



# Basics of DWI and PWI

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**Current Concepts**  
 Monterey  
 October 2005

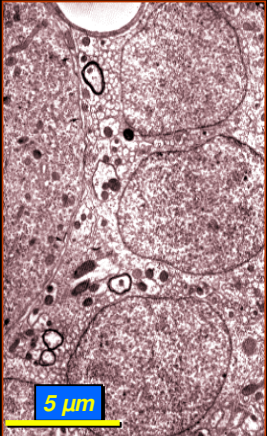

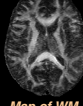
## Proton Motions and DWI

DWI measures proton motion:  
 5-20  $\mu\text{m}/\text{image}$ .

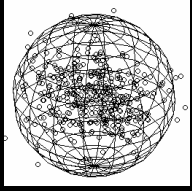
Samples  $\mu\text{environment}$ .  
 Measured as ADC.

Diffusion sensitive to:

- Water content-  
 ADC increases with PD.
- Water compartments-  
 Intracellular water slower.
- Water hindrances-  
 Ordered in WM.

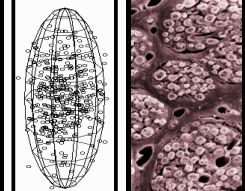
### Diffusion in an isotropic medium (GM)



similar molecular displacements in all directions

*The Apparent Diffusion is quantitated as the "ADC"  
 - Measure of translation*

### Diffusion in an ordered environment (WM)

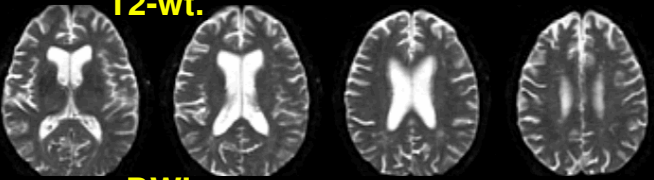


greater molecular displacement along cylinders than across

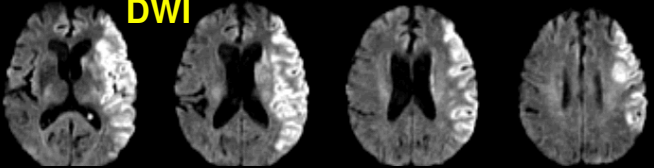
*Deviation from random paths-  
 "Fractional Anisotropy"  
 -Measure of WM integrity*

## MRI and Acute Stroke 4 hours

**T2-wt.**



**DWI**



## Practices of DWI: Artifacts

**Artifacts in EPI DWI can be serious**

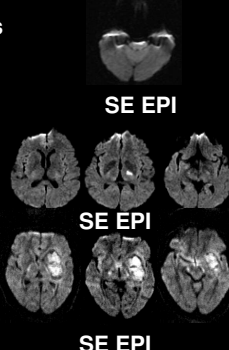
Artifacts arise from susceptibility issues

- Signal drop-outs/pile-up
- Aspect ratio distortions
- Sensitivity to shims/metal

Eddy currents effects

- Aspect ratio distortions/smearing

Use auto-shimming prior to exam  
 Avoid overly high resolutions (<192)  
 New correction algorithms  
 New Sequences for DWI

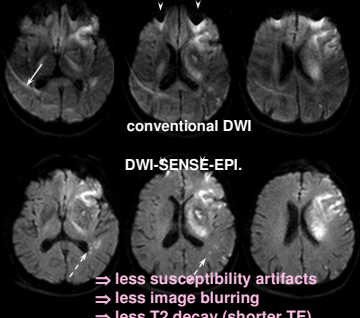


## Can DWI Be Improved?

*"Perhaps the biggest immediate improvement affecting all of MR today is the development of parallel imaging..."*

Parallel Imaging (SENSE)

