

Chest EIT: why it's difficult, and what we're doing about it?

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Andy Adler

Systems and Computer Engineering, Carleton University, Ottawa

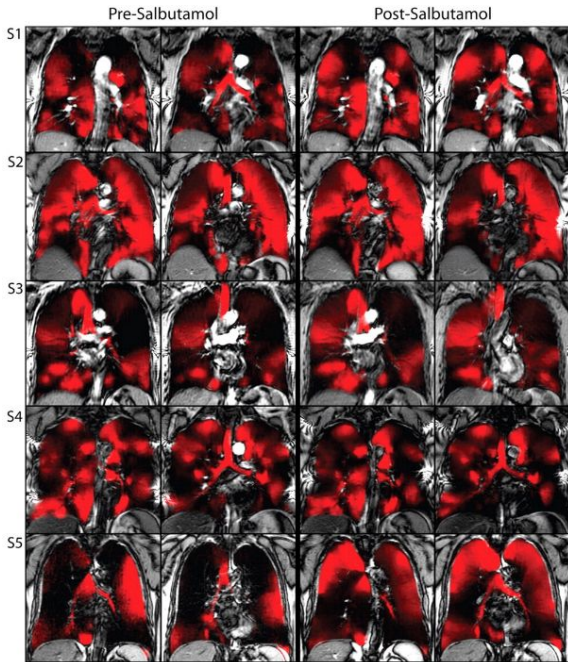
Lung Imaging

Lung Imaging

Source: Kirby et al,
Radiology 261.1 (2011)

Pre- and post-salbutamol
 ^3He MR images (red)
registered to two center
coronal thoracic ^1H MR
images (gray scale) for five
representative patients with
COPD

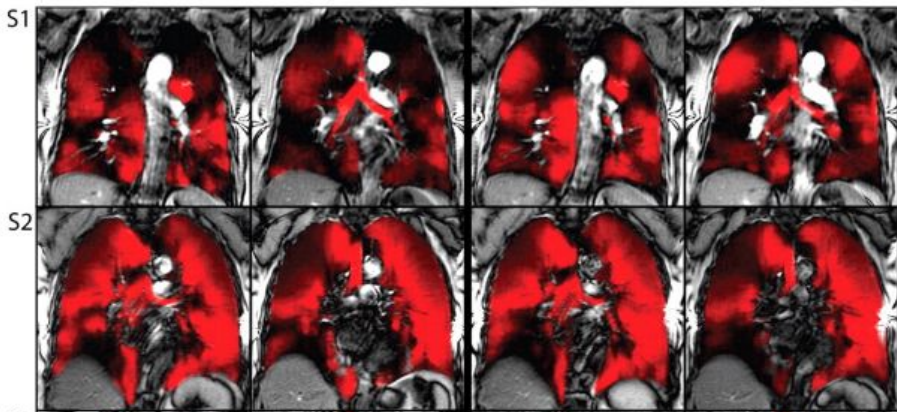
S1, S2: stage II disease,
S3, S4: stage III disease,
S5: stage IV disease.



Imaging \Rightarrow new clinical insights

Pre-Salbutamol

Post-Salbutamol

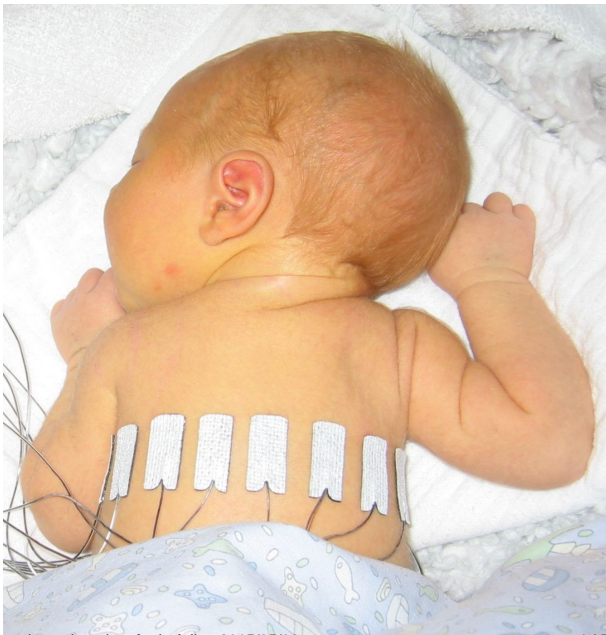


Electrical Impedance Tomography

Electrical Impedance Tomography

10-day old healthy
baby with EIT
electrodes

Source: Heinrich, Schiffmann,
Frerichs, Klockgether-Radke,
Frerichs; *Intensive Care Med*;
2006.
[eidors3d.sf.net/data_contrib/if-
neonate-spontaneous](http://eidors3d.sf.net/data_contrib/if-neonate-spontaneous)



EIT – Purpose?

EIT – Purpose?

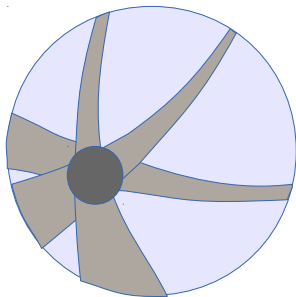
- Imaging ––

EIT – Purpose?

- Imaging --
- Monitoring ++

EIT – easy?

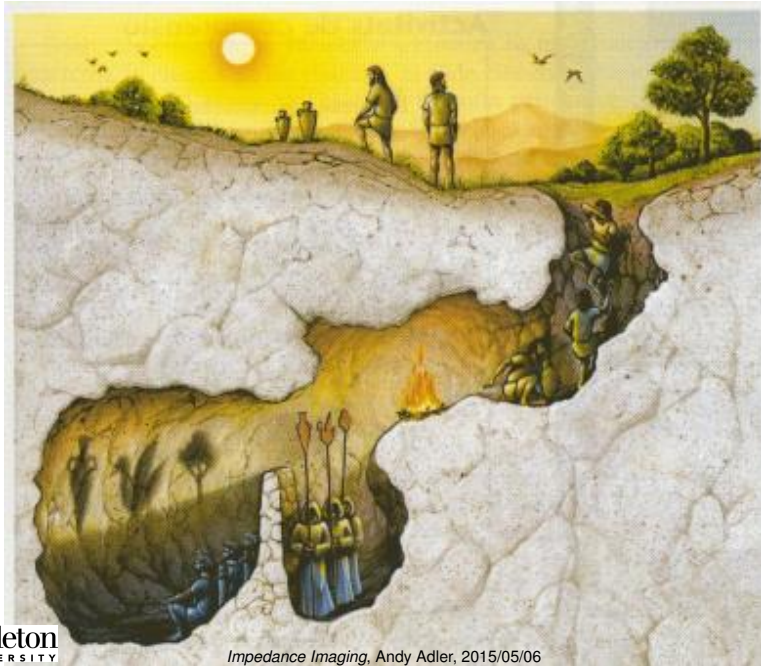
- Audio-frequency analog circuits
- Low frequency A/D converters
- mV-range voltage measurement
- Simple imaging algorithms?



EIT – easy!

Why is EIT difficult?

