

# A short guide to DXA and BMD

Terry Watson  
Freeman Hospital

How does it work?



Patient pathways



What's new?



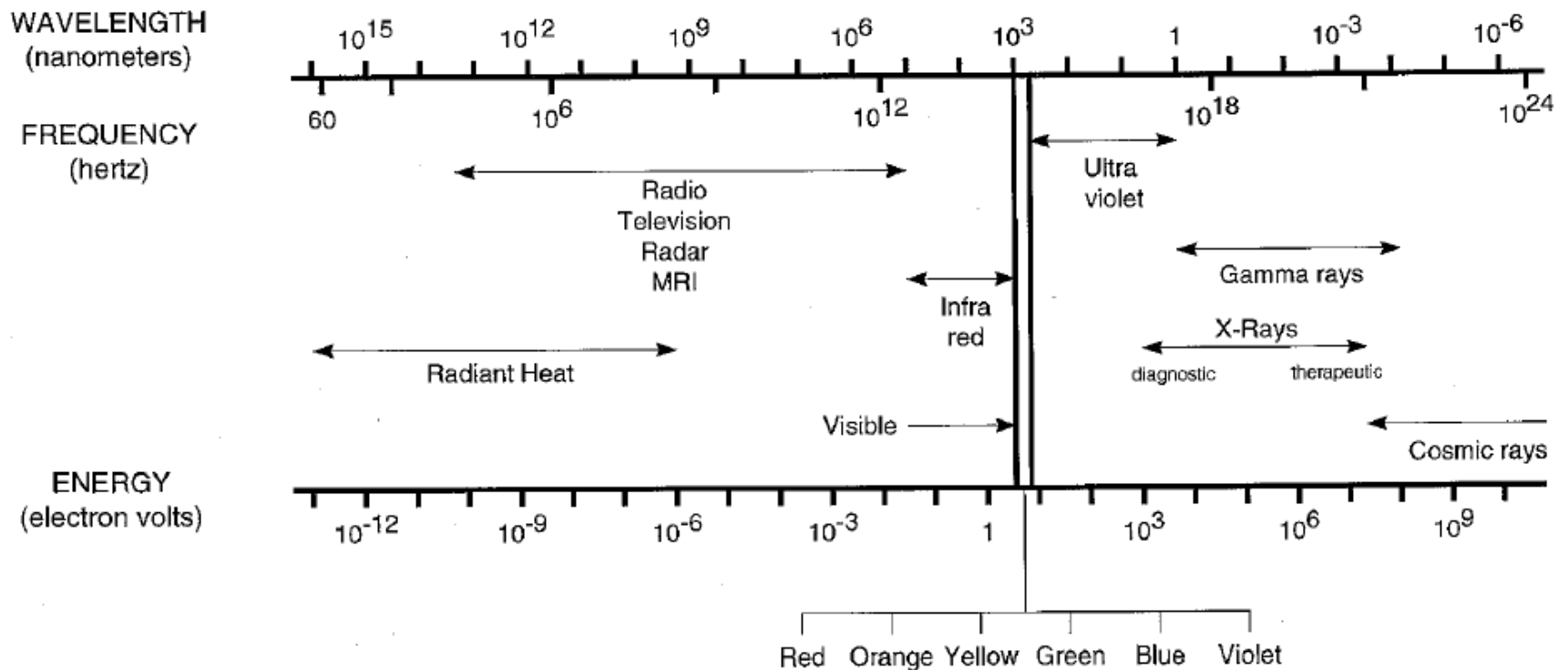
# First Principles

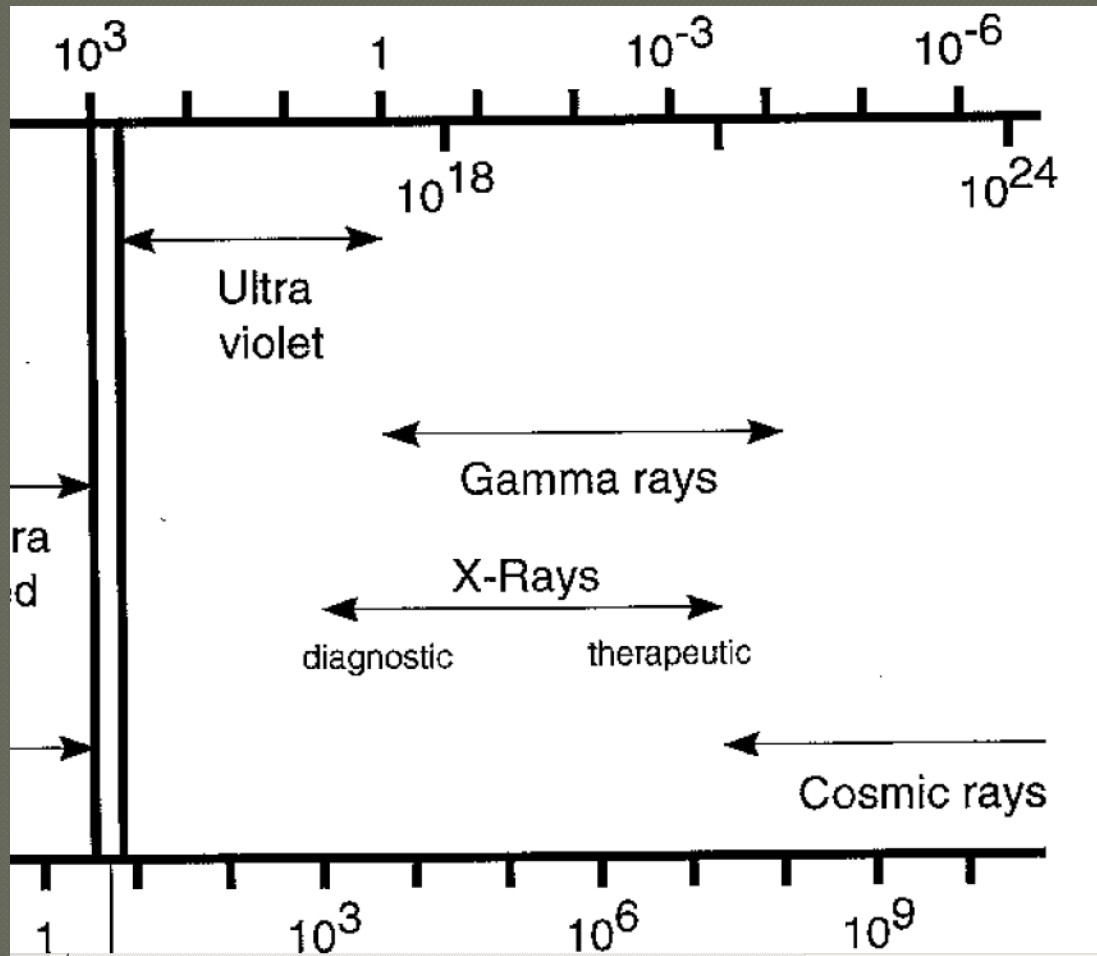
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- ◉ Why do we measure Bone Density?
- ◉ How do we measure Bone Density?
  - Number of *in vivo* methods developed
    - SPA
    - QCT
    - Ultrasound
    - DEXA(DXA)
  - Most important are the methods of measuring the 'Attenuation' of a beam of electromagnetic radiation or ultrasound.



# Electromagnetic Spectrum





# For routine diagnostic x-rays...

Hot filament:  
regulates electron  
production and  
therefore beam  
current and x-ray  
output



Rotating anode: spreads  
heat over larger area so  
increasing output

**We need** : a controlled source of high energy electrons, a suitable hard wearing small x-ray target, a means of cooling and a way to direct and control the x-rays passing through the patient





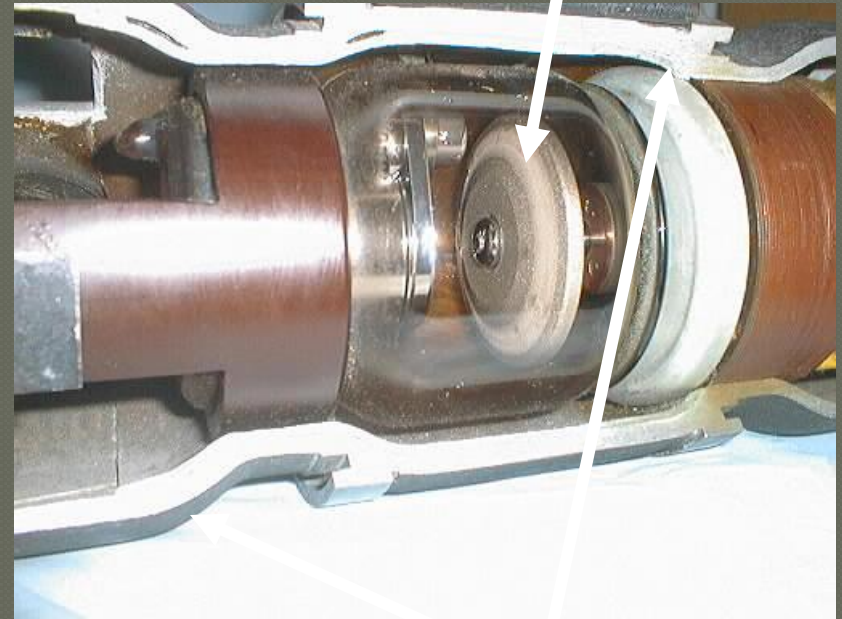
# Cut away x-ray unit

Anode disc showing damage pits



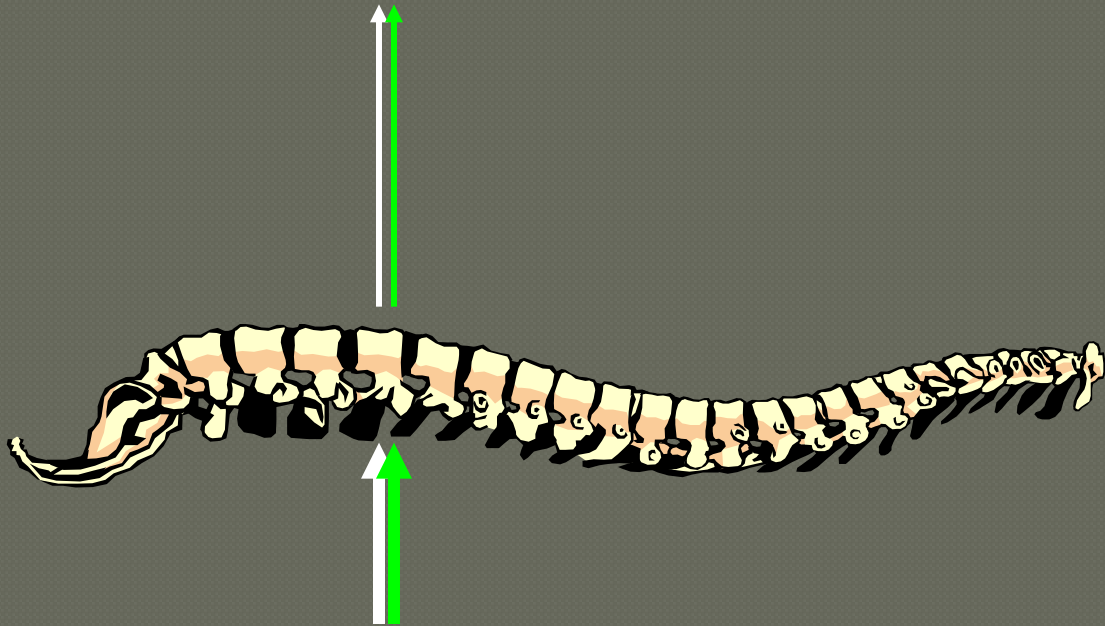
X-ray window

High voltage plugs



Radiation Shielding

DXA works by sending a narrow beam of x rays through bone. The bone density determines the amount that is absorbed by bone.



