

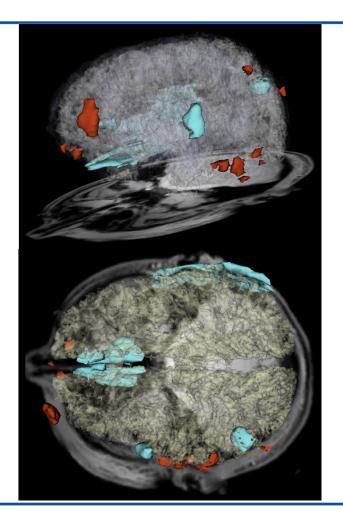
## Automatic Segmentation of Traumatic Brain Injury MRI volumes using Atlas Based Classification and 3D Slicer

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NA-MIC Tutorial Contest: Summer 2011

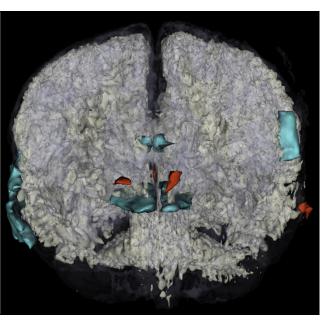


- acquire familiarity with several MR sequences commonly used for TBI imaging
- learn how to apply ABC to perform joint co-registration and automatic segmentation of TBI volumes
- acquire expertise on how to identify and characterize TBI pathology using various MRI sequences
- gain exposure to informed strategies for quantification of TBI-related edema or hemorrhage





- This tutorial assumes that you have already completed the following tutorials:
- Slicer 3 Visualization Tutorial by Sonia Pujol, available at
  - <u>http://www.slicer.org/slicerWiki/images/2/2e/Slicer3\_Data</u>
     <u>LoadingAndVisualization\_UCSF2010\_SoniaPujol.pdf</u>
- Interactive Editor Tutorial by Sonia Pujol, available at
  - <u>http://www.slicer.org/slicerWiki/images/6/69/InteractiveEditorT</u> utorial\_Slicer3.6-SoniaPujol.pdf



Edema (teal) and hemorrhage (red) in a traumatic brain injury patient, with white matter (solid beige) and gray matter (transparent hue) superposed.



This tutorial requires the installation of the **Slicer3.6 release** and the tutorial dataset. They are available at the following locations:

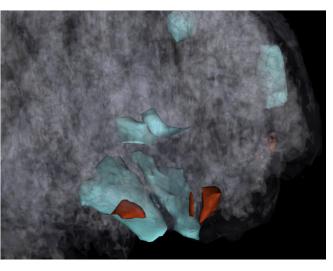
## Slicer3.6 download page

http://www.slicer.org/pages/Downloads/

### **Tutorial dataset:**

TBISegmentationData\_TutorialContestSummer2011.zip

http://www.na-mic.org/Wiki/images/7/74/TBISegmentationData TutorialContestSummer2011.zip

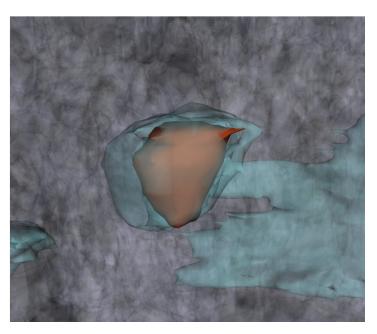


Enhanced view of frontal edema and hemorrhage several days after the insult. This type of pathology is very common in traumatic brain injury.

National Alliance for Medical Image Computing http://www.na-mic.org



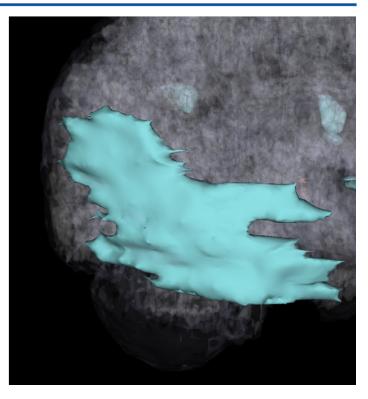
- This tutorial has been developed on Windows 7. It is compatible with the following platforms:
  - Windows XP, Windows 7
  - Linux 32, Linux 64
  - Mac/Darwin



Detailed view of parietal edema (teal) and hemorrhage. It is interesting to note how the non-hemorrhagic edemic region encases the brain portion that is actively bleeding. With recovery, it is expectable for the hemorrhage to shrink in both volume and mass.



