



# Diffusion MRI Analysis

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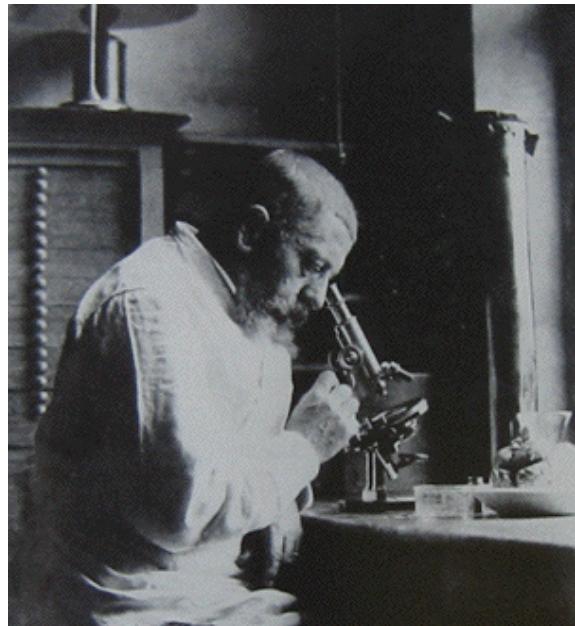
Harvard Medical School

# Brain Anatomy



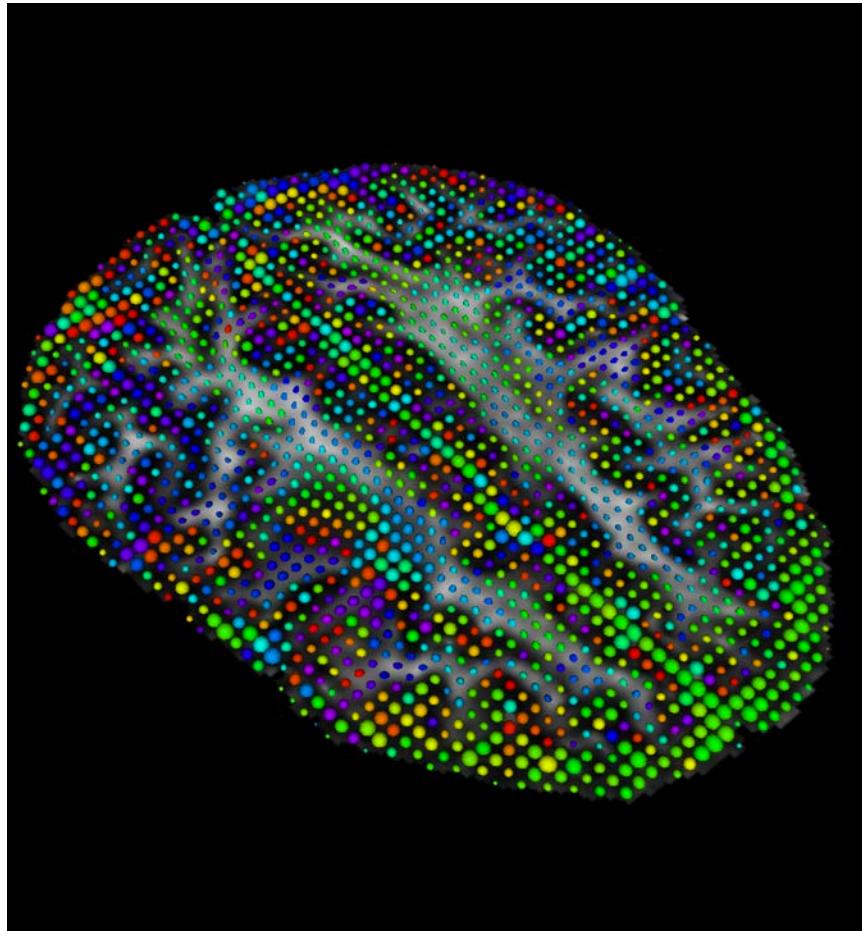
- White matter ~45% of the brain
- Myelinated nerve fibers (~ 10  $\mu\text{m}$  axon diameter)

# White Matter Exploration



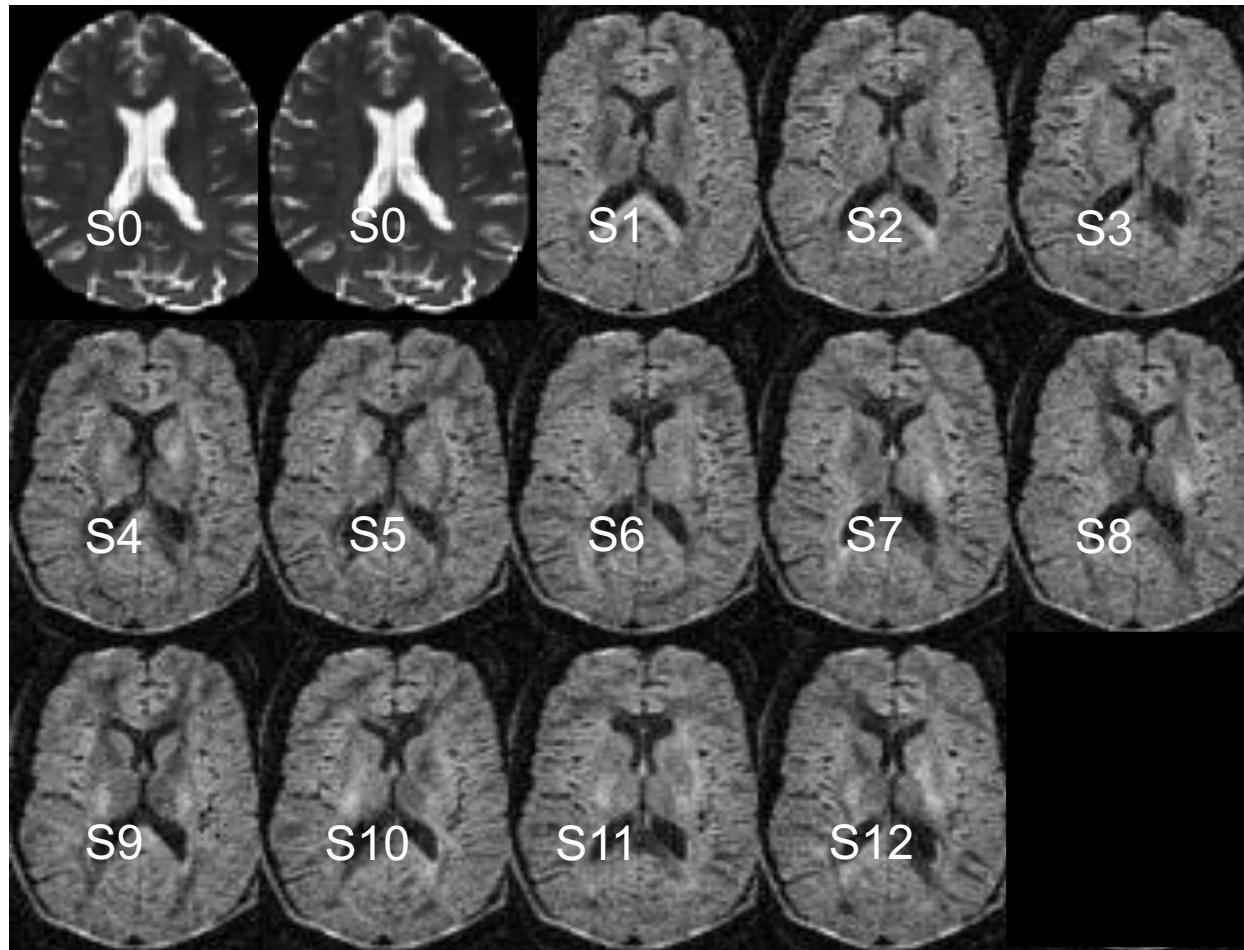
Jules Joseph Dejerine  
(*Anatomie des centres  
nerveux* (Paris, 1890-1901):  
Atlas of Neuroanatomy based  
on myelin stained preparation

# Diffusion Tensor Imaging (DTI)



- First non-invasive window on white matter anatomy
- Measurement of the motion of water molecules using MRI techniques.
- Three-dimensional reconstruction of the trajectory of white matter bundles

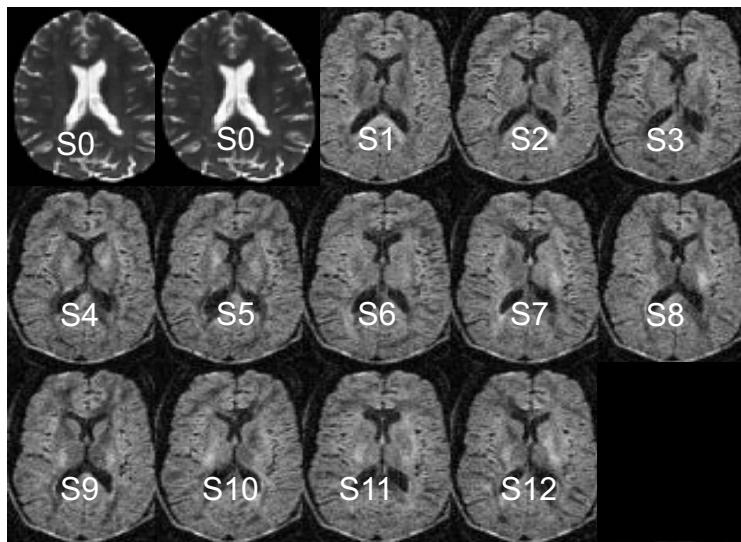
# Diffusion Weighted Imaging (DWI)



In this example, the DWI scan was acquired with 12 diffusion sensitizing gradient directions (S1-S12) and 2 non-diffusion sensitizing gradients (S0)

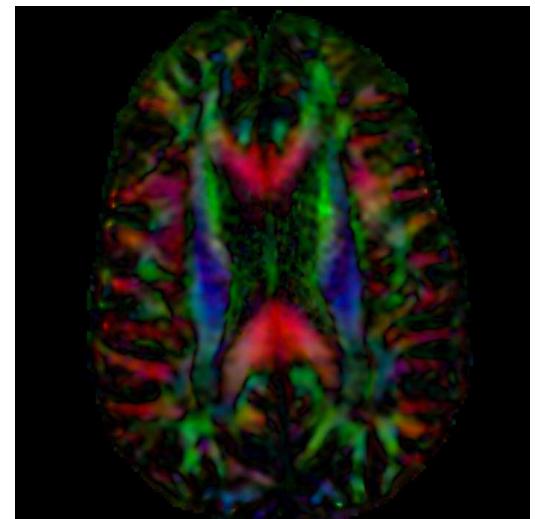
# From DWI to DTI

DWI



DWI dataset acquired with  
12 gradient and 2 baseline

DTI



DTI dataset

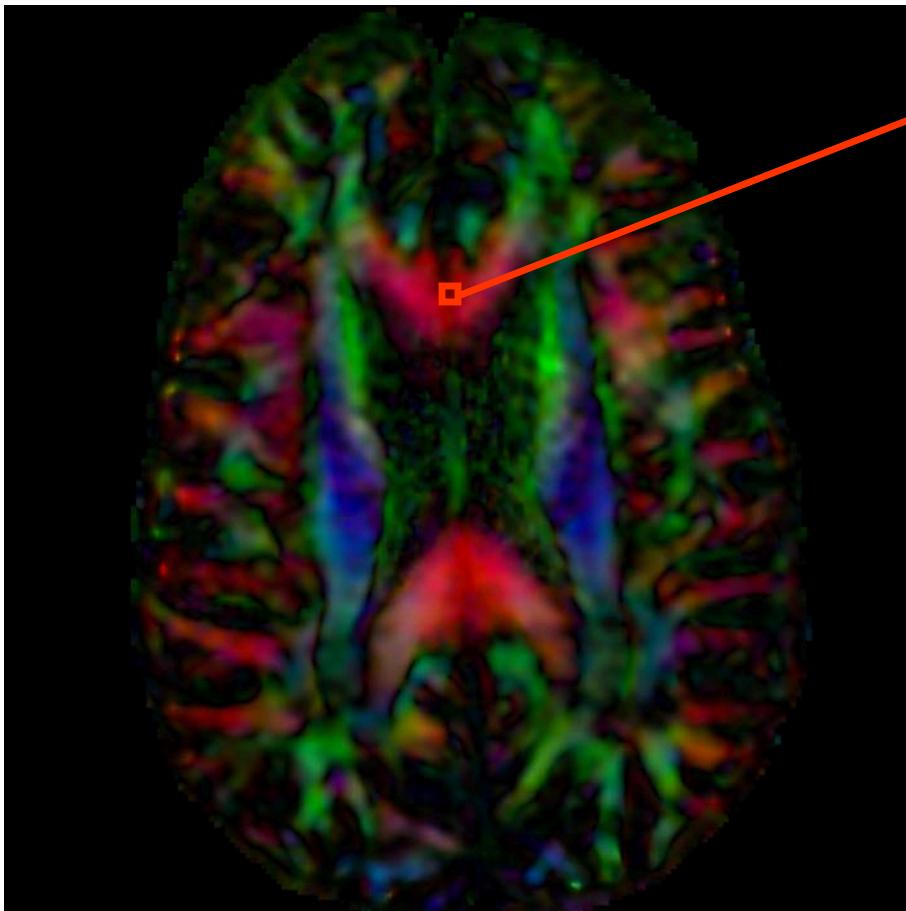


$$S_i = S_0 e^{-b \hat{g}^T \underline{D} \hat{g}_i}$$

Stejskal-Tanner (1965)

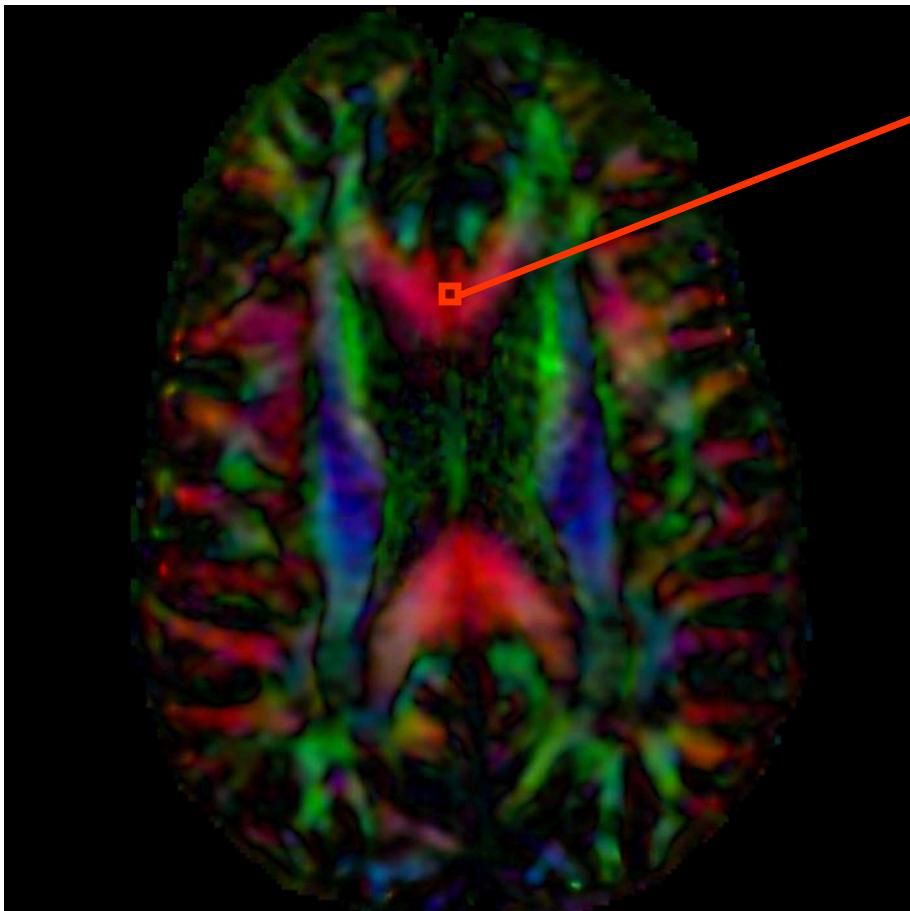
*Si: DWI volume acquired with  
ith gradient  
So: Baseline volume*

# Diffusion Tensor Imaging



$$S_i = S_0 e^{-b \hat{g}_i^T \underline{D} \hat{g}_i}$$

# Diffusion Tensor Imaging

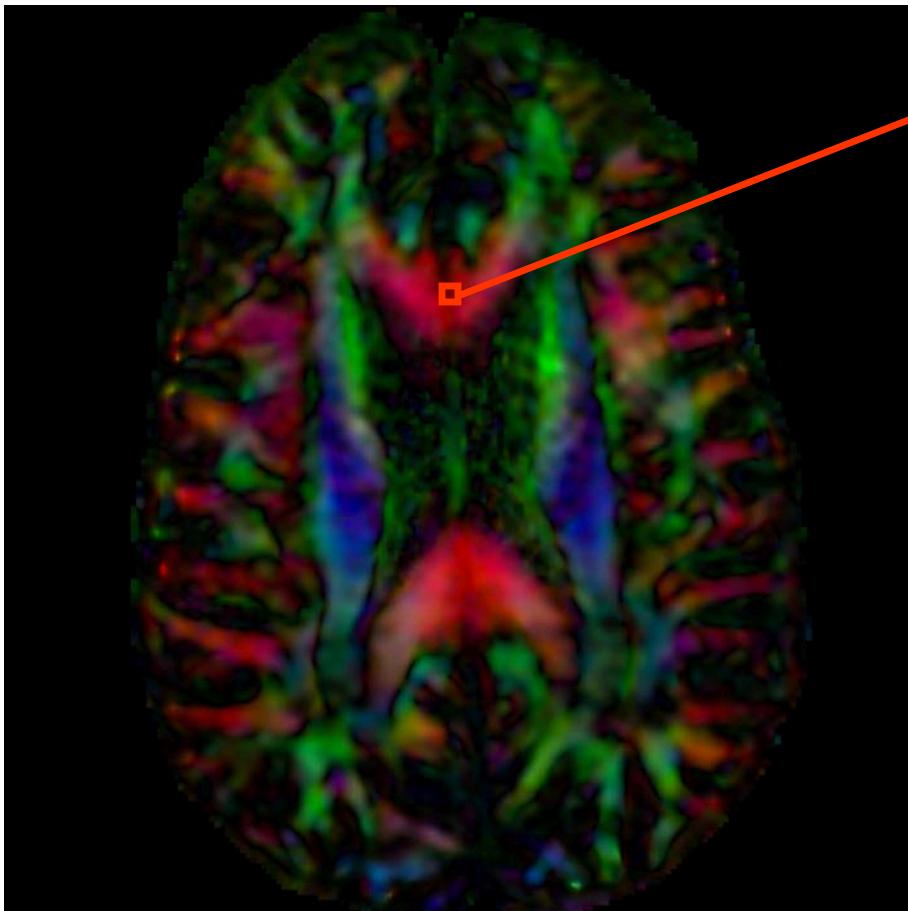


$$S_i = S_0 e^{-b \hat{g}_i^T \underline{D} \hat{g}_i}$$

↓

$$\underline{D} = \begin{bmatrix} D_{xx} & D_{xy} & D_{xz} \\ D_{yx} & D_{yy} & D_{yz} \\ D_{zx} & D_{zy} & D_{zz} \end{bmatrix}$$

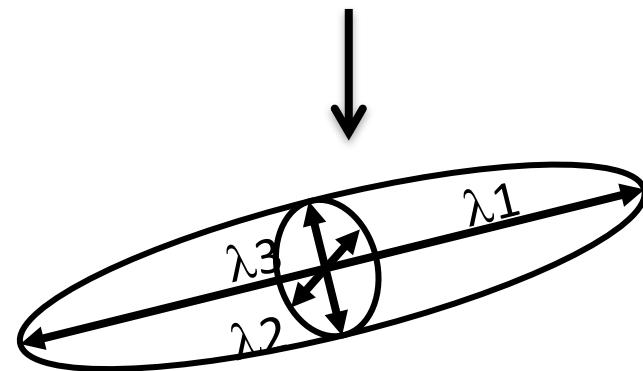
# Diffusion Tensor Imaging



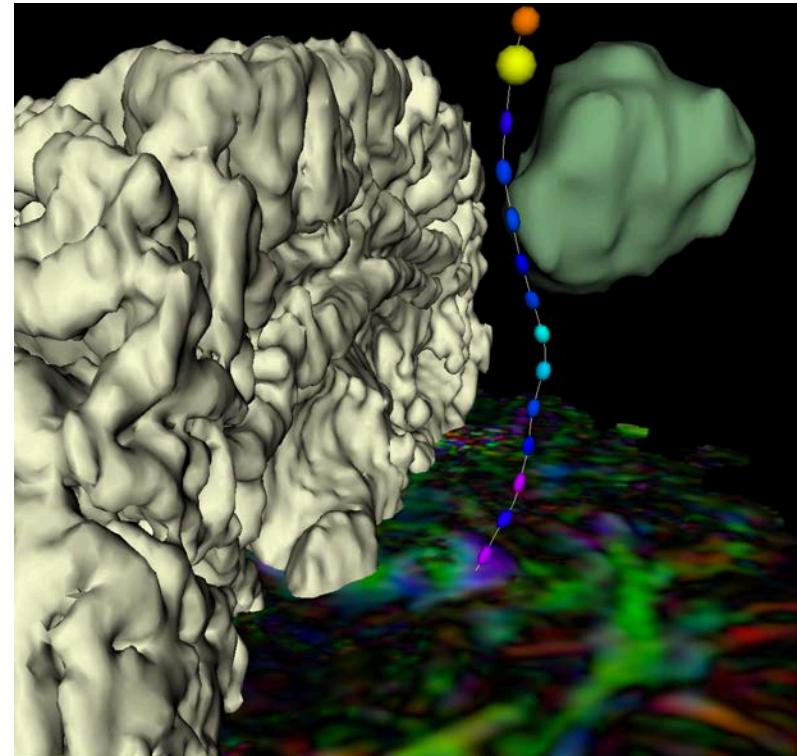
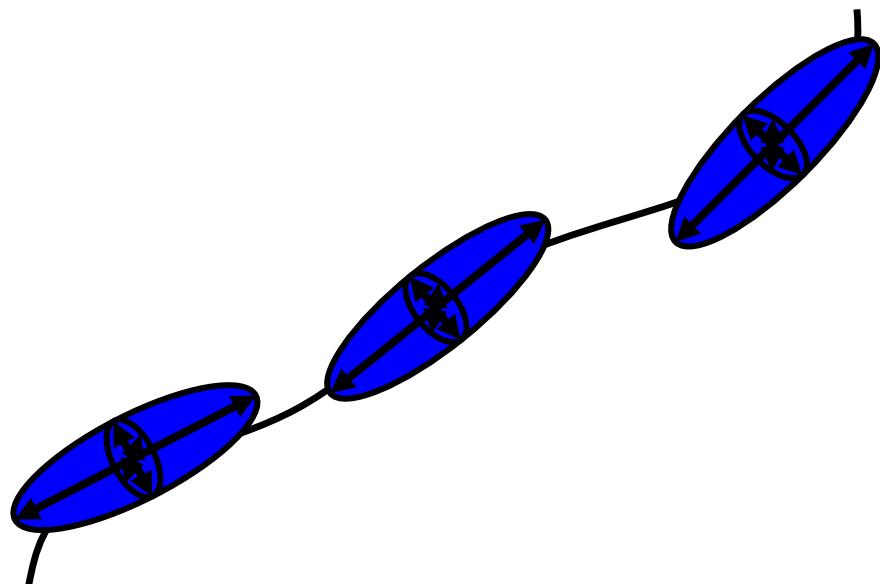
$$S_i = S_0 e^{-b \hat{g}^T \underline{D} \hat{g}_i}$$

↓

$$\underline{D} = \begin{bmatrix} D_{xx} & D_{xy} & D_{xz} \\ D_{yx} & D_{yy} & D_{yz} \\ D_{zx} & D_{zy} & D_{zz} \end{bmatrix}$$

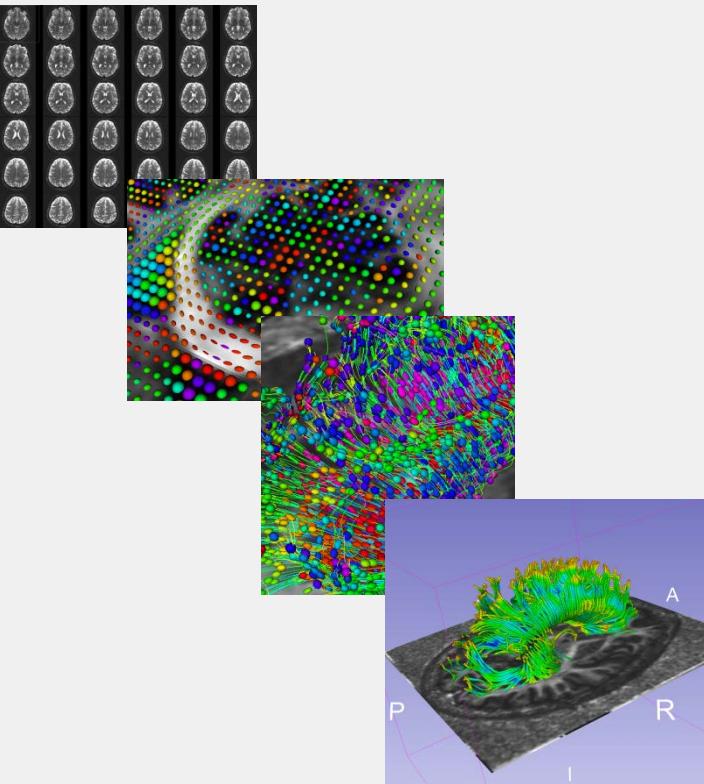


# Tractography



DTI tractography provides 3D reconstruction of the trajectory of white matter pathways

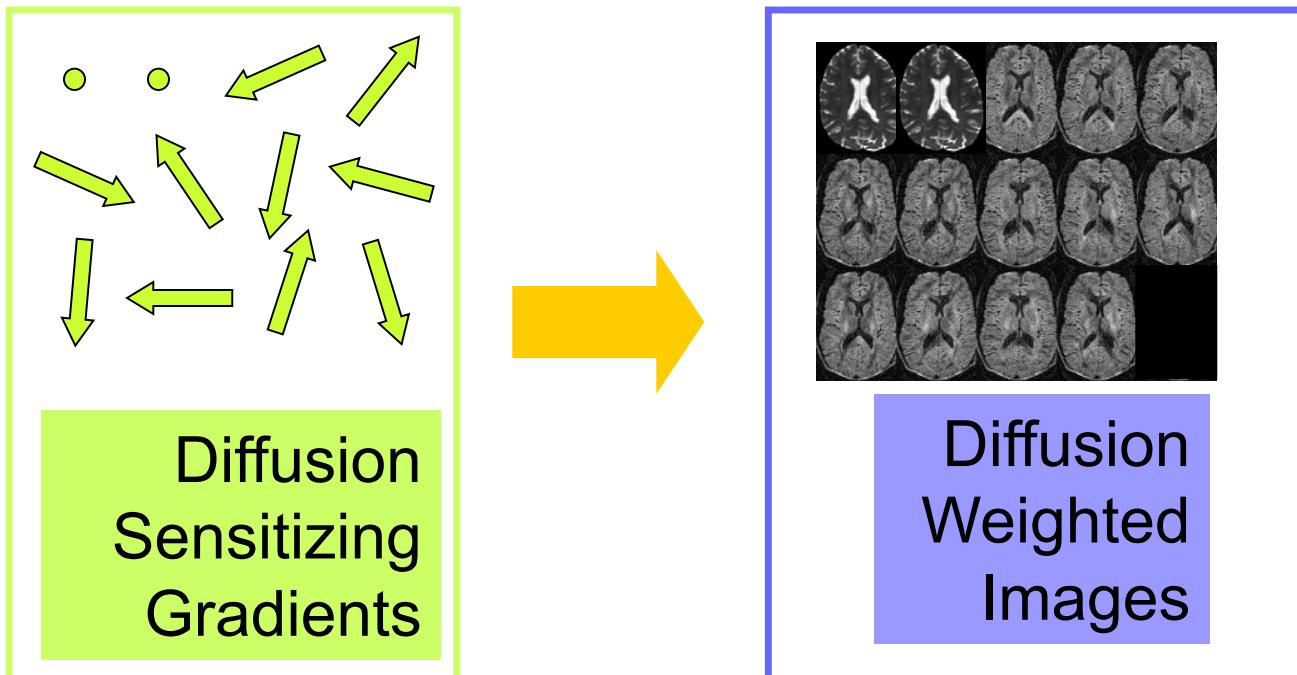
# Tutorial Outline



This tutorial is an introduction to the fundamentals of Diffusion MRI analysis, from the estimation of diffusion tensors to the interactive 3D visualization of fiber tracts.

# Tutorial Dataset

The tutorial dataset SlicerDiffusionMRITutorialData is a Diffusion Weighted MR scan of the brain acquired with 41 gradient directions and one baseline.



# 3D Slicer

The tutorial uses the 3D Slicer (Version 4.8.1, revision 26813, Stable Release) software available at:

<http://download.slicer.org>

## *Disclaimer*

It is the responsibility of the user of 3DSlicer to comply with both the terms of the license and with the applicable laws, regulations and rules. Slicer is a tool for research, and is not FDA approved.

# SlicerDMRI

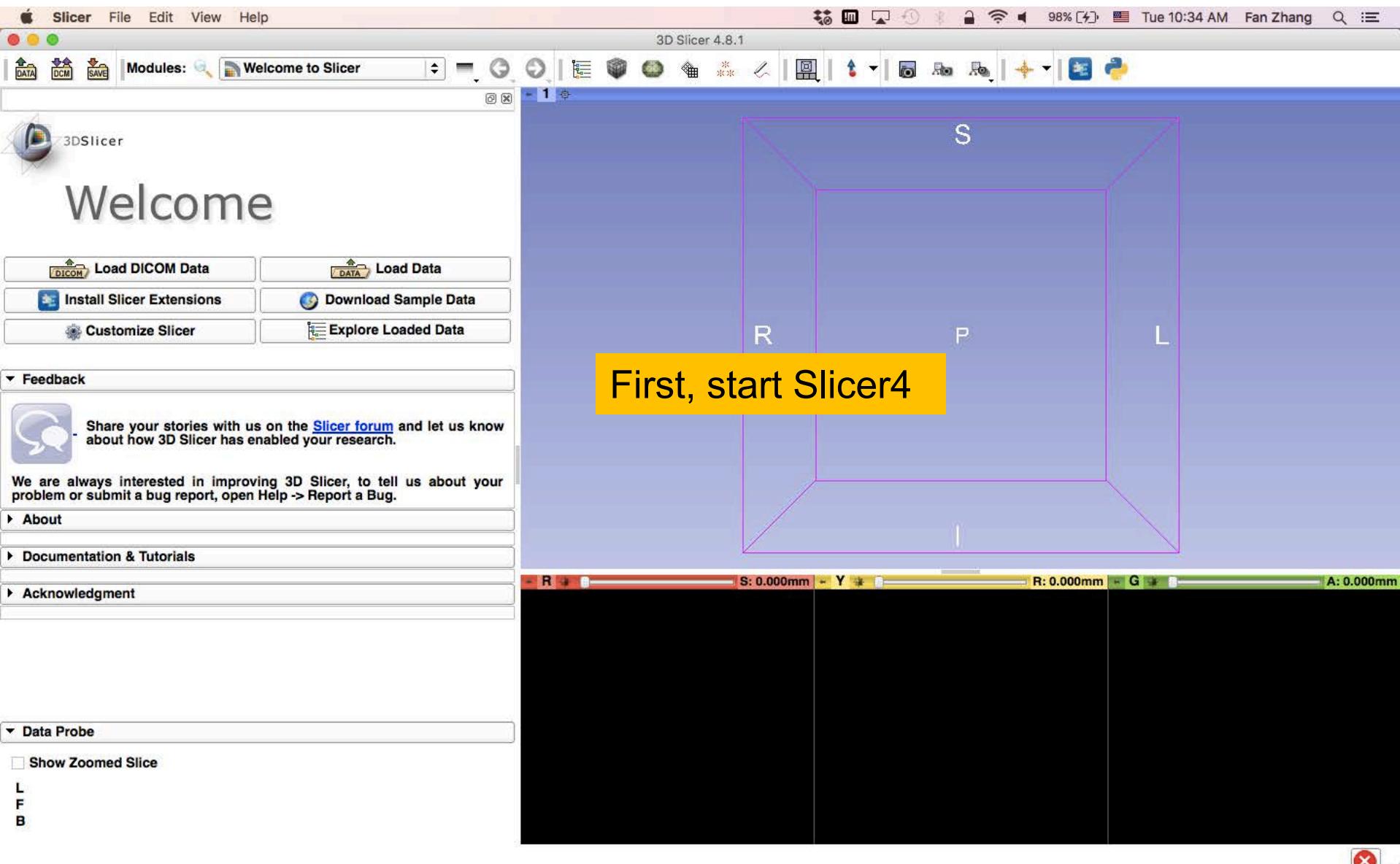
An open-source project to improve and extend diffusion magnetic resonance imaging software in 3D Slicer:

<http://dmri.slicer.org>

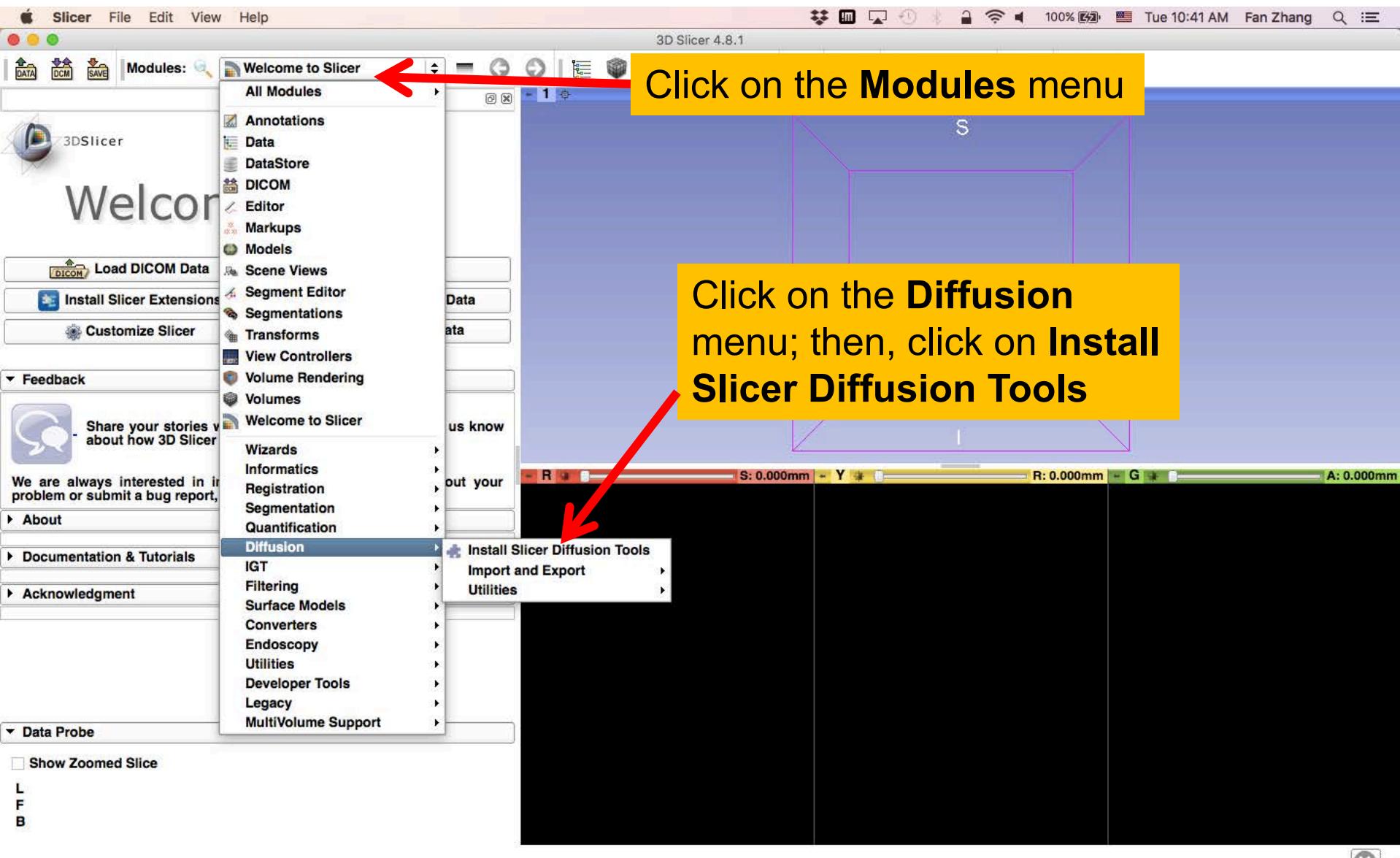
## *Disclaimer*

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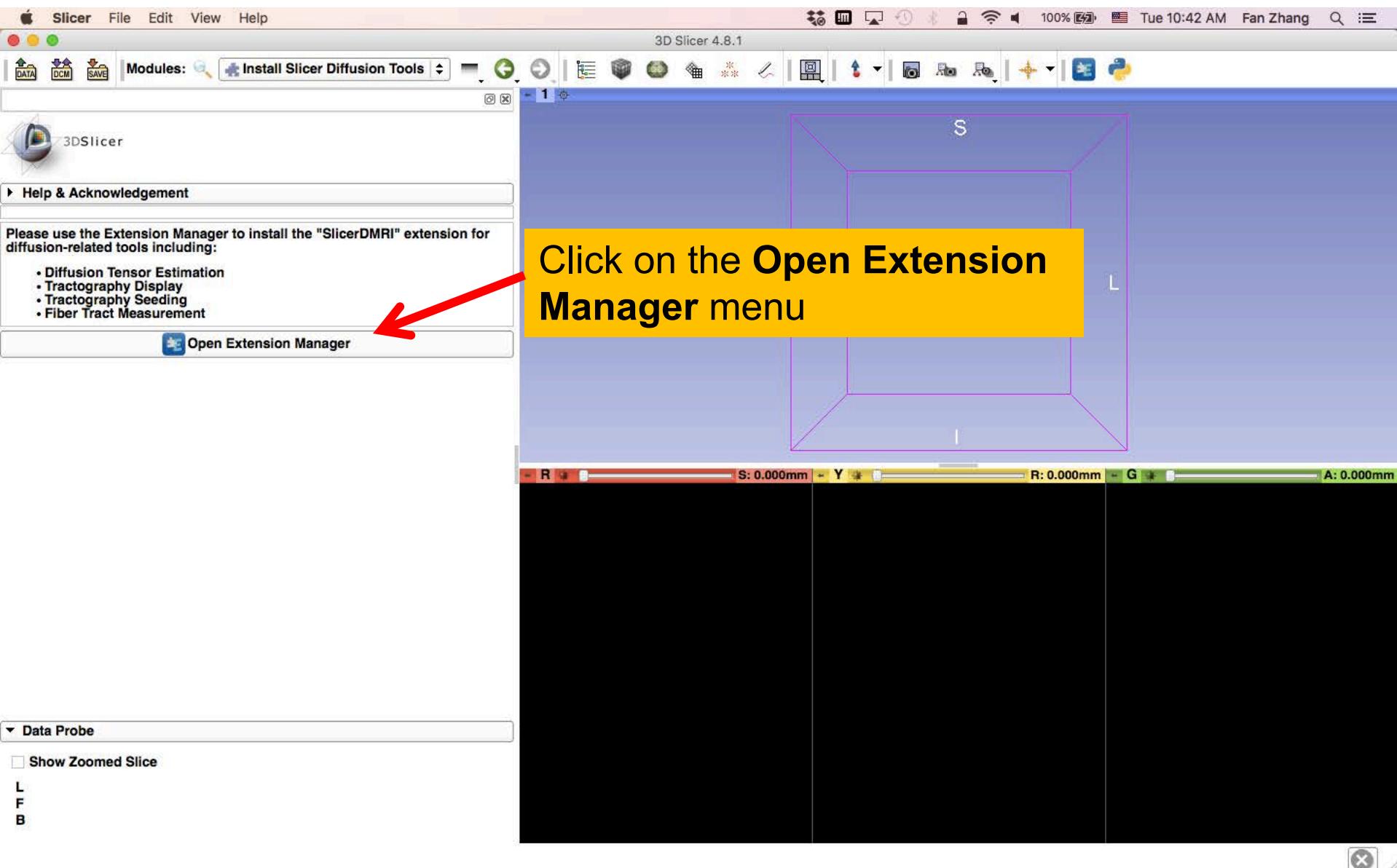
# Install SlicerDMRI



