

## Joint CI-JAI advanced accelerator lecture series Imaging and detectors for medical physics

# Lecture 2: Detectors for medical imaging

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### **Course layout**

Day	AM 09.30 – 11.00	PM 15.30 – 17.00				
Week 1						
6 <sup>th</sup> June	Lecture 1: Introduction to medical imaging	Lecture 2: Detectors for medical imaging				
7 <sup>th</sup> June	Lecture 3: X-ray imaging					
8 <sup>th</sup> June		Tutorial				
Week 2						
13 <sup>th</sup> June	Lecture 4: Radionuclides					
14 <sup>th</sup> June	Lecture 5: Gamma cameras	Lecture 6: SPECT				
16 <sup>th</sup> June	Lecture 7: PET					
Week 3						
22 <sup>nd</sup> June	Tutorial					



### **Books**

- 1. N Barrie Smith & A Webb Introduction to Medical Imaging Cambridge University Press
- 2. Edited by M A Flower Webb's Physics of Medical Imaging CRC Press
- A Del Guerra Ionizing Radiation Detectors for Medical Imaging World Scientific
- 4. W R Leo

Techniques for Nuclear and Particle Physics Experiments Springer-Verlag



# From particle physics to medical imaging

"The significant advances achieved during the last decades in material properties, **detector** characteristics and high-quality electronic system played an everexpanding role in different areas of science, such as high energy, nuclear physics and astrophysics. And had a reflective impact on the development and rapid progress of radiation detector technologies used in medical imaging." D. G. Darambara, Nucl. Inst. And Meth. A 569 (2006) 153-158

- Nuclear medicine: fertile field for applications of technology from high energy physics
- Main goal: better image quality for same dose delivered or reduce dose delivered for same image quality



## Analogue –vs– digital

Analogue imaging (films)	Digital imaging
Continuous range of possible optical densities up to some limiting value	Discrete and limited range of optical densities
Narrow exposure latitude → strict exposure requirements	Very wide exposure latitude
Little possibility of image processing	Image processing possible + needed to overcome limitations of manufacturing processes (bad pixels, spatial sensitivity variation)
Only one image display	Various image displays possible
Cheap but one use	Expensive but multiple use
High resolution	Low(er) resolution <sup>1</sup>

<sup>1</sup>Not clinically significant when choosing right matrix size and image receptor to match the application

Analogue  $\rightarrow$  digital imaging (= more quantitative information available) thanks to:

- 1. Integrated electronics
- 2. Fast computers



# Specific requirements for medical imaging detectors

• Detector for medical application = special detector with its own specifications:

Detection range	- Low energies: 18 keV for mammograms for ex.
Read-out	<ul> <li>No trigger (no bunch crossing) → self-triggering electronics or free running</li> <li>High acquisition rates &gt; GHz</li> <li>Manageable number of read-out channels</li> </ul>
Event size	<ul> <li>Can be small 1 bit ÷ 10 bytes</li> </ul>
Geometry	<ul> <li>Large area often required</li> <li>Almost no dead space</li> </ul>
Patient's requirements	<ul> <li>Meet stringent ethical requirements and regulation</li> <li>Ensure patient's comfort</li> </ul>
Market	<ul> <li>Can be large: 10<sup>3</sup>÷10<sup>6</sup> units</li> </ul>



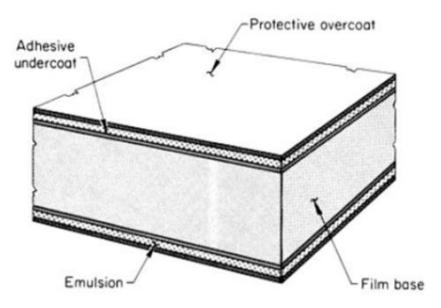
## **Digital X-ray radiology**

- Primary requirements for digital X-ray detector for radiology:
  - 1. Speed
  - 2. Area
  - 3. Spatial resolution
  - 4. Dose rate to exposure ratio
- Trade-off between:
  - 1. Spatial resolution & dynamic range
  - 2. Read-out speed & time resolution
  - 3. Contrast & exposure time



## **Radiographic films**

Ref. 3 – Chapter 2.3



- Photographic emulsion = mixture of silver halide grains of different size + gelatine
- Emulsion deposited on transparent support = base
- Emulsion coated with thin protective layer
- Limitations:
  - 1. Limited dynamic range
  - 2. Lack of digital processing