- Clinical background
- Clinical workflow
  - installation of the ABC module
  - loading and exploration of TBI data
  - understanding common MR sequences for TBI
  - automatic segmentation of TBI using ABC
  - generation of 3D models
  - lesion segmentation and model
     generation
- Conclusions

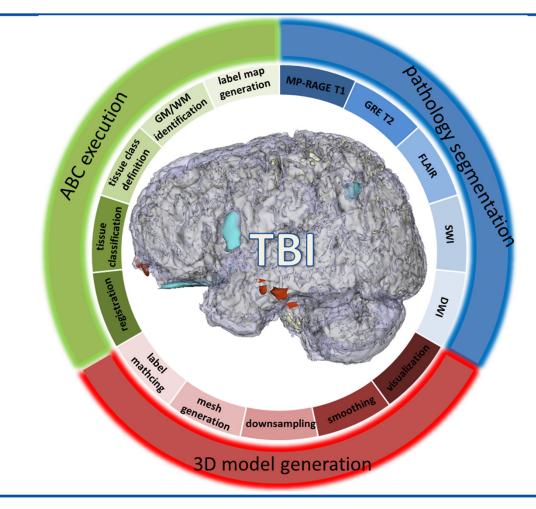


Contre-coup edema due to acceleration/deceleration forces at work during traumatic brain injury. Although contralateral with respect to the primary injury, the spatial extent of this edemic region is notable.



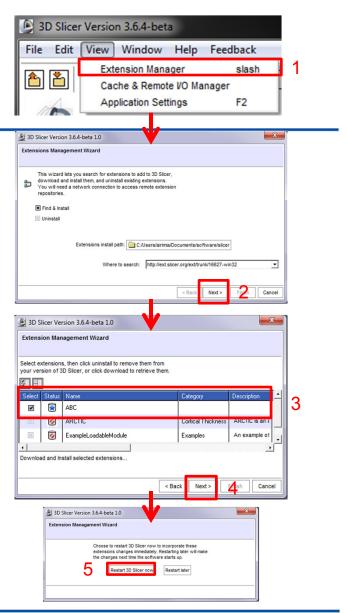
- With traumatic brain injury (TBI), there are an estimated 1.7 million cases in the United States alone every year, with an estimated 1.2 million ER visits and over 50,000 deaths.
- Over 5.3 million cases of required long-term daily assistance exist as a result of TBI, which results in health care costs of over \$60 billion/year
- Despite many innovations, progress towards patient-tailored characterization of the structural and functional substrates associated with TBI-related neural and cognitive impairment remains dissatisfactory and the relationship between neurophysiological markers of cognitive dysfunction and TBI structural damage has not been acceptably elucidated
- 3D Slicer offers a powerful and unparalleled set of tools for the exploration and quantification of TBI





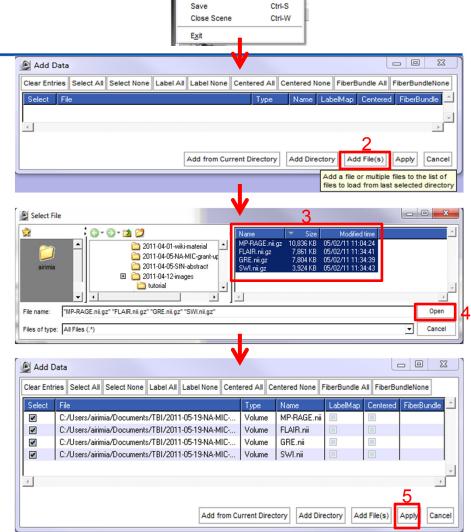


- After Slicer has been loaded, the Welcome window will appear.
- To install the ABC module:
  - 1 Click on View → Extension Manager
  - 2 Click on Next
  - 3 Select ABC from the list of modules
  - 4 Click on Next; you will be prompted to specify the local hard drive location for ABC
  - 5 After installation, you must select Restart 3D Slicer now for the changes to take effect





- After Slicer has been loaded, the Welcome window will appear.
- To load the TBI volumes associated with the case study:
  - 1 Click on File → Add Data
  - 2 Click on Add File(s)
  - 3 Navigate to the data folder and sort the volume files in descending order by size. This will ensure that all files are later processed in proper order.
  - 4 Click on Open
  - 5 In the Add Data dialog box, click on Apply



3D Slicer Version 3.6.4-beta
 File Edit View Window Help

CtrLO

Ctrl-A

Load Scene...