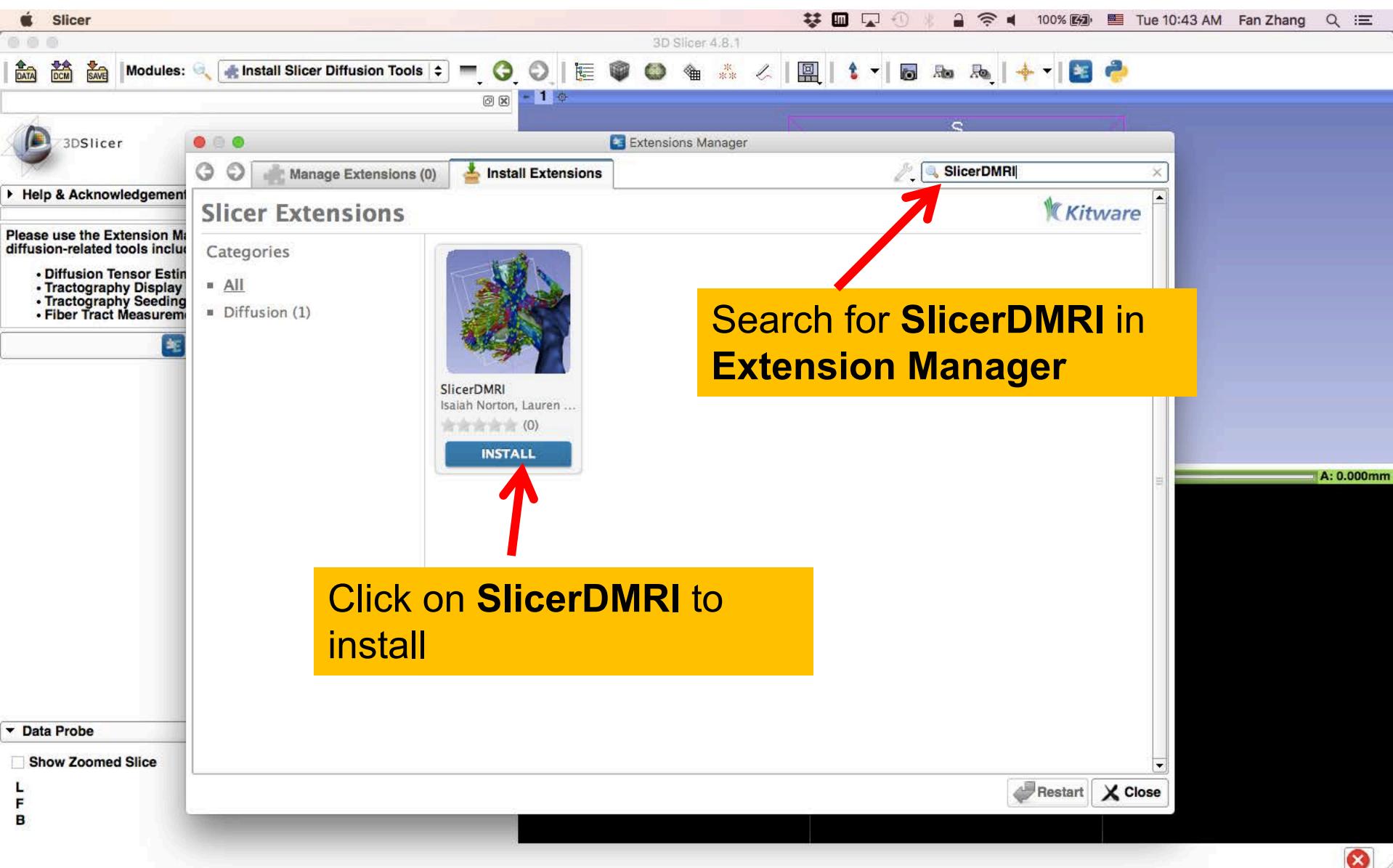
# Install SlicerDMRI



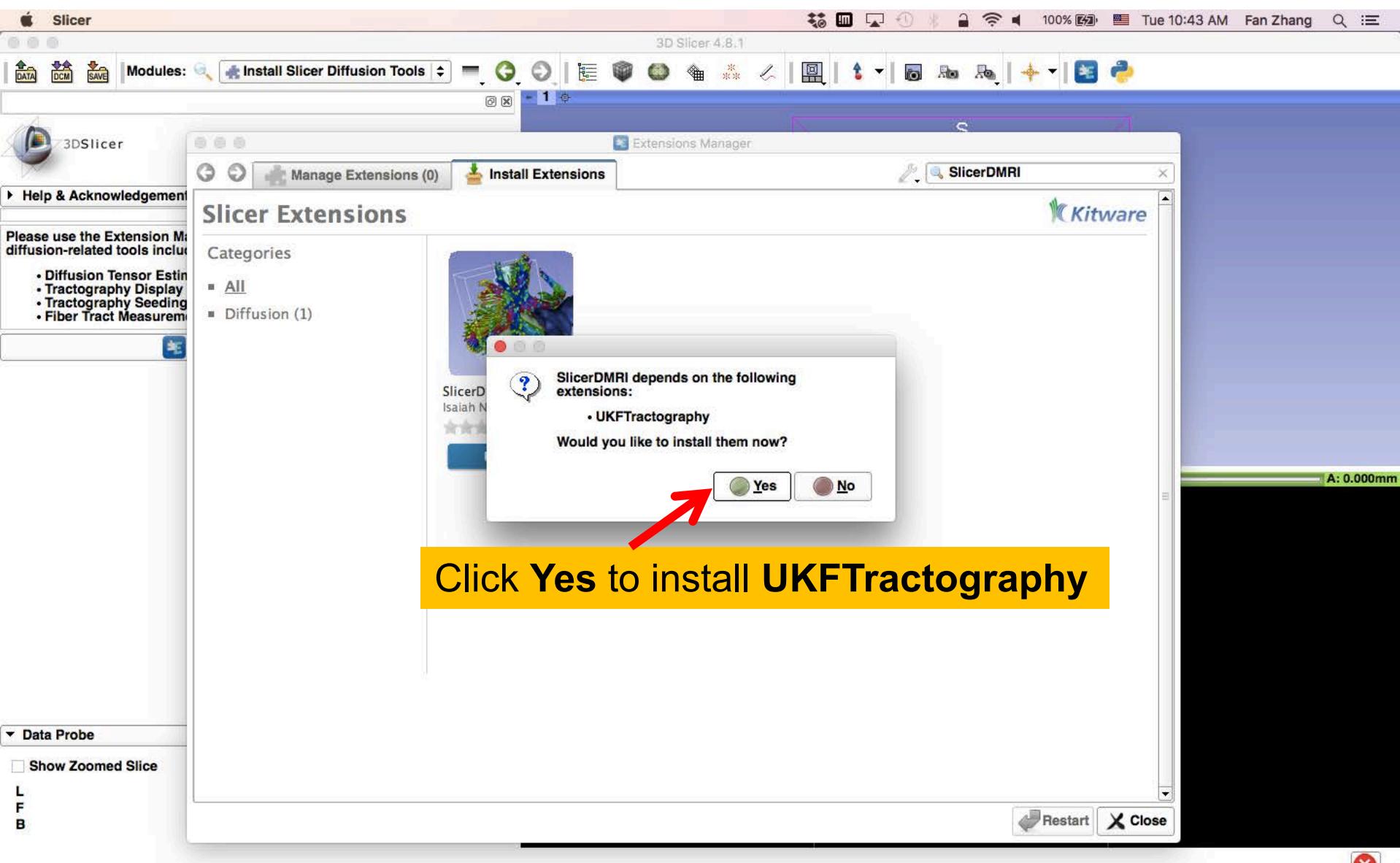
# Install SlicerDMRI



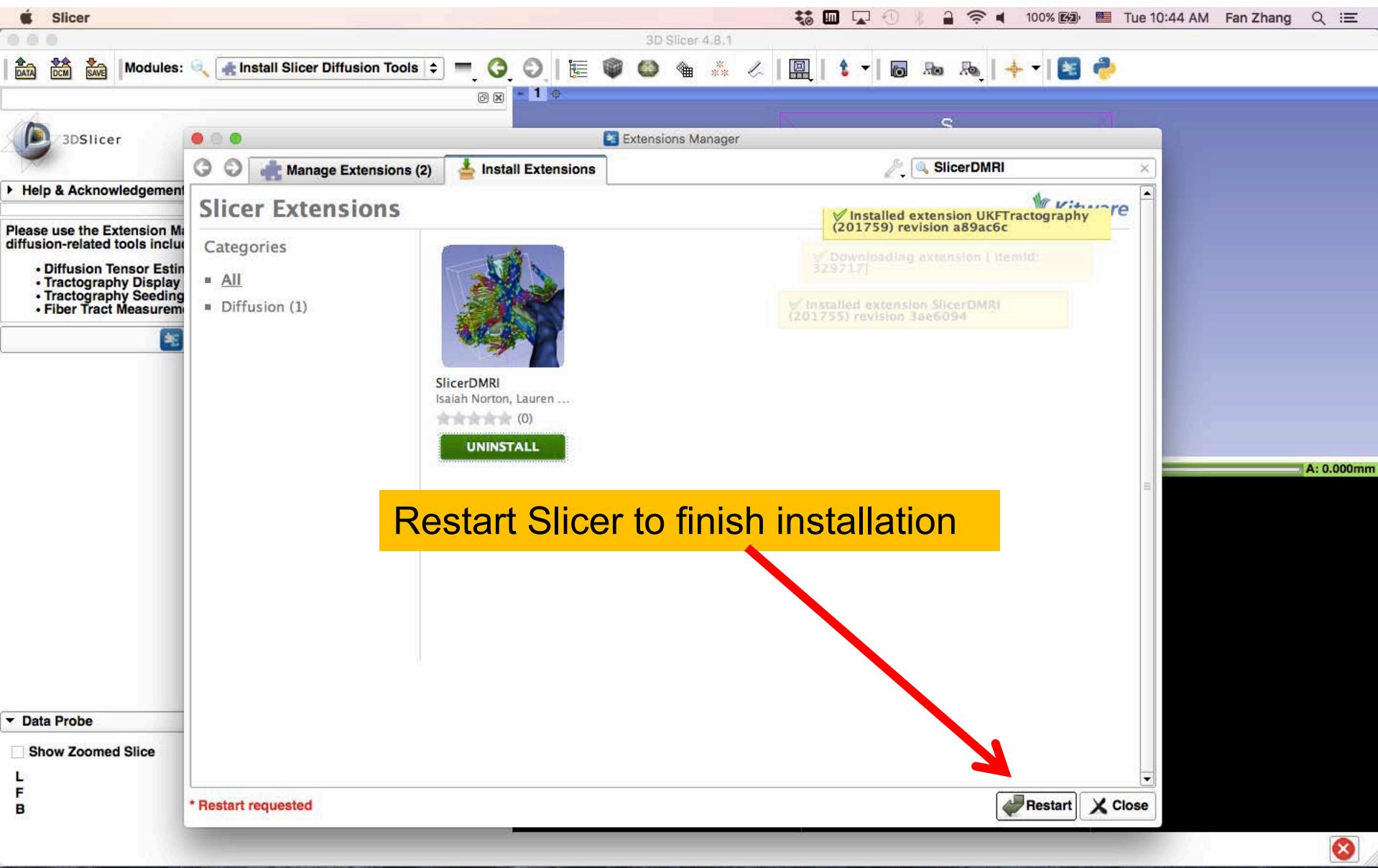
# Install SlicerDMRI



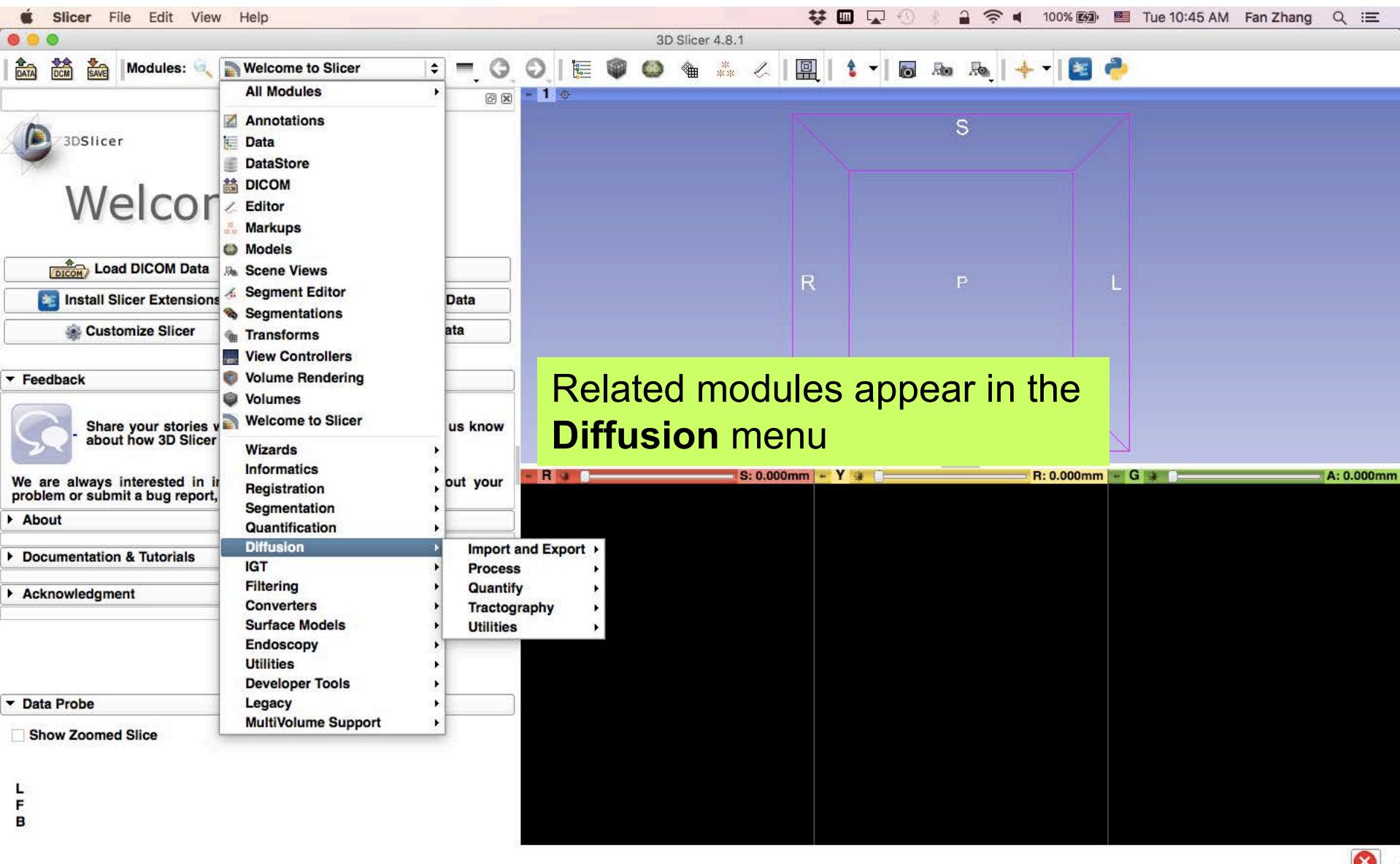
# Install SlicerDMRI



# Install SlicerDMRI



# Install SlicerDMRI

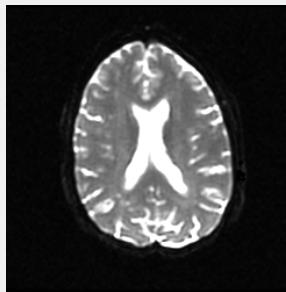


# Learning Objectives

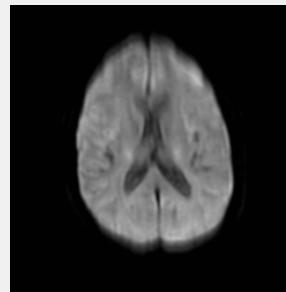
Following this tutorial, you'll be able to

- 1) Estimate a tensor volume from a set of Diffusion Weighted Images
- 2) Understand the shape and size of the diffusion ellipsoid
- 3) Reconstruct DTI tracts from a pre-defined region of interest
- 4) Interactively visualize DTI tracts seeded from a fiducial

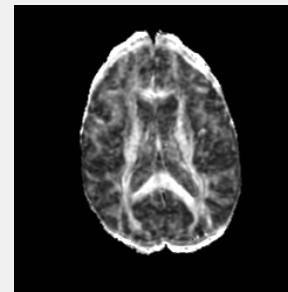
# MR Diffusion Analysis Pipeline



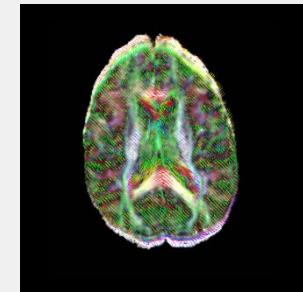
DWI  
Acquisition



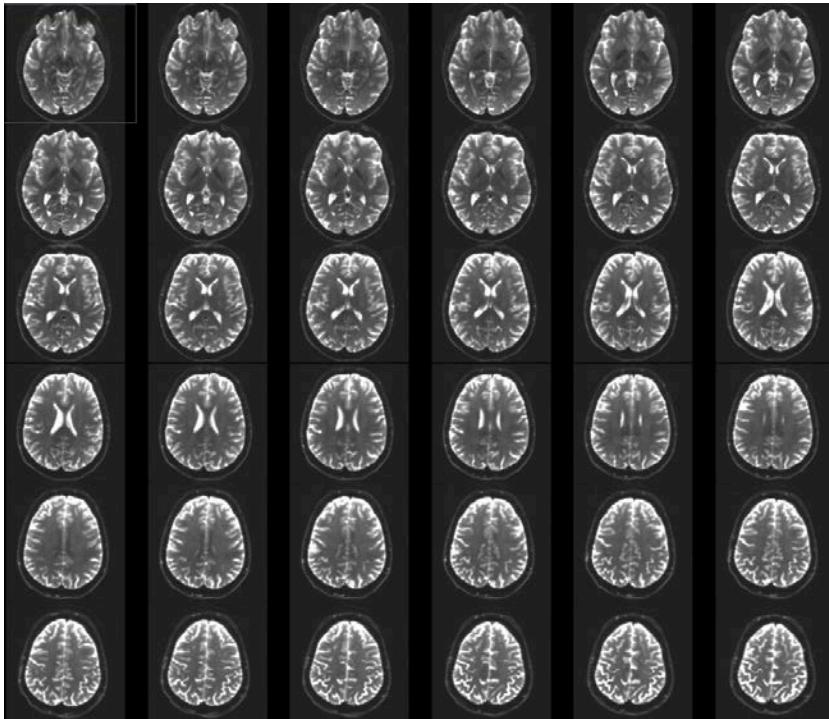
Tensor  
Calculation



Scalar  
Maps



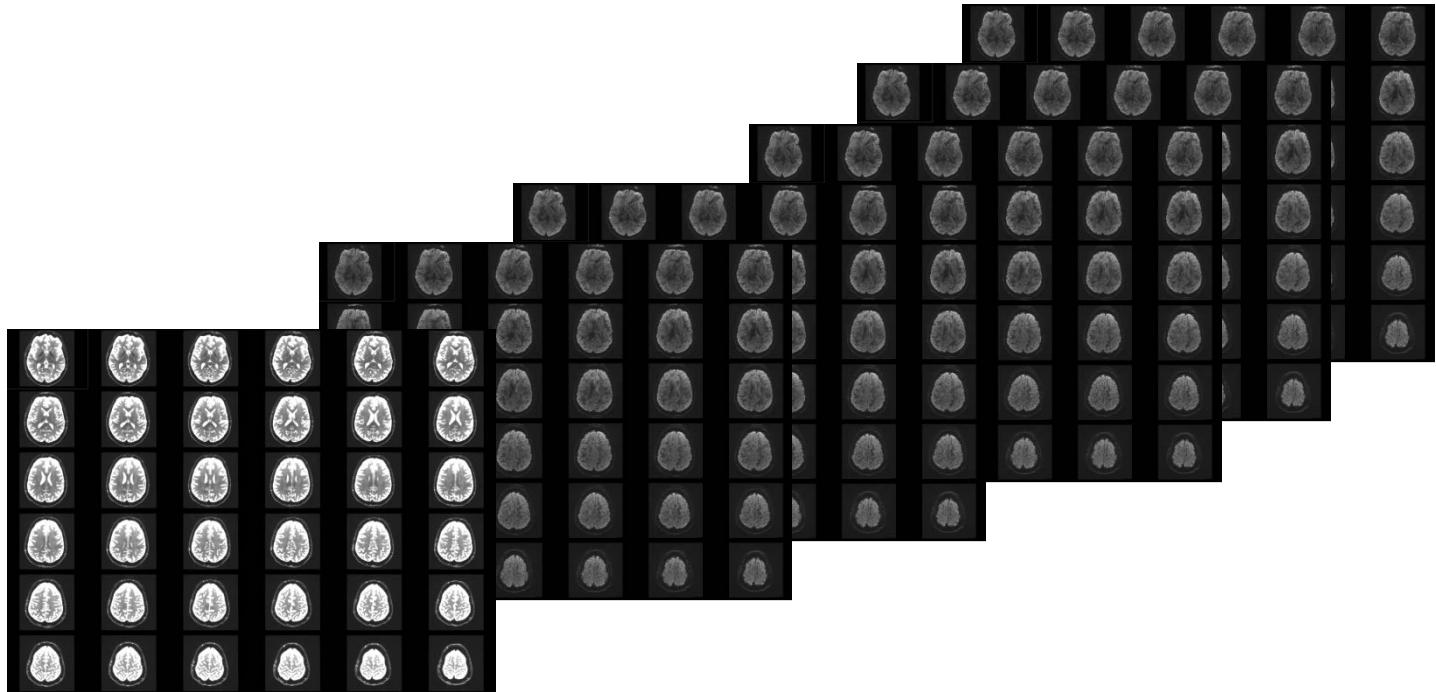
3D  
Visualization



# Part 1:

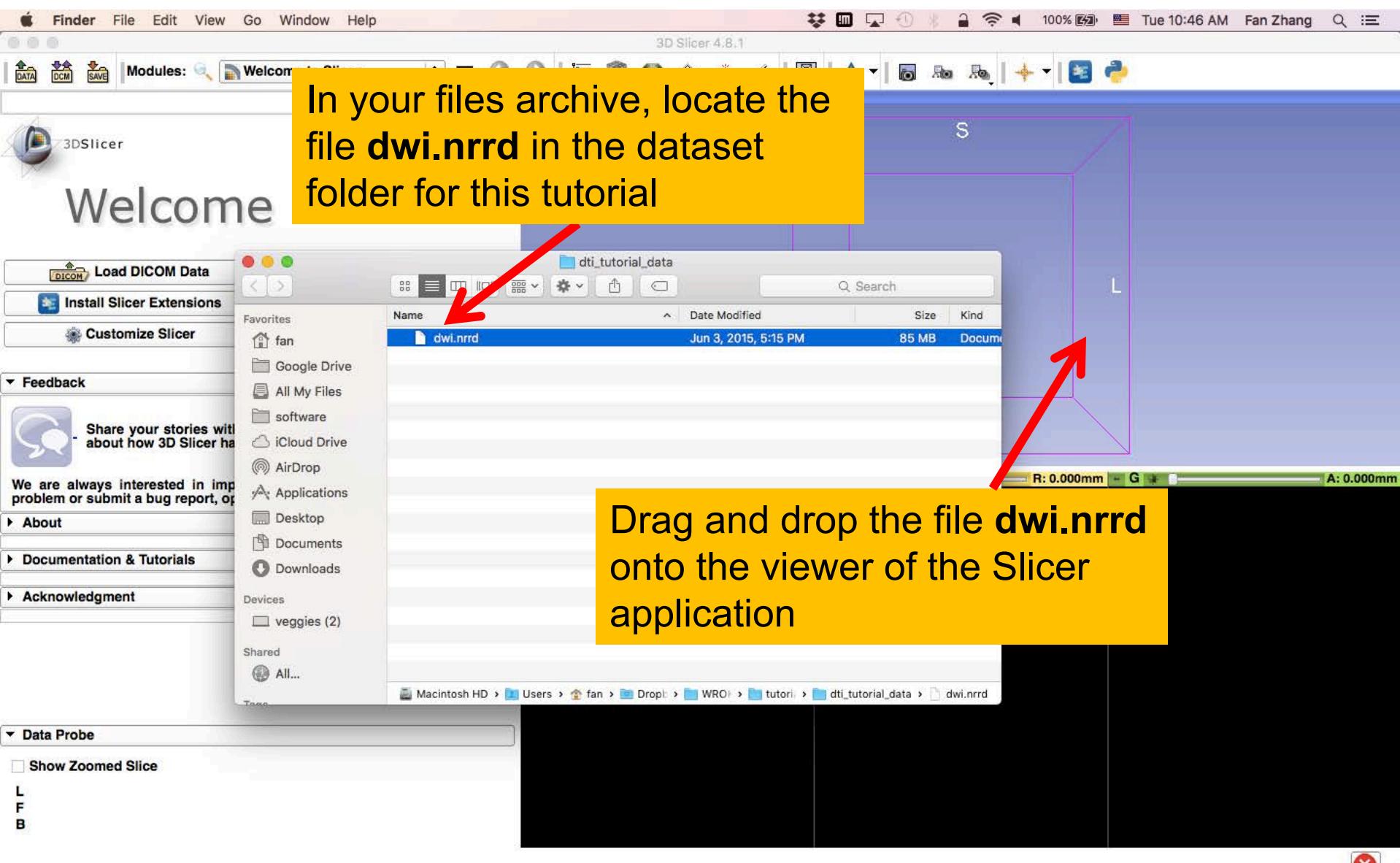
## From DWI images to Tensors

# Understanding the DWI Dataset

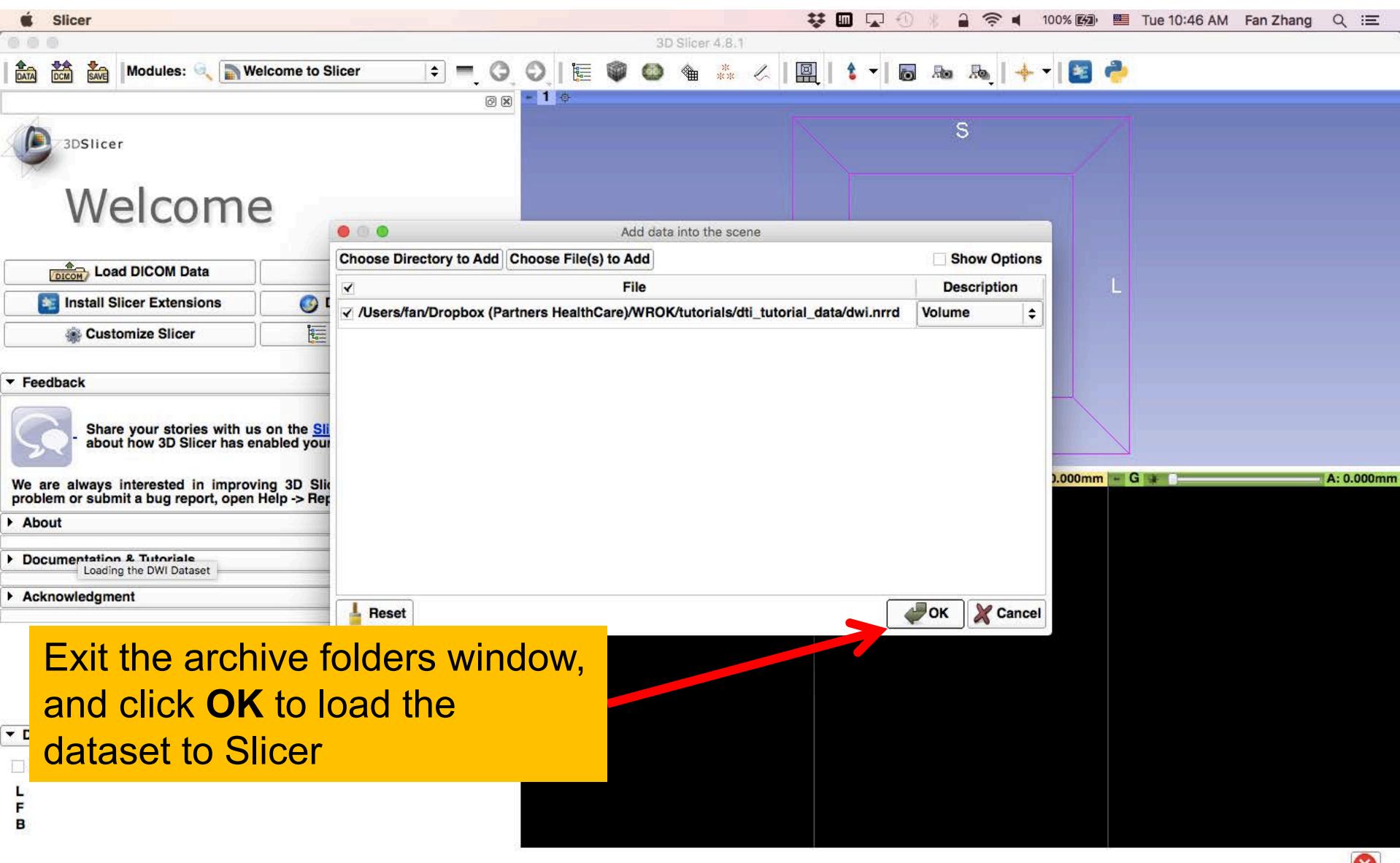


The Diffusion Weighted Imaging (DWI) dataset is composed of 41 volumes acquired with 41 different diffusion-sensitizing gradient directions, and one baseline image acquired without diffusion weighting.