Two different energies x-ray energies are used to avoid influence from soft tissue

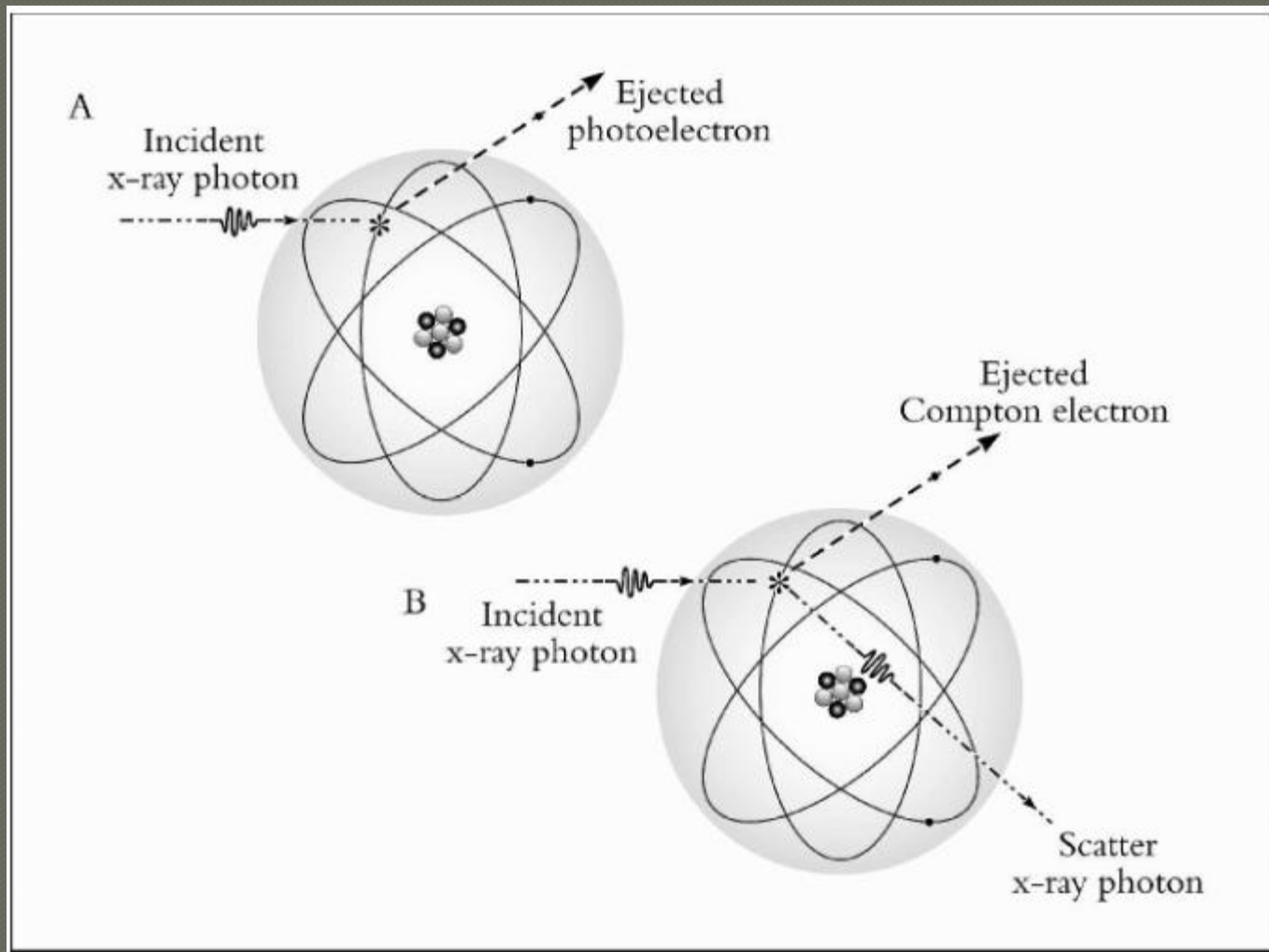
# How do we create X-rays of two distinct energies?

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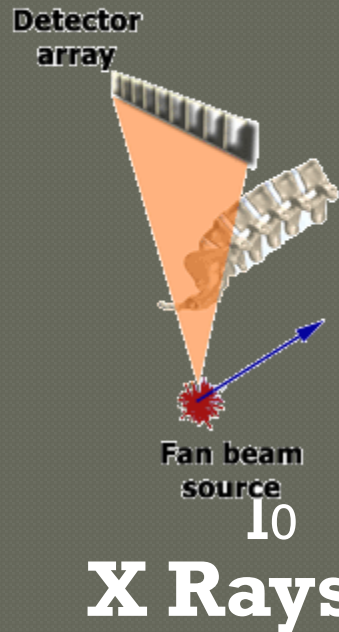
- ⦿ 'Hologic' machines – Fast switching in voltages (60 times per second)
- ⦿ 'Lunar' machines – Constant single energy X-ray transmission that is filtered by thin sheet of metal near to the tube.  
(cerenium or samarium)



# Attenuation in tissue

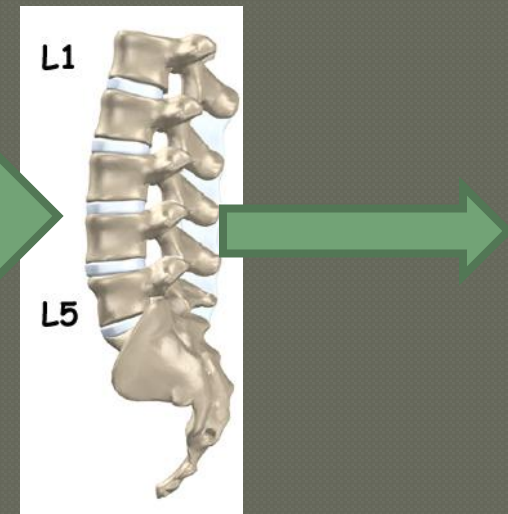


DXA works by measuring a narrow beam of x rays transmitted through bone



$I = < I_0$

**X Rays out**



Narrow x-ray beams obey a well defined exponential law of absorption

# Linear Attenuation coefficients

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- Numerical value which can be applied to describe a tissues ability to stop(attenuate) a radiation beam
- Directly proportional to the physical density of the tissue
- For DEXA the simplest model is to identify two distinct tissue types:
  - Soft Tissue
  - Bone

# For narrow beam x-rays passing through a person ...

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- Absorption depends upon the bone mineral density (BMD) ( $\text{g}/\text{cm}^2$ ) which is what we want to know
- It also depends upon soft tissue thickness and fat content which varies within and between patients.
- DXA uses two x-ray energies at each point
- Changes in absorption with energy are much greater for bone than soft tissue
- The difference in the two signals at each point when compared to soft tissue baseline gives BMD

So why is DXA useful then?

x-rays can suggest  
deficiency in bone  
mineral..

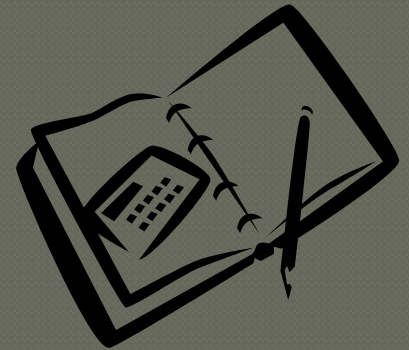


...but DXA provides  
both an image and a  
numerical assessment  
of bone density

# Specifications DXA

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- Precise measurement of Bone mineral ( $\text{g}/\text{cm}^2$ )
- Reproducible
- Scan time 30secs per region
- Patient appointment time 20 mins
- Low radiation dose

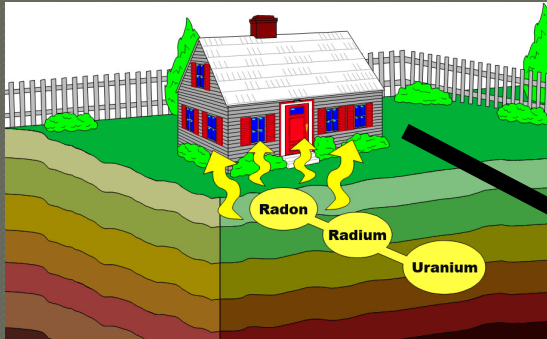


# Typical radiation dose

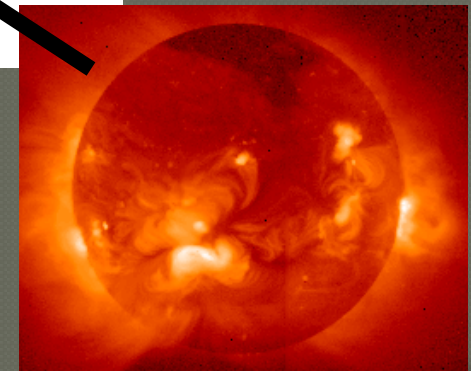
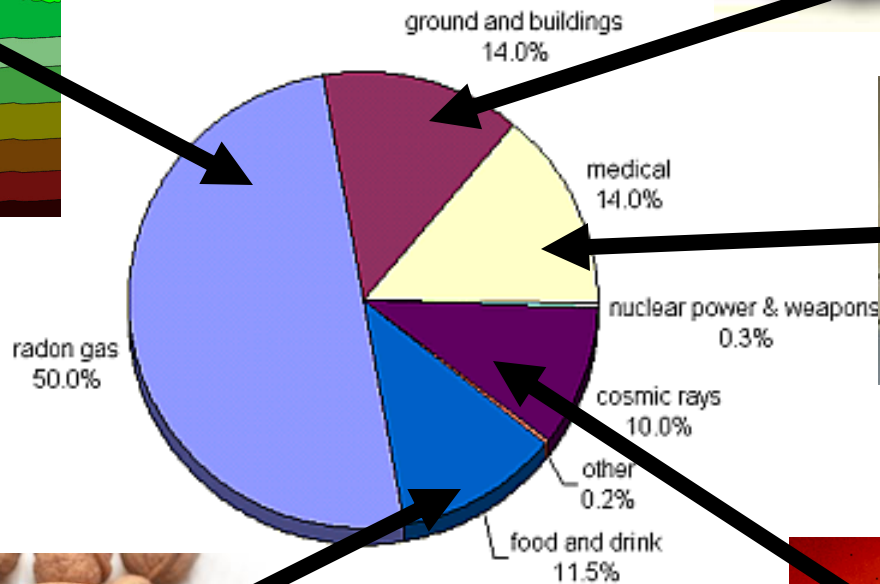
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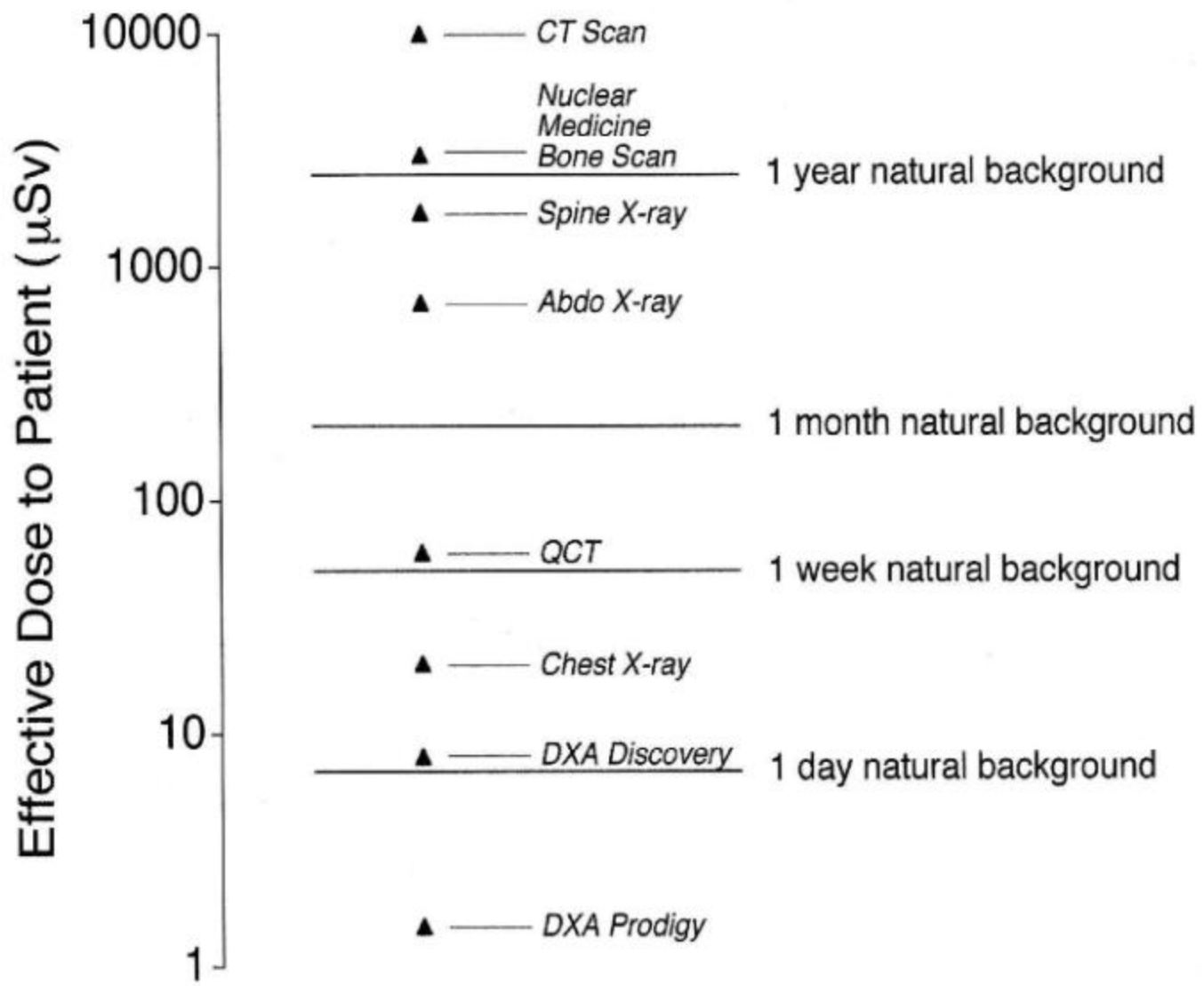
- The radiation dose of a Dexa scan to the patient is typically  $1\mu\text{Sievert}$ 
  - Includes both spine and hip images
  