## Image formation

- X-ray arriving on film interact with silver halide crystals  $\rightarrow e^-$  released
- e<sup>-</sup> migrates until trapped in a grain where neutralises silver ion
- If about four silver ions are captured and neutralised at same grain → stable configuration reached = grain said to be sensitised
- Development of film = sensitised grains are reduced to metallic silver + unsensitised grains are removed
- 1 X-ray sensitises one or more grains, while optical photons sensitise only one grain



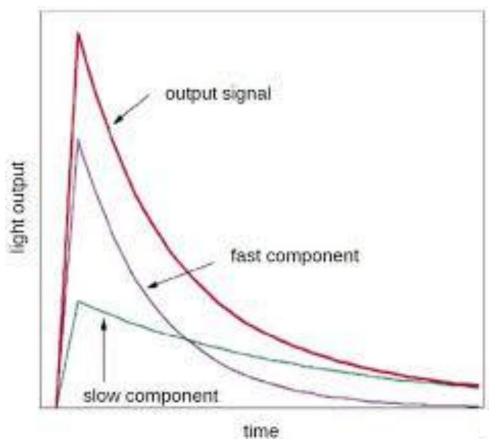
## **Scintillators**

Ref. 4 – Chapter 7

- Scintillation = emission of light generated by the passage of a particle / radiation
- Radiation loses energy = photoelectric + Compton interactions → excitation of atoms and molecules → de-excitation = emission of optical photons:
  - 1. Fluorescence: reemission occurs within  $10^{-8}$  s  $\cong$  time for atomic transition
  - 2. Phosphorescence or afterglow: excited state metastable  $\rightarrow$  reemission delayed by few  $\mu$ s to hours depending on scintillator material



## **Scintillation light**

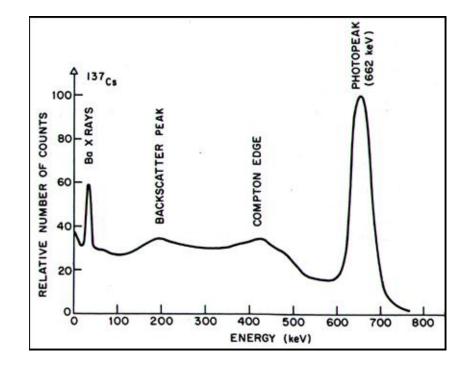


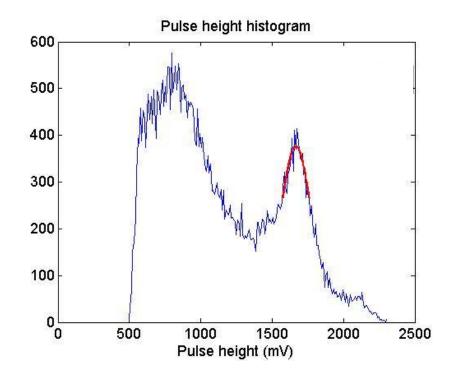
 $N = A \exp\left(\frac{-t}{\tau_f}\right) + B \exp\left(\frac{-t}{\tau_s}\right)$ N = n of photons emittedat time t $\tau_f, \tau_s = \text{decay constants of}$ fast and slow components

*A*, *B* = relative amplitudes of fast and slow components



## Example output signal: <sup>22</sup>Na source





FP7 funded ENVISION project – University of Oxford With help from University of Sussex



## **Scintillators**

	Inorganic	Organic		
Material	Mainly alkali halides with small activator impurity	Aromatic hydrocarbon compounds with benzene-ring structures		
Density	High	Low		
Atomic number	High	Low		
Stopping power	High	Low		
Light output	High	Low		
Energy resolution	High	Low		
N of photons generated	Linear <sup>1</sup>	Non linear <sup>1</sup>		
Decay time	~500 ns	Few ns or less		
Temperature dependence	Yes	No		
Hygroscopic	Usually yes	No		

<sup>1</sup>With energy of incident radiation



# Commonly used inorganic scintillators

	Nal(TI)	Ba₂F	BGO	GSO	LSO	LYSO	LaBr <sub>3</sub>
Density (g cc <sup>-1</sup> )	3.67	4.89	7.13	6.71	7.4	7.3	5.1
Effective atomic number	51	54	75	58	66	66	~50
Relative light yield	100	5 16	15	20	75	75	166
Light decay time (ns)	230	0.6 620	300	~60	~40	40	16÷30
Emission wavelength (nm)	410	195÷220 310	480	430	420	428	380
Refractive index	1.85	1.49	2.15	1.85	1.82	1.82	~1.9
Attenuation length (mm) at 511 keV	25.6	22.7	11.2	15.0	11.4	11.6	21.3