Inverse Problems . . . *Plato's cave*



Plato's cave . . . Shadows on the wall



Source: iamcriselleeee.files.wordpress.com/2013/11/cave-2.jpg

Inverse Problems

Forward Problem: *Forms* \Rightarrow *Shadows*

Inverse Problems

Forward Problem: *Forms* \Rightarrow *Shadows*

Inverse Problem: *Shadows* \Rightarrow *Forms*

Inverse Problems

Forward Problem: *Forms* \Rightarrow *Shadows*

Inverse Problem: *Shadows* \Rightarrow *Forms*

- Ill-conditioned
Sensitivity to some movements is low

Inverse Problems

Forward Problem: *Forms* \Rightarrow *Shadows*

Inverse Problem: *Shadows* \Rightarrow *Forms*

- Ill-conditioned
Sensitivity to some movements is low
- Ill-posed
Some movements don't change shadows

Inverse Problems

Forward Problem: *Forms* \Rightarrow *Shadows*

Inverse Problem: *Shadows* \Rightarrow *Forms*

- Ill-conditioned
Sensitivity to some movements is low
- Ill-posed
Some movements don't change shadows
- Noisy
Flickering light

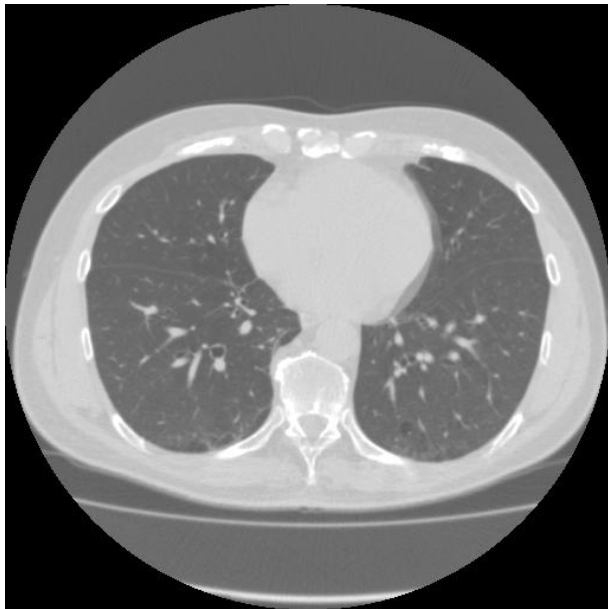
Ill-conditioned

- Low sensitivity to phenomena of interest
- High sensitivity to phenomena *not* of interest

Ill-conditioned ...

Healthy Adult
Male CT slide

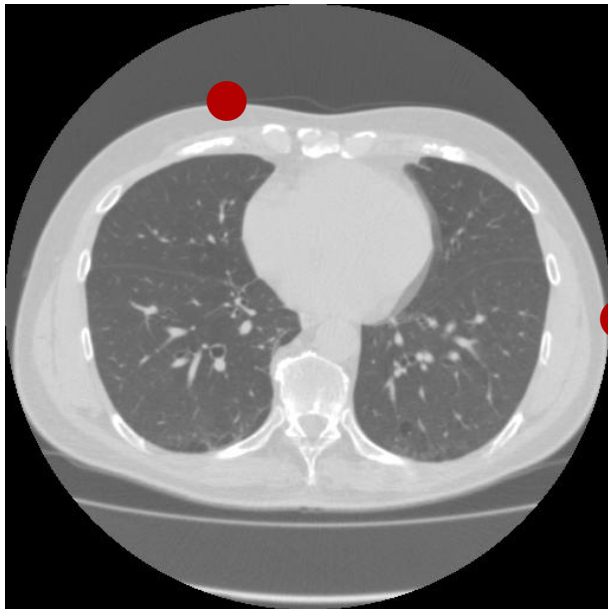
Source:
[eidors3d.sf.net/tutorial/
netgen/extrusion](http://eidors3d.sf.net/tutorial/netgen/extrusion)



Ill-conditioned ...

Healthy Adult
Male CT slide

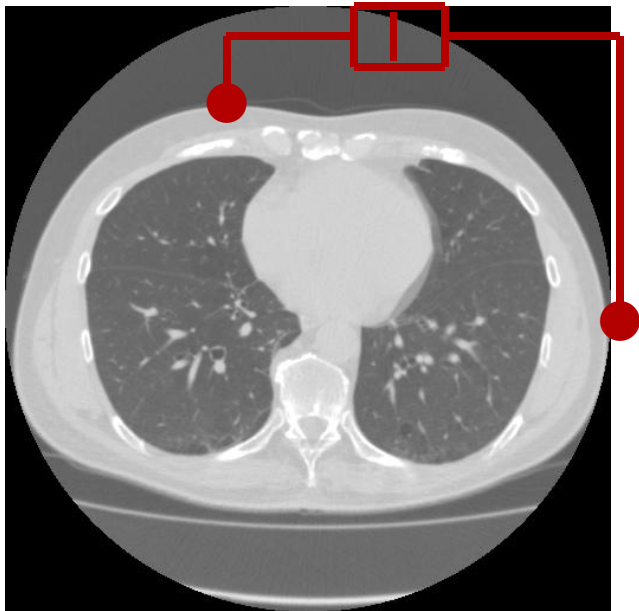
Source:
[eidors3d.sf.net/tutorial/
netgen/extrusion](http://eidors3d.sf.net/tutorial/netgen/extrusion)



Ill-conditioned ...

Healthy Adult
Male CT slide

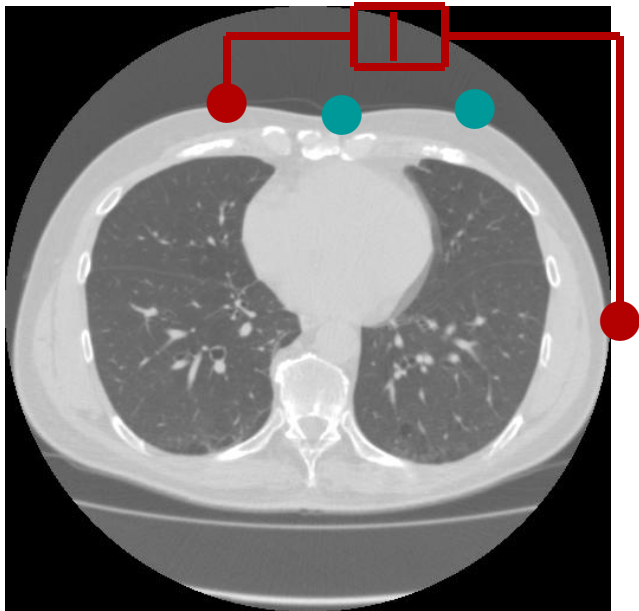
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[eidors3d.sf.net/tutorial/
netgen/extrusion](http://eidors3d.sf.net/tutorial/netgen/extrusion)



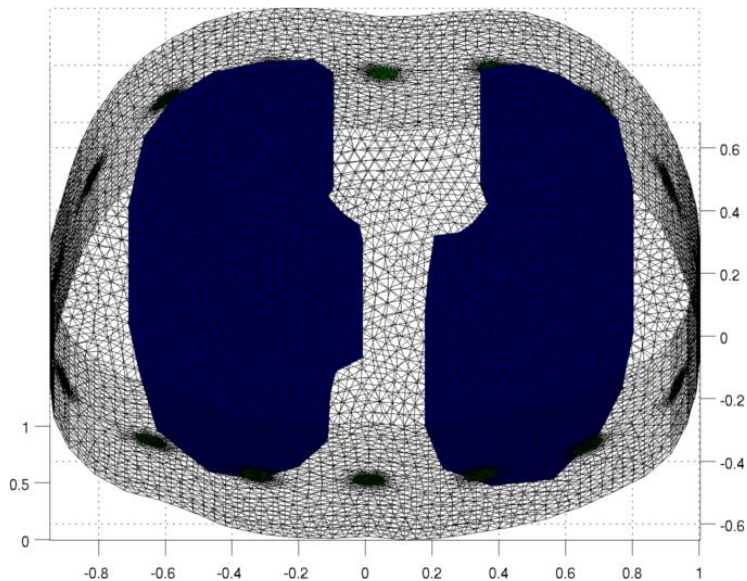
Ill-conditioned ...