- Lets put this into context

# Mean UK Background dose 2.7mSv per year



### Background Radiation in the UK





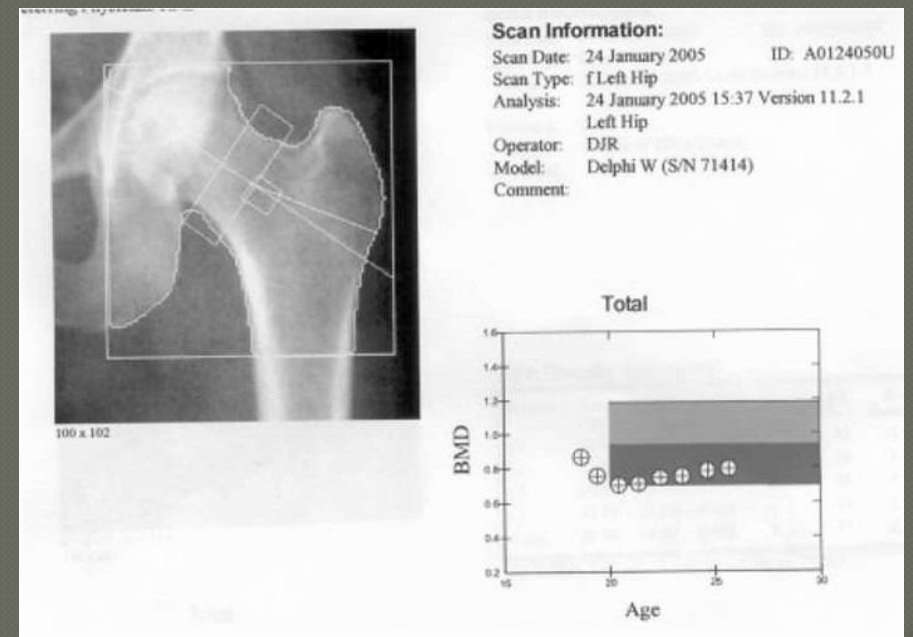
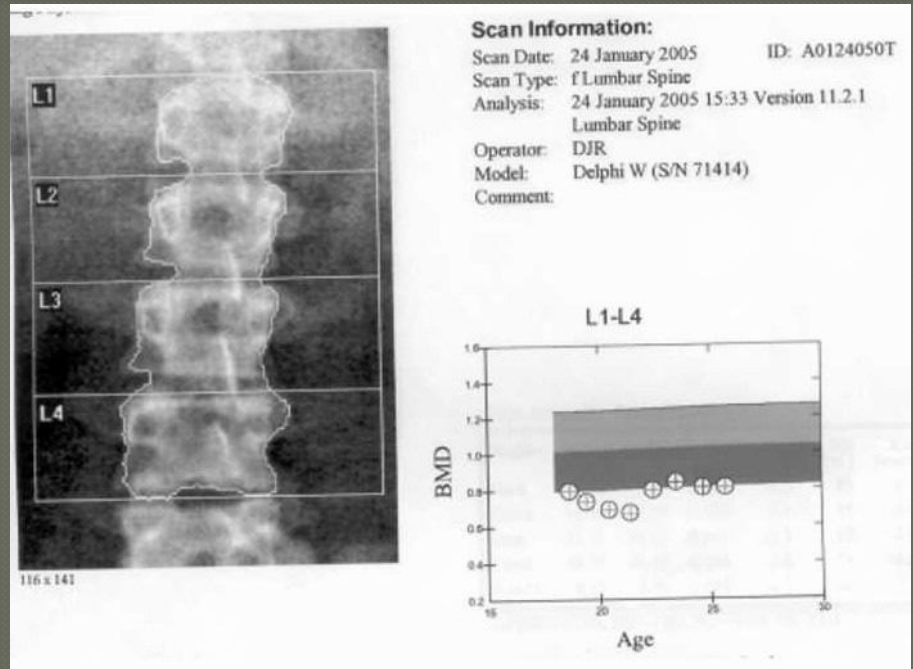
# Risk of death comparable to having a DEXA

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- Exposure to natural background radiation for 4 hours
- Smoking one-tenth of a cigarette
- Travelling 3 miles by car
- Travelling 15 miles in an airliner
- Rock climbing for 5 seconds
- Canoeing for 20 seconds
- Working in a factory for half a day
- Being a woman aged 30 for 60 minutes
- Being a woman aged 40 for 20 minutes
- Being a woman aged 50 for 8 minutes
- Being a woman aged 60 for 3 minutes
- Being a woman aged 70 for 1 minute

Why do we acquire an image if  
we are interested in the  
numbers?

The function of the image is to ensure the appropriate region of interest is measured every time...



# Good precision depends critically on ROI placement ...

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In bone densitometry, the gold standard is the ash weight of the skeletal site being tested. BMD testing is very precise and accurate compared to other tests commonly used in clinical practice, but precision and accuracy will vary according to the instrument used, the skeletal site being measured, *and the skill of the technologist.*

**So image quality is an important factor ensuring  
precision**

# The report shows.....

- 1) A skeletal image (lumbar spine or proximal hip)
- 2) Bone mineral densities ( $\text{g}/\text{cm}^2$ )
- 3) The patient result plotted against a normal population

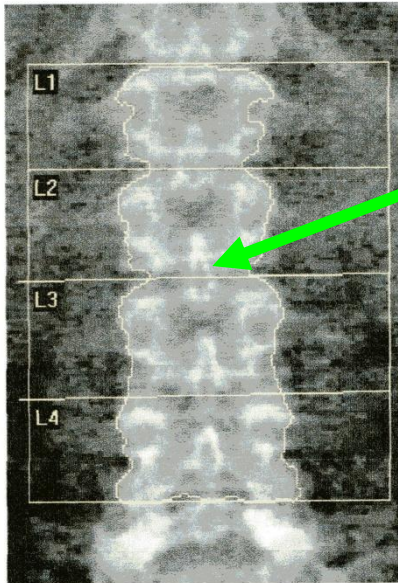
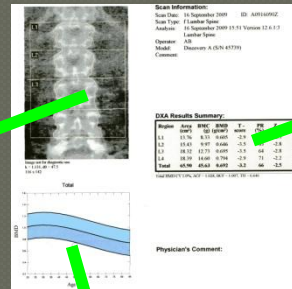


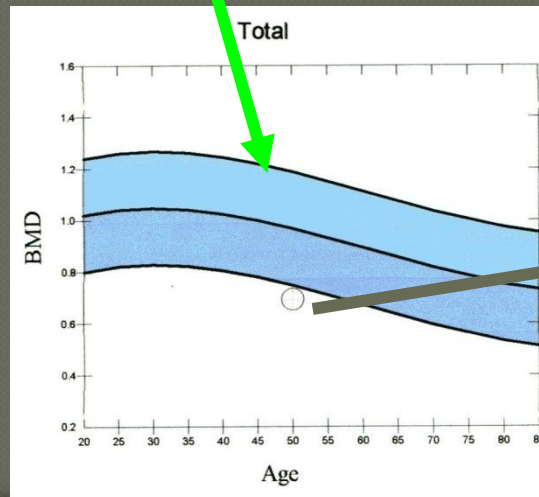
Image not for diagnostic use  
 $k = 1.131, d_0 = 47.5$   
 116 x 142



## DXA Results Summary:

Region	Area ( $\text{cm}^2$ )	BMC (g)	BMD ( $\text{g}/\text{cm}^2$ )	T-score	PR (%)	Z-score	AM (%)
L1	13.76	8.33	0.605	-2.9	65	-2.3	71
L2	15.43	9.97	0.646	-3.5	63	-2.8	68
L3	18.32	12.73	0.695	-3.5	64	-2.8	69
L4	18.39	14.60	0.794	-2.9	71	-2.2	77
<b>Total</b>	<b>65.90</b>	<b>45.63</b>	<b>0.692</b>	<b>-3.2</b>	<b>66</b>	<b>-2.5</b>	<b>72</b>

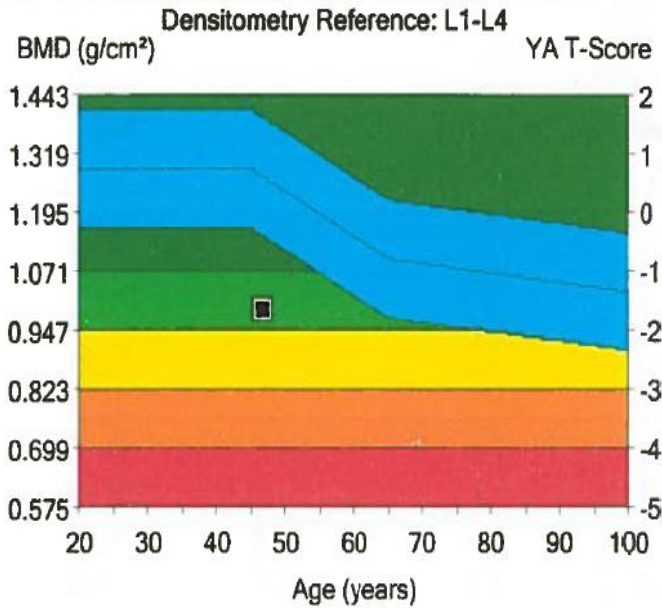
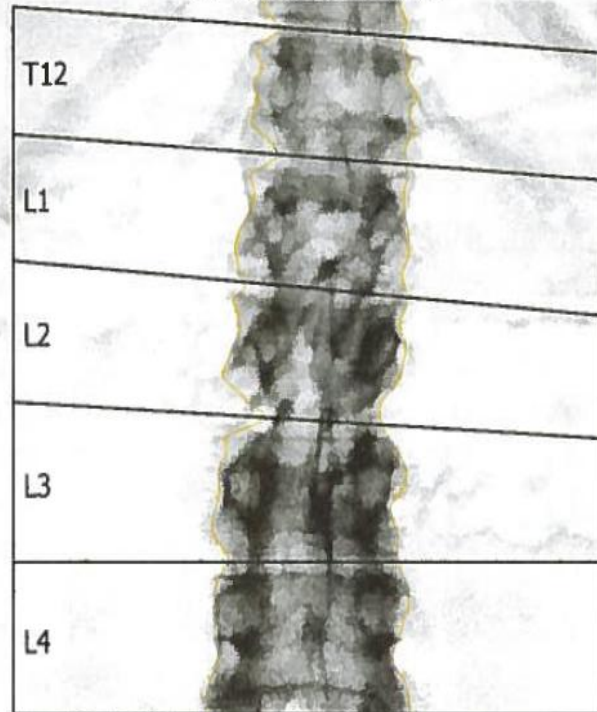
Total BMD CV 1.0%, ACF = 1.028, BCF = 1.007, TH = 6.646