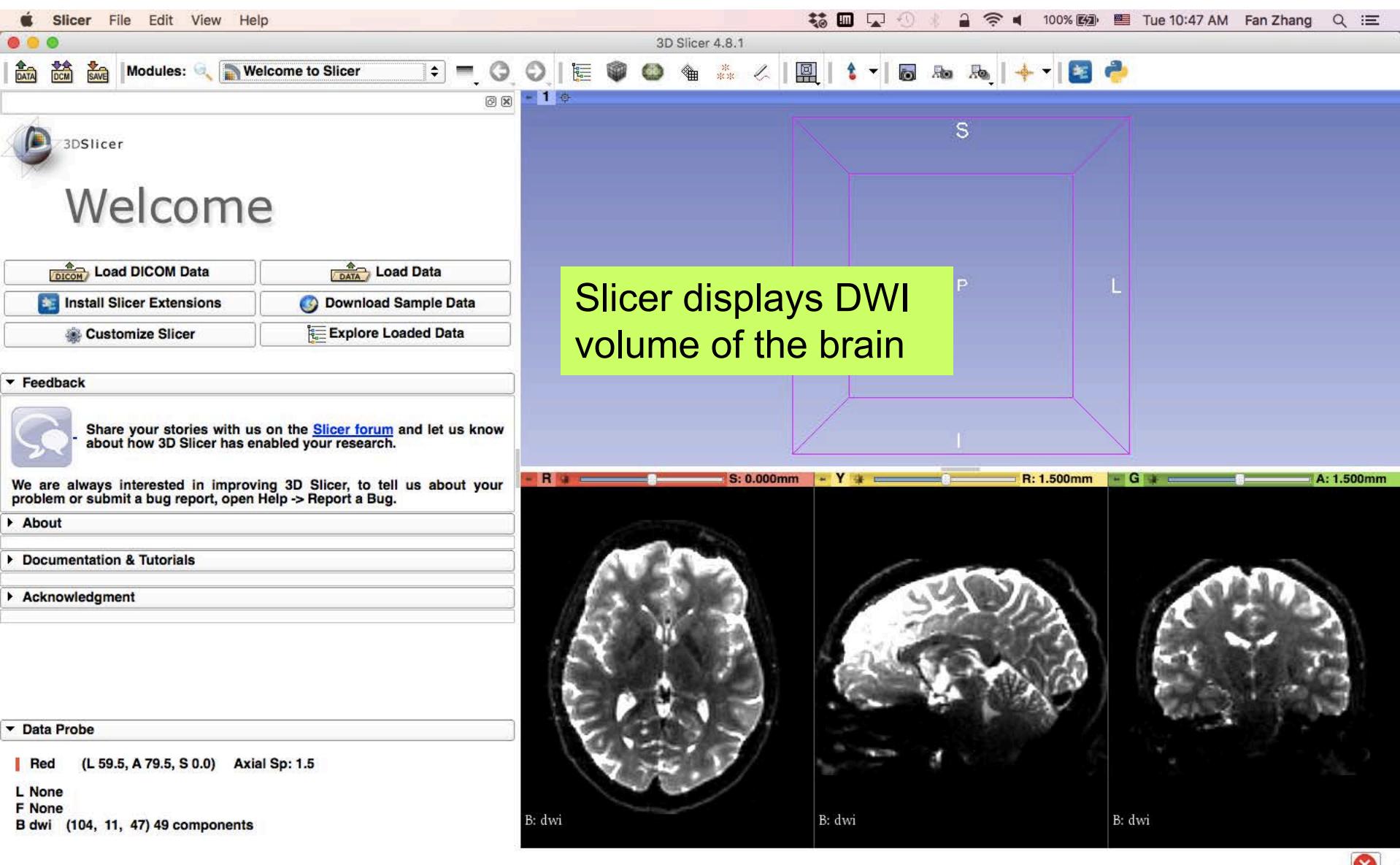
# Loading the DWI Dataset



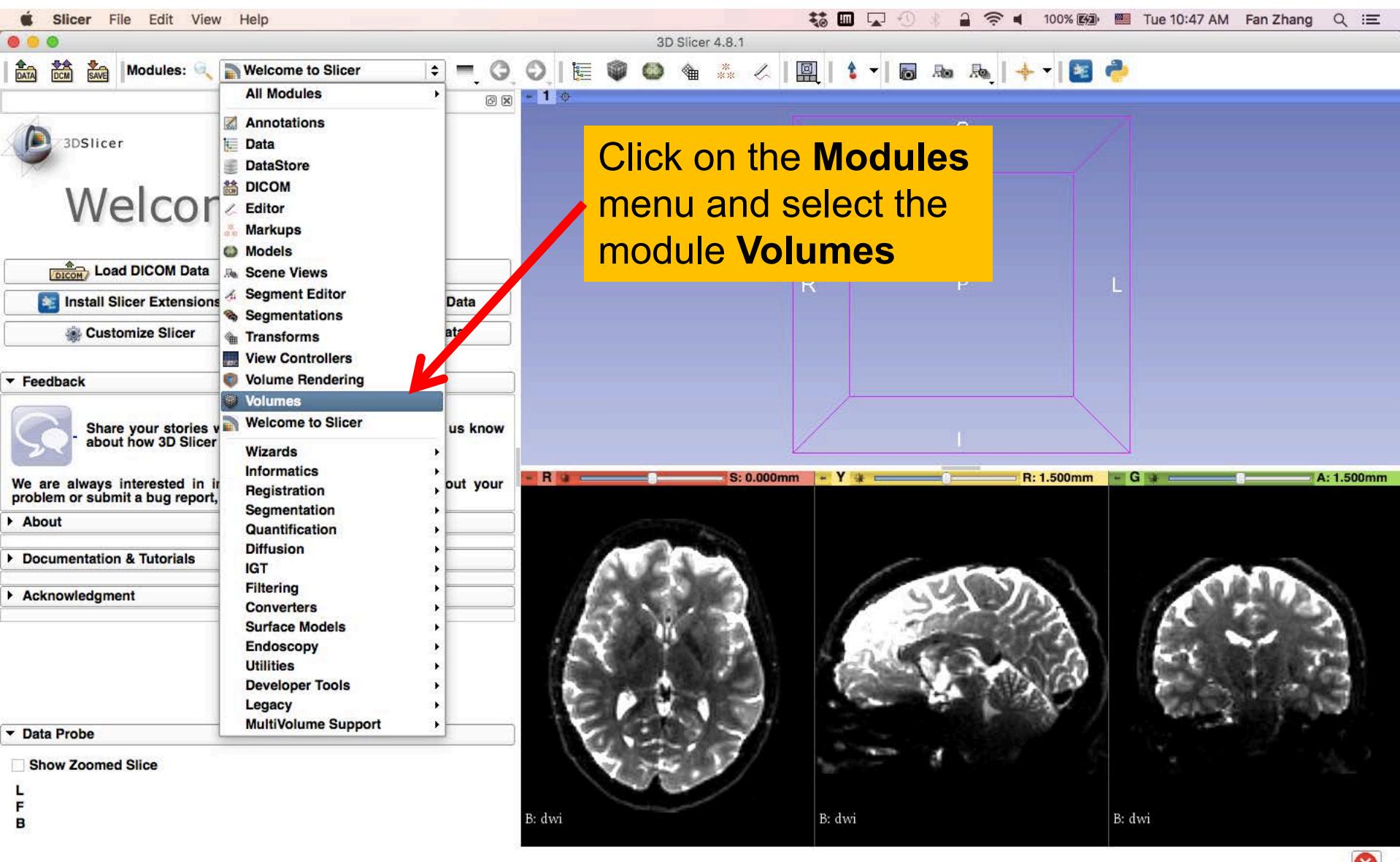
# Loading the DWI Dataset



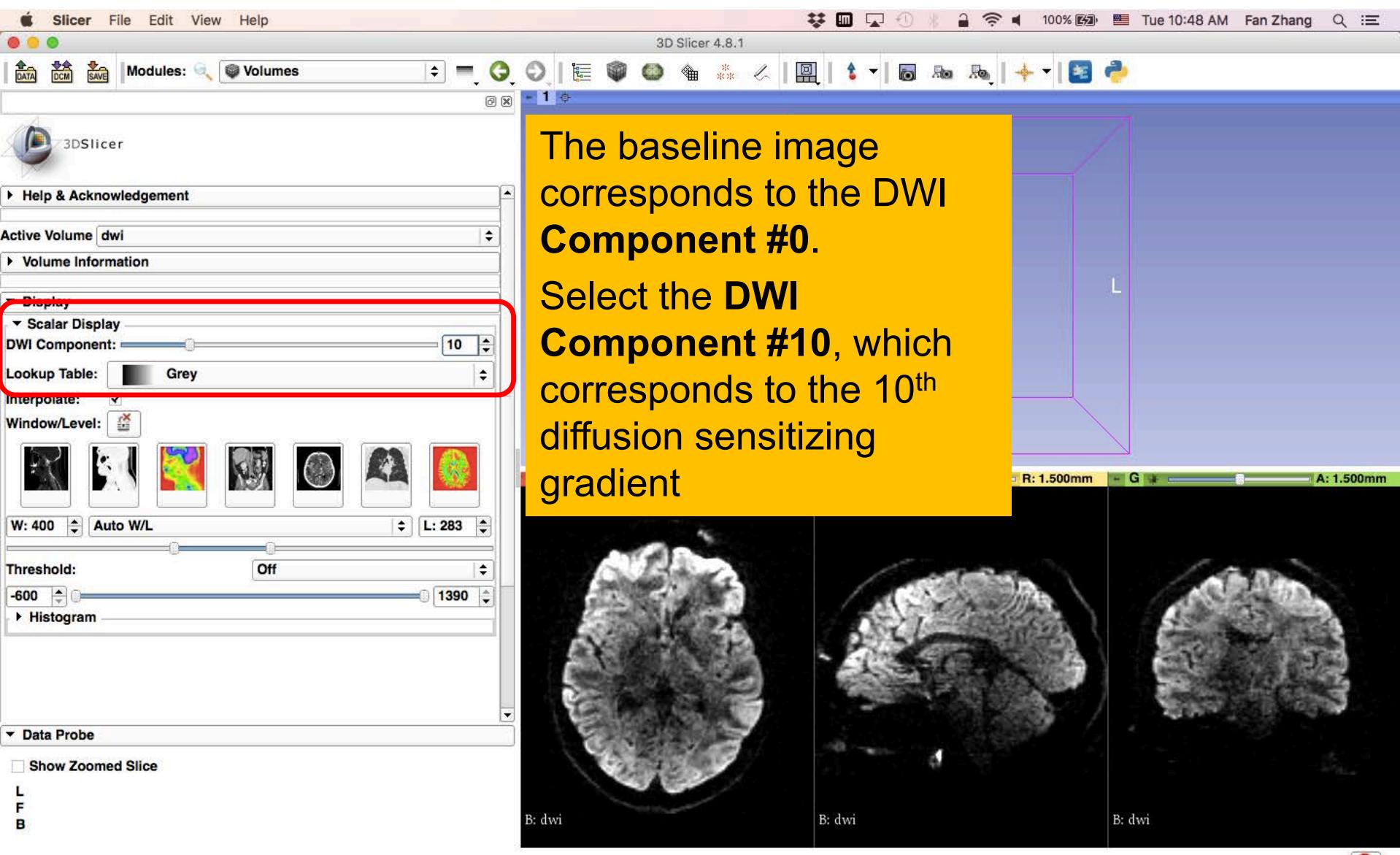
# Loading the DWI Dataset



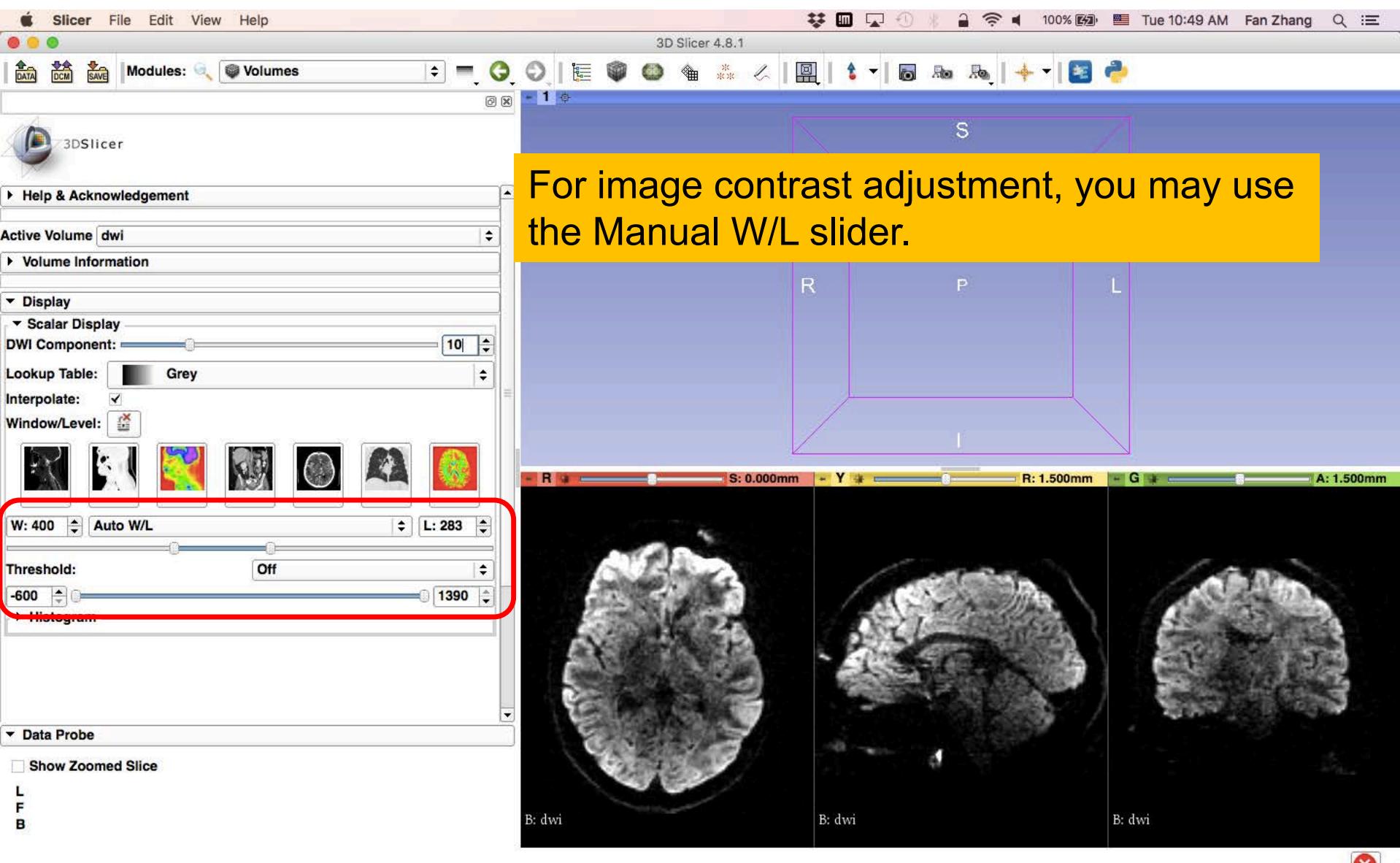
# Loading the DWI Dataset



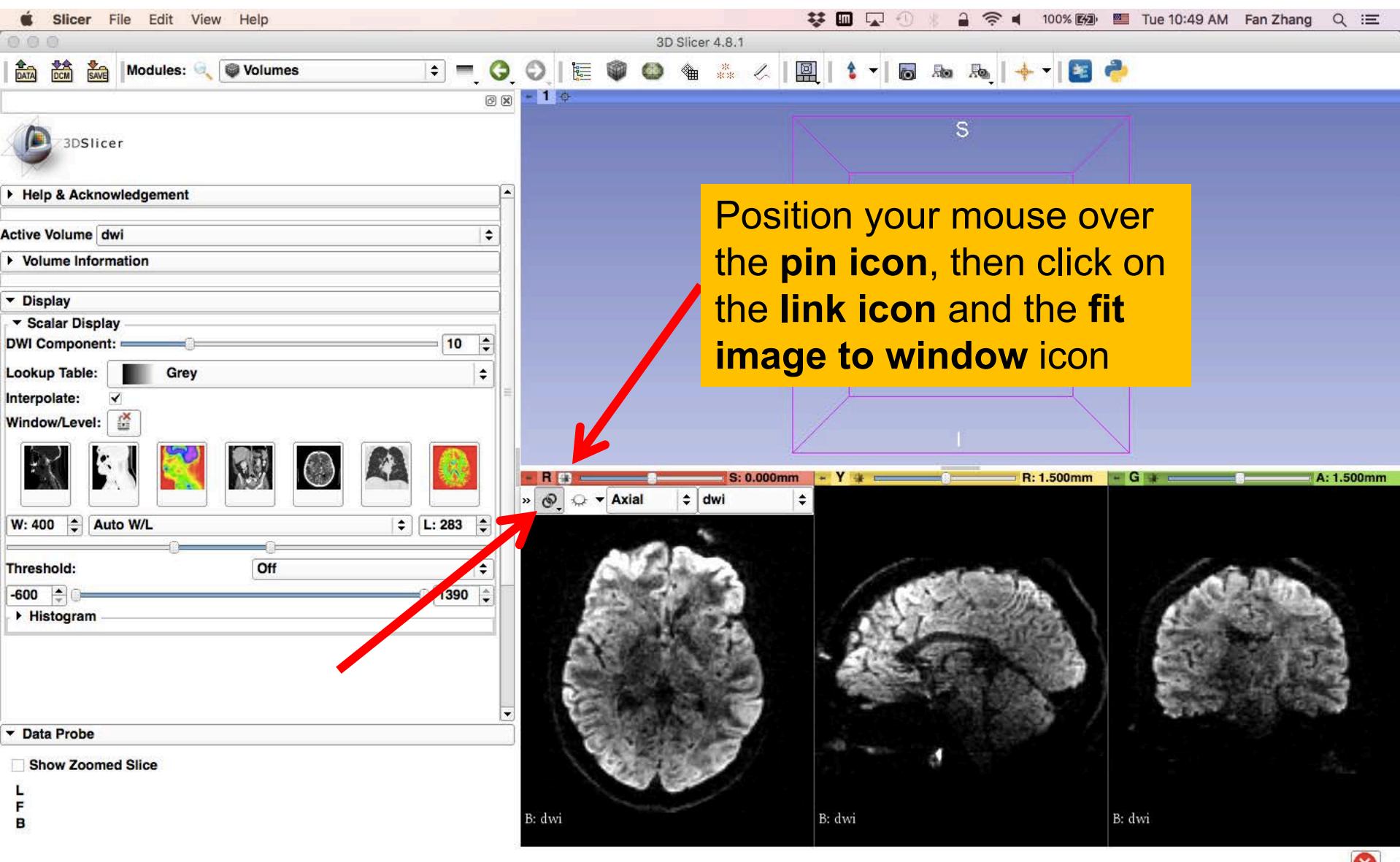
# Loading the DWI Dataset



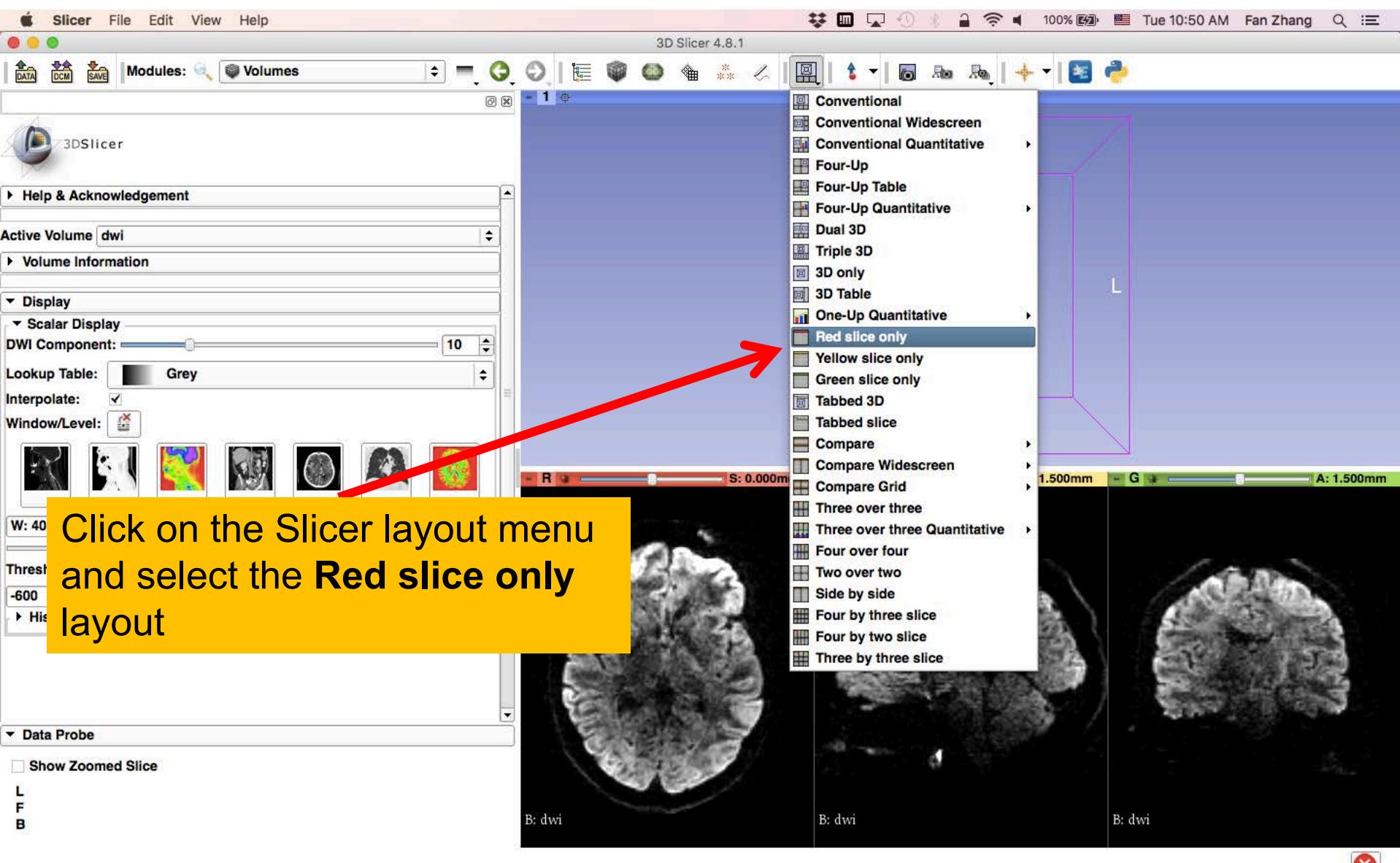
# Loading the DWI Dataset



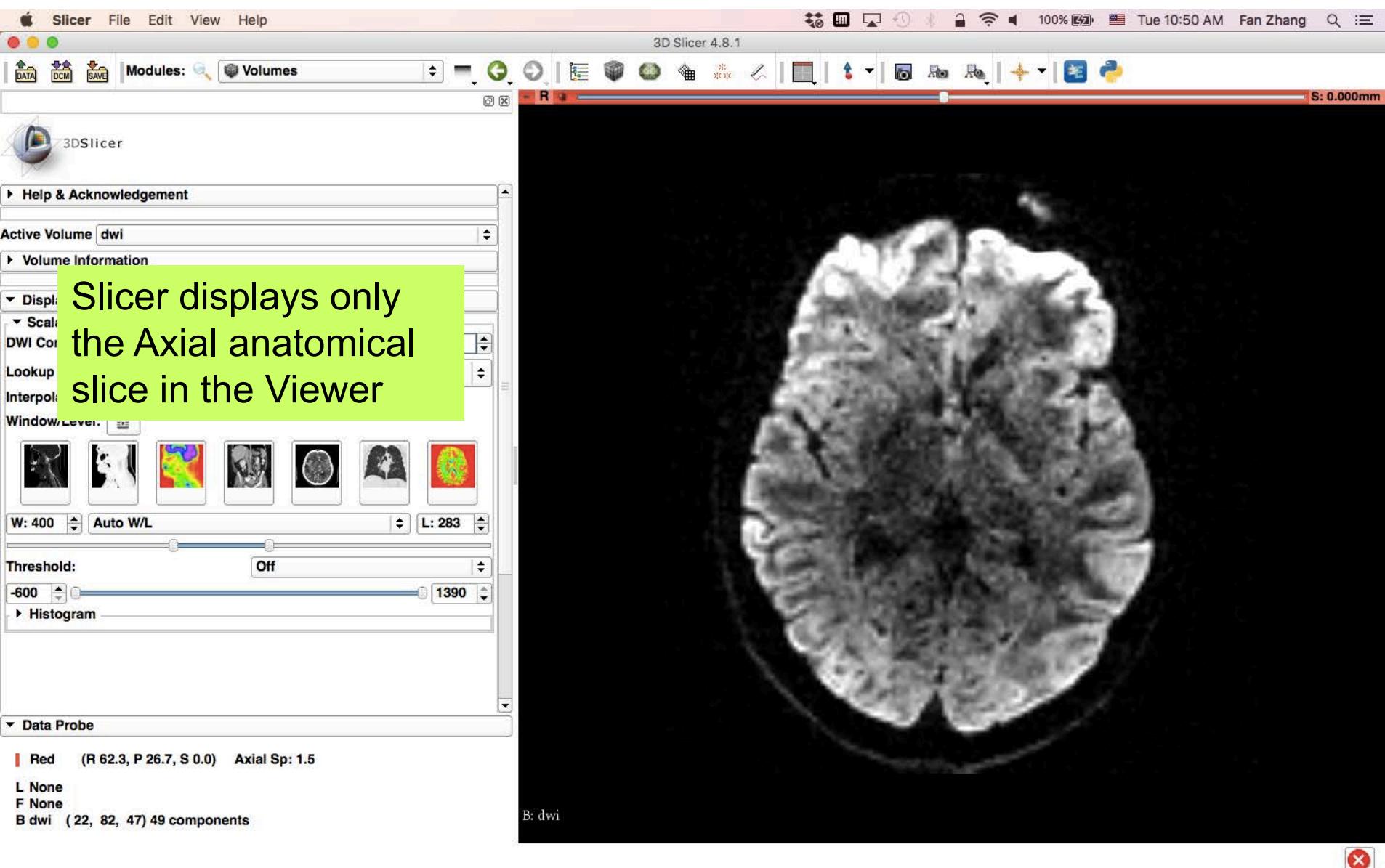
# Loading the DWI Dataset



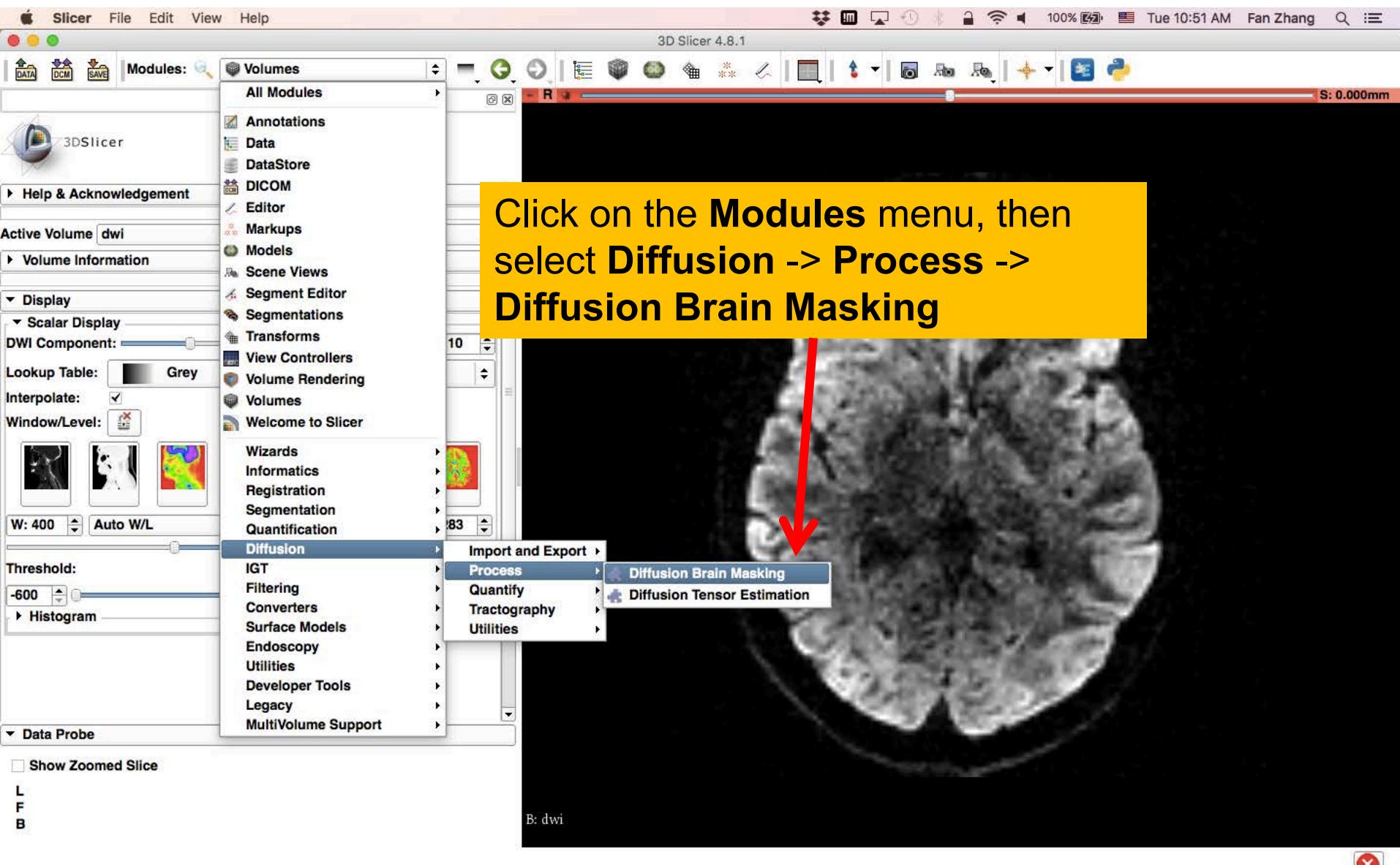
# Loading the DWI Dataset



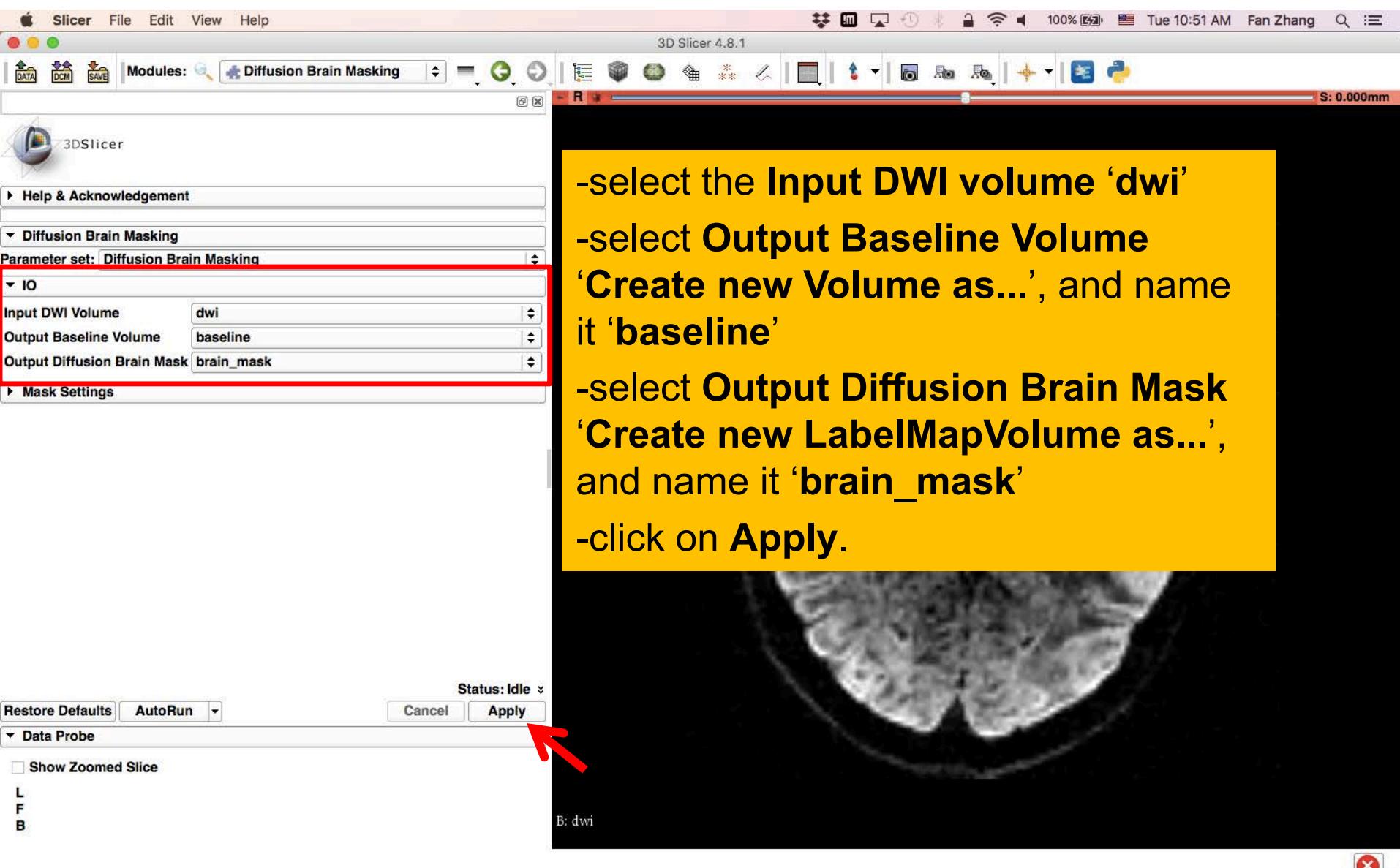
# Loading the DWI Dataset



# Creating a brain mask



# Creating a brain mask

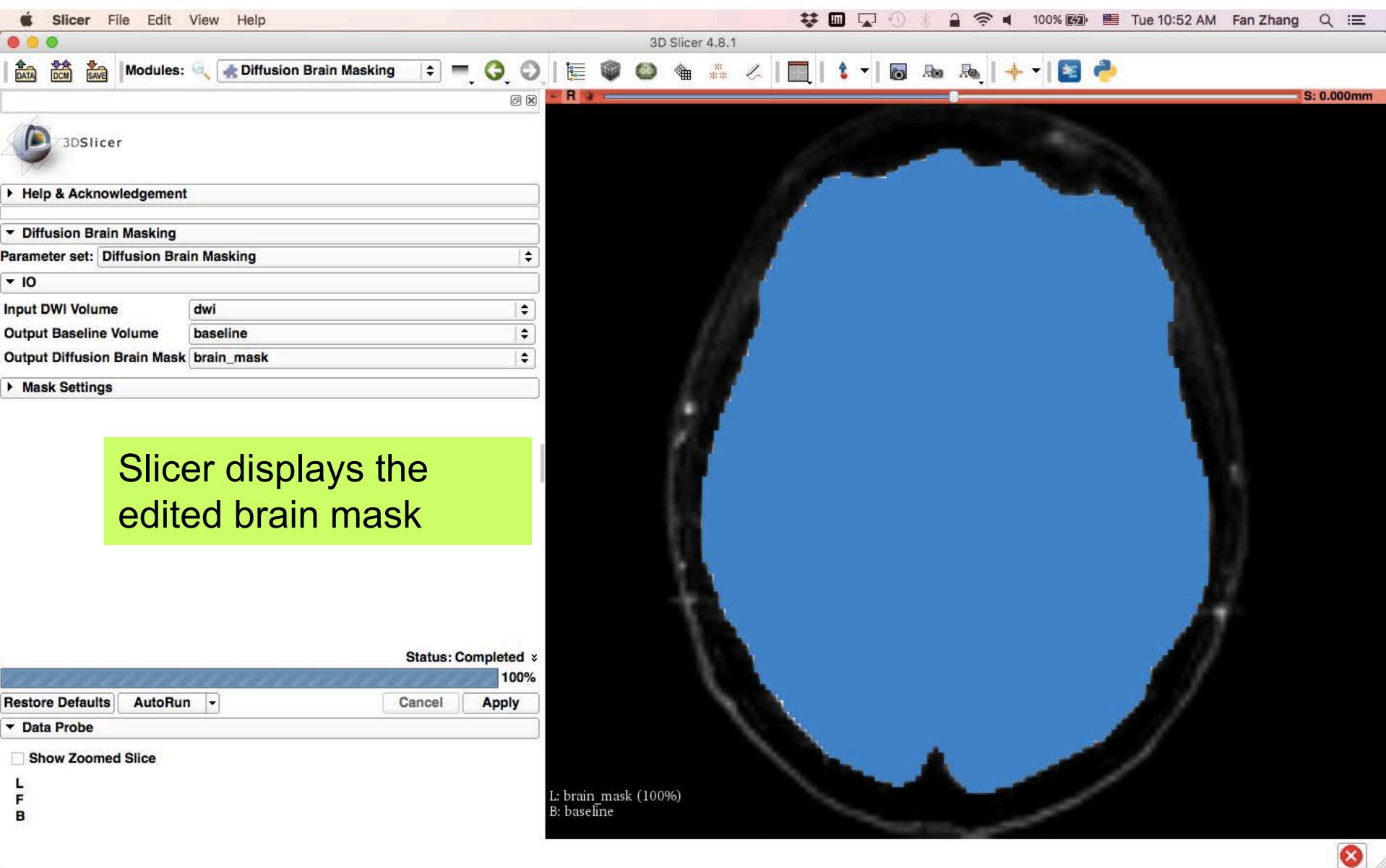


The screenshot shows the 3D Slicer 4.8.1 interface with the 'Diffusion Brain Masking' module selected. A yellow callout box on the right contains the following instructions:

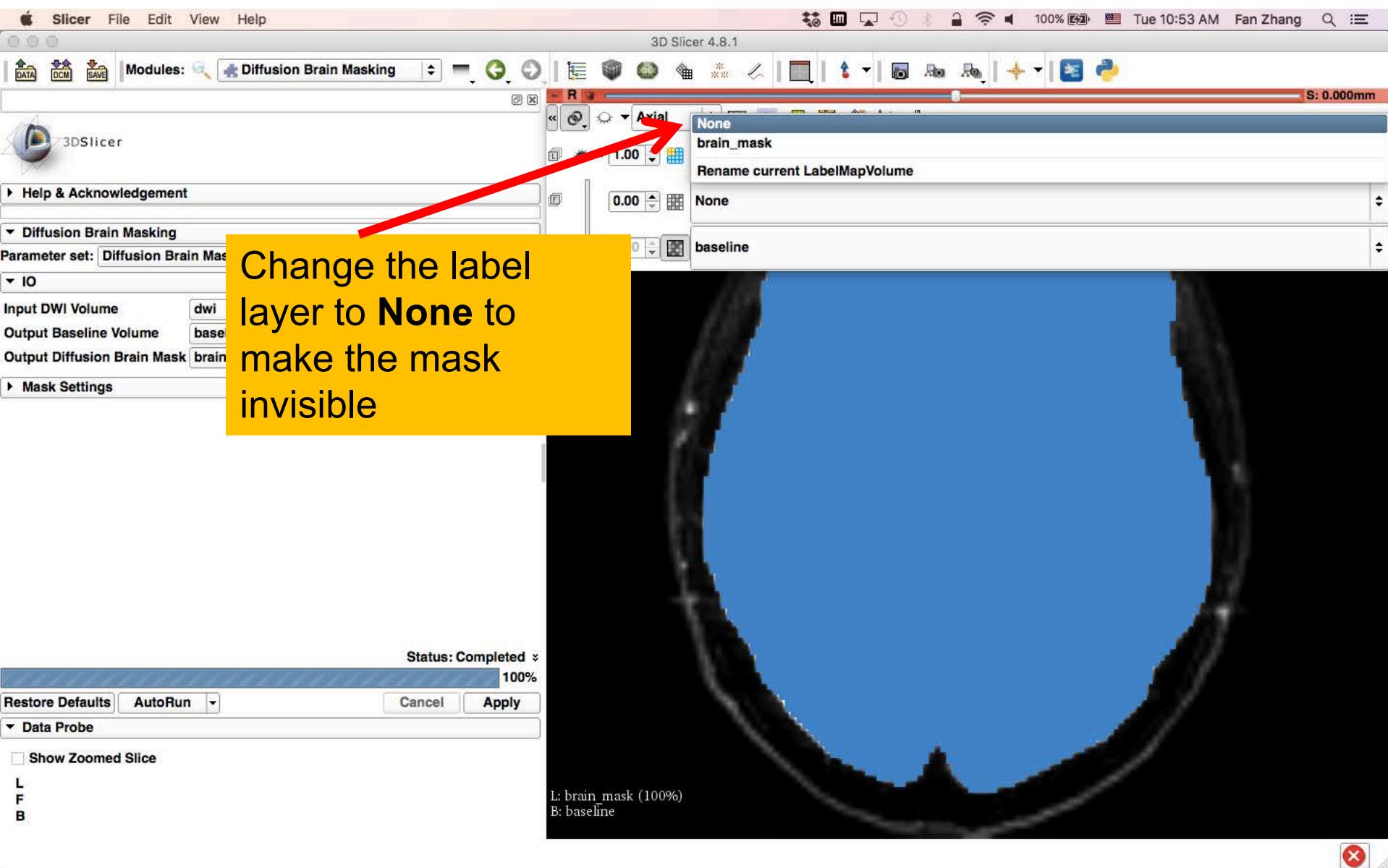
- select the **Input DWI volume 'dwi'**
- select **Output Baseline Volume 'Create new Volume as...', and name it 'baseline'**
- select **Output Diffusion Brain Mask 'Create new LabelMapVolume as...', and name it 'brain\_mask'**
- click on **Apply**.

The 'IO' parameters in the module panel are highlighted with a red box. The 'Apply' button at the bottom of the panel is also highlighted with a red arrow. The status bar at the bottom of the interface shows 'Status: Idle'.

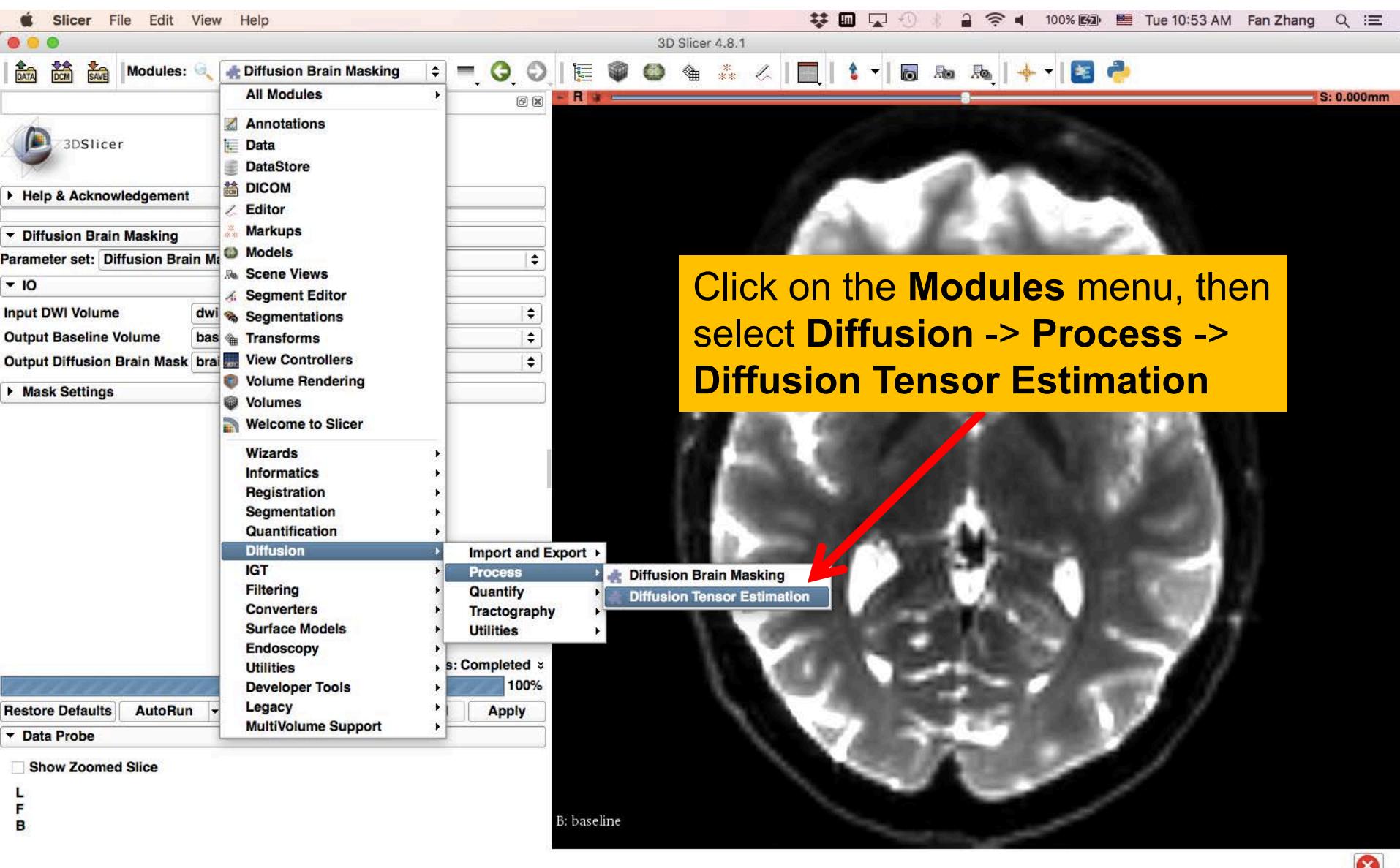
# Creating a brain mask



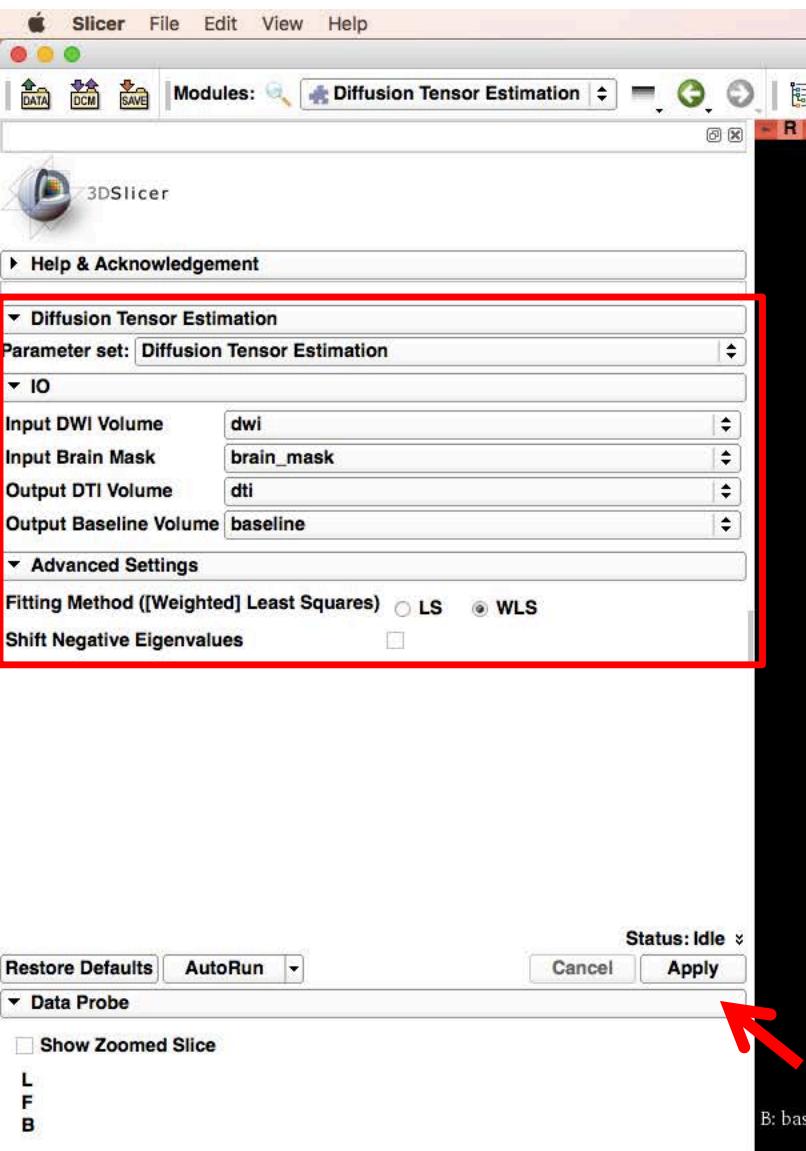
# Creating a brain mask



# Estimating the tensor

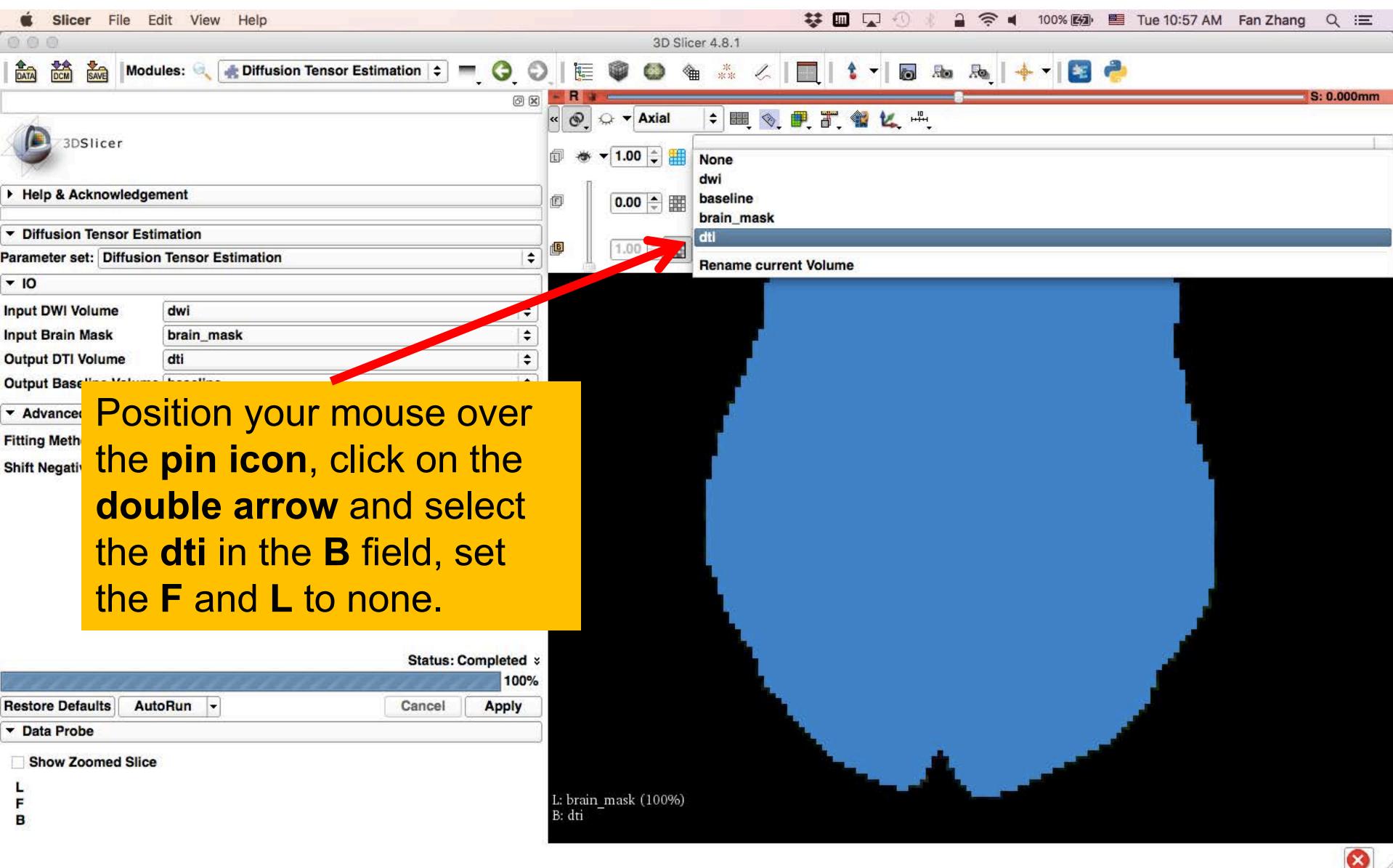


# Estimating the tensor

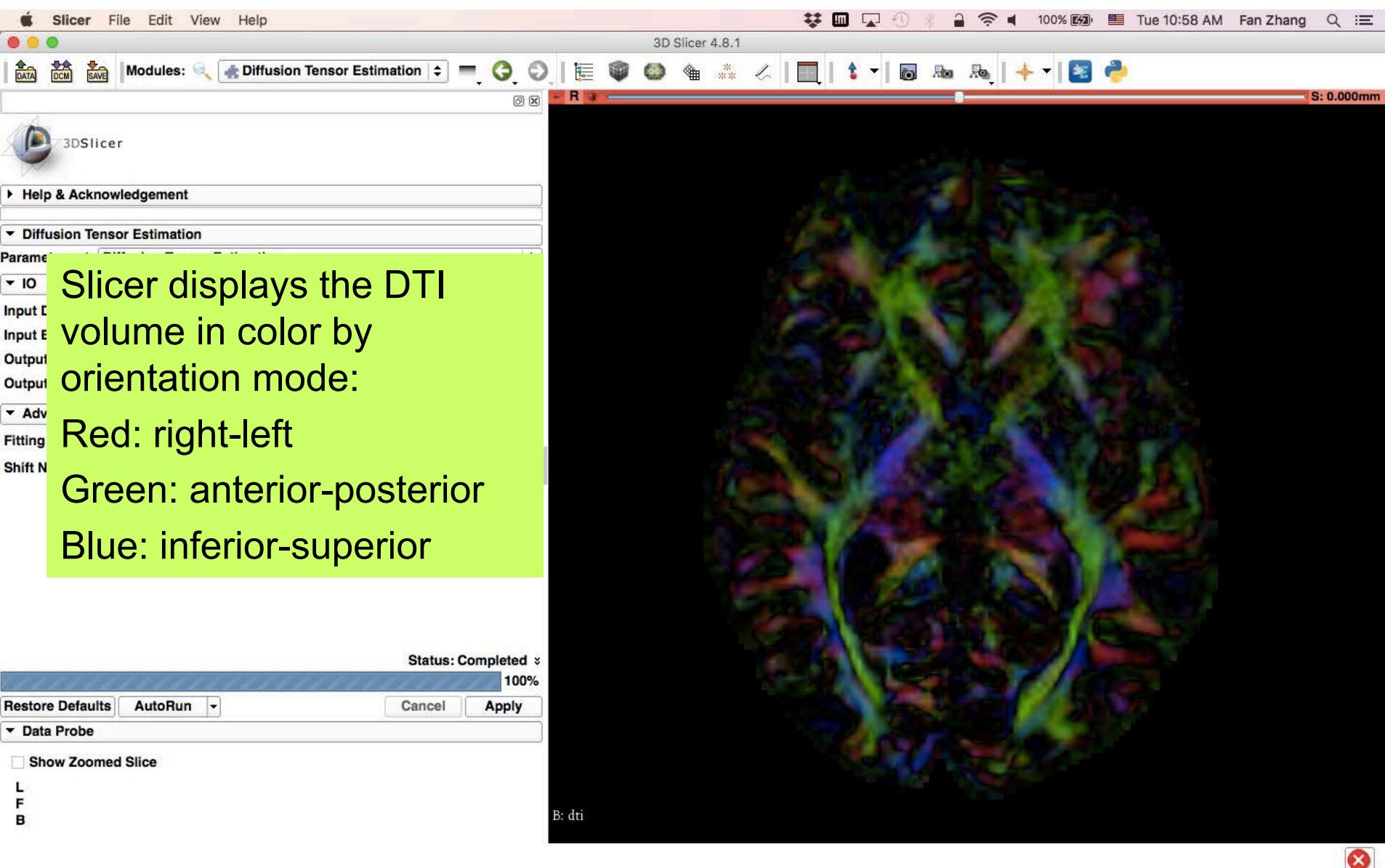


**-Set the Input DWI volume to 'dwi'**  
**-Set the Input Brain Mask to 'brain\_mask'**  
**- Select Output DTI Volume 'Create DiffusionTensorVolume as ...', and name it 'dti'**  
**-Set Output Baseline Volume to 'baseline'**  
**-Under 'Advanced Settings', set Fitting Methods to 'WLS' (Weighted Least Squares)**  
**-Click on Apply.**

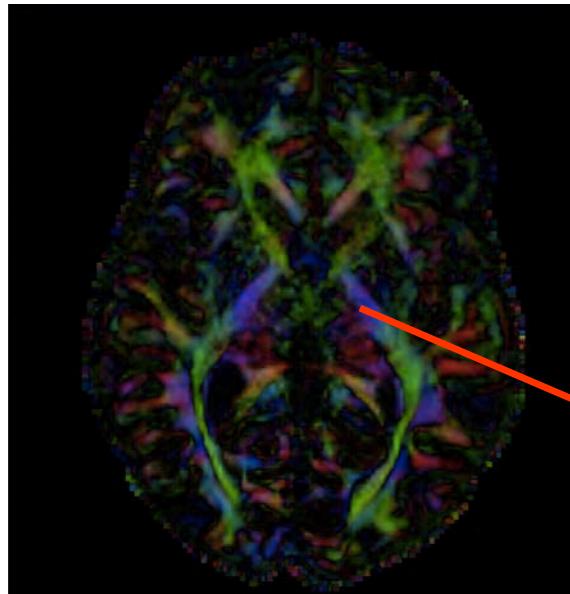
# Estimating the tensor



# Exploring the DWI Dataset



# Diffusion Tensor Data



$$S_i = S_0 e^{-b \hat{g}^T \underline{D} \hat{g}_i}$$

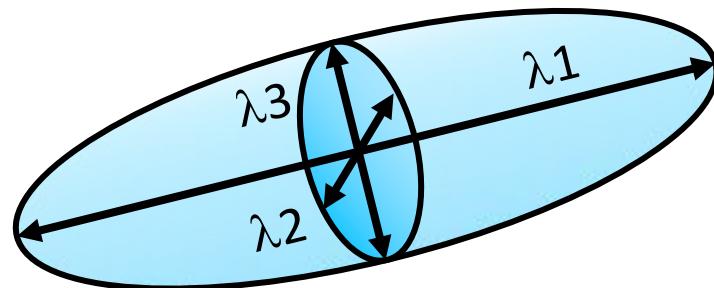
Stejskal-Tanner equation (1965)

$$\underline{D} = \begin{bmatrix} D_{xx} & D_{xy} & D_{xz} \\ D_{yx} & D_{yy} & D_{yz} \\ D_{zx} & D_{zy} & D_{zz} \end{bmatrix}$$

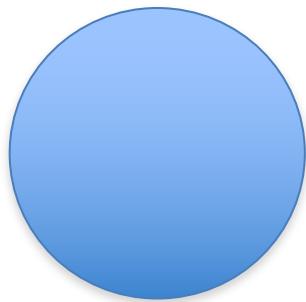
The diffusion tensor  $\underline{D}$  in the voxel (I,J,K) is a 3x3 symmetric matrix.

# Diffusion Tensor

- The diffusion tensor  $\mathbf{D}$  in each voxel can be visualized as a diffusion ellipsoid, with the eigenvectors indicating the directions of the principal axes, and the ellipsoidal proportional to the square root of the eigenvalues defining the
- Scalar maps can be derived from the rotationally invariant eigenvalues  $\lambda_1, \lambda_2, \lambda_3$  to characterize the size and shape of the diffusion tensor.

