

中華民國醫事放射學會
Taiwan Society of Radiological Technologists (TWSRT)

MRI Lecture 1

Perfusion

台北醫學大學一部立雙和醫院
影像醫學部 醫學物理師 李宜恬
2016.11.19



Brain Perfusion Imaging

- The information on the capillary microcirculation of tissue
- Quantitative measurement
 - Blood volume
 - Blood flow
 - Temporal data (transit time and time to peak)
- Two major techniques
 - Dynamic-susceptibility-contrast (DSC) MRI
 - Arterial spin labeling (ASL) MRI

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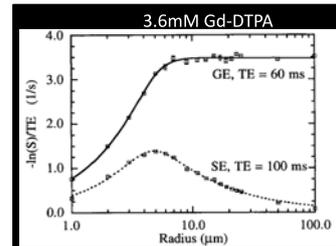
Techniques of Brain Perfusion

- Dynamic susceptibility-contrast MRI (DSC)**
 - CBF/ CBV/ MTT/K₂
- Dynamic contrast-enhanced MRI (DCE)**
 - CBF/ PS/ v_p / v_e
- Arterial spin labeling (ASL)**
 - CBF

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DSC MRI

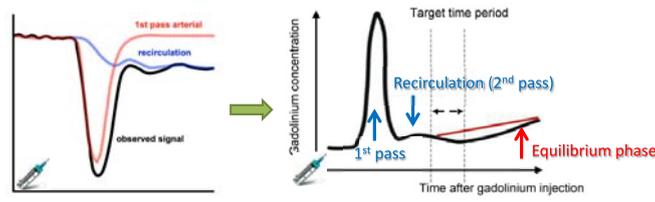
- T₂/T₂* weighted imaging**
 - T₂*W DSC-MRI is sensitive to large vessels.
- Bolus tracking**
 - Paramagnetic tracer
 - e.g. Gadolinium chelate
- Fast imaging**
 - GE-/SE-EPI
 - FLASH (SPGR)
 - Fast spin echo, etc



Boxerman et al., MRM 1995

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DSC MRI

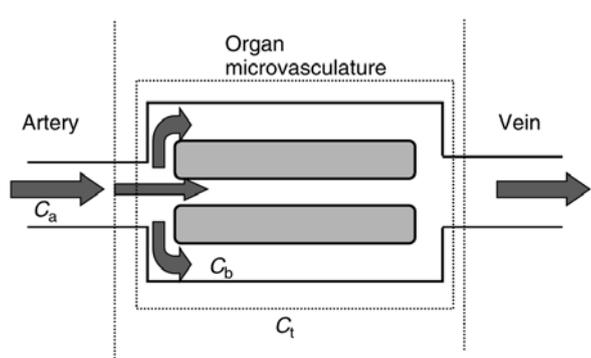


$$c_i(t) \propto \Delta R2^* = -\frac{1}{TE} \cdot \ln\left(\frac{S(t)}{S_0}\right)$$

Bang et al., Ann Neurol 2007;62:170-176

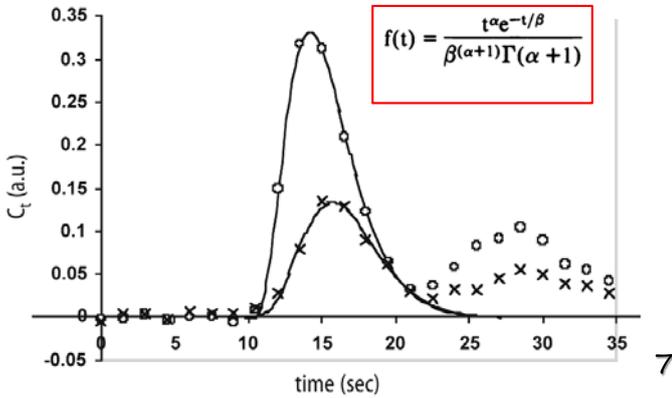
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Tracer Kinetics



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First-pass Fitting

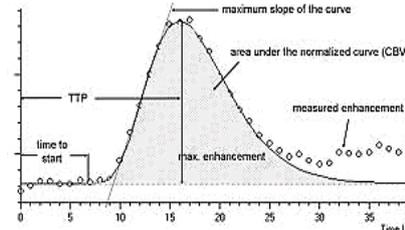
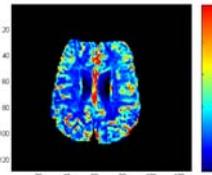


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CBV

$$CBV = \frac{K_H}{\rho} \cdot \frac{\int_0^\infty c_i(t) dt}{\int_0^\infty c_a(t) dt}$$

- Cerebral Blood Volume *VOF: Venous output function*
- Describes the blood volume of the cerebral capillaries and venules per cerebral tissue volume.
- Unit: % or ml/100g



