CHARACTERISATION OF CZT CRYSTALS FOR IMPROVED POSITION RESOLUTION AND DEFECT CORRECTION

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DETECTOR CHARACTERISATION

INTRODUCTION

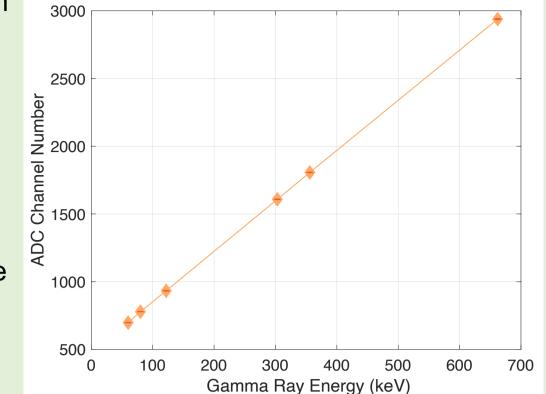
Due to its excellent gamma ray spectral performance across a broad energy range and its ability to be operational at room temperature, CdZnTe (CZT) is a seemingly ideal candidate for applications in nuclear security. Despite these obvious advantages, today's commercial detector-grade CZT does not demonstrate a high level of material homogeneity. Through extended defects such as Te inclusions, grain boundaries, dislocations and twins, the performance and production of CZT is severely limited [1].

The requirement for high-grade detector crystals in industry limits the size of crystals that can be grown, meaning that for a high detector efficiency, a multi-element array of smaller crystals is necessary. However, these crystals must endure a pre-selection testing process, which further escalates costs. So, in terms of CZT's current industrial applications, it seems that one must accept either poor performance or high costs.

DETECTOR LINEARITY

Sources with known y emission energies were used (²⁴¹Am – 60keV, ⁵⁷Co – 122keV, ¹³³Ba – 80, 302, 356keV, ¹³⁷Cs – 662keV) and matched to their ADC channel number.

It can be seen that the detector response is linear and therefore required no further correction.

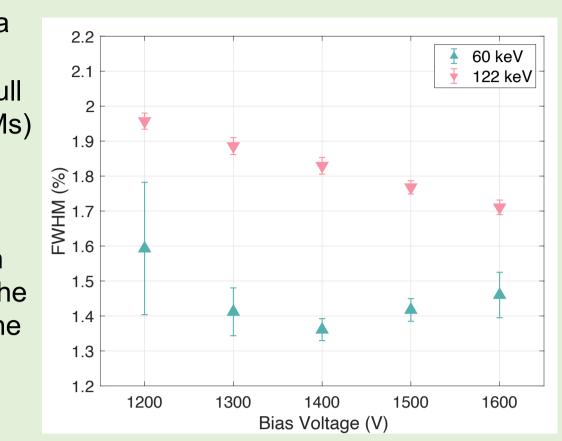


Detector characterisation measurements were performed with the ASIC readout to determine the optimal settings for further measurements across all five crystals.

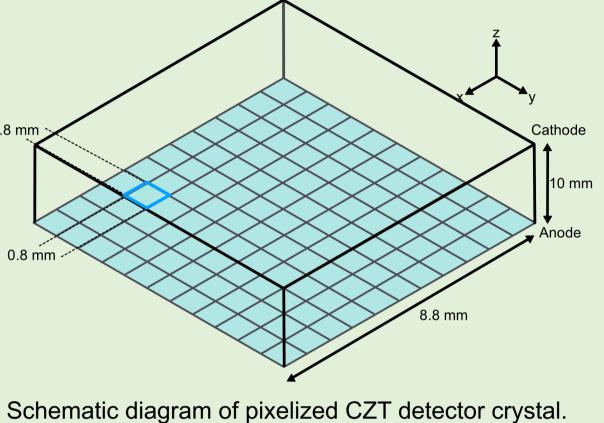
Each characterisation measurement was taken using Crystal 1. All five crystals have identical design and geometry, hence the selected characterisation parameters could be used across all further spectra captured.

ENERGY RESOLUTION AS A FUNCTION OF BIAS VOLTAGE

Spectra were captured over a range of bias voltages, photopeaks were fitted and full width half maximums (FWHMs) calculated to quantify energy resolution.

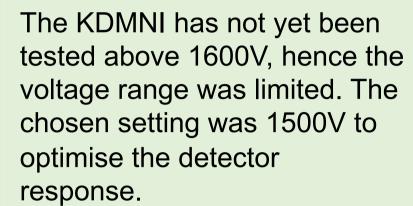


Five detector crystals of identical design and geometry were studied using precision collimated gamma scanning techniques. This allowed an investigation into the uniformity of response and areas of inhomogeneity within each crystal, as well as the identification of defects. Data taken and methods developed during this project could pave the way for the development of a detectorresponse correction, ultimately allowing for reduced CZT costs and increased efficiency.



