Exercises Instructions

# Introduction

This material is intended for practical demonstration using STIR on PET and SPECT Image Reconstruction.

Simulated data will be prepared during the exercises. These are based on 2 sets of images:

* Thorax phantom data are obtained from the recent open access article: *Tsoumpas et al 2013 Phys Med Biol*.   
  We have two respiratory gated positions of a thorax FDG PET phantom along with the corresponding CTAC image.
* Brain data are obtained from BrainWeb.  
  We have a segmented brain-map.

You will probably only want to run either the brain or the thorax data (except for the motion correction exercise which is currently only for the thorax).

The input data are stored in the folders called EX**\_\*, but you will need to run the scripts from the “main” exercises folders** (open a terminal, cd to where you extracted the exercises, and always cd back after every exercise).

We are using Python for some of the exercises. Python is an open-source interactive language, a bit like MATLAB. We provide Python scripts for the evaluation, so you should be fine. It would be best to read a Python tutorial before the course. We will use Spyder as our Python environment. You can start it from the terminal with

cd ~/exercises

spyder&

As an alternative to Python, we also provide instructions for loading sinograms and images in a display program. Note that in the text below we’re using AMIDE for display. ImageJ would work as well (see the end of the document).

See the appendices at the end of this document for some information to get started

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# Exercise: Data Simulation Brain

(Always run scripts from the exercises directory)

This is a simple simulation of a brain phantom. PSF is incorporated. Scatter is set to zero. Randoms are constant.

Read and run script:

./run\_simulation\_brain.sh

## Python evaluation

Start spyder with the evaluation script

spyder evaluate\_simulation\_brain.py&

or if spyder is running, just open the file.

## Command line evaluation

You will need to extract the sinograms in an “image” Interfile to be able to load them in AMIDE

cd working\_folder/brain

extract\_segments my\_prompts.hs

Extract as SegmentByView (0) or BySinogram (1)?[0,1 D:0]: 1

Display the extracted sinogram (e.g. my\_promptsseg0\_by\_sino.hv) using **AMIDE**. On Linux, you can do

amide my\_promptsseg0\_by\_sino.hv&

Check the central sinogram plane to see if it looks as expected.

Go back to main directory

cd ../..

We can also extract profiles through the sinogram to display these in Excel or GNUmeric or similar. An example of this is given in the script

./evaluate\_simulation\_brain.sh

which will extract the segments and create profile text files for you. Output files will be

working\_folder/brain/profile\_prompts.txt

working\_folder/brain/profile\_randoms.txt

# Exercise: Data Simulation Thorax

(Always run scripts from the exercises directory)

This is a simple simulation of a thorax phantom (2 gates). PSF is not incorporated. Scatter is simulated using STIR. Randoms are constant.

Read and run script:

./run\_simulations\_thorax.sh

## Python evaluation

Start spyder with the evaluation script

spyder evaluate\_simulation\_thorax.py&

or if spyder is running, just open the file.

## Command line evaluation

You will need to extract the sinograms in an “image” Interfile to be able to load them in AMIDE

cd working\_folder/GATE1/

extract\_segments my\_prompts\_g1.hs

Extract as SegmentByView (0) or BySinogram (1)?[0,1 D:0]: 1

cd ../GATE2/

extract\_segments my\_prompts\_g2.hs

Extract as SegmentByView (0) or BySinogram (1)?[0,1 D:0]: 1

cd ..

Import the extracted sinogram (e.g. my\_prompts\_g1seg0\_by\_sino.hv) using **AMIDE**.

Select and export the central sinogram plane and upload them online.

Subtract the two sinograms. This can be done in AMIDE or on the command line

stir\_subtract -s diff.hs GATE1/my\_prompts\_g1.hs GATE2/my\_prompts\_g2.hs

extract\_segments diff.hs

Extract as SegmentByView (0) or BySinogram (1)?[0,1 D:0]: 1

(Note: when using copy-paste of the above lines, please make sure that the “-s” option of stir\_subtract uses a minus sign. On some systems it copies as a character which looks the same but isn’t…)

How does the difference look like in sinogram space?