Tomandl et al. © RSNA, 2003 May-June
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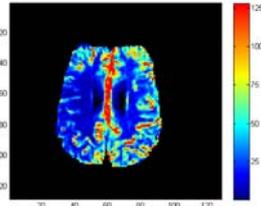
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CBF

$$C_t = AIF(t) \otimes [CBF \cdot R(t)]$$

- Cerebral Blood Flow
- Represents instantaneous capillary flow in tissue.
- Unit: ml/ 100g/ min



Hjort N et al. Stroke. 2005;36:388-397

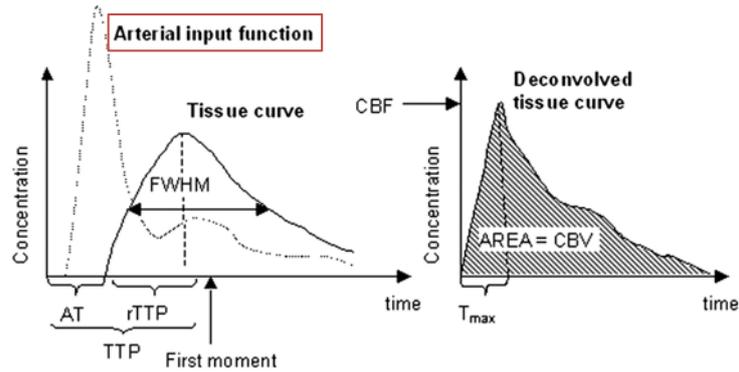
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CBF

$$C_t = AIF(t) \otimes [CBF \cdot R(t)]$$



Hjort N et al. Stroke. 2005;36:388-397

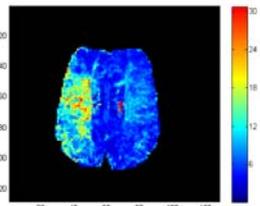
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$$MTT = CBV / CBF$$

MTT

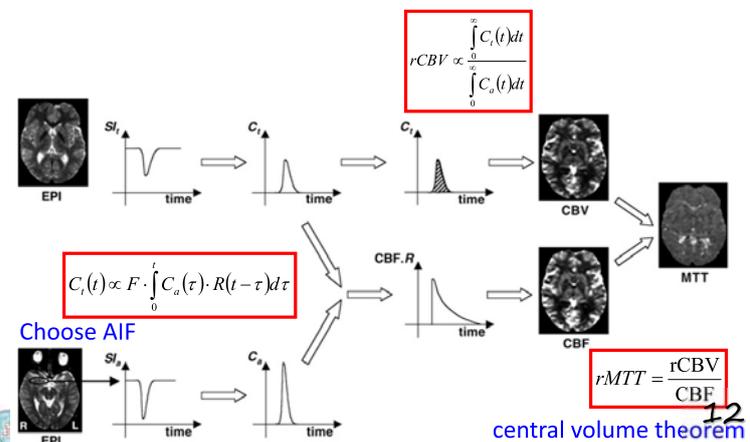
- Mean Transit Time
- Measures the length of time a certain volume of blood spends in the cerebral capillary circulation.
- Unit: min or s
- $MTT = CBV / CBF$



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Procedure of Analysis



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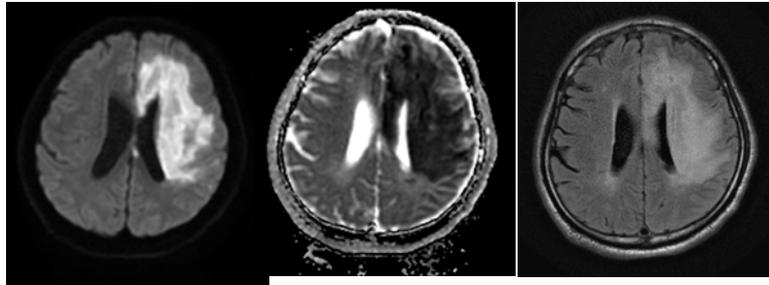


SHH Case: 吳○耀 092XX024

DWI

ADC

T2 Flair



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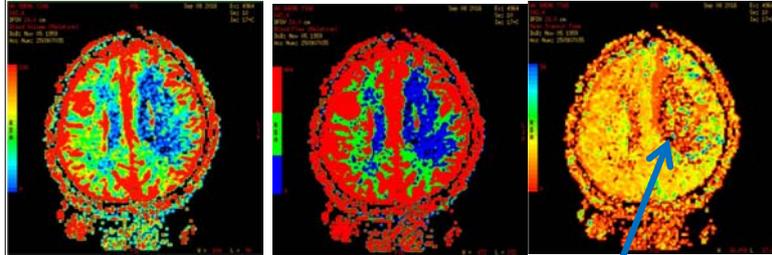
SHH Case: 吳○耀 092XX024

* GE perfusion post-processing tool:

CBV

CBF

MTT



MTT Shorten???