Healthy Adult
Male CT slide

Source:
[eidors3d.sf.net/tutorial/
netgen/extrusion](http://eidors3d.sf.net/tutorial/netgen/extrusion)



Finite Element Modelling



Finite Element Modelling

Step 1: Create Finite Element Model

```
##### THORAXMDL01
% get contours
thorax = shape_library('get','adult_male','boundary');
rlung = shape_library('get','adult_male','right_lung');
llung = shape_library('get','adult_male','left_lung');
% one could also run:
% shape_library('get','adult_male');
% to get all the info at once in a struct

% show the library image
shape_library('show','adult_male');
print_convert thoraxmdl01a.jpg '-density 100'
##### THORAXMDL02
shape = { 1, % height
         {thorax, rlung, llung}, % contours
         [4,50], % perform smoothing with 50 points
         0.04}; % small maxh (fine mesh)

elec_pos = [ 16, % number of elems per plane
            1, % equidistant spacing
            0.5]'; % a single z-plane

elec_shape = [0.05, % radius
             0, % circular electrode
             0.01 ]'; % maxh (electrode refinement)

fmdl = ng_mk_extruded_model(shape, elec_pos, elec_shape);
% this similar model is also available as:
% fmdl = mk_library_model('adult_male_16el_lungs');

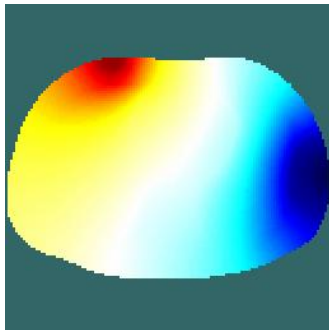
[stim,meas_sel] = mk_stim_patterns(16,1,[0,1],[0,1],{'no_meas_current'}, 1);
fmdl.stimulation = stim;

img=mk_image(fmdl,1);
img.elem_data(fmdl.mat_idx{2})= 0.3; % rlung
img.elem_data(fmdl.mat_idx{3})= 0.3; % llung

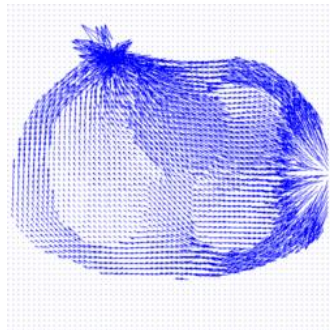
clf; show_fem(img); view(0,70);
print_convert thoraxmdl02a.jpg '-density 100'
```



Finite Element Modelling



Simulated Voltages



Voxel Currents

Finite Element Modelling

Step 2: Forward solve and current streamlines

```
img_v = rmfield(img, 'elem_data');
img_v.node_data = vh.volt(:,1);
img_v.calc_colours.npoints = 128;

PLANE= [inf,inf,0.35]; % show voltages on this slice

subplot(221);
show_slices(img_v,PLANE); axis off; axis equal
print_convert thoraxmdl03a.jpg
%
%%%% THORAXMDL04
img_v = img;
img_v.fwd_model.mdl_slice_mapper.npx = 64;
img_v.fwd_model.mdl_slice_mapper.npy = 64;
img_v.fwd_model.mdl_slice_mapper.level = PLANE;
q = show_current(img_v, vh.volt(:,1));
quiver(q.xp,q.yp, q.xc,q.yc,10,'b');
axis tight; axis image; ylim([-1 1]);axis off
print_convert thoraxmdl04a.jpg

%%%% THORAXMDL05
img_v.fwd_model.mdl_slice_mapper.npx = 1000;
img_v.fwd_model.mdl_slice_mapper.npy = 1000;
img_v.fwd_model.mdl_slice_mapper.level = PLANE;

% Calculate at high resolution
q = show_current(img_v, vh.volt(:,1));

pic = shape_library('get','adult_male','pic');
imagesc(pic.X, pic.Y, pic.img);
% imgt= flipdim(imread('thorax-mdl.jpg'),1); imagesc(imgt);
colormap(gray(256)); set(gca,'YDir','normal');
hold on

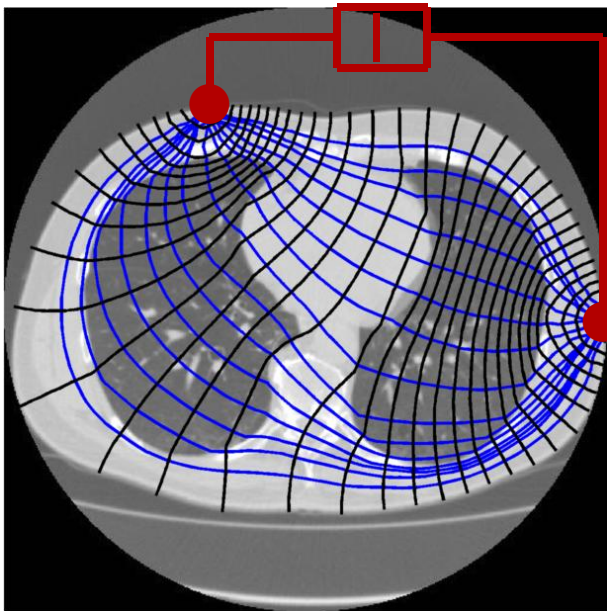
sx = linspace(-.5,.5,15)';
sy = 0.05 + linspace(-.5,.5,15)';
hh-streamline(q.xp,q.yp, q.xc, q.yc,sx,sy); set(hh,'Linewidth',2, 'color','b');
hh-streamline(q.xp,q.yp,-q.xc,-q.yc,sx,sy); set(hh,'Linewidth',2, 'color','b');

axis equal; axis tight; axis off; print_convert thoraxmdl05a.jpg
```