Z = -2.5

T = -3.2

AP Spine Bone Density



Region	1		2		3	
	BMD (g/cm <sup>2</sup> )	Young-Adult (%)	T-Score	Age-Matched (%)	Z-Score	
L1	0.815	72	-2.7	67	-3.4	
L2	1.036	86	-1.4	80	-2.2	
L3	1.102	91	-0.9	85	-1.7	
L4	0.982	82	-1.8	76	-2.6	
L1-L4	0.989	83	-1.7	77	-2.4	

# DEXA can be used to diagnose osteoporosis

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- Osteoporosis is diagnosed in adults where  $T = -2.5$  or less at the lumbar spine or hip (WHO criteria 1994)
- This may not necessarily represent a treatment threshold as a full clinical assessment is indicated prior to treatment

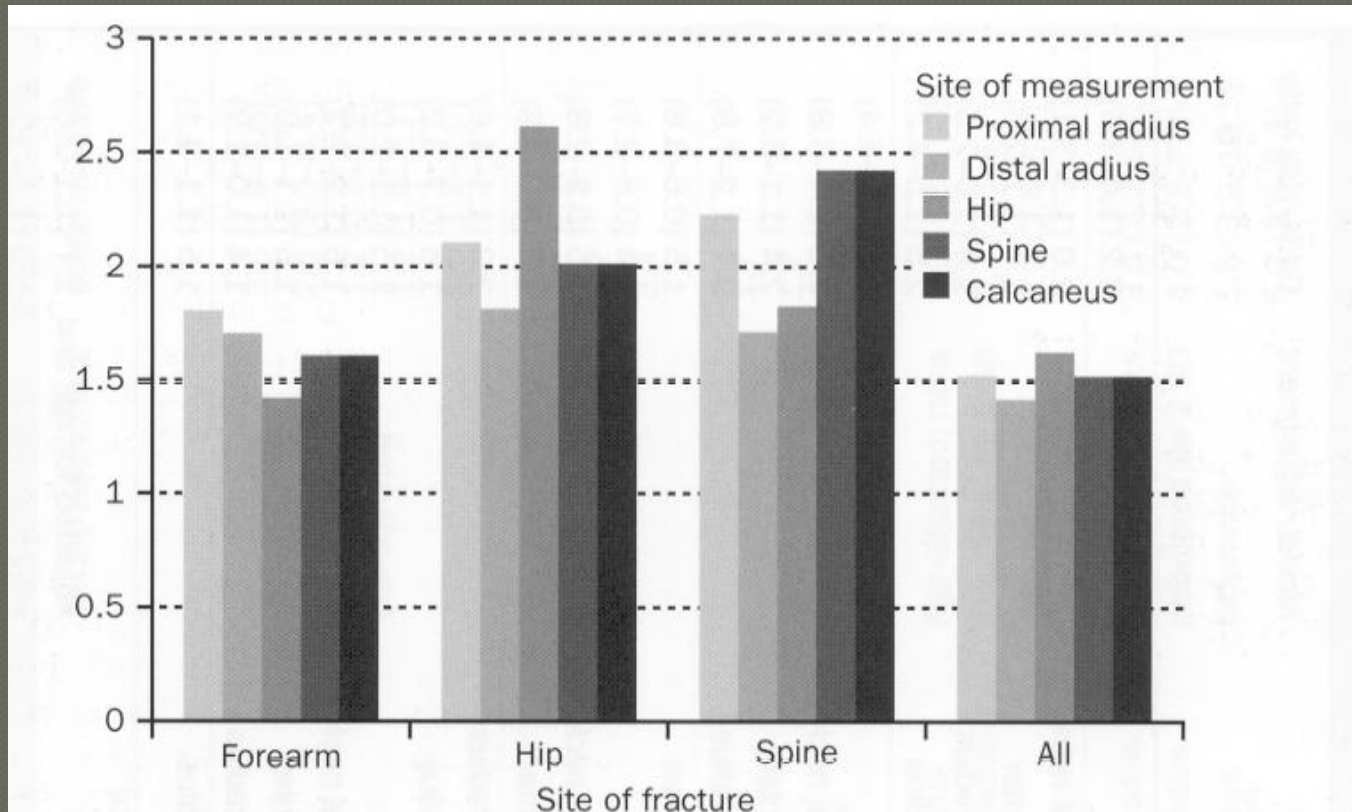
# DEXA can be used to diagnose osteoporosis

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- ⦿ Normal bone: T-score better than -1
- ⦿ Osteopenia: T-score between -1 and -2.5
- ⦿ Osteoporosis: T-score less than -2.5
- ⦿ Established (severe) osteoporosis *includes the presence of a non-traumatic fracture.*

# Risk of future fracture increases by factor of between 1.4 and 2.6 for every 1SD decrease in BMD

Marshall et al 1996



# How do we use DXA?

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- Bone Density (DXA) is a sensitive indicator of future fracture but is not very specific
- It cannot therefore be used for population screening
- It can however be used along with clinical risk factors as part of a case finding strategy

# The indications for DEXA

based upon NOS 'Local Provision for Osteoporosis' and AGO report

- Early Menopause
- Prolonged Amenorrhoea
- HRT Critical
- Vertebral Deformity
- Low Trauma Fractures
- Osteopenia on X-ray
- Long term/high dose steroids
- Eating disorders
- Chronic Liver disease
- Alcohol abuse
- Kidney dialysis
- Hyperparathyroidism
- PBC
- Hypogonadism
- Malabsorption Syndrome
- Transplant Assessment
- Growth Hormone
- JCA
- Thyroid Dysfunction
- Follow up/previous abnormal DEXA
- Other indication / trial patient

# Point of care clinical risk assessment

<http://www.shef.ac.uk/FRAX/>

Please answer the questions below to calculate the ten year probability of fracture with BMD.



## Weight Conversion:

pound:

## Height Conversion:

inch:

Country : **UK**      Name / ID :       [About the risk factors](#)

**Questionnaire:**

1. Age (between 40-90 years) or Date of birth  
Age:       Date of birth:  
 Y:  M:  D:

2. Sex       Male     Female

3. Weight (kg)     

4. Height (cm)     

5. Previous fracture       No     Yes

6. Parent fractured hip       No     Yes

7. Current smoking       No     Yes

8. Glucocorticoids       No     Yes

9. Rheumatoid arthritis       No     Yes

10. Secondary osteoporosis       No     Yes

11. Alcohol 3 or more units per day       No     Yes

12. Femoral neck BMD (g/cm<sup>2</sup>)  
Select DXA

Now available as an i-phone app !

## Calculation Tool

Please answer the questions below to calculate the ten year probability of fracture with BMD.



Country: **UK**

Name/ID:

[About the risk factors](#)

### Questionnaire:

1. Age (between 40 and 90 years) or Date of Birth

Age:

Date of Birth:

Y:

M:

D:

2. Sex

Male

Female

3. Weight (kg)

4. Height (cm)

5. Previous Fracture

No

Yes

6. Parent Fractured Hip

No

Yes

7. Current Smoking

No

Yes

8. Glucocorticoids

No

Yes

9. Rheumatoid arthritis

No

Yes

10. Secondary osteoporosis

No

Yes

11. Alcohol 3 or more units/day

No

Yes

12. Femoral neck BMD (g/cm<sup>2</sup>)

Select BMD

**BMI: 24.5**

The ten year probability of fracture (%)

without BMD

Major osteoporotic

**6.0**

Hip Fracture

**0.8**

[View NOGG Guidance](#)

### Weight Conversion

Pounds kg

### Height Conversion

Inches cm

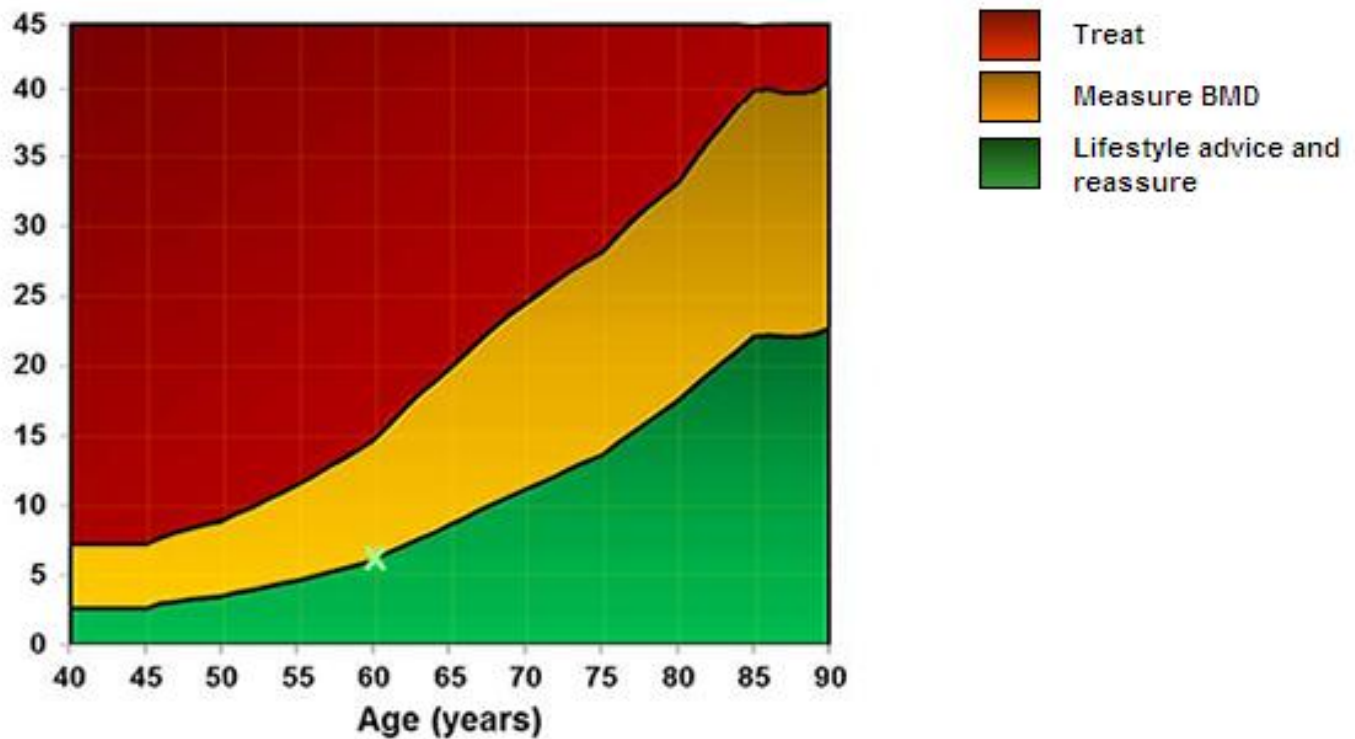
**01208987**

Individuals with fracture risk assessed since 1st June 2011



## Assessment threshold - Major fracture

10 year probability of major osteoporotic fracture  
(%)



## Calculation Tool

Please answer the questions below to calculate the ten year probability of fracture with BMD.