Import Scene.

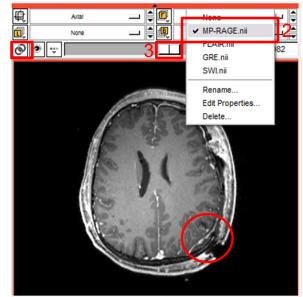
Add Volume...

Download Sample Da Add Data...

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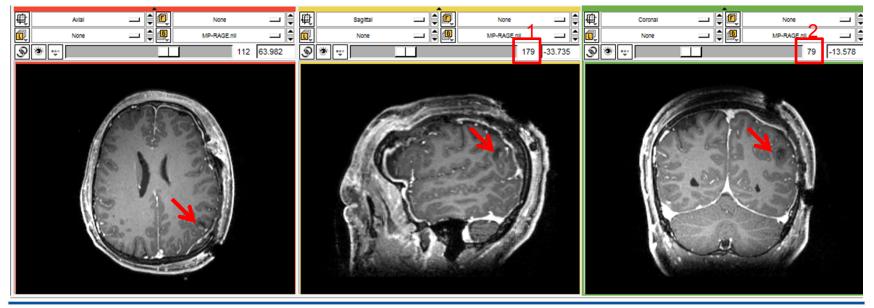


- 4 TBI data volumes are provided for this tutorial: MP-RAGE, FLAIR, GRE, and SWI.
- MP-RAGE: this is an anatomical volume acquired using a T1-weighted sequence, in which white matter has higher intensity than gray matter
  - To explore a TBI-related lesion in this patient:
    - 1 link the views using S
  - The lesion associated with open-head TBI becomes apparent as a hypo-intensity located in the parieto-occipital region (see circle to the right)

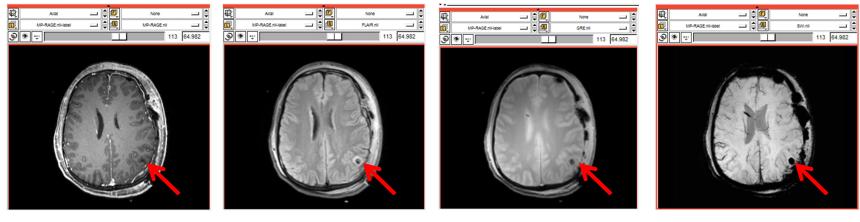




- To locate the lesion more accurately within the head:
  - 1 navigate to slice 179 in the sagittal view
  - 2 navigate to slice 79 in the coronal view
- The lesion and open head injuries are now apparent on all three views
- Similar exploratory navigation can reveal the lesion in FLAIR, GRE, and SWI volumes







MP-RAGE T1

FLAIR

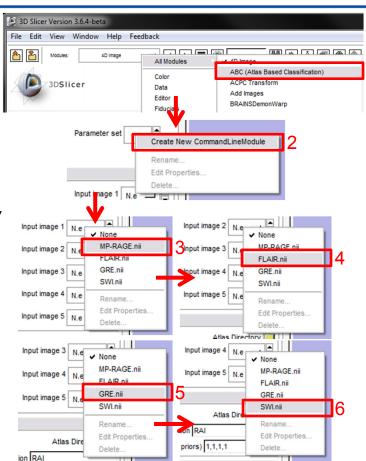
GRE T2

SWI

- Use of multiple image channels provides complementary information on pathologies
- T1 is a standard volume, with good anatomical resolution and high WM/GM contrast
- FLAIR is excellent for revealing CSF-perfused lesions as image hyper-intensities
- T2 GRE is useful for discerning areas associated with hemorrhages
- SWI is suitable for the detection of micro-bleeds and is superior to GRE in this respect

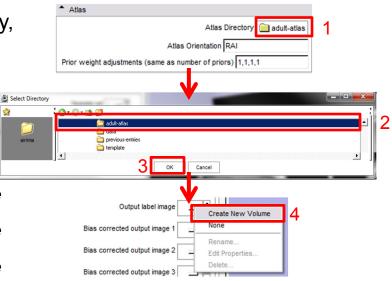


- To load ABC, click on:
  - 1 –Modules → All Modules → ABC
- The ABC interface appears. To specify input data:
  - 2 click Parameter Set → Create New Command Line Module
  - 3 click Input image 1 → MP-RAGE.nii
  - 4 click Input image 2 → FLAIR.nii
  - 5 click Input image 3 → GRE.nii
  - 6 click Input image 4 → SWI.nii





