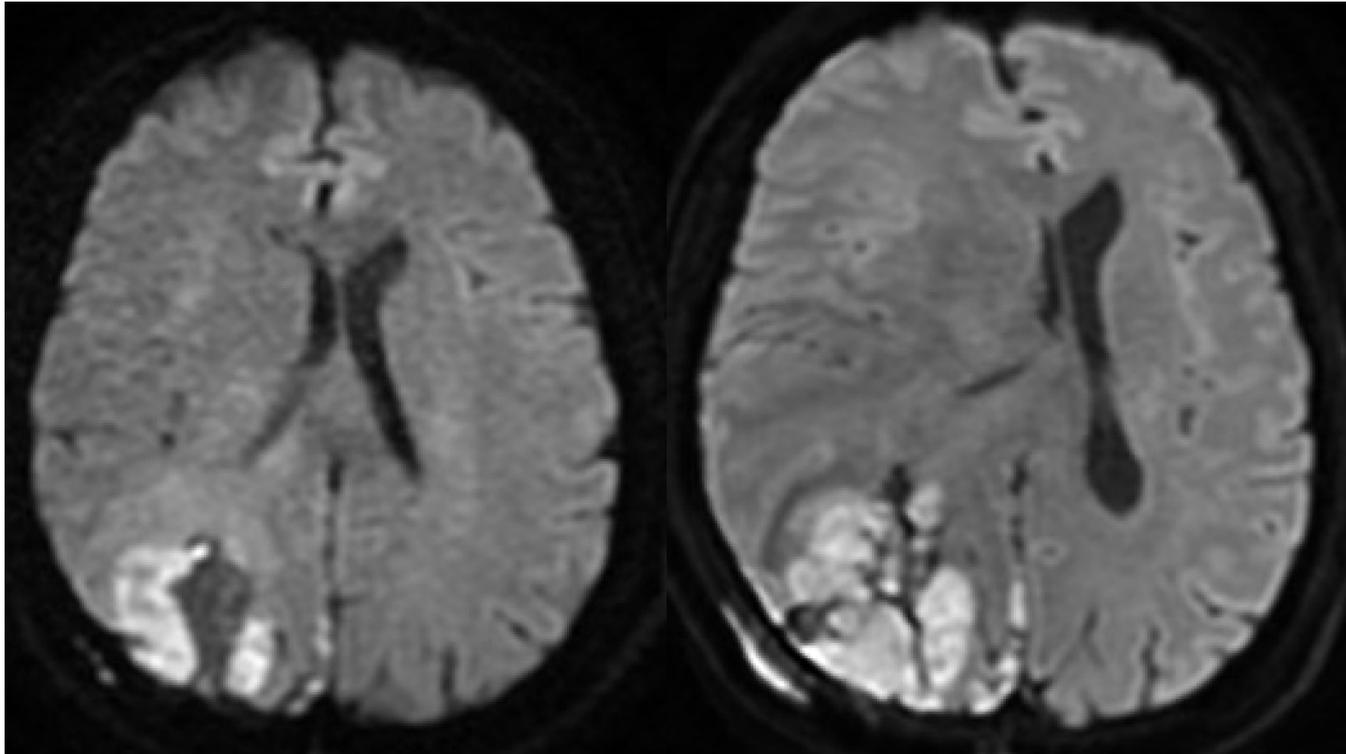
Lésion tectale + hydrocéphalie

**= Urgence vitale**

décès par hydrocéphalie = rapide ++ même si lésion causale lentement évolutive

## Gliome : diffusion reflète cellularité

Savoir quand contrôle rapproché/ prévenir clinicien = diffusion



Sarcome