- Improve Temporal Resolution
- Increased Volume Coverage/ Time
- Reduction of Motion Artifacts
- Reduce T2/T2\*-Blurring
- Diminish Off-resonance Artifacts
- Decrease SAR Problems
- Reduction of Total Acquisition Time



conventional DWI

DWI-SENSE-EPI

- ⇒ less susceptibility artifacts
- ⇒ less image blurring
- ⇒ less T2 decay (shorter TE)

## The MR PROPELLER Sequence

"Periodically rotated overlapping parallel lines with enhanced reconstruction"

**Multi-shot, multi-echo FSE (blades)**  
**Each blade passes thru k-space center**  
**Robust motion correction**  
**No distortions, no susceptibility**  
**Slower, marketed by GEMS**

PROPELLER blades, Arfanakis, et al. ISMRM, Kyoto 2004  
Motion Correction With PROPELLER MRI  
 J. Pipe, MRM 42:963-969 (1999)

## DWI "Mimics" Beyond Clinical Stroke

**Reduced ADC "cytotoxic" vs. reduced mobility.**  
 Screen for "non-stroke" events (headaches, TIA).  
 Secondary ischemia (acute trauma, venous occlusion)  
 ADC reduced in "cytotoxic" edema: CJD, pediatric diseases.  
 ADC reduced in abscess \*\* vs. highly cellular tumor cores.

**Elevated ADC "vasogenic" (usually seen also as elevated T2).**  
 MS plaques\*\* (older), edema, inflammation, necrosis.

**T2 FSE**  
**DWI**

*4 y.o. male, 3 days s/p MVA, decreasing mental status, UE weakness*  
 G. Sorensen, MGH

## Diffusion vs. Perfusion

**(Self)-Diffusion**  
 ... has zero mean, but r.m.s. displacement characterizes diffusion coefficient !

Start → End

— large diffusion coefficient  
 — small diffusion coefficient

**Perfusion**  
 Capillary microcirculation in organs. Rate of nutrient supply.

## Why does ADC of water decrease in acute stroke ?

perfusion deficit → energy deficit → failure of Na<sup>+</sup>/K<sup>+</sup> pumps → intracellular accumulation of Na<sup>+</sup> → influx of water → glial and neuronal swelling → reduction in extracellular space → Mobility hindered = ADC drop → DWI hyperintensity → eventual lysis/edema

## Single-shot EPI is Ideal for Diffusion MR

$\delta = 25 \text{ msec}$     $\Delta = 40 \text{ msec}$

**EPI Readout**

**Diffusion-Weighted images**   **ADC map**

T2-wt ( $b=0$ )   DWI ( $b=800$ )   ADC map

$$e^{-\gamma \delta G [D-\delta/3] \text{ADC}}$$

"b"

## The Principles of DWI: The Protocol

**Acquire all gradient axes.**

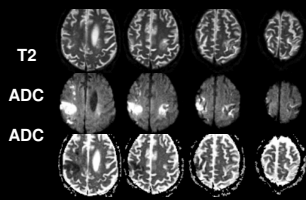
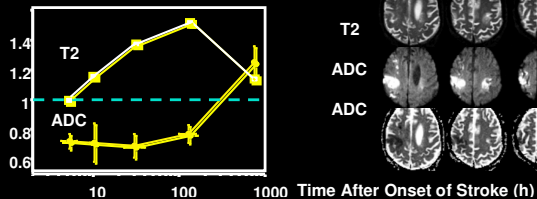
- \* Don't be fooled by WM anisotropy effects.
- \* X, Y, and Z ("trace")
- \* Off-axis directions also hot ("tensor")
- \* Note b<sub>0</sub>, DWI, AND ADC!