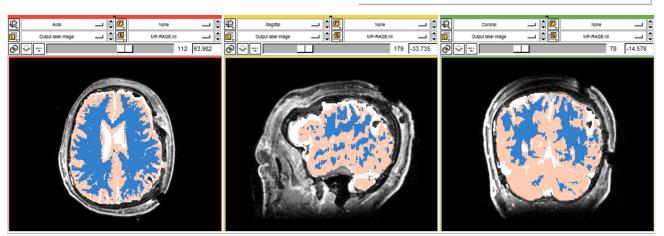
- 1 click on and locate the *adult-atlas* directory, which was included in the tutorial data set
- 2 select adult-atlas from its location
- 3 click OK to select the atlas template
- 4 click Output label image → Create New Volume
- 5 Click Bias corrected output image 1  $\rightarrow$  Create New Volume
- 6 Click Bias corrected output image 2 → Create New Volume
- 7 Click Bias corrected output image 3 → Create New Volume
- 8 Click Bias corrected output image 4  $\rightarrow$  Create New Volume

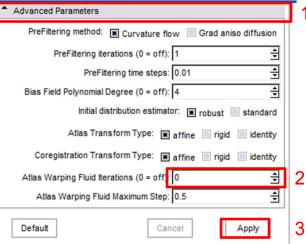


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- 1 click on the Advanced Parameters arrow
- 2 enter 0 (off) for the number of fluid iterations
- 3 click Apply to run ABC on the dataset
- ABC will now co-register the FLAIR, GRE and SWI images to the MP-RAGE image, and perform the tissue classification
- Allow up to several hours for ABC execution
- ABC produces 3 tissue categories:
  - gray matter
  - white matter
  - CSF







Now that we have obtained the label map of GM, WM and CSF using ABC, we can generate the 3D models associated with each of these three tissue types.

- 1 From the Modules drop-down menu, select All Modules
- 2 From the second column, select Model Maker

This will display the interface of the *Model Maker* module, which allows one to create 3D models in Slicer.

3D Slicer Version 3.6.4-bet	a		
File Edit View Window	Help Feedback		
Modules: UBC (A	All Modules	4D Image	Fast Rigid registration
-		ABC (Atlas Based Classification)	FastMarchingSegmentation
3DSlicer	Color	ACPC Transform	FetchMI
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$\overline{\gamma}$	Editor	BRAINSDemonWarp	Fiducial Registration
	Fiducials	BRAINSFit	FiducialSeeding
Help & Acknowledgement	Measurements	BRAINSROIAuto - Foreground masking tool	Fiducials
ABC (Atlas Based Classific	Models	BRAINSResample	Gaussian Blur
	ROI	BRAINSVectorDemonWarping	Gradient Anisotropic Diffusion
	SlicerWelcome	BSpline to deformation field	GradientAnisotropicFilter
	Slices	Binarize Map	Grayscale Fill Hole
	Transforms	Cast Image	Grayscale Grind Peak
Input Images	VolumeRendering	-	
	Volumes	ChangeTracker	Grayscale Model Maker
	Wizards	CheckerBoard Filter	Histogram Matching
	Informatics	ClipModel	IA_FEMesh
	Registration	CollectFiducials	IGTRecorder
	Registration	Color	IGTToolSelector
	Segmentation  Quantification	CongealingCLI	Image Label Combine
	Diffusion	Convert Fiducials to Labelmap	Joint Rician LMMSE Image Filter
	Diffusion	Create Single ROI file	Label Map Smoothing
	IGT •	Create a DICOM Series	LabelStatistics
	Filtering	CropVolume	Labelmap Seeding
•	Surface Models	Curvature Anisotropic Diffusion	Linear registration
▲ Atlas	Converters •	Data	MRI Bias Field Correction
	Endoscopy	Dicom to Nrrd Converter	Mask Image
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<ul> <li>Output Images</li> </ul>	Utilities	EMSegmenter	Model Maker
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		Endoscopy	ModelIntoLabelVolume
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		Expert Automated Registration	Models
Bi	as corrected output image	Explode Volume Transform	Multiply Images
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Bi	as corrected output image	Extract Skeleton	NeuroNav
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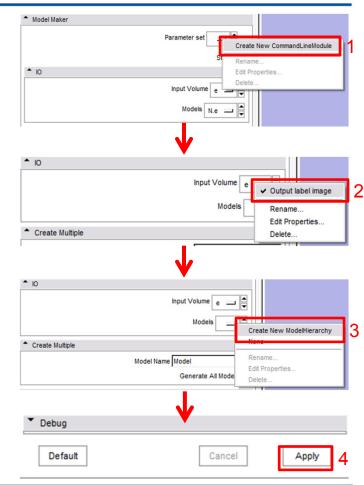