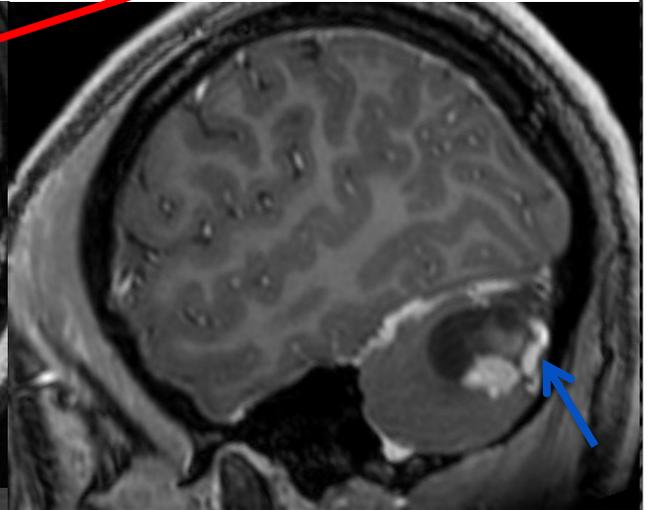
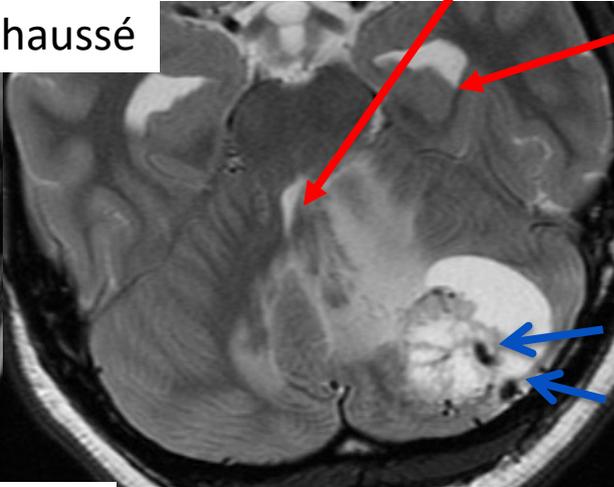
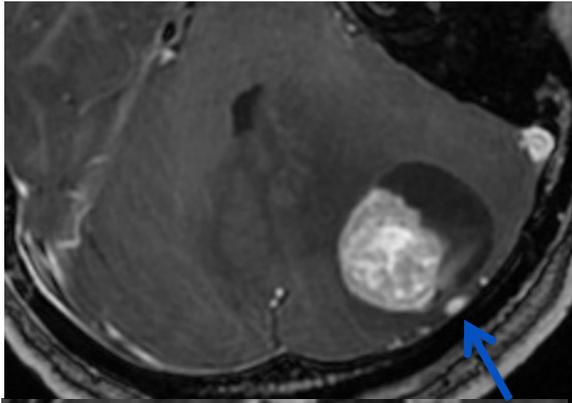
13 jours après

Chercher, voir, décrire vx = risque chirurgical +++

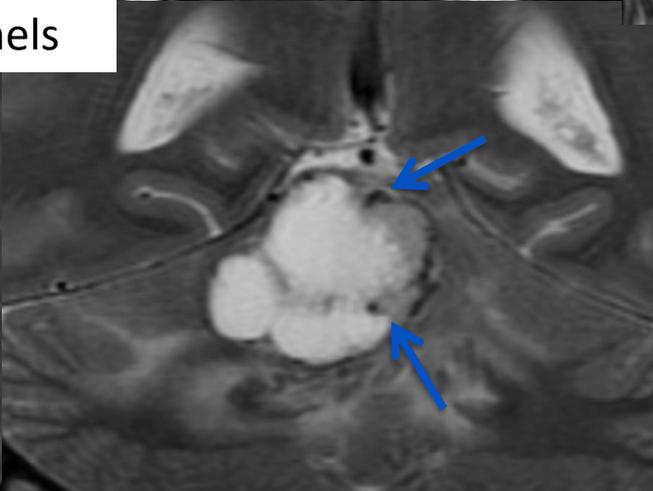
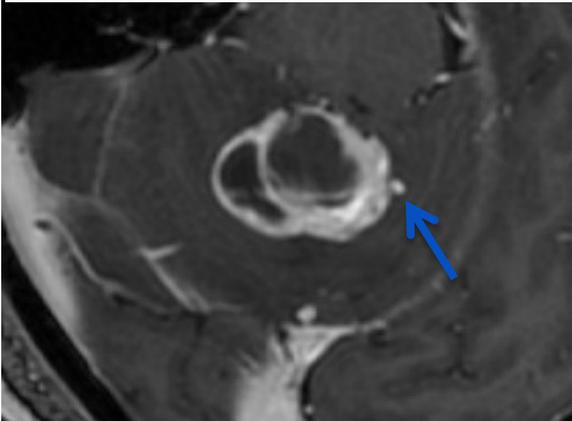
Hémangioblastomes