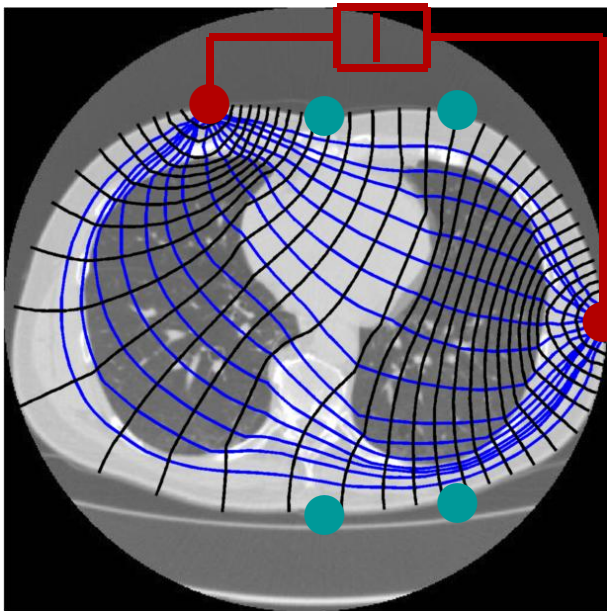
Thorax Propagation

CT Slice with
simulated current
streamlines and
voltage
equipotentials



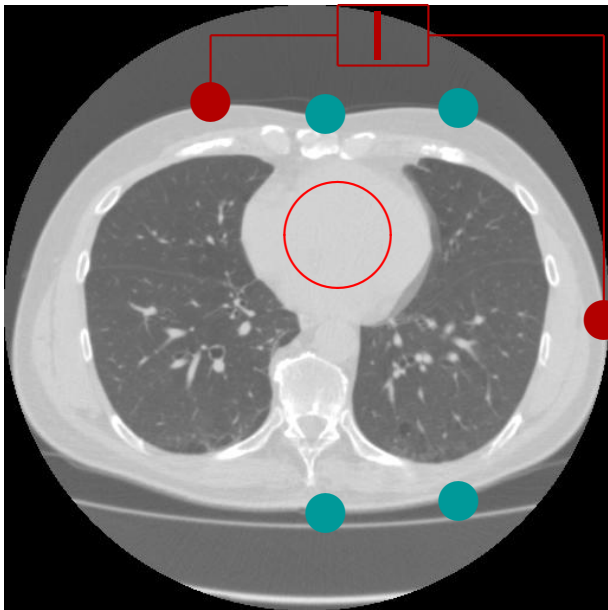
Thorax Propagation

CT Slice with
simulated current
streamlines and
voltage
equipotentials



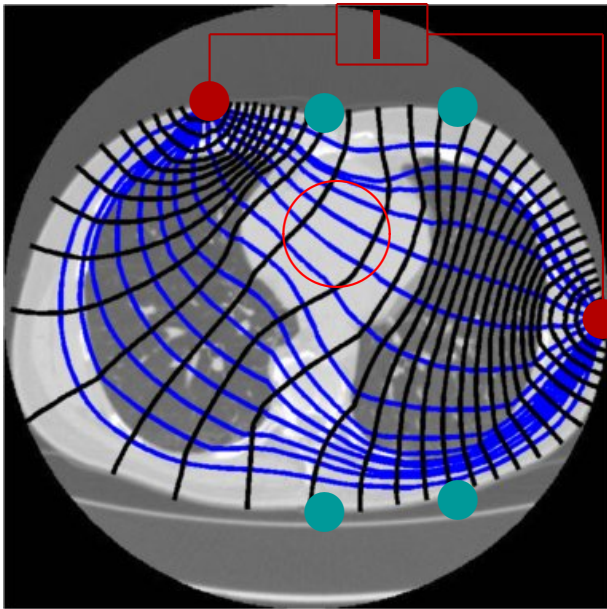
Changing Conductivity

Heart receives
blood (diastole)
and is more
conductive

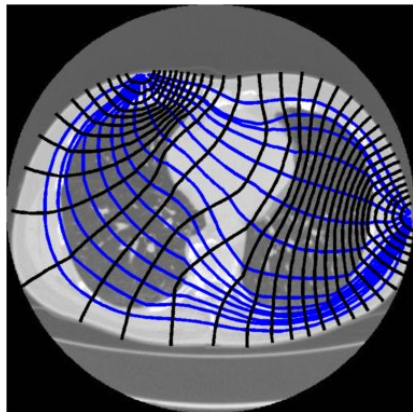
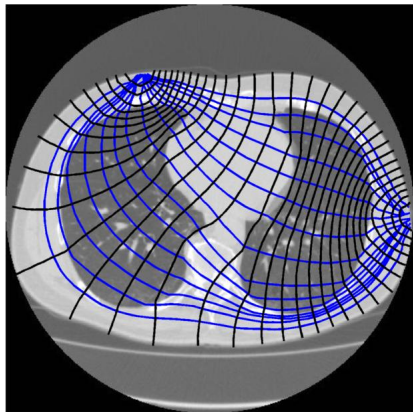


Changing Conductivity

Heart receives
blood (diastole)
and is more
conductive



Ill-conditioned: low sensitivity



Finite Element Modelling

Step 3: Change conductivity and re-solve

```
select_fcn = inline('(x-0.1).^2+(y-0.3).^2<0.2^2','x','y','z');
memb_frac = elem_select( img.fwd_model, select_fcn);
img-mk_image(fmdl,1);
img.elem_data = 1 + memb_frac*2.0;
img.elem_data(fmdl.mat_idx(2))- 0.3; % rlung
img.elem_data(fmdl.mat_idx(3))- 0.3; % llung
show_fem(img)
img_v = img;
% Stimulate between elecs 16 and 5 to get more interesting pattern
img_v.fwd_model.stimulation(1).stim_pattern = sparse([16;5],1,[1,-1],16,1);
img_v.fwd_solve.get_all_meas = 1;
vh = fwd_solve(img_v);

img_v = rmfield(img, 'elem_data');
img_v.node_data = vh.volt(:,1);
img_v.calc_colours.npoints = 128;

PLANE= [inf,inf,0.35]; % show voltages on this slice
```

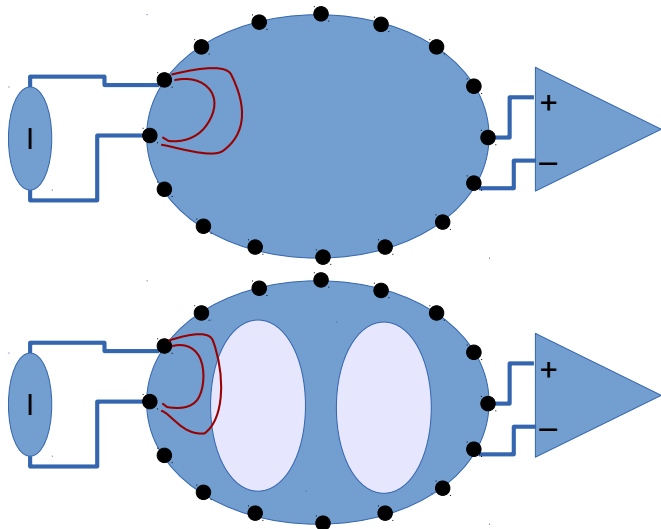
```
clf
imagesc(pic.X, pic.Y, pic.img); colormap(gray(256)); set(gca,'YDir','normal');
hh-streamline(q.xp,q.yq, q.xc, q.yc,sx,sy); set(hh,'Linewidth',2);
hh-streamline(q.xp,q.yq,-q.xc,-q.yc,sx,sy); set(hh,'Linewidth',2);

[x y] = meshgrid( linspace(pic.X(1), pic.X(2),size(imgs,1)), ...
                 linspace(pic.Y(2), pic.Y(1),size(imgs,2)));

hold on;
contour(x,y,imgs,31);
hh= findobj('Type','patch'); set(hh,'Linewidth',2)

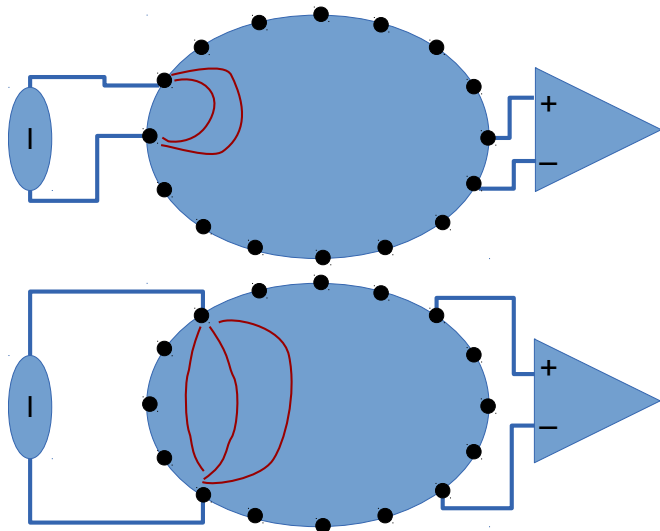
hold off; axis off; axis equal; %ylim([50,450]);
print_convert thoraxmdl07a.jpg
```