To generate the 3D models:

- 1 From the *Parameter Set* drop-down menu, select *Create New Command Line Module*
- 2 From the *Input Volume* drop-down menu, select *Output label image*
- 3 From the *Models* drop-down menu, select *Create New Model Hierarchy*

• 4 – Click Apply

These steps will create 3D models for WM, GM and CSF.

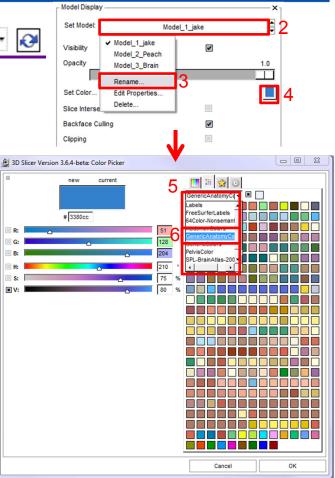






- 1 From the tool bar, click on the Models button
- 2 From the Set Model drop-down menu, select Model\_1\_jake
- 3 From the Set Model drop-down menu, select Rename and type WM; repeat steps 1-2 to rename Model\_2\_Peach as GM and Model\_3\_Brain as CSF
- 4 Set the model to WM as in step 2, and click on the color box;
- 5 In the Color Picker dialog box, click on the color drop down menu
- 5 Select Generic Anatomy Colors as the color map; this will assign a color scheme in accordance with Slicer conventions.





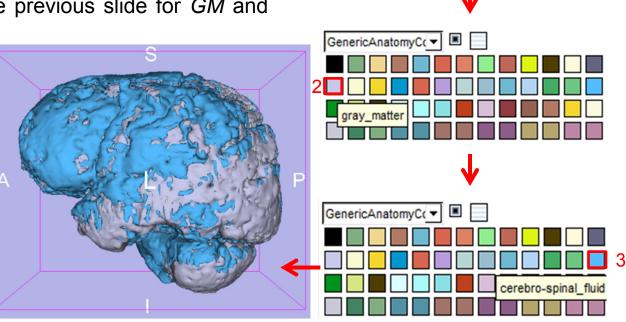


Now that we have selected an appropriate color scheme, we need to assign suitable colors to each tissue type.

 1 – In the Color Picker dialog box, select beige for WM

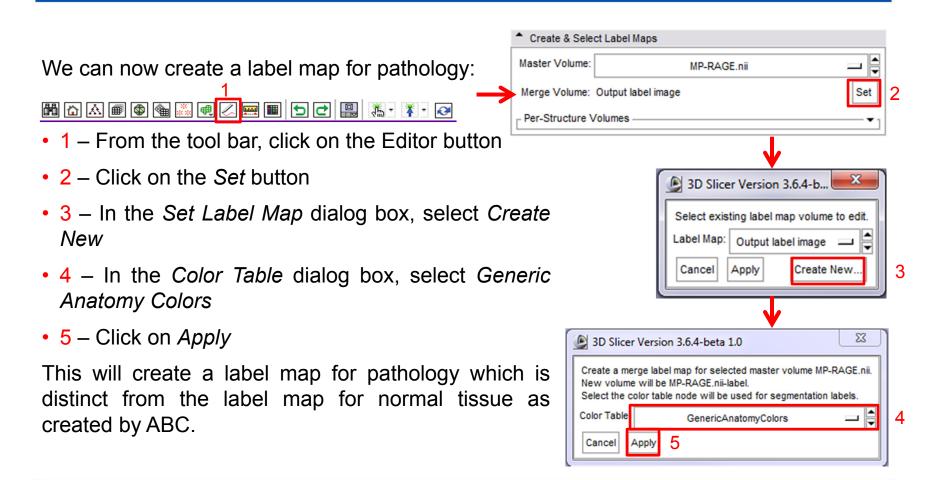
Repeat steps 2-5 on the previous slide for *GM* and *CSF* 

- 2 For *GM*, click on purple as shown
- 3 For *CSF*, click on blue as shown. The 3D model should now look as to the right. Note the variable CSF thickness.



