



# *Slicer3 Training Compendium*

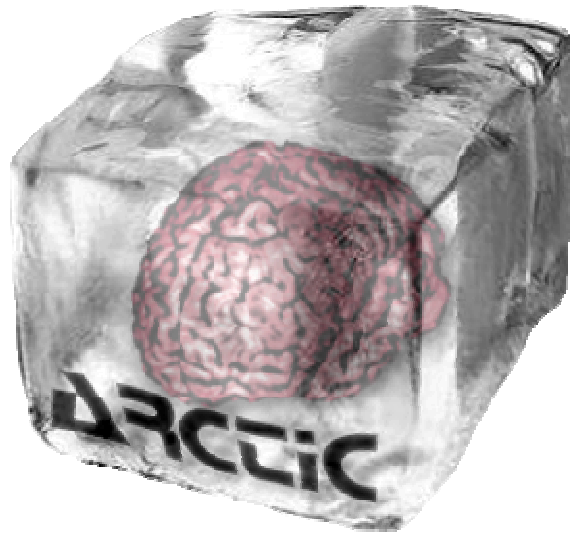
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Slicer3 Training Tutorial

# ARCTIC

(Automatic Regional Cortical ThIckness)



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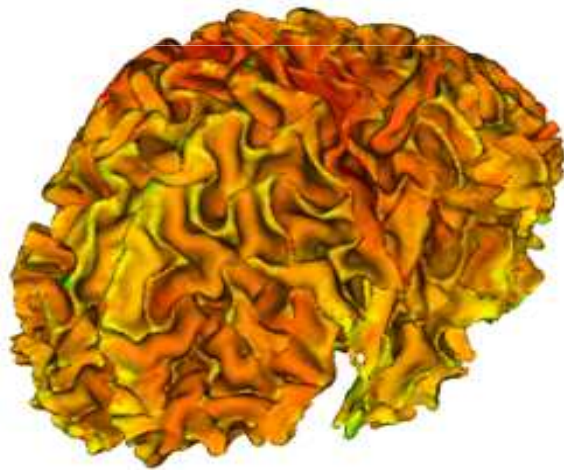


# Learning Objective

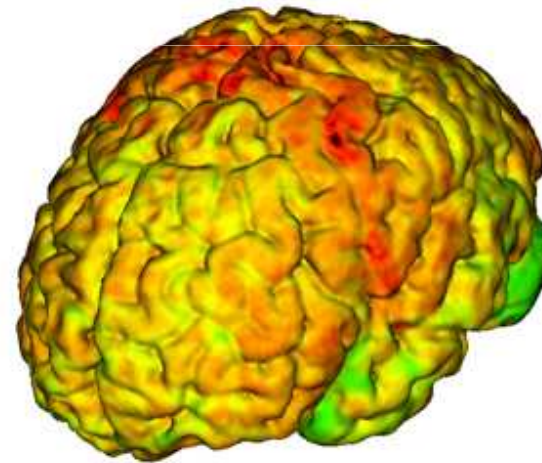
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Following this tutorial, you will be able to perform an individual analysis of regional cortical thickness.

You will learn how to **load input volumes**, **run the end-to-end module ARCTIC** to **generate cortical thickness information** and **display output volumes**.



*Cortical thickness on WM surface*



*Cortical thickness on GM surface*



# *Prerequisites*

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This tutorial assumes that you have already completed the tutorial **Data Loading and Visualization**.

Tutorials for **Slicer3** are available at the following location:

- **Slicer3** tutorials

<http://www.na-mic.org/Wiki/index.php/Slicer3.2:Training>



# Materials

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This tutorial requires the installation of Slicer3, BatchMake, BatchMake wrappers, the tutorial dataset and the external modules. They are available at the following locations:

- Slicer3 download page (***Slicer 3.2***)  
<http://www.slicer.org/pages/Downloads>
  - BatchMake download page (***BatchMake 1.0.6***)  
<http://www.batchmake.org/batchmake/resources/software.html>
  - BatchMake wrapper download page (***ARCTIC\_BatchMake\_Wrapper\_1.0***)
  - Tutorial dataset download page(***ARCTIC\_Tutorial\_example\_1.0***)
  - External modules download page (***ARCTIC\_Executables\_1.0***)  
<http://www.nitrc.org/projects/arctic/>
  - Atlas download page(***UNC\_Pediatric\_Brain\_Atlas***)  
<http://www.insight-journal.org/midas/item/view/2277>
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**Disclaimer:** *It is the responsibility of the user of Slicer to comply with both the terms of the license and with the applicable laws, regulations, and rules.*



# *Materials: Tutorial dataset*

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The tutorial dataset (*ARCTIC\_Tutorial\_example\_1.0*) is a ZIP file.

Unzip this file somewhere in your computer.

An “*ARCTIC\_Tutorial\_example\_1.0*” folder will be created, containing:

- A pediatric case: T1-weighted and T2-weighted images.
- An “ARTIC-Results/” directory, in which results of the tutorial example will be saved.



# *Materials: External modules*

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The executables are in a ZIP file (*ARCTIC\_Executables\_1.0\_linux32/64*).

Unzip this file somewhere in your computer.

An “*ARCTIC\_Executables\_1.0\_linux32/64*” folder will be created, containing executables needed to perform the cortical thickness analysis.

To add the executables as Slicer3 external modules:

- Open Slicer3
- Go to View → Application Settings → Module Settings
- Click on the “add a preset” button
- Select the “*ARCTIC\_Executables\_1.0*” folder and confirm
- Close Slicer3



# *Materials: Atlas*

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The atlas and its related files are in a ZIP file (*UNC\_Pediatric\_Brain\_Atlas*).

Create a “pediatric-atlas-4years-sym-T1-RAI” folder somewhere in your computer.

Unzip the ZIP file in this new folder.

The “pediatric-atlas-4years-sym-T1-RAI” folder will thus contain the atlas and its related files.

You can then unzip all the images (gunzip command).



# Prerequisites

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## Add the executables in the PATH.

**-tcsh usage :** setenv PATH ARCTIC-Executables-Directory/:Slicer3D-Plugins-Directory/:Batchmake-Directory/:\${PATH}

**-bash usage :** export PATH=ARCTIC-Executables-Directory/:Slicer3D-Plugins-Directory/:Batchmake-Directory/:\${PATH}

Notice : To execute ARCTIC within Slicer3, it is not necessary to add "Slicer3D-Plugins-Directory/" in the PATH.

## Set ARCTIC environment variable

**-tcsh usage :** setenv BatchmakeWrapper\_Dir Batchmake-Wrapper-Directory/

**-bash usage :** export BatchmakeWrapper\_Dir=Batchmake-Wrapper-Directory/

### WITH:

ARCTIC-Executables-Directory/ : Downloaded folder (ARTIC\_Executables\_1.0)

Slicer3D-Plugins-Directory/ : Directory containing Slicer3 plugins

Release: "Slicer3Dir"/lib/Slicer3/Plugins

Compiled Version: "Slicer3Dir"/Slicer3-build/lib/Slicer3/Plugins

Batchmake-Directory/ : Directory containing BatchMake application

Batchmake-Wrapper-Directory/ : Downloaded folder (ARTIC\_Batchmake\_Wrapper\_1.0)





# Overview

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- 1- Pipeline overview
- 2- Input images
- 3- Pipeline description
- 4- Output images and organisation
- 5- Execution within Slicer
- 6- Example with tutorial dataset
- 7- Command line execution



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# *Pipeline Overview*

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All the tools used in the current pipeline are Slicer3 modules, some of them being UNC external modules. The user can thus perform a regional cortical thickness analysis on an individual subject within Slicer3.

**Two different modes** can be used, depending on the input images:

- Raw images (T1-weighted, T2-weighted, PD)
- Tissue segmentation label image



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## What you need...

### *Raw images*

T1-weighted image  
Tissue segmentation atlas directory

#### Optional

T2-weighted image  
PD-weighted image  
Atlas raw image + its parcellation  
Case parcellation image

