

**DIGITAL IMAGE:** Stored in RAM as a 2-D array:

gamma-camera: 32x32, 64x64, 128x128, 256x256

CT, MRI, US: 256x256, 512x512

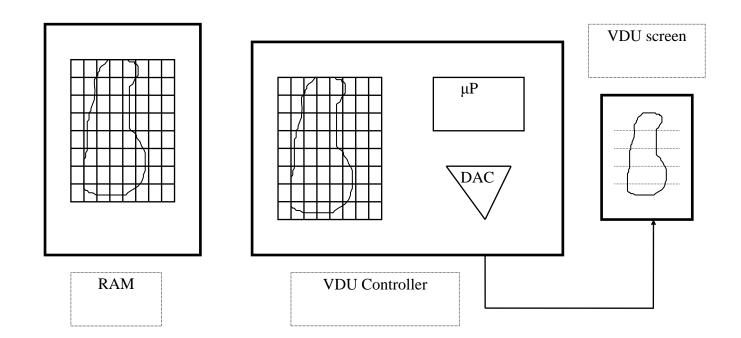
DSA: 512x512, ..., 2048x2048.

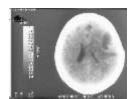
#### **Image depth:**

• Number of quantization levels,

• It's the 3<sup>rd</sup> dimension, e.g. 512x512x8, where 8 is the number of bits, or 2<sup>8</sup>.

## **IMAGE DISPLAY** (1)





#### **IMAGE DISPLAY(2)**

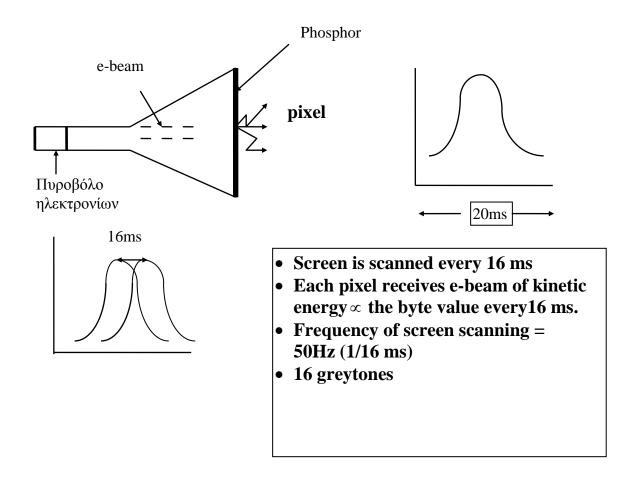
#### **VDU-CONTROLLER:**

- μP reads byte
- Transfers byte to level e.g. for 16 greytones (0-15)

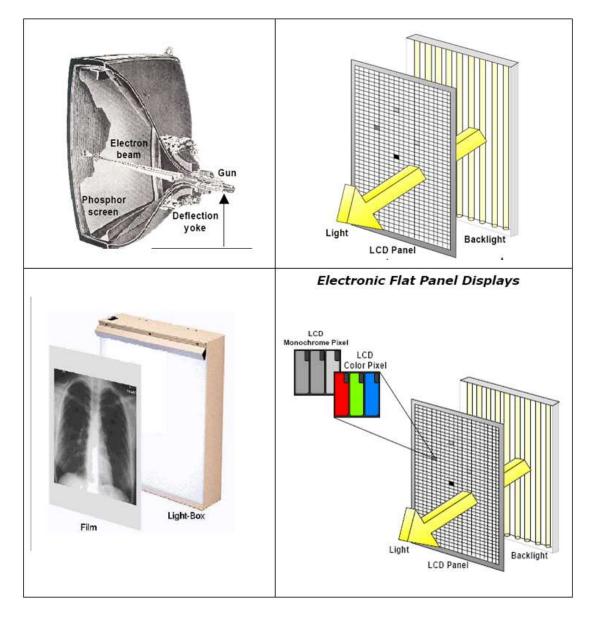
15	150	240	: 16	0	9	15
39	80	62	<del></del>	2	5	4
0	33	230		0	2	14

- Calculates level (e.g. 5 x 8 µV D.A.C.)
- Applies dc Voltage to VDU's electron gun

#### **IMAGE DISPLAY (3)**



Active Matrix LCD display screens consist of a matrix of thin film transistors (TFT), one TFT per pixel. TFTs are micro-switching transistors (and associated capacitors) that are arranged in a matrix on a glass substrate to control each picture element (or pixel). Switching on one of the TFTs will activate the associated pixel. By adjusting the voltage applied on the TFT, in small increments, e.g. 1/256 or 1/1024 increments, it is possible to create the gray-scale effect. For colour display, 3 sub-pixels of RGB colors are combined to form a color pixel (<a href="http://www.practical-home-theater-guide.com/lcd-display.html">http://www.practical-home-theater-guide.com/lcd-display.html</a>).



