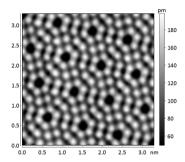


#### **Spatial Calibration**

For image analysis to produce meaningful results, the spatial calibration of the image must be known. If the data acquisition parameters can be read from the image (or parameter) file then the spatial calibration of the image can be determined.



For simplicity and clarity, spatial calibration will not be indicated on subsequent images.

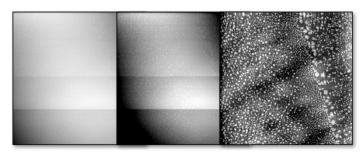
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With Scanning Probe Microscope images there is no guarantee that the sample surface is level (so that the z values of the image are, on average, the same across the image).

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#### **SPM Image Display**

By treating each scan line of an SPM image independently, anomalous jumps in the apparent height of the image (produced, for example, in STMs by abrupt changes in the tunnelling conditions) can be corrected for.



Raw image

Compensated for tilt

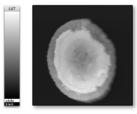
Line-by-line compensation

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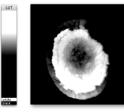
#### **Image Processing**

Image processing means changing all or some of the pixel values in an image, usually with the aim of making some feature(s) of the image more easily 'visible'.

The most trivial example would include changing the colour used to represent each pixel value — the look-up table (LUT).



default greyscale

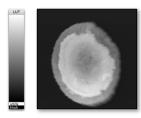


increased contrast

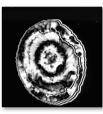
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#### **Image Processing**

The LUT does not have to be a linear, or even monotonic. A non-linear mapping between pixel value and displayed colour can often reveal unexpected detail in the image.







default greyscale

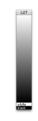
zebra greyscale

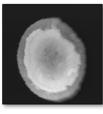
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#### **Image Processing**

Changing the LUT is reversible, as it is only the mapping between pixel values and display colours that is changed.

Taking a differential – replacing each pixel with the value of the local differential of the surface with respect to some direction - is irreversible in the sense that integrating doesn't (necessarily) get you your original image back.







greys → z values

greys  $\rightarrow \frac{\partial z}{\partial x}$  values

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#### **Kernel Filters**

Processing is often carried out using a kernel filter which uses an  $n \times n$  matrix of numbers. The kernel matrix is applied to every pixel in an image in turn.



1 1 1 1 1 1

1 1 1

Central value in matrix aligned with target pixel

The elements of the kernel matrix are multiplication values that are applied to a target pixel and its neighbouring pixels. The target pixel is replaced with the normalised sum of these products, and then the process is repeated for the next (overlapping) set of pixels.

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#### **Kernel Filters**

The simplest kernel filters use 3x3 matrices...