### *Segmented image*

Raw image  
Tissue segmentation label image

#### Optional

Atlas raw image + its parcellation  
Case parcellation image

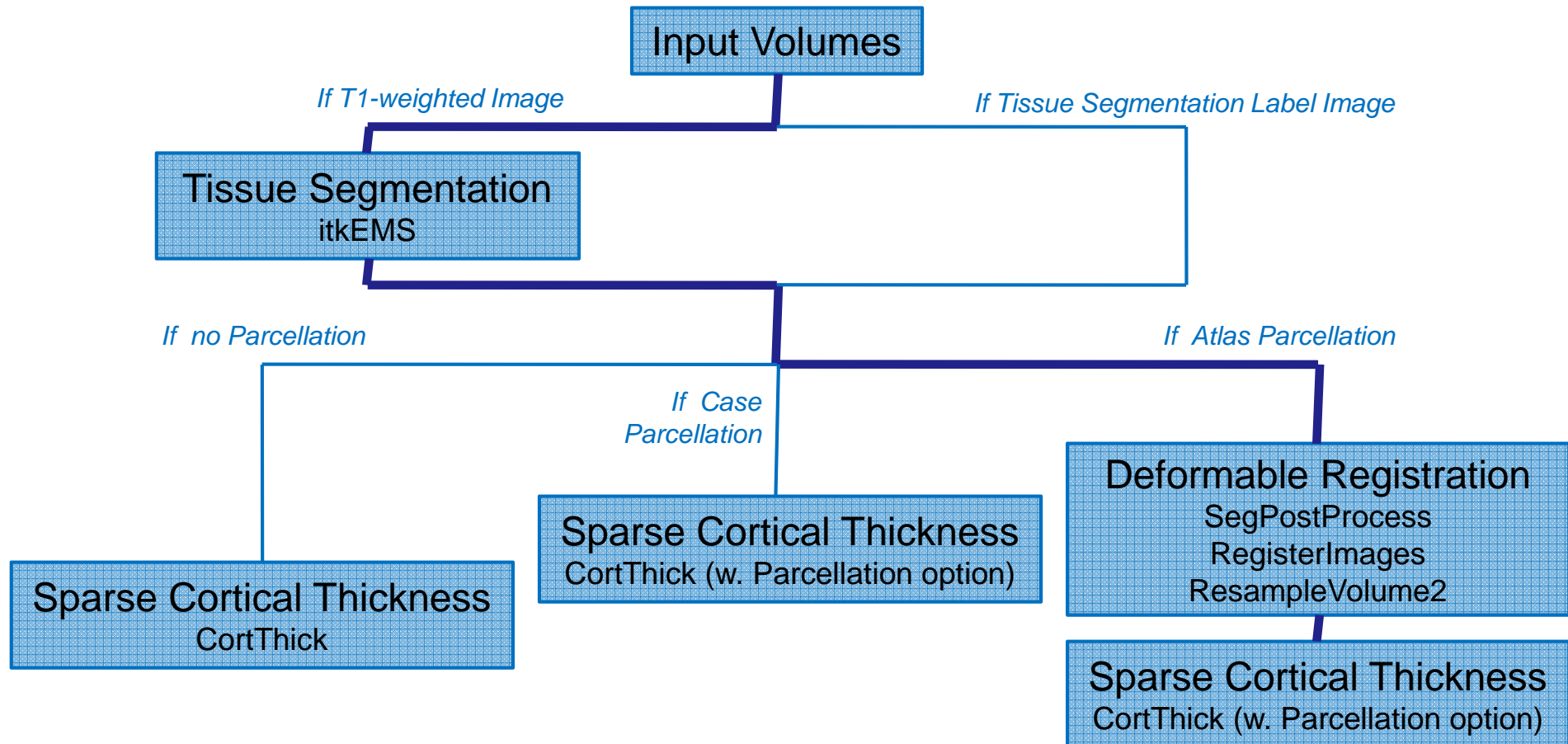


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# Pipeline Description





# *Pipeline Description*

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## **1. Tissue segmentation**

Module : itkEMS (UNC Slicer3 external module)

## **2. Regional atlas deformable registration**

### 3.1. Skull stripping

Module : SegPostProcess (UNC Slicer3 external module)

### 3.2. Deformable registration of T1-weighted atlas

Module : RegisterImages (Slicer3 module)

### 3.3. Applying transformation to its parcellation map

Module : ResampleVolume2 (Slicer3 module)

## **3. Sparse and asymmetric Cortical Thickness**

Module : CortThick (UNC Slicer3 module)

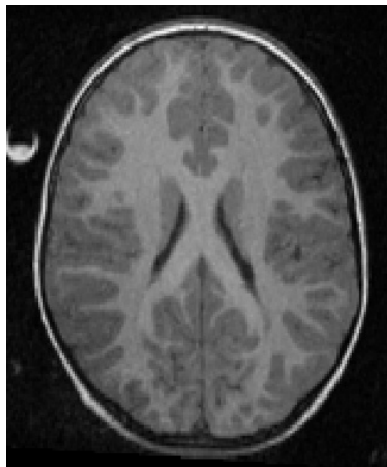


# Pipeline Description

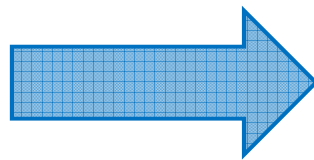
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## Tissue segmentation (itkEMS external module)

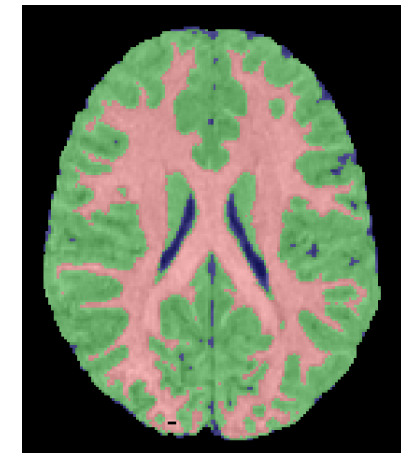
Probabilistic atlas-based automatic tissue segmentation via an Expectation-Maximization scheme. ItkEMS also performs an intensity inhomogeneity correction of the input image that removes gradual variations in the image intensities mainly due to RF coil imperfection



*Input\_T1-Image.gipl*



*Image\_corrected\_EMS.gipl*



*Image\_labels\_EMS.gipl*

# *Pipeline Description*

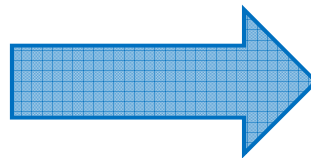
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## **Skull Stripping (SegPostProcess external module)**

This step is performed using the previously computed tissue segmentation label image.



*Image\_corrected\_EMS.gipl*



*Image\_corrected\_EMS\_stripped.gipl*

# Pipeline Description

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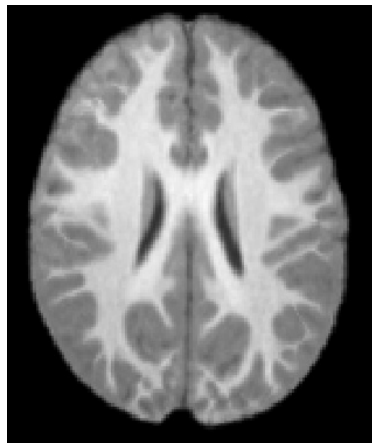
## Deformable registration of T1-weighted atlas (RegisterImages module)

B-spline pipeline registration.

A transformation file is created and will be used by the next step.

### Module link

[http://www.na-mic.org/Wiki/index.php/ITK\\_Registration\\_Optimization#Pipeline\\_Registration](http://www.na-mic.org/Wiki/index.php/ITK_Registration_Optimization#Pipeline_Registration)



*Atlas.gipl*



*AtlasRegistered\_Image\_corrected\_EMS\_stripped.gipl*



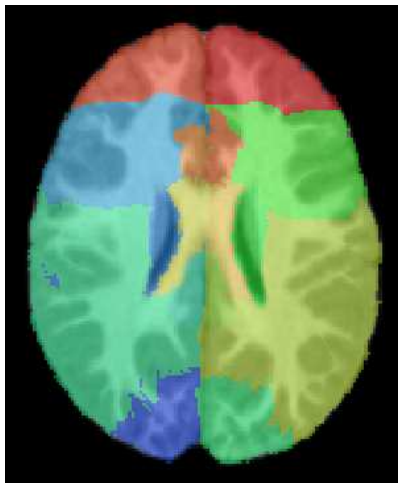
# Pipeline Description

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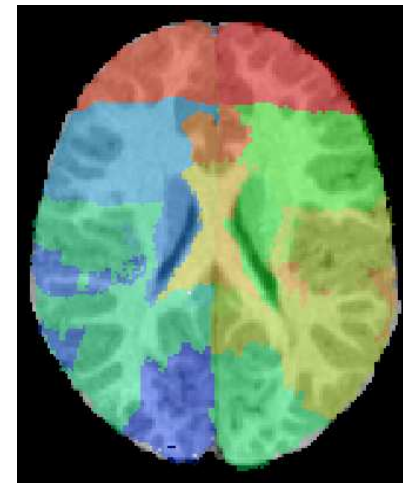
## Applying transformation to the atlas' parcellation map (ResampleVolume2 module)

**Module link :**

<http://slicer.spl.harvard.edu/slicerWiki/index.php/Modules:ResampleVolume2-Documentation>



*Parcellation.gipl*



*ParcellationRegistered\_Image\_corrected\_EMS\_stripped.gipl*

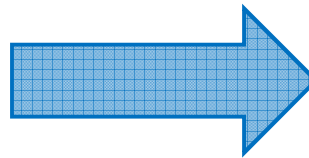
# Pipeline Description

## Cortical Thickness (CortThick external module)

Sparse and asymmetric local cortical thickness



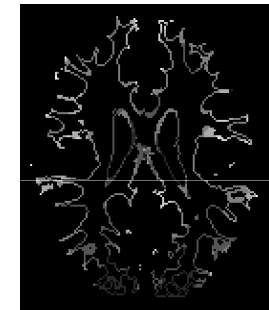
*Image\_labels\_EMS.gipl*



Label	Average	Std Dev	Nb Of Elem
1	2.96	1.81	1214
2	3.8	1.79	2113
3	2.93	1.89	1128
4	4.09	1.8	1796
5	3.9	2.52	897
6	4.15	1.93	9
7	4.31	1.76	90
8	3.39	1.41	2772
9	2.81	1.61	1479

*Regional Cortical Thickness information*

*Optional Outputs*



*WM\_AvgBoundary.gipl*



*GM\_AvgBoundary.gipl*



# *Pipeline Description*

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## **Volume information (ImageStat and ImageMath modules)**

Both modules are used to generate volume information in the following files :

- TissueSegmentationVolumes.csv  
White matter, gray matter and CSF volumes.
- ParcellationMapVolumes.csv (if the parcellation image is provided)  
White matter, gray matter and CSF volumes per lobe.