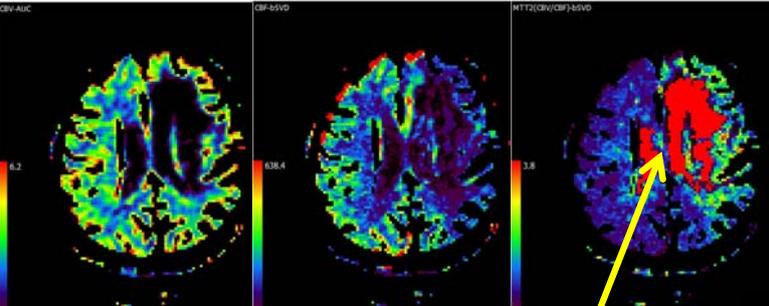
SHH Case: 吳○耀 092XX024

* PMA toolbox:

CBV

CBF

MTT



MTT Prolong!!!

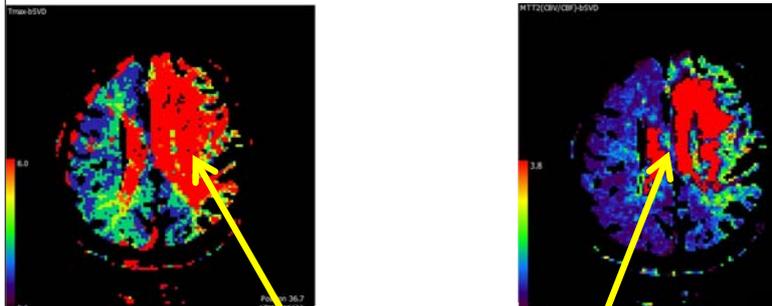


SHH Case: 吳○耀 092XX024

* PMA toolbox:

Tmax

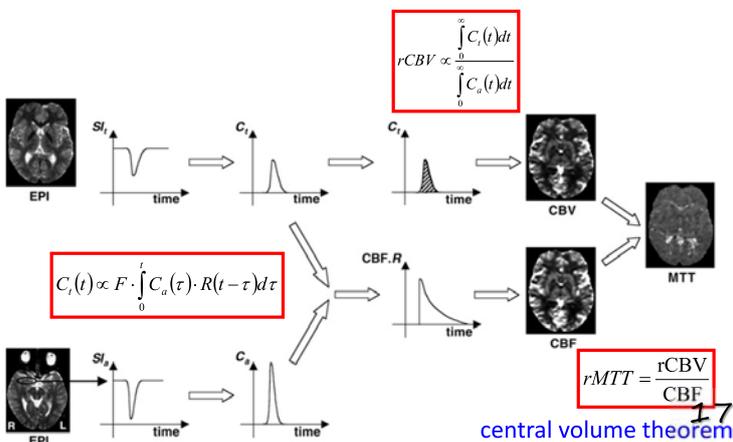
MTT



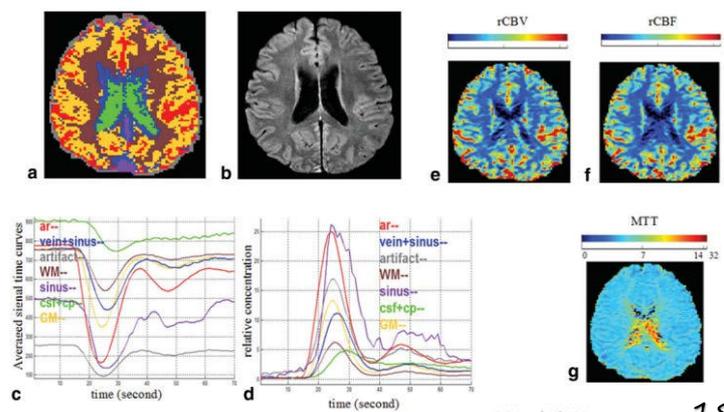
Tmax Delay!!!

MTT Prolong!!!