Country: UK

Name/ID: female aged 60

[About the risk factors](#)

### Questionnaire:

1. Age (between 40 and 90 years) or Date of Birth

Age:

60

Date of Birth:

Y:

M:

D:

2. Sex

 Male Female

3. Weight (kg)

70

4. Height (cm)

169

5. Previous Fracture

 No Yes

6. Parent Fractured Hip

 No Yes

7. Current Smoking

 No Yes

8. Glucocorticoids

 No Yes

9. Rheumatoid arthritis

 No Yes

10. Secondary osteoporosis

 No Yes

11. Alcohol 3 or more units/day

 No Yes12. Femoral neck BMD (g/cm<sup>2</sup>)

Select BMD

Clear

Calculate

**BMI: 24.5**

The ten year probability of fracture (%)

without BMD

Major osteoporotic

14

Hip Fracture

2.2

[View NOGG Guidance](#)

### Weight Conversion

Pounds → kg

Convert

### Height Conversion

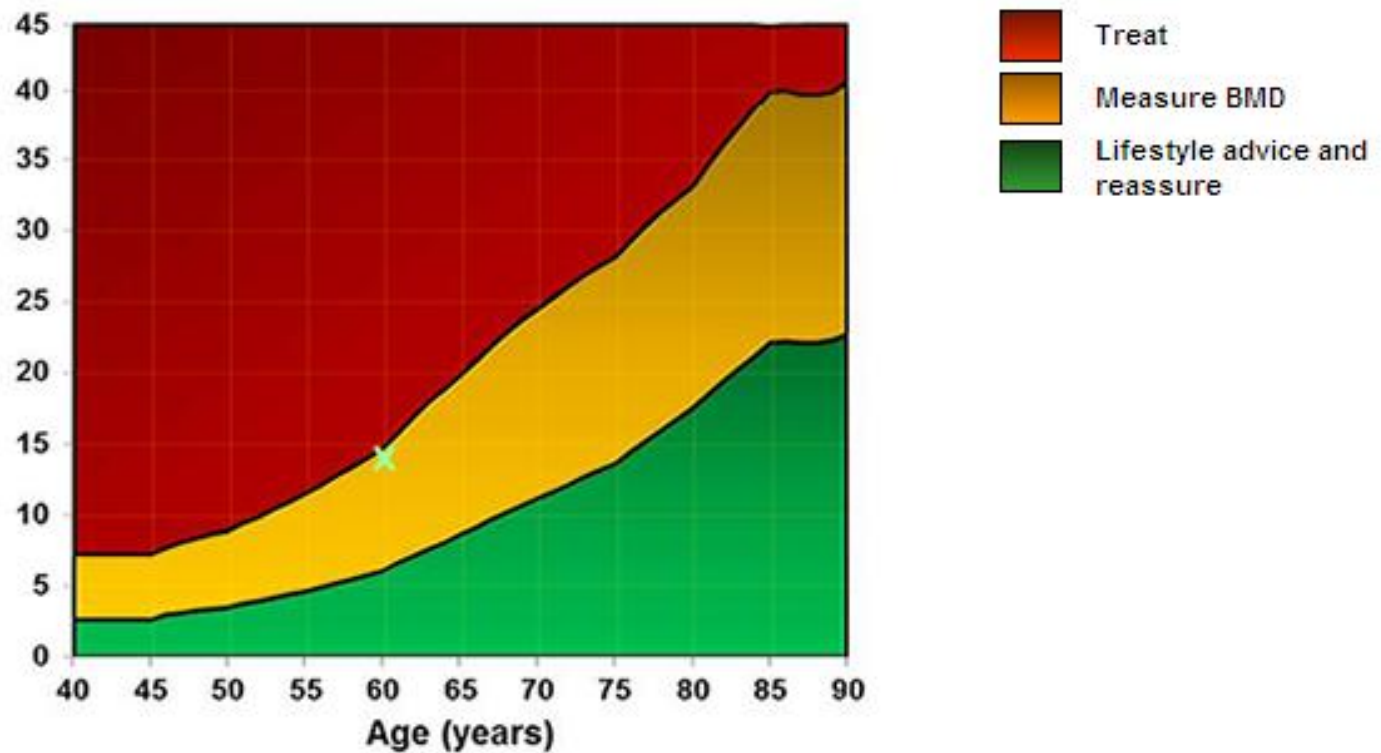
Inches → cm

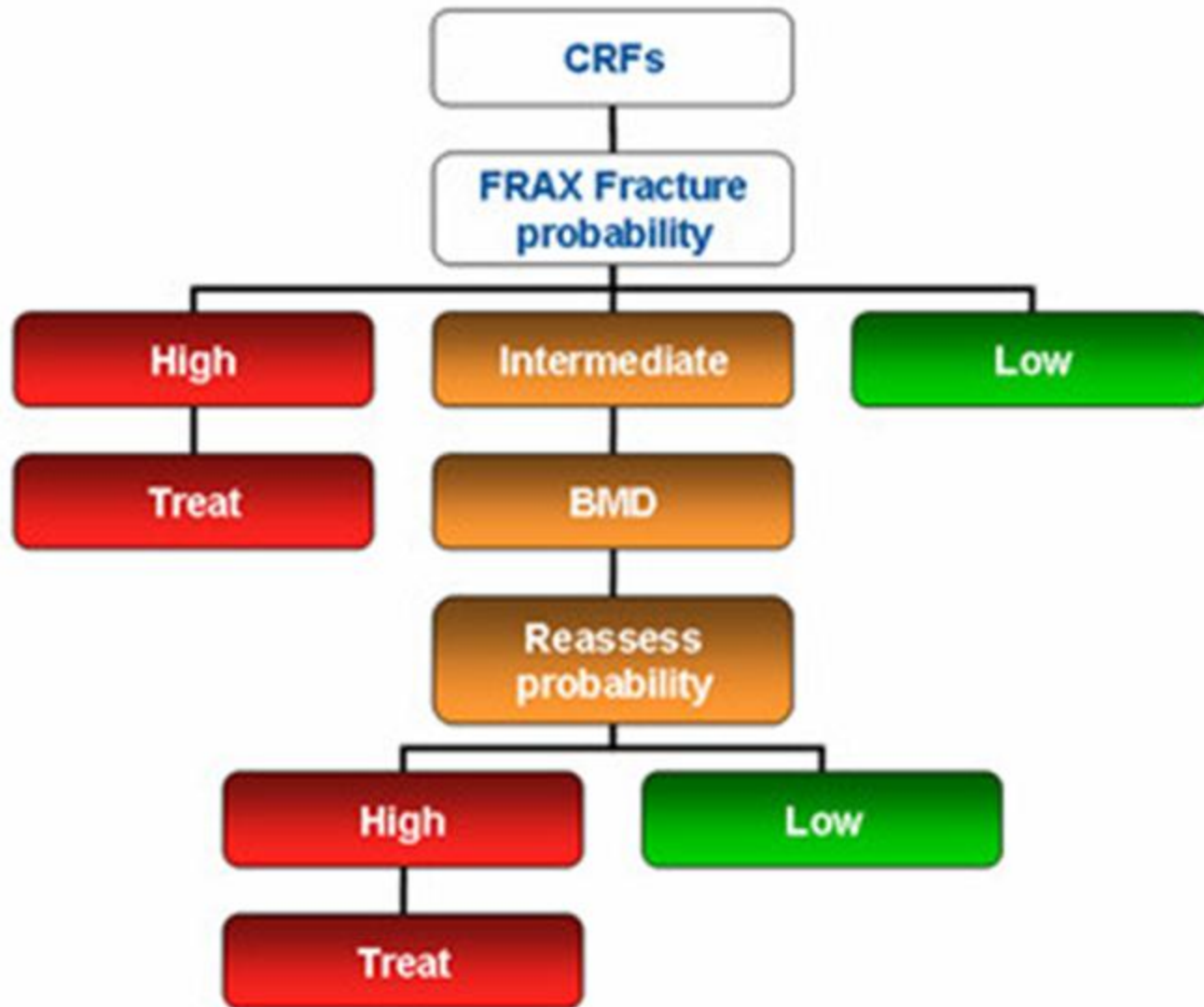
Convert

**01208987**Individuals with fracture risk  
assessed since 1st June 2011

## Assessment threshold - Major fracture

10 year probability of major osteoporotic fracture (%)





## Calculation Tool

Please answer the questions below to calculate the ten year probability of fracture with BMD.



Country: UK

Name/ID: female aged 60

[About the risk factors](#)

### Questionnaire:

1. Age (between 40 and 90 years) or Date of Birth

Age:

60

Date of Birth:

Y:

M:

D:

2. Sex

 Male Female

3. Weight (kg)

70

4. Height (cm)

169

5. Previous Fracture

 No Yes

6. Parent Fractured Hip

 No Yes

7. Current Smoking

 No Yes

8. Glucocorticoids

 No Yes

9. Rheumatoid arthritis

 No Yes

10. Secondary osteoporosis

 No Yes

11. Alcohol 3 or more units/day

 No Yes12. Femoral neck BMD (g/cm<sup>2</sup>)

GE-Lunar

0.715

T-score: -2.3

Clear

Calculate

**BMI: 24.5**

The ten year probability of fracture (%)  
with BMD

Major osteoporotic

20

Hip Fracture

5.2

[View NOGG Guidance](#)

### Weight Conversion

Pounds kg

Convert

### Height Conversion

Inches cm

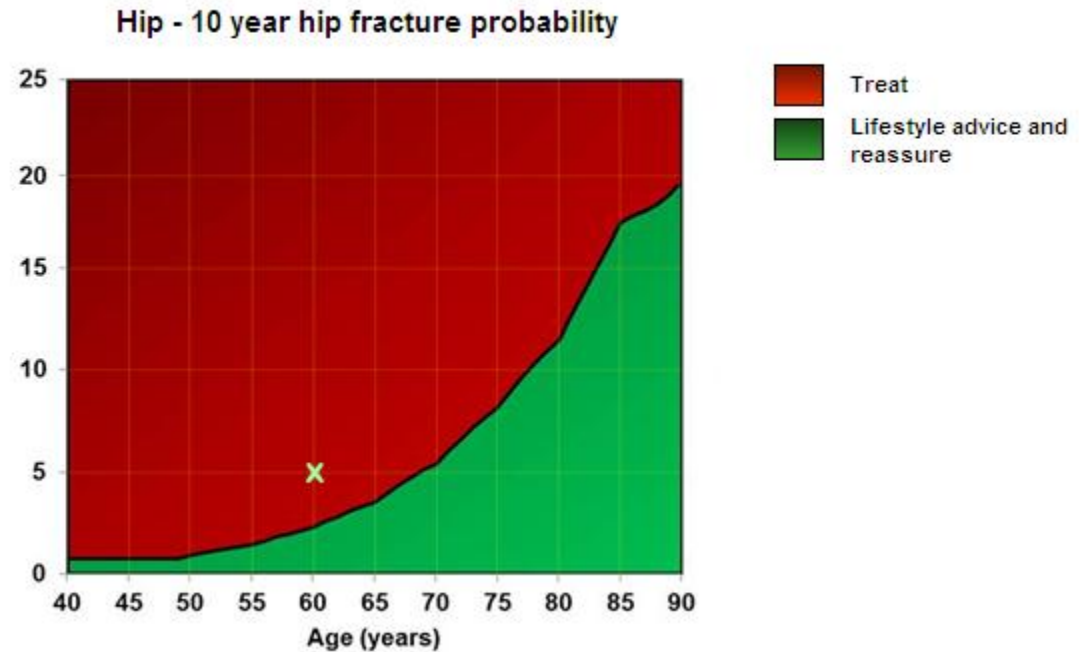
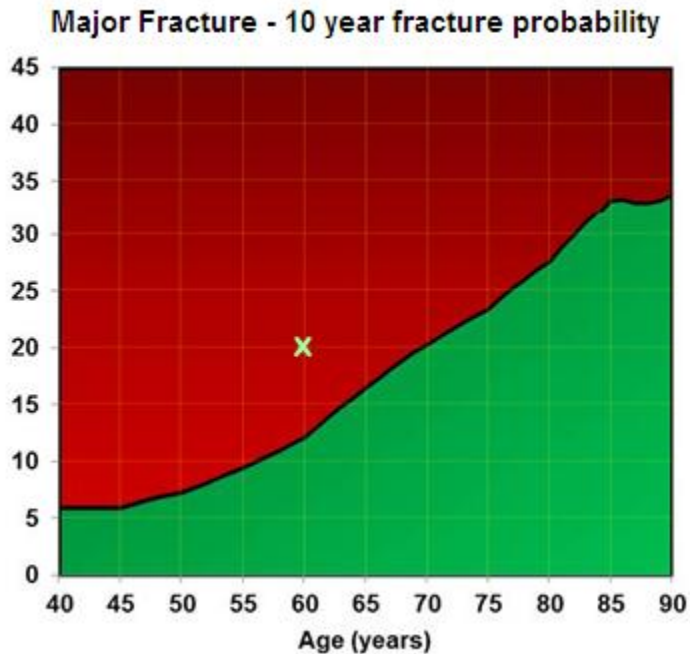
Convert

**01208987**

Individuals with fracture risk  
assessed since 1st June 2011



## Intervention Threshold



The low cost of generic alendronate, which has a broad spectrum of anti-fracture efficacy, makes this the first line treatment in the majority of cases.