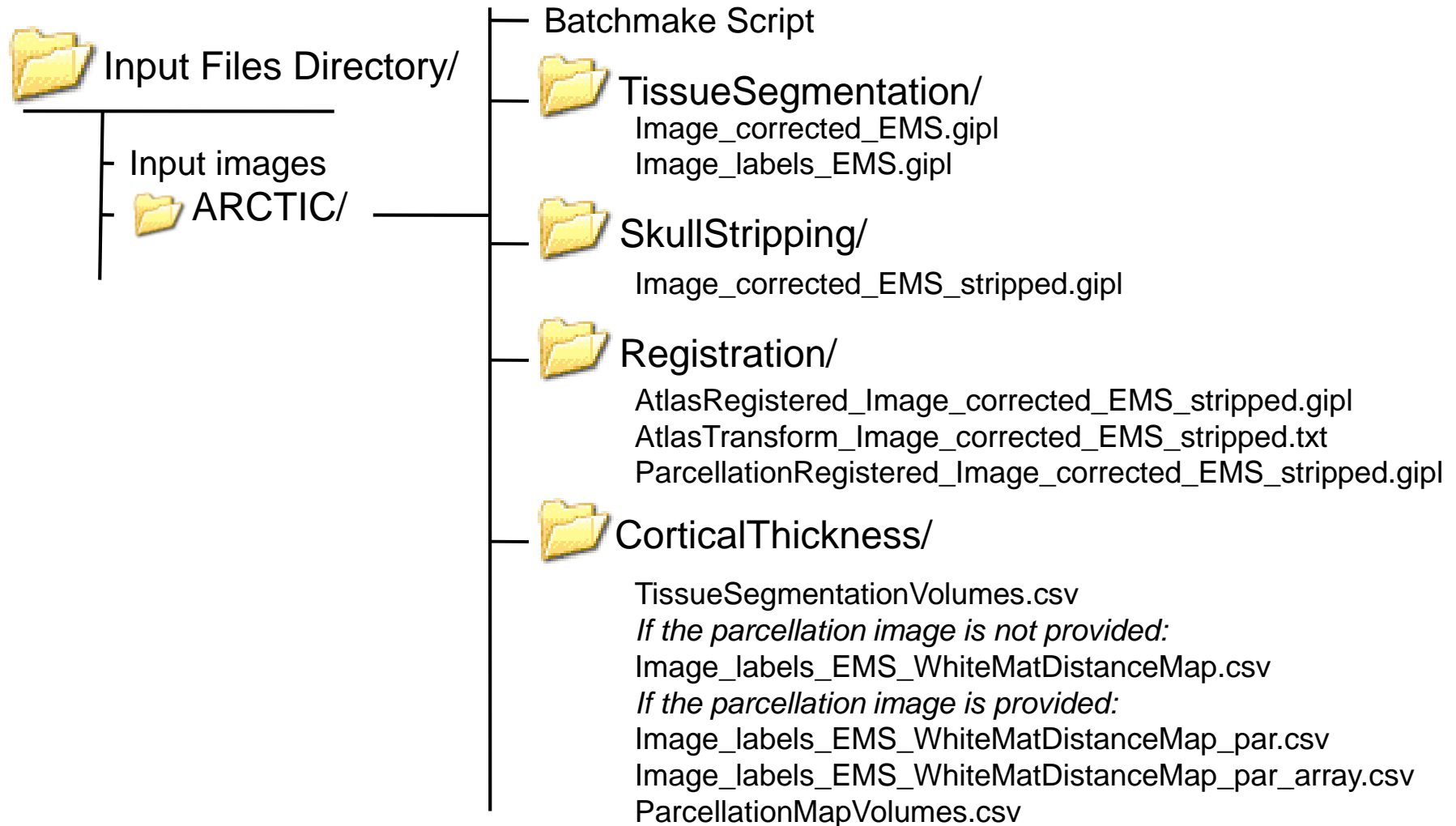
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# Output and Organisation

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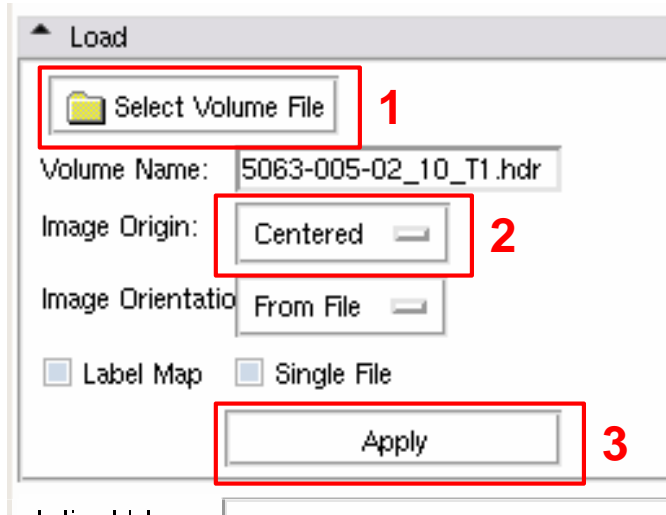
# *Execution within Slicer*

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- **Load input images**
- Demonstration with « Raw Images »
- Demonstration with « Segmented Image »



# Demonstration : Load the input images

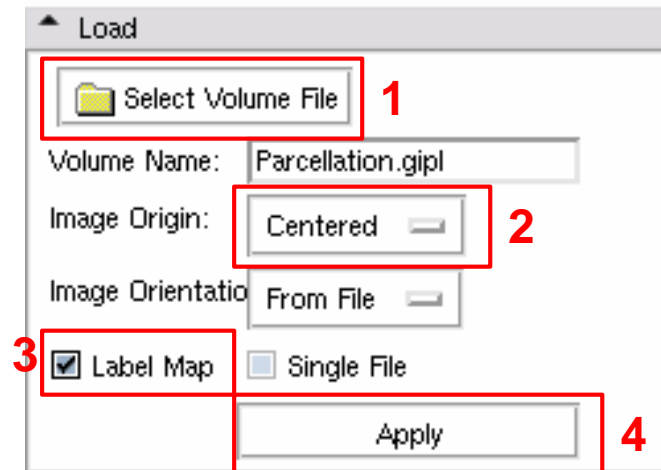


How to load raw images (case and atlas)?

- 1- Select the image in the browser
- 2- Set the image origin as « centered »
- 3- Click on « Apply » to load

How to load parcellation and label images?

- 1- Select the image in the browser
- 2- Set the image origin as « centered »
- 3- Check the « label map » button
- 4- Click on « Apply » to load





# *Demonstration in Slicer*

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- Load input images
- **Demonstration with « Raw Images »**
- Demonstration with « Segmented Image »
- Parcellation option
- Advanced parameters



# Demonstration : "Raw Images" Input 1

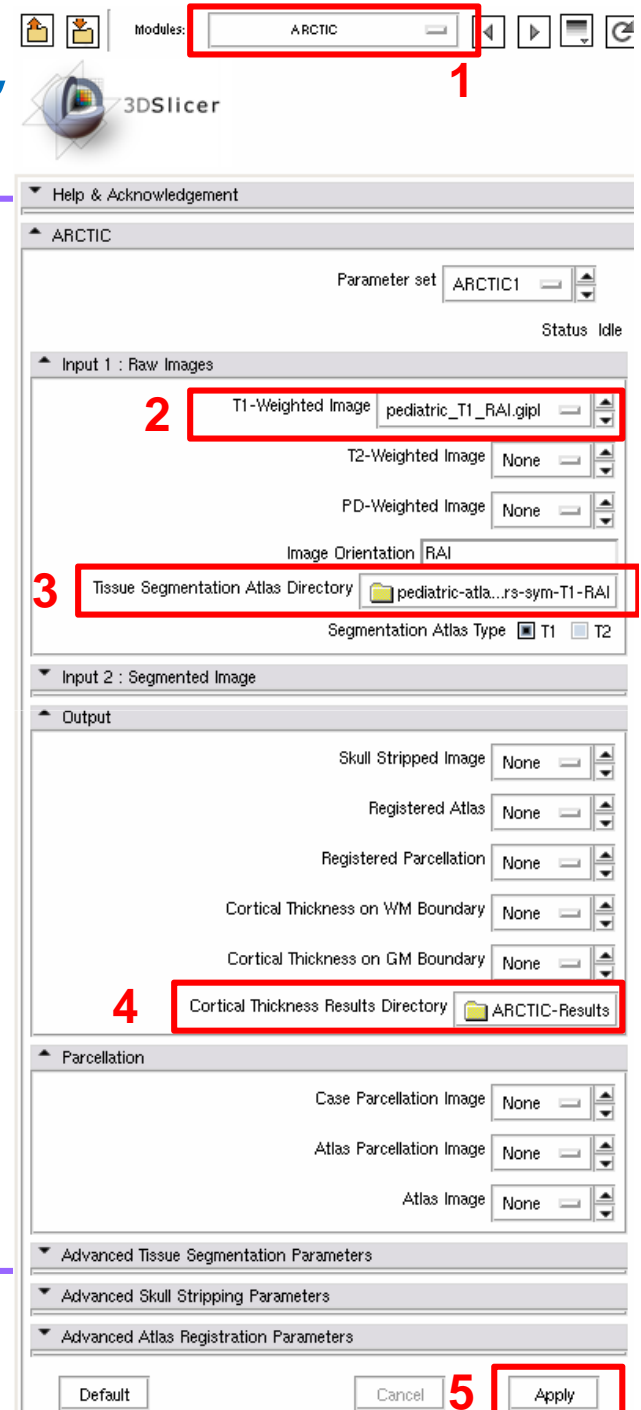
1- Select the « ARCTIC » module (in All Modules)