**DWI**   **ADC**

reduced ADC due to anisotropy   Trace ADC

## ADC vs T2-wt Shine-Thru

	T2	DWI	ADC
• hyperacute (< 6hrs)	iso	high	low
• acute (6hrs to 4 d)	high	high	low
• subacute (4 to 10 d)	high	iso/h	iso
• chronic (>10 d)	high	iso/l	high



## The Principles of DWI: *The Protocol 2004* The impact of new MR technology

Better SE-EPI

Image unwarping (postprocessing)  
Dual-echo (TRSE)

Parallel Imaging for DWI  
SENSE-DWI

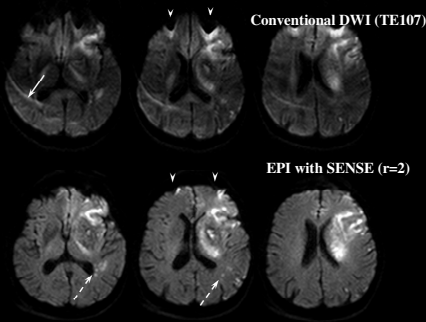
FSE-based DWI

PROPELLER (and Turbo-PROP)

Spiral-based DWI

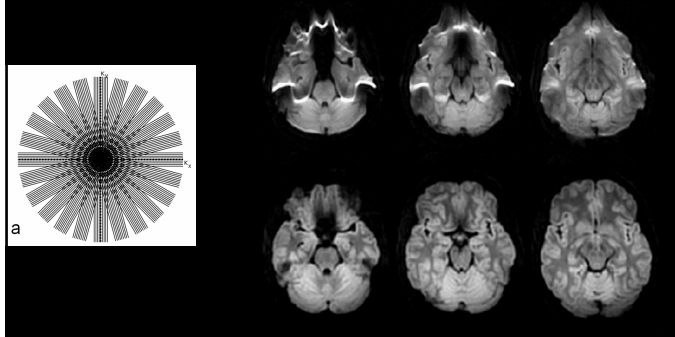
SNAILS

## The Principles of DWI: *The Protocol 2004* SENSE-EPI for DWI

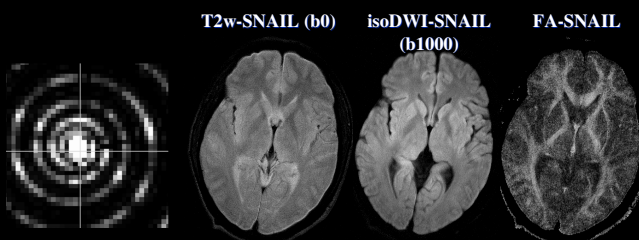


**Advantages:**  
Image distortions  
Short TE's  
Less blurring  
Increased conspicuity  
*Hi-res!*

## The Principles of DWI: *The Protocol 2004* PROPELLER (and Turbo-PROP)



## The Principles of DWI: *The Protocol 2004*



512x512, 8 averages, 64-leaf, b800, 23 minutes

## Integrated MRI for Acute Stroke

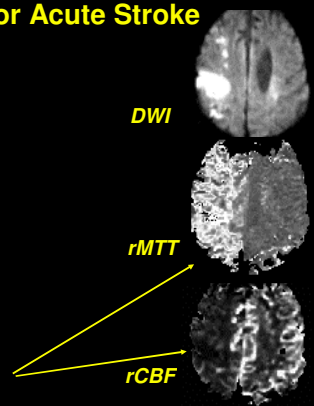
### What is PWI?

Collect serial EPI images  
Inject contrast agent:

Perfused brain - dark  
Ischemic brain - bright

Hemodynamics tracked  
Time-resolved transit  
Correlate with DWI, T2

Perfusion deficit present:  
(low rCBF, long rMTT)

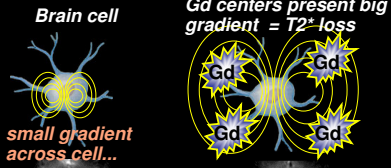




## Why Do We Call PWI "Perfusion"?

Gd shortens T1...  
However...  
Gd also shortens T2\*.