Mass effect + hydrocéphalie

Nodule mural + kyste non réhaussé



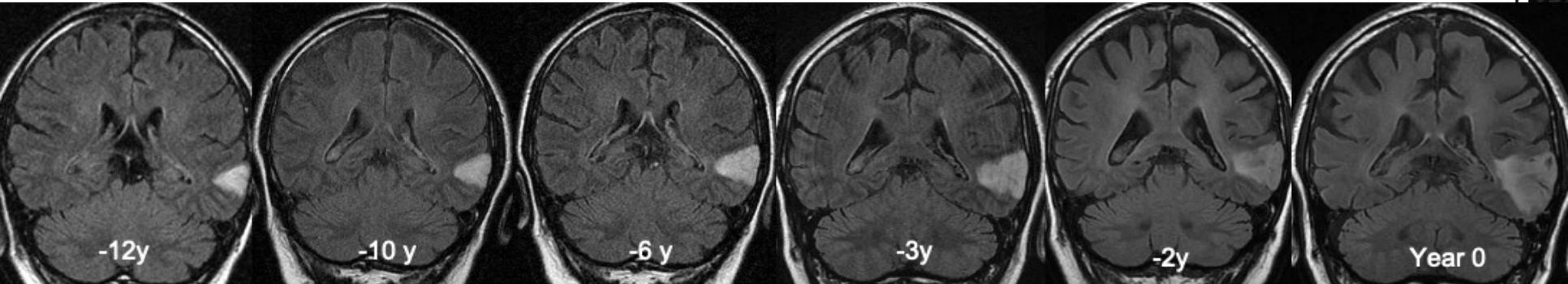
Nodule + kystes intralésionnels



Flow voids

## Gliome bas grade, lésion lentement évolutive

Evolution = IRM ancienne, pas juste précédente.



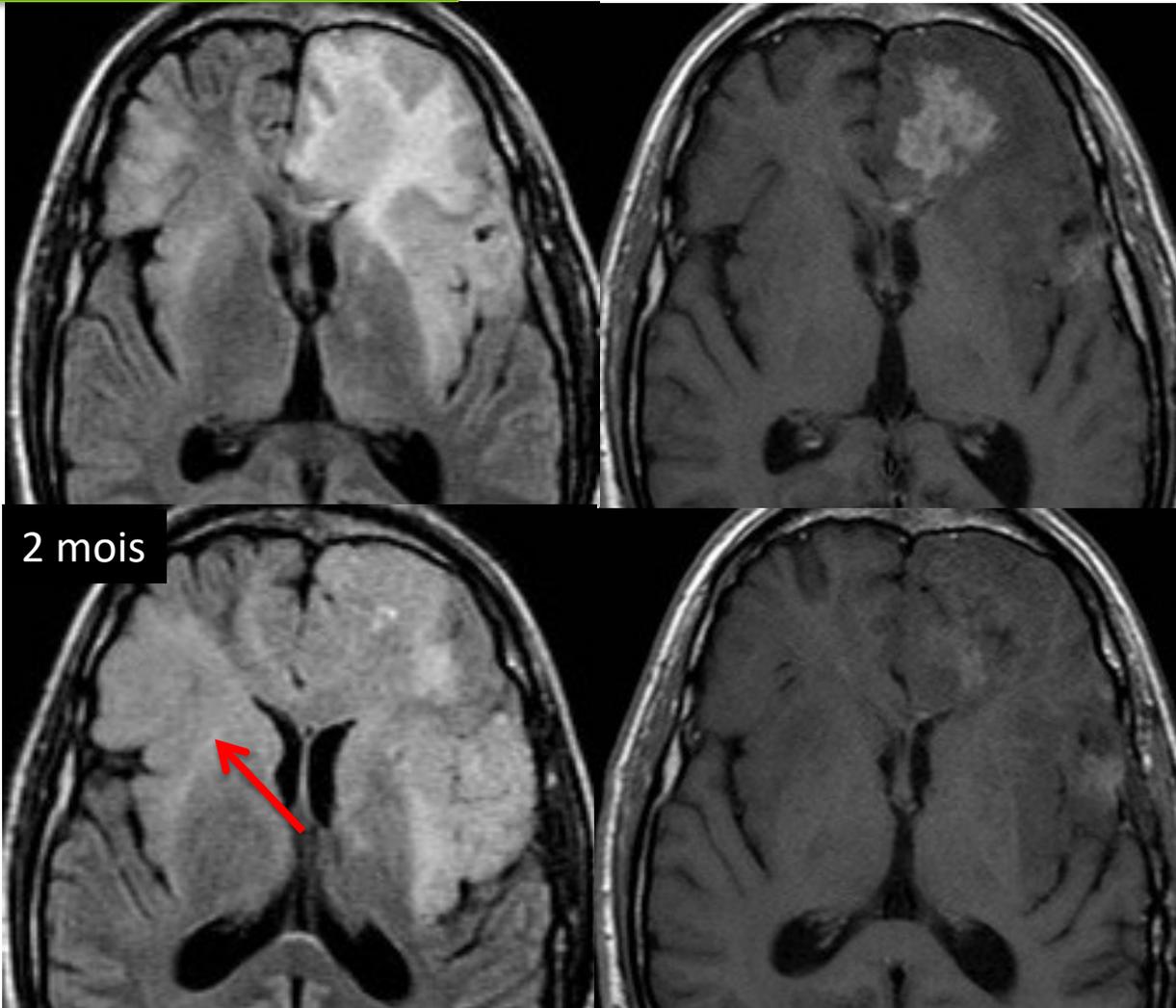
GBG pris pour dysplasie

IRM toujours comparées avec précédente.