GenericAnatomyC( -

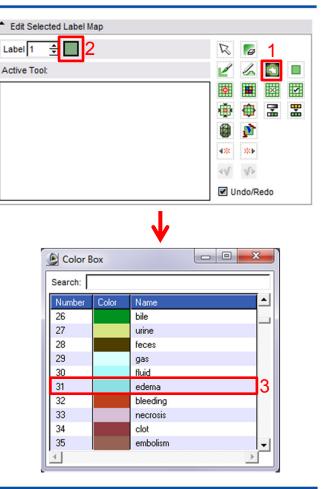






Now we can assign labels for the two pathology types, namely edema and bleeding:

- 1 From the label edit bar, click on *Level Tracing*
- 2 Click on the color button next to the Label
- 3 In the Color Box dialog box, select 31: edema





- 1 In the Foreground menu, select FLAIR.nii
- 2 Click on *the* tool bar menu
- 3 select *Red slice only*

#### Layout

 4 – Use the level tracing tool to segment the edema using the FLAIR volume.

2 🖳 🦾 🕻 🕻 🐼 Axial None MP-RAGE.nll-label ✓ MP-RAGE.nii ✓ I Conventional layout ଚ FLAIR.nii GRE nii Conventional widescreen layout SWI.nii Ø Four-up layout Output label image 88 Dual 3D layout Bias corrected output image 1 Bias corrected output image 2 Triple 3D layout Bias corrected output image 3 3D only layout Bias corrected output image 4 MP-RAGE.nii-label Red slice only layout 3 Rename... Yellow slice only layout Edit Properties... Green slice only layout Delete Tabbed 3D layout Tabbed slice layout Compare layout Compare widescreen layout Side-by-side lightbox layout

Toggle GUI panel visibility

Toggle GUI panel L/R

close

Recall that edema is hyperintense in this sequence modality, and that it appears in slices 105-117. The hypointense region in the middle is actually hemorrhage, but it will be segmented separately in following steps so we need not worry about it for now.

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Once the edema has been segmented, we can proceed to the segmentation of hemorrhage. Viewing the label map and the underlying image at the same time can be accomplished by adjusting label map opacity.

- 1 From the red slice image options menu, click on Adjust label map opacity
- 2 Use the slider to select a convenient opacity
- 3 In the Color Box dialog box, select 32: bleeding

This step ensures that 3D Slicer color labeling conventions are respected.



Label 1

Active Tool:

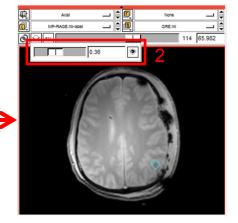
GREN

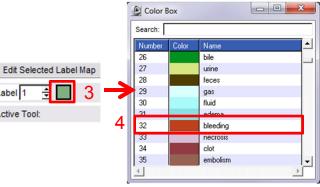
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MP-RAGE nil-label

114 65.982







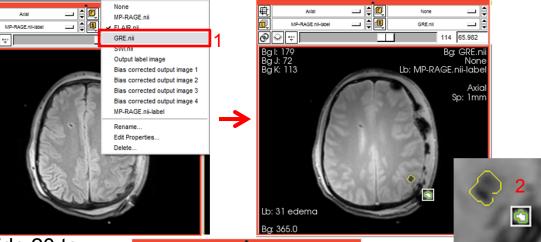
We can now segment bleeds.

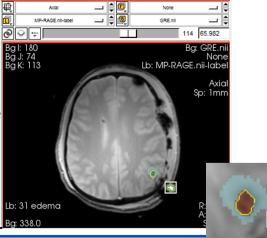
• 1 – From the red slice background menu, click on *GRE.nii* 

 2 – Use the label map opacity slider as shown on the previous slide as well as the level

tracing tool as demonstrated on slide 20 to obtain a convenient view of the hemorrhage in relation to the surrounding edema.

3 – Segment the bleeding in slices 106-116. The label map should now be similar to the inset shown to the right. Note that hemorrhage is surrounded by edemic region which does not bleed.





# Create pathology 3D models

Once pathology has been segmented, their 3D models can be created.

- 1 From the tool bar menu, select the Conventional layout
- 2 For Input Volume, select MP-

*RAGE.nii-label.* This is the label map for the pathology.