2- Add the T1-weighted image

3- Set the Tissue Segmentation Atlas Directory for the tissue segmentation

4- Set the output directory

5- Click on the « Apply » button to process the data

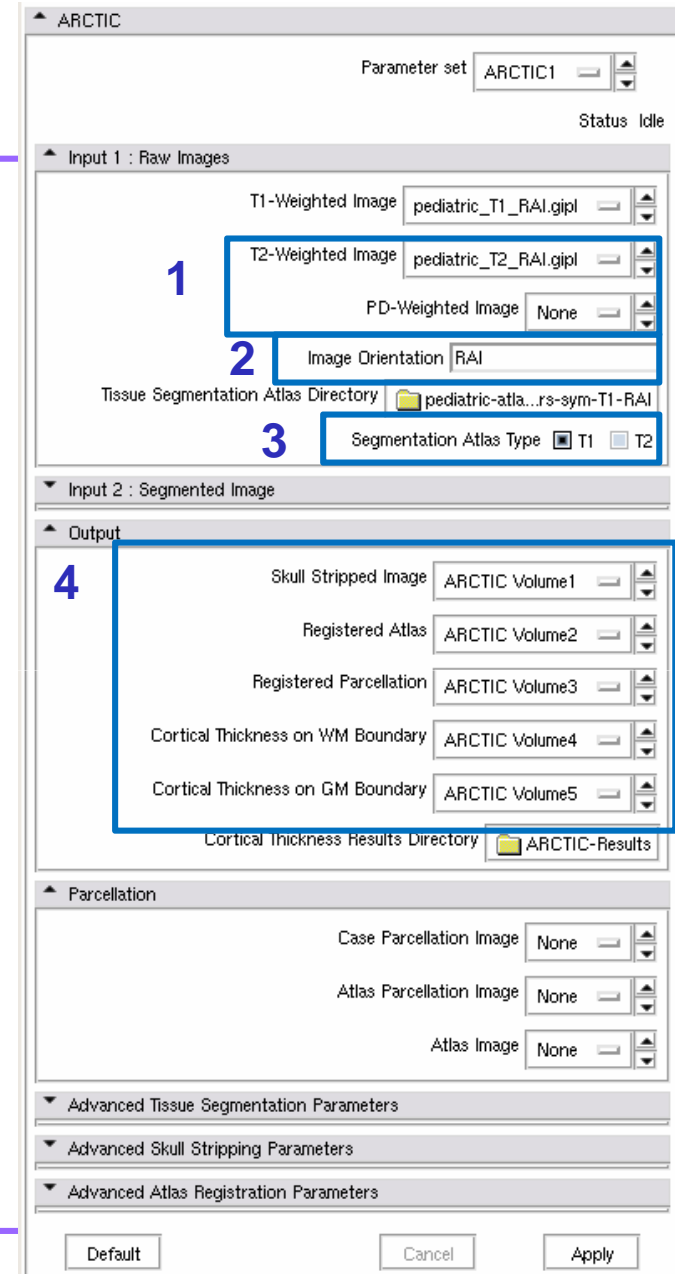




# Demonstration : "Raw Images" Input 1

## Verifications / Options

- 1- If available, set the T2 and/or PD-weighted images to improve the tissue segmentation
- 2- Check that the atlas has the same orientation than the input images
- 3- Check the tissue segmentation atlas type (T1-weighted or T2-weighted image)
- 4- Set the output images to be displayed in Slicer (« Create a new volume » instead of « None »)





# *Demonstration in Slicer*

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- Load input images
- Demonstration with « Raw Images »
- **Demonstration with « Segmented Image »**
- Parcellation option
- Advanced parameters



# Demonstration : "Segmented Image" Input 2

- 1- Select the « ARCTIC » module (in All Modules)
- 2- Set the tissue segmentation label image and check the related tissue labels
- 3- Set its raw image (T1-weighted, T2-weighted, PD-weighted)
- 4- Set the output directory
- 5- Click on the « Apply » button to process the data

The screenshot shows the ARCTIC module interface in 3DSlicer. The 'Modules' dropdown at the top is set to 'ARCTIC' (1). The 'Parameter set' is 'ARCTIC1' and the status is 'Idle'. Under 'Input 2 : Segmented Image', the 'Tissue Segmentation Image' is set to 'None' (2). Below it, 'White Matter Label 1', 'Grey Matter Label 2', and 'CSF Label 3' are listed. The 'Raw Image' is also set to 'None' (3). In the 'Output' section, the 'Cortical Thickness Results Directory' is set to 'ARCTIC-Results' (4). At the bottom, the 'Apply' button is highlighted (5).





# Demonstration : "Segmented Image" Input 2

## Options

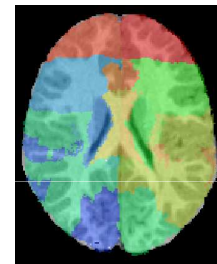
1- Set the output images to be displayed in Slicer ( « Create a new volume » instead of « None » )



Skull Stripped Image



Registered Atlas



Registered Parcellation



Cortical Thickness on WM Boundary



Cortical Thickness on GM Boundary

The screenshot shows the ARCTIC software interface. The 'Output' section is highlighted with a blue box and contains the following settings:

- Skull Stripped Image: ARCTIC Volume1
- Registered Atlas: ARCTIC Volume2
- Registered Parcellation: ARCTIC Volume3
- Cortical Thickness on WM Boundary: ARCTIC Volume4
- Cortical Thickness on GM Boundary: ARCTIC Volume5

Other visible settings include:

- Parameter set: ARCTIC1
- Status: Idle
- Input 1: Raw Images
- Input 2: Segmented Image
  - Tissue Segmentation Image: None
  - White Matter Label: 1
  - Grey Matter Label: 2
  - CSF Label: 3
  - Raw Image: None
- Parcellation
  - Case Parcellation Image: None
  - Atlas Parcellation Image: None
  - Atlas Image: None
- Cortical Thickness Results Directory: ARCTIC-Results

Buttons at the bottom: Default, Cancel, Apply.



# *Demonstration in Slicer*

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- Load input images
- Demonstration with « Raw Images »
- Demonstration with « Segmented Image »
- **Parcellation option**
- Advanced parameters



# Parcellation options

## Parcellation options

*If you want to perform a lobar cortical thickness analysis, choose between the two possibilities*

**a-** Add a parcellation image which is defined in the input coordinate space (« Case Parcellation Image »)

**b-** Add the atlas raw image **and** its parcellation, defined in the atlas coordinate space (« Atlas Parcellation Image »)

The screenshot shows the ARCTIC software interface with the following settings:

- Parameter set: ARCTIC1
- Status: Idle
- Input 1: Raw Images
- Input 2: Segmented Image
  - Tissue Segmentation Image: None
  - White Matter Label: 1
  - Grey Matter Label: 2
  - CSF Label: 3
  - Raw Image: None
- Output
  - Skull Stripped Image: ARCTIC Volume1
  - Registered Atlas: ARCTIC Volume2
  - Registered Parcellation: ARCTIC Volume3
  - Cortical Thickness on WM Boundary: ARCTIC Volume4
  - Cortical Thickness on GM Boundary: ARCTIC Volume5
  - Cortical Thickness Results Directory: ARCTIC-Results
- Parcellation
  - Case Parcellation Image: None (labeled 'a')
  - Atlas Parcellation Image: Parcellation.gipl (labeled 'b')
  - Atlas Image: template.gipl
- Advanced Tissue Segmentation Parameters
- Advanced Skull Stripping Parameters
- Advanced Atlas Registration Parameters

Buttons at the bottom: Default, Cancel, Apply



# *Demonstration in Slicer*

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- Load input images
- Demonstration with « Raw Images »
- Demonstration with « Segmented Image »
- Parcellation option
- **Advanced parameters**



# Advanced parameters

## Tissue segmentation parameters

**a-** Filter options: specifies smoothing parameters prior to the segmentation

**b-** Priors weighting the tissue classes for the segmentation

**c-** Atlas warping options:

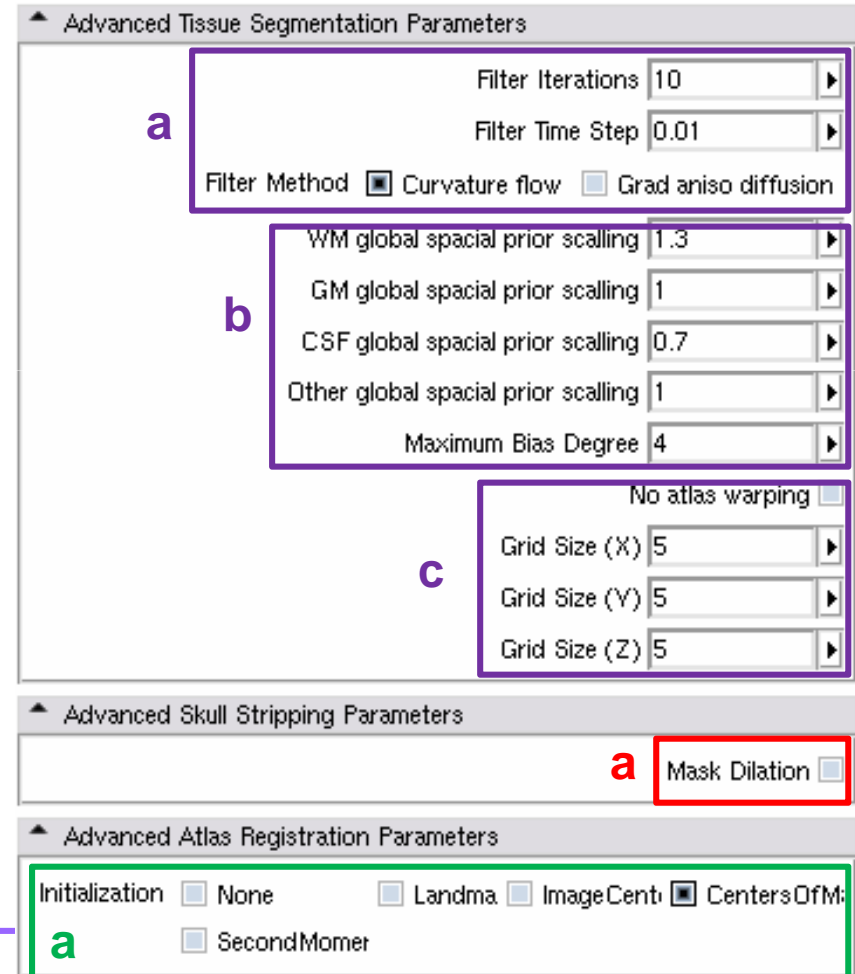
- No atlas warping:
- Unchecked by default: atlas to subject B-Spline registration is performed
- Checked: atlas to subject affine registration is performed instead of the warping
  - Grid size X,Y,Z: grid controls points for atlas warping