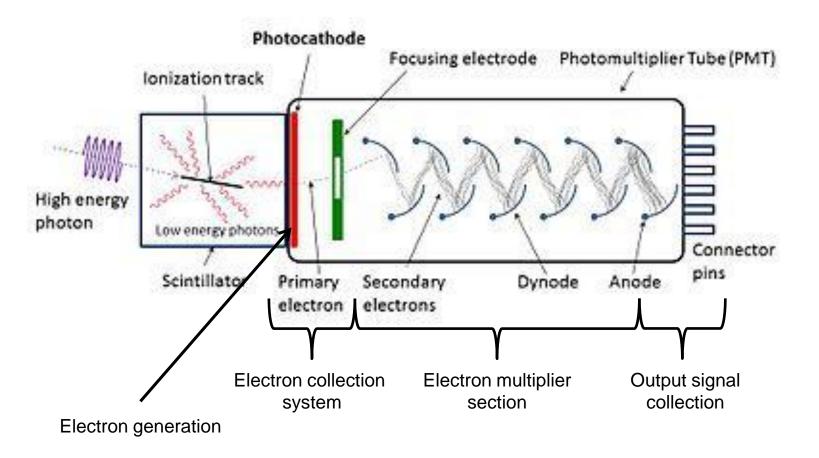
### **Photodetectors**

- Convert light into electric signal
- Various types:
  - Photomultiplier tubes (PMTs)
  - Photodiodes (PD) / Avalanche Photodiodes (APDs)
  - Silicon Photomultipliers (SiPMs)
  - Vacuum Phototriodes (VPTs)



## **Photomultipliers tubes**

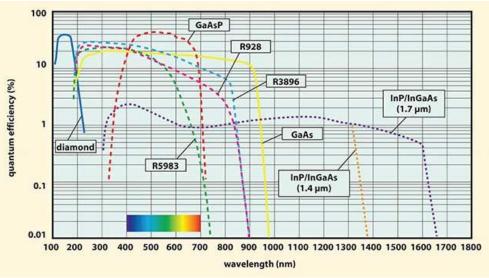
Ref. 4 – Chapter 8





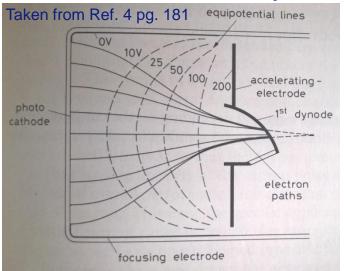
## **PMT: the various stages**

#### Photocathode



- Quantum efficiency  $\eta(\lambda)$ :  $\eta(\lambda) = \frac{N_{pe}}{N_{\gamma}}$
- Typical η are 10%÷40% for λ between 200÷800 nm

#### Electron collection system

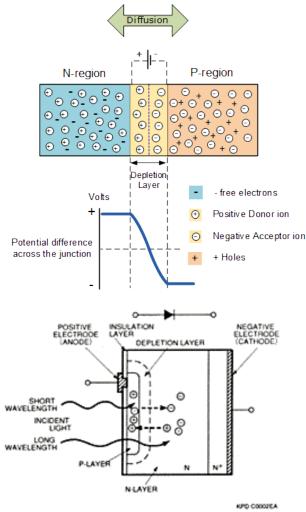


#### **Electron multiplier section**

- Conventional PMTs 10÷14 stages = gain up to 10<sup>7</sup>
- Dynodes kept at increasingly higher V



## Photodiodes (PDs)



- *p*-*n* junction in a silicon substrate wafer
- Incident light creates e-h pairs in the junction
- No intrinsic amplification → small output signal that requires post amplification →
  - Better for applications with large number of optical photons produced → no amplification
  - Amplification introduces noise, especially if fast signal required

## Avalanche Photodiodes (APDs)

- APD = PD operated near breakdown voltage = higher operating bias 100÷200 V
- High internal electric field → internal avalanche multiplication effect:
  - $\rm e^-$  generated is accelerated by high electric field  $\rightarrow$  creates free carriers by impact ionisation
- Internal gain = bigger output signal:
  - 1. Gain usually  $10^2$

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2. Gain of  $10^3$  can be achieved using special doping techniques but requires very high voltage  $\sim 1500$  V