Each crystal has a 8.8×8.8×10 mm³ volume; segmented into 11×11 pixels with 0.8 mm pitch.

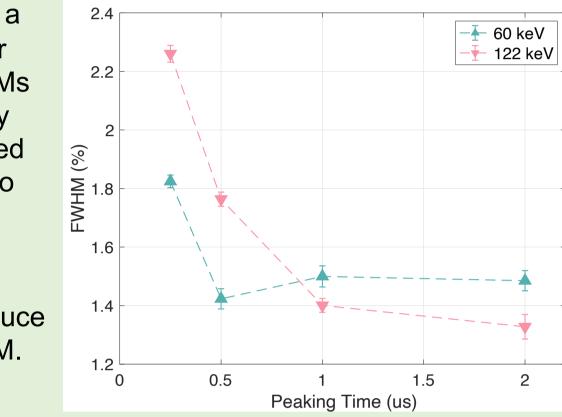
The Kromek DMatrix Nuclear Imager (KDMNI) is a fully integrated, photon counting, pixelated, 12-bit energy discriminating CZT detector consisting a CZT detector module coupled to ASICs (which provide energy and timing information), ADC converters, and an FPGA [2].



ENERGY RESOLUTION AS A FUNCTION OF PEAKING TIME

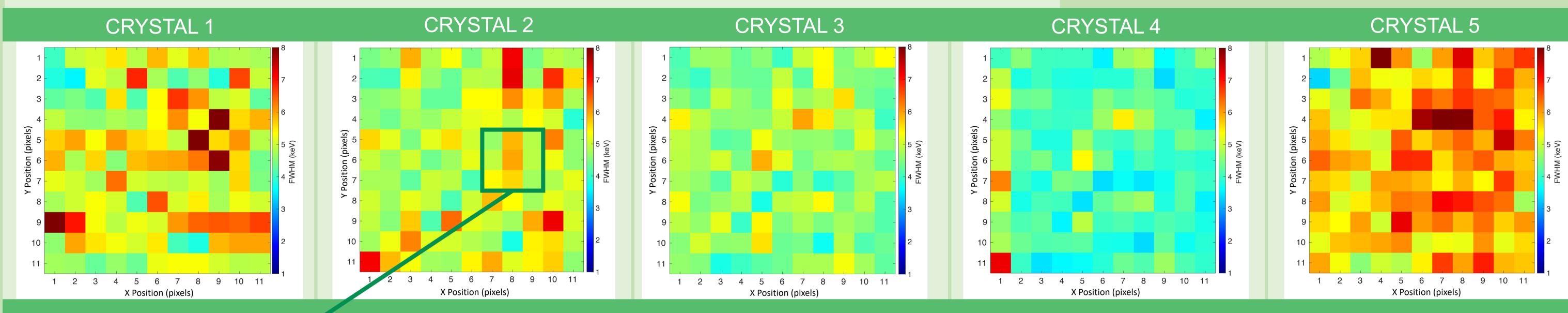
Spectra were captured over a range of peaking times, their photopeaks fitted and FWHMs calculated to quantify energy resolution. The KDMNI limited the range of peaking times to 2μ s and below.

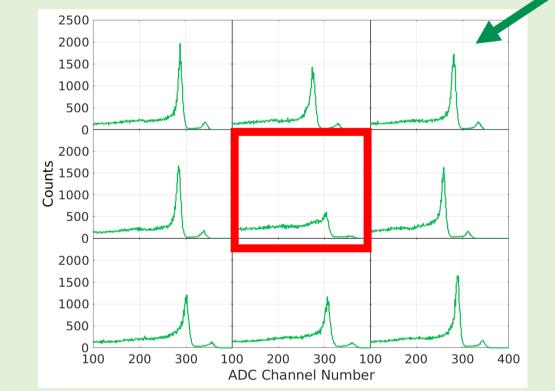
A peaking time of 2μ s was selected in order to best reduce ballistic deficit [2] and FWHM.



