Cardiac NM

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Nuclear Cardiology

• RNV
  – Tc-99m labelled RBCs

• MPS
  – TI-201
  – Tc-99m MIBI
  – Tc-99m tetrofosmin
  – Rb-82

• Others
  – I-123 MIBG
  – C-11 fatty acids
  – Tc-99m HL91
  – F-18 FDG
Is there any money in it

- In E.U. about 300,000 cardiac studies per year
- In USA about 1,000,000 cardiac studies per year
- Turn over of nuclear medicine radiopharmaceuticals $41,000,000 pwe annum
The perfect agent

- High myocardial uptake
- Uptake proportional to blood flow
- Low background
- Good dosimetry
- Washout or no washout? But stable distribution from injection to imaging
TI-201

• Thallium-201 is a metallic element in group IIIA of the periodic table
• Biological similarities to potassium
• Monovalent cation
• Cyclotron produced
• Decays by electron capture (EC) to mercury
• Physical half-life is 73.1 hours
• Biological half-life is 10 days
General principles

• Use as much activity as you can justify for dosimetry not finances
• Do not exceed recommended maximum activity of Tc-99m added to vial
• Image at correct time post injection
TI-201

- Emits predominantly mercury x-rays at 69-83KeV (88%)
- Principle photo peaks at 135 and 167 keV(12%)
- No in-house preparation and no quality control required before injection
- First pass extraction is 60-70% (stress injection), 80-90% (rest injection)
- Maximum myocardial uptake is approximately 3.7-4% of injected dose
- Extraction decreases at higher flow rates
TI-201

• Accumulation and retention depends on coronary blood flow and cellular viability
• Redistributes
• Critical organs: Ovaries, kidneys, bone surface
• Myocardial uptake has two components: early component (80% with half life of 4 hours) and delayed component (20% with half life of 40 hours)
• Disappearance from the blood compartment rapid and has two components: 92% disappears with a half life of 5 minutes and 8% with a half life of 40 hours
• Lung uptake proportional to LVEDP
TI-201

• Requires stringent imaging protocol especially stress images must be started by 5 minute p.i. not to loose sensitivity but at that time patient may not have recovered from stress
• Whole body effective dose is 0.22mSv/MBq (80MBq study =18mSv)
• Excretion: Faeces (80%), urine (20%)
TI-201

- **Limitations:** Relatively long physical half life [high radiation burden], relatively low injected activity [low-signal to noise ratio, sub optimal images (obese), low counts levels [impairs high quality ECG-gating SPECT], relatively low energy emission [low resolution images and attenuation by soft tissues]
Tc-99m MIBI

- Lipophilic monovalent cation
- Generator produced
- Emits gamma rays at 140 KeV
- Physical half-life is 6 hours
- The biological half-life is 680 +/- 45 minutes
- Favourable myocardium to background radiation for myocardial imaging
Tc-99m MIBI

- High energy photons decreases the problems of photon attenuation
- Distribution depends on plasma membrane and mitochondrial membrane potentials
- Once accumulated, it is bound in a relatively stable fashion
- Redistribution is negligible or partial [10-15%] (does not redistribute to a degree that can be imaged clinically)
Tc-99m MIBI

- First pass extraction fraction is approximately 60% (rest injection), 40% (stress injection)
- 1-2% of the injected dose localizes to the myocardium at rest
- Higher dose compensates for lower extraction
- Flexible imaging protocols
- At higher flow rates there is a plateau in extraction and at low flow rates extraction reduces
Tc-99m MIBI

- Whole body effective dose is 0.009mSv/MBq at rest, 0.008mSv/MBq at stress
- In a 1000MBq study the effective dose will be 8.7mSv
- The primary route of excretion is hepatobiliary (33%). Need to wait 60 mins p.i. for non interference of liver uptake
- Critical organs: Gall bladder, kidneys and colon
- Preparation takes longer (includes boiling in water bath for 20 mins minimum)
Tc-99m tetrofosmin (where different from Tc-99mMIBI

- Hepatic uptake is lower than with Tc-Sestamibi and it also clears more quickly therefore can start imaging at 30mins pi
- First pass extraction fraction is about 54%
- The myocardial extraction plateaus at higher flow rate
- Overestimates flow at low flow rates
Tc-99m tetrofosmin

- Whole body effective dose is 0.008 mSv/MBq at rest, 0.007 mSv/MBq at stress
- In a 1000MBq study the effective dose will be 7.5 mSv
- Preparation requires no boiling in water bath
Other agents

• Tc-99m Teboroxime
  – Rapid uptake and clearance imaging must be finished within 12 minutes p.i. needs triple headed camera

• Tc-99m Neot
  – Like MIBI but killed rats!!