Non-conductive lungs decrease sensitivity



Relative sensitivity = $2.8\times$

Own goal: adjacent stimulation and measurement



Relative sensitivity = $20.6\times$

Finite Element Modelling

CODE: Adjacent vs. Skip 4

```
extra={'ball','solid ball - sphere(0,0,1;0.1);'}
fmdl= ng_mk_cyl_models(2,[16,1],[.05],extra);
fmdl.stimulation = mk_stim_patterns(16,1,[0,1],[0,1],{ },1);
img= mk_image(fmdl,1);          vh = fwd_solve(img);
img.elem_data(fmdl.mat_idx(2)) = 2;  vi = fwd_solve(img);
s1 = norm(vh.meas - vi.meas);

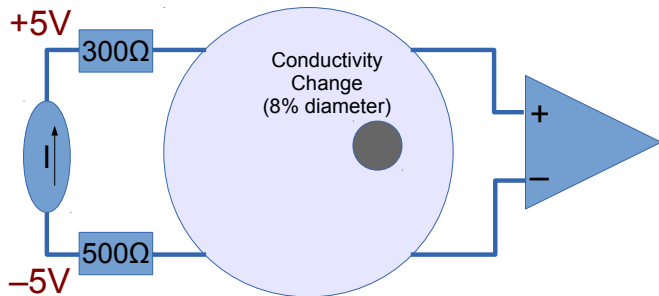
fmdl.stimulation = mk_stim_patterns(16,1,[0,5],[0,5],{ },1);
img= mk_image(fmdl,1);          vh = fwd_solve(img);
img.elem_data(fmdl.mat_idx(2)) = 2;  vi = fwd_solve(img);
s2 = norm(vh.meas - vi.meas);

disp(s2/s1)
```

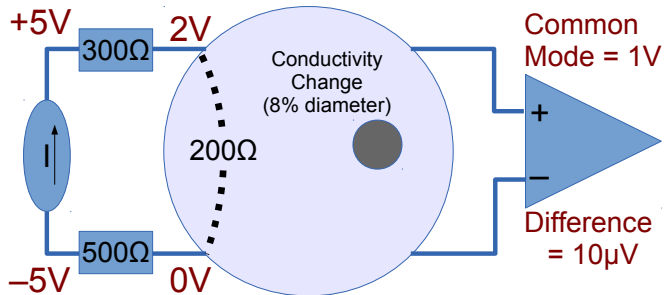
Ill-conditioned

- Low sensitivity to phenomena of interest
- High sensitivity to phenomena *not* of interest

EIT Instrumentation



EIT Instrumentation



Amplifier Requirements

- Common Mode Rejection Ratio (CMRR)
- $\text{CMRR} = \frac{1\text{V}}{10\mu\text{V}} = 10^5 = 100 \text{ dB}$

Amplifier Requirements

- Common Mode Rejection Ratio (CMRR)
- $CMRR = \frac{1V}{10\mu V} = 10^5 = 100 \text{ dB}$



Low Cost Low Power Instrumentation Amplifier

AD620

FEATURES

Easy to use

- Gain set with one external resistor
(Gain range 1 to 10,000)
- Wide power supply range ($\pm 2.3 \text{ V}$ to $\pm 18 \text{ V}$)
- Higher performance than 3 op amp IA designs
- Available in 8-lead DIP and SOIC packaging
- Low power, 1.3 mA max supply current

Excellent dc performance (B grade)

- 50 μV max, input offset voltage
- 0.6 $\mu\text{V}/^\circ\text{C}$ max, input offset drift
- 1.0 nA max, input bias current
- 100 dB min common-mode rejection ratio ($G = 10$)

Low noise

- 9 nV/ $\sqrt{\text{Hz}}$ @ 1 kHz, input voltage noise
- 0.28 μV p-p noise (0.1 Hz to 10 Hz)

Excellent ac specifications

- 120 kHz bandwidth ($G = 100$)
- 15 μs settling time to 0.01%

CONNECTION DIAGRAM

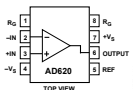


Figure 1. 8-Lead PDIP (N), CERDIP (Q), and SOIC (R) Packages

PRODUCT DESCRIPTION

The AD620 is a low cost, high accuracy instrumentation amplifier that requires only one external resistor to set gains of 1 to 10,000. Furthermore, the AD620 features 8-lead SOIC and DIP packaging that is smaller than discrete designs and offers lower power (only 1.3 mA max supply current), making it a good fit for battery-powered, portable (or remote) applications.

The AD620, with its high accuracy of 40 ppm maximum

100 dB min common-mode rejection ratio

Amplifier Requirements



Low Cost Low Power Instrumentation Amplifier

AD620

FEATURES

Easy to use
Gain set with one external resistor
(Gain range 1 to 10,000)
Wide power supply range (±2.3 V to ±18 V)
Higher performance than 3-opp amp IA designs
Available in 8-lead DIP and SOIC packages
Low power, 1.3 mA max supply current
Excellent dc performance (B grade)
50 μ V max. input offset voltage
0.6 μ V/°C max. input offset drift
1.0 nA max. input bias current
100 dB min. common-mode rejection ratio (CMRR = 10)
Low noise

CONNECTION DIAGRAM



Figure 1. 8-Lead PDIP (N), CDIP (Q), and SOIC (R) Packages

PRODUCT DESCRIPTION

The AD620 is a low cost, high accuracy instrumentation amplifier that requires only one external resistor to set gains of 1 to 10,000. Furthermore, after 0.1% tolerance 0.1% μ V/°C and

100 dB min common-mode rejection ratio

Amplifier Requirements



Low Cost Low Power
Instrumentation Amplifier

AD620

FEATURES

- Easy to use
- Gain set with one external resistor
(Gain range 1 to 10,000)
- Wide power supply range (±2.3 V to ±18 V)
- Higher performance than 3 op amp IA designs
- Available in 8-lead DIP and SOIC packages
- Low power, 1.3 mA max supply current
- Excellent dc performance (B grade)
 - 50 μ A max. input offset voltage
 - 0.6 μ V/°C max. input offset drift
 - 1.0 nA max. input bias current
 - 100 dB min common-mode rejection ratio (G = 10)
- Low noise

CONNECTION DIAGRAM

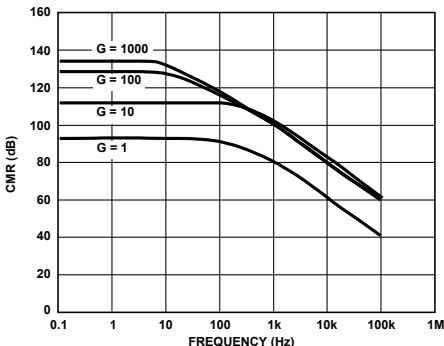


Figure 1. 8-Lead PDIP (N), CDIP (Q), and SOIC (R) Packages

PRODUCT DESCRIPTION

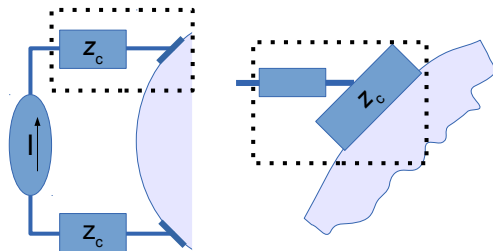
The AD620 is a low cost, high accuracy instrumentation amplifier that requires only one external resistor to set gains of 1 to 10,000. For information, refer to [AD620](#) Evaluation Kit, Rev. 0, and [AD620](#) and [AD620](#) Datasheets.