Procedure of Analysis

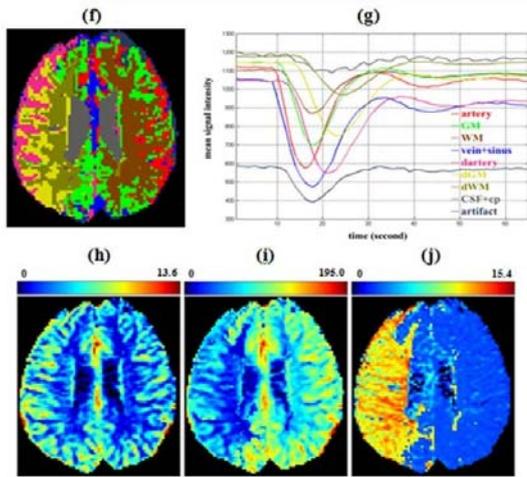


Tissue Classification



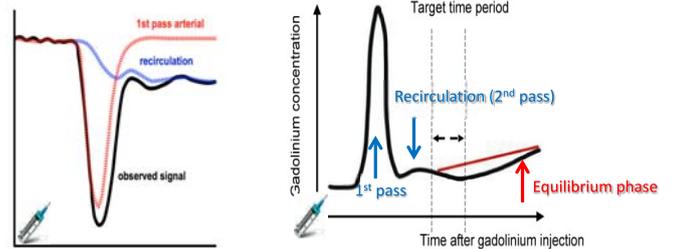
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Tissue Classification



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DSC-MRI for measuring Vascular permeability (K_2)

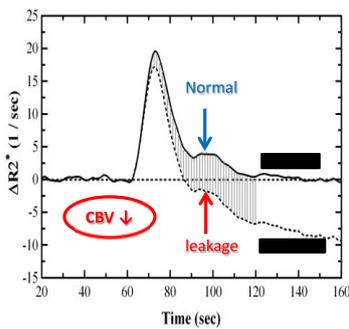


Bang et al., Ann Neurol 2007;62:170-176

- Equilibrium phase: wash-in = wash-out (Dynamic equilibrium)

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Detect contrast agent leakage by DSC-MRI



Boxerman et al., Am J Neuroradiol 2006; 27:859-67

- BBB damage will approach baseline faster and may fall below it. (T1 effect of contrast accumulation in parenchyma)
- The effect of BBB disruption can be modeled by comparing with normal tissue.
- The measurement of permeability, often referred to as K_2 , can be derived.

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Methods for Measuring Leakage by DSC-MRI

- Preloading Dose (Opts) + Weiskoff model
- Dual echo (spin and gradient echo, SAGE) + Vonken model

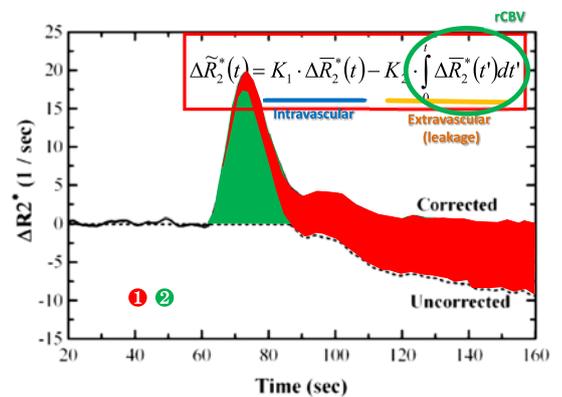
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Preloading Dose

- Procedure
 - Dose: 0.05 -0.1 mmol/kg
 - Infusion
 - Wait for about 5 - 6 minutes.
- T1 effect decreases while dose increases.
 - Sufficient dose may correct rCBV perfectly.
 - Too high dose may collide the assumptions of Weiskoff model.

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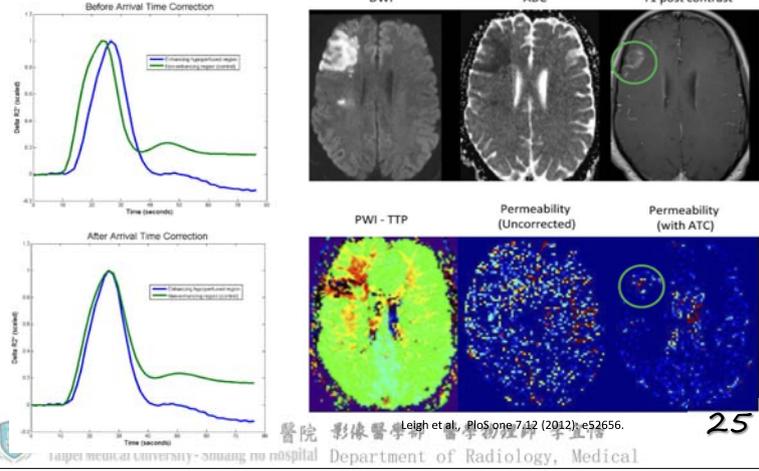
Weiskoff Model