• Tc-99m furifosmin
  – Rumoured to be even more deadly than Neot
Practical differences

- **TI-201**
  - Single injection
  - Low energy X-rays
  - Patient must be stressed next to camera
  - Maybe better looking at viability

- **Tc-99m MIBI/TF**
  - Double injection
  - Higher energy gamma rays
  - Patient can be stressed away from camera
  - Better for gating
Lung TI-201 uptake proportional to LVEDP
Protocol for TI-201

- **Stress**: 0 min
- **Stress imaging**: 5 min
- **Rest/redistribution imaging**: 3-4 hours

74-105 MBq TI-202
Protocol for Tc-99m MIBI/TF

one day

Stress            Stress imaging        Rest imaging

0 min       45-60 min                           2hours            45-60 min
Fatty meal
Fatty meal

250 MBq
Tc-99m
MIBI/TF

750 MBq
Tc-99m
MIBI/TF
Protocol for Tc-99m MIBI/TF
two days

Stress            Stress imaging

0 min       45-60 min                                                   45-60 min

Fatty meal

500 MBq
Tc-99m
MIBI/TF

>24
hours

Rest imaging

45-60 min

Fatty meal

500 MBq
Tc-99m
MIBI/TF
Why perform a stress test

- At stress, blood flow in the heart increases.
- Cannot occur in stenotic vessels.
- Less flow down these arteries (defect on scan).
- Returns to normal at rest (no defect on scan).
Stages of the Bruce protocol
Pharmacological stress

• Why might pharmacological stress be so useful?
• Not all patients can perform physical stress
  – The elderly
  – Arthritic
  – PVD, diabetes
  – Anaemic
  – Unfit
Pharmacological stress

• Main agents used
• A2 receptors
  – Dipyridamloe
  – Adenosine
• B1 receptors
  – Dobutaomine
  – Arbutamine
Why do vasodilators work

• Coronary artery vasodilataion is normal response to stress
• Increases flow during short diastole
• Mediated via cAMP
• Atherosclerotic vessels do not dilate
• Steal of blood from these vessels
• >50% drop in perfusion seen on SPECT (>90% needed for ecg)
How Vasodilators work
Effect of Dipyridamole on blood flow
Adenosine

- Works on A2 receptors directly
- Plasma half life 20-60 seconds
- Vasodilates via cAMP
- Contraindications
  - Asthma
  - 3rd degree heart block
- Relative contraindications
  - Wheezy bronchitis
  - 2nd degree heart block
Dobutamine

- Works at beta 1 receptors
- Increase stroke volume > HR
- No need to reach Max predicted HR
- Contra-indications
  - Recent VF
  - Allergy to dobutamine
  - Critical aortic stenosis
- Relative contra-indications
  - Recent VT
Acquiring the image

• As per previous talk
• Remember need to have good quality image for diagnosis
• Look for patient movement correct or re-scan
• Look for attenuation
  – Breast-ant wall women
  – Diaphragm inf wall men
Inferior attenuation in a man

Image to left shows unattenuated image with apparent inferior defect in inferior wall. Above is the same patient after attenuation correction.
Quantitative cardiac SPECT

- Compares results of patient’s scan with normal data base
- Controlled for gender, age, (occ race) tracer injection time and activity, also stress/rest
- Assigns >-2.5 s.d as defect
- Assigns improvement of >+1.5 s.d as reperfusion
Smei-quantification: Forming a bulls eye plot

Sequential rings of data used to build the bulls eye plot
Polar “bullseye” plots - Emory
MPS has high NPV

- Myocardial perfusion scintigraphy has a high negative predictive value
- Even if stenosis on angiography a normal MPS means low risk
- Therefore MPS can be used to determine who gets treatment
- If MPS used in all patients in EU savings would be 4 billion Euros/year-EMPIRE
Costs of investigating chest pain—EMPIRE

