# **X-Ray Computed Tomography (XRCT)**



- The recent development in image analysis techniques and software has provided the chance to use techniques such as XRCT in a wide range of applications.
- It can be utilised with particular software for characterising concrete microstructures including air pores (porosity) and aggregate distribution.
- XRCT is a non-destructive technique that can obtain cross-sectional images of the assessed sample and the computer processes the results, displaying them as 2D images.
- 3D images can be reconstructed using stacks of the 2D images within supplementary software.
- The scan is carried out by passing x-rays through a solid body/material while it is rotating where the scan starts from the top down to the bottom.
- The x-rays are detected after they have passed through the body and their intensity is measured, e.g. beams that have passed through less dense constituents are much less attenuated than those that have passed through high density solid matter.
- A computer is used to reconstruct this information where the relative density of the constituents is represented according to grey scale contrast on the image
- High density/ x-ray attenuation is represented by brighter pixels and low density by darker pixels.
- XRCT can be used for studying/evaluating concrete quality including the distribution, shape, orientation and size of key features such as aggregates and pores, as well as crack length and propagation



Example XRCT scan and sequence of image analysis techniques (Masad et al, 1999)

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## <u>As an example</u>

- A study was carried out in order to examine the effect of pre-coating/treating methods on the macro-scale (rounded) porosity caused by entrapment,
- XRCT was conducted using a Phoenix Nanotom 180NF XRCT system (see next Figure; GE Sensing and Inspection Technologies, GmbH, Wunsdorf, Germany) with a maximum electron acceleration energy of <u>110 kV</u>, and acquiring 1440 projection <u>images.</u>
- During the scan the detector size was set to provide a spatial resolution of <u>34.6 μm</u> and the scan length was approximately 45 min per sample.
- Images were reconstructed, after each set of projections, using the back projection algorithm in the 'datos | x rec' software.
- No corrections were required for beam hardening or sample displacement artefacts. All post-scanning image analysis was conducted using ImageJ software v1.44p.
- The tested sample was a 30 mm diameter cylinder cored from a >28-day-old 100 mm before being wrapped in a nylon sheet prior to scanning.
- Infranview image software was used to initially convert all X-ray images to 8-bit greyscale before scaling to 34.6 μm resolution.
- In order to eliminate any potential edge effects, only the middle 540 images in the stack were taken and the remaining 172 images at the top and bottom were removed.
- Number of process were carried out based on ImageJ software v1.44p







X-Ray Computed Tomography (XRCT) Phoenix Nanotom 180NF XRCT system (A) threshold sample where R = rubber aggregate, N = natural aggregate, A = air voids and C = cement paste. (B) Binary thresholded image and (C) Corresponding 3D reconstruction where yellow represents air voids and blue represents rubber particles.

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### Principles of SEM

- SEM is a microscopic imaging technique that uses a high energy electron beam to scan a specimen under vacuum.
- Electrons are accelerated down a column towards the sample at energies ranging from a few hundreds to tens of thousands of electron volts.
- When the beam leaves the column, it is deflected using scan coils into a scanning pattern that is focused to a spot on the sample
- SEM can use different analysis techniques involving the secondary electron (SE) detector, back-scattered electron (BSE) detector, and Energy Dispersive X-ray Spectroscopy (EDS).
- In SE detection mode, the SEM can provide very high-resolution images of sample surface morphology down to nano scale resolution.
- This is obtained by analysing the signals resulting from the interactions between the electron beam and atoms at or near the surface of the sample.
- SEM micrographs usually reveal the surface structure of the tested sample (surface morphology) as they have a large depth of field yielding a characteristic three-dimensional appearance due to the very narrow convergence angle of the electron beam.
- SE can provide a wide range of magnifications from 10 times to more than 5× 105 times



- Back-scattered electrons (BSE) are reflected from just beneath the sample surface by elastic scattering, and are commonly used in analytical SEM along with the spectra from EDS analysis.
- BSE micrographs can provide information about the distribution of different elements in the sample due to the strong relation between the intensity of the BSE signal and the atomic number of the specimen, i.e. high atomic number samples produce higher contrast/appearance up to 5-10 nm.
- Characteristic X-rays are emitted when the electron beam ejects an inner shell electron from the sample, causing a higher-energy electron to fill the shell and release residual energy in the form of photons at x-ray wavelength.
- The energy of these characteristic X-rays are used to identify the elemental composition and their spatial distribution across a flat (polished) sample surface