#### PDs –vs– APDs

	PDs	APDs
Output signal	Small	Large
Spectral sensitivity	Red/green	Red/green
Response time	Fast	Fast
Bias	10÷100 V	$100 \div 200 \text{ V}$ ~1500 V for special APDs <sup>1</sup>
Gain	$G_{PD} = 1$	$G_{APD} = 10^2 \times G_{PD} @ 100 \div 200 V$ $G_{APD} = 10^3 \times G_{PD} @ \sim 1500 V$
Sensitivity to temperature	Yes	Yes
Area	Few mm across	Few mm across
Cost	Cheap	More expensive

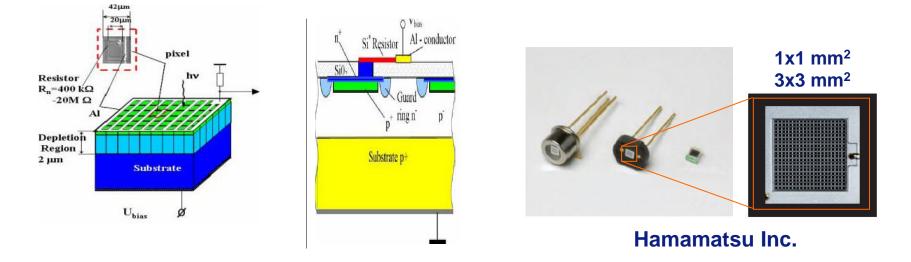
<sup>1</sup>For APDs doped with special doping techniques to obtain very high gain

 $Signal_{PD}(1), Signal_{APD}(10^2 \div 10^3) < Signal_{PMT}(10^5 \div 10^8)$ 

 $\rightarrow$  Limited application to nuclear imaging so far



### **SiPMs**



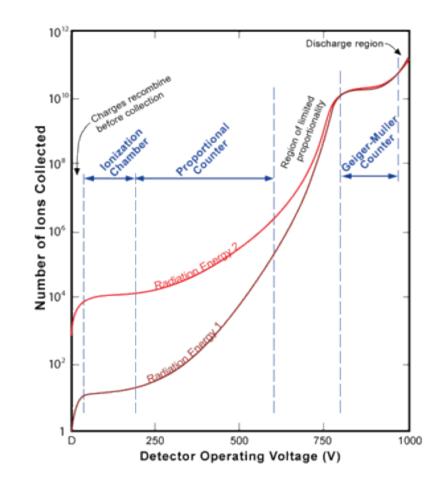
- Array of Silicon Photodiodes on common substrate each operating in Geiger mode
- SiPMs have high speed (sub ns) and gain (10<sup>6</sup>) at low bias (< 100 V) and work in high magnetic fields (7 T)</li>



### **Gaseous detectors**

Ref. 4 – Chapter 6

- Volume filled with gas mixture with applied electric field
- Electron ion pairs created
- Number of pairs collected depends on voltage applied → various regions:
  - 1. Recombination
  - 2. Ionisation
  - 3. Proportional
  - 4. Limited proportionality
  - 5. Geiger-Muller





# Gaseous detectors in medical imaging

Ref. 2 – Chapter 5.3.5 and Ref. 3 – Chapter 6.3

- Disadvantages:
  - Higher energy threshold for production of ionisation than other detectors  $\rightarrow$  less ionisation produced for same incident energy  $\rightarrow$  lower energy resolution
  - Low  $Z \rightarrow$  low stopping power

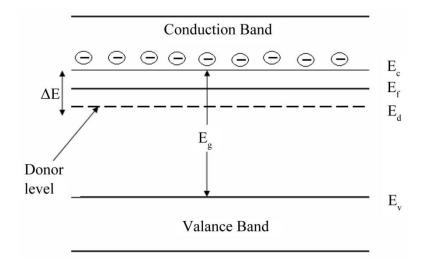
 $\rightarrow$  Limited application in medical imaging



## **Semiconductor detectors**

Ref. 4 – Chapter 10, Ref. 3 – Chapters 3.3 and 6.4

 Crystalline material with energy band structure = valence, conduction bands and 'forbidden' energy gap



- Semiconductor detectors –vs– gaseous detectors
  - Advantages:
    - Only few eV needed to create e—h pair in Si and Ge → for given energy greater ionisation produced than in gas → higher energy resolution
    - 2. Higher Z than gas  $\rightarrow$  greater stopping power
  - Disadvantages:
    - 1. More expensive