## Skull stripping parameters

**a-** Check to apply a dilation of the mask (necessary if the tissue segmentation has a low quality)

## Atlas registration parameters

**a-** Different initialization methods





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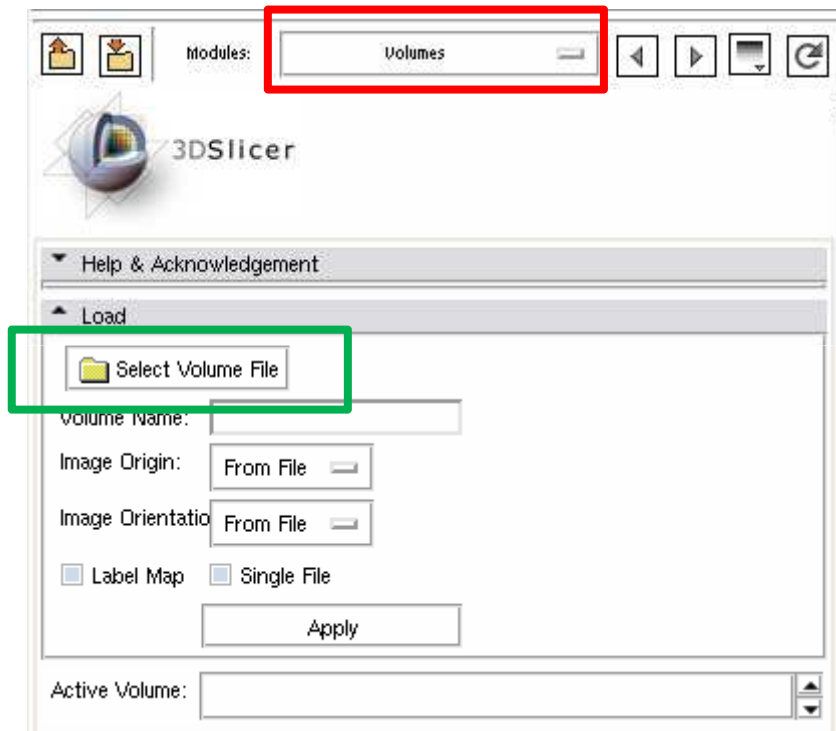


# *Example with tutorial dataset*

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- Load input images
- Run ARCTIC

# Load input images

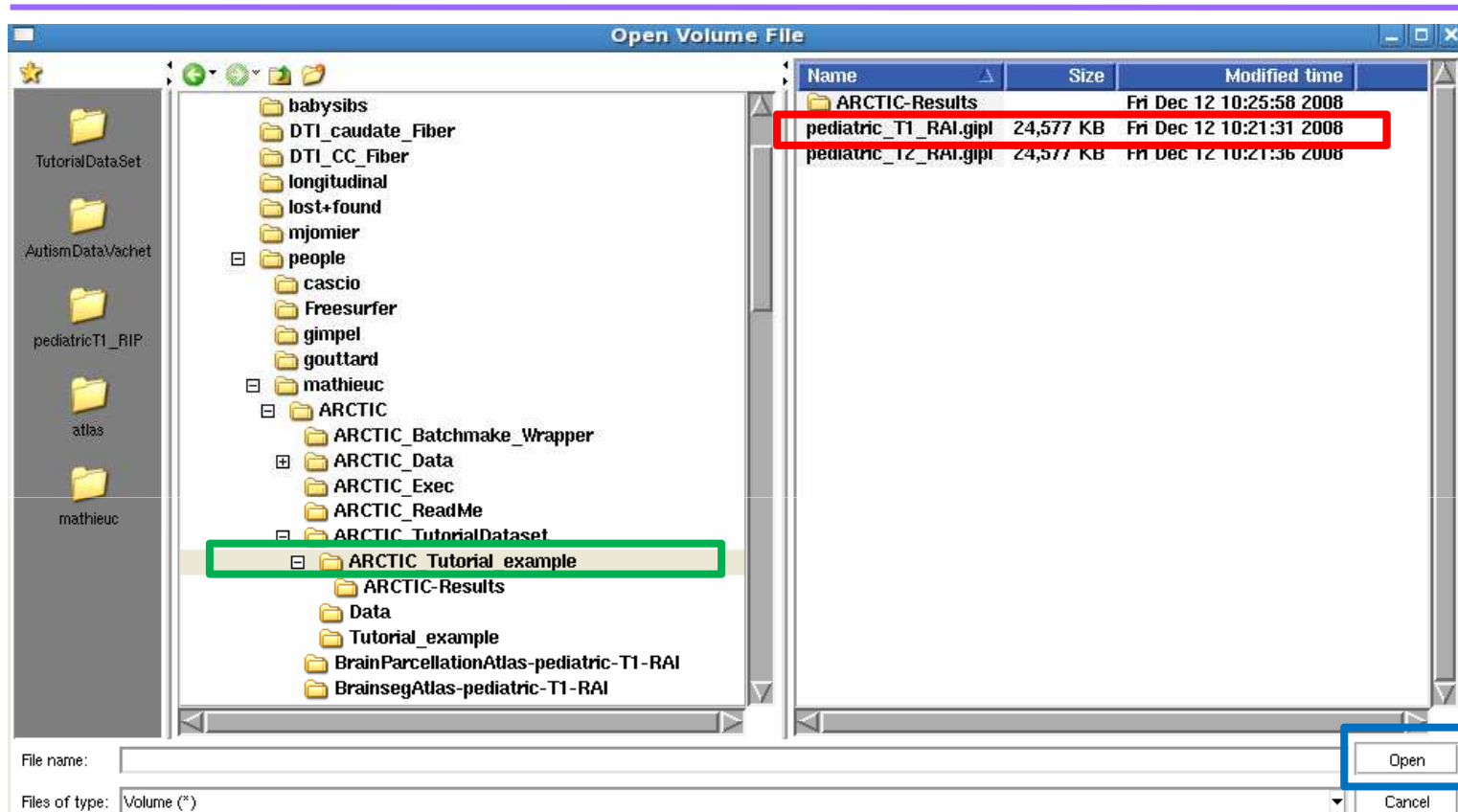


In Slicer, select the module « **Volumes** » to load the input images.

Then click on the « **Select Volume File** » button to load the images.



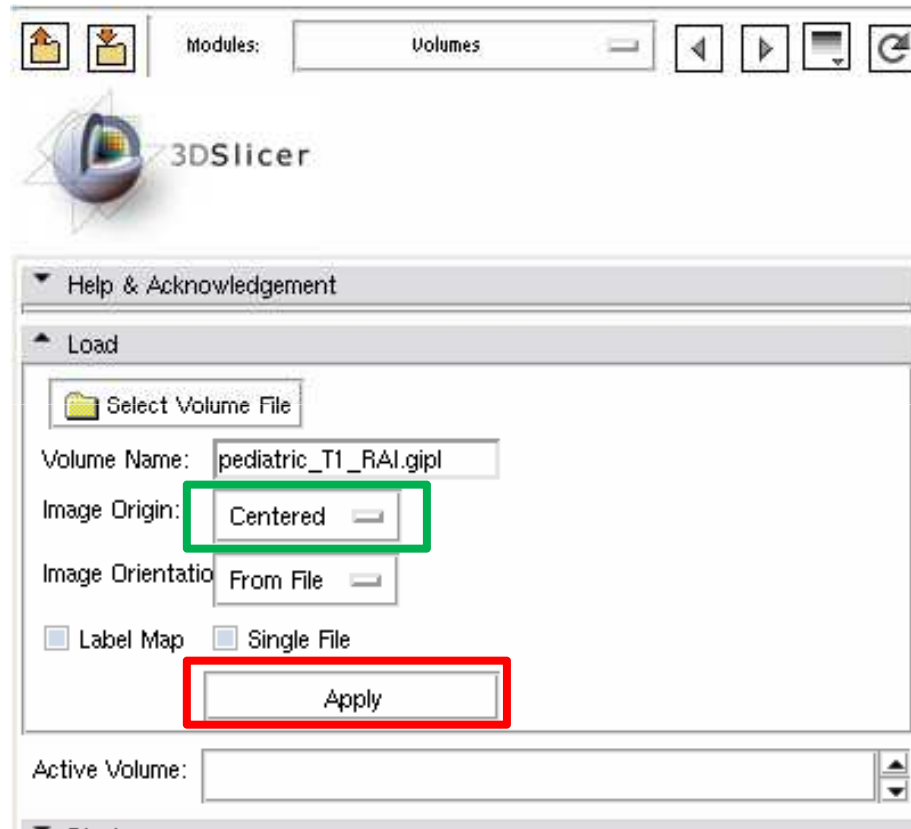
# Load input images



A new window 'Open Volume File' is now open. Select the « **ARCTIC\_Tutorial\_example** » directory. Select the « **pediatric\_T1\_RAI.gipl** » file in the Data directory and click on « **Open** ».



# Load input images

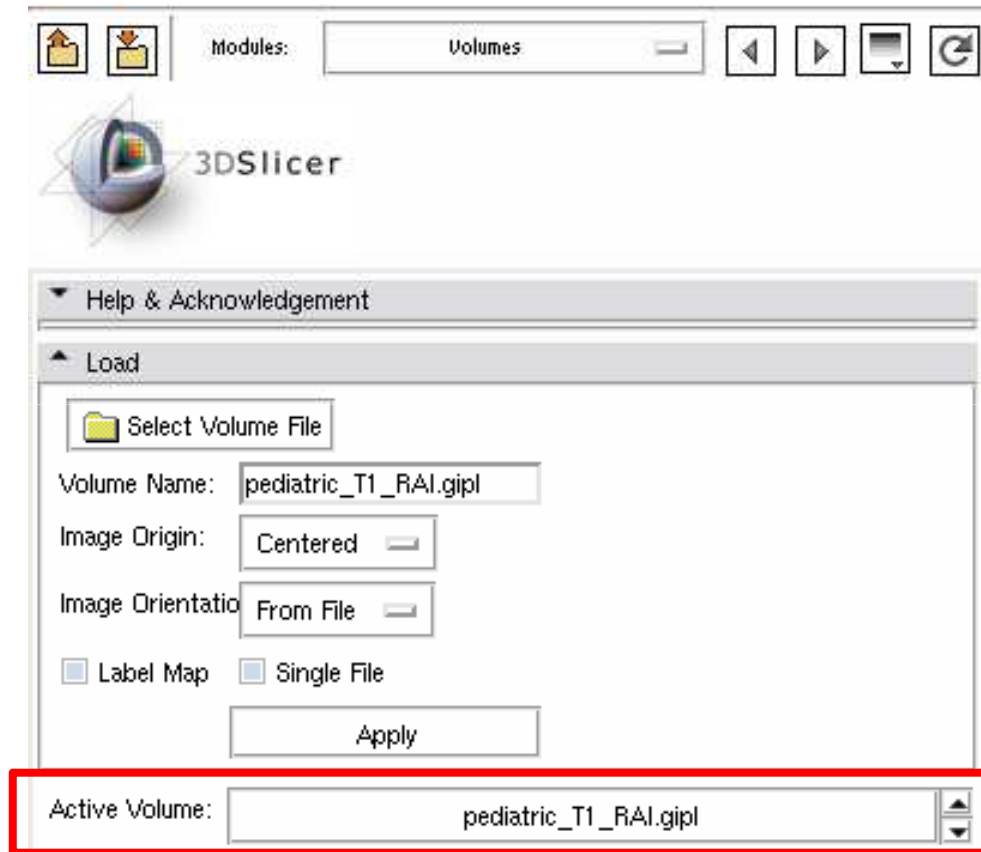


Now, select the Image Origin as « **Centered** ».