IDENTIFYING DEGRADED PIXELS AND CRYSTAL UNIFORMITY 3

A single detector crystal was placed into the Kromek DMatrix Nuclear Imager and spectra were captured with gamma sources of known emission energies (²⁴¹Am – 60keV, ⁵⁷Co – 122keV, ¹³⁷Cs – 662keV). The photopeaks were fitted and FWHMs were calculated, allowing one to plot FWHM uniformity maps of all 121 0.8x0.8 mm² pixels of the crystal. Using these, pixel spectra demonstrating low energy tailing and/or broadening could be identified for further investigation. Below are detector uniformity maps corresponding to ⁵⁷Co.



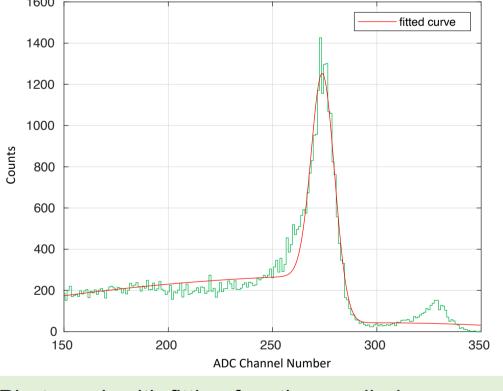


Pixel cluster 1 – a 3x3 cluster of raw pixel

spectra located on detector Crystal 2.

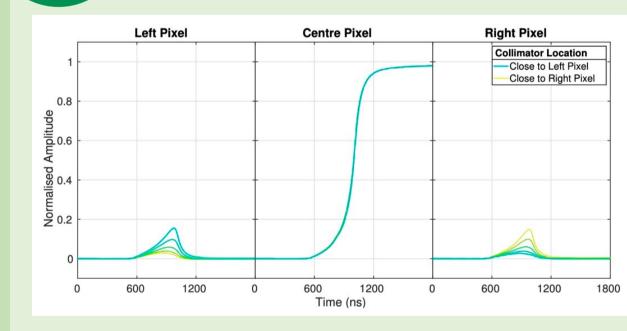
In comparison to its neighbours, the central pixel in Pixel Cluster 1 highlighted on the left suffers significant charge loss, resulting in low energy tailing and a less defined photopeak.

One cause of this could be a Te inclusion which obstructs the electron cloud transportation through the crystal, resulting in a charge loss proportional to the number of electrons crossing this defect's geometrical area [3].



Photopeak with fitting function applied.

PULSE SHAPE ANALYSIS



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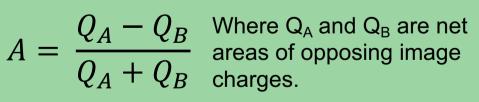
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Left: average charge collection signal and neighbouring transient image charges for 5 collimator positions across a central pixel [4].

Digital readout signals from the collimator system designed at the University of Liverpool allow the output of average signals across collimator position. Due to their position dependence,

the ratio between neighbouring image charge signals can be parameterised to give an indication of defect position.

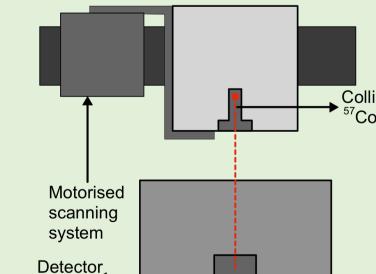
• These asymmetry parameters can then be used to determine the position of



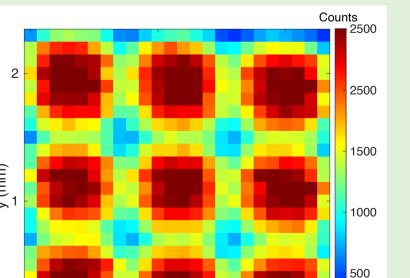
gamma interactions without the need for collimated position information. This methodology has been used to achieve sub-pixel position resolution [4].

Spectra were created and compared across the 0.8 mm pixel width based upon the known-collimator position and the PSA-determined position. Below, it's visible that these PSA approximations reflect collimated spectra well, and therefore these methods could be used to develop a defect correction method

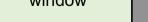
COLLIMATED SCANNING



The crystals were then separated from their ASIC stack and added into a digitizing system designed at the University of Liverpool, Collimated enabling signal readout of a 3x3 pixel cluster. Collimated gamma ⁷Co sou sources (²⁴¹Am, ⁵⁷Co, ¹³⁷Cs) were then attached to stepper motors (left) to gather charge collection and transient signals as a function of interaction position [4]. A 0.5 mm collimator was used to minimise beam divergence - a primary source of error in these