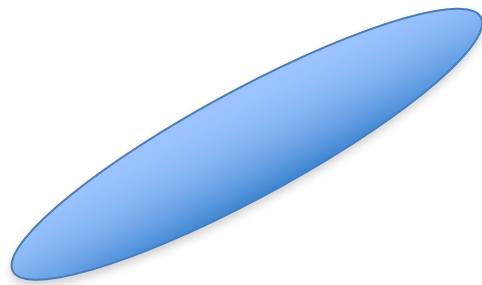


# Diffusion Tensor Shape



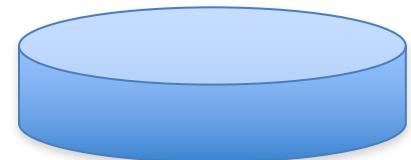
$$\lambda_1 = \lambda_2 = \lambda_3$$

Isotropic media  
(Cerebrospinal  
Fluid, gray matter)



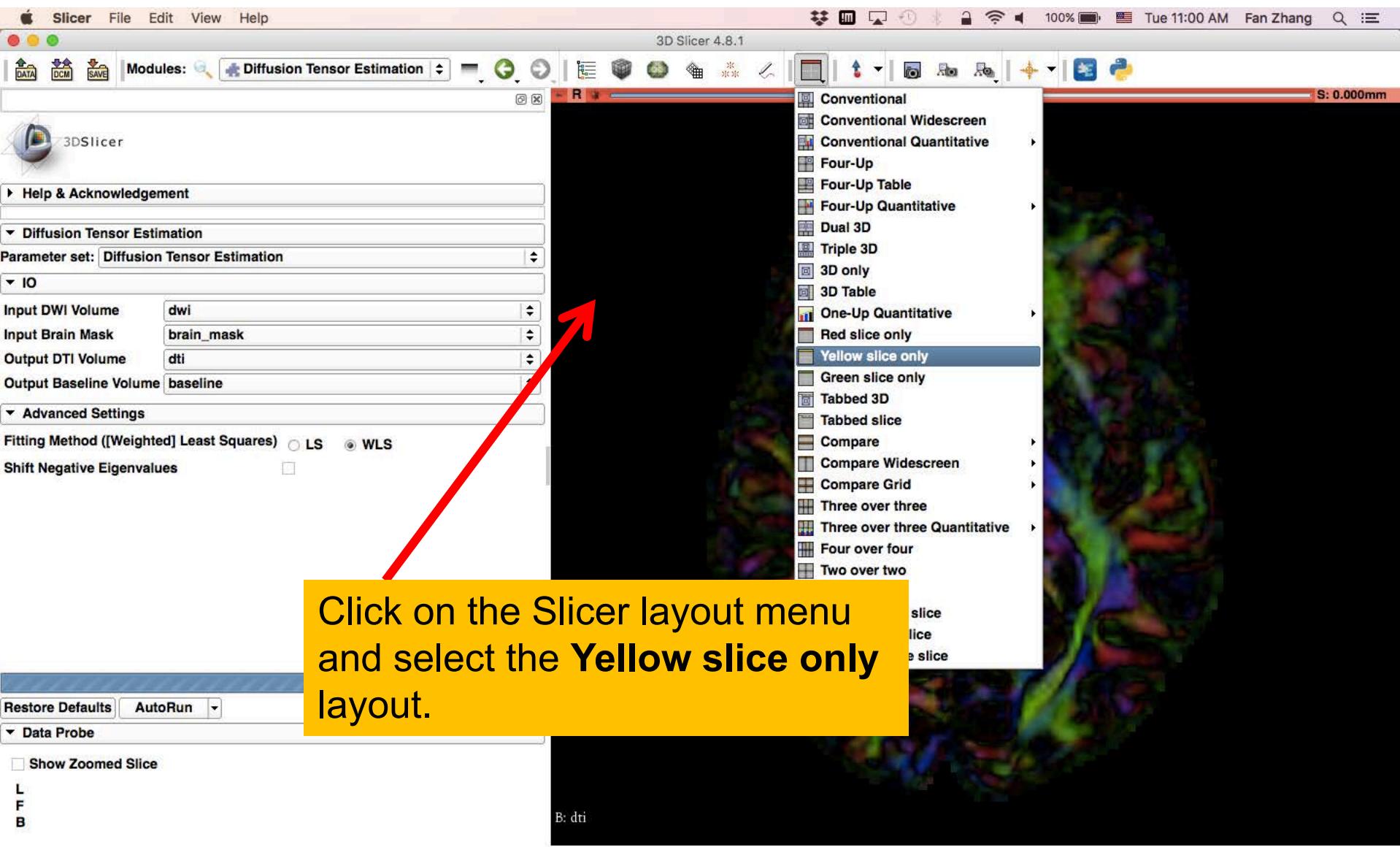
$$\lambda_1 \gg \lambda_2, \lambda_3$$

Anisotropic media  
(white matter)



$$\lambda_1 \sim \lambda_2 \gg \lambda_3$$

# Exploring the DWI Dataset



# Corpus Callosum

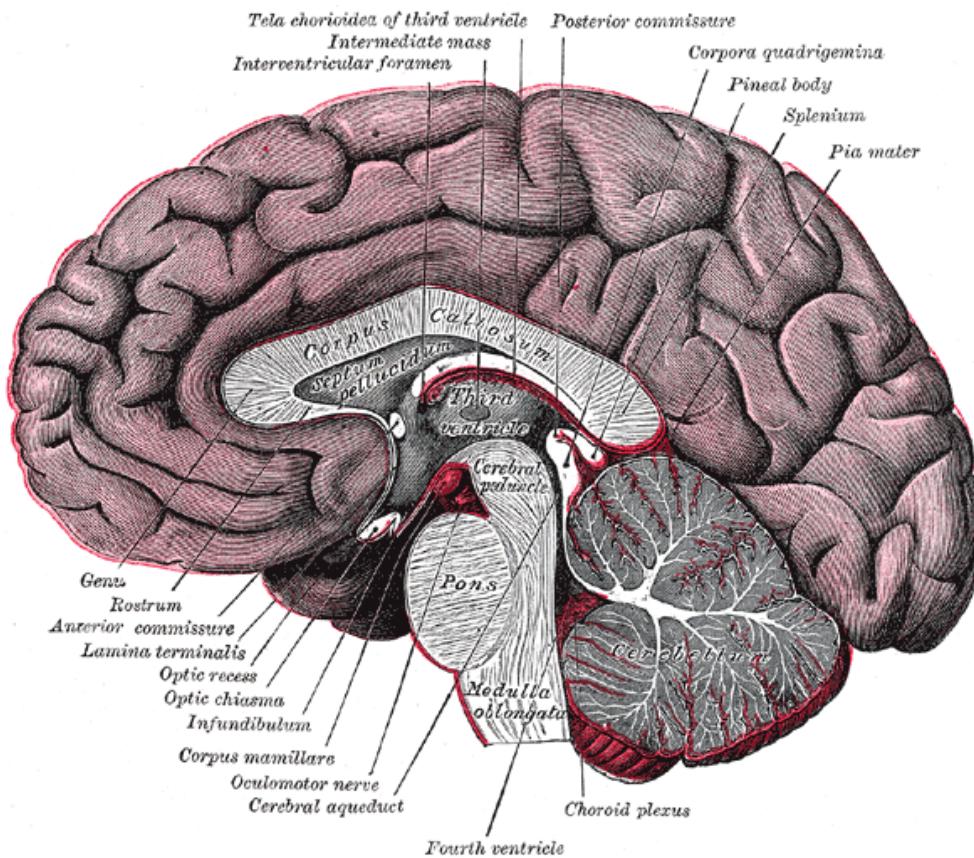
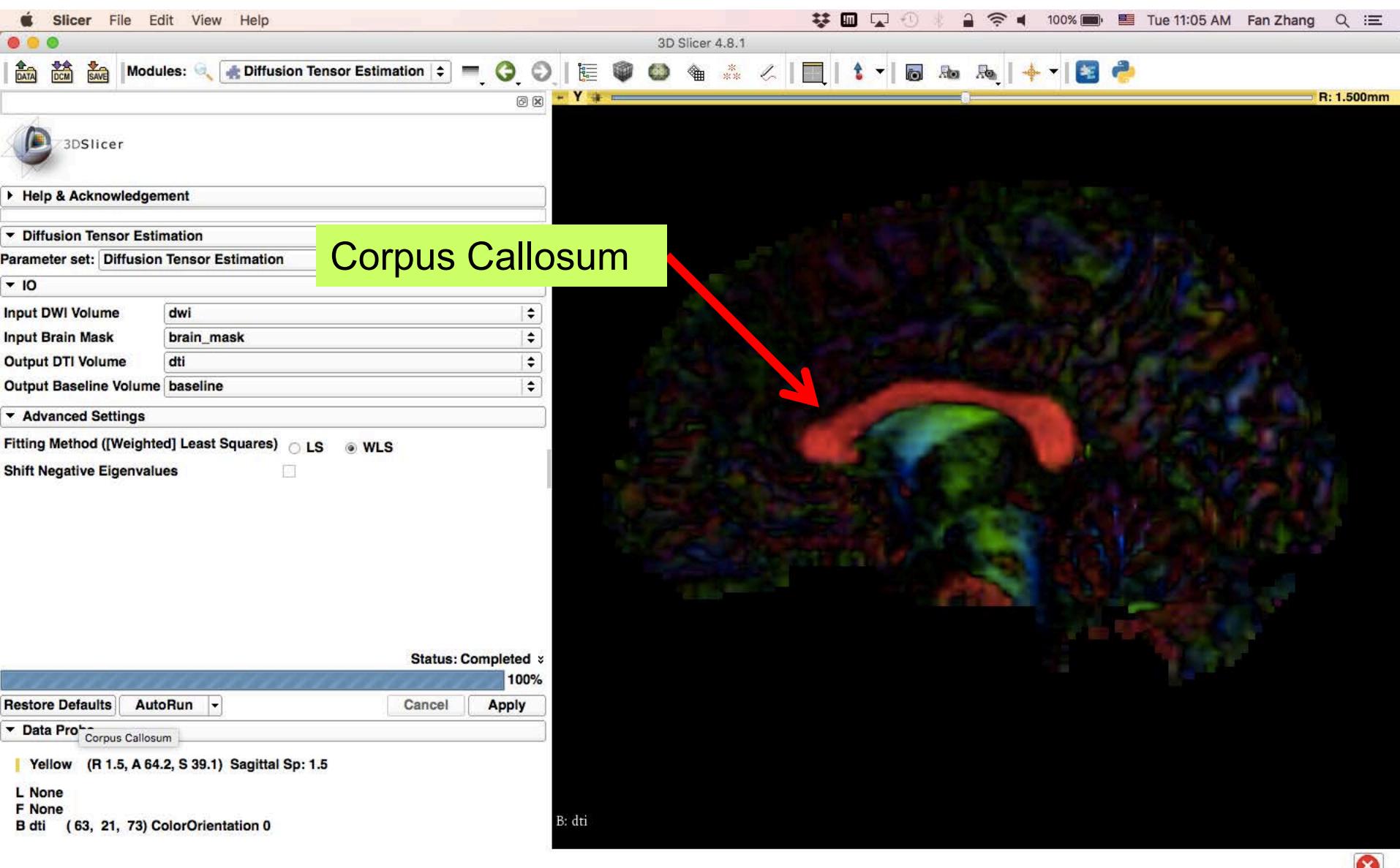


Image from Gray's Anatomy

The corpus callosum is a broad thick bundle of dense myelinated fibers that connect the left and right hemisphere. It is the largest white matter structure in the brain

# Corpus Callosum

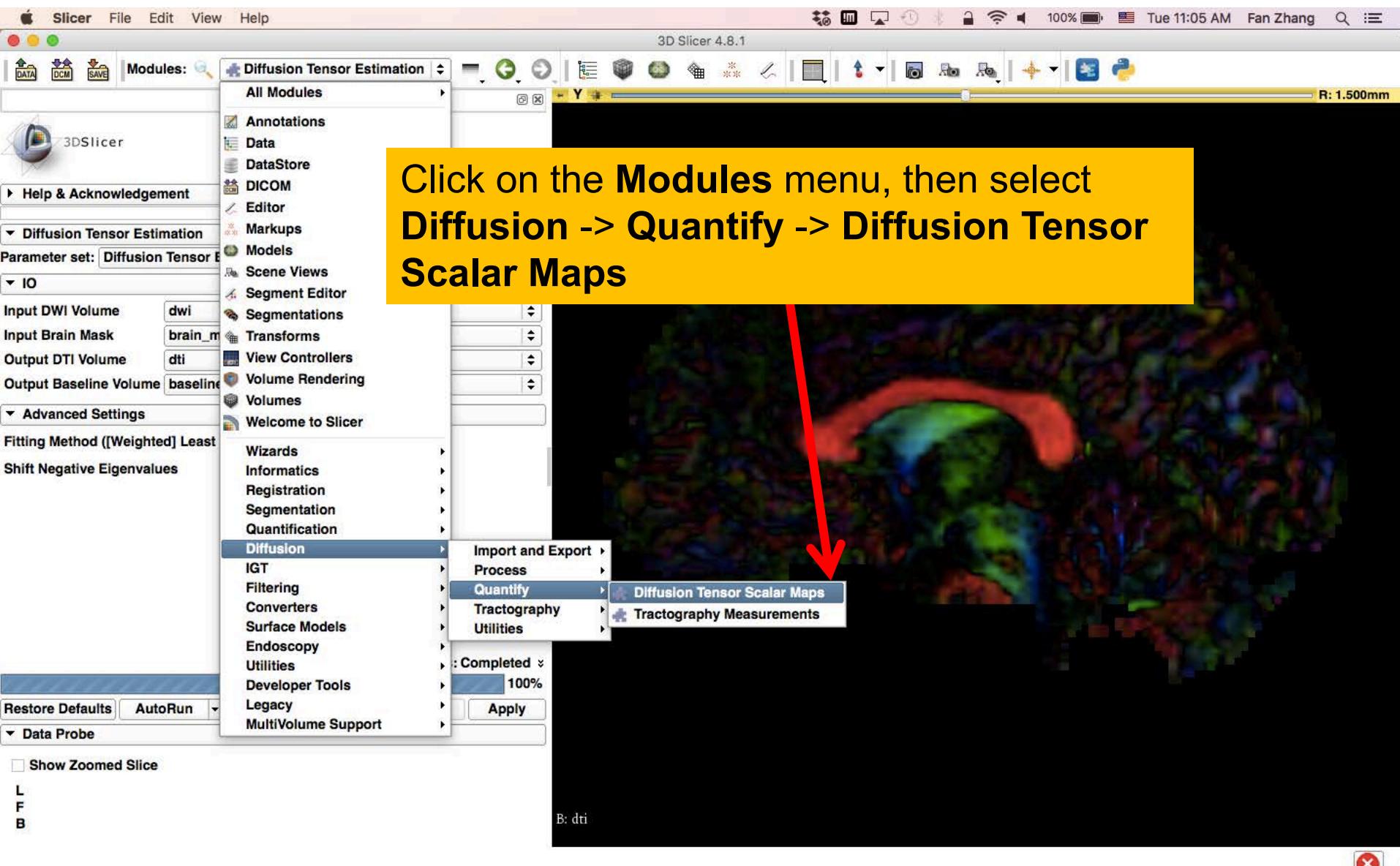


# Characterizing the Size of the tensor: Trace

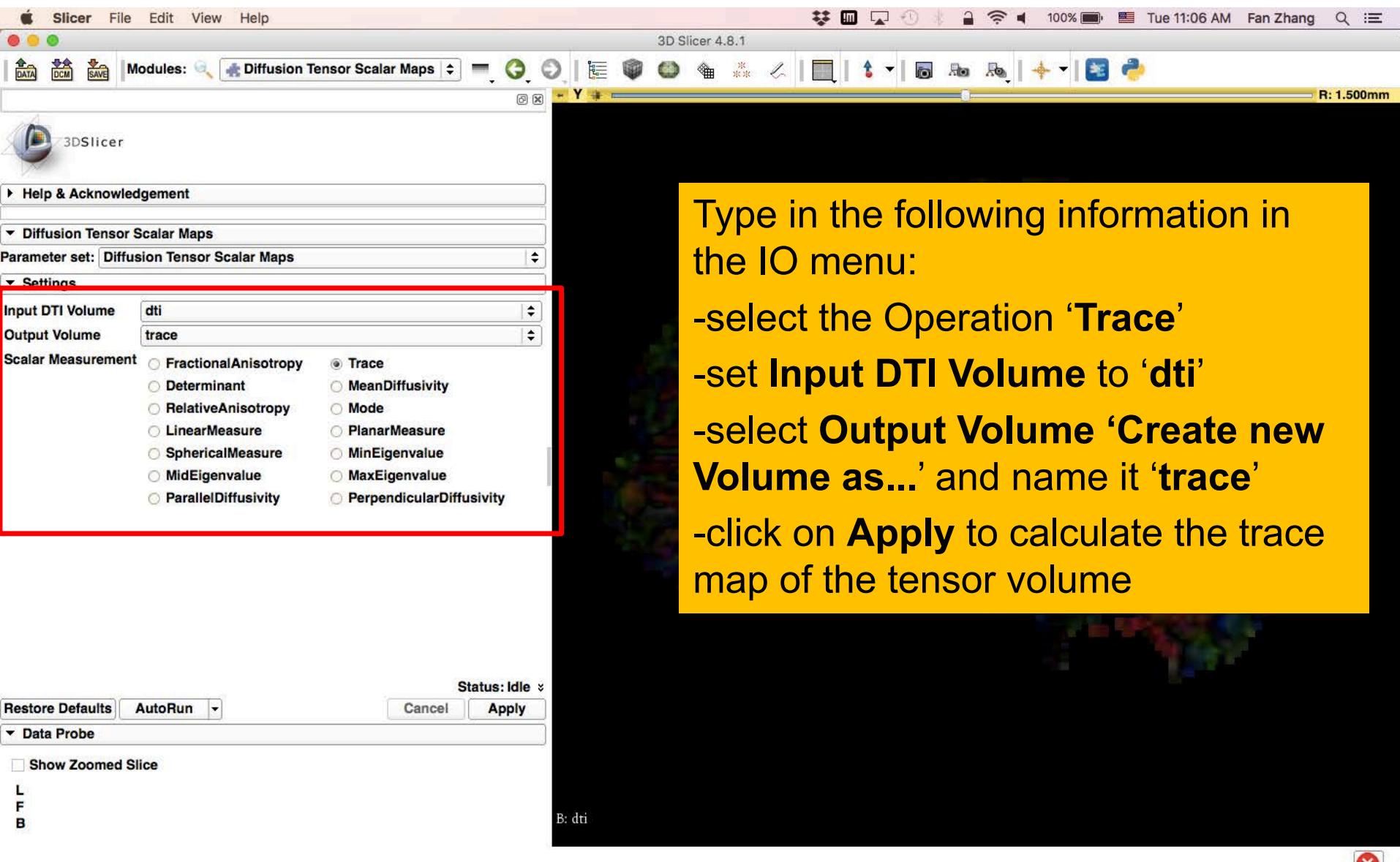
$$\text{Trace}(D) = \lambda_1 + \lambda_2 + \lambda_3$$

- $\text{Trace}(D)$  is intrinsic to the tissue and is independent of fiber orientation, and diffusion sensitizing gradient directions
- $\text{Trace}(D)$  is a clinically relevant parameter for monitoring stroke and neurological condition (degree of structural coherence in tissue)
- $\text{Trace}(D)$  is useful to characterize the size of the diffusion ellipsoid

# Trace



# Trace

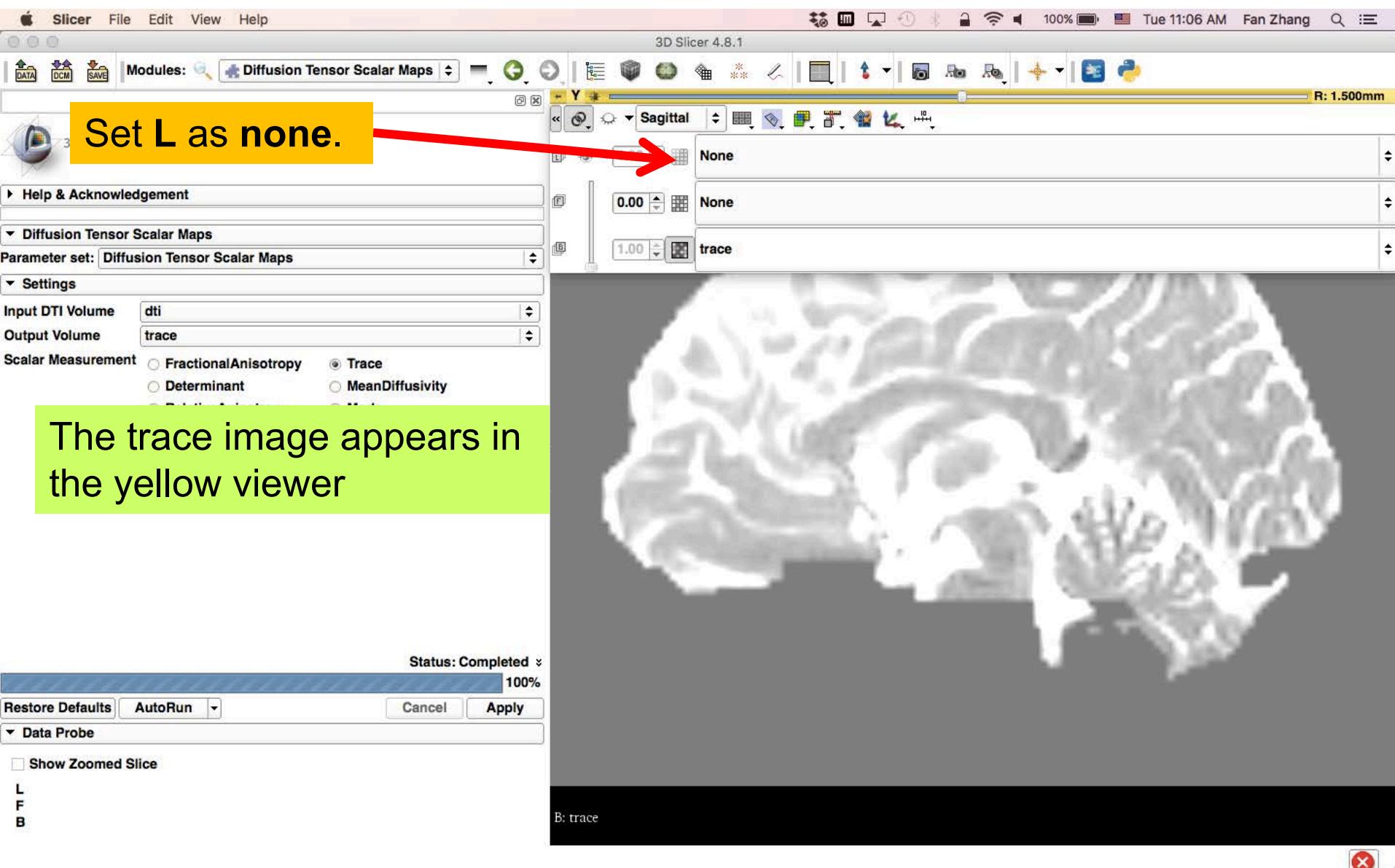


The screenshot shows the 3D Slicer interface with the 'Diffusion Tensor Scalar Maps' module selected. A red box highlights the 'Scalar Measurement' section, which includes fields for 'Input DTI Volume' (set to 'dti') and 'Output Volume' (set to 'trace'). The 'Scalar Measurement' list contains various options, with 'Trace' selected. The main workspace shows a 3D volume rendering of a brain with colored fibers.

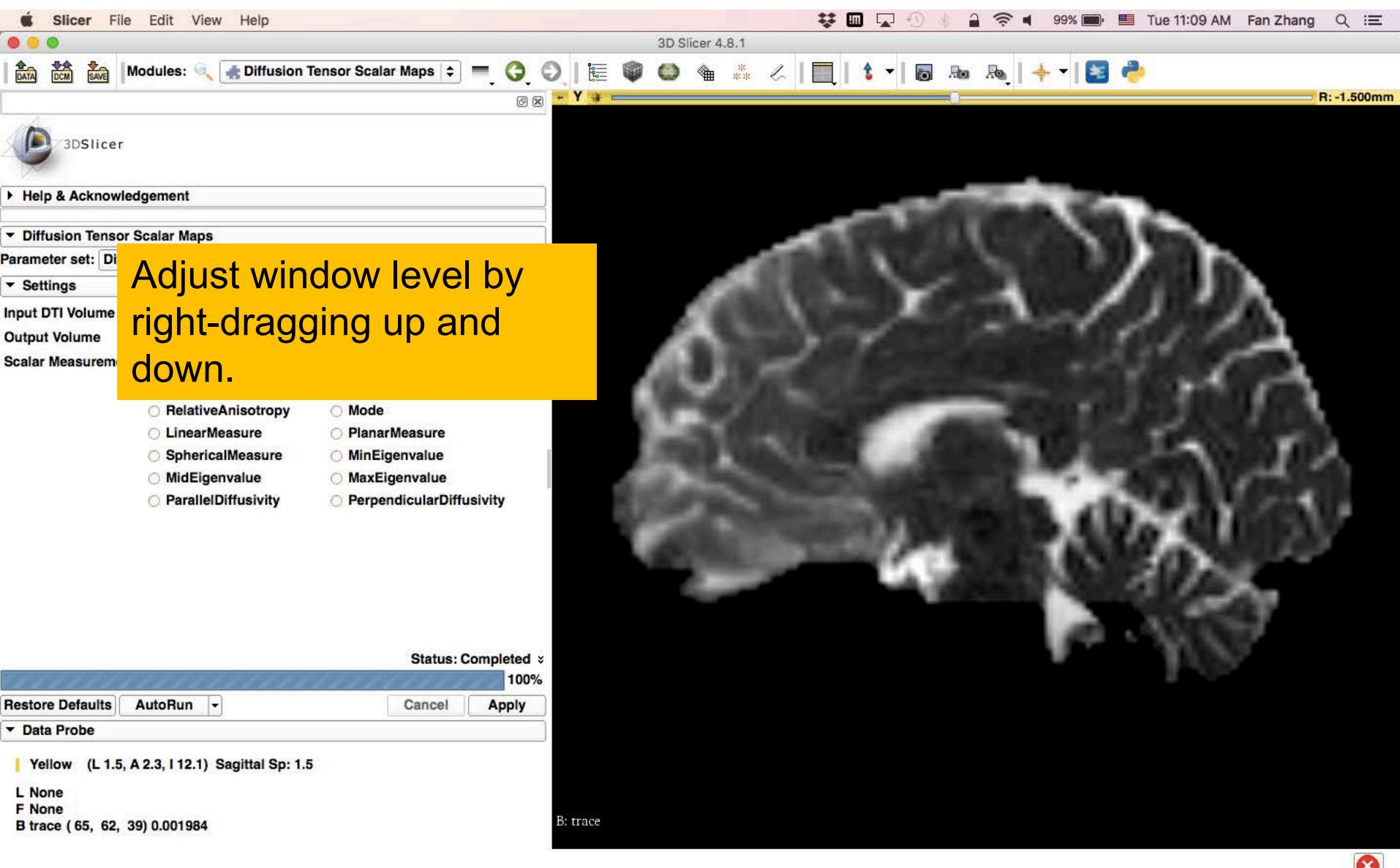
Type in the following information in the IO menu:

- select the Operation '**Trace**'
- set **Input DTI Volume** to '**dti**'
- select **Output Volume 'Create new Volume as...'** and name it '**trace**'
- click on **Apply** to calculate the trace map of the tensor volume

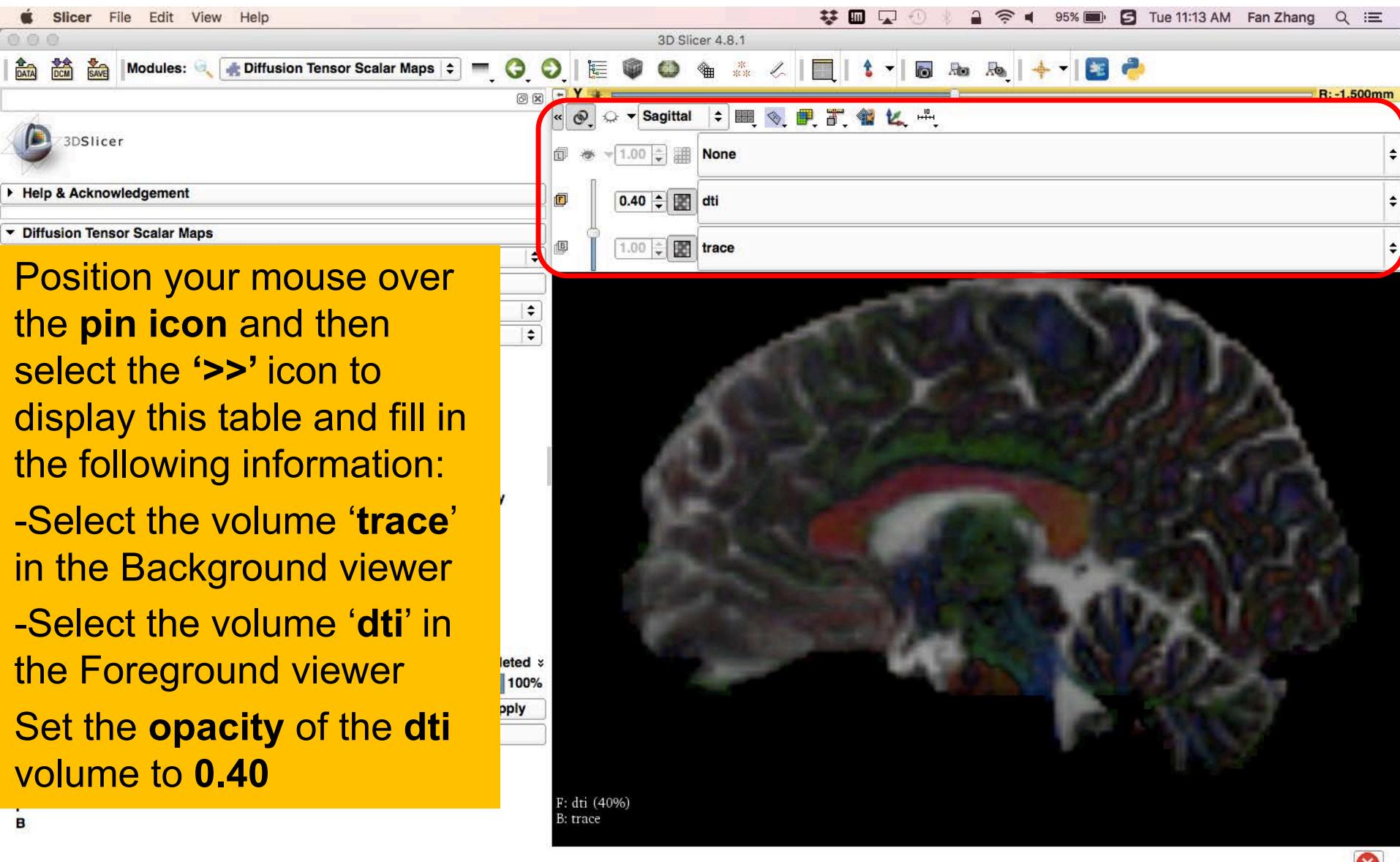
# Trace



# Trace



# Trace



# Trace

Position your mouse within the region of the Corpus Callosum and observe the trace values in the Data Probe

Parameter set: Diffusion Tensor Scalar Maps

Input DTI Volume: dti

Output Volume: trace

Scalar Measurement:  Trace

FractionalAnisotropy

Determinant

RelativeAnisotropy

LinearMeasure

SphericalMeasure

MidEigenvalue

ParallelDiffusivity

MeanDiffusivity

Mode

PlanarMeasure

MinEigenvalue

MaxEigenvalue

PerpendicularDiffusivity

Status: Completed 100%

Restore Defaults AutoRun Cancel Apply

Trace

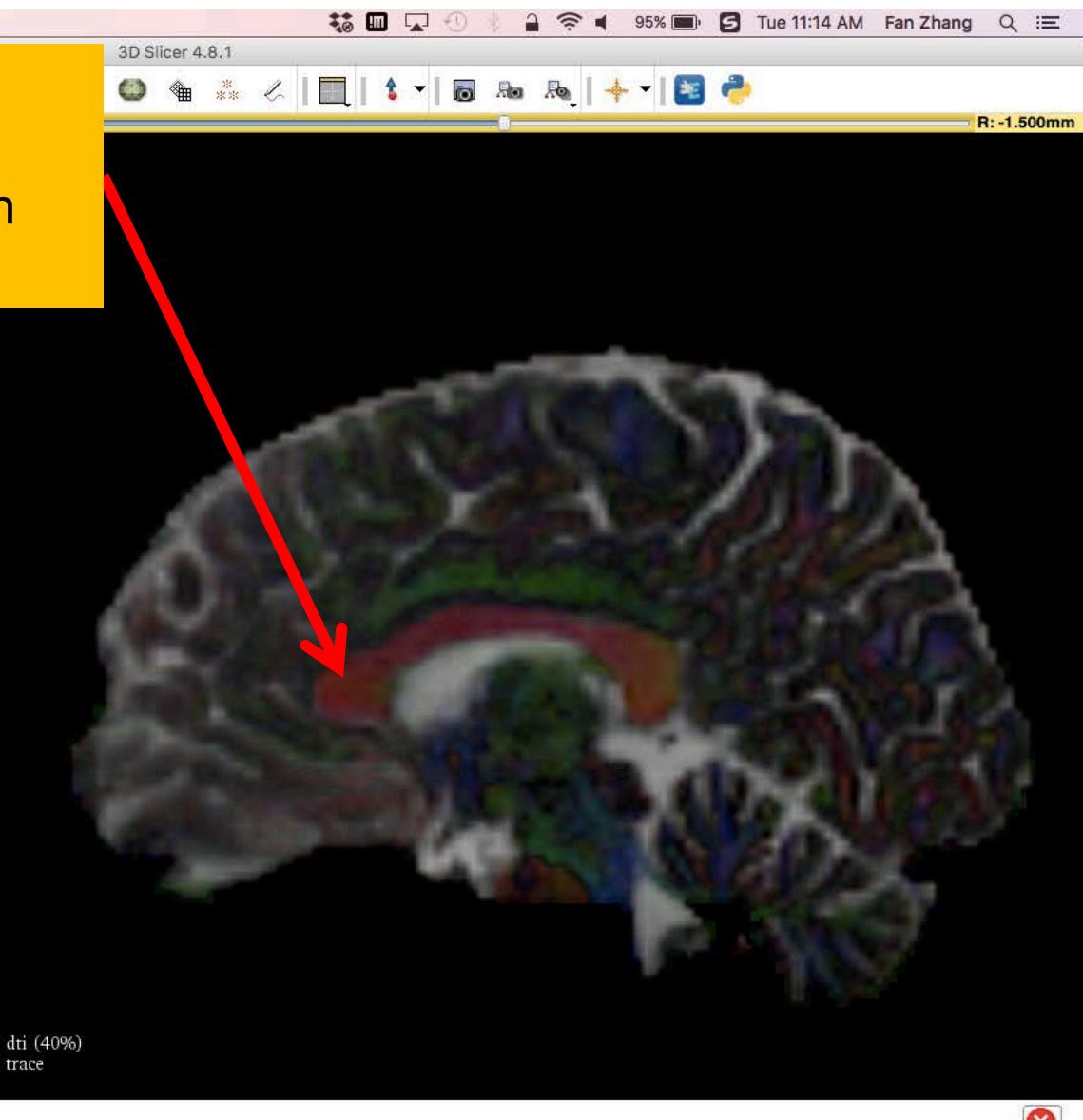
Data Probe

Yellow (L 1.5, A 33.9, S 8.4) Sagittal Sp: 1.5

L None

F dti (65, 41, 53) ColorOrientation 0

B trace (65, 41, 53) 0.001909



A red arrow points from the text "Position your mouse within the region of the Corpus Callosum and observe the trace values in the Data Probe" to the corpus callosum region of the brain scan. The brain scan is a multi-colored 3D rendering of the brain's internal structures. The data probe status bar at the bottom left shows "Yellow (L 1.5, A 33.9, S 8.4) Sagittal Sp: 1.5" and "B trace (65, 41, 53) 0.001909".

# Trace

Note how the Trace values are fairly uniform in both white and gray matter, even if the tissues are different in structure.

Input DTI Volume: dti  
Output Volume: trace

Scalar Measurement:  Trace  
 FractionalAnisotropy  
 Determinant  
 RelativeAnisotropy  
 LinearMeasure  
 SphericalMeasure  
 MidEigenvalue  
 ParallelDiffusivity  
 Trace  
 MeanDiffusivity  
 Mode  
 PlanarMeasure  
 MinEigenvalue  
 MaxEigenvalue  
 PerpendicularDiffusivity

Status: Completed 100%

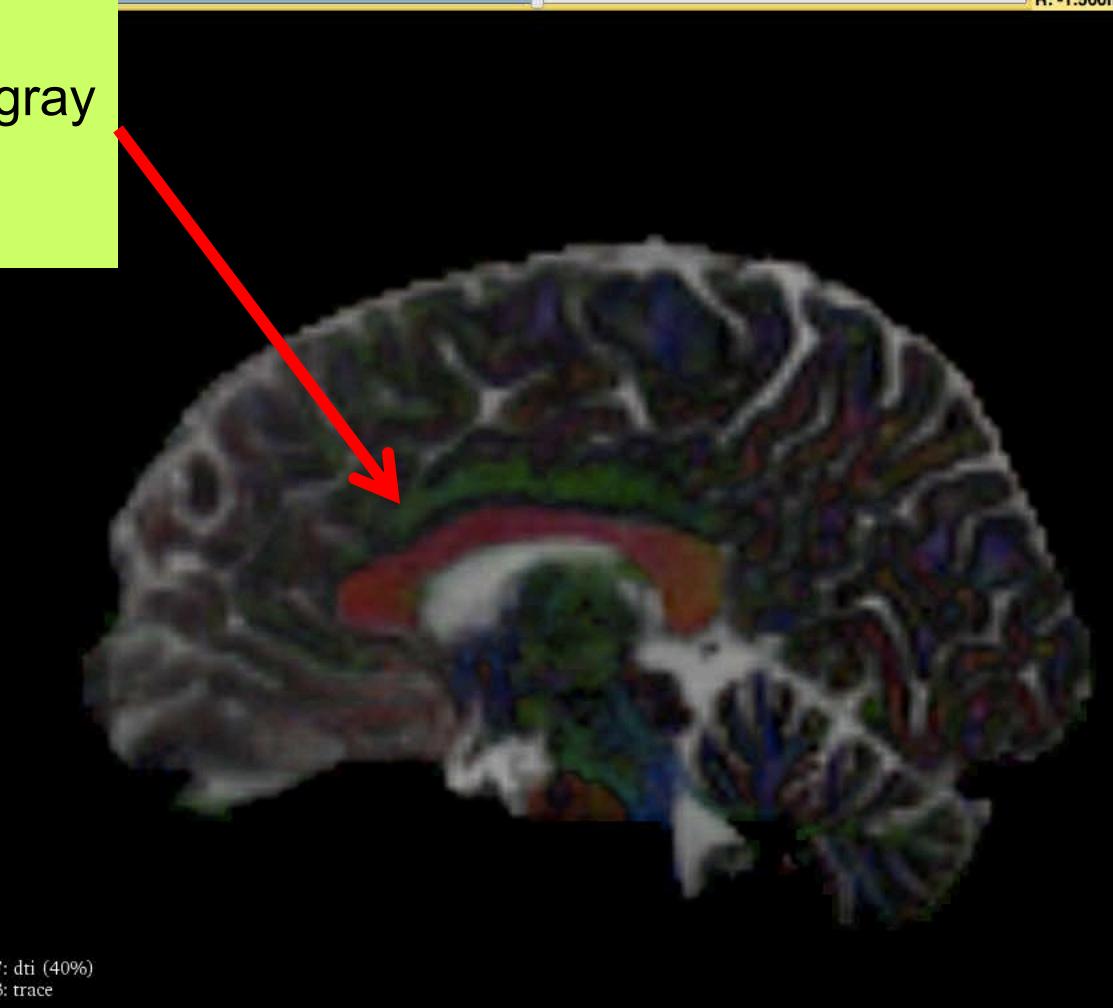
Restore Defaults AutoRun Cancel Apply

Yellow (L 1.5, A 31.3, S 19.1) Sagittal Sp: 1.5

L None  
F dti (65, 43, 60) ColorOrientation 0  
B trace (65, 43, 60) 0.002775

3D Slicer 4.8.1

R: -1.500mm

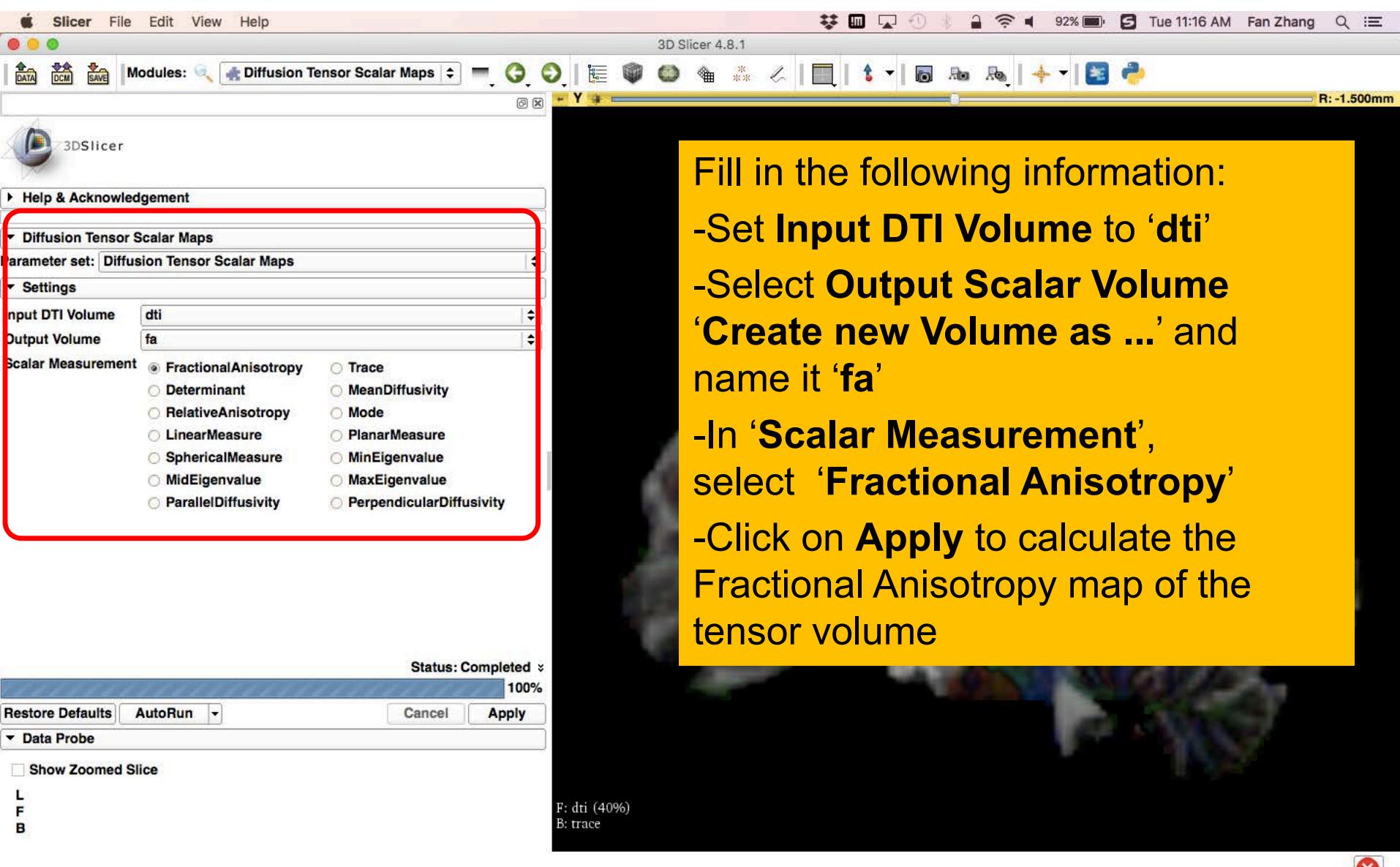


# Scalar Maps: Fractional Anisotropy

$$FA(D) = \frac{\sqrt{(\lambda_1 - \lambda_2)^2 + (\lambda_1 - \lambda_3)^2 + (\lambda_2 - \lambda_3)^2}}{\sqrt{2} \sqrt{\lambda_1^2 + \lambda_2^2 + \lambda_3^2}}$$

- FA(D) is intrinsic to the tissue and is independent of fiber orientation, and diffusion sensitizing gradient directions
- FA(D) is useful to characterize the shape (degree of 'out-of-roundness') of the diffusion ellipsoid
- Low FA:  → High FA: 

# Fractional Anisotropy



3D Slicer 4.8.1

Fill in the following information:

- Set **Input DTI Volume** to 'dti'
- Select **Output Scalar Volume** 'Create new Volume as ...' and name it 'fa'
- In '**Scalar Measurement**', select '**Fractional Anisotropy**'
- Click on **Apply** to calculate the Fractional Anisotropy map of the tensor volume

3DSlicer

Help & Acknowledgement

Diffusion Tensor Scalar Maps

Parameter set: Diffusion Tensor Scalar Maps

Settings

Input DTI Volume: dti

Output Volume: fa

Scalar Measurement:

- FractionalAnisotropy
- Trace
- Determinant
- MeanDiffusivity
- RelativeAnisotropy
- Mode
- LinearMeasure
- PlanarMeasure
- SphericalMeasure
- MinEigenvalue
- MaxEigenvalue
- MidEigenvalue
- ParallelDiffusivity
- PerpendicularDiffusivity