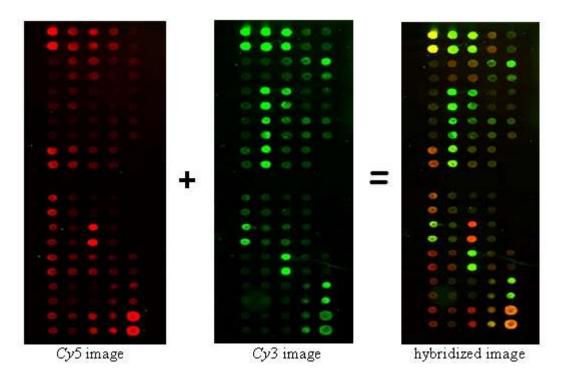
#### Creation of microarray images

cDNA microarrays consist of thousands (e.g. 6000) of individual cDNA sequences printed in a high-density array on a glass microscope slide using a robotic arrayer

(see <a href="http://www.bio.davidson.edu/courses/genomics/chip/chipQ.html">http://www.bio.davidson.edu/courses/genomics/chip/chipQ.html</a>).

For two mRNA samples (extracted from 2 classes such as normal-abnormal cases), the two samples or targets are reverse-transcribed into cDNA, are labeled using different fluorescent dyes (e.g., the red-fluorescent dye Cy5 and the green-fluorescent dye Cy3), and are then mixed and hybridized with the arrayed cDNA sequences on the slide.

After this hybridization, the slides are imaged using a scanner which makes fluorescence measurements for each dye (red and green). Microarray images (16-bit) can be acquired using a number of devices, including a laser scanning confocal microscope, or scanner, and a charge coupled device (CCD) camera.



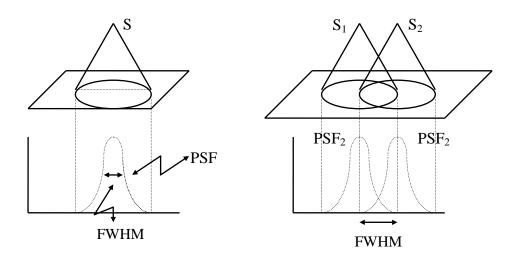
# **IMAGE QUALITY**



#### **Image quality (1)**

- Medical Image, it's related to the clarity of the specific information sought for in the image by the observing physician.
- objective image quality features for assessing image quality:

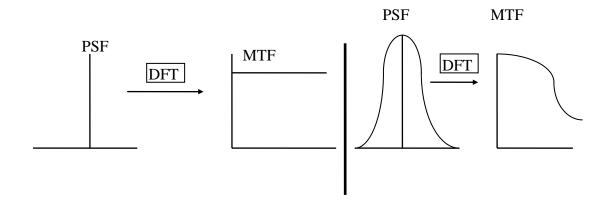
**Sharpness- Contrast - Noise** 

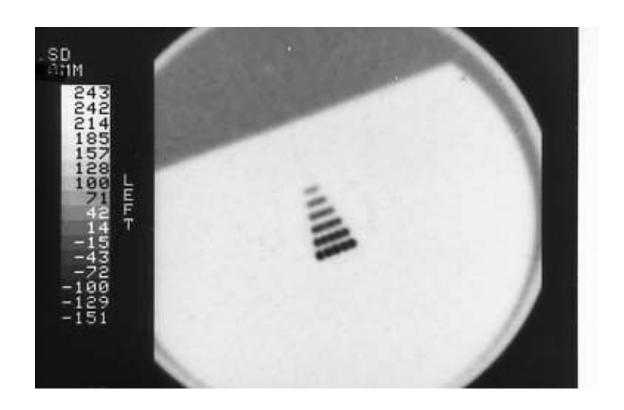


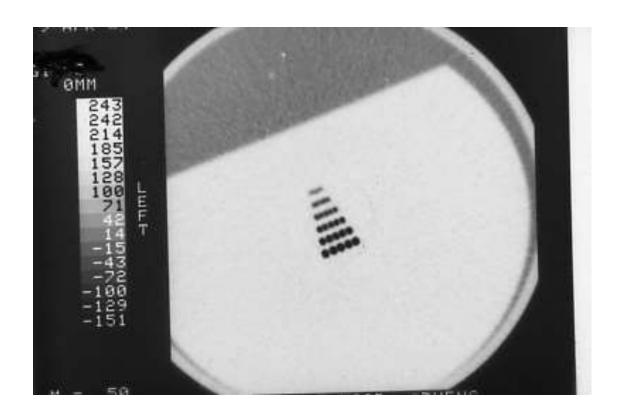
#### **Image quality (2)**

Image Sharpness: concerns mainly the ability to distinguish image detail. Image sharpness is assessed by the image spatial resolution, i.e. the ability of the imaging system to distinguish (i.e. clearly display) two small high contrast dots close to each other.

Quantitatively, the spatial resolution is determined by the smallest distance between two distinguishable high contrast dots or by the magnitude of the FWHM (Full Width at Half Maximum) or by the LSF (the number of distinguishable lines per cm) or by the MTF (Modulation Transfer Function).