1 1 1

1 4 1 smooth 1 1 1

sharpen

gradient

-1 -1 -1

-1 9 -1 -1 -1 -1

-2 -1 0

-1 0 1

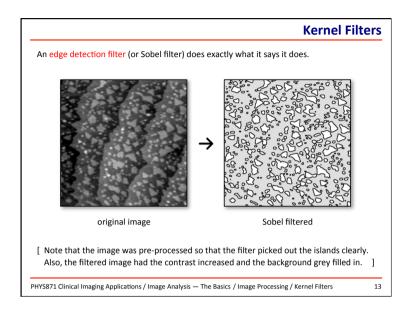
0 1 2

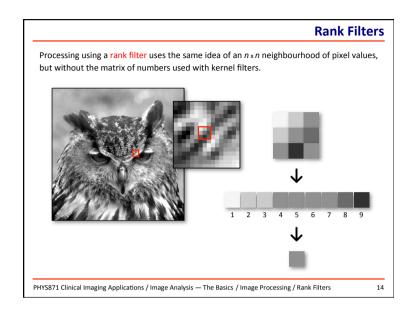


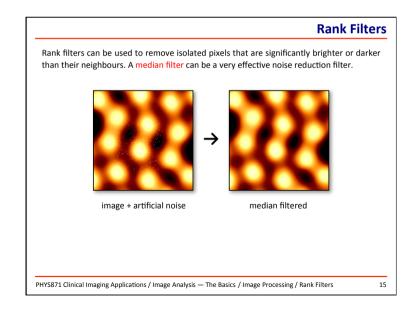


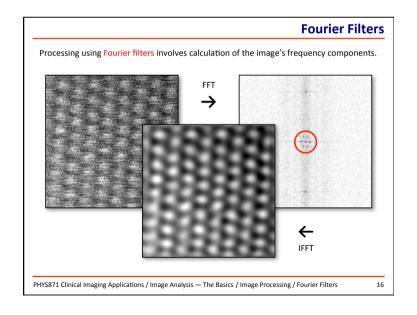


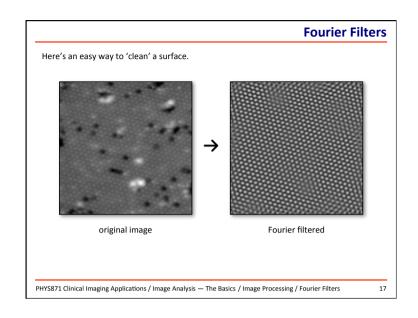
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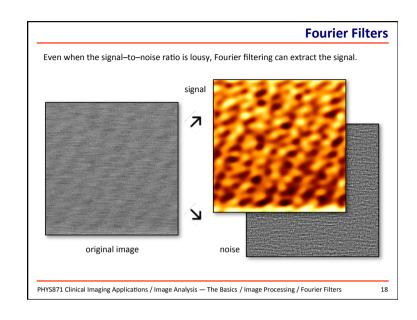


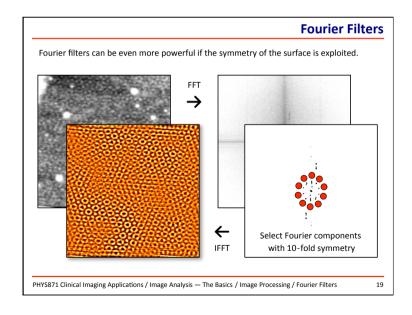


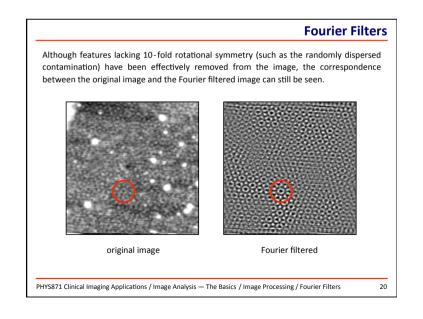




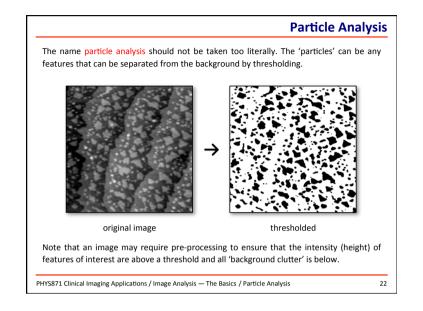


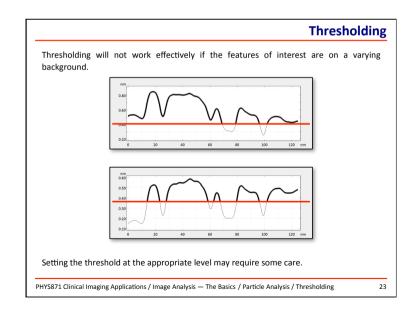


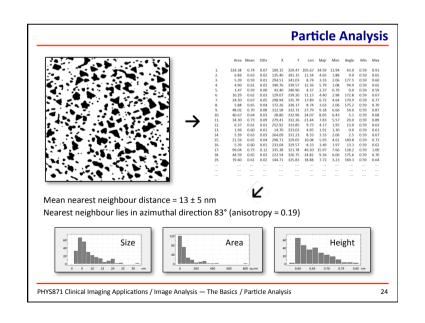




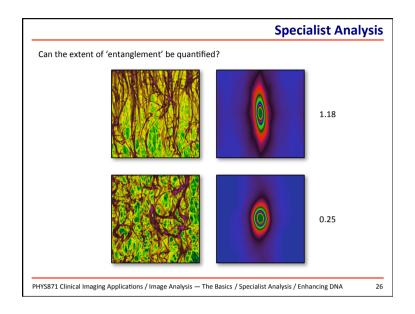
# Image Analysis Image analysis means extracting quantitative information that is derived from the pixel values in an image. Rather than being used as an intermediate step in image processing, the FFT can be a valuable source of quantitative information. Fourier transform strongest components The FFT maxima occur at a spatial frequency of 2.33 per μm (→ period = 0.43 μm).







# **Specialist Analysis** When "seeing the wood for the trees", or in this case the adsorbate for the substrate, computers can find the task much harder than an eye/brain combination. The property of the DNA strands that allows them to be separated from the background clutter is their curvature (the second differential of height wrt transverse distance).



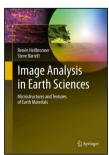
#### **Image Analysis**

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If you'd like to know more about the principles behind digital image processing and image analysis, or maybe you want to get your teeth into some of the maths, then try...

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The examples used have been drawn from the field of earth sciences rather than medical sciences, but the same principles apply.



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