100 dB min common-mode rejection ratio

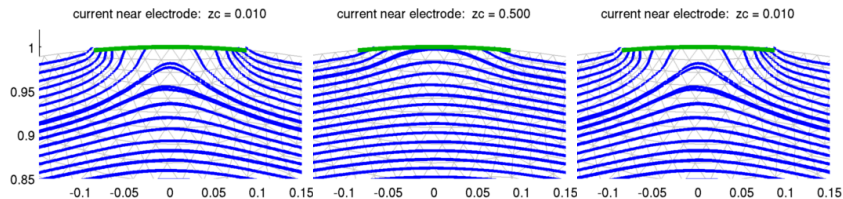
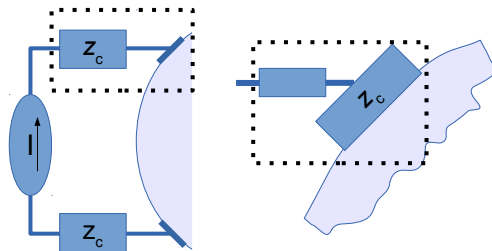


00775-0-016

What is contact impedance?



What is contact impedance?



More challenges

- Crosstalk
- Amplifier input impedance
- ...

More challenges

- Crosstalk
- Amplifier input impedance
- ...

Question:

- Geophysicists wont reuse a “hot” electrode for a few minutes. It becomes polarized. Why do we?

Inverse Problems: EIT

Why is EIT difficult?

Inverse Problems: EIT

Why is EIT difficult?

What are we doing about it?

Inverse Problems: EIT

Why is EIT difficult?

What are we doing about it?

We = the EIT community

Inverse Problems: EIT

Why is EIT difficult?

What are we doing about it?

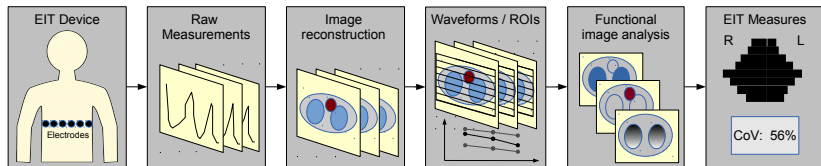
We = the EIT community

- Imaging --
- Monitoring ++

What are we doing about it?

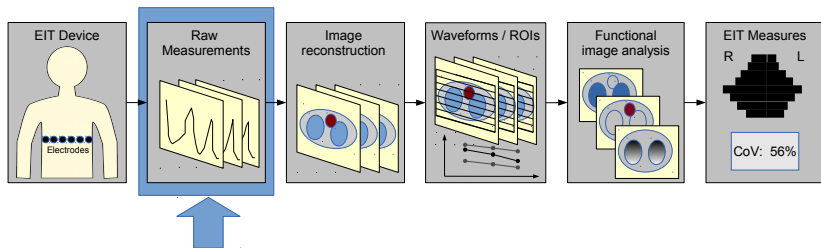
- Understanding EIT's sensitivity and limitations
- Developing novel measurement protocols and analysis methodologies

Analysis “pipeline”



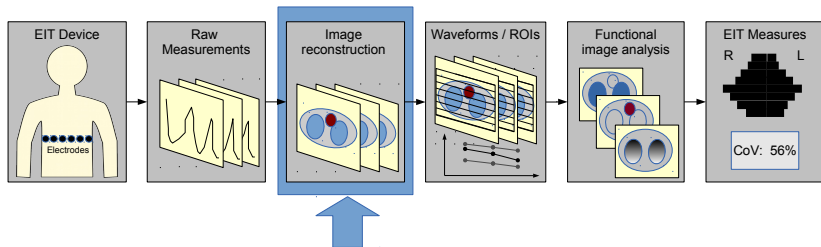
(Thanks to Inéz Frerichs for helping elaborate this concept)

Raw Measurement



- Where to place electrodes?
- “Off-plane” contributions?
- Frame rate?
- Stimulation / Measurement patterns?

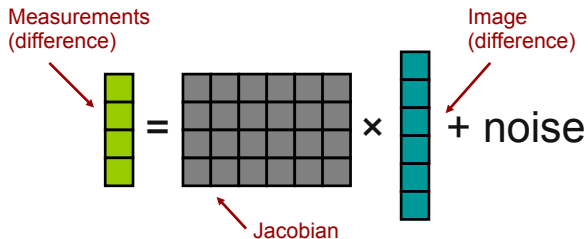
Raw Image Reconstruction



- Selection of reference/baseline
- Mixed absolute/difference algorithms
- Electrode position/movement
- Electrode errors
- Data Quality