### Gliome bas grade

Croissance lente puis accélération 6 mois avant transformation anaplasique

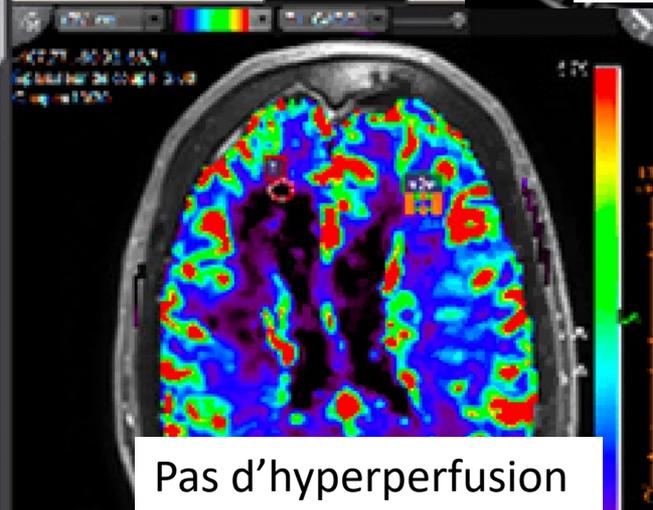
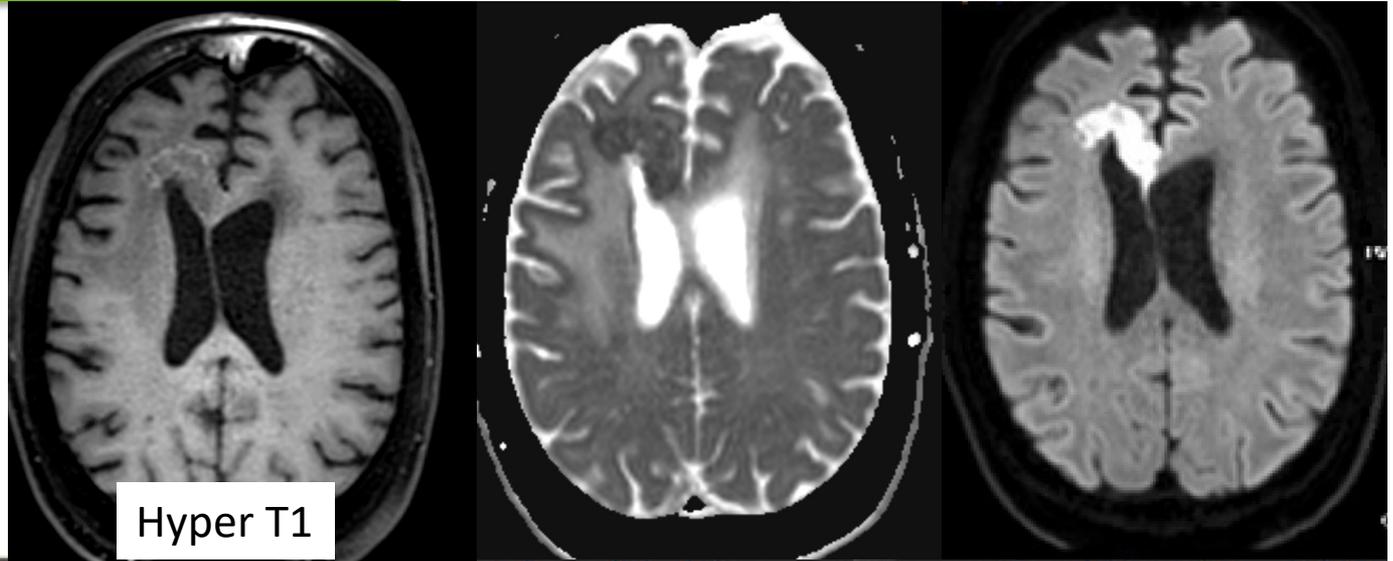
## Pseudoréponse