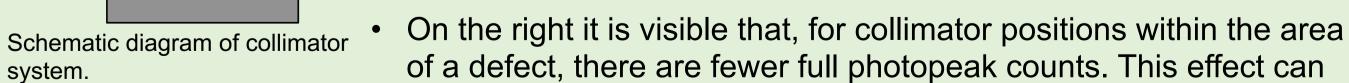


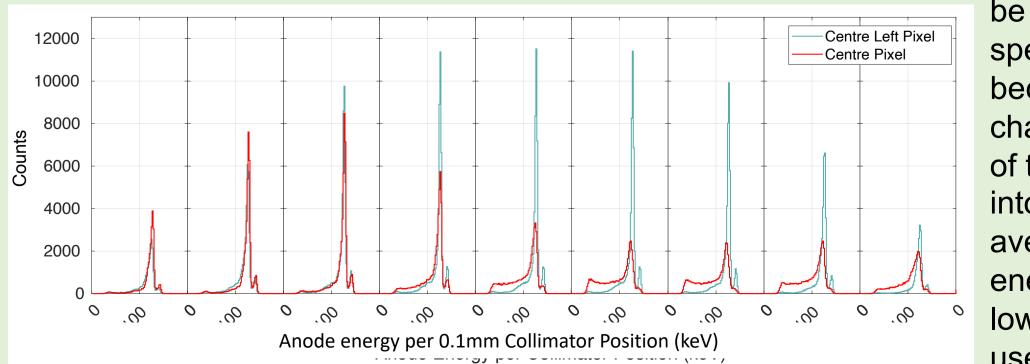
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system.

measurements.





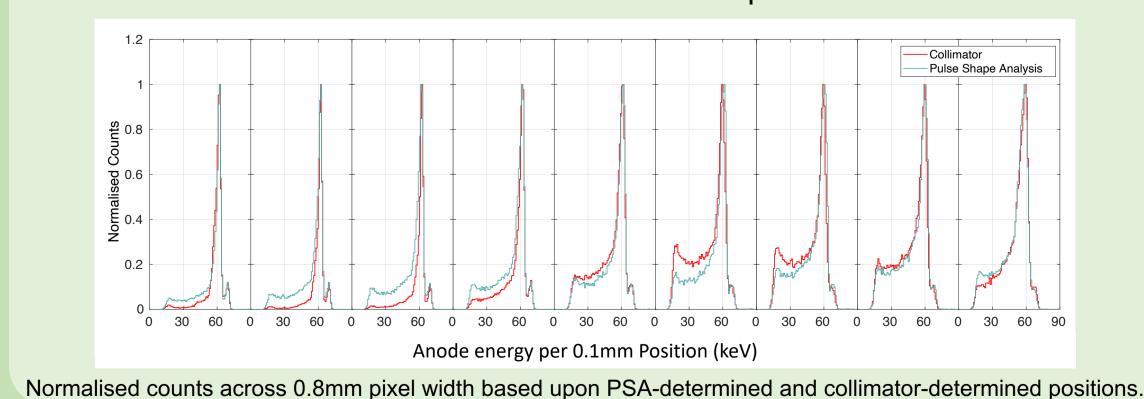
Comparison of energy spectra using ⁵⁷Co as a function of collimator position across the widths of the centre left and centre pixel in pixel cluster 1 located on crystal 2.

be studied further in the spectra below, where it becomes apparent that charge loss is the cause of this. Similarly, further into the defect, the average photopeak energy begins to shift

Comparison of typical cluster from lower – this effect can be crystal 4 (top) to degraded cluster from used in the development crystal 2 (bottom) intensity maps of a correction method. showing number of full photopeak events as a function of collimator position.

REFERENCES

[1] Bolotnikov, A., & al., 2010. Te Inclusions in CZT Detectors: New Method for Correcting their Adverse Effects. *IEEE Transactions on Nuclear Science*, April, 57(2), pp. 910-919. [2] McAreavey, L. & al., 2017. Characterisation of a CZT detector for dosimetry of molecular radiotherapy. *Journal of Instrumentation*, Volume 12. [3] Bolotnikov, A.E, & al., 2013. Characterisation and evaluation of extended defects in CZT crystals for gamma-ray detectors. Journal of Crystal Growth, Volume 379, pp. 46-56. [4] Rintoul, E., & al., 2022. Sub-Voxel Identification of Gamma-Ray Interaction Positions within a Pixelated CZT Detector through Signal Analysis. In Review.



FUTURE WORK

 Collimated side scans of the detector have been performed to quantify detector response as a function of depth of interaction. This will allow position resolution improvement and defect correction through the 10mm depth of the detector using rise time analysis.

Data has been taken for multiple degraded clusters at a range of energies to be used in the development and implementation of a universal defect correction method, improving the suitability of CZT detector crystals for nuclear security applications.