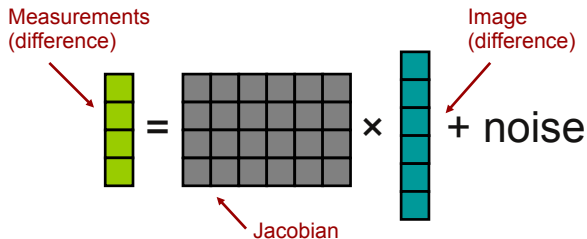
Reconstruction in Pictures

- Forward Problem

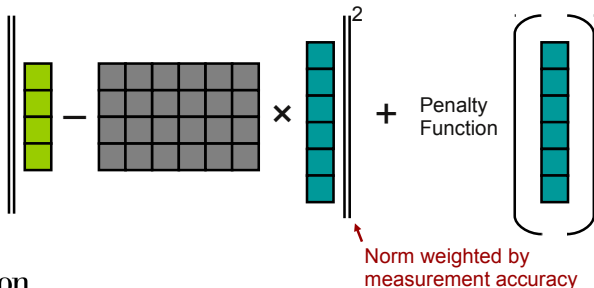


Reconstruction in Pictures

- Forward Problem

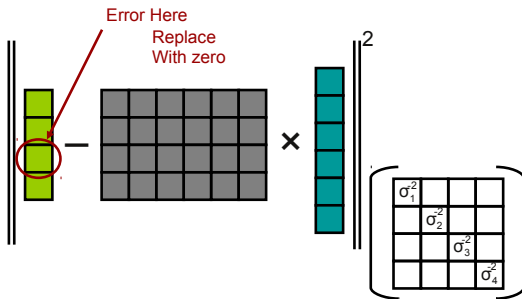


- Linear Solution: Minimize norm



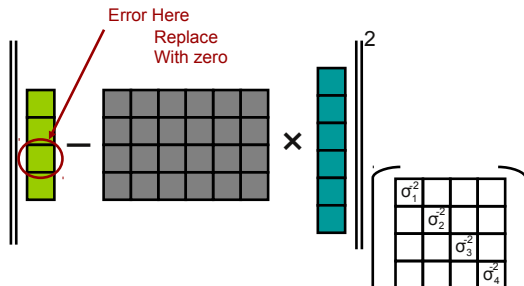
Reconstruction with Data Errors

“Traditional”
Solution

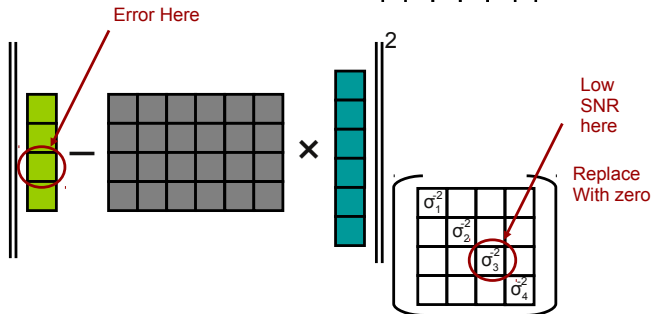


Reconstruction with Data Errors

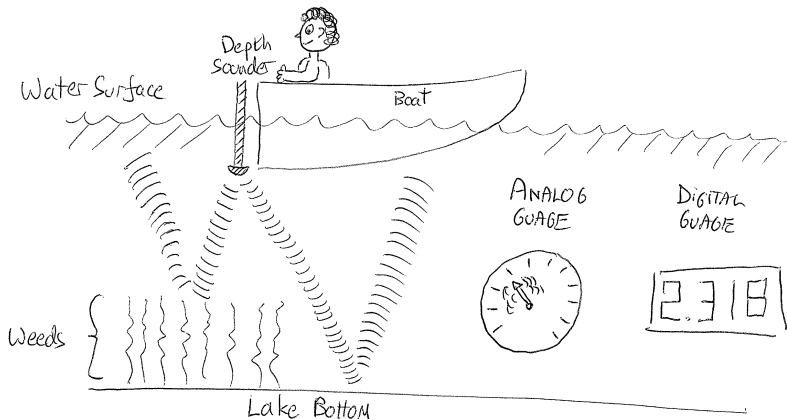
“Traditional”
Solution



Error Model
Solution



Data Quality

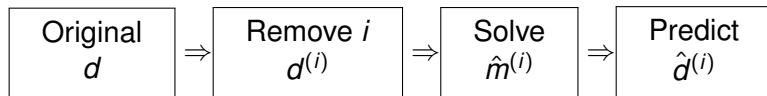


Depth Sounder – with analog and digital gauges

Data Quality Measure: Concept

Concept: High Quality Data are Consistent

Idea: predict each data point from all others

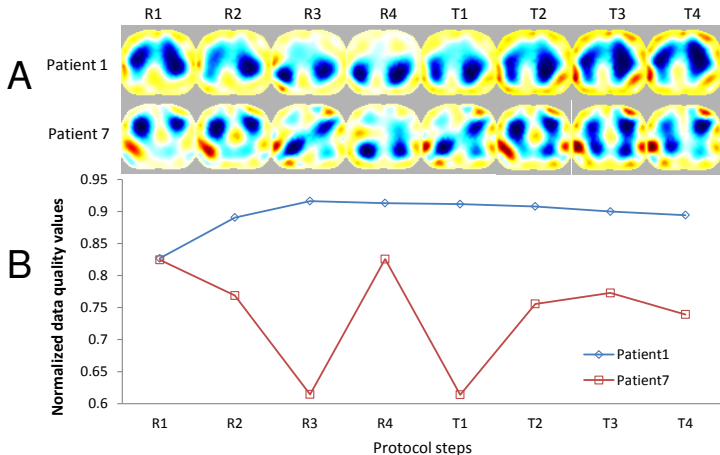


Calculate error

$$\epsilon_i = d_i - \hat{d}_i^{(i)}$$

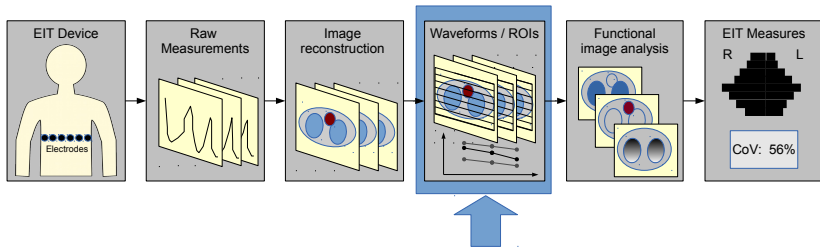
Example: Data quality measures

(R1–R4 — recruitment: PEEP \uparrow , T1–T4 — titration: PEEP \downarrow).



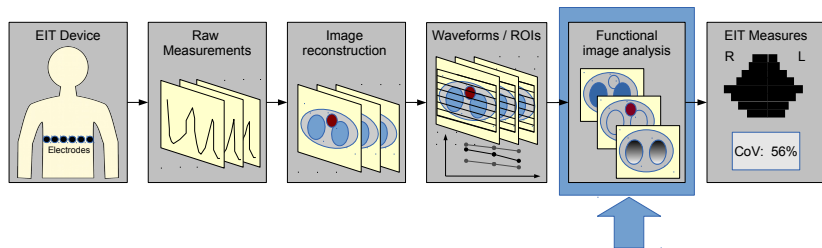
Clinical data (Wolf, *Pediatr Crit Care Med*, 2012) and data quality metric for each stage of the protocol. A: EIT, B: data quality.

ROIs and Waveforms



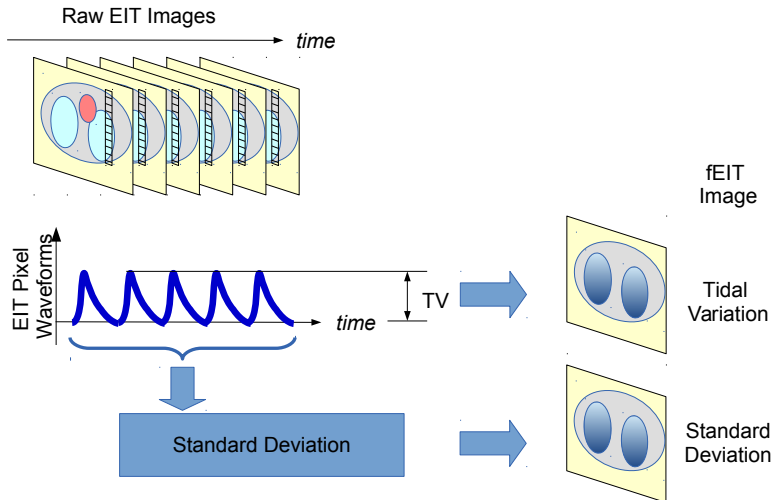
- Lung ROIs
- Anterioposterior direction