Bevacizumab  
Avastin

Disparition pdc= 0  
MAIS infiltration tumorale ↑

# Bevacizumab= avastin

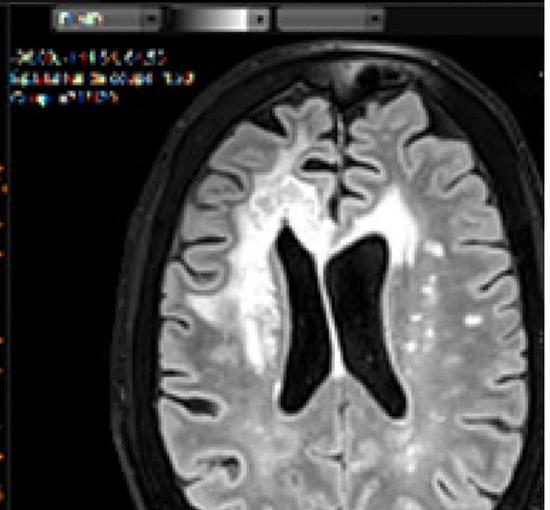
Nécrose gélatineuse / de coagulation



Pas d'hyperperfusion



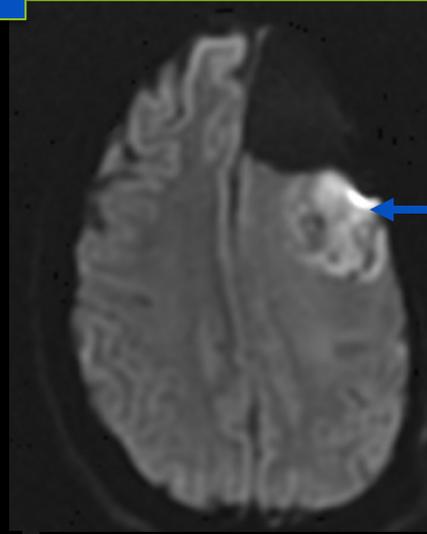
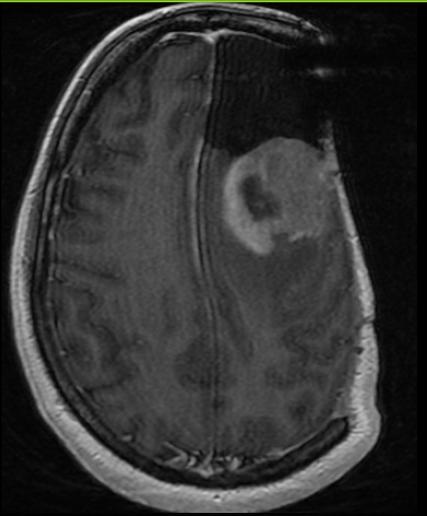
Non rehaussée



# Bevacizumab

## Nécrose gélatineuse / de coagulation

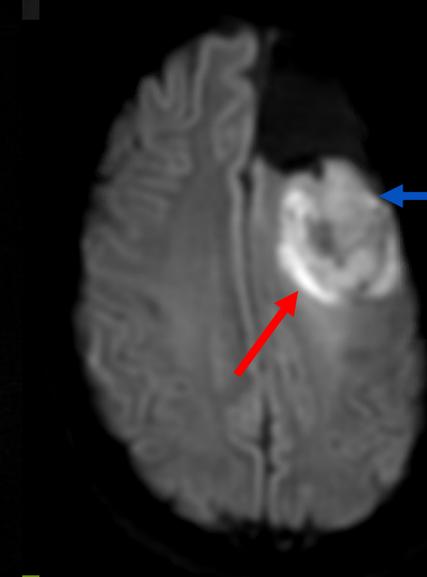
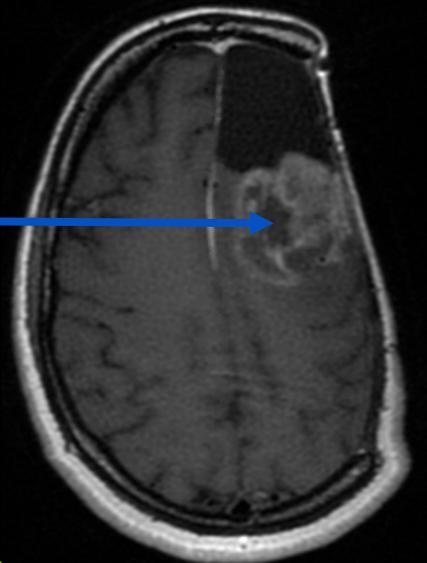
Avant BV