Status: Completed 100%

Restore Defaults AutoRun Cancel Apply

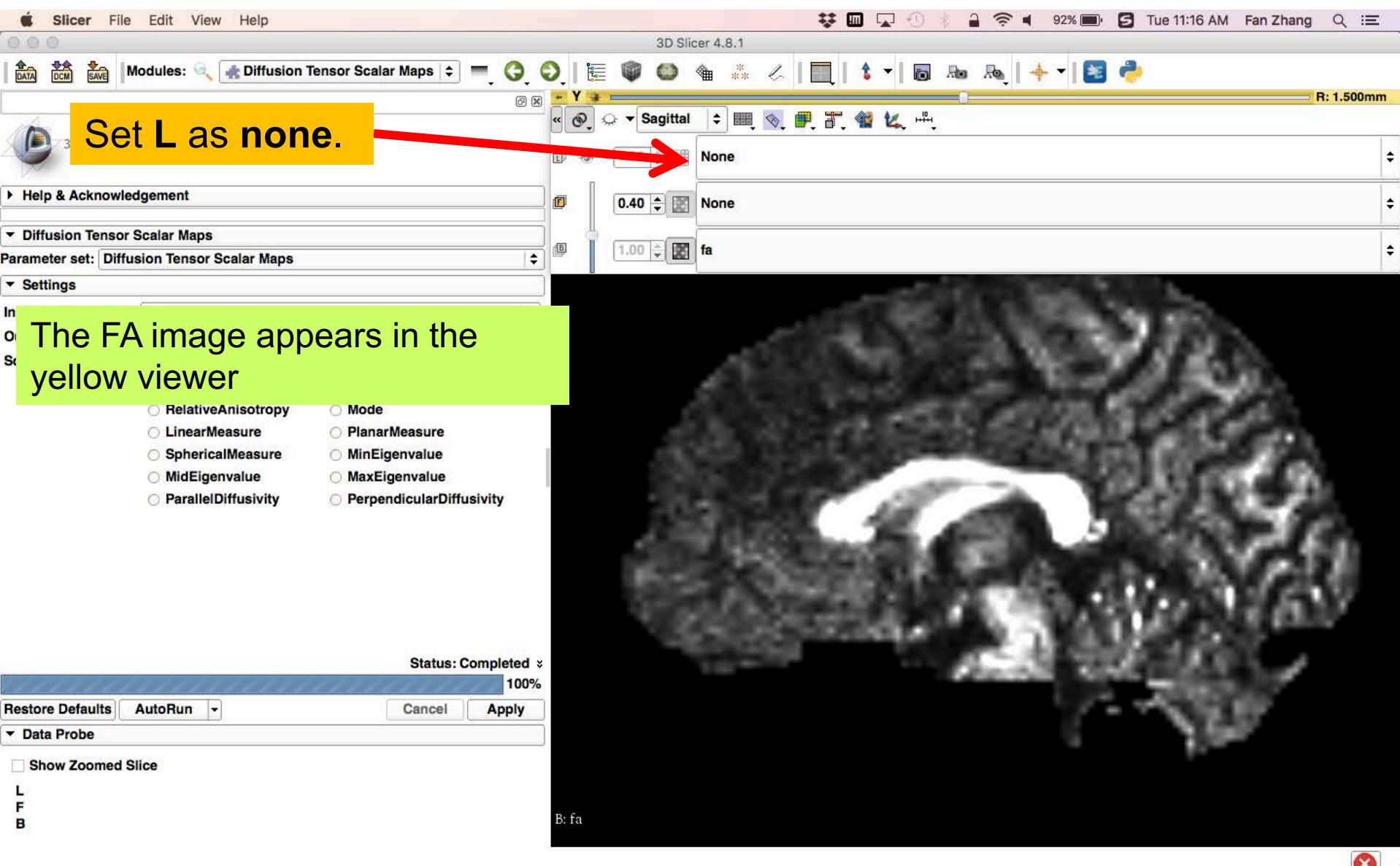
Data Probe

Show Zoomed Slice

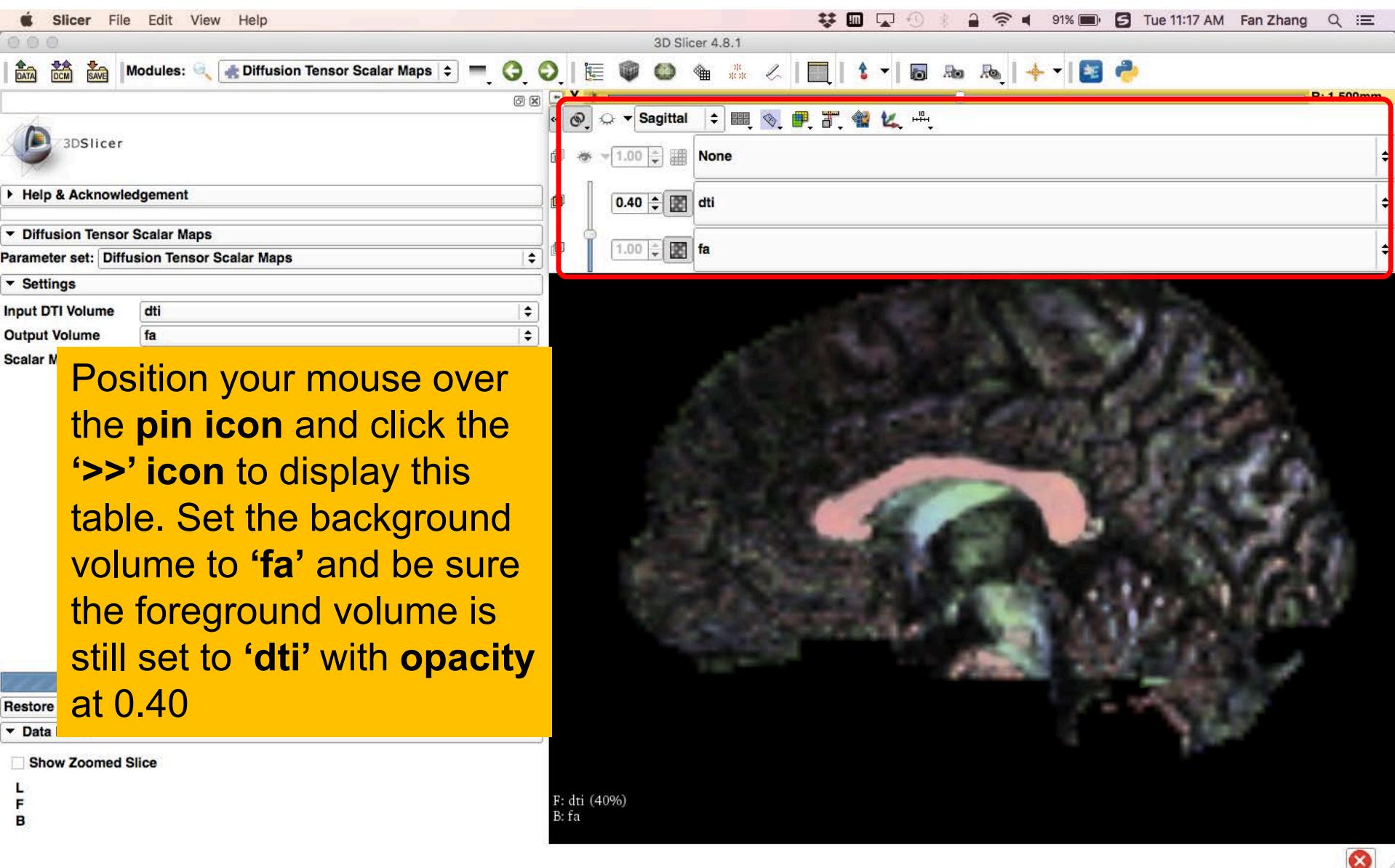
L F B

F: dti (40%)  
B: trace

# Fractional Anisotropy



# Fractional Anisotropy



# Fractional Anisotropy

Explore the FA values in the Corpus Callosum and in adjacent gray matter areas. Note how the FA values are high in the white matter areas, and low in gray matter regions

Output Volume: fa

Scalar Measurement:  FractionalAnisotropy  Trace  
 Determinant  MeanDiffusivity  
 RelativeAnisotropy  Mode  
 LinearMeasure  PlanarMeasure  
 SphericalMeasure  MinEigenvalue  
 MidEigenvalue  MaxEigenvalue  
 ParallelDiffusivity  PerpendicularDiffusivity

Fractional Anisotropy

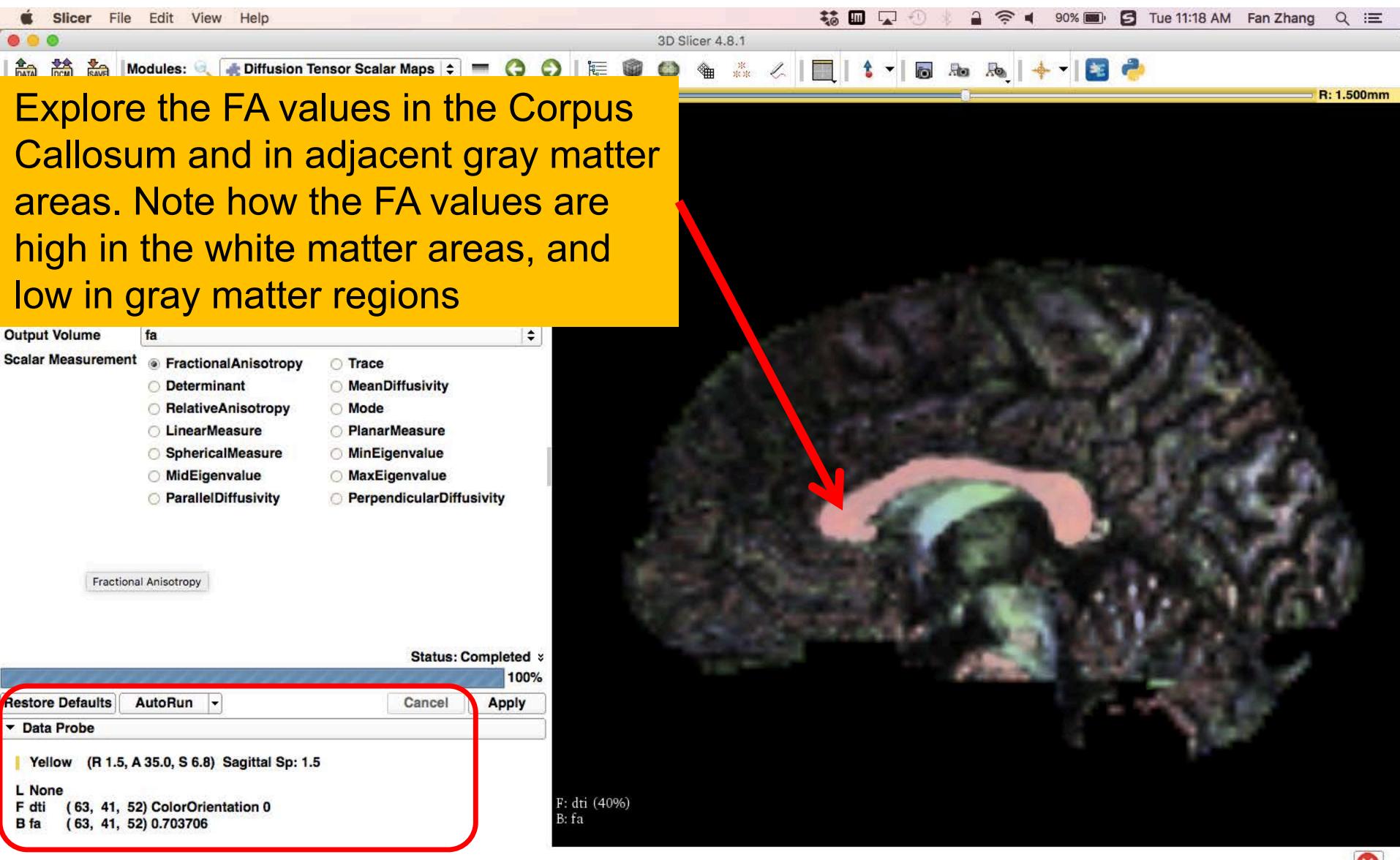
Status: Completed 100%

Restore Defaults AutoRun

▼ Data Probe

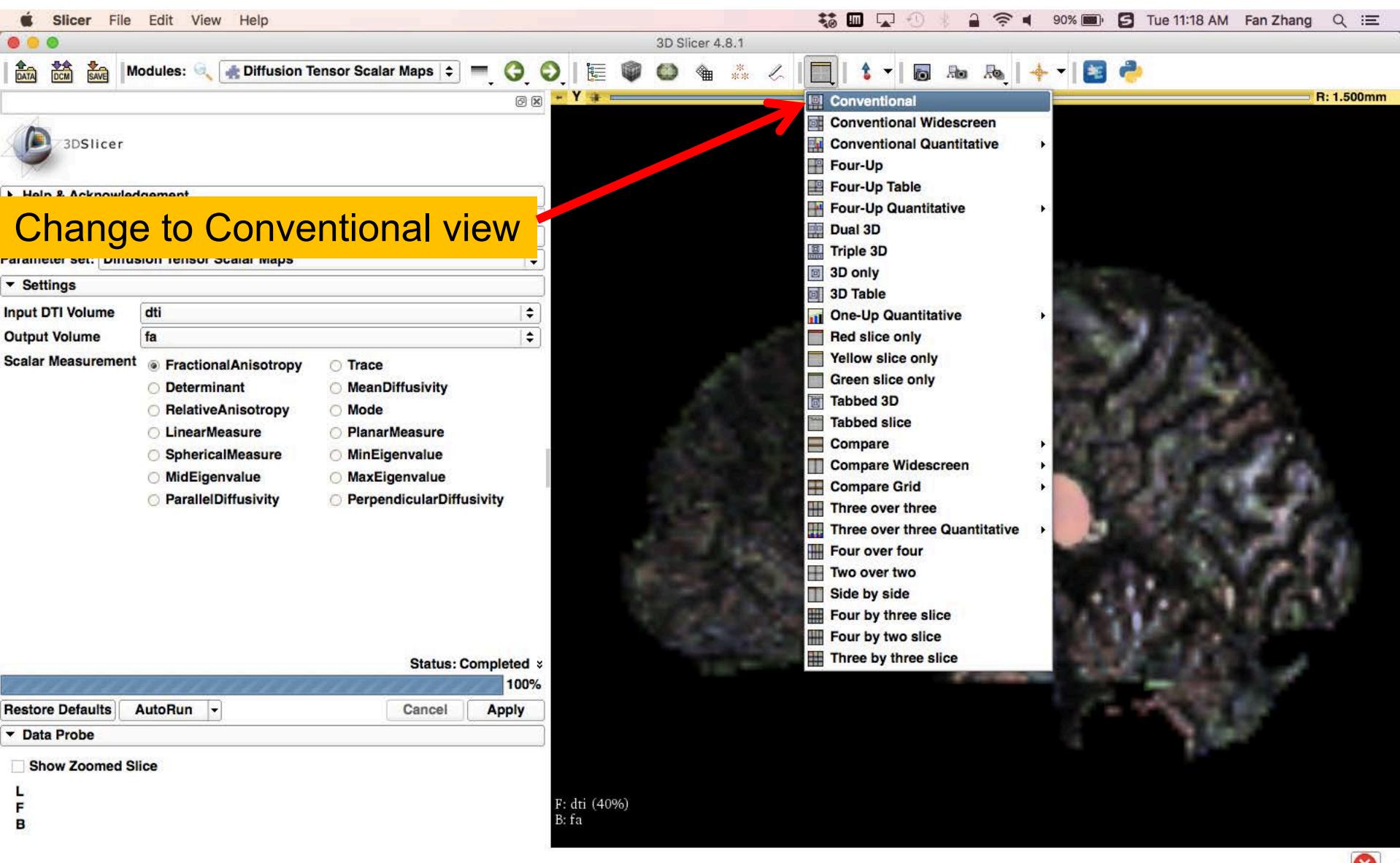
Yellow (R 1.5, A 35.0, S 6.8) Sagittal Sp: 1.5

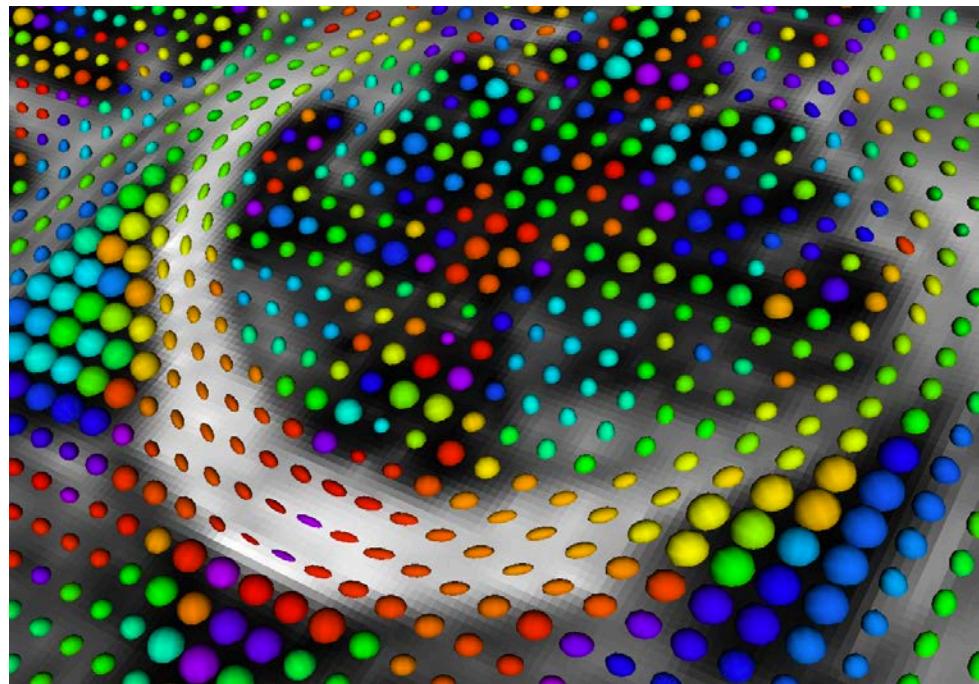
L None  
F dti (63, 41, 52) ColorOrientation 0  
B fa (63, 41, 52) 0.703706



A red arrow points from the text "Explore the FA values in the Corpus Callosum and in adjacent gray matter areas." to the corpus callosum region in the 3D brain scan. The 3D Slicer interface is visible, showing the brain scan in a sagittal plane. The control panel on the left shows the scalar measurement is set to 'fa'. The status bar at the bottom indicates the status is 'Completed' at 100%.

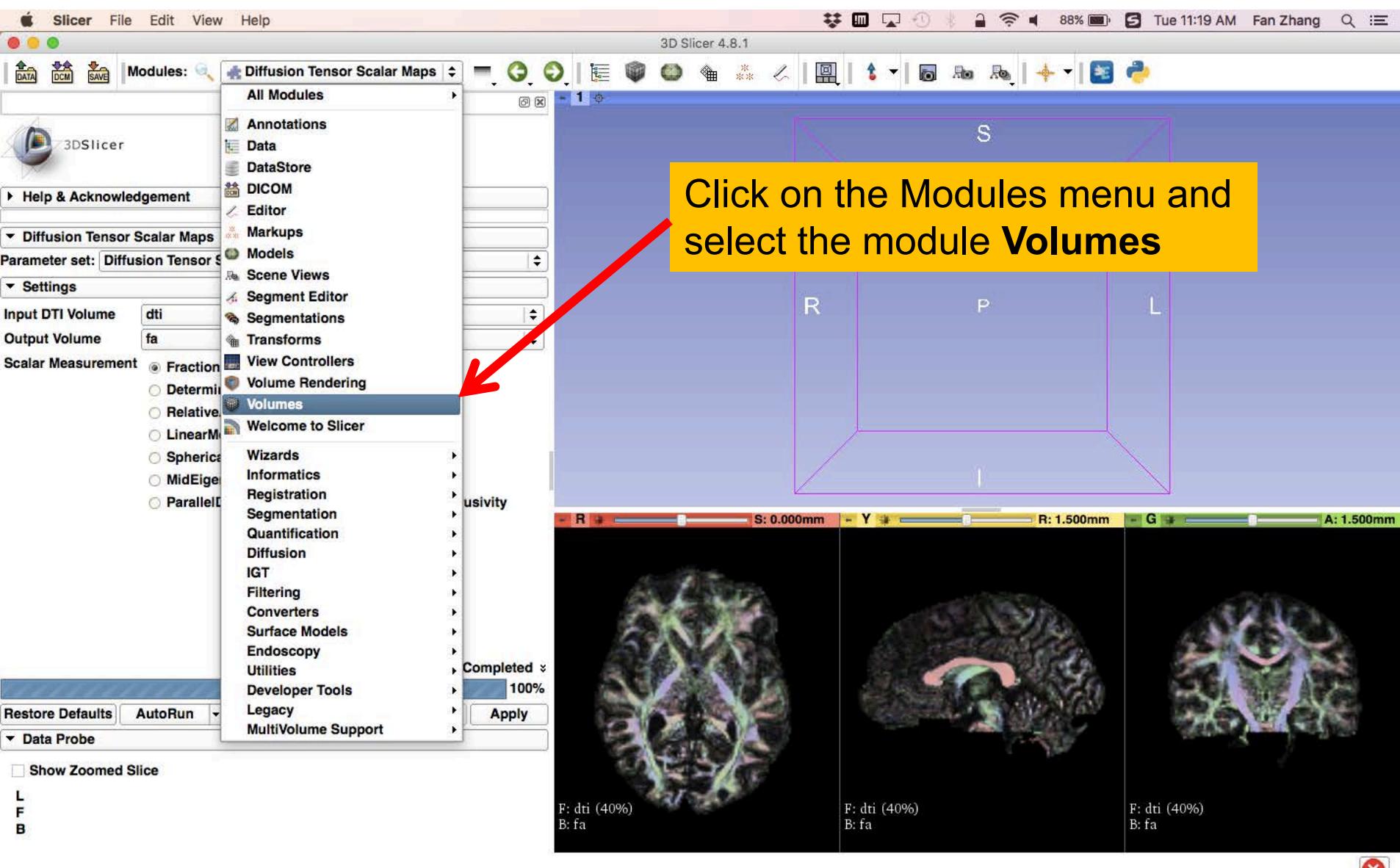
# Fractional Anisotropy



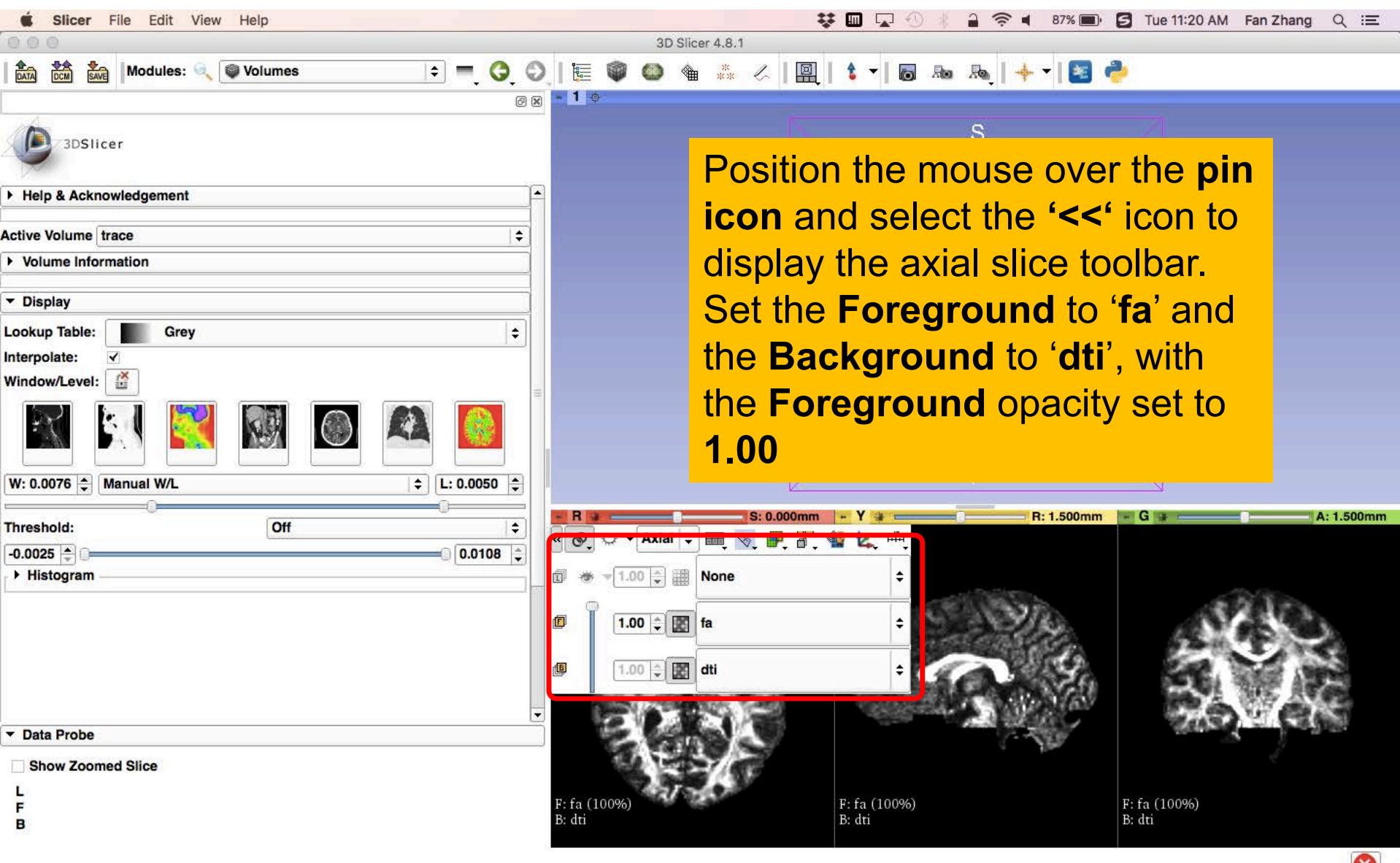


## Part 2: Visualizing the tensor data

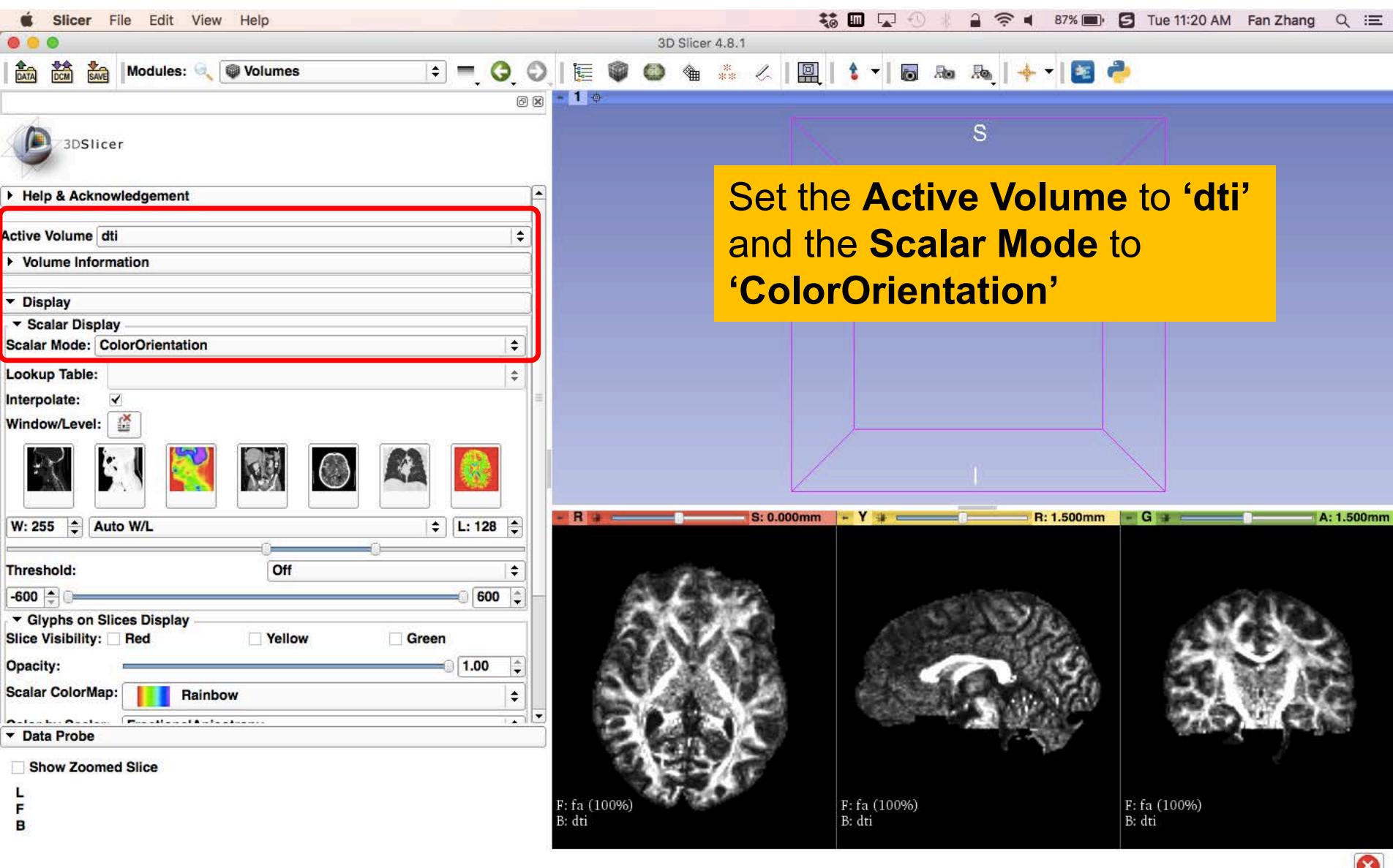
# 3D Visualization: Glyphs



# 3D Visualization: Glyphs



# 3D Visualization: Glyphs



# 3D Visualization: Glyphs

3D Slicer 4.8.1

Scalar Mode: ColorOrientation

Lookup Table:

Interpolate:

Window/Level:

W: 255 Auto W/L L: 128

Threshold: Off

-600 600

**Glyphs on Slices Display**

Slice Visibility:  Red  Yellow  Green

Opacity: 1.00

Scalar ColorMap: FullRainbow

Color by Scalar: ColorOrientation

Scalar Range:

Glyph Type: Ellipsoids

Scale Factor: 50.00

Spacing: 5.00

**Data Probe**

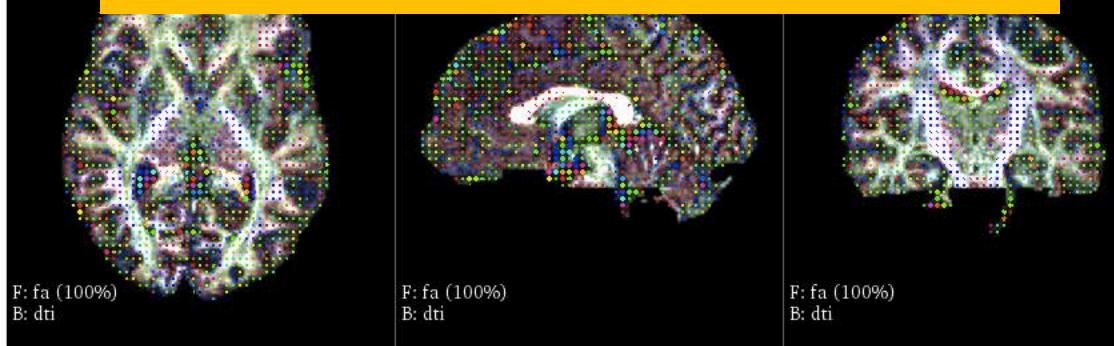
Show Zoomed Slice

L F B

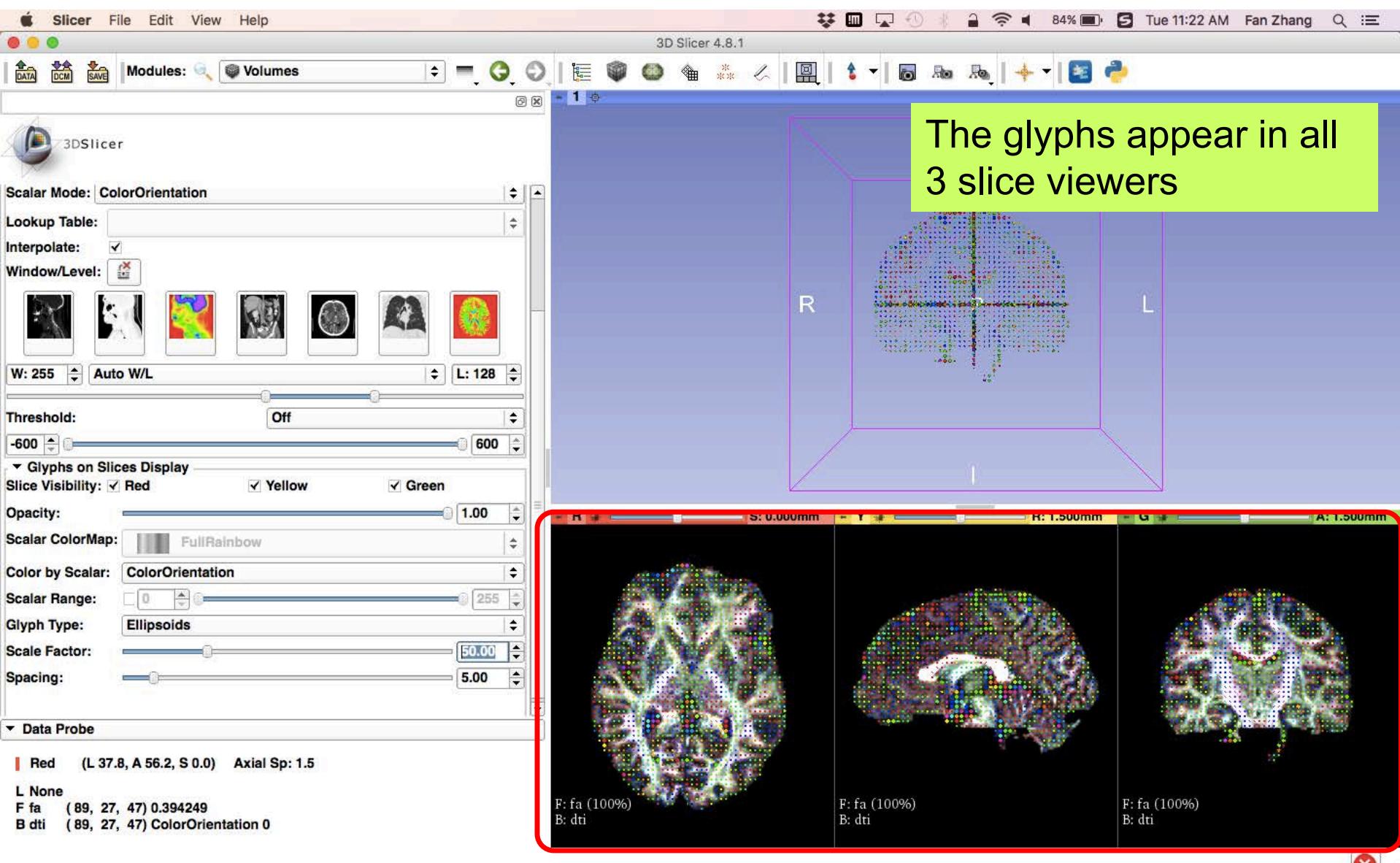
**S**

Scroll down the module panel and in the **Glyphs on Slices Display** section:

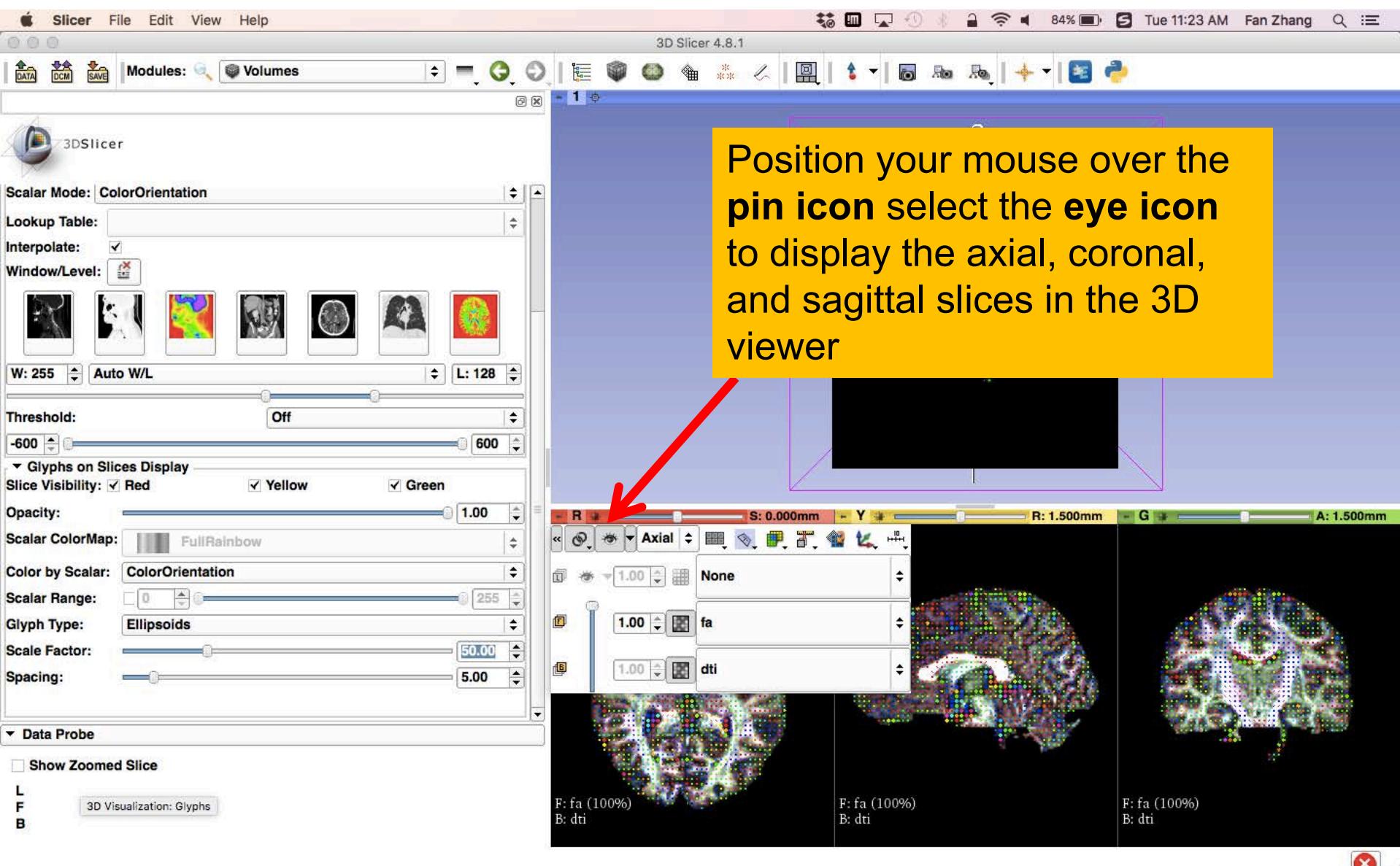
- Check off the option for **Red, Yellow, and Green Slice Visibility**
- Set the **Color by Scalar** parameter to '**ColorOrientation**'
- Set the **Glyph Type** to '**Ellipsoids**'