## **Signal formation**

• Ionising radiation  $\rightarrow e-h$  pair  $\rightarrow$  Charge carriers =  $e^-$  in conduction band +  $h^+$  in valence band

$$J = e n_i (\mu_e + \mu_h) E$$

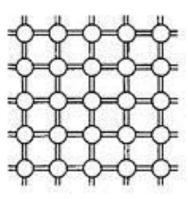
- J =current density
- $n_i =$  electron / hole concentration
- $\mu_e$ ,  $\mu_h$  = electron and hole mobility
- E = electric field
- If material fully depleted = solid-state ionisation chamber  $\rightarrow$  very high detection efficiency for each e-h pair

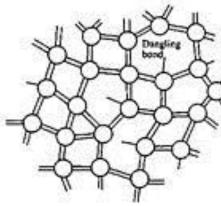


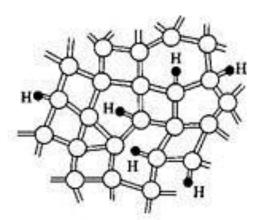
## **Amorphous semiconductors**

http://chemwiki.ucdavis.edu/Textbook\_Maps/Inorganic\_Chemistry\_Textbook\_Maps/Map%3A\_Inorganic\_Chemistry\_y\_(Wikibook)/Chapter\_10%3A\_Electronic\_Properties\_of\_Materials%3A\_Superconductors\_and\_Semiconductors/1 0.7\_Amorphous\_semiconductors

- Disordered or glassy form
- Two most studied: a-Si and a-Se







Crystalline Si

Amorphous Si a-Si

a-Si:H

Advantage: can be produced in large areas



## **Semiconductor materials**

	Si	a-Si	a-Se	Ge	CdZnTe
Atomic number	14	14	34	32	48,30,52
Average atomic number	14	14	34	32	49.1
Density (g/cm <sup>3</sup> )	2.33	2.30	4.30	5.32	5.78
Bandgap (eV)	1.12	1.80	2.30	0.66	1.572
$e^-$ mobility $\mu_e$ (cm <sup>2</sup> /Vs)	1400	1	0.005	3900	1000
$e^{-}$ lifetime $\tau_e$ (s)	>10 <sup>-3</sup>	7x10 <sup>-9</sup>	10 <sup>-6</sup>	>10 <sup>-3</sup>	3÷5x10 <sup>-6</sup>
h <sup>+</sup> mobility $\mu_h$ (cm <sup>2</sup> /Vs)	480	0.005	0.14	1900	50÷80
$h^+$ lifetime $\tau_h$ (s)	2x10 <sup>-3</sup>	4x10 <sup>-6</sup>	10 <sup>-6</sup>	10 <sup>-3</sup>	10 <sup>-6</sup>
$\mu_e \tau_e (\mathrm{cm}^2/\mathrm{V})$	>1.4	7.0x10 <sup>-9</sup>	5.0x10 <sup>-9</sup>	>3.9	3÷5x10 <sup>-3</sup>
$\mu_h \tau_h (\mathrm{cm}^2/\mathrm{V})$	0.96	2.00x10 <sup>-8</sup>	1.40x10 <sup>-7</sup>	1.90	5÷8x10 <sup>-5</sup>
Ionisation energy (eV/e-h pair)	3.62	4.00	20÷60 <sup>1</sup>	7.90	4.64
Dielectric constant	11.7	11.7	6.6	16.0	10.9
Resistivity ( $\Omega$ cm)	<104	10 <sup>12</sup>	<b>10</b> <sup>12</sup>	46	3x10 <sup>10</sup>

<sup>1</sup>Depends on ionising energy



## Si planar detectors

Ref. 3 – Chapter 6.4

- Si material
  - Advantages:
    - Spatial resolution down to few  $\mu\text{m}$  for microstrips & pixels
    - Excellent energy resolution = few keV @ room temperature
  - Disadvantages:
    - Low  $Z \rightarrow$  low stopping power
- Two different planar configurations:
  - 1. Strip detectors = separate read-out strips
  - 2. Pixel detectors = one of two electrodes segmented in small area cells called pixels