1=ecg, 2=ecg+MPS, 3=MPS+angio, 4=angio
Risk stratification

• Men more at risk than women at any age
• Smokers more at risk
• High cholesterol at risk
• Diabetics at high risk (may also be without symptoms)
• Poor haemodynamic response to stress
Risk stratification

• Advantages of MPS
  – Gives good information on function
  – Accurate
  – Reproducable
  – Understandable
  – Quantifiable (Bullseye)
Risk stratification-MPS

• The risk of cardiac event is linked to volume of myocardium at risk
• If more than 1 coronary artery territory involved, greater the risk
• If LVEF reduced the greater the risk
• LV dilatation increase risk
• Lung uptake increases risk
Risk assessment using TI-201

• Kaul et al JACC, 1988
• Identified main risk factors from TI-201 MPS
  – Highest risk Inc lung activity
  – Then > 1 coronary art ter involved
  – Then reduction in LVEF
  – Then size of defects
Using MPS in practice

Patient at risk of IHD

Low risk MPS
Medical Treatment

High risk MPS
Coronary angiography

Revascularisation
Showing improvement in ischaemic myocardium
Change in LVEF after CABG

Use of Gated LVEF before and after CABG

New work looking at gated SPECT un
same context
Where is nuclear cardiology best?

- Gate keeper for angiography
- Diabetes
- Women
Diabetes and silent ischaemia

• Araz et al Acta Diabet 20004
• Reviewed 116 type II diabetics
• 15% had ischaemic changes on MPS
• 10% patients no chest pain
• Recommended screening all type II diabetics with HBA1> 9%
Women

- In 1970 Common belief women did not have IHD
- Did not explain major cause of death in women in Europe
- Symptoms atypical (may be food related)
- Risk factors diabetes, hypertension, obesity
- Racial factors – Africans & South Asians worse if then more to Europe or North America
Women and heart disease

• Not able to do physical stress so easily
• Problems of breast attenuation in MPS
• Despite this clear evidence that a normal MPS in women= low risk of cardiac event and abnormal MPS = rate of cardiac event similar to men (Hachamovitch et al JACC 1996, Berman et al JACC 2003)
A woman with extensive ischaemia
Looking at attenuation

- Review the cine mode planar acquisition
- See if attenuation obvious from some angles
- Also should affect the same site at stress and rest (in women breast must be in same position)
- Differential of a fixed defect MI or attenuation
- A-C does appear to introduce new artefacts
- However gated study if moves it was attenuation if akinetic MI most likely
Gated SPECT MPS ?attenuation
Gated MPS SPECT ischaemia
Quantitative gated studies

• Programmes such as QGS
• Used to analyse LVEF, EDV and ESV
• Also can provide imaging of wall motion
• Reduced regional wall motion can be due to
  – Severe ischaemia
    • Hibernating
    • Stunning
  – Infarct
Problems of plotting 4D on 2D
Hibernation

• Defined as chronic severe ischaemia
• Hypoxic tissue
• Not able to burn lactate through Kreb cycle
• Able to survive on phosphorylation of glucose alone
• Cannot contract
• So (especially in delayed imaging) some perfusion but no movement seen on gated SPECT
Antero-septal hibernation
Assessment of cardiac function

• Use of gated studies
• Improve specificity of scans
• Look for wall motion abnormalities-Infarct, ischaemia, stunning and hibernation
• Look for dyskinesia – ventricular aneurysm
• Global function to look at prognosis
Cardiac MUGA

Ejection Fraction = 56% (Two View Manual)

<table>
<thead>
<tr>
<th>Emptying</th>
<th>Filling</th>
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</thead>
<tbody>
<tr>
<td>MaxRate (EDU/sec)</td>
<td>5.5092</td>
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<tr>
<td>ApgRate (EDU/sec)</td>
<td>2.7895</td>
</tr>
<tr>
<td>Time to peak</td>
<td>38</td>
</tr>
</tbody>
</table>

1/3 Filling Fraction: 0.6000

Volume Curve

Beat Histogram

Heart Rate: 97
Last Useful Frame: 31
MS/Frame: 19
Beat Window: 40%
Forward Framing
Total Beats: 482
Accepted: 432
Rejected: 0
R-R time (ms)
Max: 705
Min: 505
Mean: 614
MUGA vs ECHO