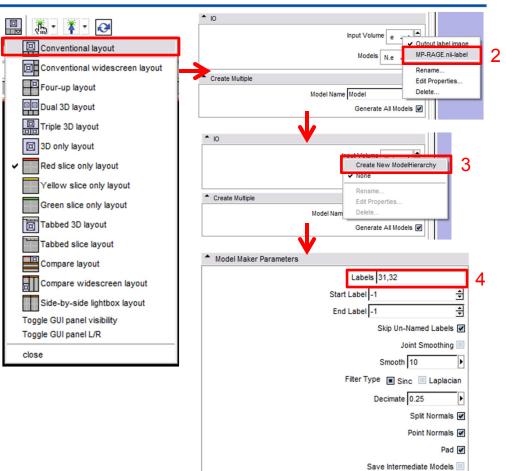
• 3 – Under Models, select Create

New Model Hierarchy

• 4 – Under Model Maker Parameters,

type "31,32" in the Label input box.

• 5 – Click on Apply





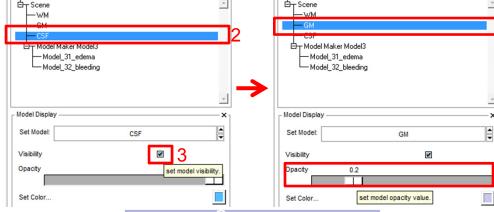
Hierarchy & Display

Model Hierarch

We can now visualize the full model, which includes both pathology and healthy-looking tissue.

☆ & # \$ 

- 1 On the tool bar, click Models
- 2 Under Hierarchy and Display, select CSF



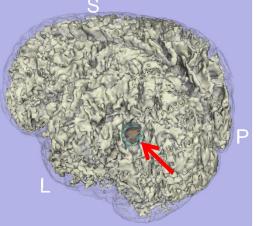
Hierarchy & Displa

5

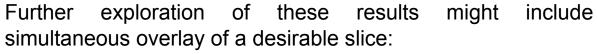
Model Hierarchy

- 3 Under *Visibility*, uncheck the toggle box
- 4 Under *Hierarchy and Display, select GM*
- 5 Decrease the opacity to 0.2

This will allow us to visualize the edema and hemorrhage in relationship to the rest of the brain, as shown to the right.



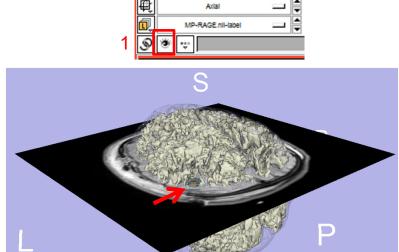


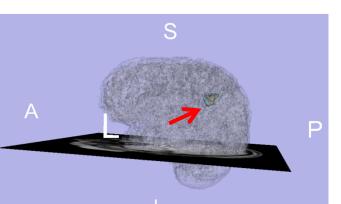


1 – On the red box tool bar, toggle slice visibility (left below)

One can also display the pathology with GM/WM transparency (see figure to the right below)