In women who are intolerant of alendronate or in whom it is contraindicated, other bisphosphonates, denosumab, strontium ranelate or raloxifene may provide appropriate and cost-effective treatment options

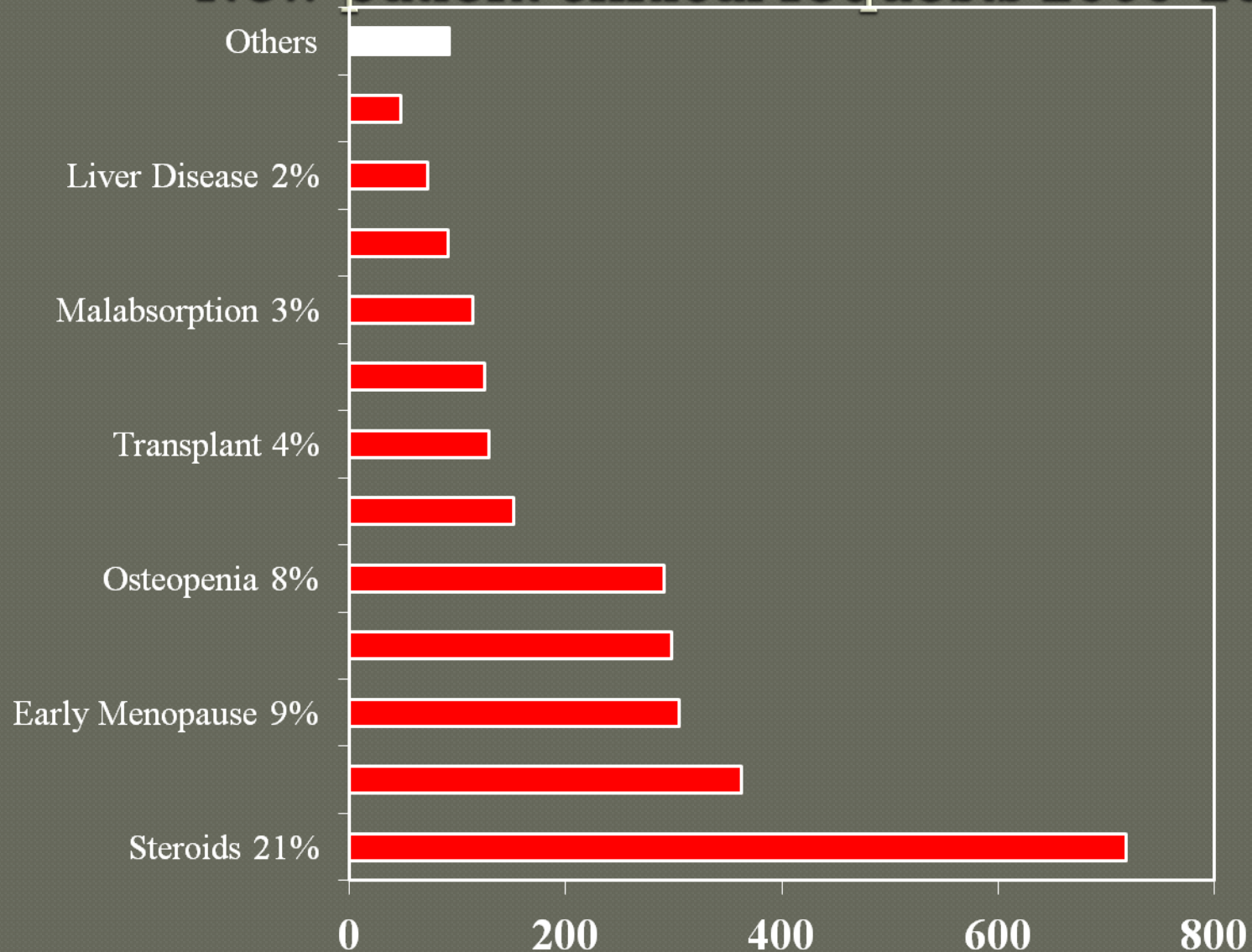
# Dexa requests at FH

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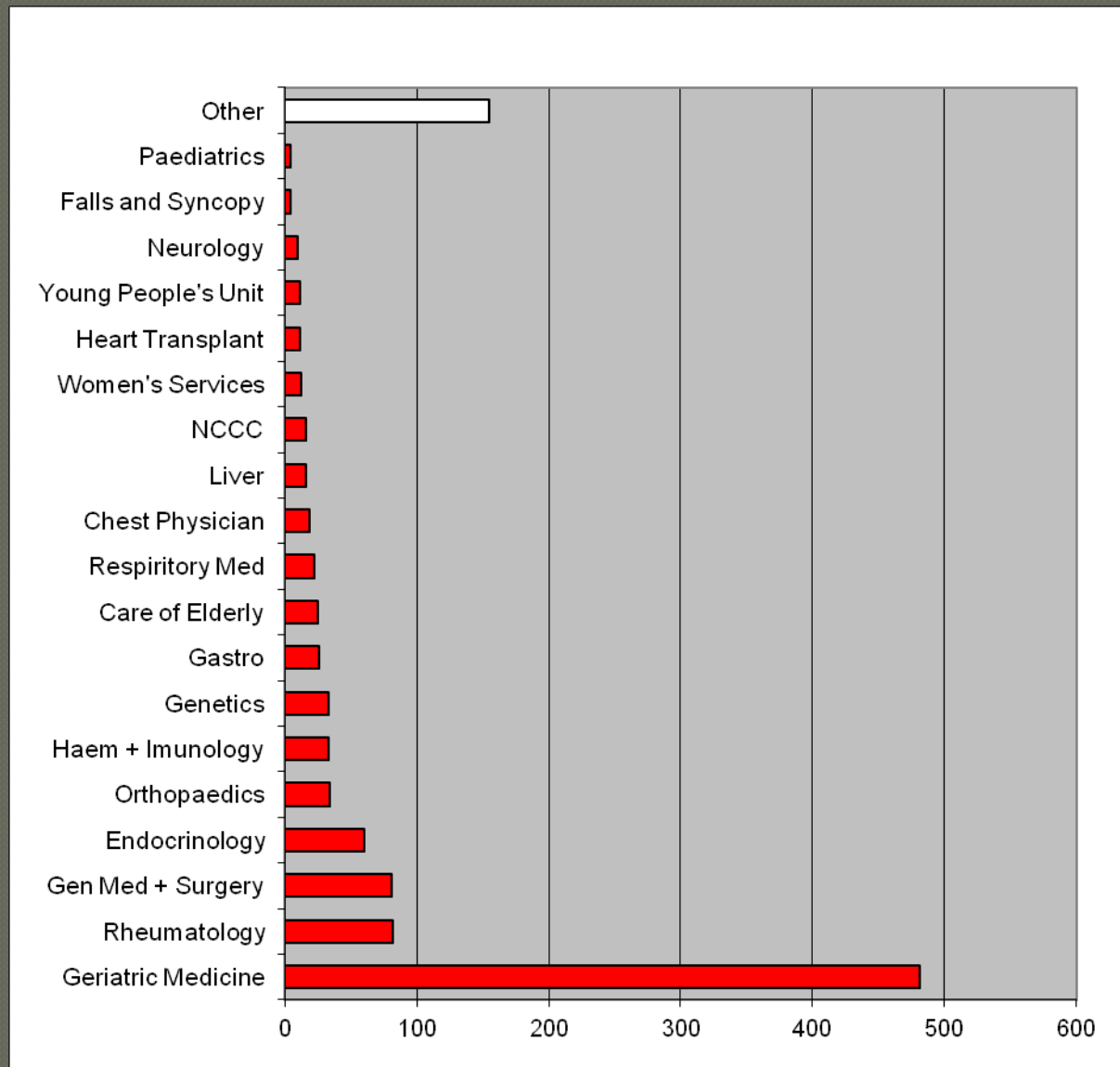
How many new patients?

How many review patients?

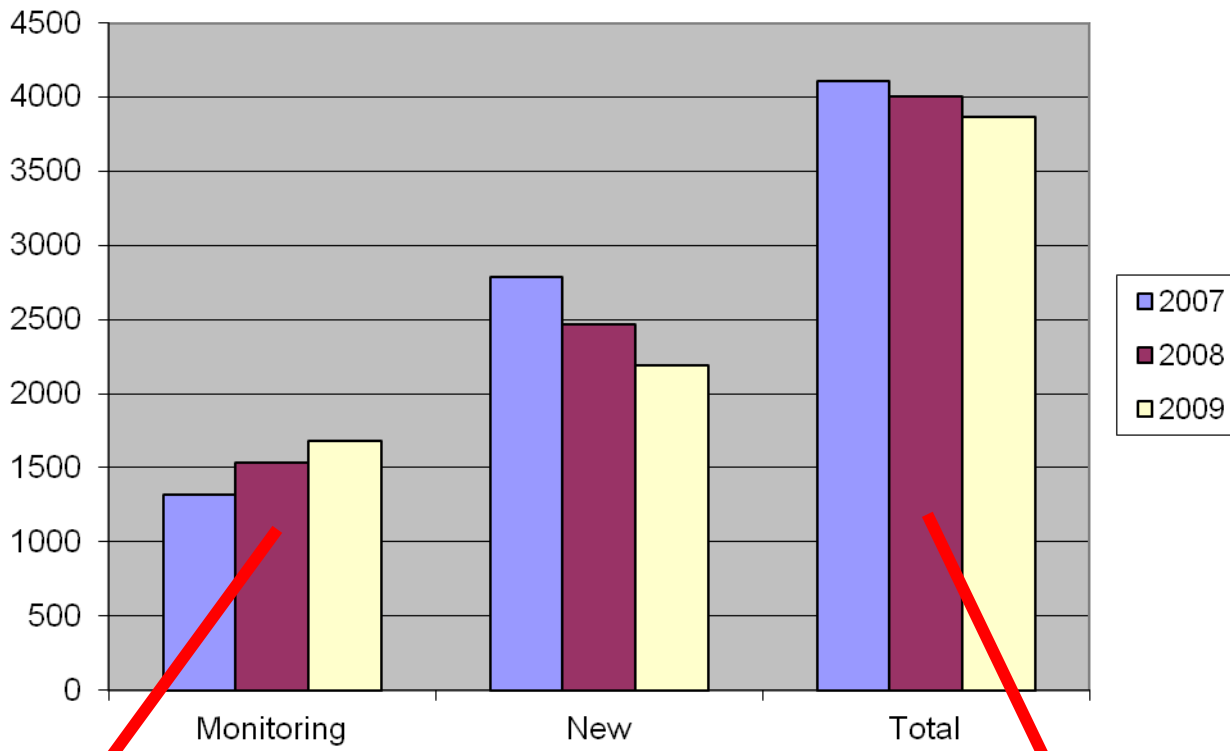
# New patient clinical requests 2000-2001