And click on « **Apply** ».



# Load input images



The first image is now loaded.

You can check it in the « **Active Volume** » widget.



# *Load input images*

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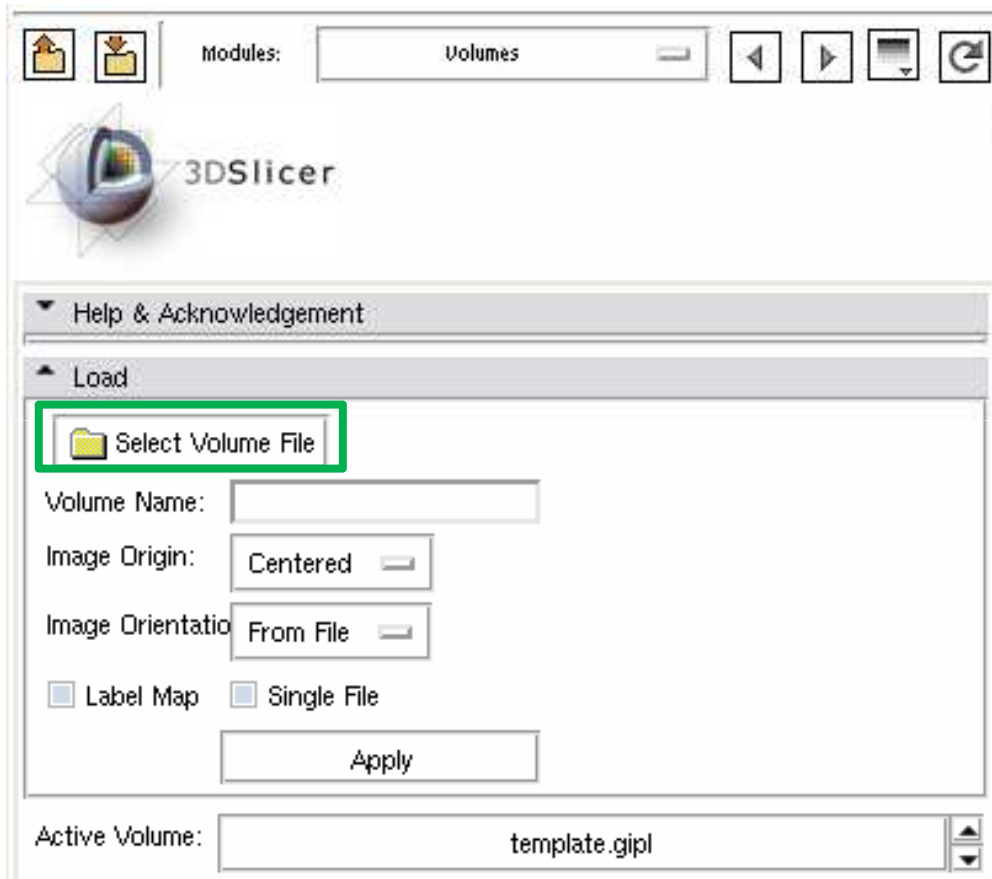
Apply the same steps to load the T2-weighted and atlas images.

One can find the T2-weighted image in the same directory than the T1-weighted one.

The atlas image, named « `template-stripped.gipl` » is in the « `pediatric-atlas-4years-sym-T1-RAI` » directory.

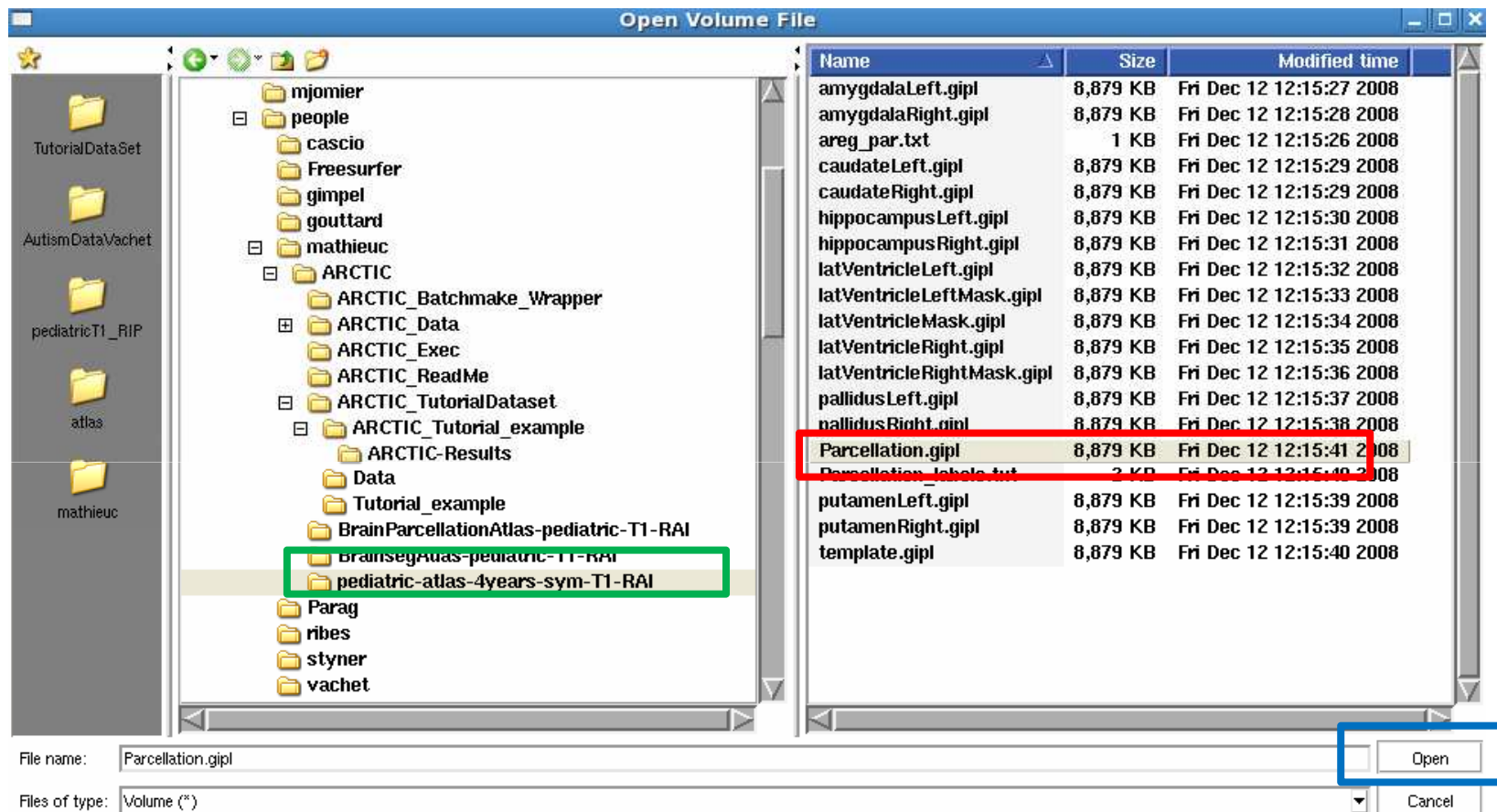


# Load input images



Now we will load the parcellation image.  
Click on the « **Select Volume File** » button to load the parcellation.

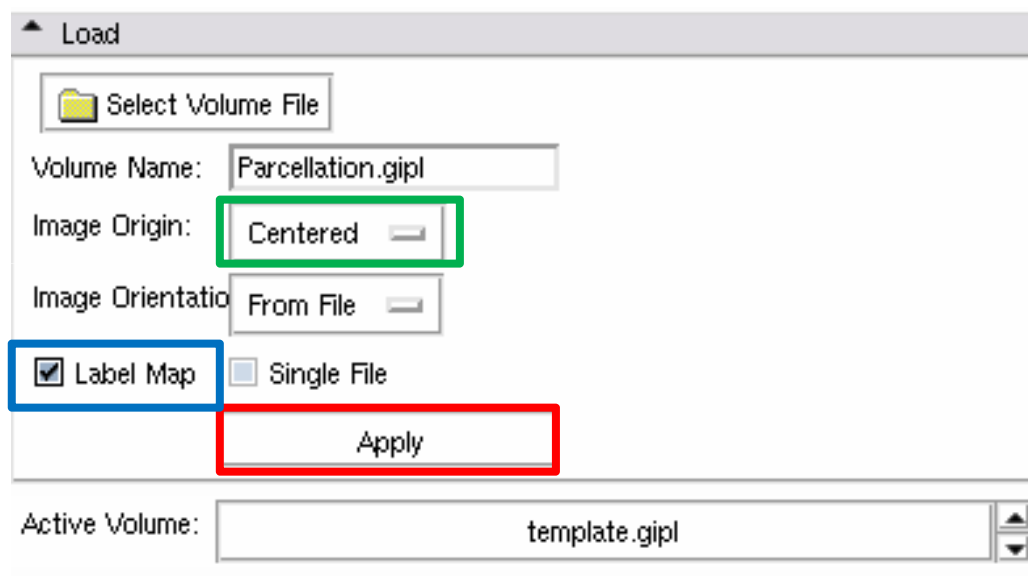
# Load input images



A new window 'Open Volume File' is now open. Select the « **pediatric-atlas-4years-sym-T1-RAI** » directory . Select the « **Parcellation.gipl** » file and click on « **Open** » .