DWI: hypercellularité

Après BV

CEL ↓

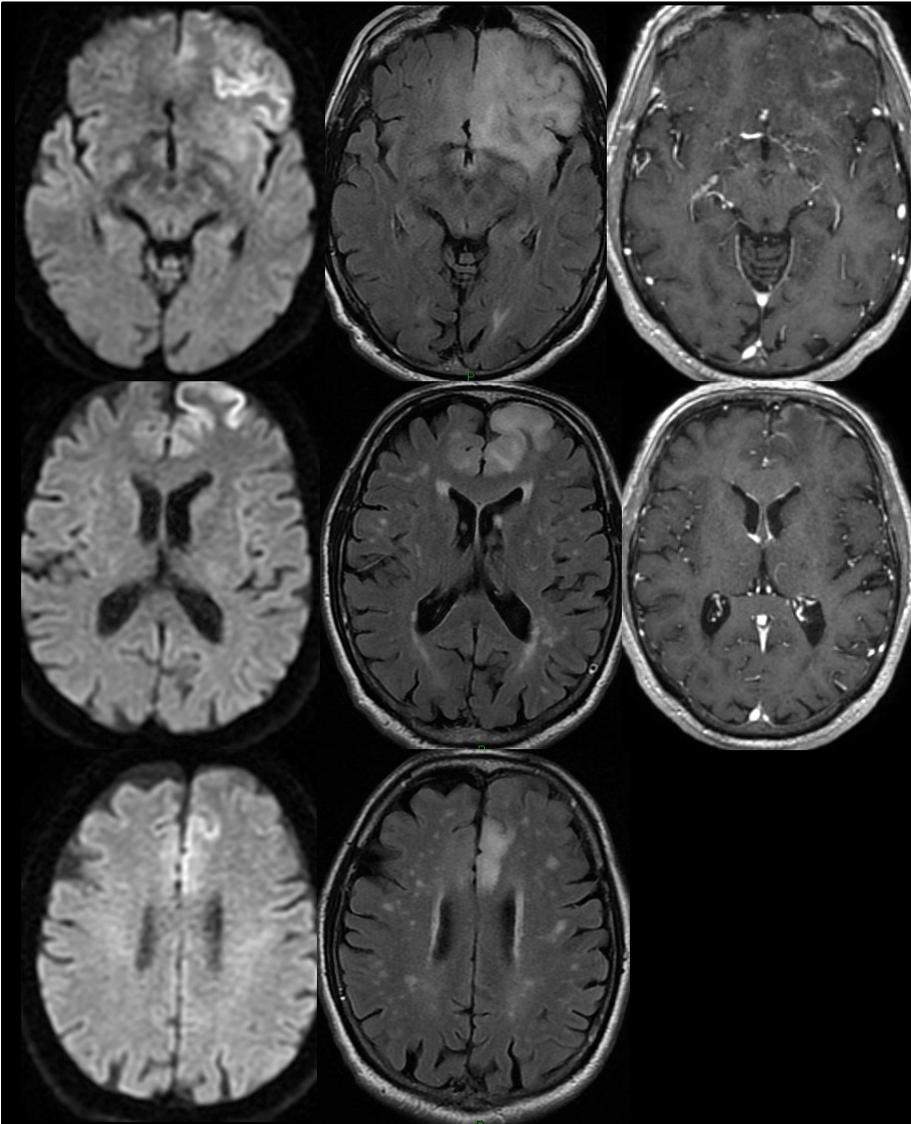


DWI: hypercellularité  
+ Nécrose gélatineuse

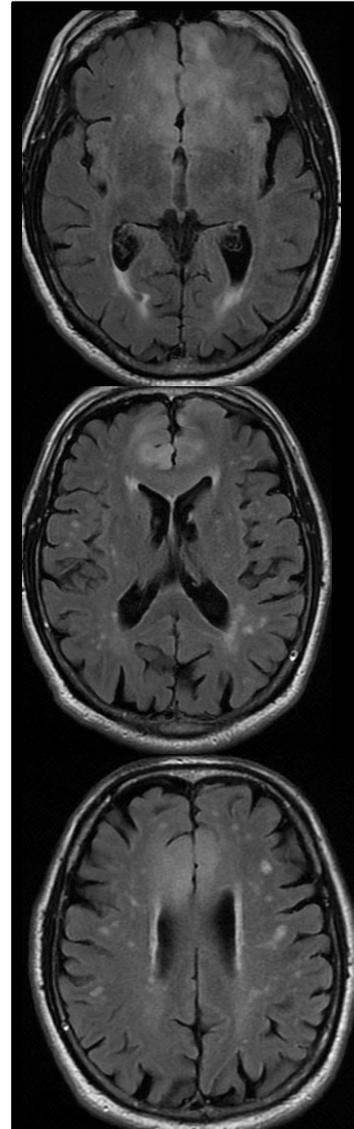
= plus hyperintense que  
L'hypersignal  
d'hypercellularité