# Clinical requests by NUTH speciality snapshot July 2008



### Patient Numbers for DXA Freeman Hospital



Min 2 years or about 5% change in BMD

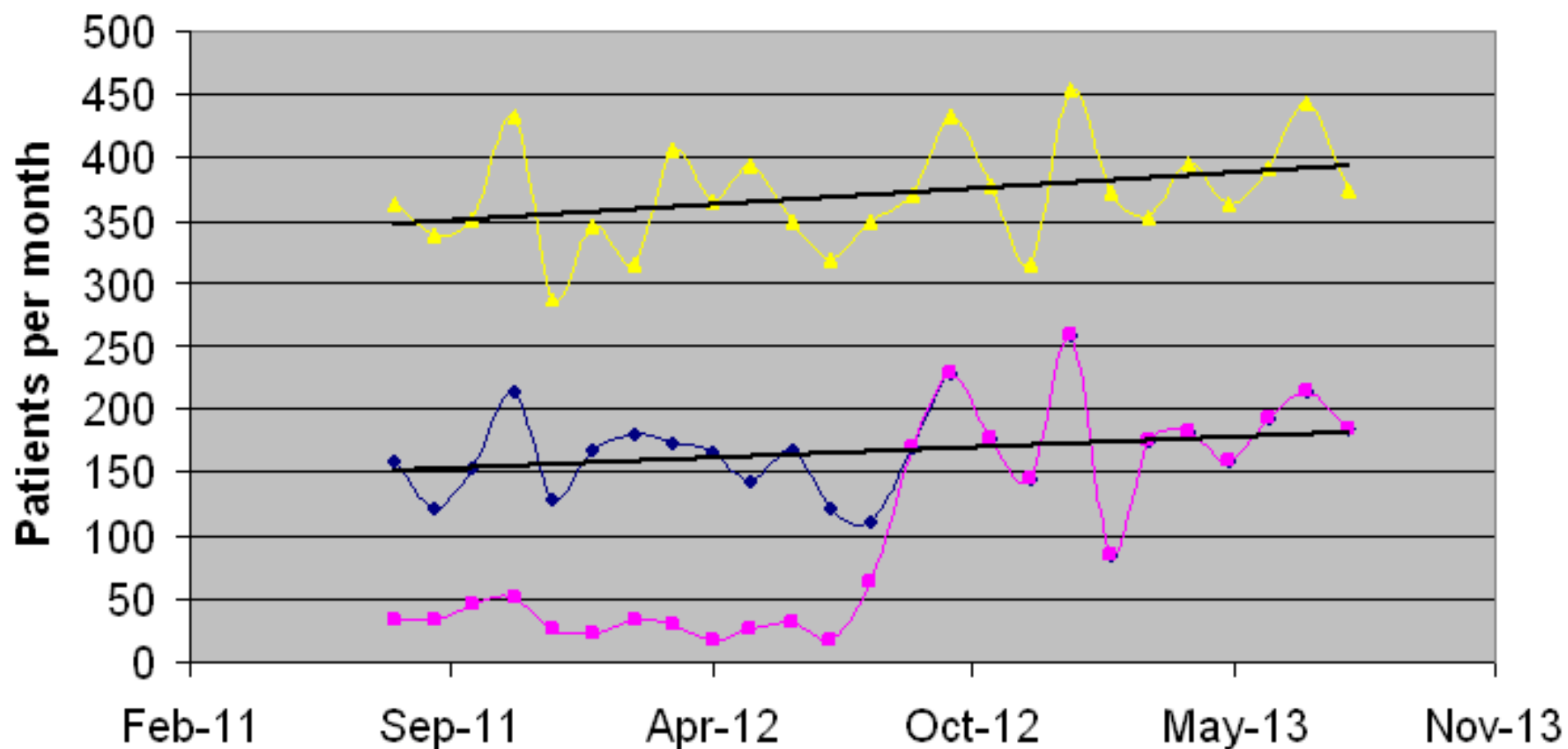
Broadly consistent with NOS guidelines

# DXA Freeman Patient numbers Aug 2011 - Aug-13 (25 Months)

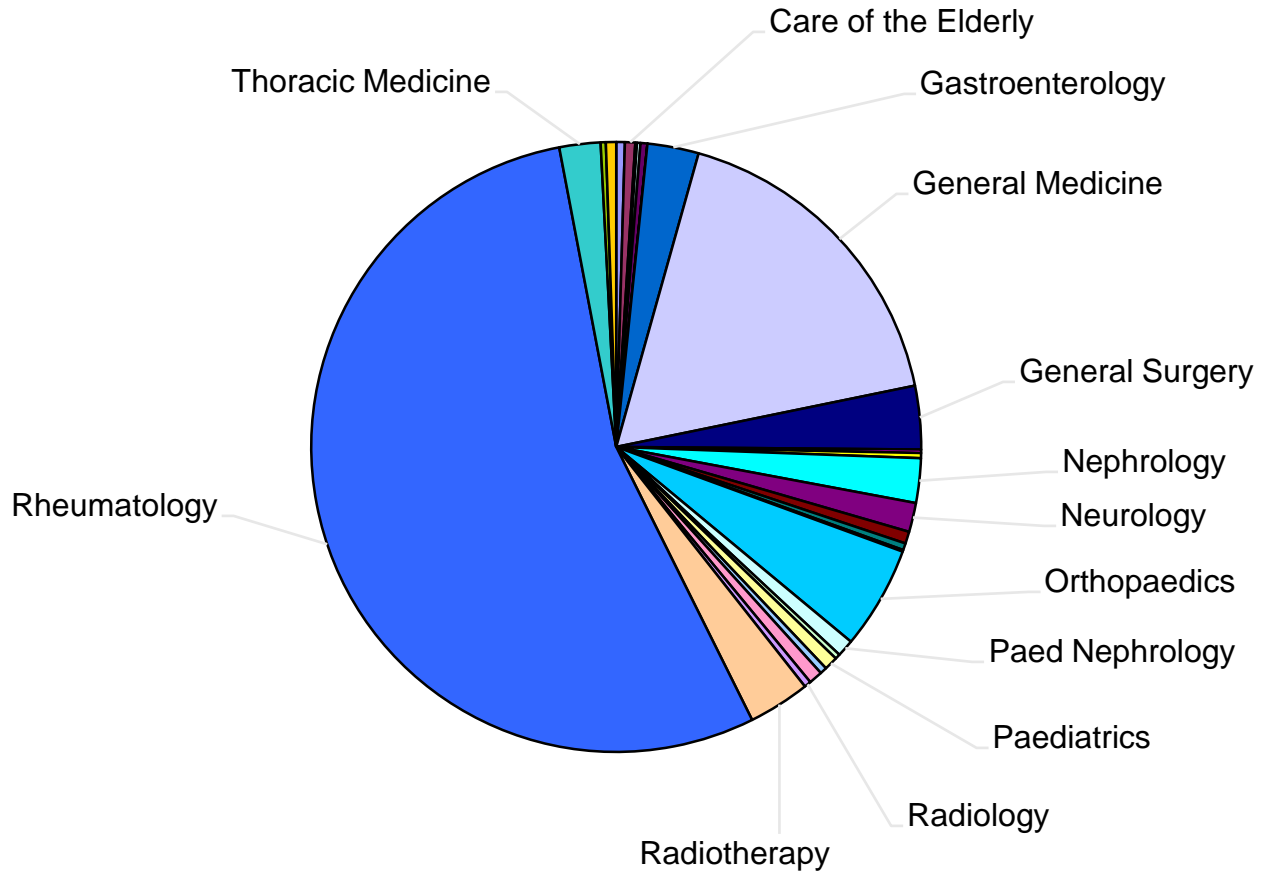
Total Scans (Yellow) : Extra 40 scans per month (11%) over period

Bone Clinic (Blue): Extra 30 scans per month (19.5%) over period

One Stop (Pink): new bone clinic patients only up to Aug 2012



# First Quarter 2013



# Case study

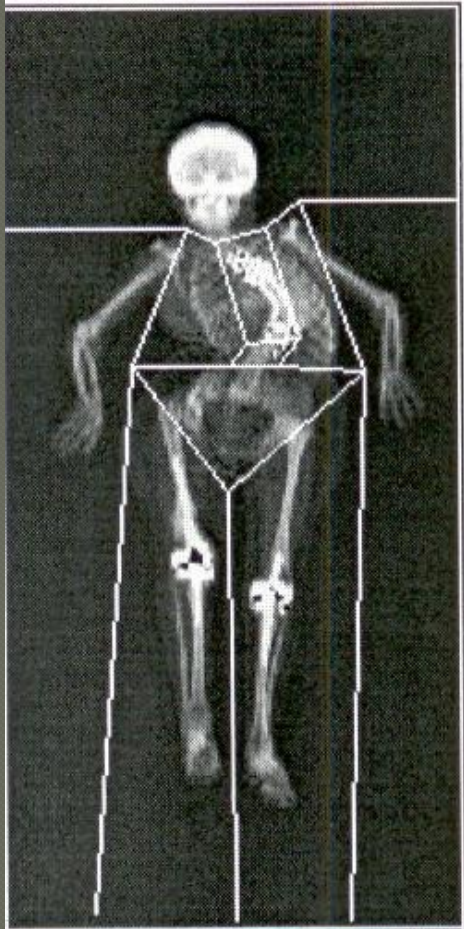
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- A young women with severe scoliosis

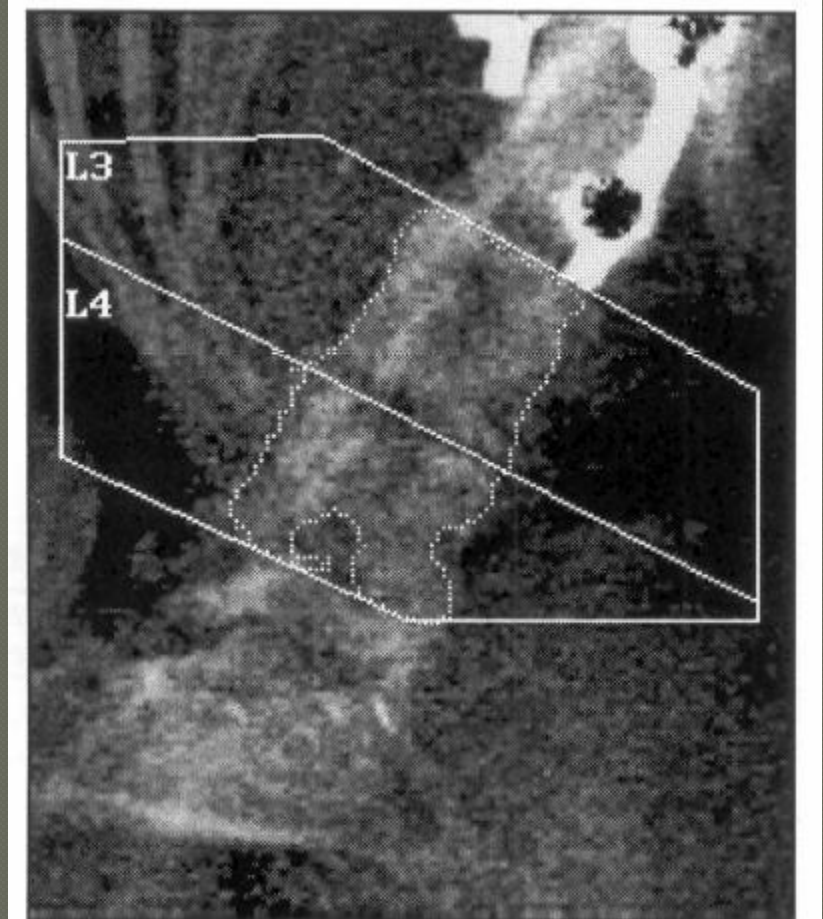
# Relevant History

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- Young woman aged 31
- Severe Scoliosis
- Multiple joint replacements
- Limited Mobility



1 0000 10 00 5007



# Findings

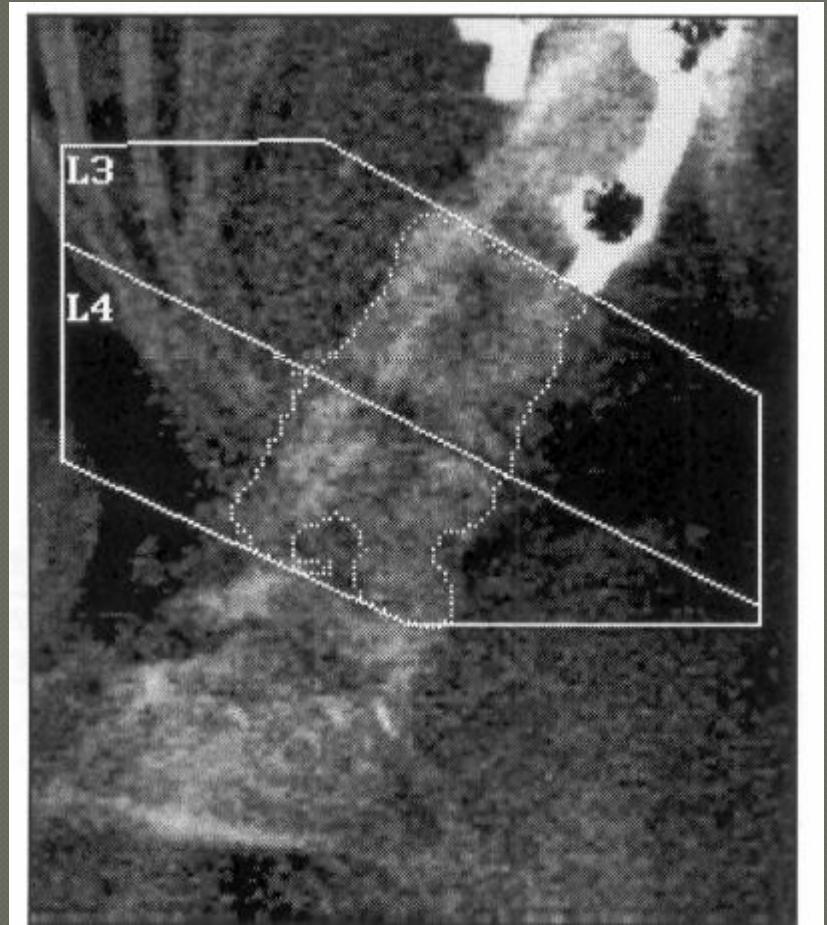
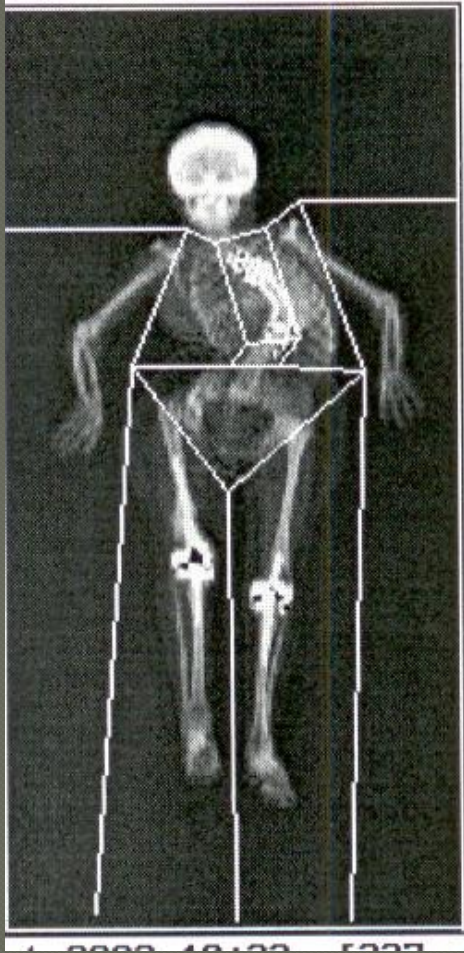
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- Measured at 26 years and 32 years
- $Z(L3/L4) = -2.0$  and  $-2.5$
- $Z(WB) @32: +8.73$  [influenced by metal]
- BMD(pelvis) @32: close to expected
- BMD(arms) @32: 20-30% below expected