Boxerman et al., AJNR, 2006

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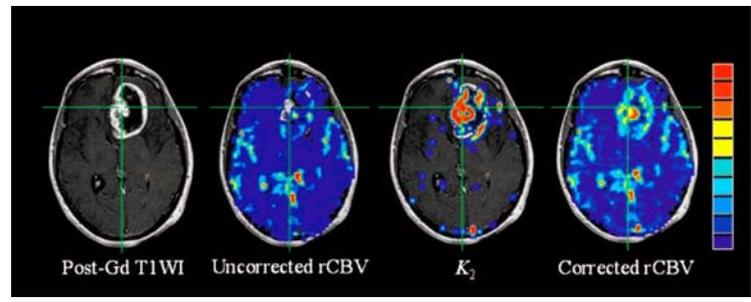
Arrival time correction (ATC)



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Weisskoff Model Correction

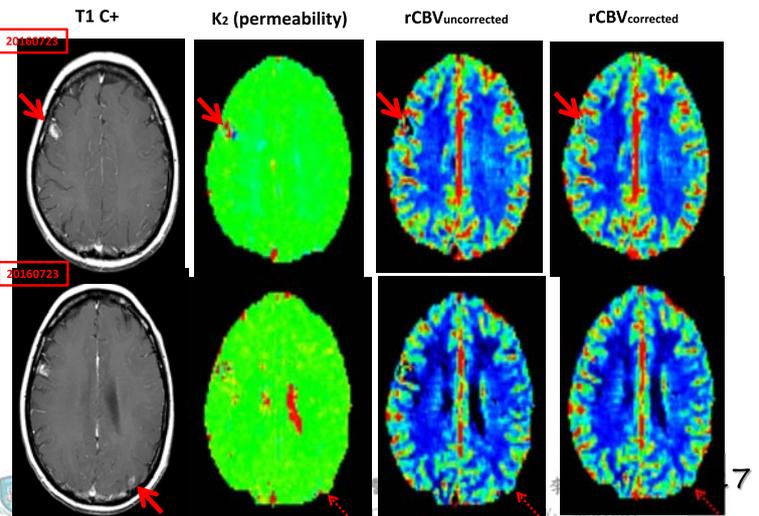


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Boxerman et al., AJNR, 2006

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Lung meta with chemotherapy

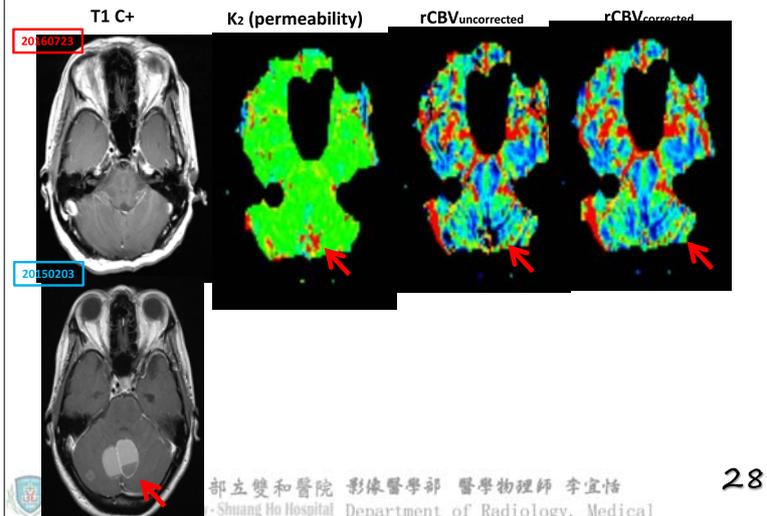


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Lung meta with chemotherapy

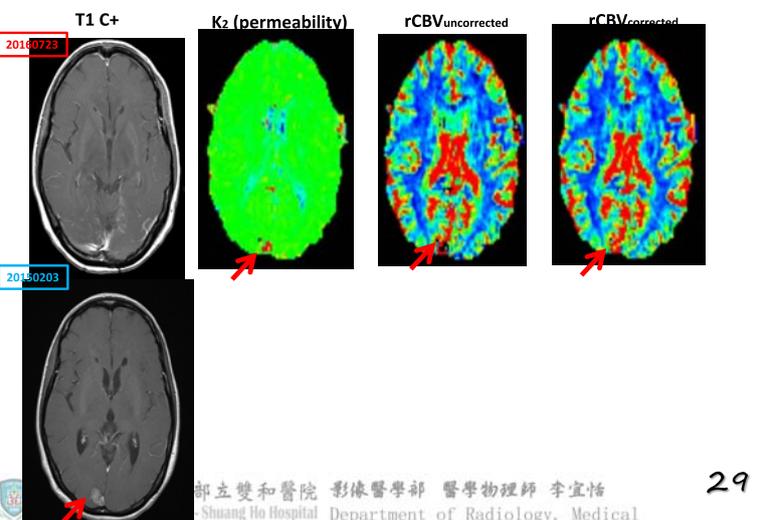


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Lung meta with chemotherapy



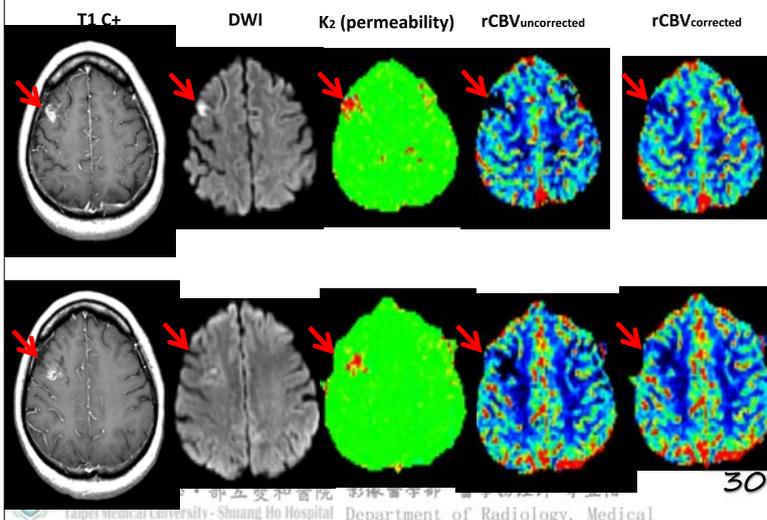
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ICA stenosis & normal perfusion with infarct



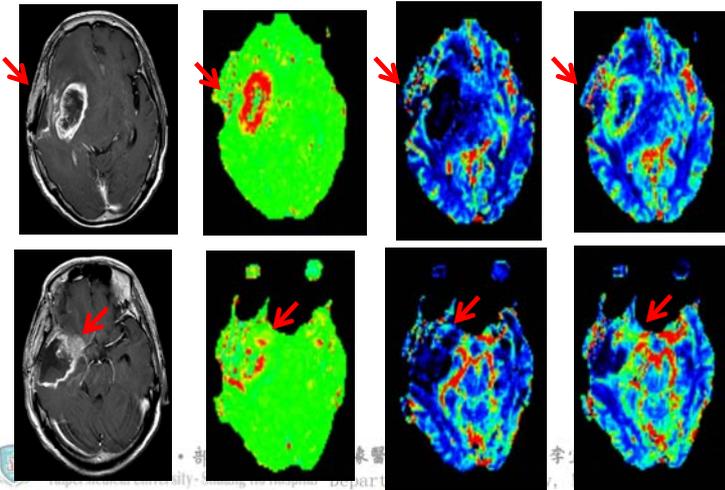
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Radiation necrosis or recurrent tumor?

T1 C+ K₂ (permeability) rCBV_{uncorrected} rCBV_{corrected}



1

ASL MRI