# Etat de mal

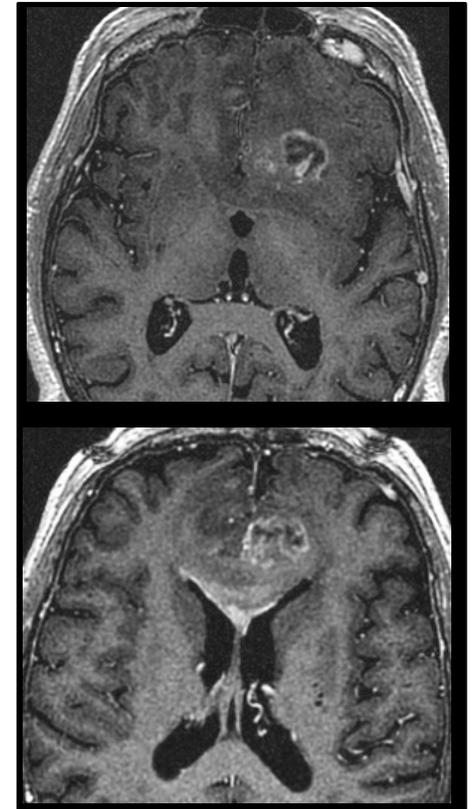
Crise



2 mois

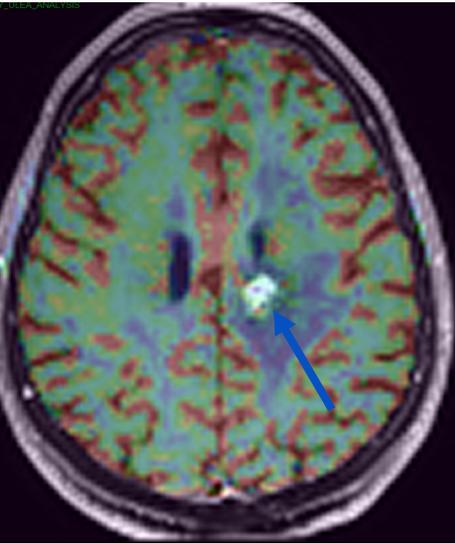


1 mois

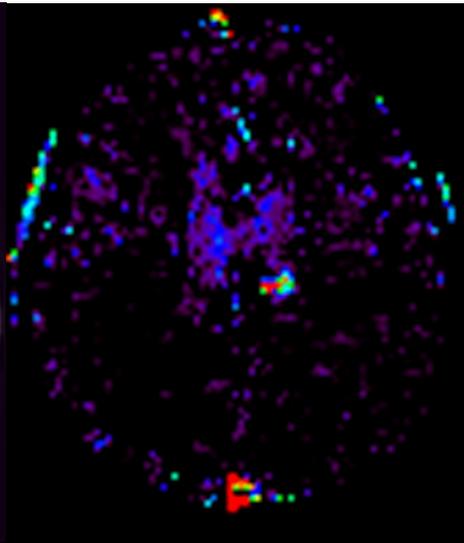


## Radionécrose

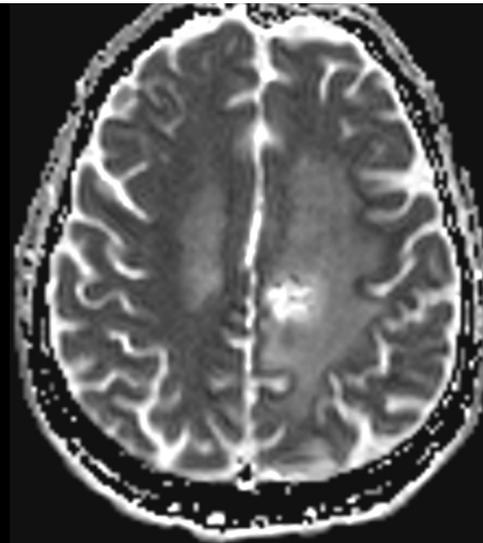
## Glioblastome : Nouvelle lésion réhaussée