On a GRE-EPI image,  
a bolus of  
0.1mmol/kg Gd  
can decrease T2\*  
by 50%-80%



GRE  
EPI  
TE60

At bolus  
peak



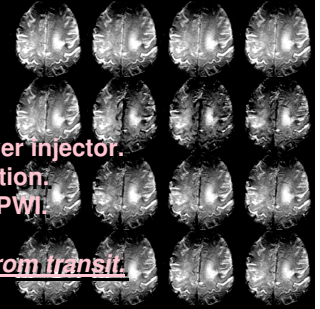
## PWI Protocol

GRE-EPI images acquired  
during bolus injection.

12 slices every 2 sec.  
40 phases over 80 sec.  
0.1 mmol/kg Gad - Power injector.  
1.5mm per pixel resolution.  
as for T2, FLAIR, DWI, PWI.

Maps of rCBV, rMTT, rCBF from transit.

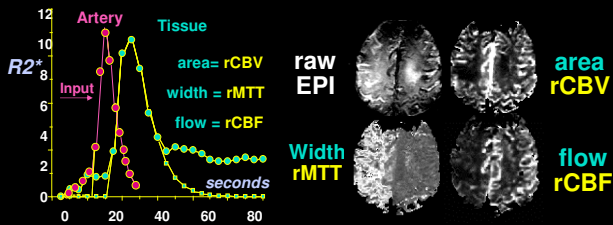
Arterial input needed.



Bolus  
transit

## How Does PWI Map Hemodynamics?

2. From the SI behavior map, model the transit dynamics  
by a "gamma variate" analysis. This yields parametric  
maps of the vascular volume, transit, and flow...



## The Diffusion - Perfusion Mismatch

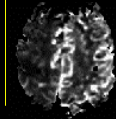
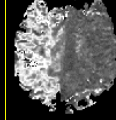
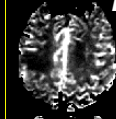
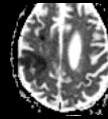
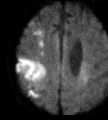
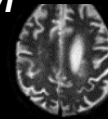
DWI

PWI

T2-wt  
Early stroke  
not seen

Diffusion-wt  
Clear depiction of  
lesion

Apparent Diffusion  
Acute stroke has low  
ADC

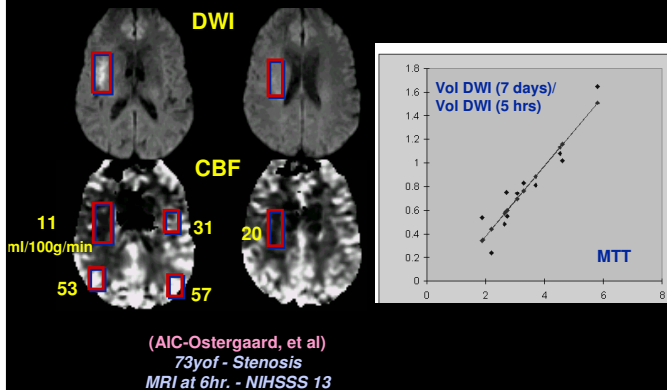


Blood Volume  
Lesion has  
reduced CBV

Mean Transit  
large perfusion  
deficit

Blood Flow  
Reduced flow  
around lesion

## Is There a Need for CBF Quantitation?

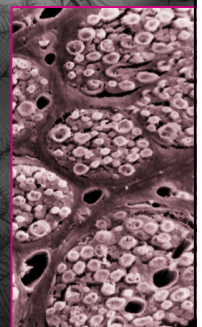


## Proton Motions are Ordered by WM

DWI White Matter "Anisotropy" is Altered with  
Growth, Aging, Disease...

Motion of water hindered in a restricted  
environment (inside neuronal fibers).  
Complex motion described by a "tensor".  
Number, integrity, myelination of fibers  
alters the measured tensor.  
Cognitive performance appears to be  
correlated with anisotropy.

How will DTI impact neuroimaging?