- Arterial spin labeling uses **arterial blood water** as an endogenous contrast agent.
- Blood is "**tagged**" or **magnetically inverted** which changes its magnetic properties and its effect on MR signal.
- Create paramagnetic tracer to suppress MR signal wherever arterial blood is delivered.
- Can be used to **quantify CBF** (cerebral blood flow) in arterioles and capillaries.

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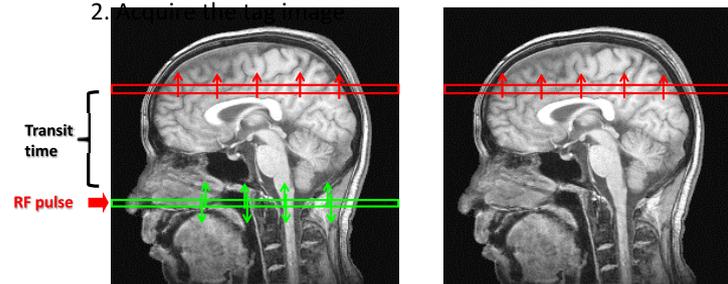
ASL CBF MAP

- **Pulsed Arterial Spin Labeling (PASL)**
 - A volume of blood is labeled upstream of the region of interest by a short RF pulse
- **Continuous Arterial Spin Labeling (CASL)**
 - Increase the delivered RF energy
 - Two sets of transmitter and receiver coils

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(Pseudo) Continuous ASL

1. Tag inflowing arterial blood by magnetic inversion
2. Acquire the image



Tagging

Control

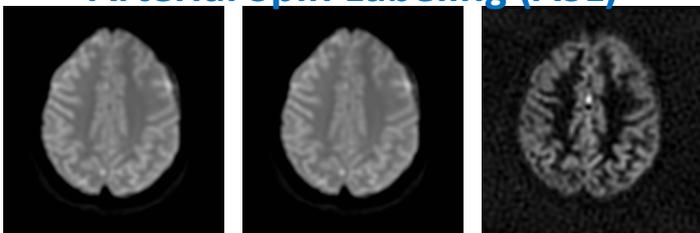
Imaging slice

Tagging plane

Courtesy from CGMH Prof. Liu Ho-Ling

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Arterial Spin Labeling (ASL)