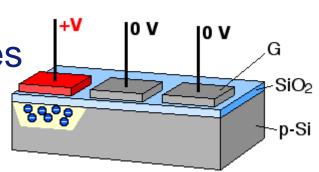
## **Charge-Coupled Devices (CCDs)**

Ref. 2 – Chapter 2.8.3.2

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- Integrated circuit = series of electrodes deposited on Si substrate + depletion region below = 2D array of MOS capacitors = 2D array of potential wells
- Ionising radiation → photoelectron created and trapped in wells = charge distribution = image
- Image read out shifting the charge from one well to next until it reaches digitising electronics





## CCDs Pro and Cons

- Advantages:
  - Transfer efficiency very high = approaching 100%
  - Read-out times of 30 frames/s possible with 1000x1000 pixels
- Disadvantages:
  - Small sensitive area limited by engineering to 5 cm.
     Possible solutions:
    - Separate arrays may be 'patched' together but 'patch' lines may be visible on images
    - Demagnification optical system  $\rightarrow$  better solution
  - Low detection efficiency for photons with E > 30 keVand charged particles  $\rightarrow$  used coupled with scintillators

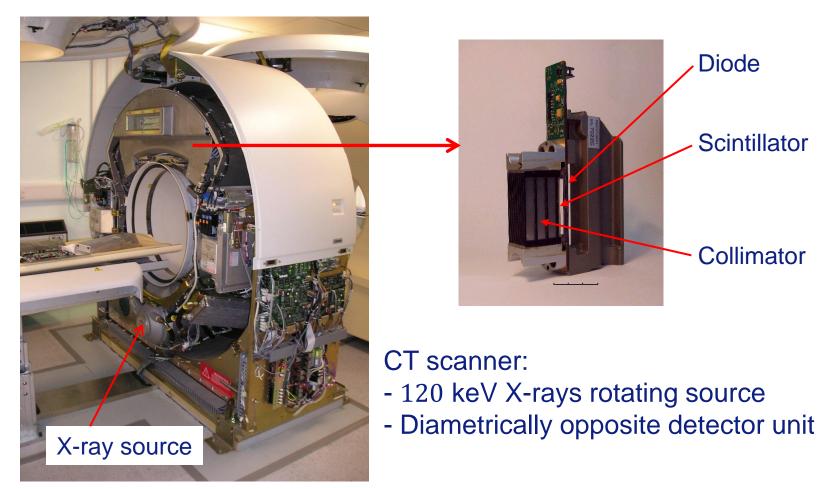


## **Flat panels**

- Flat panel detector composed of:
  - 1. a-Si:H pixel detector:  $\cong \mu m$  thin a-Si:H sensor coupled to array of thin film transistor (TFT) = high sensitivity to visible light, low sensitivity to X-rays
  - 2. Coating:  $400 \div 500 \ \mu m$  thick layer of phosphor or scintillator = high sensitivity to X-rays
- Incident X-ray → converted into green light by the coating → green light converted into electric signal by the a-Si:H pixel detector
- Used in general radiography, mammography, fluoroscopy



#### **CT** scanners





#### **SPECT scanners**

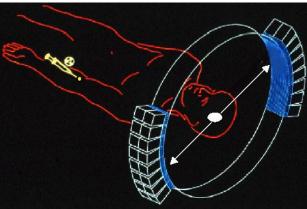


- <sup>99</sup> Tc<sup>m</sup> injected into body  $\rightarrow$  one 141 keV  $\gamma$
- SPECT scanner: two or three rotating detector heads = gamma camera

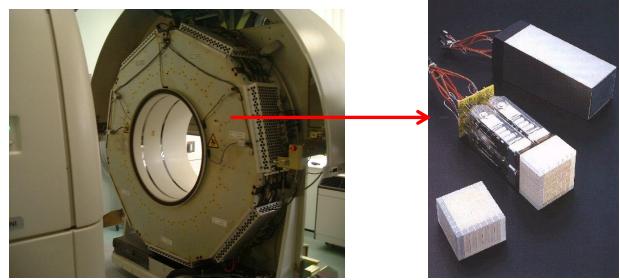




### **PET scanners**



- −  $β^+$  emitter injected into body → two back-to-back 511 keV γ from e<sup>+</sup> decay
- PET scanner: ring of detectors



Courtesy Mike Partridge (Oxford)



## Imaging systems for X-ray radiotherapy

#### • Cone Beam CT (CBCT)

- CT scanner but with X-ray beam diverging (cone)
- Electron Portal Imaging Devices (EPIDs):
  - Used to check patient set-up and field placement
  - Measure therapeutic beam exiting the patient
  - Detectors:
    - Camera based
    - Liquid-ionising-chamber based
    - a-Si flat-panel based