# Load input images

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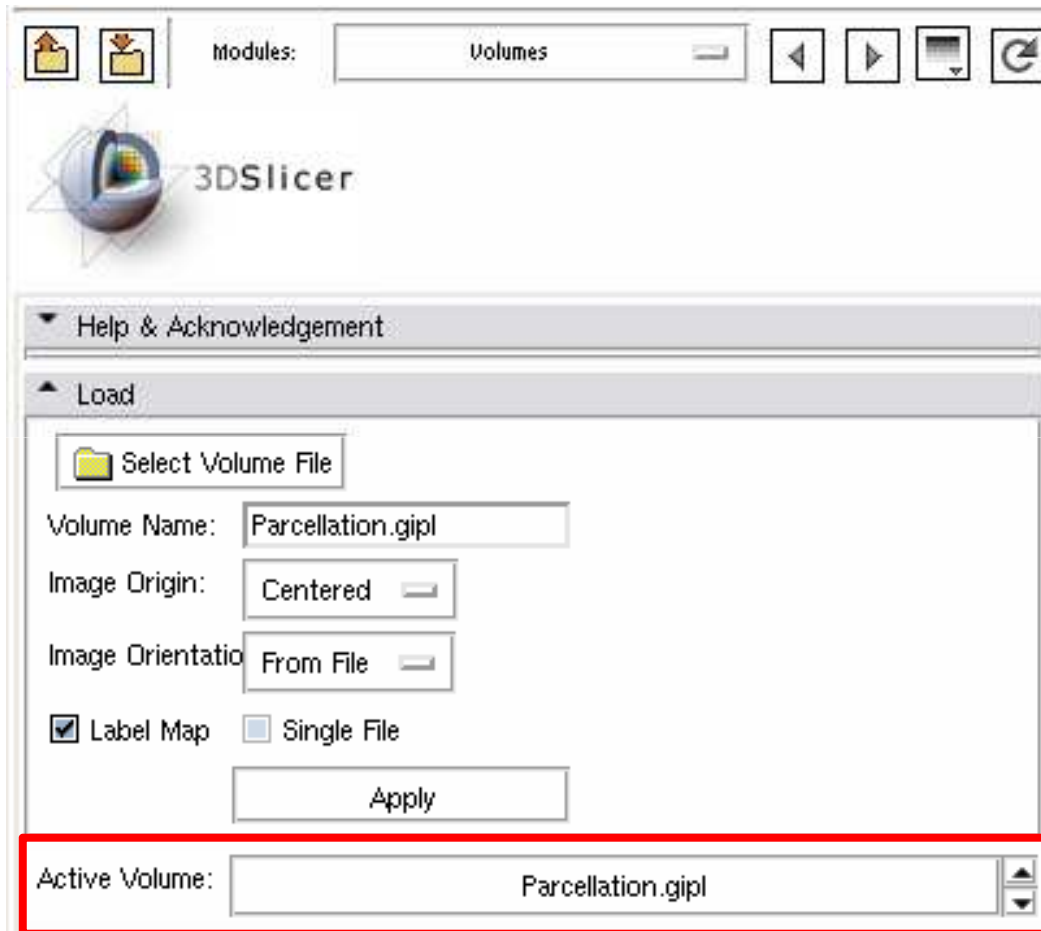
Now, select the Image Origin as « **Centered** ».

Then, check the « **Label Map** » case to load the parcellation as a label image.

And click on « **Apply** ».



# Load input images



The dataset is now loaded.

You can check it in the « **Active Volume** » widget while displaying the 4 images.





# *Example with tutorial dataset*

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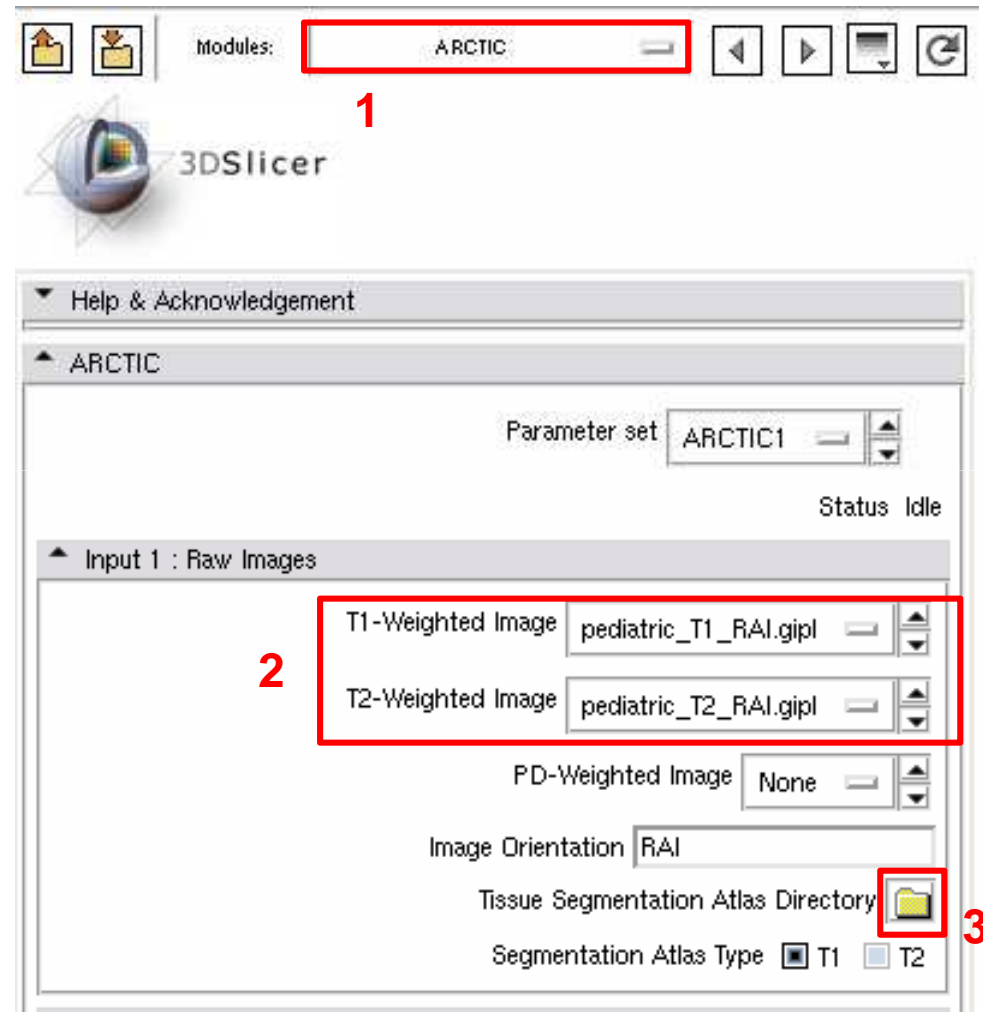
- Input images loading
- ARCTIC execution

# Module execution

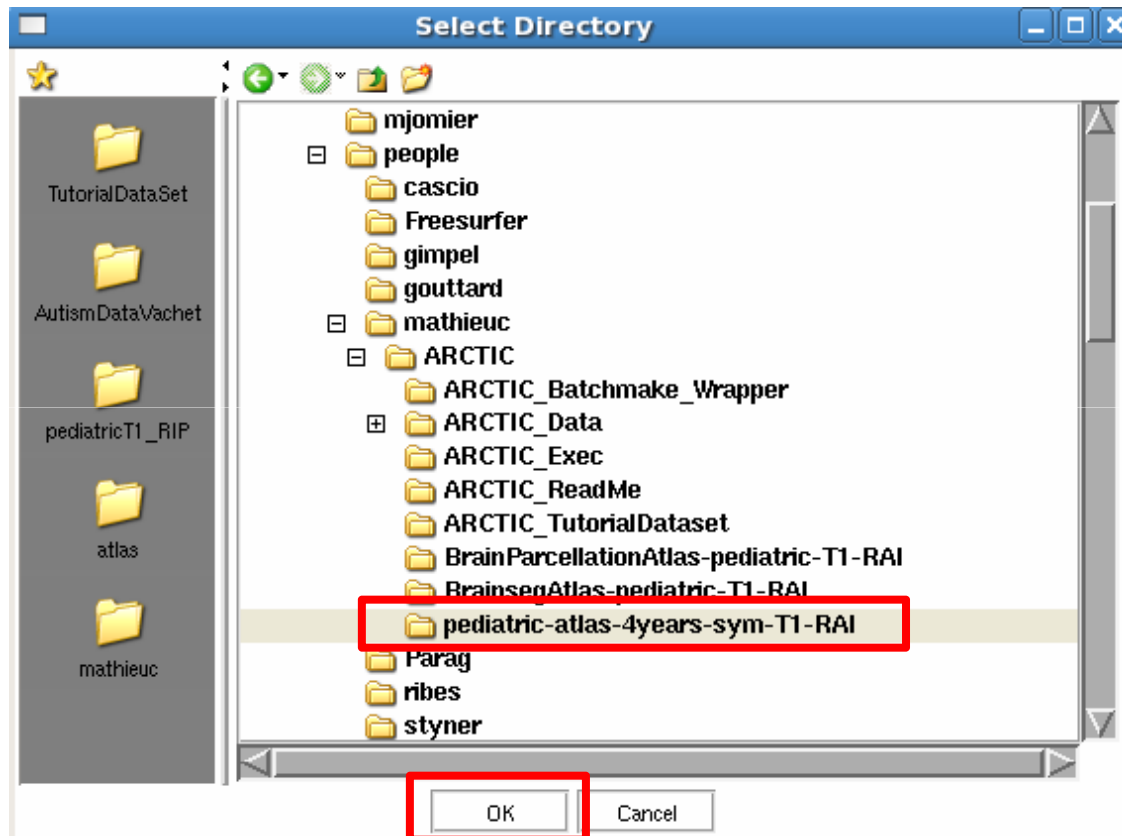
1- Select the « **ARCTIC** » module (in All Modules)

2- Set the T1 and T2-weighted images (**pediatric\_T1/T2\_RAI.gipl**)

3- Click on the « **Tissue Segmentation Atlas Directory** » button



# Module execution



A new window is now open to select the tissue segmentation atlas.