## DTI is Not all Myelin!

**DTI is related to...**

- Fiber diameter and neuronal density,
- Status of the myelination in neurogenesis,
- Degree of (de)myelination in aging/disease,
- Macrostructural features such as intravoxel fiber-tract coherence/crossings

## In DTI, We Measure ADC Along All Axes

ADC vs. Direction = ADC Tensor

$$\underline{ADC} = \begin{bmatrix} ADC_{xx} & ADC_{xy} & ADC_{xz} \\ ADC_{xy} & ADC_{yy} & ADC_{yz} \\ ADC_{xz} & ADC_{yz} & ADC_{zz} \end{bmatrix}$$

## Diffusion Tensor Imaging DTI: The EPI Sequence

$(G_x, G_y, G_z) = \{(1,1,0), (0,1,1), (1,0,1), (-1,1,0), (0,-1,1), (1,0,-1)\}$   
 SE-EPI TR6000, TE107, 128x128, b0-1000, 6 axes, 3.5min/NEX

Diffusion-Weighted images (b=0, b=800) in xy, yz, xz, -xy, -yz, -xz planes. Tensor calc leads to Trace ADC and FA maps.

## DW Images to ADC Maps to Eigenvalues

$$S(\underline{b}) = S_0 \exp\left(-\sum_{i=1}^3 \sum_{j=1}^3 b_{ij} ADC_{ij}\right)$$

$$\underline{ADC} = \begin{bmatrix} ADC_{xx} & ADC_{xy} & ADC_{xz} \\ ADC_{xy} & ADC_{yy} & ADC_{yz} \\ ADC_{xz} & ADC_{yz} & ADC_{zz} \end{bmatrix}$$

$$\underline{ADC} = \begin{bmatrix} \lambda_1 & 0 & 0 \\ 0 & \lambda_2 & 0 \\ 0 & 0 & \lambda_3 \end{bmatrix}$$

## Eigenvalues to FA Maps

Fractional Anisotropy = FA = Created from  $\lambda_1, \lambda_2, \lambda_3$

- 0 (sphere) < FA < 1 (cylinder)
- 0.05 (GM) < FA < 0.8 (splenium)

$$FA = \frac{3\sqrt{D:D}}{2\sqrt{D:D}}$$

$$D:D = (\lambda_1 - \langle D \rangle)^2 + (\lambda_2 - \langle D \rangle)^2 + (\lambda_3 - \langle D \rangle)^2$$

$$D:D = \lambda_1^2 + \lambda_2^2 + \lambda_3^2$$

Isotropic, 128x128, 25FOV, 2mm slice  
37 slices, 13 min, 6NEX

## DTI Protocol

**Parameters (CV-i Signa 1.5T)**

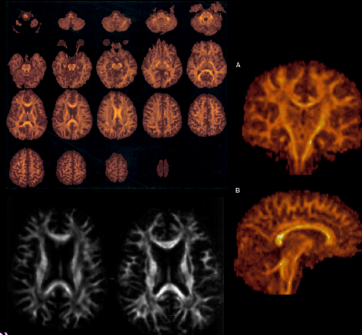
- $\delta = 20\text{ms}; \Delta \approx 34\text{ms}; G_d = 1.4\text{G/cm}$
- Gradients ( $G_x, G_y, G_z$ ) =  $\{(1,1,0), (0,1,1), (1,0,1), (-1,1,0), (0,-1,1), (1,0,-1)\}$
- TE/TR/TI = 106/6000/2100ms
- FOV = 24cm; slice 2/0
- matrix 128x128.