- MUGA
  - Accurate LVEF
  - Need special equipment and trained staff
  - Reproducable
  - Uses radiation

- ECHO
  - Needs special equipment and trained staff
  - No radiation
  - Can see valves as well
  - LVEF not so accurate
Prognostication with gated studies

- Reduction of LVEF below 50% is abnormal
- Reduction of LVEF below 40% means significant reduction in LV function
- Reduction of LVEF below 30% means symptoms
- Reduction of LVEF below 20% means symptoms at rest
- Reduction of LVEF below 10% means transplant or death
- If ventricular aneurysm all these are worse
Note large defect and spalying of walls to apex suggesting aneurysm
Both LVEF and LV size predicts survival

Death rate by LVEF

Why PET?

• Improved resolution - not really required in cardiology
• Improved sensitivity – this may be important-financially as reduced acquisition time
• Improved attenuation correction-good
• Look at metabolism-could be very good
F-18 FDG

• Most commonly available PET radiopharmaceutical
• Uptake dependent on glucose drive in cells
• Related to hypoxia
  – Glucose to lactate (no O₂) = 4ATP per molecule
  – Glucose to Krebs cycle =32 ATP per molecule
• So ischaemic tissue needs lots of glucose but remember in diabetics competitive uptake so need blood glucose <6mmol/l
Viability

• In patients with severe ischaemia identification of viable heart very important
• Revascularisation may result in improvement in flow and over time return of function
• Prognosis dependent on LVEF in these patients is viable myocardium not improved 32% 1 year survival vs 7% if improved (Dy Carli et al JTCS 1998)
• F-18 FDG can find that viability
Perfusion and viability

• Viability found with F-18 FDG (where uptake may be increased in severe ischaemia)

• How about perfusion
  – N-13 H$_3$
  – Rb-82

• However though not perfect can use single photon for perfusion
SPET and PET
FDG in action

• Wu et al Kyoto JNM 2007
• Looked at 41 patients with severe IHD
• Using TI-201/FDG PET mismatches identified (heart divided into 17 segments)
• 394 viable segments per heart were identified in 31 patients
• 29 had CABG, 76% of these had an improvement of >5% in LVEF
FDG in viability

- Slart et al Groningen JNC 2006
- 213 segments in 31 patients (17 segments per patient) were imaged with F-18 FDG before and just after CABG
- An increase in F-18 FDG uptake of more than 50% post surgery predicted improvement in LVEF in 93% of cases (specificity 85%)
Comparing NH$_3$ and F-18 FDG

- Slart et al Groningen EJNMMI 2006
- Used combination of NH3 and F-18 FDG PET to determine areas of viability (reduced NH$_3$ with normal or raised F-18 FDG) or normal NH$_3$ but raised F-18 FDG)
- 47 patients with severe IHD
- In 90% of these PET predicted an improved LVEF post surgery
What can we see

• What follows is a series of perfusion maps of perfusion and viability
• Available on line from Bringham and Women’s hospital in Boston
• Only illustrative examples
Normal

Flow

Metabolism
Infarct
Viability
Hibernation
Cardiomyopathy
In reality what it looks like
Improving the system

• Can we use PET to provide a better service
• Rb-82 may allow for faster more accurat imaging
• Possibly all can be performed in a (one stop shop)
• Especially with 64 or 128 slice CT
Rb-82 for perfusion

- Bateman et al Kansas JNC 2006
- 112 patients had Tc-99m MIBI and Rb-82 perfusion studies
- 4 blinded readers compared results with CABG (stenosis of 70%)
- Rb-82 had accuracy of 89% compared with 79% for SPECT (p=0.003)
- Rb-82 better in men and the obese
SPECT vs Rb-82 PET

J Machac NYU
AC vs PET
Proposed scheme for viability

AC CT

Rb-82 injection and scan
adenosine stress

AC-CT

F-18 FDG injection

AC-CT

CT angiogram

F-18 FDG injection

FDG scan

0    15    30    45    60    90    120    135    150 mins

18mSv without CT angiogram
42mSv with CT angiogram
Dedicated cardiac SPECT-CT camera
New technology
2 minute acquisition
400MBq Tc-99m MIBI
SPECT-CT
Cardiac PET-CT