Go back to main directory

cd ..

We can also extract profiles through the sinogram to display these in Excel or similar. You could run

./evaluate\_simulation\_thorax.sh

to extract the segments and create profiles for you.

# Preparation for Image reconstruction

For the thorax reconstruction exercise, we first need to generate a new simulation data set. This time just a single slice to speed things up. Scatter is also set to zero for simplicity here.

./run\_simulation\_single\_slice.sh

Output is in working\_folder/single\_slice

# Exercise: Image reconstruction part 1

(Always run scripts from the exercises directory)

You will need to have run the corresponding simulation script from the previous section. Output is in working\_folder/single\_slice or working\_folder/brain

We will now look at EMML, OSEM and OSEM with PSF. A sample script is provided to generate results

./run\_reconstruction\_thorax.sh

Or

./run\_reconstruction\_brain.sh

This will run EMML for 240 iterations and OSEM for 240 iterations. It will also continue OSEM from there and write images at every subiterations. Finally, PSF reconstruction will also be performed. The script generates some differences images and launches AMIDE.

You could decide to run only a subset of these exercises and modify the script for yourself.

Sample questions to address:

* Is it worth running EMML? Why not simply use OSEM?
* Does this depend on the number of iterations that you use? And post-filtering?
* When using PSF reconstruction, can you see effects discussed during the lecture?

# Exercise: Image reconstruction part 2 (MAP)

This exercise needs results from the previous step (as the MAP reconstruction starts from an OSEM image in this exercise). Output is in working\_folder/single\_slice or working\_folder/brain

We will now look at OSL and OSSPS with a Quadratic Prior. A sample script is provided to generate results

./run\_reconstruction\_thorax\_MAP.sh

or

./run\_reconstruction\_brain\_MAP.sh

This will run OSL and OSSPS (continuing from a previous OSEM image).

Sample questions to address:

* Do OSL and OSSPS generate the same results?
* Does this depend on the penalty factor? Noise level? Iteration number? Initialisation (try to remove the initial estimate for instance).

# Exercise: Adding Poisson noise

(advanced exercise. You will have to run the brain simulation and reconstruction scripts first.)

We can make the simulation more realistic by adding noise to the data. An example would be

poisson\_noise -p my\_noisy\_data my\_prompts.hs 0.1 1

Run poisson\_noise to understand what these arguments mean. Execute a similar command in working\_folder/brain to create a noisy simulation. Check if you added “enough” noise (use extract\_segments to display for instance). Adjust the reconstruction parameter files to use your new noisy data (input) and change the filename used for the output. Run some reconstructions to see what noisy does to your images.

# Exercise: Scatter Correction

(Always run scripts from the exercises directory)

There are 4 example scatter estimation scripts which can be used to investigate different questions about scatter.

**N = 0**

Ideal (i.e. correct attenuation map, scatter simulation matches with how the data was generated) scatter correction (using 3 scatter correction loops)

**N = 1**

Calculate scatter by using a smaller energy window than that simulated. This will demonstrate if the scaling technique works. We have selected 425keV for the lower energy window (original is 350keV).

**~~N = 2~~**(this is currently not available)

~~Scatter correction using the scatter estimation from the first gate for both gates. This will demonstrate how sensitive is scatter in choosing different but adjacent gates.~~

**N = 3**

Perform reconstruction & attenuation correction by using wrong attenuation map. In the particular exercise we have assigned bone attenuation value to lung attenuation value for the first gate. Then we use this wrong attenuation map located at the first gate to correct for attenuation and scatter for each gates.