Schematic diagram of an SEM (Suzuki, 2002)





#### **Scanning process and image formation**

- Typical electron guns are usually fitted with a W (Tungsten) filament or LaB6 (Lanthanum Hexaboride) cathode.
- Tungsten is often used in thermionic electron guns because it has the highest melting point and lowest vapour pressure of all metals, thereby allowing it to be heated for electron emission, and modest cost.
- Other types of electron emitters include Field Emission Guns (FEG), which may be of the coldcathode type using tungsten single crystal emitters or the thermally assisted Schottky type.
- The electron gun/FEG, that produce the electron beam as shown in the next Figure, typically operate at an accelerating voltage of between 0.2 kV to 40 kV.
- The beam is focused by one or two condenser lenses to a spot ranging between 0.4 nm to 5 nm in diameter.
- The scan is carried out by passing the beam through pairs of scanning coils or pairs of deflector plates in the electron column, typically in the final lens, which deflect the beam in the x and y axes so that it scans in a raster fashion over a rectangular area of the sample surface.
- Once the electron beam interacts with the sample, the electrons lose energy by repeated random scattering and absorption within a teardrop-shaped volume of the specimen known as the interaction volume, which extends from less than 100 nm to around 5 µm into the surface. The size of the interaction volume depends on the electron's landing energy, the atomic number of the specimen and the specimen's density.



- The energy exchange between the electron beam and the sample results in the reflection of high-energy electrons by elastic scattering, emission of secondary electrons by inelastic scattering and the emission of electromagnetic radiation, each of which can be detected by specialized detectors.
- The beam current absorbed by the specimen can also be detected and used to create images of the distribution of specimen current.
- Electronic amplifiers of various types are used to amplify the signals, which are displayed as variations in brightness on a computer monitor Each pixel of computer video-memory is synchronised with the position of the beam on the specimen in the microscope, and the resulting image is therefore a distribution map of the intensity of the signal being emitted from the scanned area of the specimen.

### As an Example

- Pre-coated/Treated methodologies on the microstructure of PRC and SCRC including the ITZ micro-porosity and other features, and the bonding characteristics between the rubber particles and the cement paste, a microstructural analysis was carried out.
- For this purpose, a Philips XL30 Environmental SEM (Figure 3 28) with Field Emission Gun (ESEM-FEG) equipped SEM was used along with an Oxford Instruments Inca model Energy Dispersive X-Ray spectrometer (EDS) with 133-eV resolution of the Mn Kα peak at Full Width Half Maximum (FWHM).



• The obtained micrographs were recorded using an Everhart-Thornley type Secondary Electron (SE) detector and a Back Scattered Electron (BSE) detector supplied by K. E. Developments used in high vacuum (hivac) SEM mode.



Philips XL30 Environmental SEM

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#### As an Example

- The scanned samples were taken from the end-third of failed flexural test prisms from the same positions in order to reduce the possibility of post-yield damage and to be representative to a high extent.
- Samples were prepared by fracture cross section followed by vacuum cold-mounting in epoxy resin.
- The cross sectional surface was then polished with wet SiC polishing wheels (P240, P400, P800, P1200) before washing with industrial methylated spirit (IMS), followed by further polishing with 6 μm and then 1 μm diamond paste.
- Polished samples were Pt sputter coated in order to provide high-resolution imaging whilst still allowing light element EDS analysis, as shown below

Scanned cross-section resin-mounted ITZ samples







- The ImageJ software tool v1.44p was used to analyse five representative micrographs that were randomly taken for each sample in order to statistically quantify the width of interfacial gap void formation (due to de-bonding).
- Ten measurements were recorded for each image such that the mean interfacial gap void width is the average of 50 readings.
- The interfacial gap voids were identified by thresholding using the same thresholding algorithm as was applied for the XRCT analysis that previously mentioned.

# **Further reading**

- Khalid Najim, Matthew Hall 'Crumb rubber aggregate coatings/ pre-treatments and their effects on interfacial bonding, air entrapment and fracture toughness in self-compacting rubberised concrete (SCRC). Materials and Structures 46(12):2029-2043.
- Khalid Najim 'Determination and Enhancement of Mechanical and Thermo-physical Behaviour of Crumb Rubber-modified Structural Concrete. PhD thesis, The University of Nottingham, 2012