Control — Labeling = ΔM

- ΔM is usually about 1% of the control signal, so many averages are usually required.
- (Tagging duration 2s + PLD 2.5s + image acquisition 0.6s) X 2 X 32 = 6 mins

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Potential Confounds in CBF Quantification with ASL Imaging

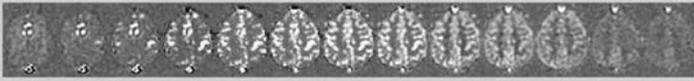
- Intra-arterial signal effects
- Transit/ Trailing time effects
- Relaxation effects

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Intra arterial signal effects

- Perfusion signal could be contaminated by the signal coming from the labeled intra-arterial blood, which has not reached the capillary/ tissue site for exchange.

Without Gradient Crushers



With Gradient Crushers

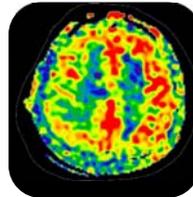


TI = 0.26 0.29 0.35 0.55 0.75 0.95 1.15 1.35 1.55 1.85 2.15 2.45 2.75 s

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Intra arterial signal effects

- Perfusion signal could be contaminated by the signal coming from the labeled intra-arterial blood, which has not reached the capillary/ tissue site for exchange.



Intra-arterial blood (Slow flow)?
L't hyperperfusion?

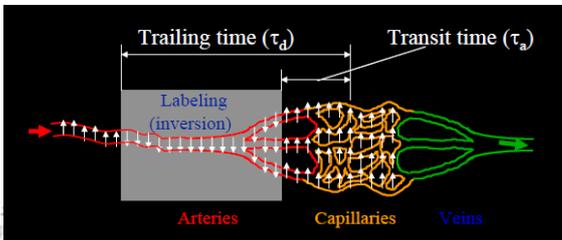
R't hypoperfusion?
Tagging artifact (Stent cases)

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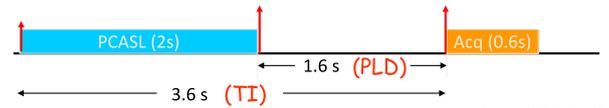
Transit/ Trailing Time Effects

- Transit/trailing time – the time for the leading/ trailing edge of the labeled blood to reach the capillary/tissue exchange site.
- Caused by
 - Gap between the inversion slab and image slices
 - Blood Traveling time from arterials to capillaries.

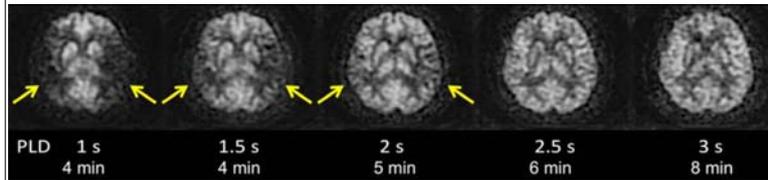


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Post Labeling Delay



Courtesy CGMH Prof. Liu Ho-Ling



- Improved CBF quantitation on long PLD using PCASL at 3 T in a patient with bil Moyamoya disease

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Taipei Medical University · Shuang Ho Hospital Department of Radiology, Medical Neuroimaging Clin.N Am. 2011 May; 21(2): 285-301

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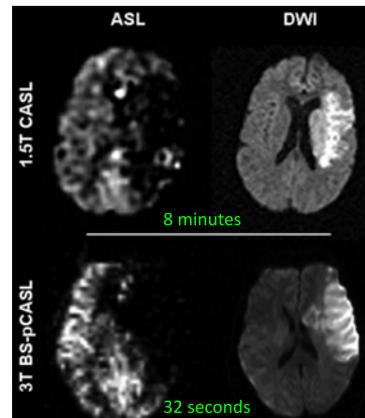
Solution to Transit Delay

- High main magnetic field
 - Extended ASL (eASL)
 - Multiple PLD
 - Velocity selective ASL (VASL)
- } Longer PLD

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1.5T VS 3T

- Higher magnetic field
 - Inherent high SNR: Less repeated acquisition => shorter imaging time
 - Longer blood T₁: Allow longer post labeling delay

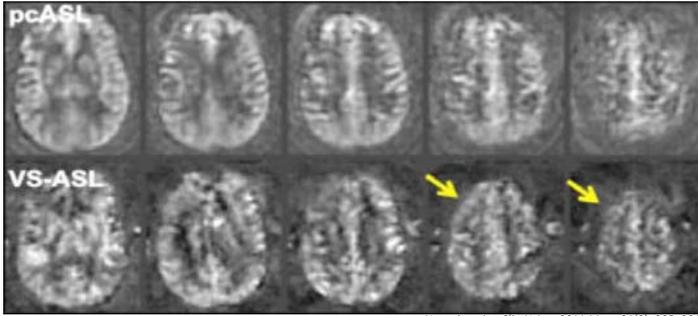


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Taipei Medical University · Shuang Ho Hospital Department of Radiology, Medical Neurotherapeutics. 2007 July; 4(3): 346-359.