**For each slice**

- 2 images with b=0 and IR
- for CSF suppression (unwarping)
- 4-6 averages with b=800s/mm<sup>2</sup>

## Clinical DTI: Protocols and Parameters

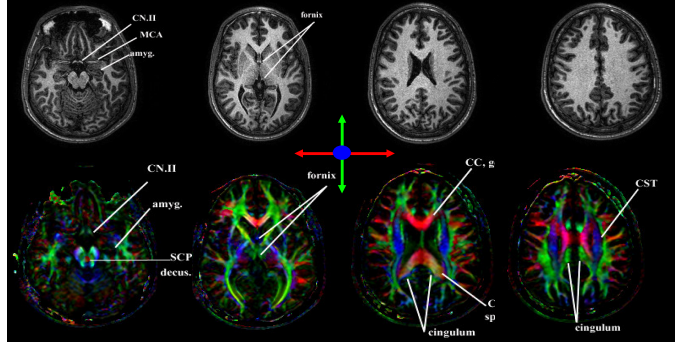
TR 10-20 RR (if gated), TR10000  
 TE min full (107)  
 FOV24, 96x96  
 Slice 2.5-5mm  
 $\delta$  (duration) 18ms  
 $\Delta$  (separation) 49ms  
 G (grad amp) 40 mT/m  
 b max 1300 sec/mm<sup>2</sup>  
 NEX (b0) 2-4  
 NEX (b1300) 64 (64x1)  
 Slices 39  
 Scan time 14-17minutes



$$"b" = e^{-\gamma \delta^2 G^2 [\Delta - \delta/3] ADC}$$

Jones, et al.  
 Neuroimage 17, 2002

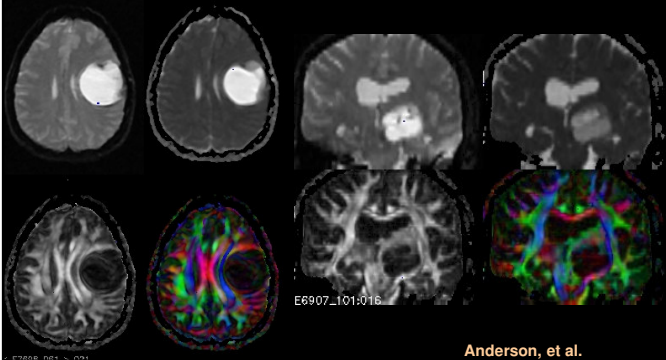
## FA "Atlas" of the Brain



S. Mori, JHU

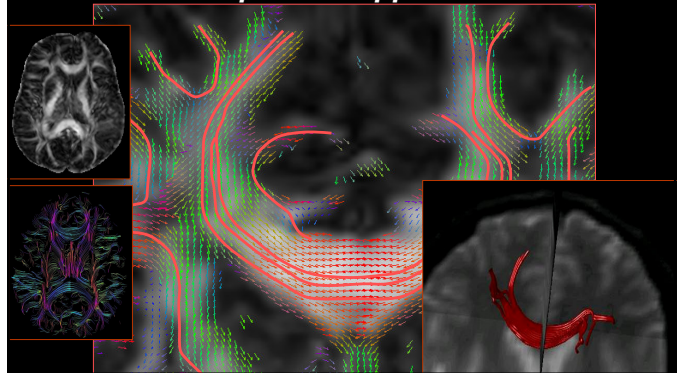
## FA Mapping Applications

Tumor Displacement, Infiltration, or Degradation of WM



Anderson, et al.

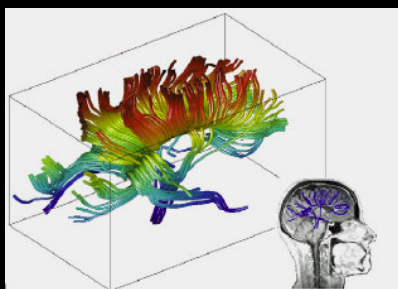
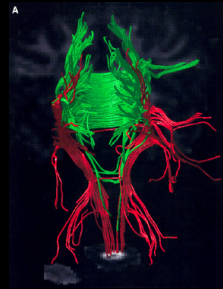
## FA Maps, Vectors and Fiber Tracking: Concepts and Applications



## Clinical DTI: Visualization

Streamtubes overlaid onto FA

Stochastic trajectories

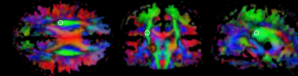


Jones, et al.  
 Neuroimage 17, 2002

Tench, et al.  
 MRM 48, 2002

## DTI and Reading

Torkel Klingberg, Maj Hedehus\*, Russ Poldrack,  
 Mike Moseley\*, John Gabrieli, Gayle Deutsch,  
 Bob Dougherty, WT Siok, Roland Bammer\*, Brian Wandell.