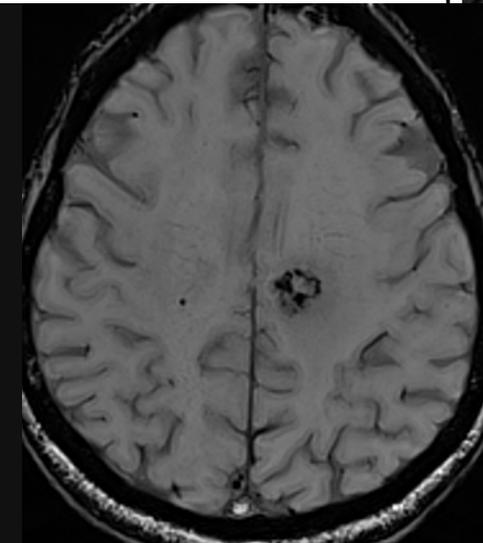
rCBV  
= pas d'hyperperfusion



K2 = perméabilité  
= leakage

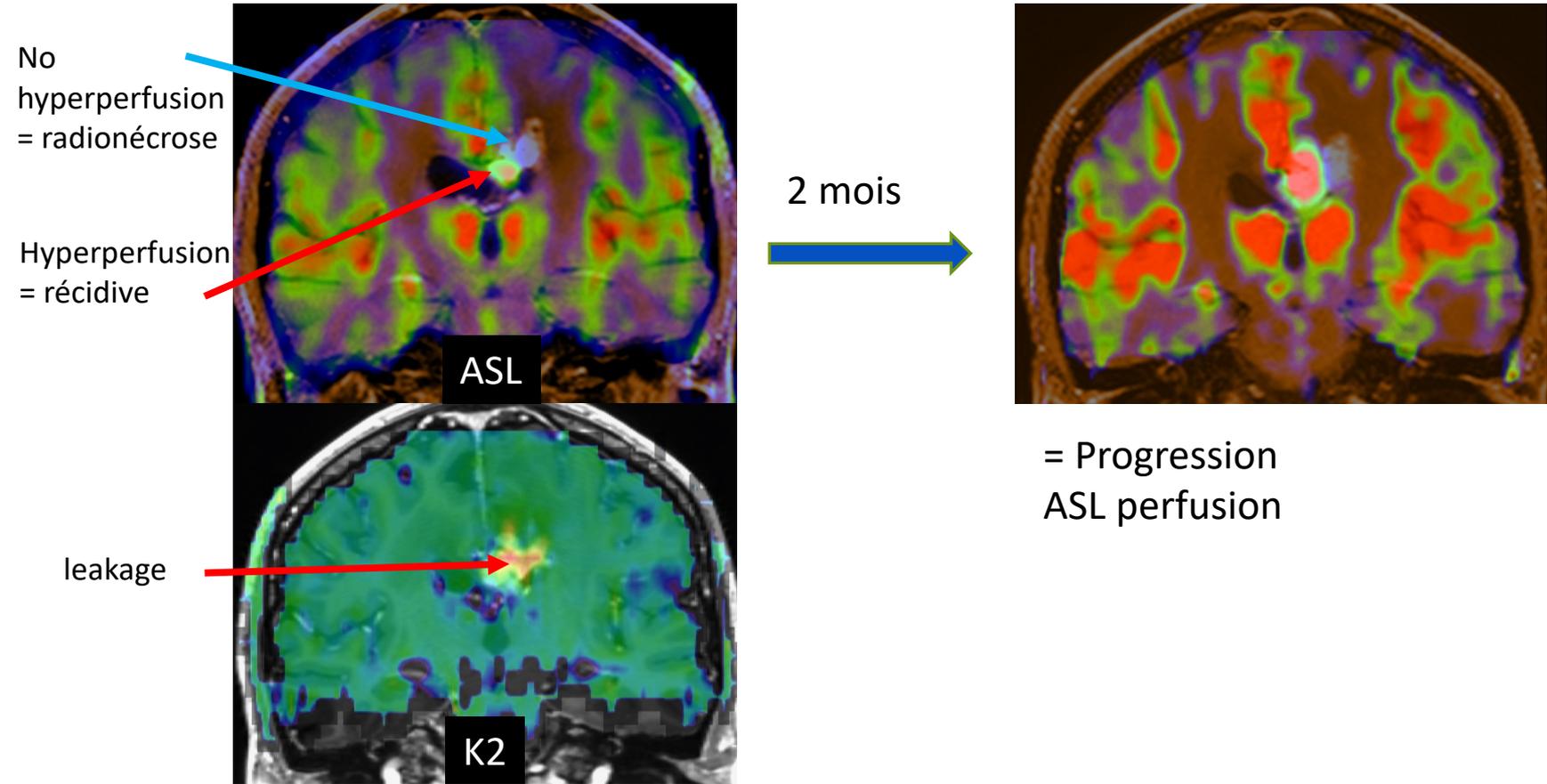


ADC ↑↑  
= nécrose  
≠ masse tumorale



Micro-hémorragie  
= fréquent dans radionécrose

# Aspects intriqués



- Ne pas oublier les **séquences conventionnelles et diffusion pour le diagnostic positif de lésion tumorale ou pseudotumorale**
- **Multimodalité** : caractérisation lésions, évolutivité  
Attention à la zone d'analyse
- Quelle que soit la lésion : retentissement (hydrocéphalie, engagement), vx
  - Pseudotumeurs : penser T2
- Suivi, connaître les traitements et leur action. Aspects intriqués