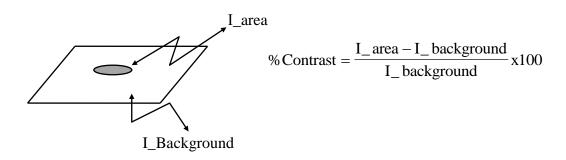






## **Image quality (3)**

Image contrast, concerns the ability to distinguish image detail of low contrast with its surrounding background.



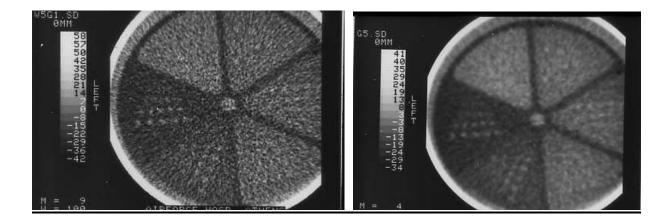
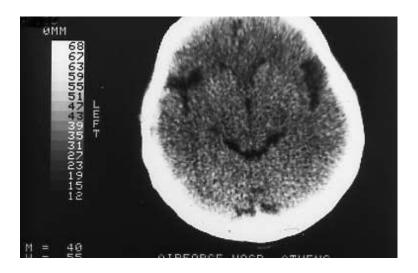
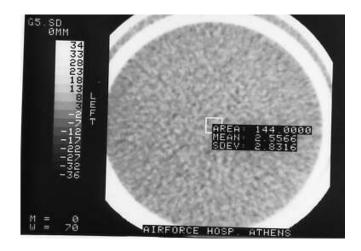


Image noise, is of statistical nature and signal dependent but, without great loss of accuracy, it can be considered additive and white. The noise contribution to each pixel of the medical image is not known but an assessment of the overall noise contribution to the image can be obtained from:

Noise = 
$$\sqrt{\frac{\sum_{i=1}^{N} (x_i - \overline{\mu})^2}{N}}$$







#### Overall image quality may be approximately evaluated by:

$$Image\_Quality = K \frac{(Sharpness)^2(contrast)^2}{Noise\_Power\_Spectrum}$$

```
%Program 0
% Read & plot image
function []=Program 1();
clc;echo off;close all;
A=[30,31,12,9,
    17,12,25,10,
    12, 8, 17, 9,
    31,12,26,22];
A=double(A); B=A;
disp('original image matrix'); disp(A);
image depth=31;tones=8;
x=size(A,1); y=size(A,2);
disp('plotted image matrix');
for i=1:x
   for j=1:y
        ival=A(i,j);
            tone ival=(tones-1)*(double(ival)-0)/(31-0);
        B(i,j) = tone ival;
    end;
end;
disp(round(B));
close all;
clear A; clear B;
A=imread('C:\Program Files\MATLAB\R2006b\work\Images\Pelvis.BMP');
% A=imread('C:\Program
Files\MATLAB\R2006b\work\Images\01254520 256X256.bmp'); %01254520 256X
256.bmp, AA1a.BMP
A=double(A);
x=size(A,1); y=size(A,2);
sprintf('x= f y=f',x,y);
%expand image to 0-255
max A=max(max(A)); min A=min(min(A));
for i=1:x
    for j=1:y
        val=A(i,j);
            tone ival=(val-min A) *(255/(max A-min A));
        B(i,j) = tone ival;
    end;
end;
max B=max(max(B)); min B=min(min(B));
[min_A max_A min B max B]
% %====== PLOT IMAGES ==========
colormap('gray');
subplot(2,2,1);imagesc(A, [0 255]);xlabel('Original Image');
axis equal; axis([1 size(A,2) 1 size(A,1)]);
subplot(2,2,2);imagesc(B, [0 255]);xlabel('Optimally plotted image');
axis equal;axis([1 size(B,2) 1 size(B,1)]);
```

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#### Output

original image matrix

30 31 12 9

17 12 25 10

12 8 17 9

31 12 26 22

plotted image matrix

7 7 3 2

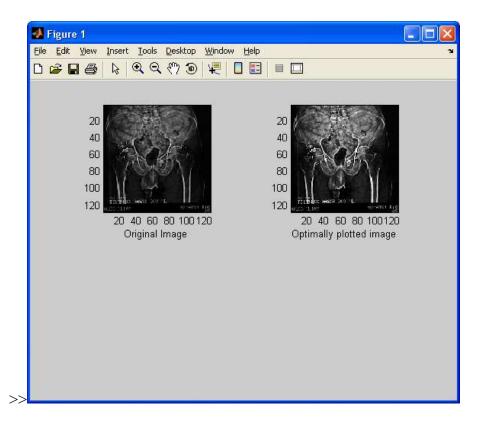
4 3 6 2

3 2 4 2

7 3 6 5

ans =

1.0000 186.0000 0 255.0000



# %Labwork: % 1) construct your own routine for scalling an image to any % interval between 0 and 255 % 2)construct a 20x20 pixels ROI at the center of an image % and calculate the mean value and standard deviation by using % your own basic code and by verifying results with matlab's respective %functions % 3)Plot histograms of the original and a modified (scaled) image. Histogram is a graph of the number of pixels (y-axis) per gray-value (x-axis)

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