• 2 – Under Hierarchy and Display, set WM opacity to 0.2



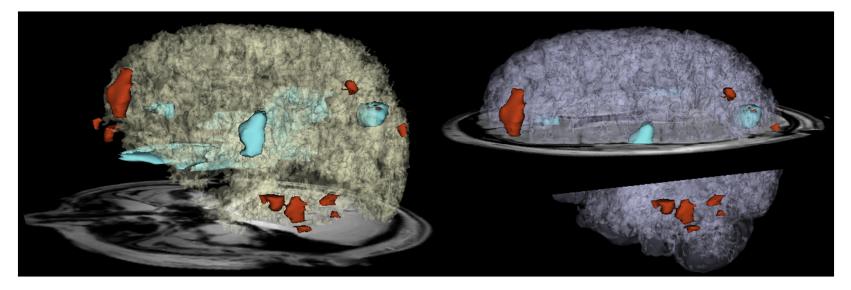


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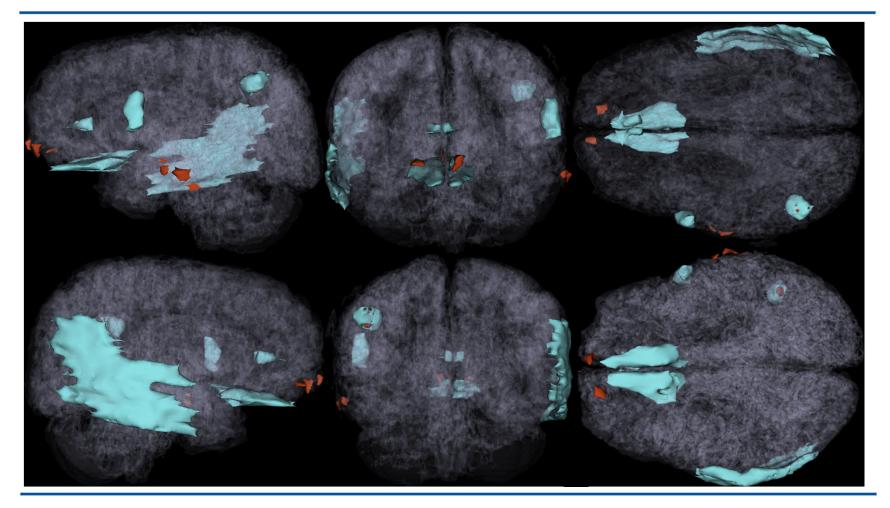
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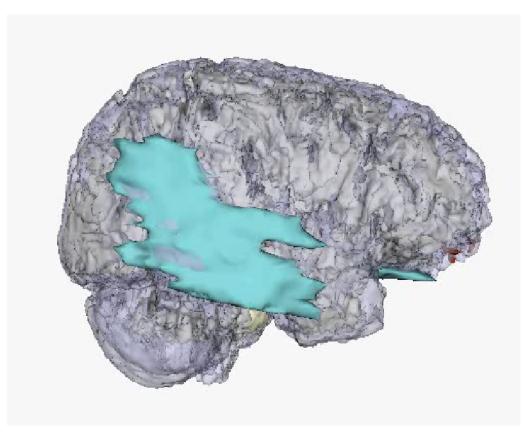
Segmentation of remaining lesions and hemorrhages in this subject can be accomplished using the tools and workflows demonstrated in this tutorial. Sample images of this undertaking are illustrated below.













- 3D Slicer offers powerful methodologies for the visualization of pathology due to traumatic brain injury
- Use of multiple MR image channels greatly enhances the ability to study and understand TBI structure/extent
- ABC is a robust algorithm to perform joint coregistration and automatic segmentation of TBI
- Completion of this tutorial allows one to acquire useful expertise on how to identify and characterize TBI
- Use of 3D Slicer can offer informed strategies for quantification of TBI-related edema or hemorrhage and for improved insight of clinical relevance



The members of the UCLA TBI team would like to convey our special gratitude to Dr. Ron Kikinis for his excellent support, direction and leadership of the concerted efforts that have made this work possible.



#### **National Alliance for Medical Image Computing** NIH U54EB005149 (PI: Ron Kikinis MD; TBI sub-award: Jack Van Horn PhD)



## National Institutes of Health

#### National Institute of Neurological Disorders and Stroke

NATIONAL INSTITUTE OF INCLURAL OGICAL DISORDERS AND STROKE NIH P01NS058489 (PI: Paul Vespa MD, FAAN, FCCN)



#### Laboratory of Neuro Imaging

University of California, Los Angeles

Director: Arthur W. Toga PhD



- A Irimia, MC Chambers, JR Alger, M Filippou, MW Prastawa, B Wang, DA Hovda, G Gerig, AW Toga, R Kikinis, PM Vespa, JD van Horn (2011) Comparison of acute and chronic traumatic brain injury using semi-automatic multimodal segmentation of MR volumes *submitted*
- A Irimia, MC Chambers, M Filippou, JR Alger, MW Prastawa, B Wang, S Gouttard, SMA Pujol, SR Aylward, DA Hovda, G Gerig, AW Toga, R Kikinis, PM Vespa, JD van Horn (2011) Threedimensional calculation and quantification of morphometric and volumetric cortical atrophy indices of widespread clinical use from MRI volumes of traumatic brain injury using 3D Slicer *Proceedings of the 41<sup>st</sup> Annual Meeting of the Society for Neuroscience (SfN 2011), Washington, DC (this work was honored with the 1<sup>st</sup> Prize in the Fine Science Tools Research Contest of the Brain Research Institute at UCLA)*
- A Irimia, JD van Horn, MC Chambers, MW Prastawa, S Gouttard, PM Vespa, DA Hovda, JR Algers, SMA Pujol, G Gerig, SR Aylward, AW Toga, R Kikinis (2011) Automatic multimodal MR image segmentation for the clinical assessment of traumatic brain injury in 3D Slicer *Proceedings of the 17<sup>th</sup> Annual Meeting of the Organization on Human Brain Mapping (OHBM 2011), Quebec City, Canada*



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