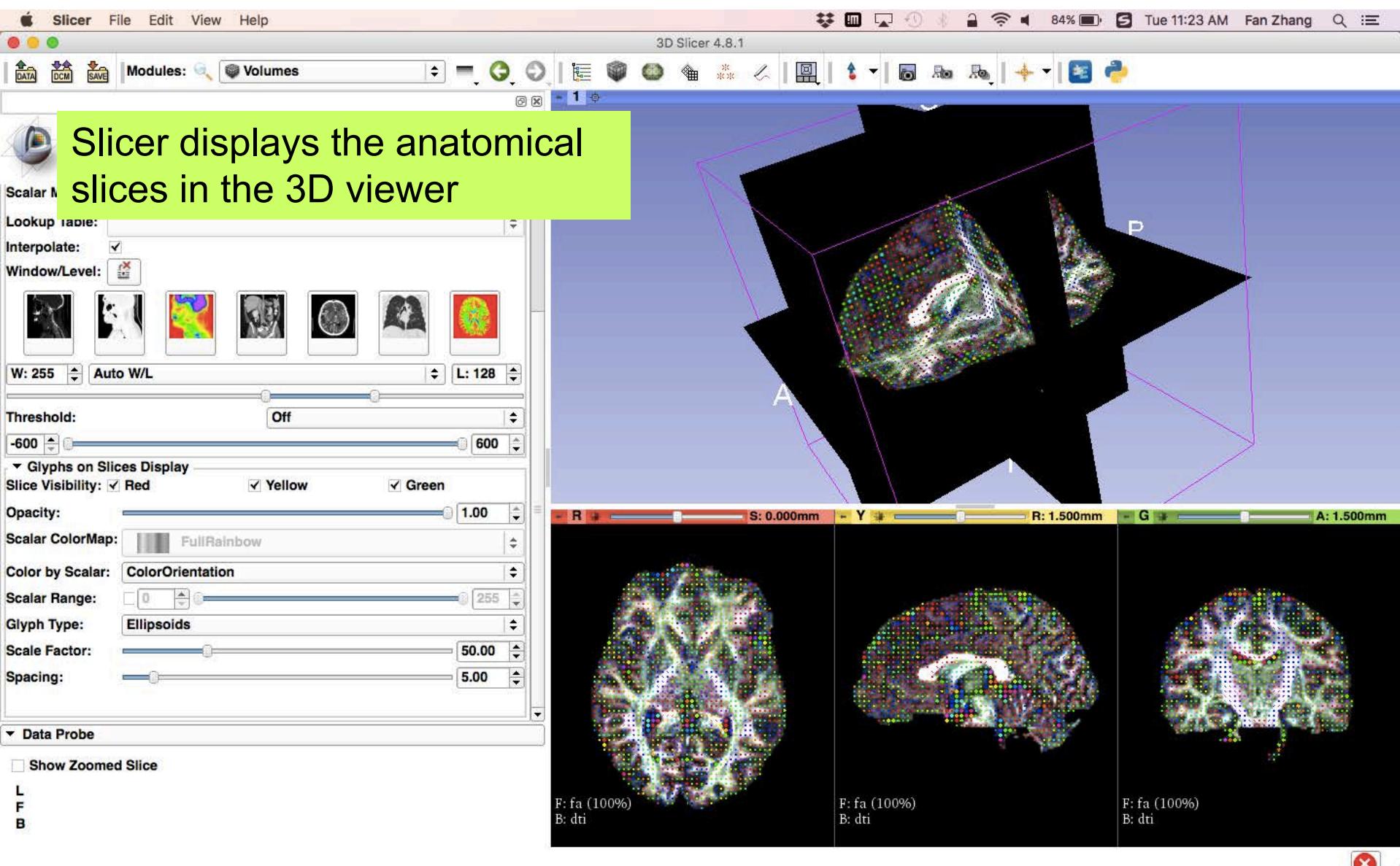
# 3D Visualization: Glyphs



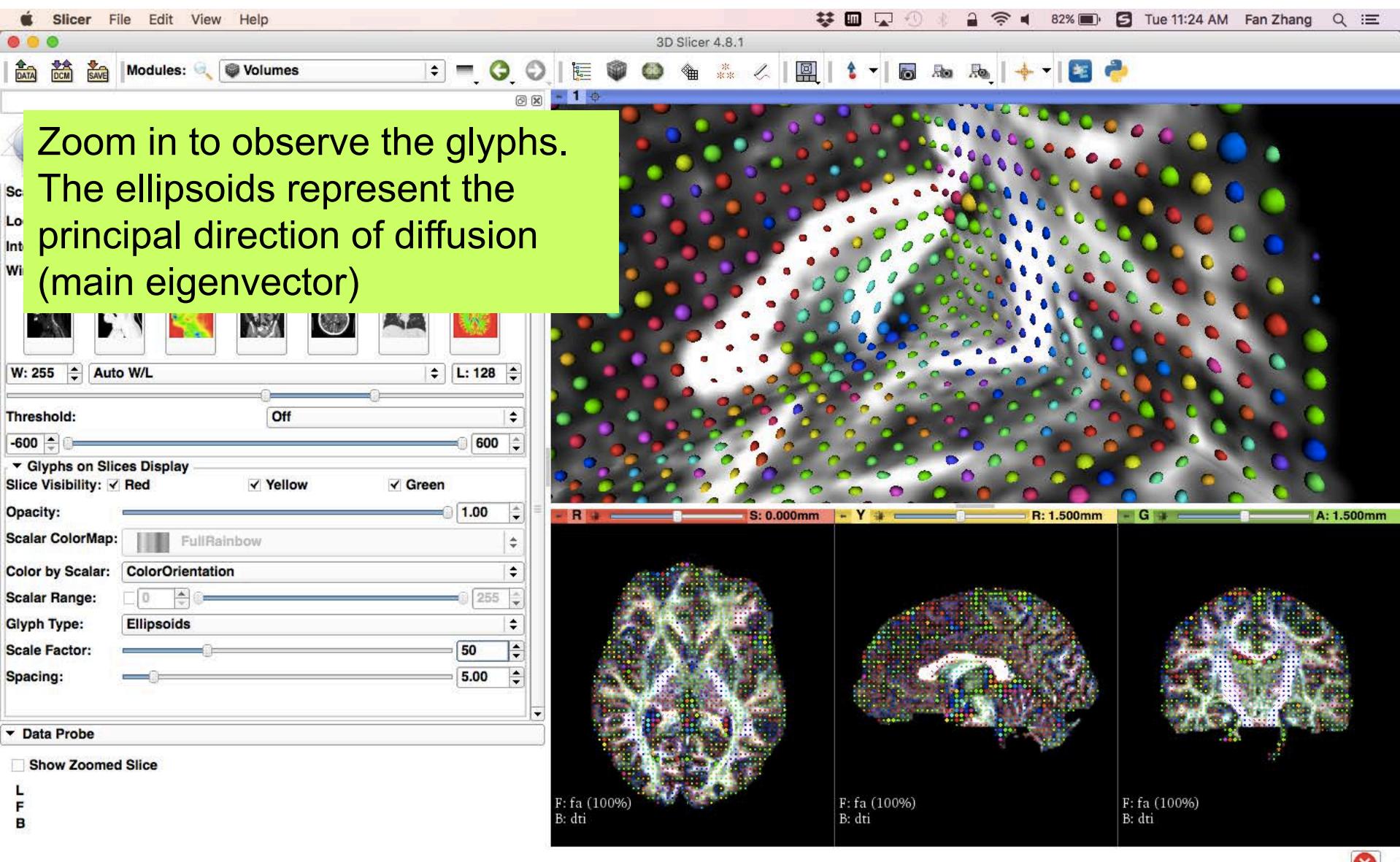
# 3D Visualization: Glyphs



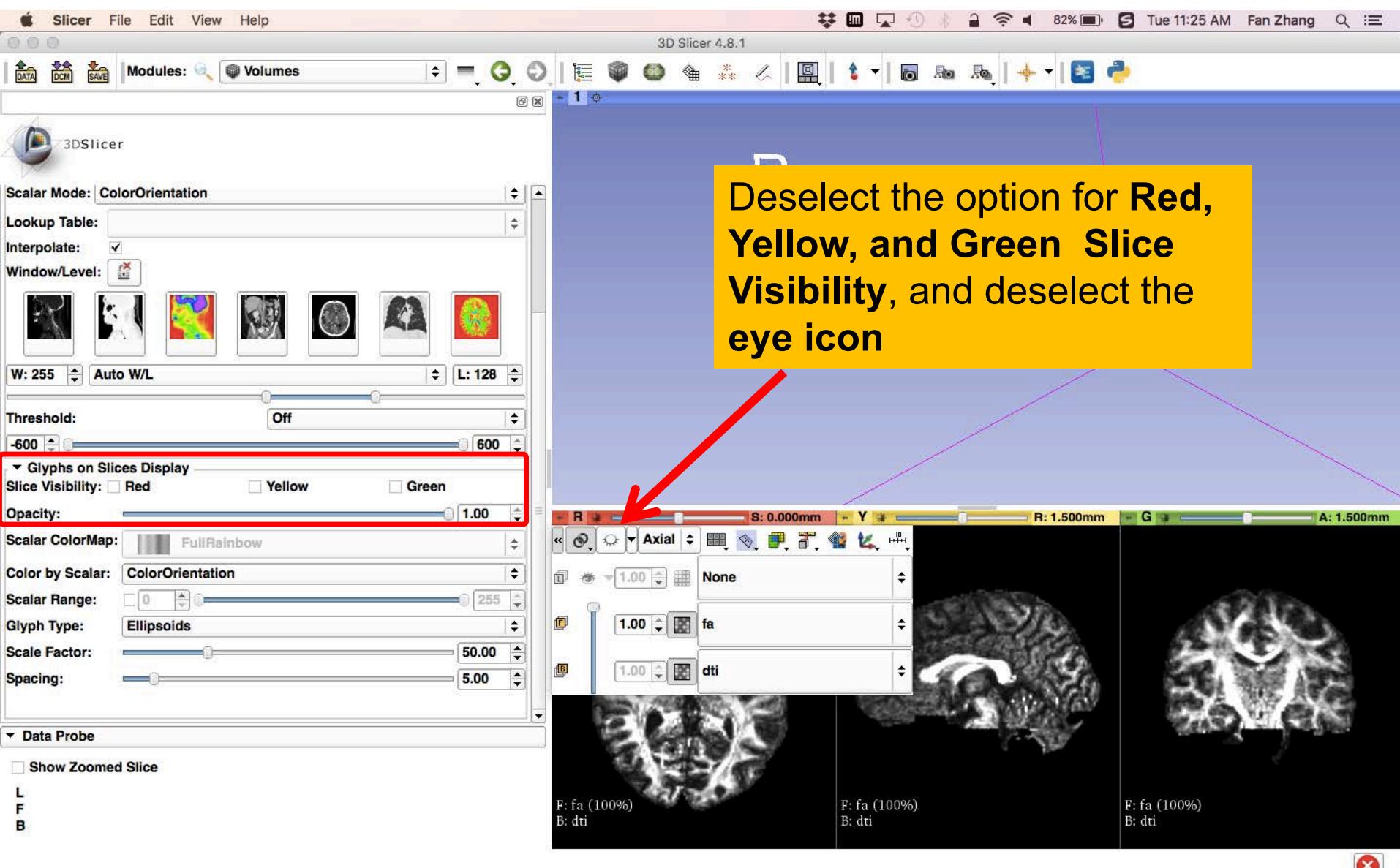
# 3D Visualization: Glyphs



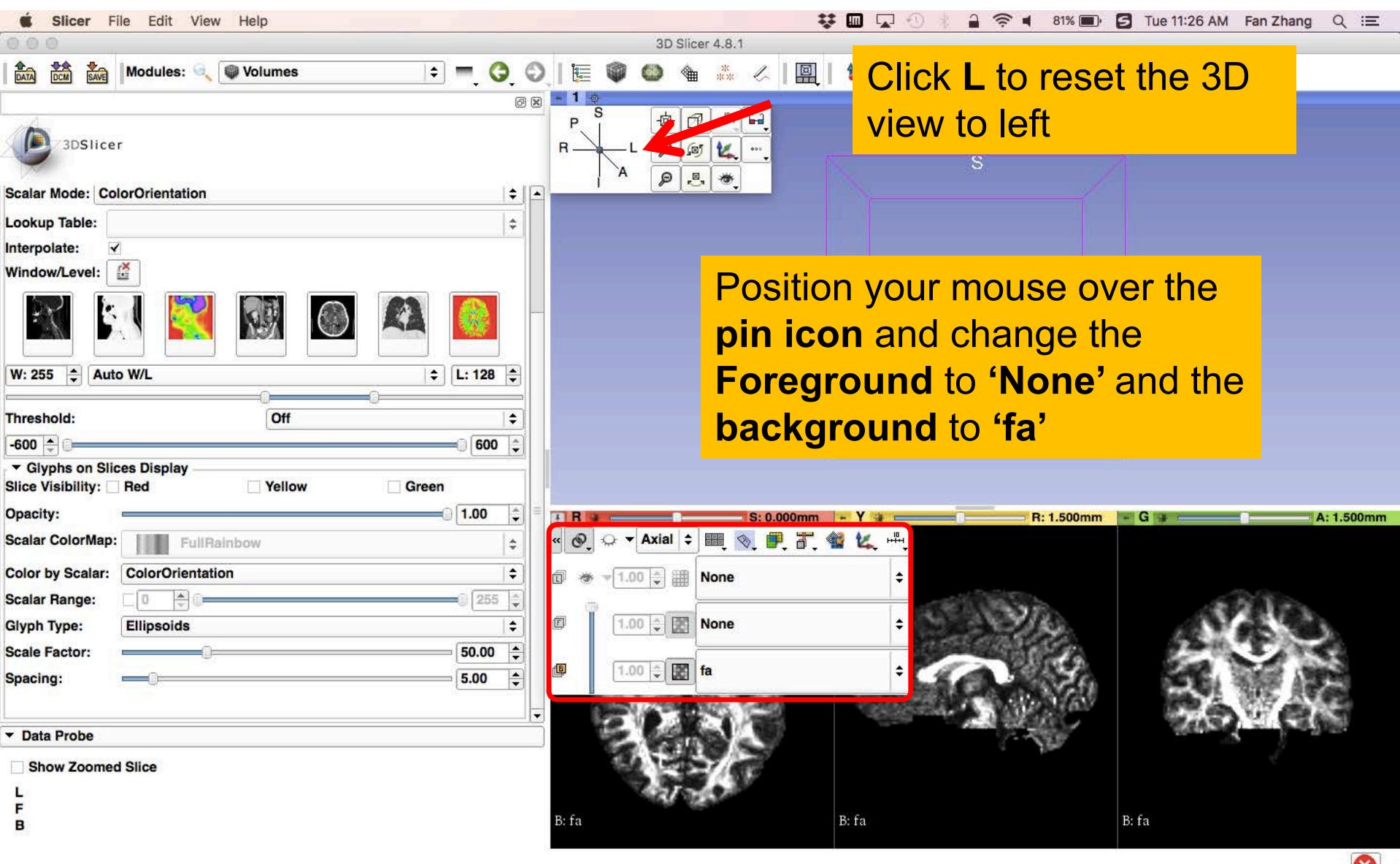
# 3D Visualization: Glyphs

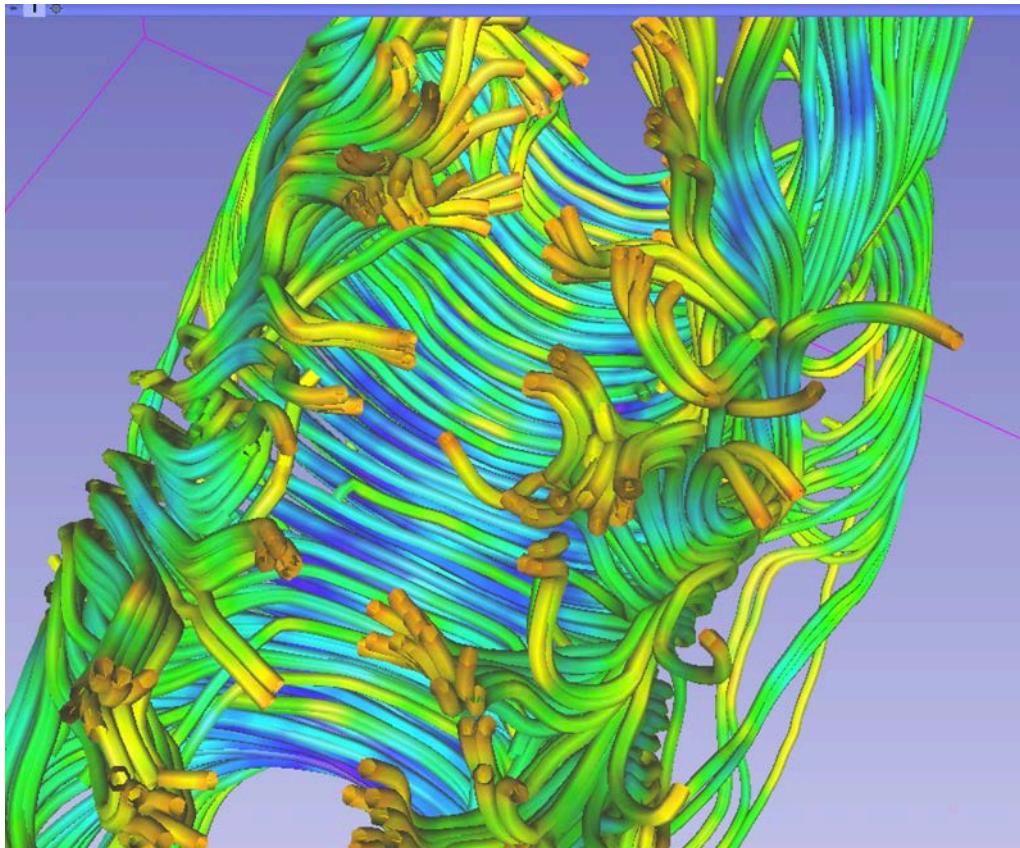


# Diffusion MRI tractography



# Diffusion MRI tractography



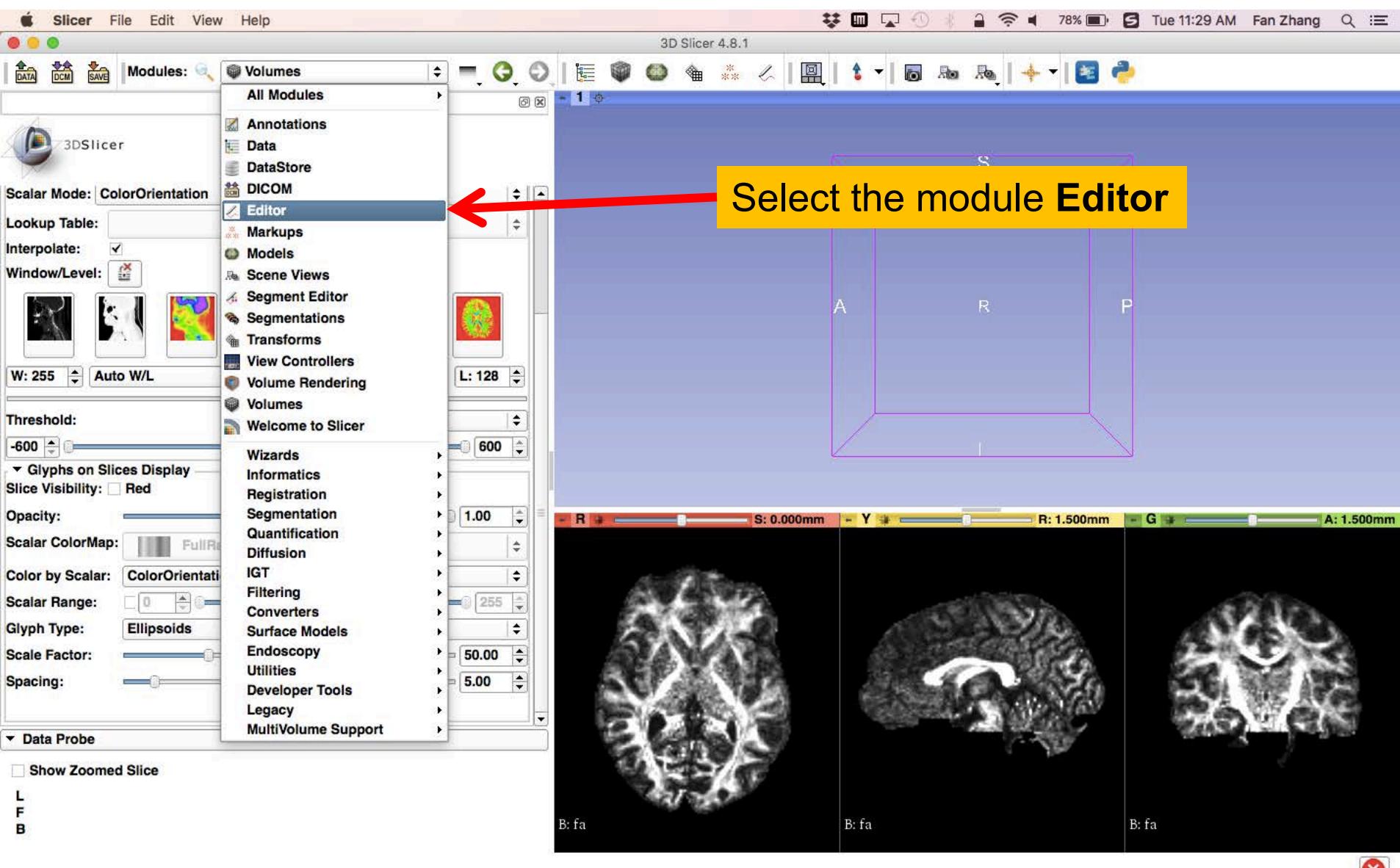


## Part 3: From tensors to tracts

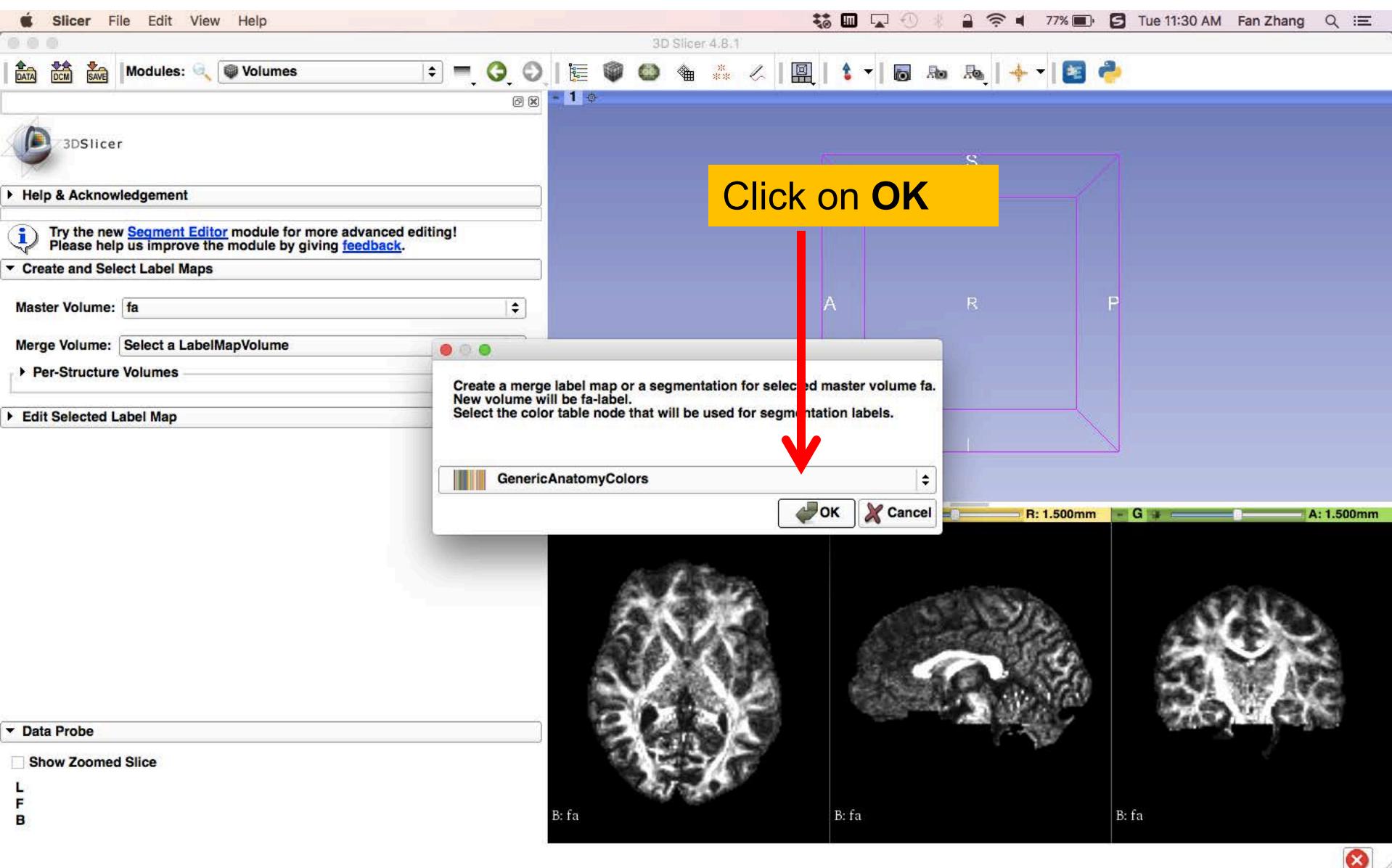
# DTI tractography

- Definition of a region of interest (ROI) for seeding tract in an FA map (Editor module)
- Single-tensor tractography (Tractography Interactive Seeding module)
- Fiducial-seeding tractography (Tractography Interactive Seeding module)

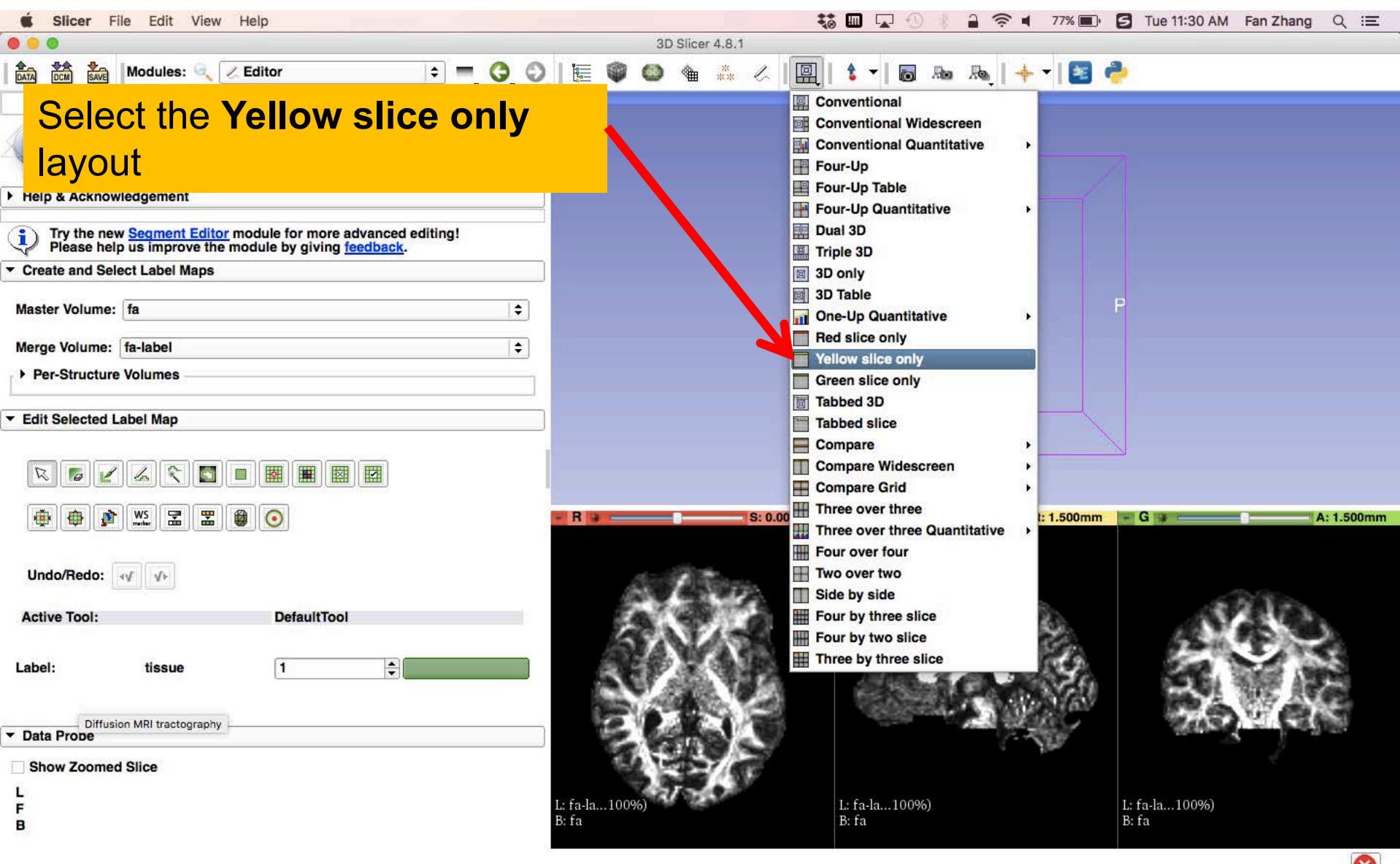
# Diffusion MRI tractography



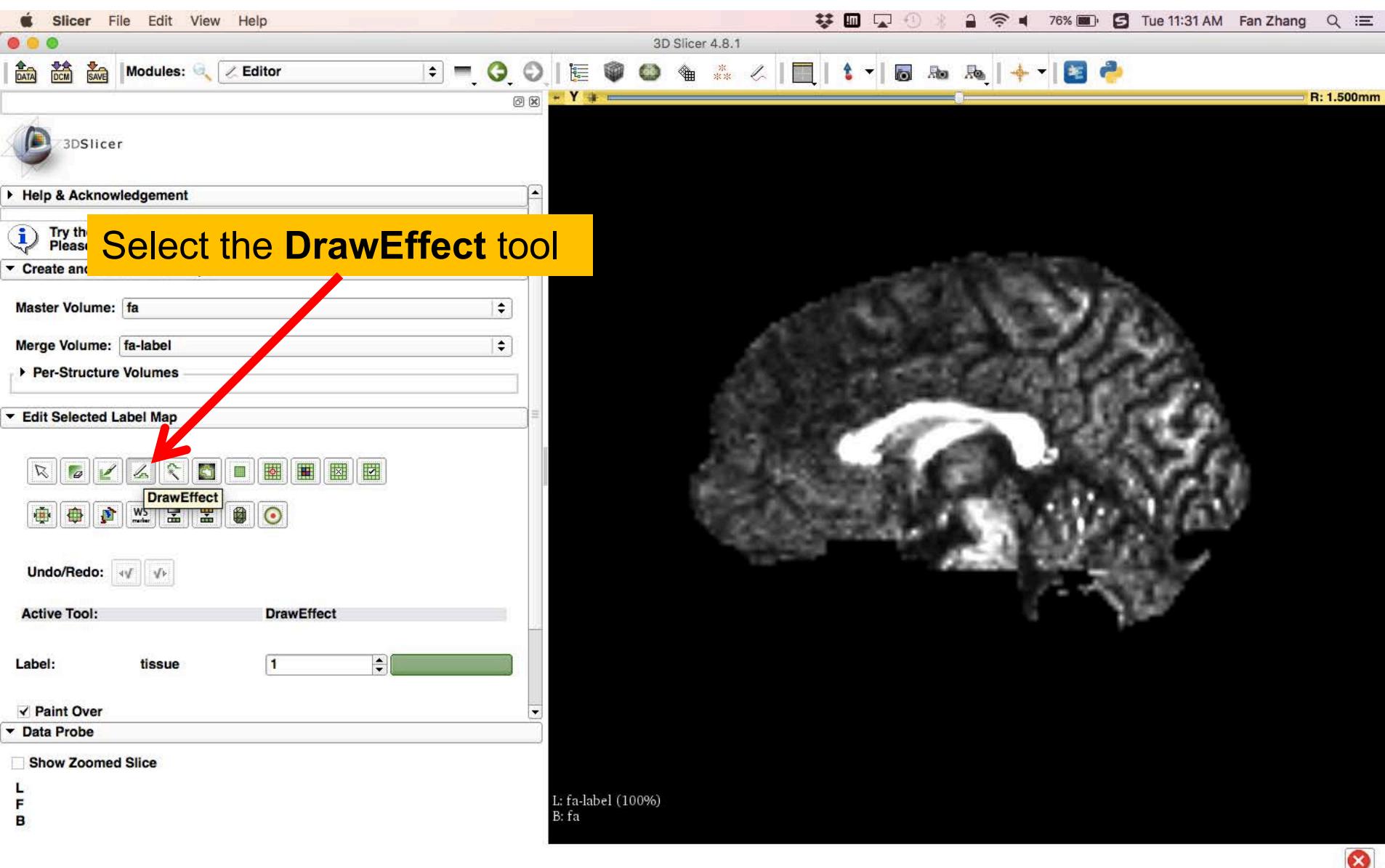
# Diffusion MRI tractography



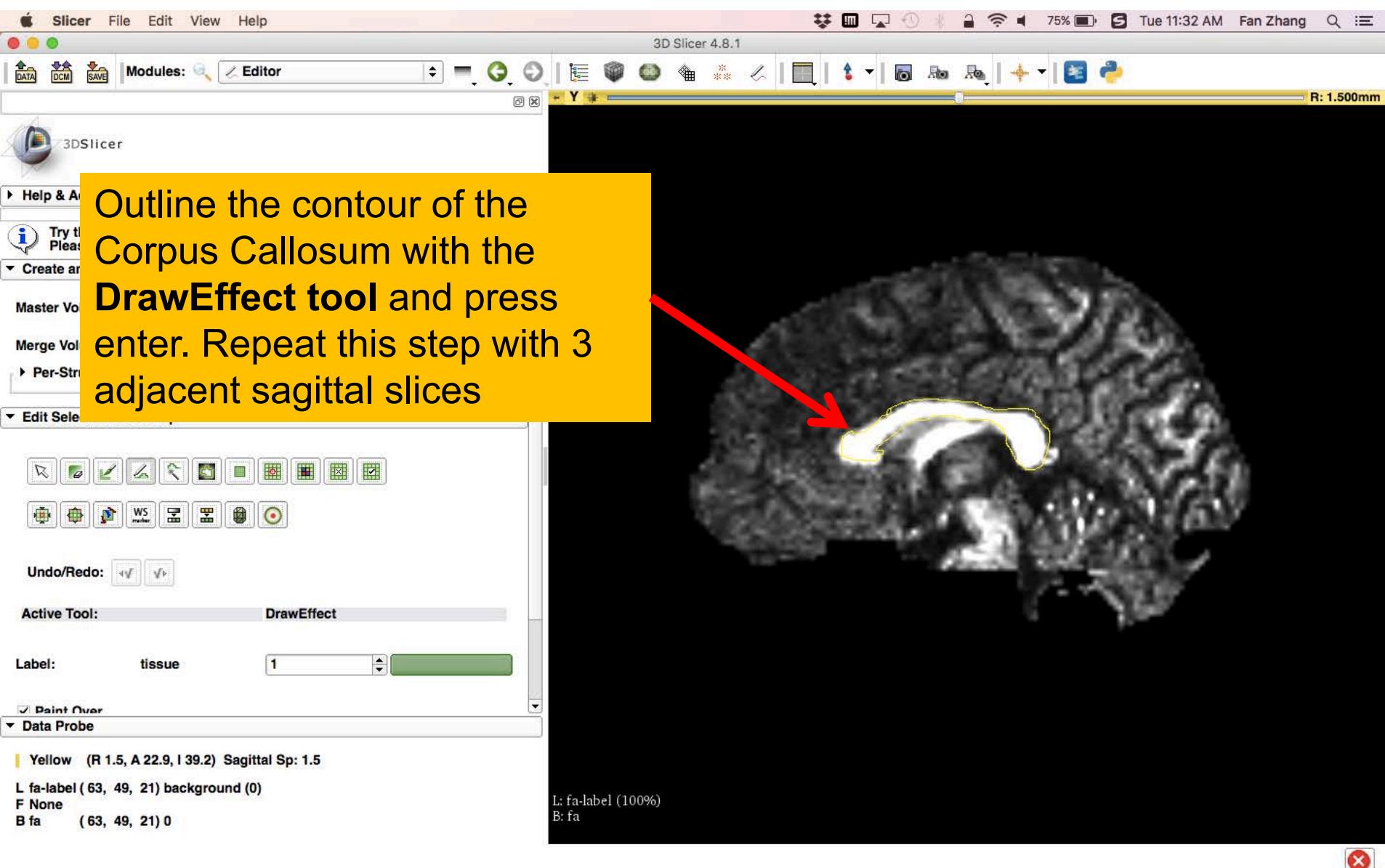
# Diffusion MRI tractography



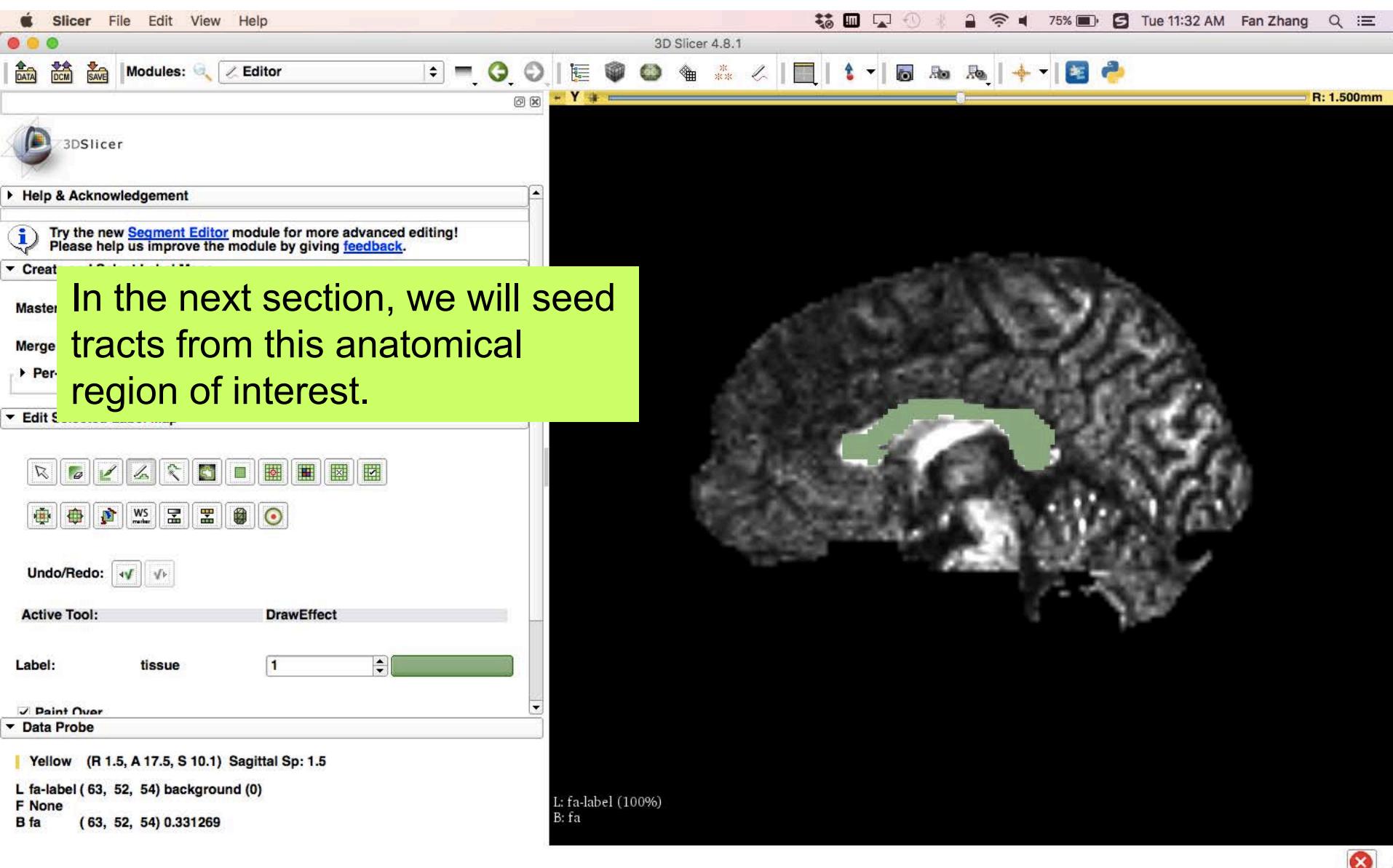
# Diffusion MRI tractography



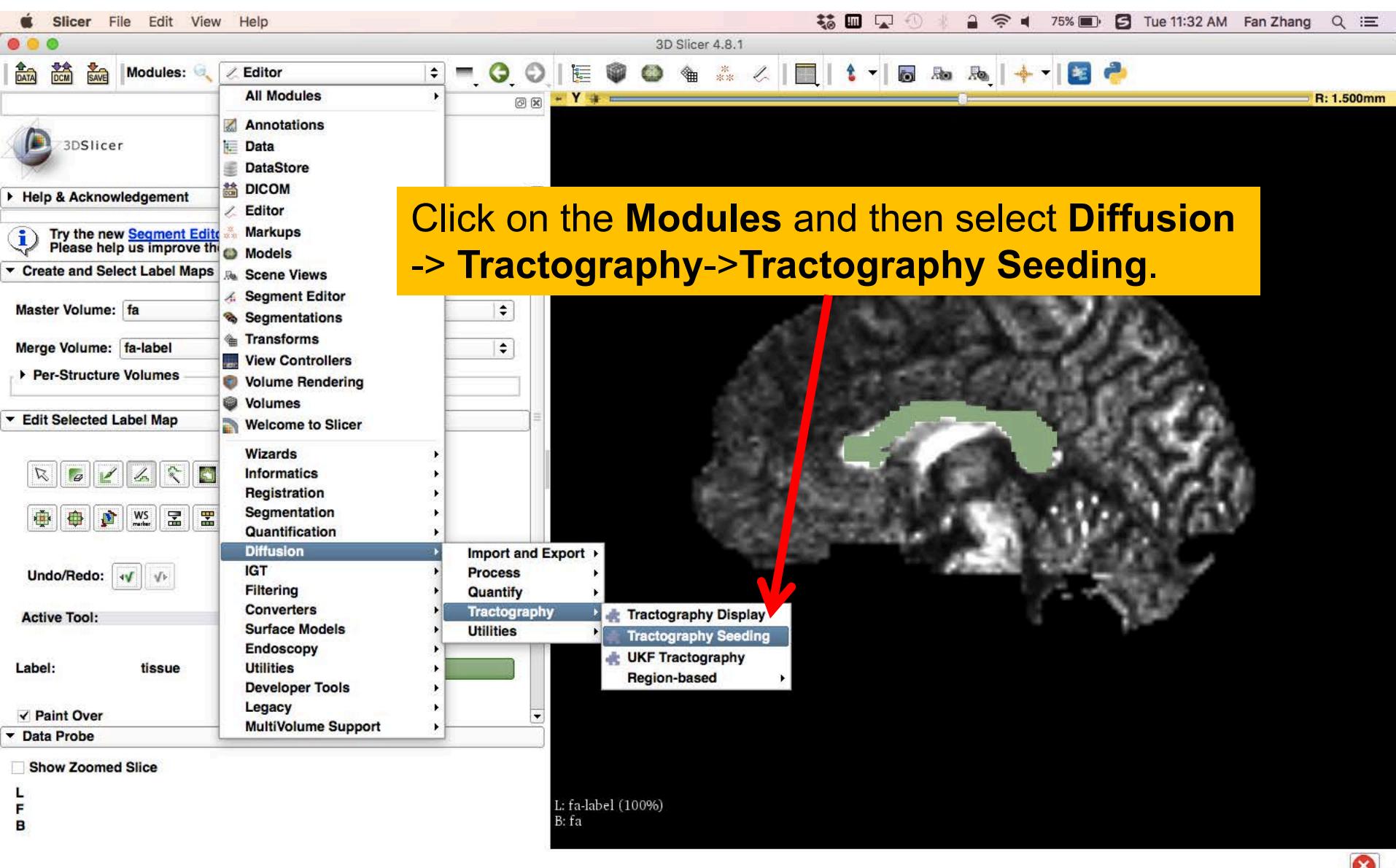
# Diffusion MRI tractography



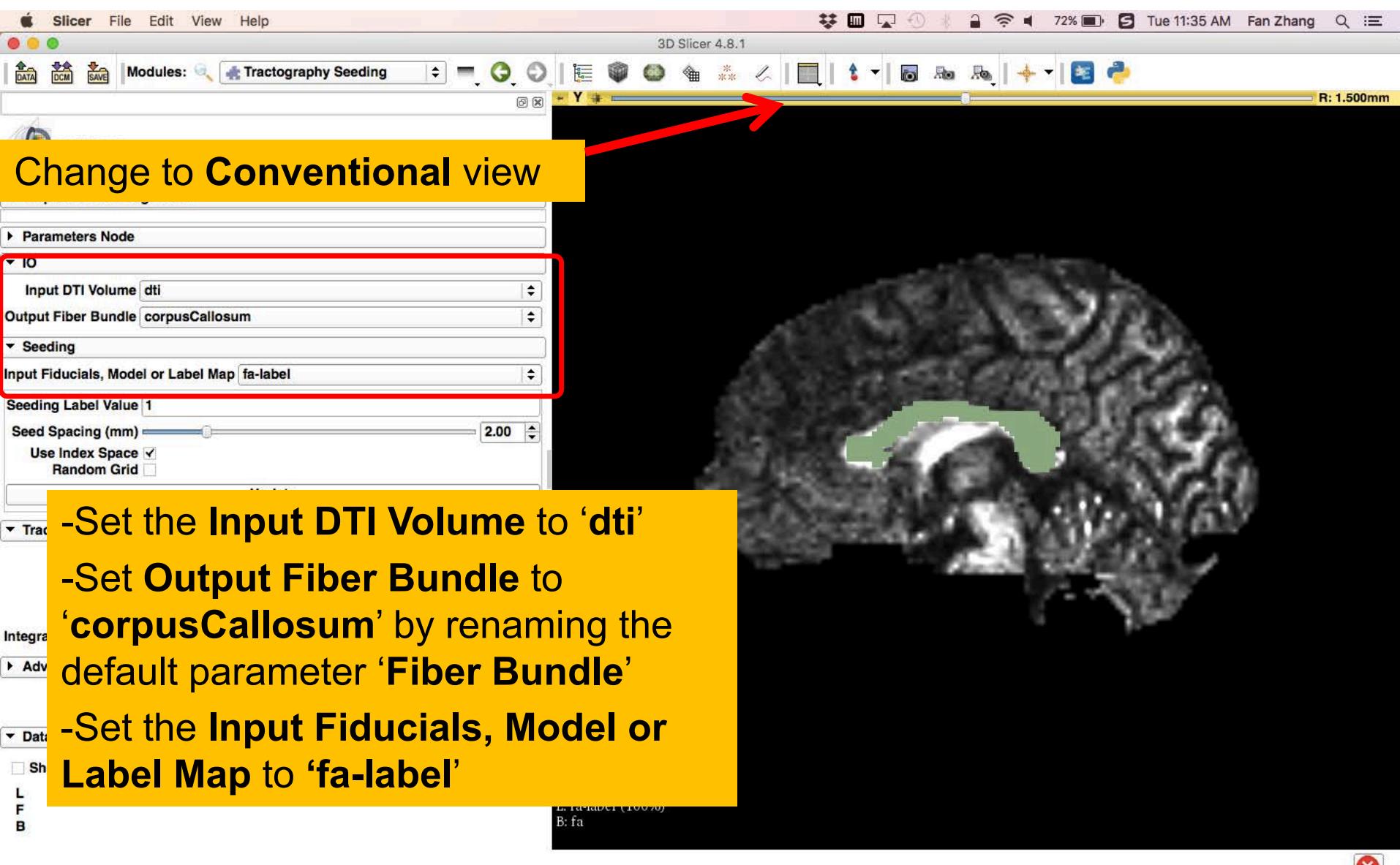
# Diffusion MRI tractography



# Diffusion MRI tractography



# Step1: I/O



Change to **Conventional** view

**Parameters Node**

**IO**

- Input DTI Volume: dti
- Output Fiber Bundle: corpusCallosum
- Seeding**
- Input Fiducials, Model or Label Map: fa-label
- Seeding Label Value: 1
- Seed Spacing (mm): 2.00
- Use Index Space
- Random Grid

- Set the **Input DTI Volume** to 'dti'  
- Set **Output Fiber Bundle** to '**corpusCallosum**' by renaming the default parameter '**Fiber Bundle**'  
- Set the **Input Fiducials, Model or Label Map** to 'fa-label'

L: Ta-Ra-CT (100%)  
B: fa

# Step 2: Seeding parameters

3D Slicer 4.8.1

Select the default Tractography Seeding parameters:

- Threshold Type: FractionalAnistropy
- Seeding Threshold:0.30
- Stopping Threshold: 0.25