You should run the scripts as folllows (you can try reading it but these scripts are relatively complicated):

./run\_scatter\_0.sh

The scripts make the following files in working\_folder/GATE1 (and similar in working\_folder/GATE2)

* input\_g1.hs: “measured” sinogram after randoms correction
* my\_scatter.hs: sinogram output of the simulation (i.e. ground truth)
* scatter\_estimate\_run0.hs etc: sinogram output of the iterative scatter estimation
* FDG\_g1.hv: input of the simulation (i.e. ground truth image)
* FBP\_recon\_with\_scatter\_correction\_run0.hv: FBP reconstruction of the scatter corrected data

Example questions to answer:

* How close is the scatter estimate in the ideal case of a simulation and how does this effect the image reconstruction? (run0)
* How different is the scatter (and its estimate) between gate 1 and gate 2? (run0)
* How different is the scatter (and its estimate) if you have a wrong estimate of the energy dependence of the detection efficiency? (run1)
* What happens to the scatter estimate and the reconstructed image if you use the wrong attenuation image? (run3)

## Python evaluation

2 Python scripts are provided as a starting point for investigating the results.

* evaluate\_scatter\_run0.py which reads results from run\_scatter0.sh and displays them comparing with the truth (i.e. simulation input and simulation scatter output) for GATE1
* evaluate\_scatter\_run3.py reads results from run0 and run3 and displays them (also for GATE1)

## Evaluation using AMIDE

Use **AMIDE** to visualize your FBP\_recon\_with\_scatter\_correction\_run0.hv for each gate and the original simulated image. Use maximum display value 25.

Can you display the subtraction (e.g. using stir\_subtract) between the two gates?

How much motion do you see in the reconstructed images?

Extract sinograms for display with AMIDE for each of the two gates, e.g.:

cd working\_folder

extract\_segments GATE2/scatter\_estimate\_run0.hs

extract\_segments GATE2/my\_scatter\_g2.hs

Can you display the difference of the two sinograms e.g. using stir\_subtract –s? Can you see motion in the original sinograms? Scatter sinograms?

# Exercise: Motion Correction

(Always run scripts from the exercises directory. This exercise depends on the output of run\_simulations\_thorax.sh)

There are 2 scripts:

run\_MCIR\_0.sh

Correct for motion using valid motion vectors and the previously calculated scatter background

run\_MCIR\_1.sh

Do not correct for motion

Read and run scripts, e.g.:

./run\_MCIR\_0.sh &

./run\_MCIR\_1.sh &

This will run everything in the “background”, i.e. you will be immediately have the prompt back. It takes about 30seconds to complete each reconstruction. If nothing happens, you can confirm the scripts run OK:

less working\_folder/MCIR/MCIR.log

less working\_folder/noMC/noMC.log

(quit less by pressing q)

Each team uses AMIDE to visualize the corresponding images (e.g. working\_folder/MCIR/MCIR\_16.hv or a later iteration 32, 48, 64). Choose the iteration depending on the computational speed of the computer. Select a transverse slice showing the myocardium (preferably z=12.15mm). Maximum scale value: 25.

# Appendices

## Installing STIR

You will need the updated source made available via the web for this course (see the STIR website). You also need a display program, we recommend AMIDE. You will then need to unpack the zip file and add the directory to your path such that they can be found when typing a command.

If you have Linux (or vGate), use the installation script. Otherwise you need to build STIR yourself, install it, and add it to your path

Open a terminal and type something like this all on one line (adjust to where your files are):

* Linux/MacOS:  
  PATH=~/STIRShortCourse/bin/:$PATH
* Cygwin  
  PATH=~/STIRShortCourse/bin/:/cygdrive/c/Program\ Files\ \(x86\)/amide/bin/:$PATH
* Windows command terminal (if Cygwin not available)  
  PATH %HOMEDRIVE%%HOMEPATH%\Documents\STIRShortCourse\bin;c:\Program Files x86\amide\bin;%PATH%

After installing, try to type in your terminal

forward\_project

You should see a usage message. If you get an error, you probably didn’t set-up your path correctly.

