

### X-ray imaging

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#### **Röntgen's first "medical imaging"**







#### Improvements....

# System-specific image processing



# Adaptive auto-windowing algorithm



#### **Basic principles**











absorption vs phase contrast imaging



d and ß as a function of energy in keV for biological tissue



#### absorption vs phase contrast imaging

#### different contrast generation





#### absorption vs phase contrast imaging

#### different contrast quality





absorption

#### phase contrast

http://www.medphys.ucl.ac.uk/research/acadradphys/researchactivities/pci.htm

### absorption imaging: simulations

- high resolution voxel models of breast
- created from CT-scans of anatomical breast specimens
- voxel size: 60 x 60 x 60 μm<sup>3</sup>
- segmentation in different tissues:
  - adipose
  - glandular
  - skin

- using brilliant undulator radiation: beam geometry, spectral angular flux,...
- simulation of absorption and scattering processes with Geant4-Software-Toolkit









#### absorption imaging: contrast reduction





#### absorption imaging: contrast reduction

0.2 mGy average glandular dose at ~10<sup>11</sup> photons





#### absorption imaging: contrast enhancement



- geometrical limitation of radiation (aperture)
- anti-scatter grid
  ideal: focused
  anti-scatter grid

Anti-Scatter grid is placed directly on the film /screenfilm-system / detector





#### absorption imaging: applied dose





## b) Collective effective dose from medicine





#### absorption imaging: applied dose

criterion = signal-difference-to-noise ratio versus averaged-glandular dose



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#### absorption imaging: applied dose





#### dual energy



Lung perfusion dual energy CT, courtesy of LMU

Actual: dual energy CT with **contrast** media for e.g. lung perfusion

- → New diagnostic tool
- → at least two scans
- ➔ high dose
- ➔ registration problems
- ➔ non-optimal image information due to overlapping spectra
- →Limitations for suitable markers

With especially designed "monoenergetic" radiation low dose individualised medicine will be developed together with MAP II

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#### dual energy









**Gout Visualization** 





#### coherent scatter imaging

- Differentiation between tissue structures
- most notably between malign and benign carcinomas by
  - different scatter intensity caused by change in the collagen structure of malign tissue
  - different scatter distribution
- Differentiation between malign and benign tissue without the need for biopsy would be a great benefit



Image of scatter distribution in tumour







#### fluorescence imaging: experiments







#### fluorescence imaging: experiments



**Figure 8.** Reconstructed GNP distribution and location within the PMMA phantom using experimental data shown in figure 7.

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#### fluorescence imaging: background



max. sensitivity at 1mGy / 5mm pixel / CNR=5 ~ **10µg/ml** ~ **100 times** more sensitive than transmission-CT

(simulation by B. Müller, HZM/MAP)



#### **Physicists' impacts**

- new **methods**:
  - phase-contrast with low-brillance sources
  - absorption/x-ray fluorescence with high-brillance sources
- pushing the **limits**:
  - dose reduction
  - lowering the concentration of contrast media
  - enhancing the sensitivity
  - enforcing persoanlized medicine
- **cooperations** between medicine and physics:
  - possibly comparative studies between UKE and my group (magnetic particle imaging vs X-ray fluorescence imaging
  - new detectors for medicine (Erika!!)