Click **Update** to generate tractography

3D Slicer

DATA DCM SAVE Modules: Tractography Seeding

Help & Acknowledgement

Parameters Node

IO

Input DTI Volume: dti

Output Fiber Bundle: corpusCallosum

Seeding

Input Fiducials, Model or Label Map: fa-label

Seeding Label Value: 1

Seed Spacing (mm):

Use Index Space  Random Grid

Update

Tractography Parameters

- Threshold Type: Fractional Anisotropy
- Seeding Threshold: 0.30
- Stopping Threshold: 0.25
- Integration Step Length (mm): 0.500mm

Advanced Options

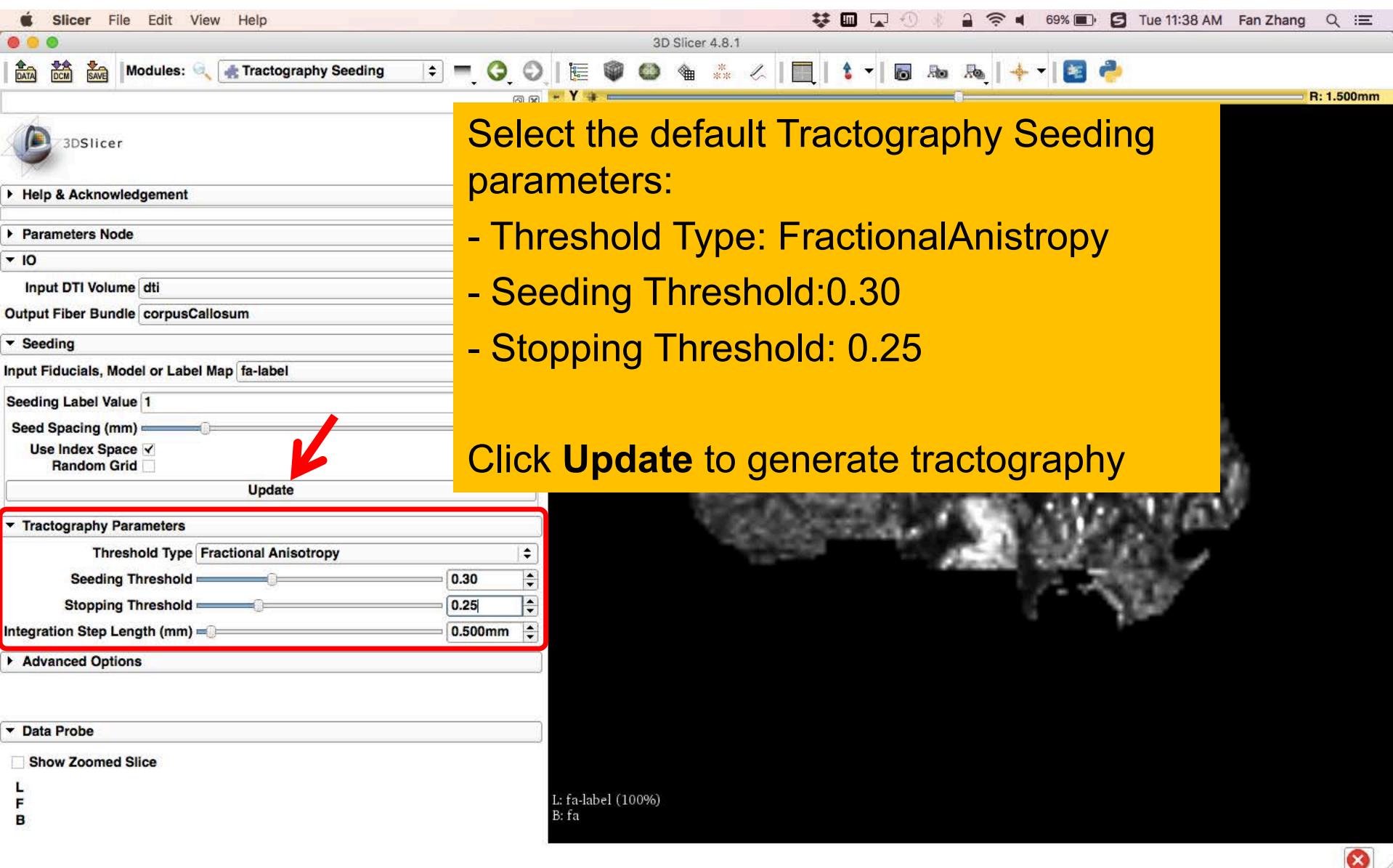
Data Probe

Show Zoomed Slice

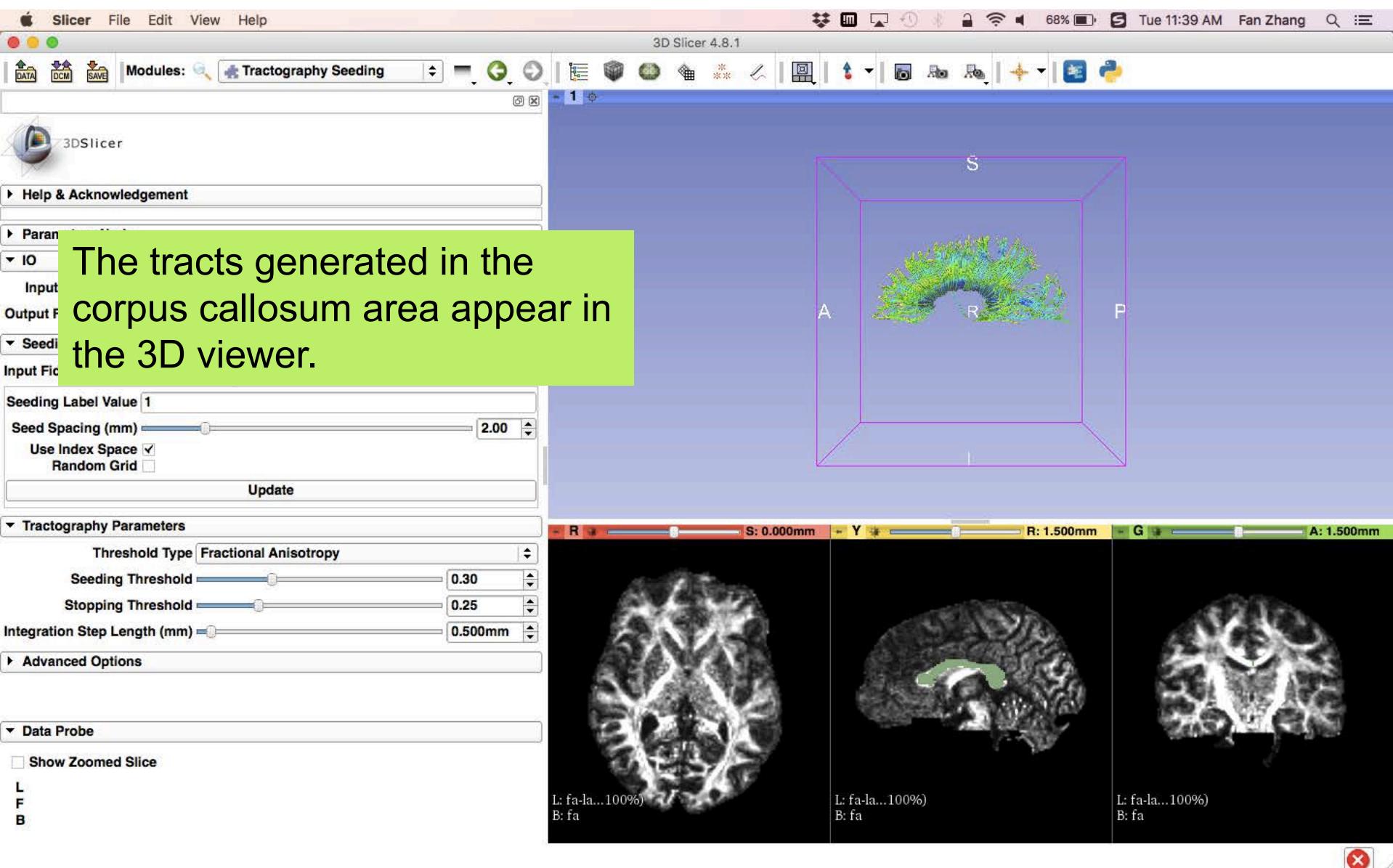
L F B

R: 1.500mm

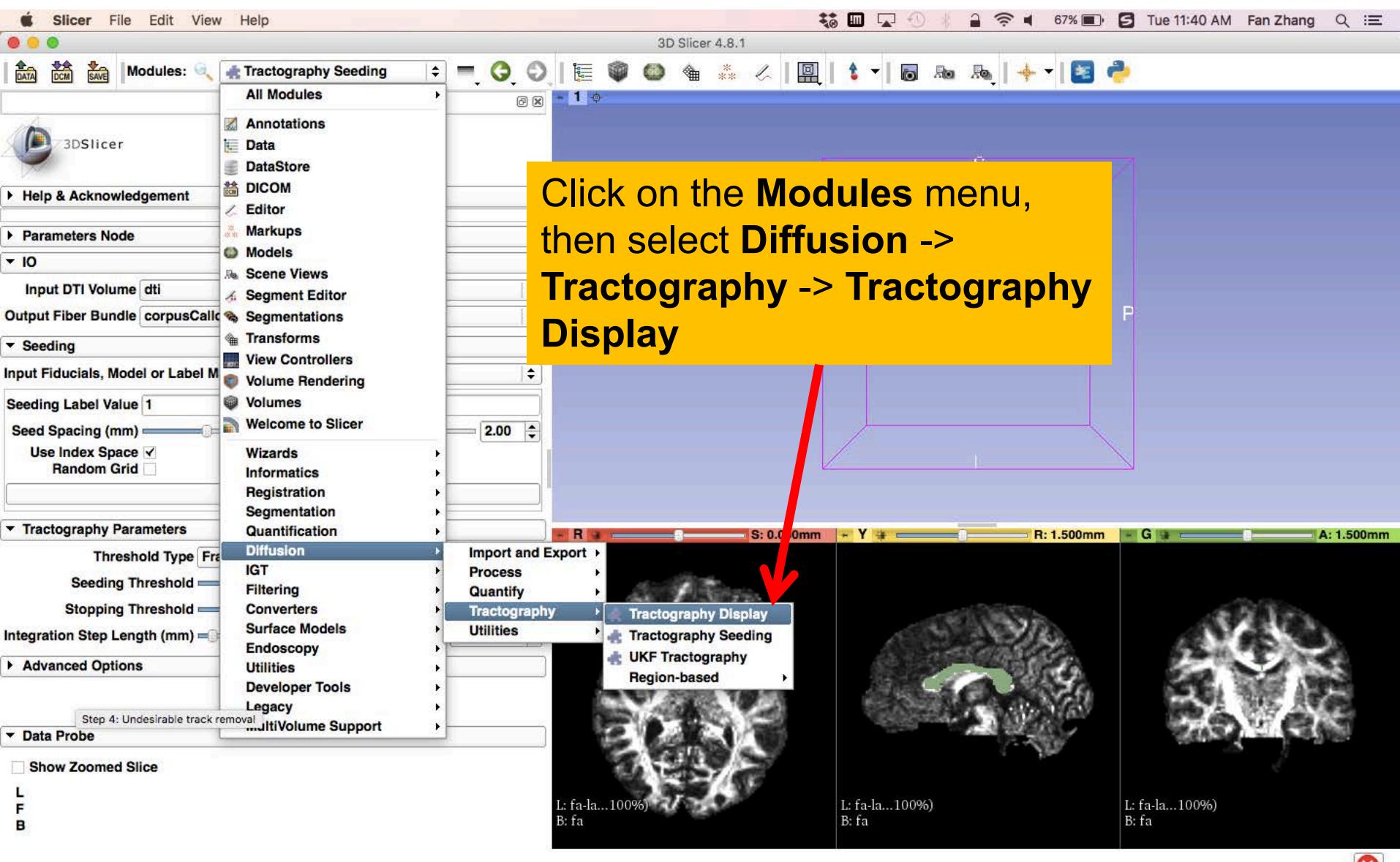
L: fa-label (100%)  
B: fa



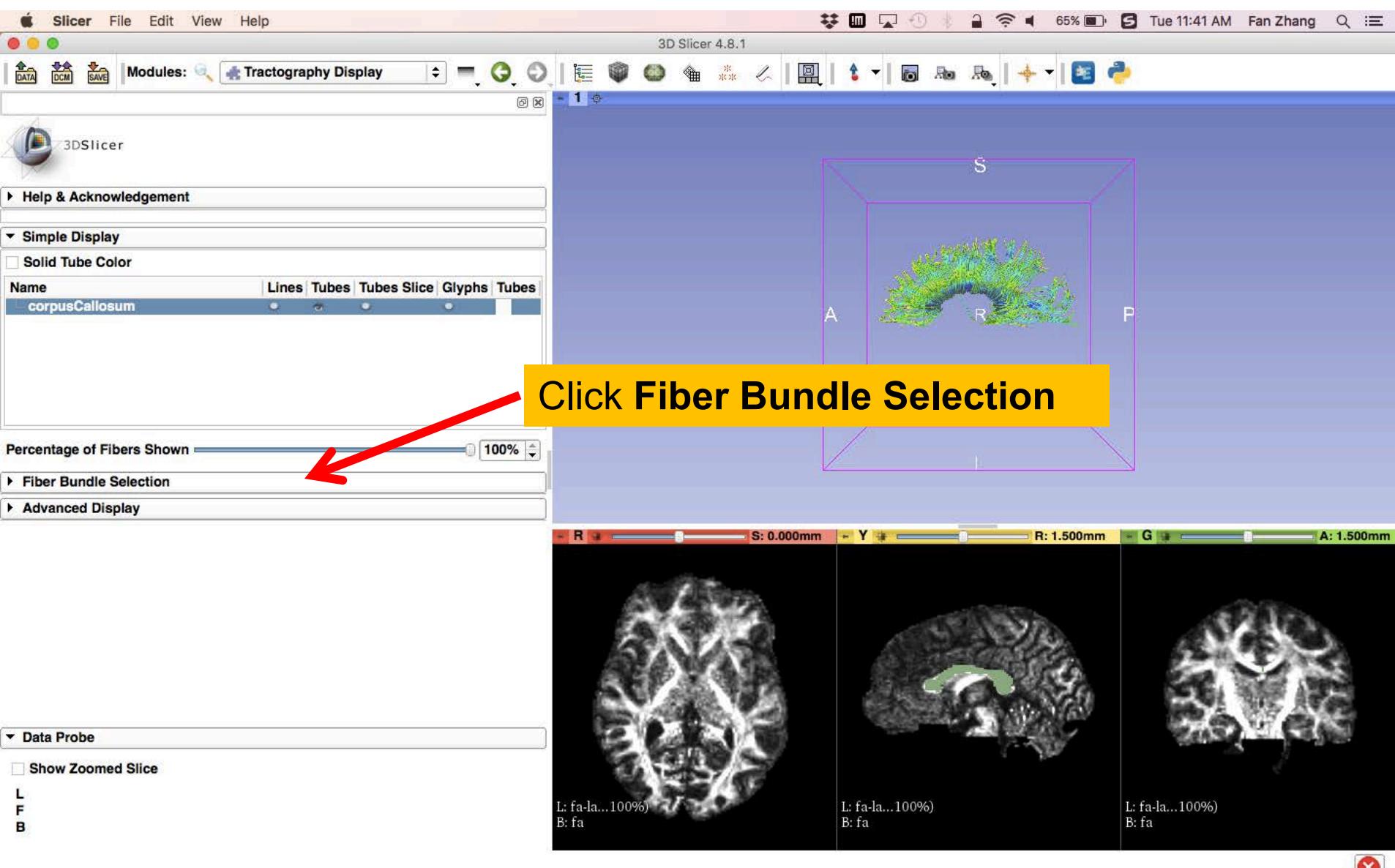
# Step 3: Generate Tracts



# Step 4: Undesirable track removal



# Step 4: Undesirable track removal



# Step 4: Undesirable track removal

Slicer File Edit View Help

3D Slicer 4.8.1

DATA DCM SAVE Modules: Tractography Display

3DSlicer

Help & Acknowledgement

Simple Display

Solid Tube Color

Name: corpusCallosum

Percentage of Fibers Shown: 100%

Fiber Bundle Selection

ROI for Fiber Selection: ROI Node

Disable ROI (radio button)

Positive ROI (radio button)

Negative ROI (radio button)

Interactive ROI Updates (checkbox)

ROI Visibility (checkbox)

Extract Bundle From ROI: None

Update corpusCallosum From ROI (checkbox)

Confirm update (checkbox)

Enable Interactive Edit (checkbox)

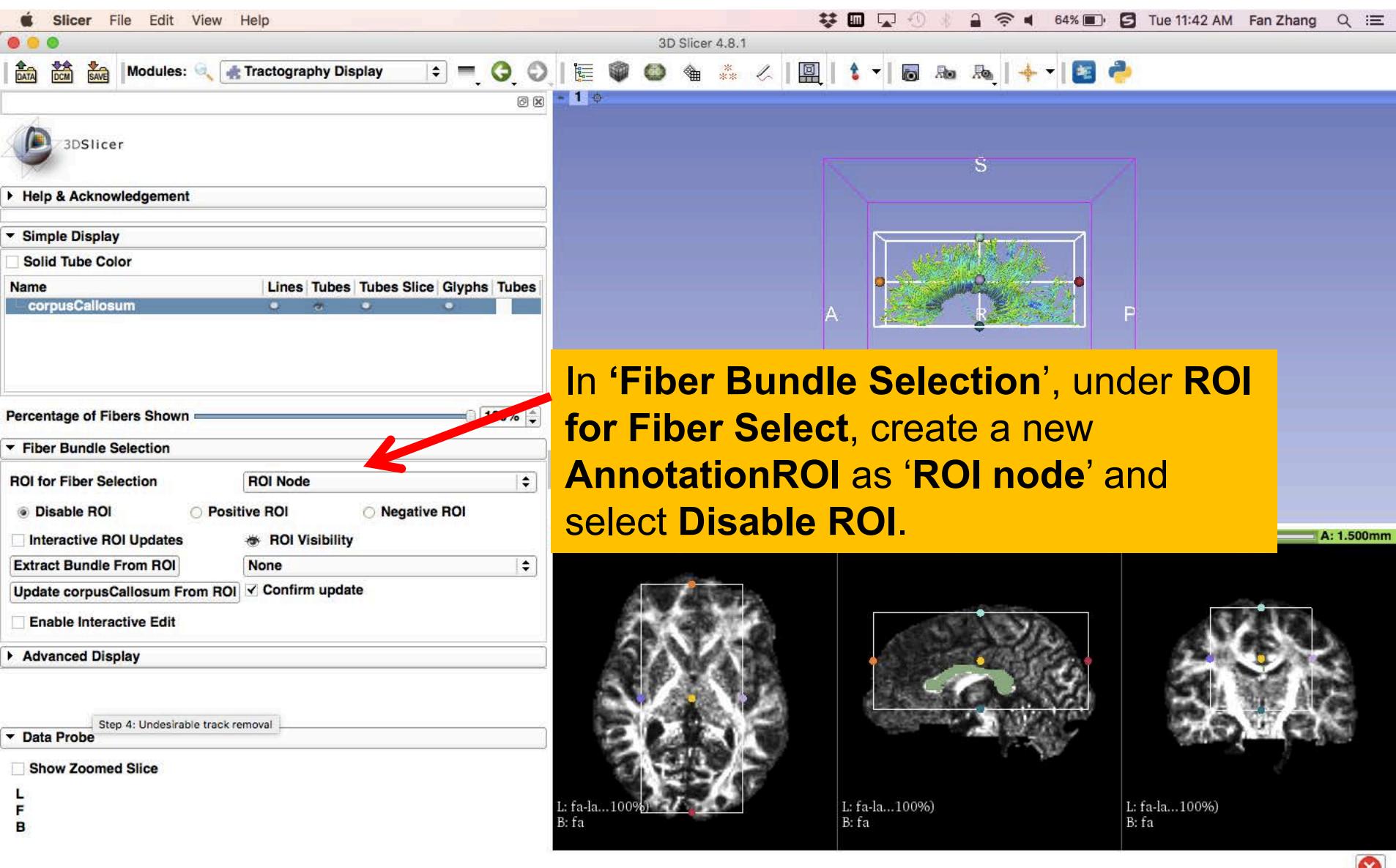
Advanced Display

Step 4: Undesirable track removal

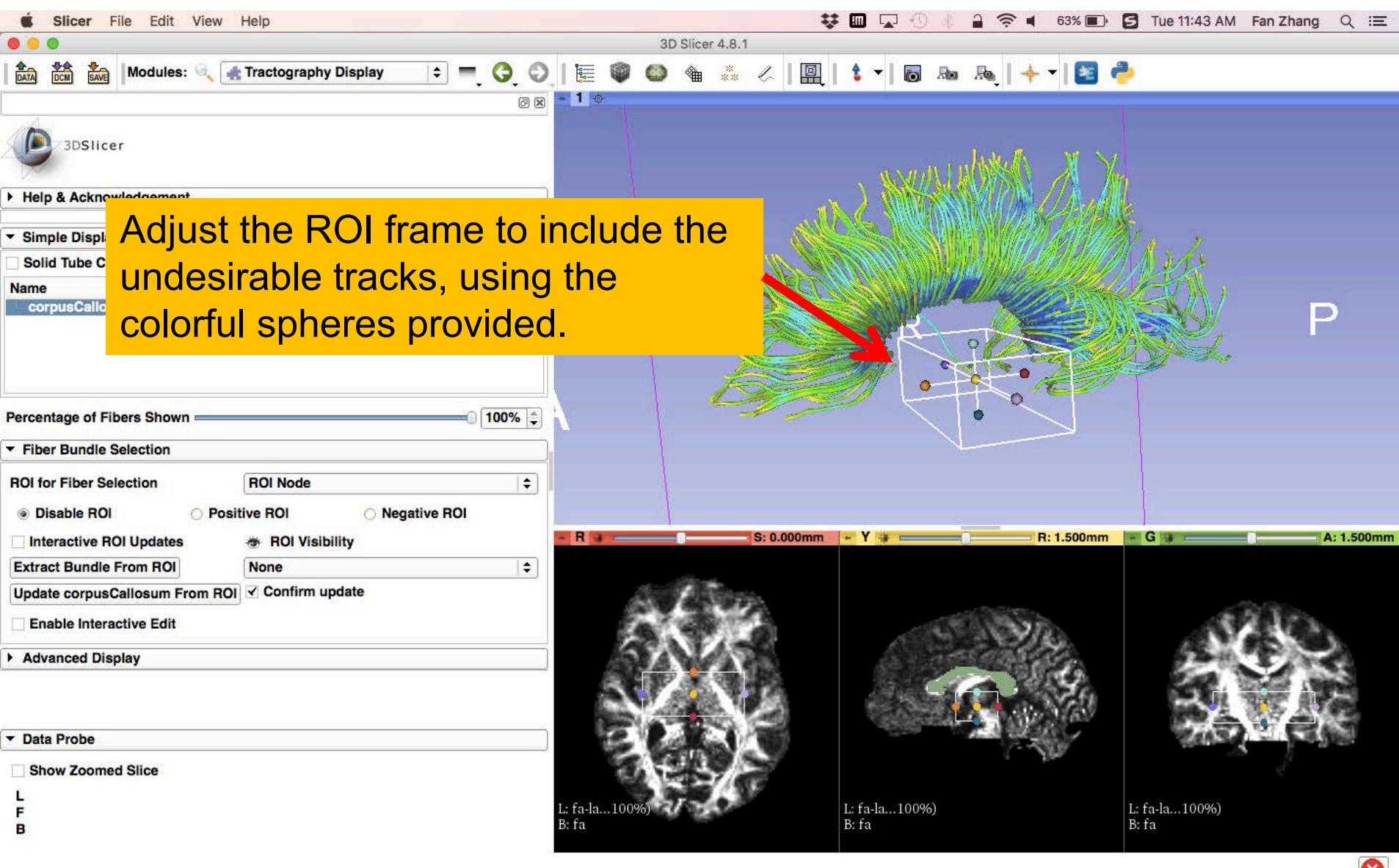
Show Zoomed Slice (checkbox)

L F B

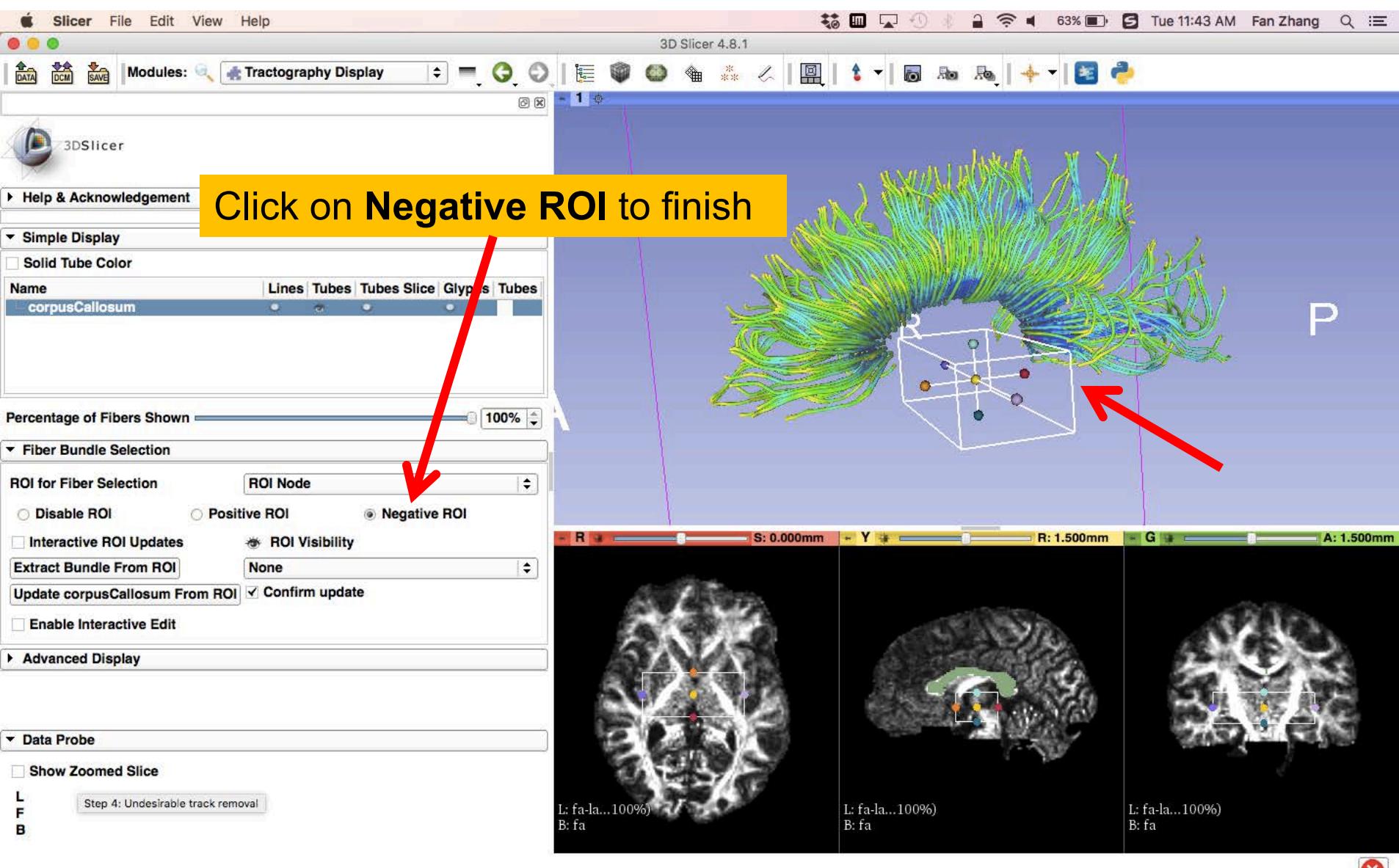
In 'Fiber Bundle Selection', under **ROI for Fiber Select**, create a new **AnnotationROI** as 'ROI node' and select **Disable ROI**.