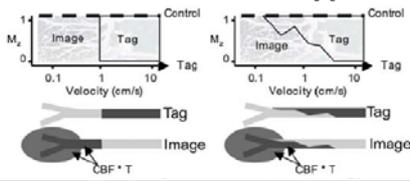
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VASL

A patient with Moyamoya disease and bilateral MCA occlusion

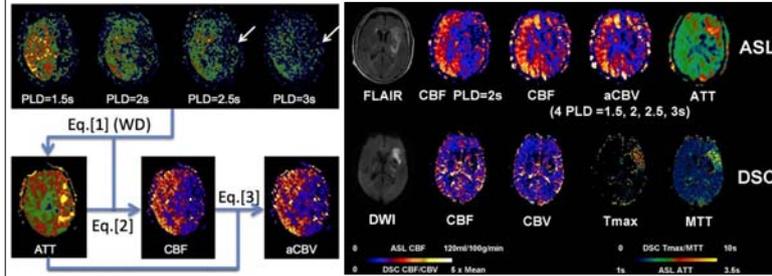


Neuroimaging Clin N Am. 2011 May; 21(2): 285-301



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Multiple PLD



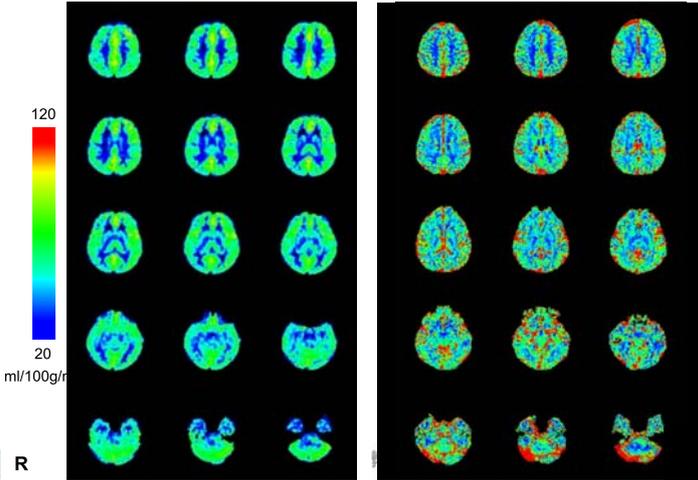
- PCASL at 4 PLDs (PLD = 1.5, 2, 2.5, 3 s) were acquired within 4.5 min in 24 patients with acute MCA infarct
- Highly significant correlations between pCASL and DSC CBF ($r=0.70$, $p=0.0001$)
- ASL ATT correlate with DSC Tmax ($r = 0.66$, $p = 0.0005$) and MTT ($r = 0.59$, $p = 0.0023$) in leptomeningeal MCA territories

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3D ASL

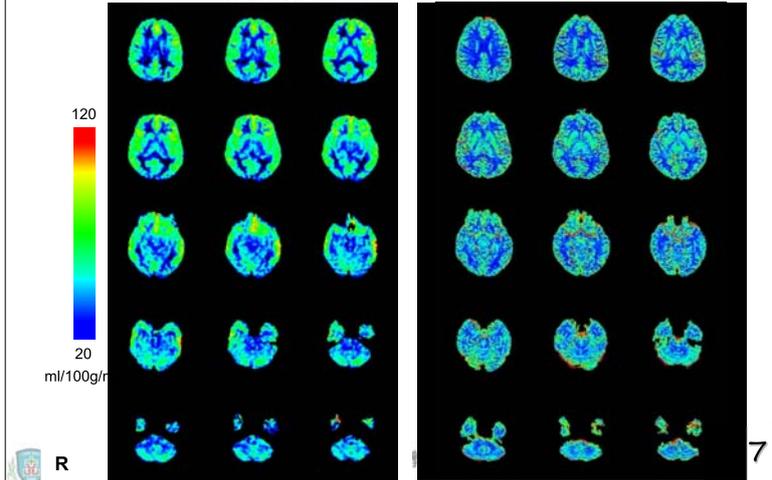
GE-DSC



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3D ASL

SE-DSC



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Conclusions

- ASL allows perfusion imaging **without contrast injection**
- ASL allows repeated measurements
- Limitations of ASL include:
 - SNR
 - Coverage
 - Delay time variations

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