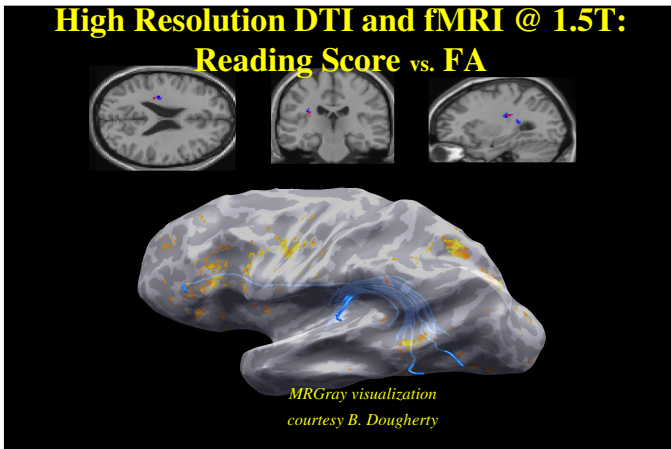
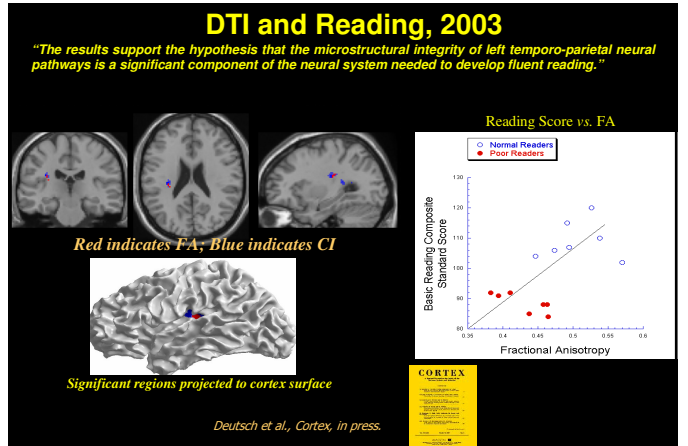
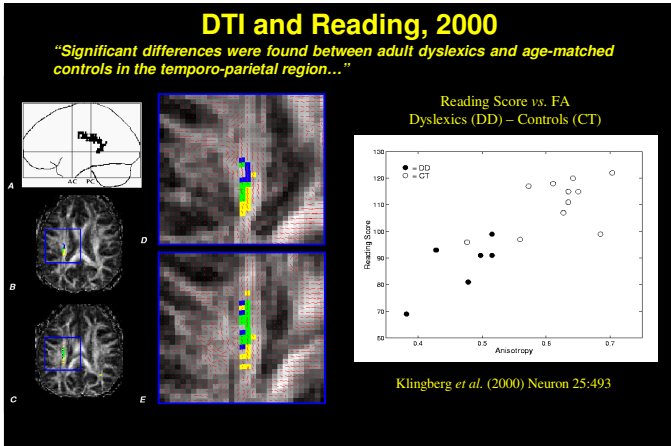
\*Center for Advanced MR Technology at Stanford  
<http://www-radiology.stanford.edu/research/RR.html>

Stanford University



School of Medicine





### Summary

DWI ideal for all neuro screening.

DTI is a subset of DWI - opens new doors.

FA sensitive to WM development /status.  
Does WM FA reflect status of regional GM metabolism - "Use it or lose it"?  
Would learning increase local FA?

Why does DTI correlate with performance?  
"Integrity" ala FA means what?  
Effective neural density?  
Structural defects or potholes?