Search and select the « **pediatric-atlas-4years-sym-T1-RAI/** » folder.

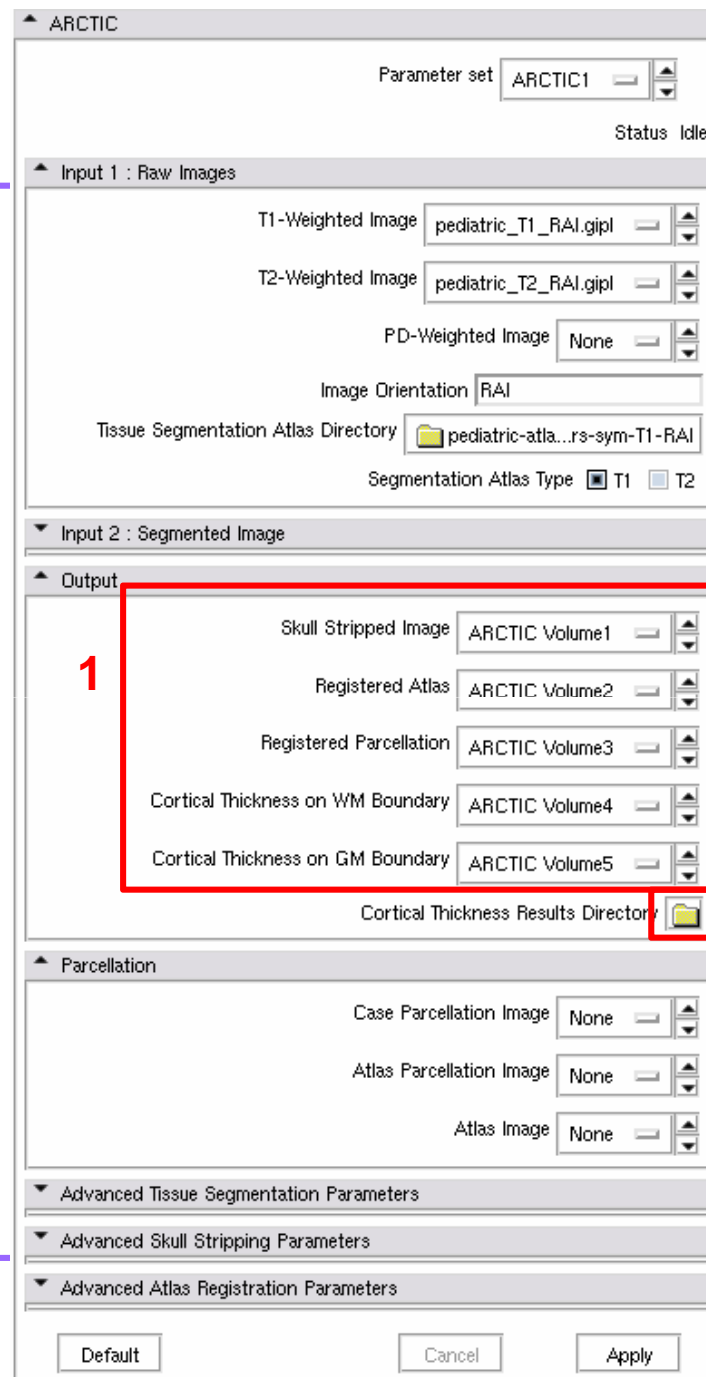
Click on the « **OK** » button to confirm.



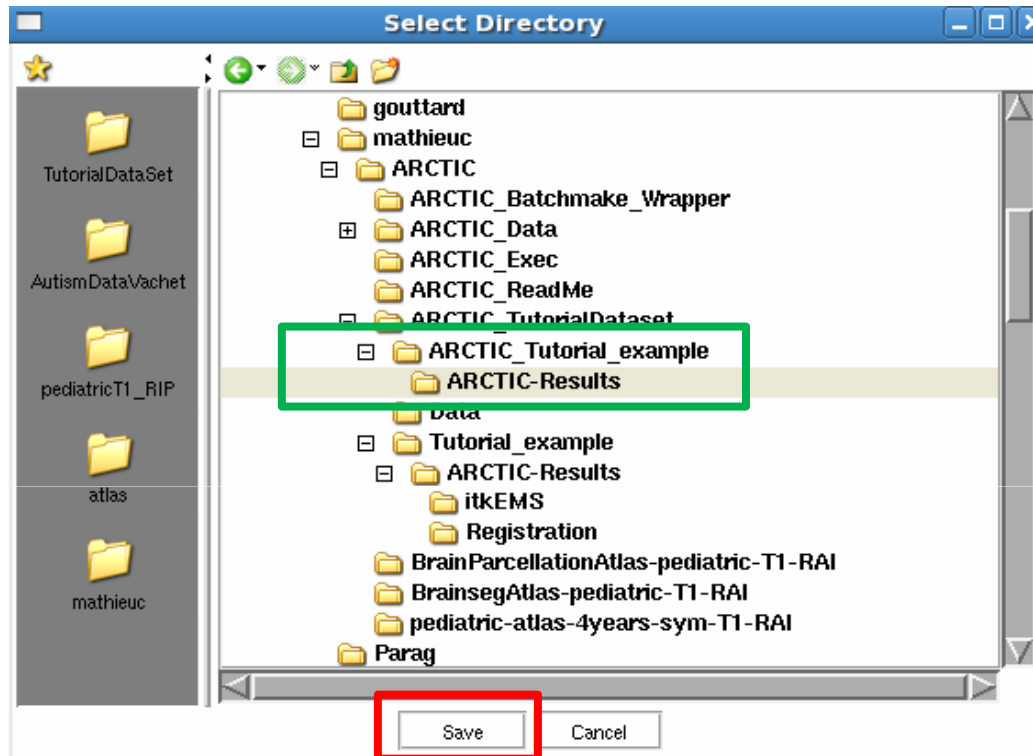
# Module execution

1- Select « **Create a new volume** » to display output images

2- Click on the « **Cortical Thickness Results Directory** » button



# Module execution



Select the « **ARCTIC-Results** » folder in the ARCTIC\_Tutorial\_Example directory

Click on the « **Save** » button to confirm your choice.



# Module execution

- 1- Add the « **Parcellation.gipl** » as Atlas Parcellation Image, and the « **template-stripped.gipl** » as Atlas Image
- 2- Click on the « **Apply** » button to start the process.

ARCTIC

Parameter set: ARCTIC1

Status: Idle

Input 1 : Raw Images

- T1-Weighted Image: pediatric\_T1\_RAI.gipl
- T2-Weighted Image: pediatric\_T2\_RAI.gipl
- PD-Weighted Image: None
- Image Orientation: RAI
- Tissue Segmentation Atlas Directory: pediatric-atla...rs-sym-T1-RAI
- Segmentation Atlas Type:  T1  T2

Input 2 : Segmented Image

Output

- Skull Stripped Image: ARCTIC Volume1
- Registered Atlas: ARCTIC Volume2
- Registered Parcellation: ARCTIC Volume3
- Cortical Thickness on WM Boundary: ARCTIC Volume4
- Cortical Thickness on GM Boundary: ARCTIC Volume5
- Cortical Thickness Results Directory: ARCTIC-Results

Parcellation

- Case Parcellation Image: None
- Atlas Parcellation Image: Parcellation.gipl
- Atlas Image: template.gipl

Advanced Tissue Segmentation Parameters

Advanced Skull Stripping Parameters

Advanced Atlas Registration Parameters

Default Cancel Apply

# Module execution

---

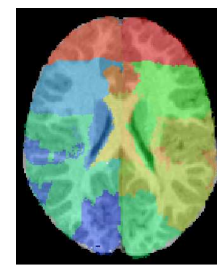
Once the execution is finished, several images are displayed within Slicer. You can compare your images with the following ones to perform a quick quality control.



*Skull Stripped Image  
ARCTIC Volume1*



*Registered Atlas  
ARCTIC Volume2*



*Registered Parcellation  
ARCTIC Volume3*



*Cortical Thickness on WM  
Boundary  
ARCTIC Volume4*



*Cortical Thickness on GM  
Boundary  
ARCTIC Volume5*



# Overview

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- 1- Pipeline overview
- 2- Input images
- 3- Pipeline description
- 4- Output images and organisation
- 5- Execution within Slicer
- 6- Example with tutorial dataset
- 7- **Command line execution**





# *Use the command line*

---

## « Raw Images » Mode

### *Global analysis*

```
ARCTIC --T1 Image_T1.gipl --segAtlasDir TissueSegmentationAtlasDirectory/
```

### *Lobar cortical thickness analysis*

If the atlas raw image and its parcellation are provided:

```
ARCTIC --T1 Image_T1.gipl --segAtlasDir TissueSegmentationAtlasDirectory/  
--atlas Atlas.gipl --atlasParcellation Parcellation.gipl
```

If the case parcellation image is provided:

```
ARCTIC --T1 Image_T1.gipl --segAtlasDir TissueSegmentationAtlasDirectory/  
--caseParcellation CaseRegisteredParcellation.gipl
```



# *Use the command line*

---

## « Segmented Image » Mode :

### *Global analysis*

```
ARCTIC --label TissueSegmentationImage.gipl --rawImage Image_T1.gipl
```

### *Lobar cortical thickness analysis*

If the atlas raw image and its parcellation are provided:

```
ARCTIC --label TissueSegmentationImage.gipl --rawImage Image_T1.gipl  
--atlas Atlas.gipl --atlasParcellation Parcellation.gipl
```

If the case parcellation image is provided:

```
ARCTIC --label TissueSegmentationImage.gipl --rawImage Image_T1.gipl  
--caseParcellation CaseRegisteredParcellation.gipl
```



# *Conclusion*

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Slicer3 toolkit provides an accessible and versatile platform to conduct image processing of MRI data, in this case, regional cortical thickness analysis using ARCTIC.

Thanks to this tutorial you are now ready to perform a regional cortical thickness analysis on your own dataset.



# *Acknowledgements*

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