## File extensions

.hv: Interfile header for image (volume)

.ahv: (ignore) old-style Interfile header for image

.v: raw data of image (in floats)

.hs: Interfile header projection data (sinograms)

.s: raw data of projection data (in floats)

.par: STIR parameter file

.sh: Shell script (sequence of commands)

.bat: Windows batch file

.log: log file (used to record output of command)

## Python:

We use Spyder as a nice IDE for Python (actually iPython which is a slightly friendlier version of Python). You need only minimal knowledge of Python for this course, but of course it would be good to read-up a bit (see below).

You will normally work by loading an example script in Spyder in the editor, executing it bit by bit, and then editing it to do some more work. Useful shortcuts for in the editor (these are in Windows-style, including the usual copy-paste shortcuts Ctrl-C and Ctrl-V).[[1]](#footnote-1)

* F9 executes the currently highlighted code.
* CTRL + <RETURN> executes the current cell (menu entry Run -> Run cell). A cell is defined as the code between two lines which start with the agreed tag #%%.
* SHIFT + <RETURN> executes the current cell and advances the cursor to the next cell (menu entry Run -> Run cell and advance).

And here are some useful ipython "magic" commands that you can use in the ipython console on the right (but not in the scripts). Most of these are identical to what you would use in the terminal.

* change to a new directory  
  cd some\_dir/another\_subdir
* change back 2 levels up  
  cd ../..
* print current working directory  
  pwd
* edit a file  
  edit FBP.par
* list files in current directory  
  ls \*.hs
* Running system commands from the ipython prompt can be done via an exclamation mark  
  !FBP2D FBP.par

One thing which might surprise you that in Python *indentation is important[[2]](#footnote-2).* You would write for instance

for z in range(0,image.shape[0]):

plt.figure()

plt.imshow(image[z,:,:])

# now do something else

Here is some suggested material on Python (ordered from easy to quite time-consuming).

* The official Python tutorial. Just read Section 1, 3, a bit of 4 and a tiny bit of 6.  
  <https://docs.python.org/2/tutorial/>
* Examples for matplotlib, the python module that allows you to make plots almost like in MATLAB  
  <https://github.com/patvarilly/dihub-python-for-data-scientists-2015/blob/master/notebooks/02_Matplotlib.ipynb>
* You could read bits and pieces of Python the Hard Way  
  <http://learnpythonthehardway.org/book/index.html>
* Google has an online class on Python for those who know some programming. This goes quite in depth and covers 2 days.  
  <https://developers.google.com/edu/python/?csw=1>

## STIR commands for evaluation:

The following is a list of commands that can be used during the exercises

extract\_segments projdata.hs

convert projection data into an (Interfile) image e.g. for display

list\_projdata\_info projdata.hs

Writes basic information about geometry etc.

Use without arguments for extra options.

list\_image\_info image.hv

Writes basic information about image geometry and values.

list\_image\_values prof.txt input\_image \

min\_plane max\_plane min\_row max\_row min\_col max\_col

(note: the backslash “\” is used in shell scripts for “line continuation”, i.e. when everything does not fit on one line)

list\_image\_values writes values to a text file (for import in Excel et al).

Indices need to be in the STIR convention (plane starts from 0, col,row are

centred around 0). Use list\_image\_info to find ranges.

Note: there is currently a bug in list\_image\_values that row (x) and column (y)

have to be given in that order (i.e. it's z,x,y while should have been z,y,x)

## Image display

Several display programs can be used. AMIDE reads the interfile volumes directly. ImageJ and others can use import of raw floats (i.e. the .v file).Settings are for instance.

Image type: 32-bit Real

Width ?

Height: ?

Offset: 0

Number of images ?

Gap between images: 0

White is 0: Ticked

Little endian: Ticked

You will have to find the data sizes from the header (the .hv file), or by using list\_image\_info.

1. Text from <http://www.southampton.ac.uk/~fangohr/blog/spyder-the-python-ide.html> [↑](#footnote-ref-1)
2. The amount of indentation is not important, as long as you are consistent (i.e. it doesn’t matter if you use 2 or 4 spaces, but you cannot mix them). [↑](#footnote-ref-2)