# Conclusions?

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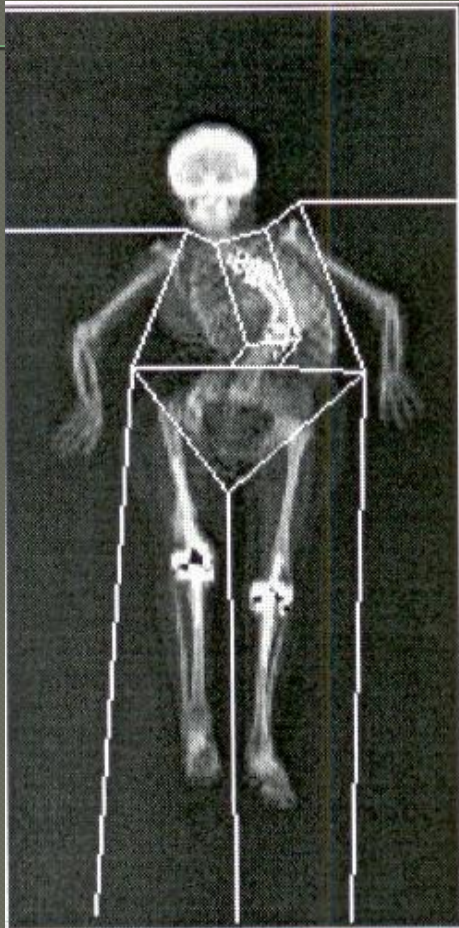
- L3/4 may not be representative
- Whole body BMD not diagnostic
- Some evidence of localised bone loss
- Utility of forearm measurements?
- Interpret all BMD with caution



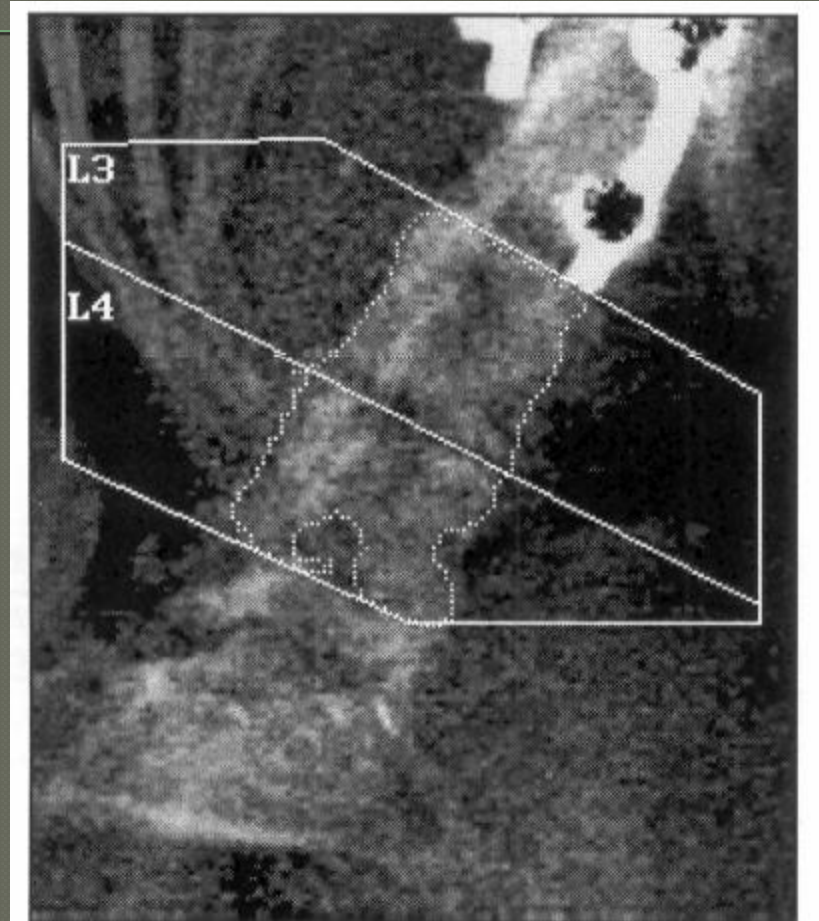
# Conclusions?

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- L3/4 may not be representative
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1 8888 18 100 5027



# Conclusions?

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- L3/4 may not be representative
- Whole body BMD not diagnostic
- Some evidence of localised bone loss
- Utility of forearm measurements?
- Interpret all BMD with caution



Utility of forearm measurements?  
Interpret all BMD with caution

# Other patient pathways?



# Emergency Admissions Unit (W43)

Dr Ashley Price Dr Soumya Kumble

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- 50-60 patients per day admitted
- Variety of conditions
- Indications for DXA / treatment?
- 62 admissions over 3 days Jan 2009
- FRAX tool
- N=21 reassurance, N=14 DXA; N=11 Treat
- Need largely unmet at present
- Very early days

# Radiology Reports

Dr Geoff Hide; Sister Sharon Abdy

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- ◉ Spine x-rays may identify vertebral fracture
- ◉ Not always followed up by GP
- ◉ Indications for DXA / treatment?
- ◉ Local audit suggests several per month
- ◉ Supported by data from Glasgow
- ◉ Pathway to DXA / Treatment unclear

# GP Referrals

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- ◉ Co-ordinated case finding strategy
- ◉ Led by single GP practice
- ◉ Selected patients offered DXA at FRH
- ◉ Technical report produced
- ◉ Service level agreement (SLA) in place
- ◉ Approx 20 patients per month
- ◉ On-going

# What's new

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- **Vertebral Fracture Assessment**
  - Total spine x-ray at same time as BMD
  - Lower dose than conventional spine x-rays
- **Electronic requesting (soon)**
  - RVI scoliosis assessments now electronic
- **Electronic reports (eventually)**

# A patient who has already fractured could fracture again...

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- fragility fracture (45+)
- no trauma (standing height or less)
- includes vertebral deformity
- independent risk for further fracture
- an indication for treatment without BMD RCP Guidelines 2001
- **Pre-existing fracture and low BMD greatly increases risk of future fracture**

# May identify pre-existing fractures by semi-quantitative grading...

**Normal  
(Grade 0)**



**Mild fracture  
(Grade 1, ~20-25%)**

**Moderate fracture  
(Grade 2, ~25-40%)**

**Severe fracture  
(Grade 3, ~40%)**

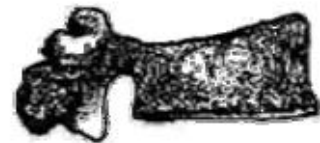
**Wedge fracture**



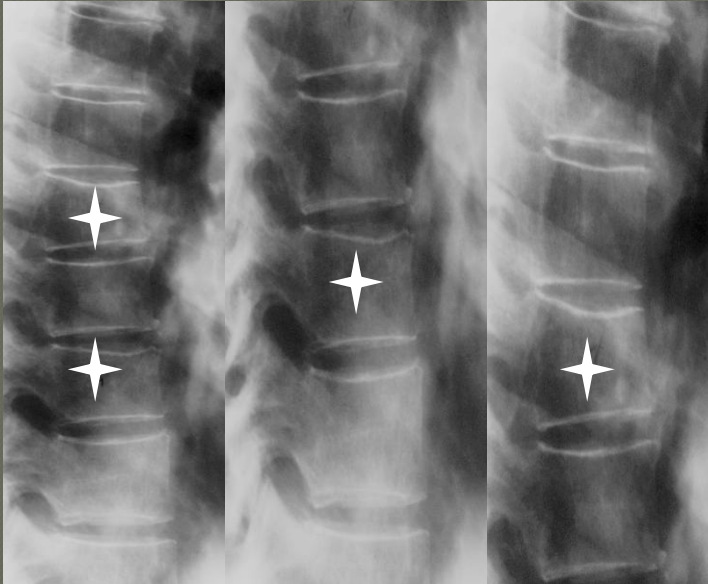
**Biconcave fracture**



**Crush fracture**

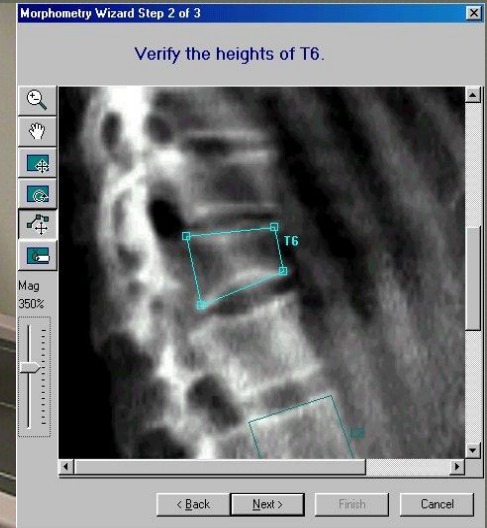


# Lateral spine x-rays allow visual grading against the grading chart...



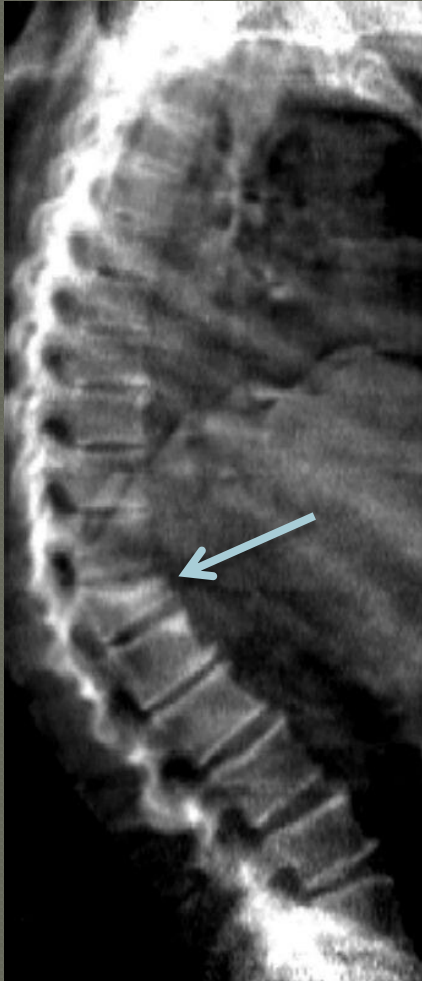
or a quantitative assessment  
measuring vertebral heights directly

# ...or we may identify and grade vertebral fractures using DXA



- So BMD and vertebral morphometry together one stop shop!

# Advantages of morphometry by DXA



- semi-automated alignment
- semi-automated positioning
- better penetration (100-140kVp)
- single view for whole spine
- dose about 1/50<sup>th</sup> (0.006mSv)
- Includes BMD assessment
- single or dual energy options
- able to see most vertebra?

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Why did the X-ray only take 1  
suitcase away on holiday?

It was travelling light!

Thankyou

Any Question?