function EIT – fEIT

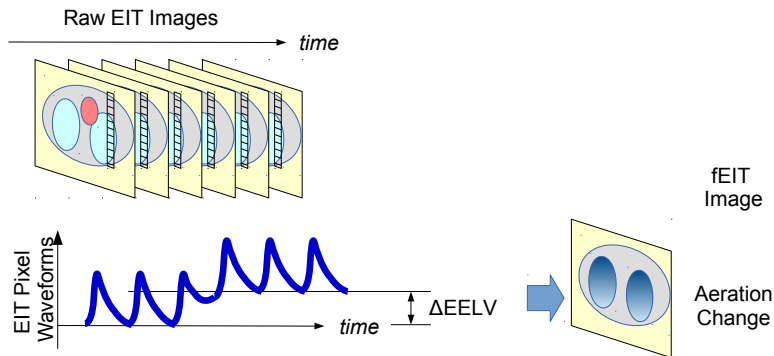


- From time sequence – generate image representing specific physiology

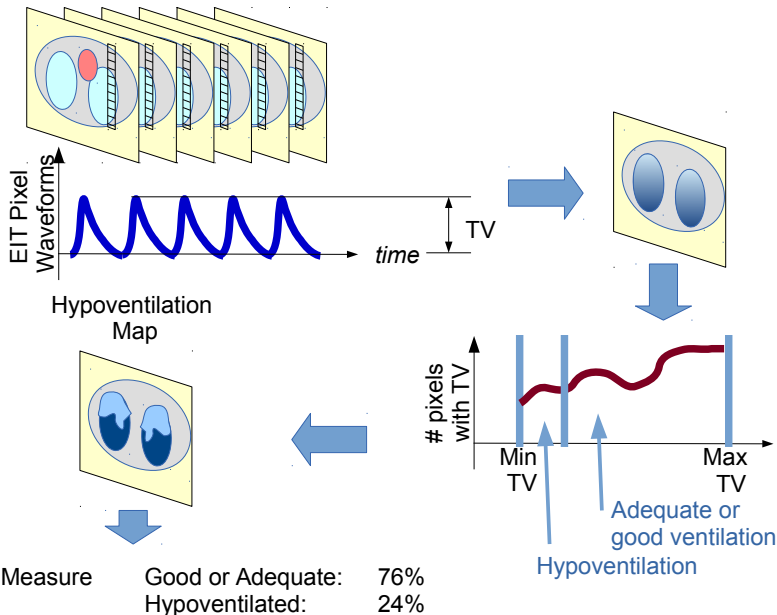
functional EIT (fEIT) images (Tidal Variation)



functional EIT (fEIT) images (Aeration Change)



functional EIT (fEIT) images (Hypo-ventilated lung)



functional EIT (fEIT) images (Hypo-ventilated lung)

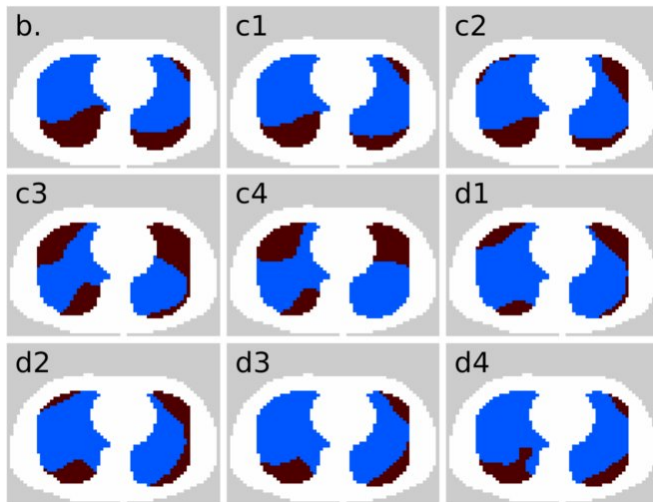
Calculate TV and threshold

CODE:

```
path= 'DATA/STUDYNAME/SUBJECT_1/YYYYMMDD/Eit/Viasys/'; dd= dir([path,'*.get']);
for idx = 1:length(dd);
    vv= eidors_readdata([path,dd(idx).name]);
    vr = mean(vv,2);
    img= inv_solve(imdl, vr, vv);
    img.calc_colours.backgnd=[1,1,1]*.8;
    ROI0 = ones(1,size(img.elem_data,1));
    [einsp,eexpi]= find_frc( img, ROI0, 13, '', 2);
    imgc = calc_slices(img); imgc(imgc==NaN) = 0;
    TV   = imgc(:, :, eexpi) - imgc(:, :, einsp);
    TV   = mean(TV,3).*ROI; % Air is +
    out = -ROI/2; % blue lungs
    out( (TV<.2*max(TV(:))&(ROI==1) ) ) = 1;
    out = calc_colours(out, img); out(55:64, :)= [];
    subplot(3,3,idx); image(out); axis off
    text(3,7,dd(idx).name(6:7), 'FontSize', 16);
end

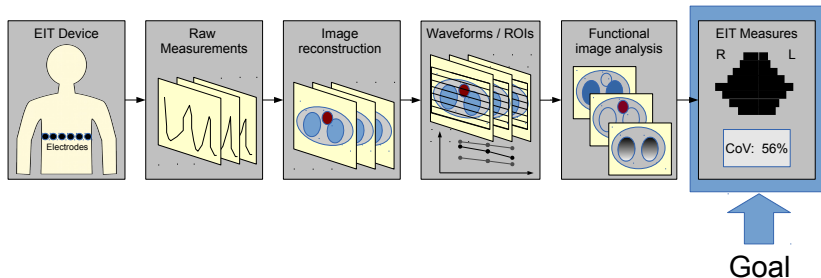
opt.vert_cut = 10; opt.vert_space = 5;
opt.horz_cut = 10; opt.horz_space = 5;
print_convert('analyse_step04a.jpg',opt);
```

fEIT image: Hypo-ventilated lung



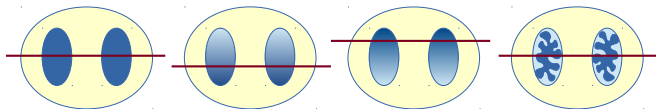
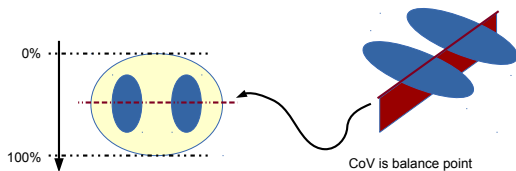
c1–c4 — recruitment: PEEP \uparrow , d1–d4 — titration: PEEP \downarrow

EIT Measures



- Average regional fEIT measures
- Characterizations of the spatial distribution of ventilation
- Examination-specific measures

#2A Measure of vertical distribution of ventilation



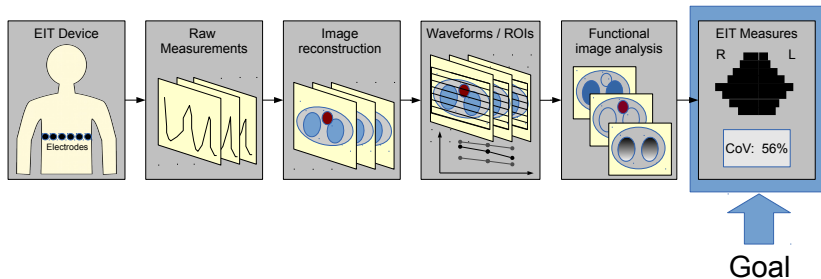
Measures of the anteroposterior distribution of ventilation

1. Center of Ventilation
2. Upper to Lower Ventilation Ratio¹

¹not recommended

Why is it difficult?
What are we doing about it?

EIT Measures



- Average regional fEIT measures
- Characterizations of the spatial distribution of ventilation
- Examination-specific measures
 - EIT measures using simultaneously measured signals
 - EIT measures using specific examination

Thank you



Traffic jam on the way to Carleton

Abstract: Currently, the most successful application of EIT is for imaging the thorax, where large movements in conductivity contrasting air and blood can be imaged over time. EIT imaging is difficult due to its low sensitivity to contrasts deep in the body, because of the diffuse nature of current propagation. EIT is thus sensitive to electrode properties, data quality, hardware imperfections and patient movement. To address these issues, several innovative strategies to analyze and interpret these data have been developed of the past few years. We introduce the concept of an EIT analysis pipeline, with stages of: measurements, raw images, waveforms, fEIT images, and measures. We will discuss recent progress in imaging the chest with EIT, and the image generation and interpretation strategies which are required.