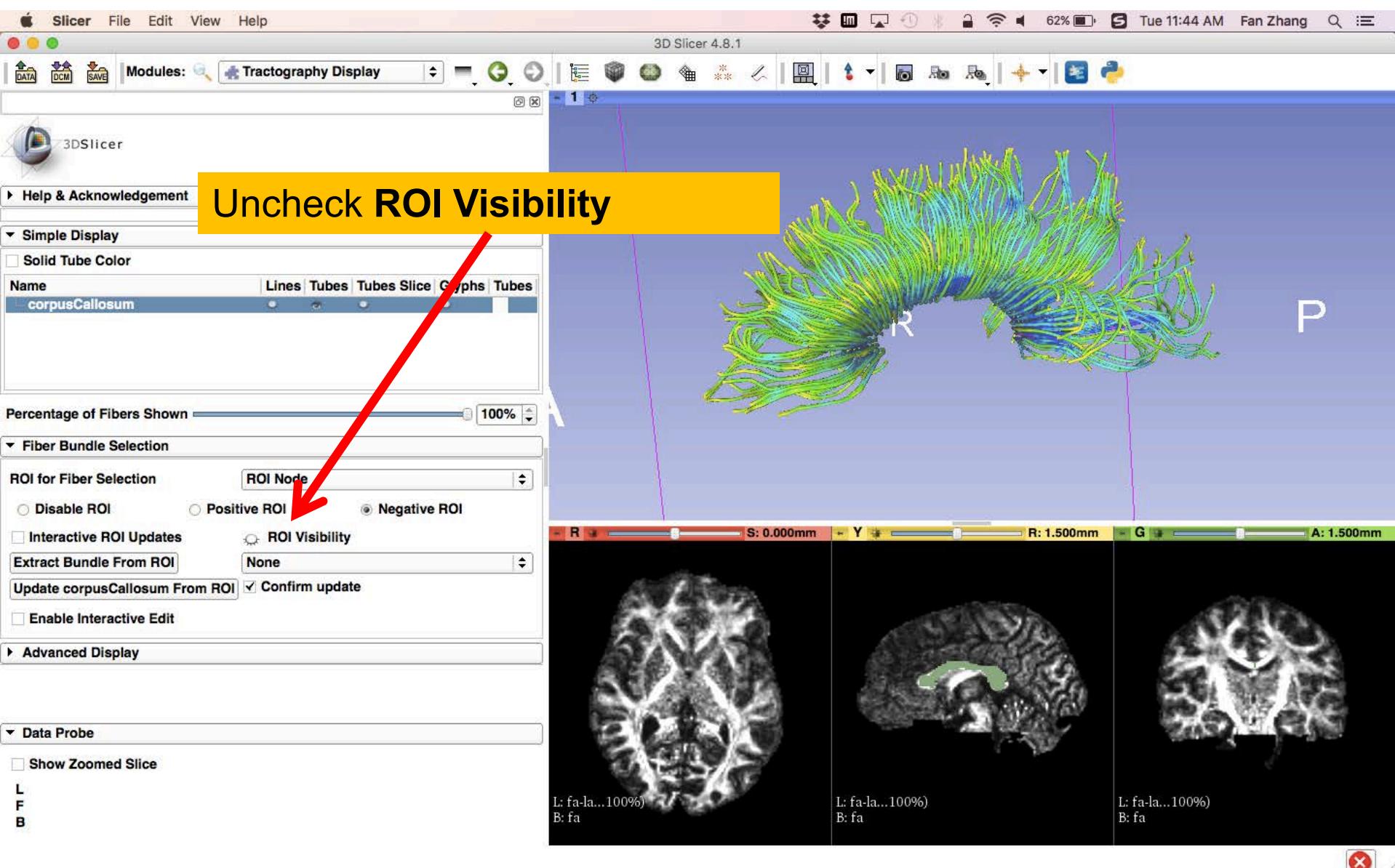
# Step 4: Undesirable track removal



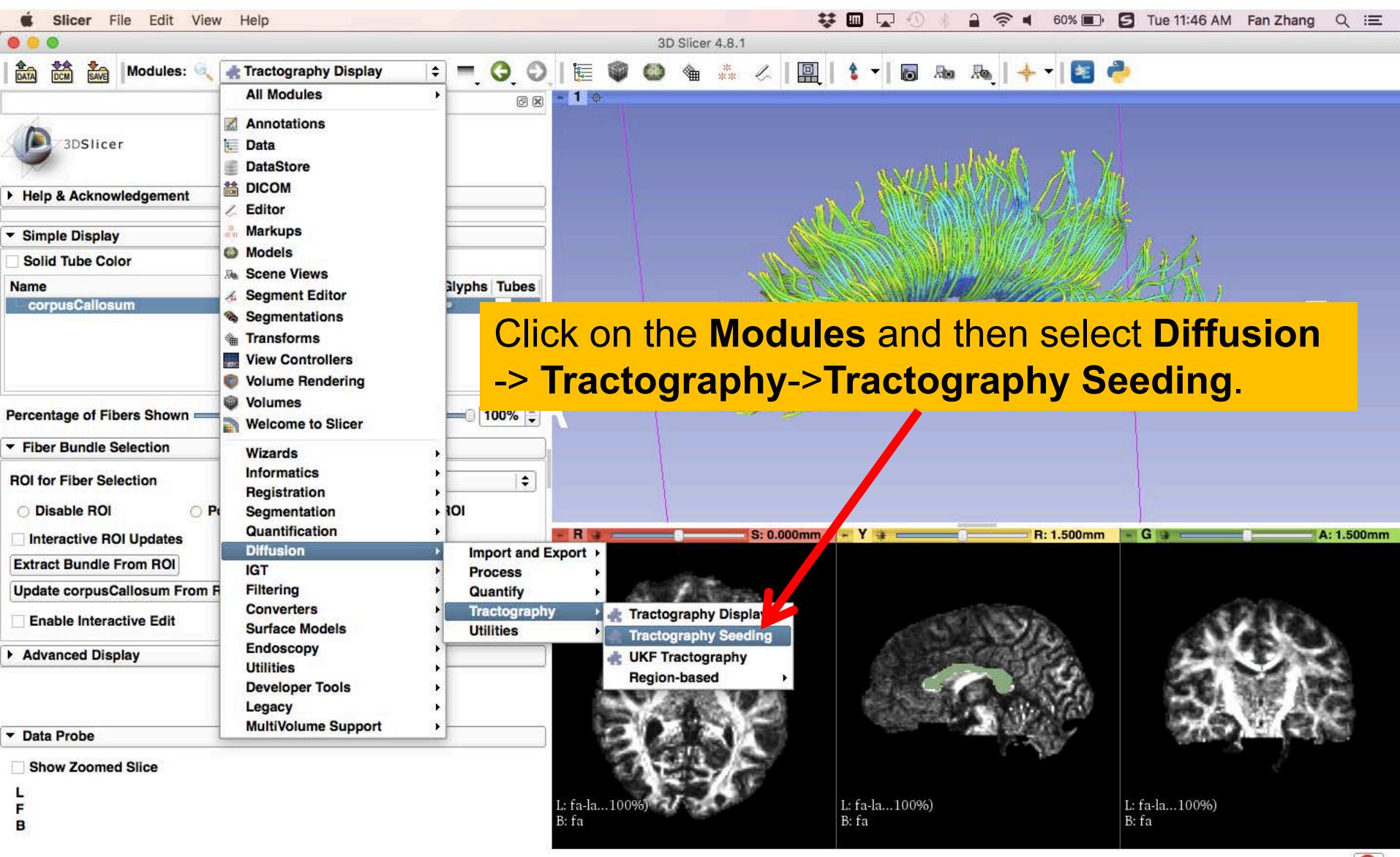
# Step 4: Undesirable track removal



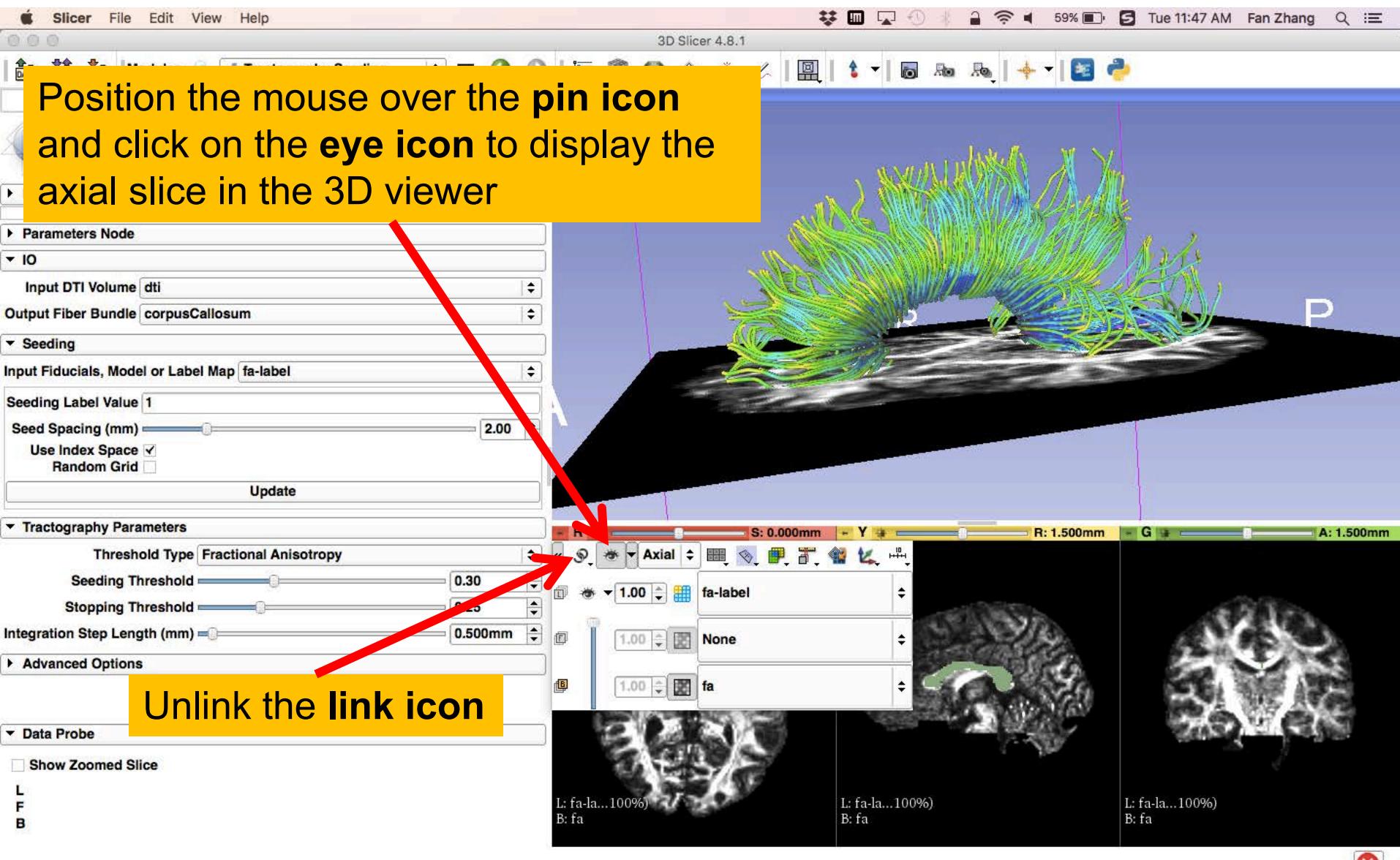
# Step 4: Undesirable track removal



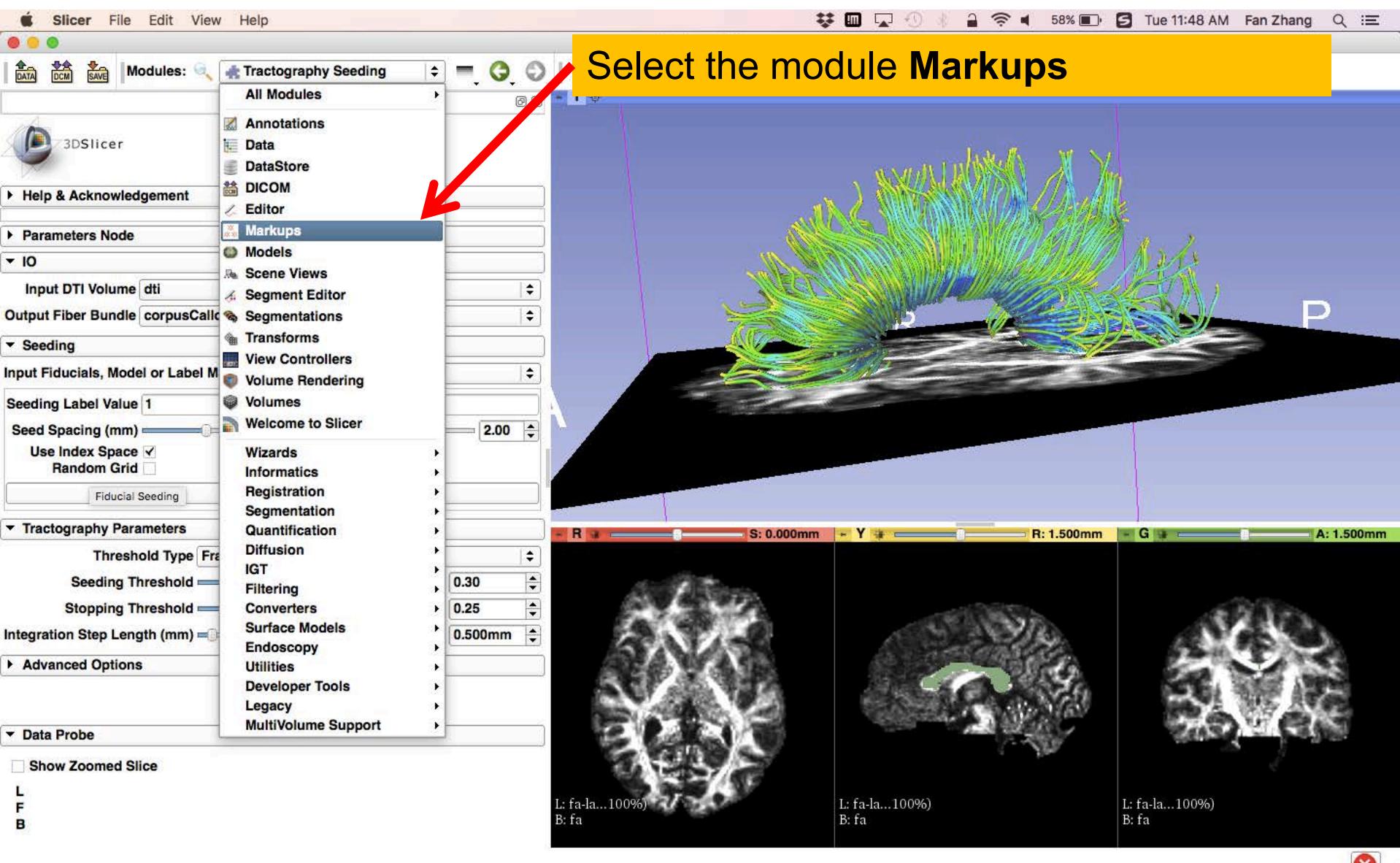
# Fiducial Seeding



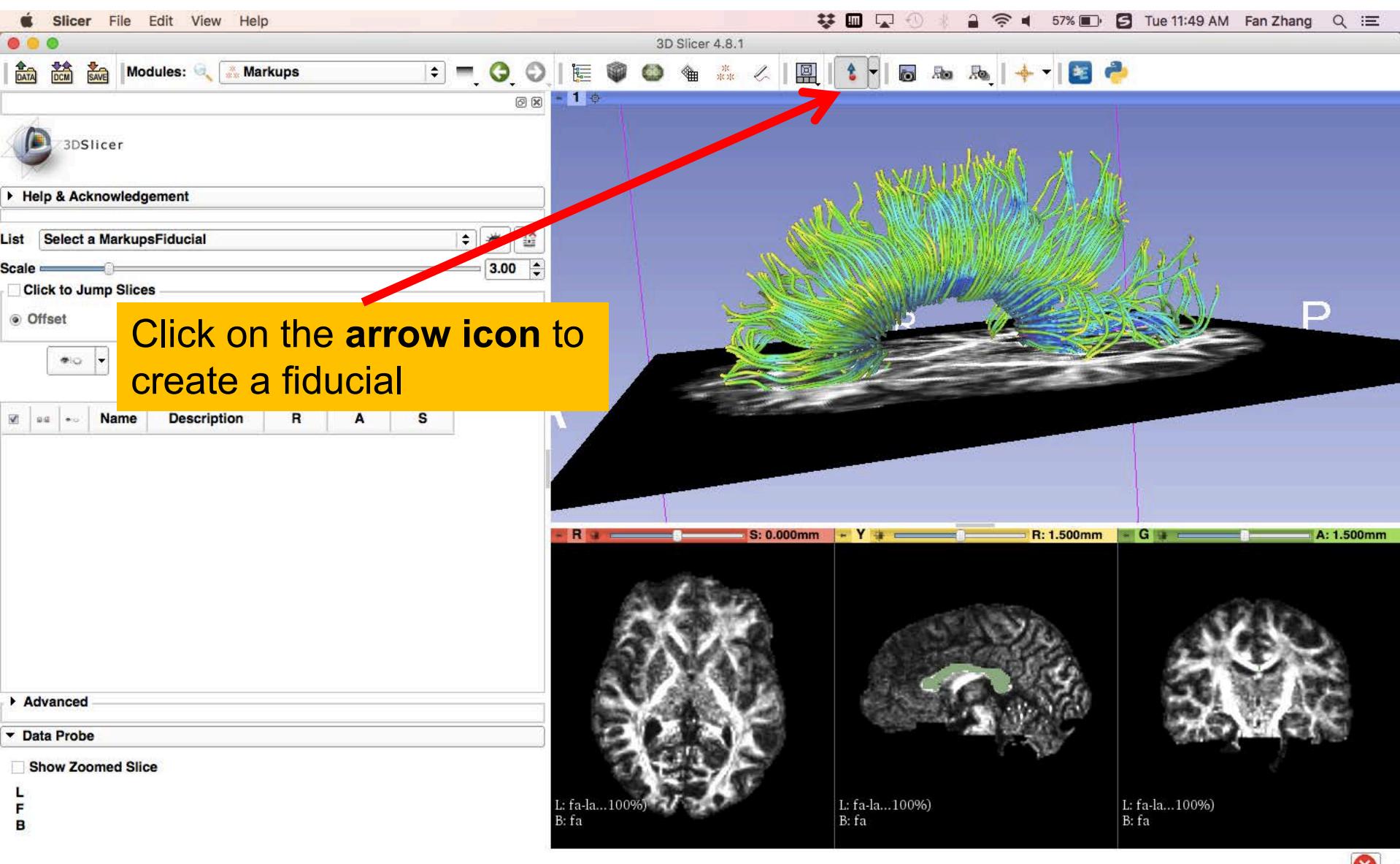
# Fiducial Seeding



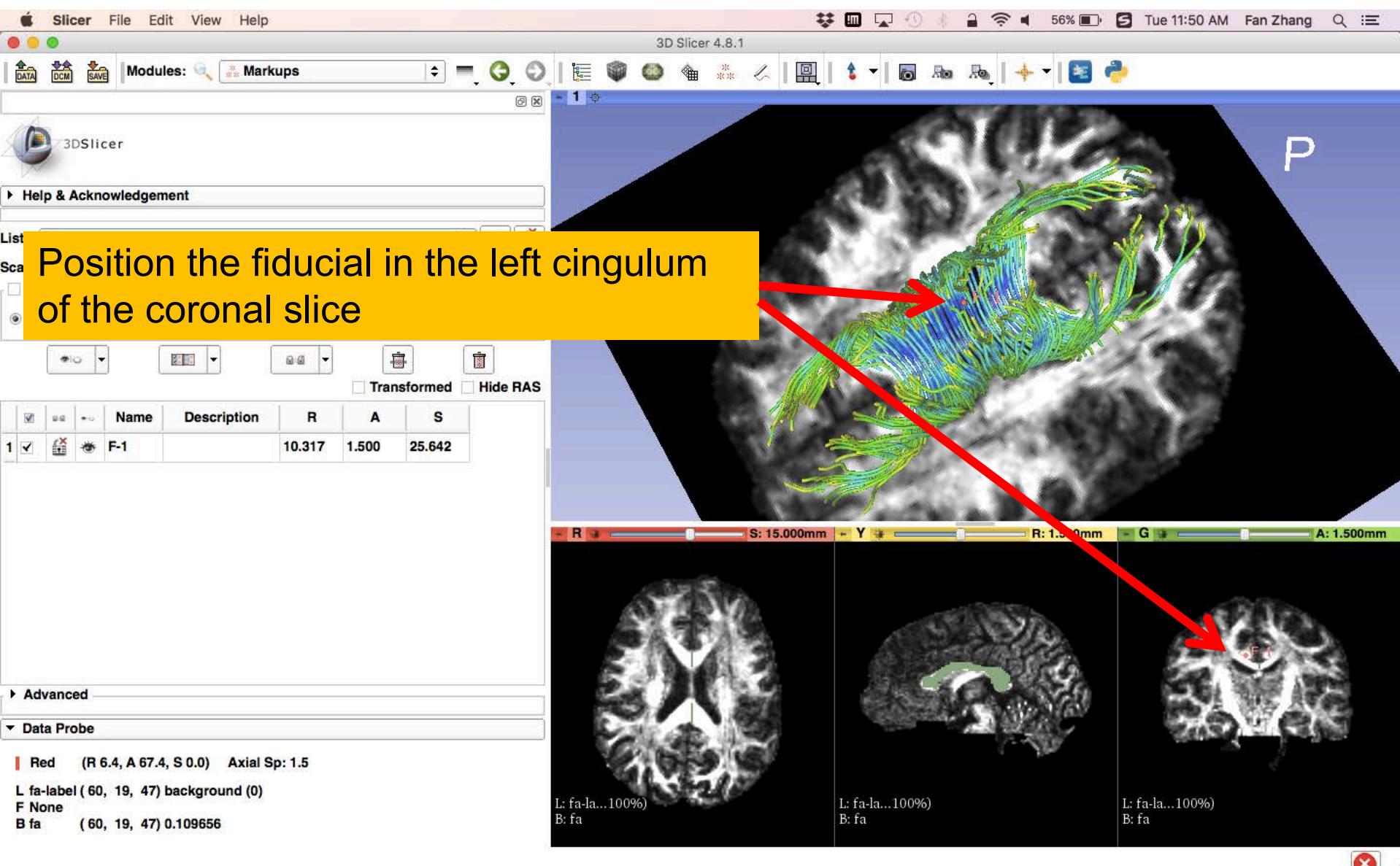
# Fiducial Seeding



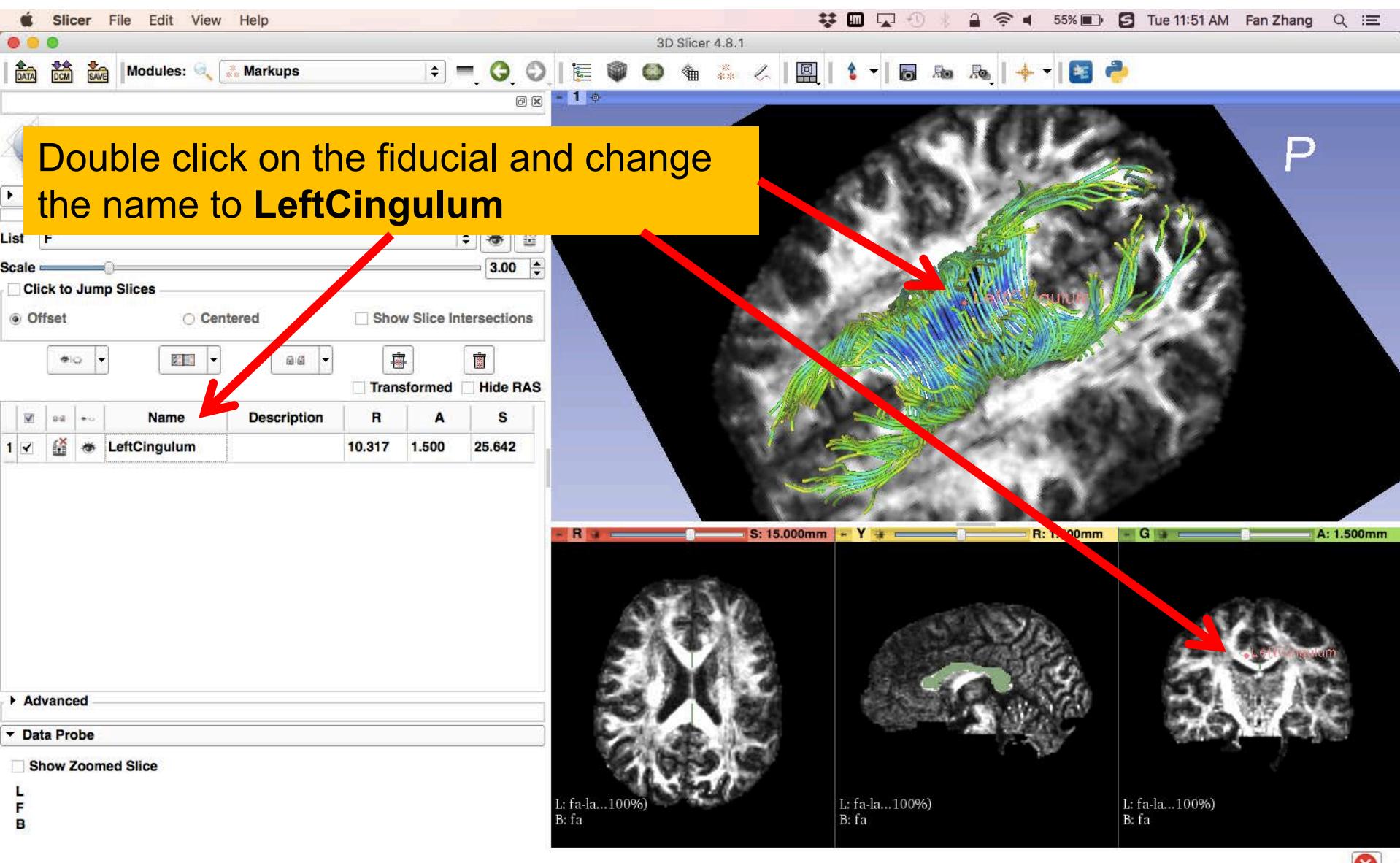
# Fiducial Seeding



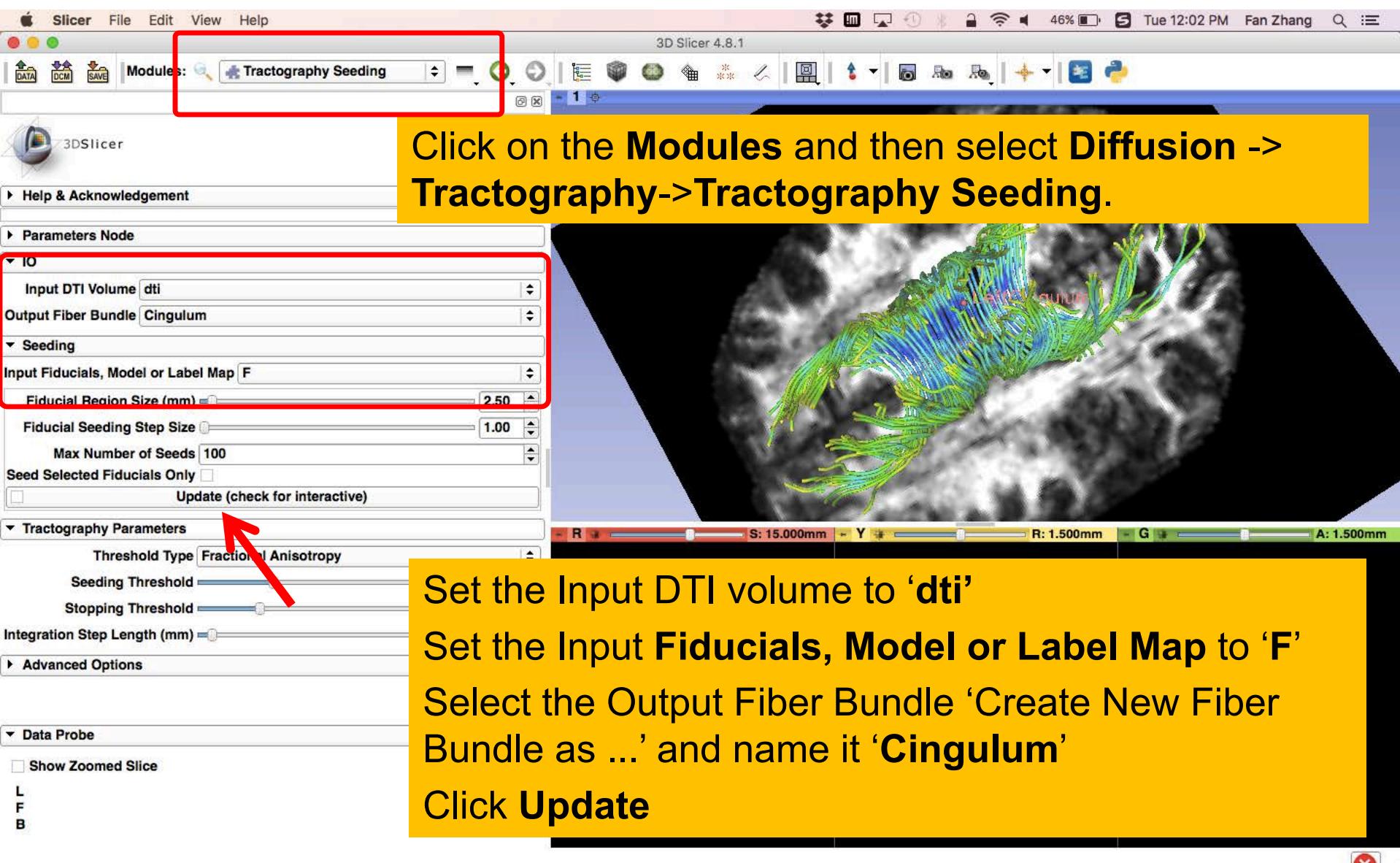
# Fiducial Seeding



# Fiducial Seeding



# Fiducial Seeding



Click on the **Modules** and then select **Diffusion -> Tractography->Tractography Seeding**.

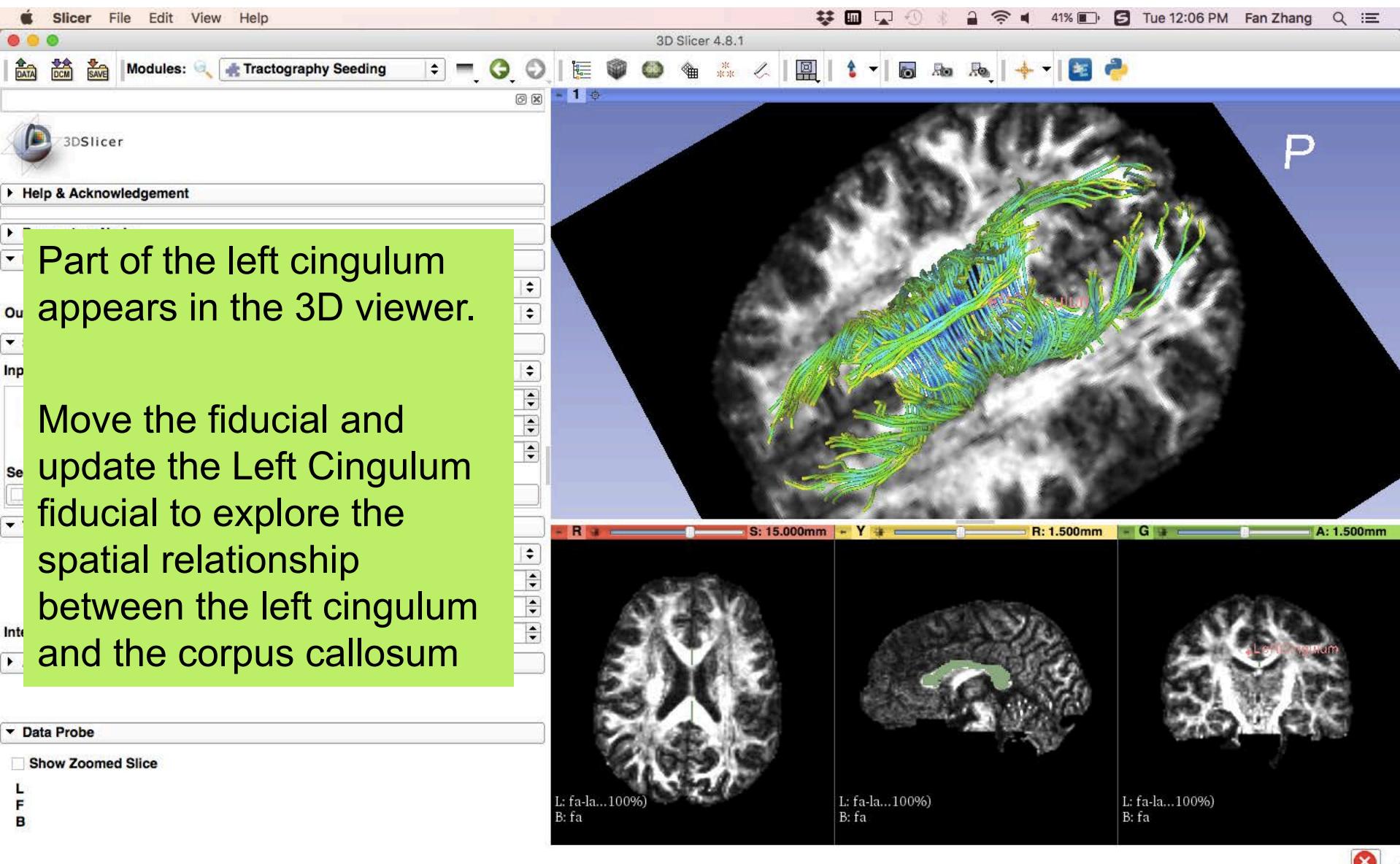
Set the Input DTI volume to 'dti'

Set the Input **Fiducials, Model or Label Map** to 'F'

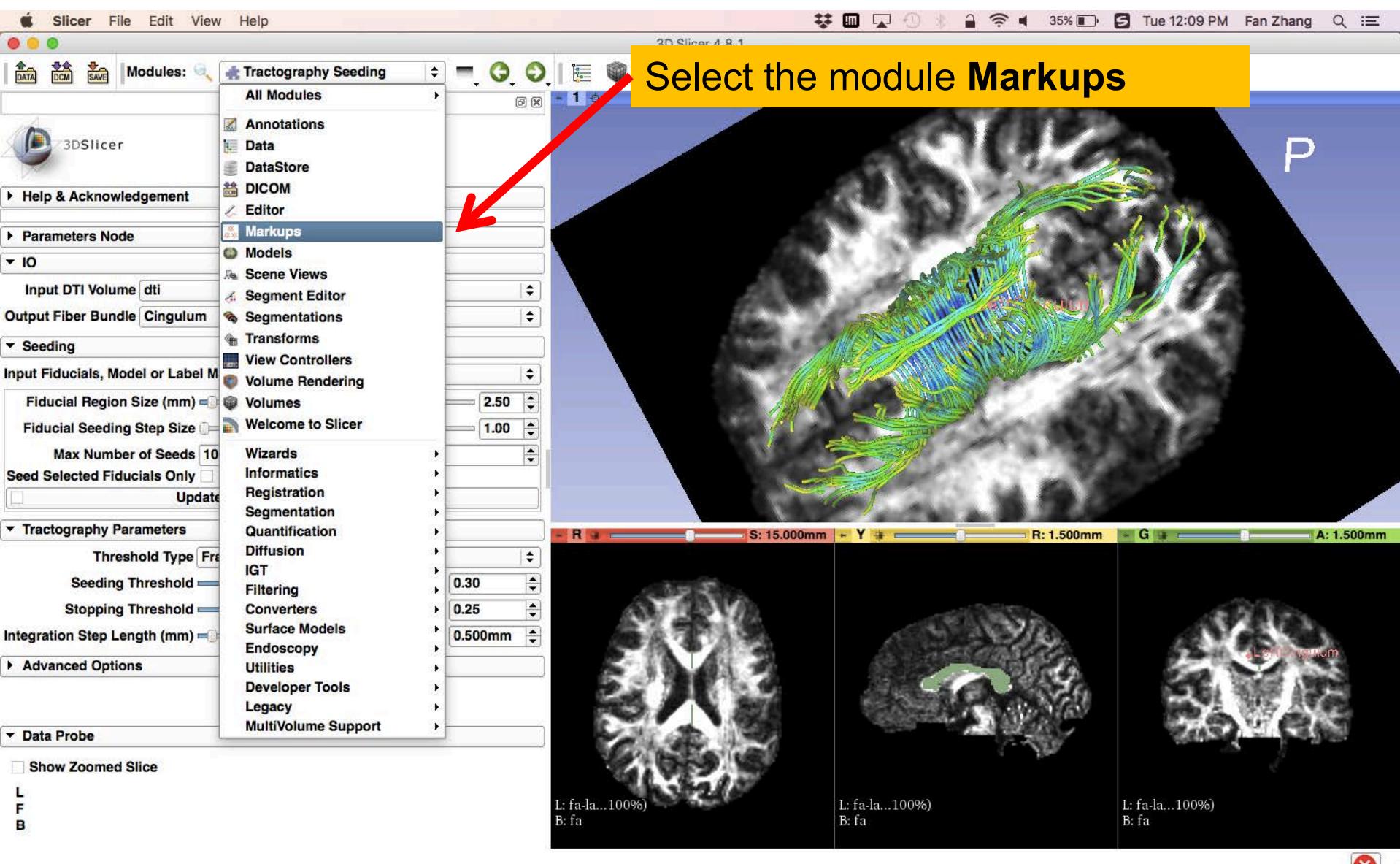
Select the Output Fiber Bundle 'Create New Fiber Bundle as ...' and name it '**Cingulum**'

Click **Update**

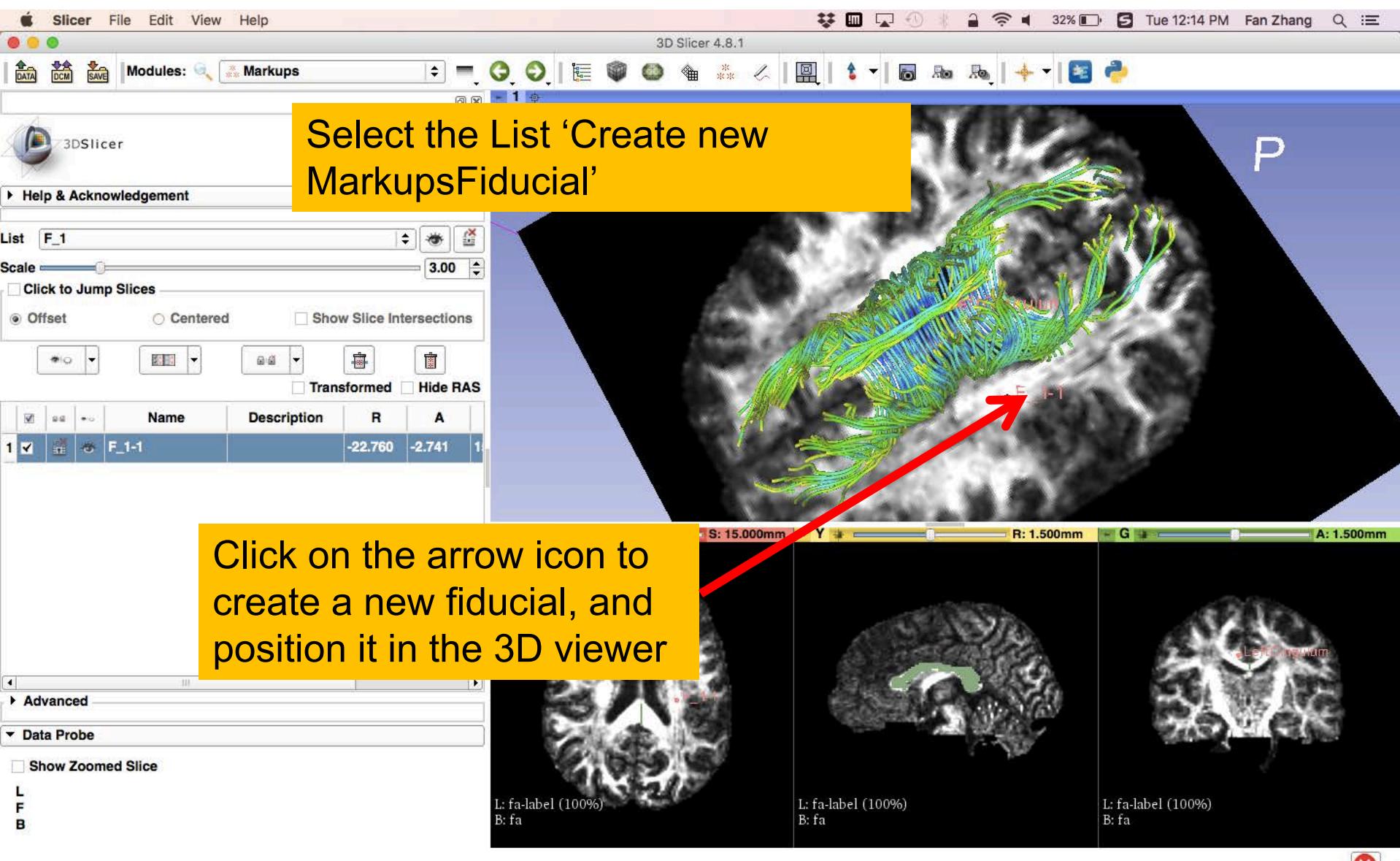
# Fiducial Seeding



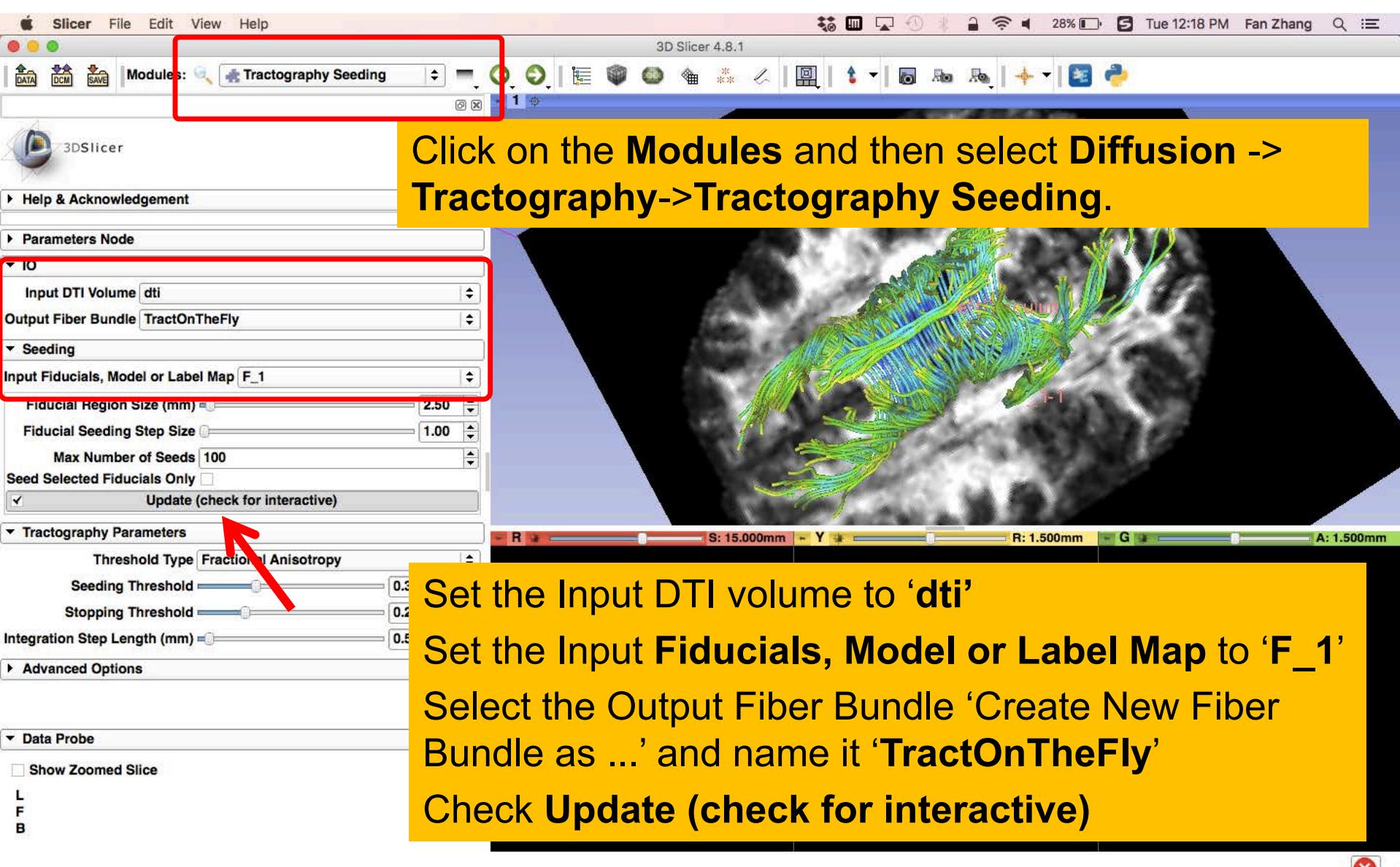
# Tractography 'on-the-fly'



# Tractography 'on-the-fly'



# Tractography 'on-the-fly'



Click on the **Modules** and then select **Diffusion -> Tractography->Tractography Seeding**.

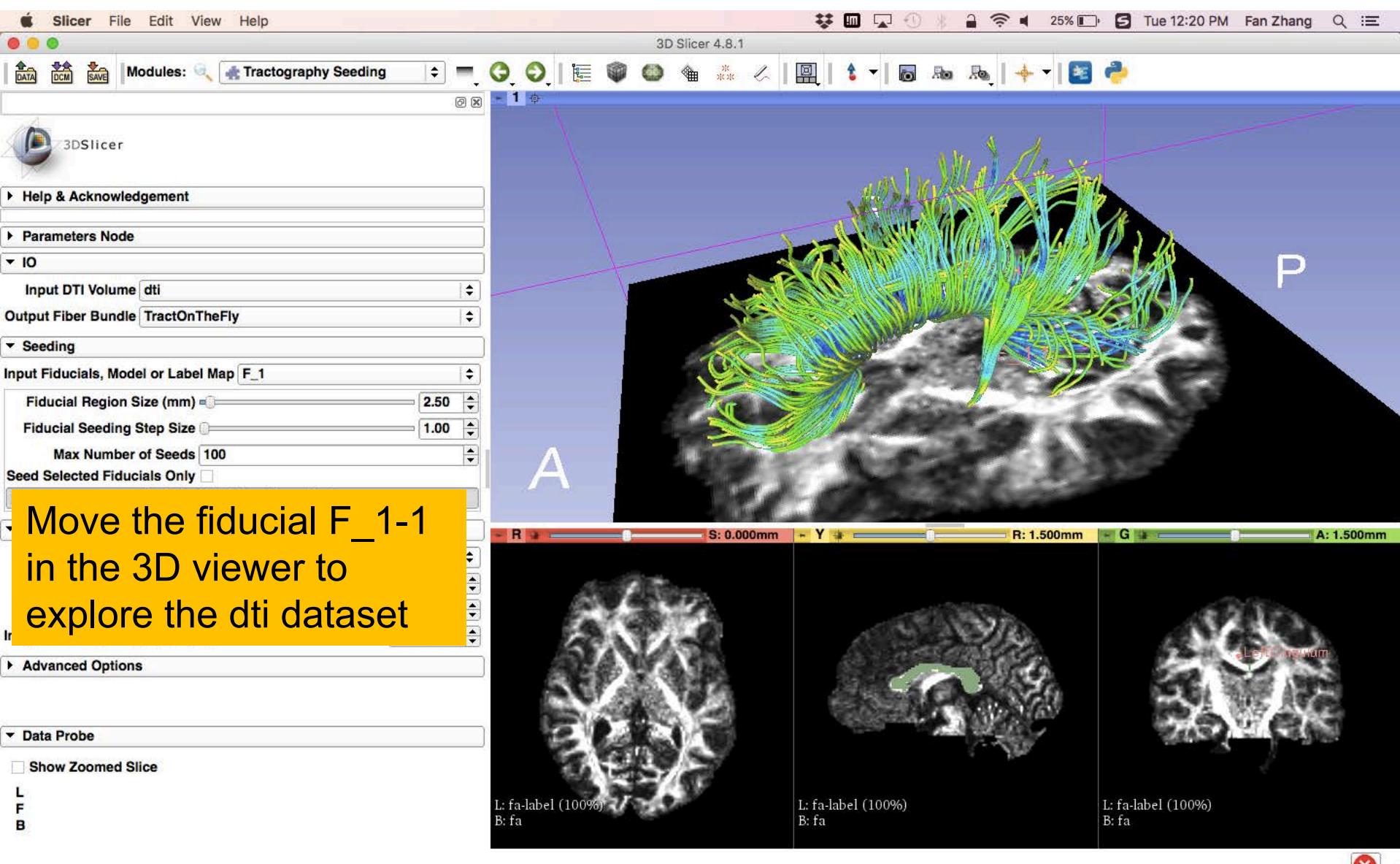
Set the Input DTI volume to '**dti**'

Set the Input **Fiducials, Model or Label Map** to '**F\_1**'

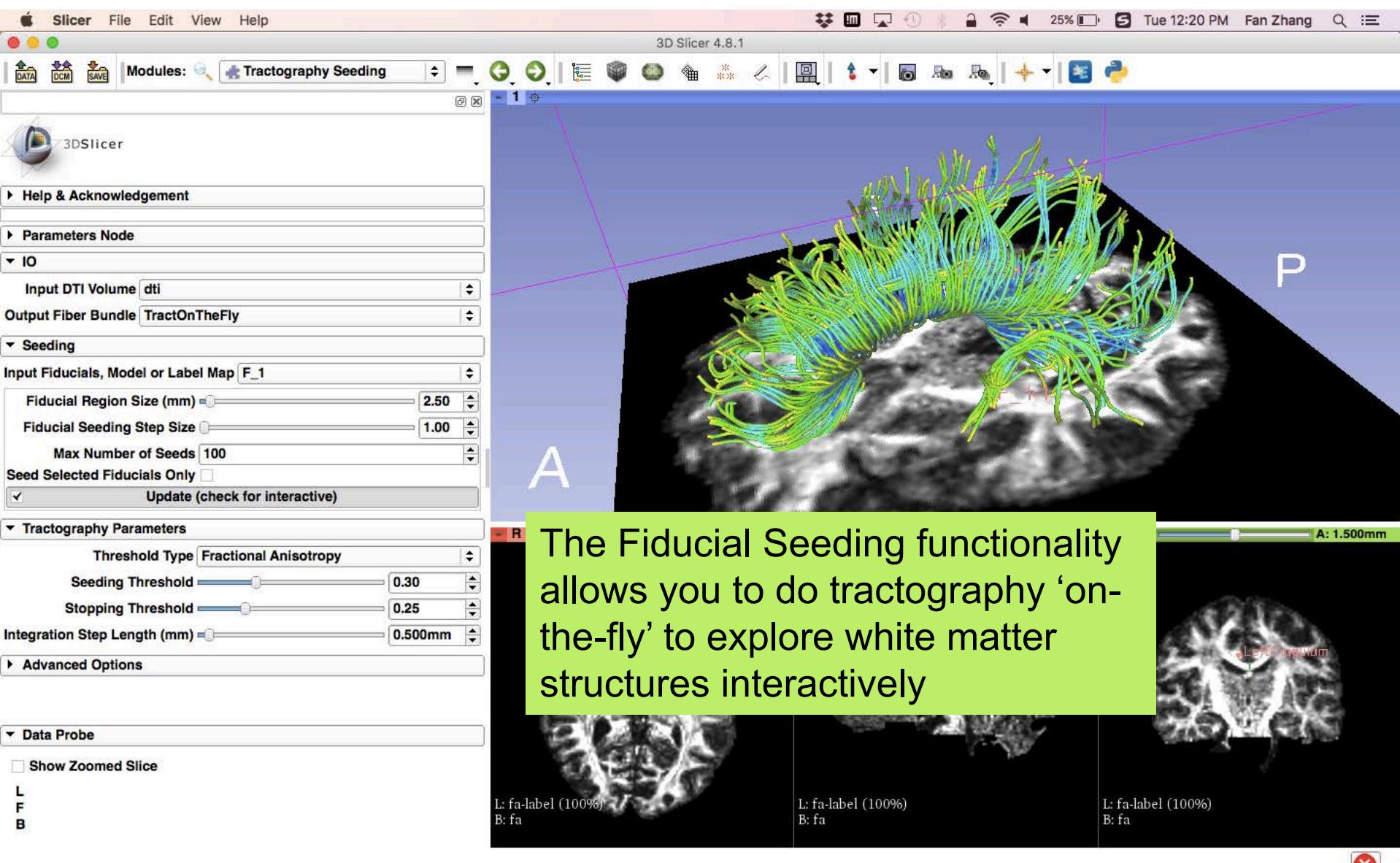
Select the Output Fiber Bundle 'Create New Fiber Bundle as ...' and name it '**TractOnTheFly**'

Check **Update (check for interactive)**

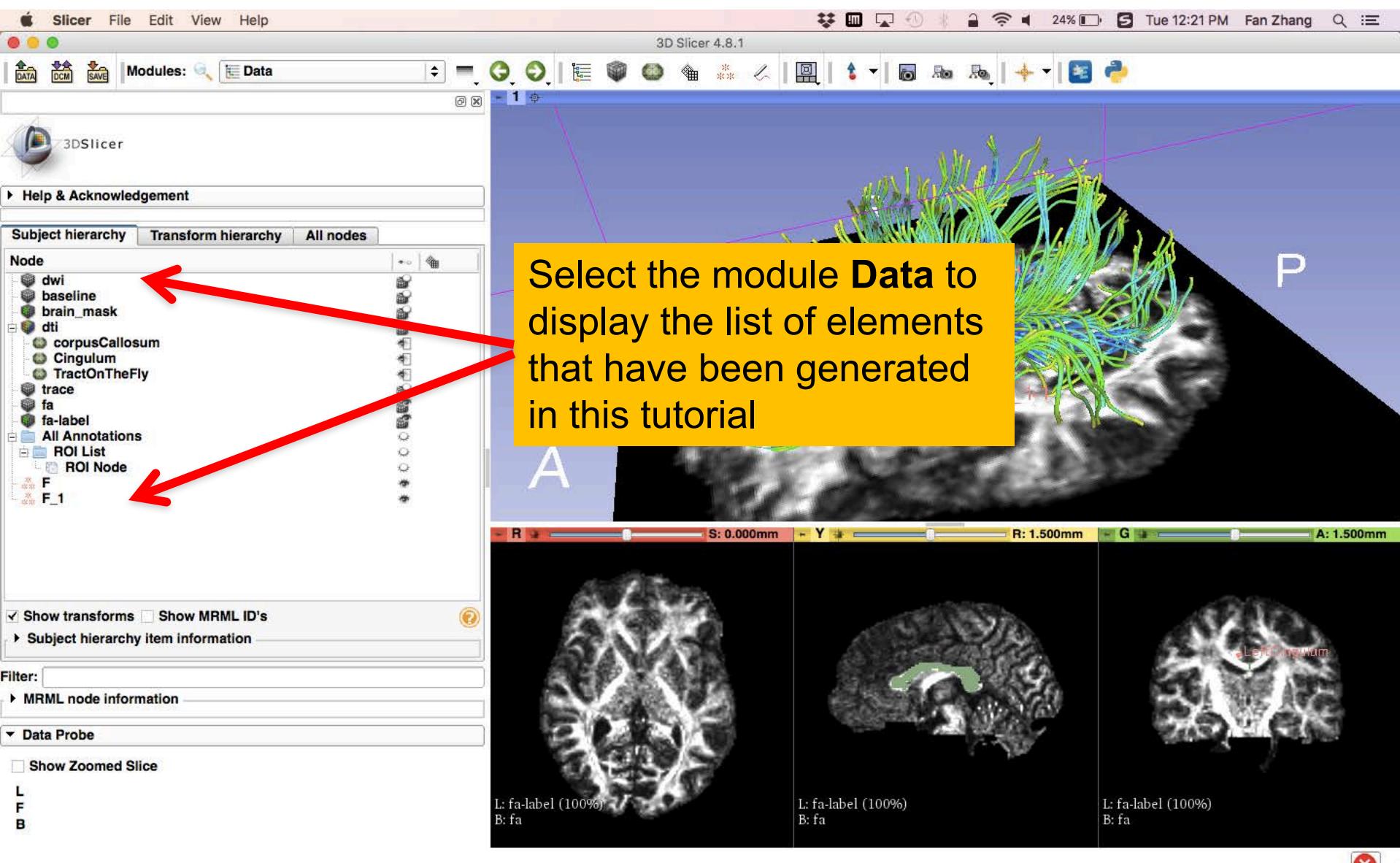
# Tractography 'on-the-fly'



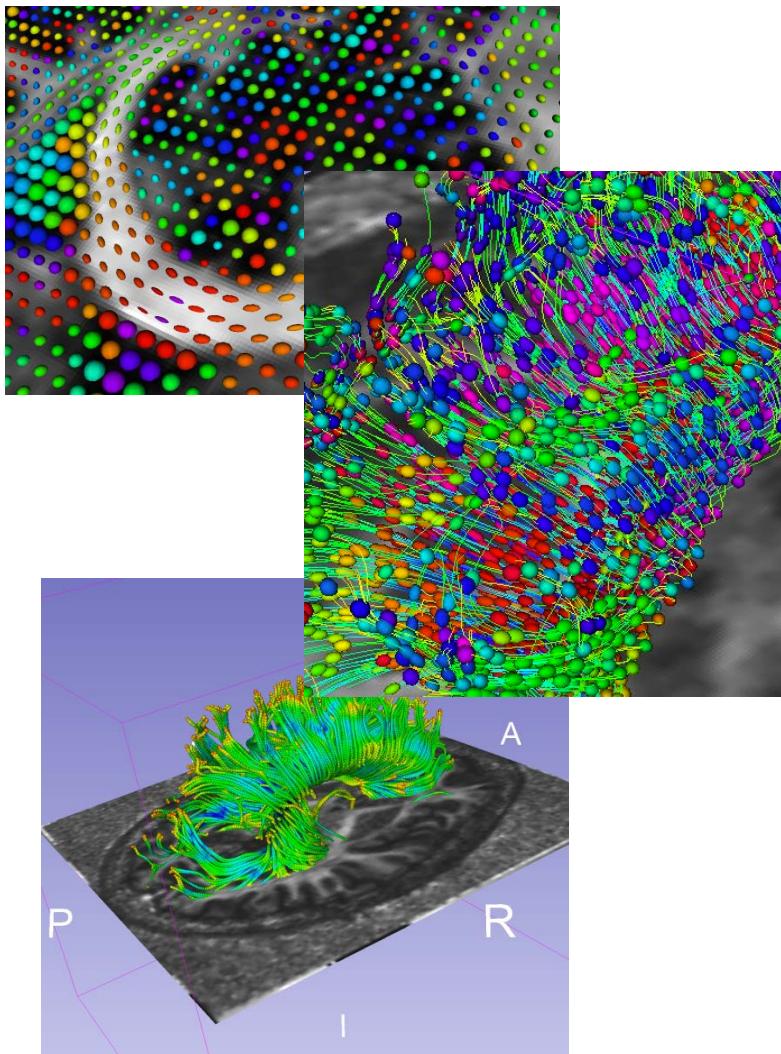
# Tractography 'on-the-fly'



# DTI Analysis



# Conclusion



This tutorial guided you through the different steps of a Diffusion MR analysis pipeline, from tensor estimation to 3D tracts visualization, for exploring and studying the 3D architecture of the brain white matter.

# Acknowledgments



- **Open Source Diffusion MRI Technology For Brain Cancer Research** NIH U01CA199459
- **National Center for Image Guided Therapy (NCIGT)**  
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NIH P41EB015902
- **Fan Zhang, Ph.D.**  
Brigham and Women's Hospital, Harvard Medical School

