

Various Approaches for Charge R3A-28 Collection Efficiency Determination via Hecht Relation on CdTe/CdZnTe Detectors



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Pulse High Spectrum Analysis

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Introduction and Theory

High resistivity CdTe and CdZnTe semiconductors are very promising materials for room temperature X-ray and gamma-ray detectors with high charge collection efficiency (CCE) and mobility-lifetime (μτ) product [1]. The higher CCE and μτ parameters detector has the more usable material it is. One of the most widespread method for determination of $\mu\tau$ parametr is from collected charge-voltage dependency via single carrier Hecht equation [2]:

$$\mathbf{CCE} = \frac{\mu \tau V}{L^2} \left(1 - exp\left(\frac{L^2}{\mu \tau V}\right) \right)$$
 (H)

which can be set by alpha particle collected charge-voltage dependency. But high energy particles generate enormous amount of electron-hole pairs which leads to plasma effect – inner screening of internal field at the location of a particles absorption [3]. Due to this effect modified Hecht equation can be used [4]:

$$\mathbf{CCE} = \frac{\mu\tau(V+Vi)}{L^2} \left(1 - exp\left(\frac{L^2}{\mu\tau(V+Vi)}\right)\right)$$
(MH).

The other modification can be set due to surface recombination velocity:

$$\mathbf{CCE} = \frac{1}{1 + \frac{S}{\mu(\frac{V}{T} + \varepsilon_S)}} \frac{\mu \tau V}{L} \left(1 - exp\left(\frac{L^2}{\mu \tau V}\right) \right)$$
(SH),

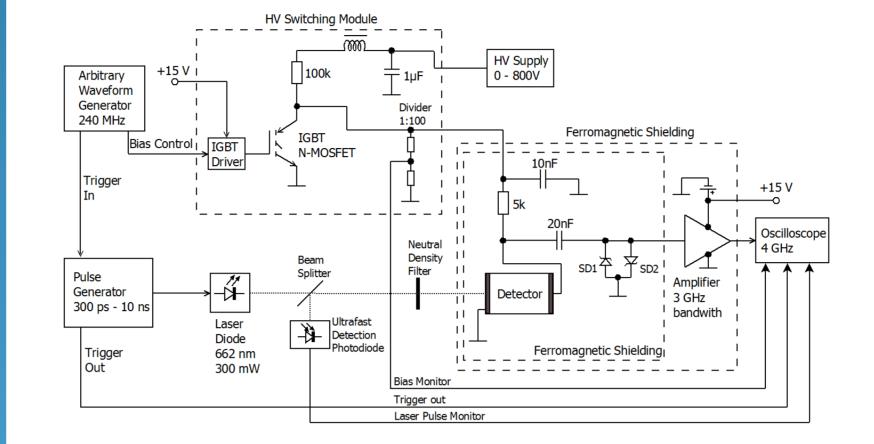
($\varepsilon_{\rm s}$ – characteristic contribution of band banding to a mean field at the surface)

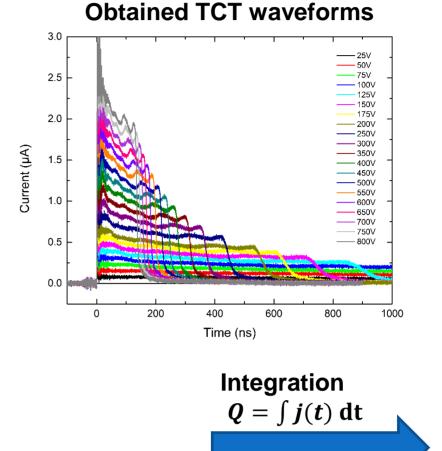
Experiment

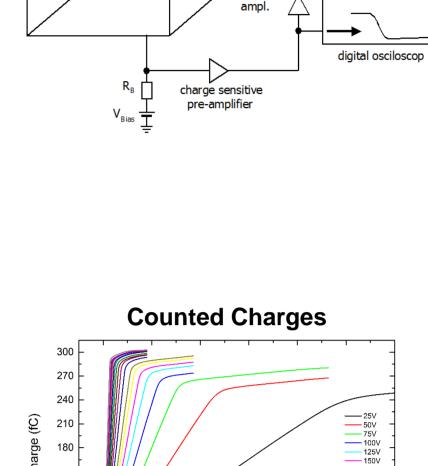
Five methods have been used to set collected charge-voltage dependency:

- ²⁴¹Am α (5,5 MeV) particle pulse height spectrum analysis (Alpha) [4]
- ²⁴¹Am γ (59,6 keV) and RTG (17,6 keV) pulse height spectrum analysis (Gamma), (RTG)
- Laser (660 nm) pulse height spectrum analysis (L-TChT)
 - Laser (660 nm) Transient Current Technique (L-TCT) [3]

Transient Current Technique (TCT)



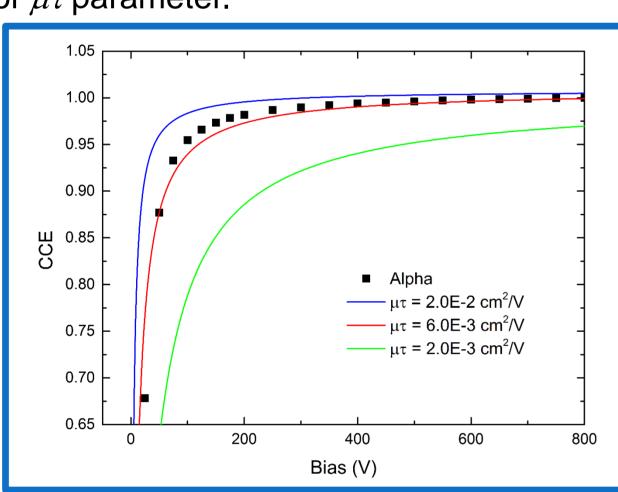




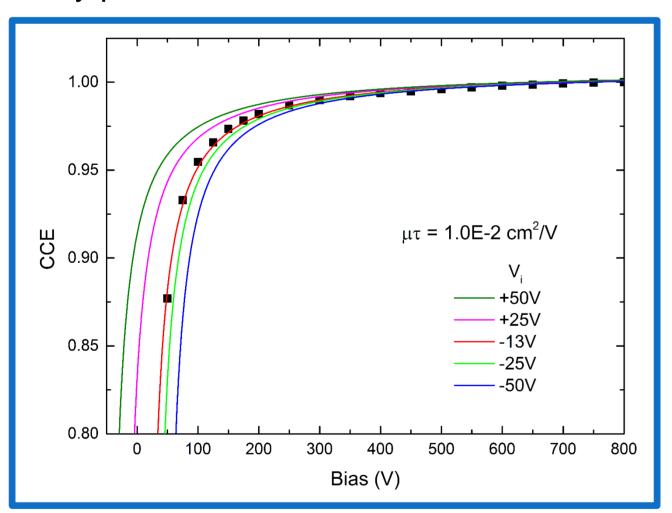
Results and Discussion

fitting Influence parameters obtained Alpha particle data of CdZnTe sample (common parameter L = 3,15 mm).

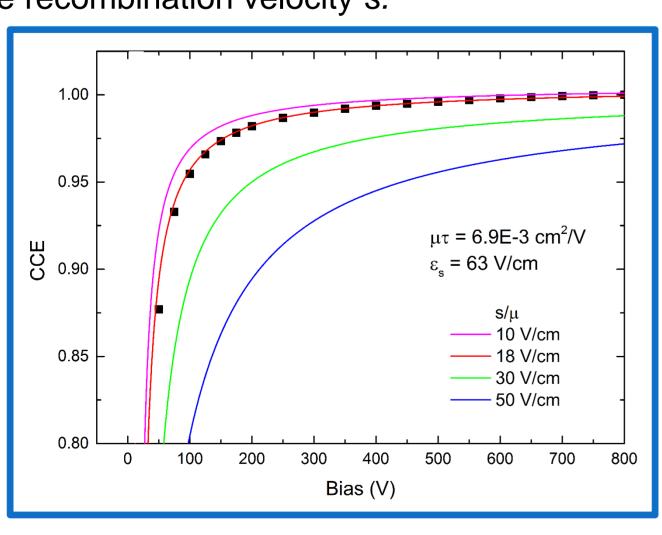
Single carrier Hecht equation (H). Influence of different values of $\mu\tau$ parameter.



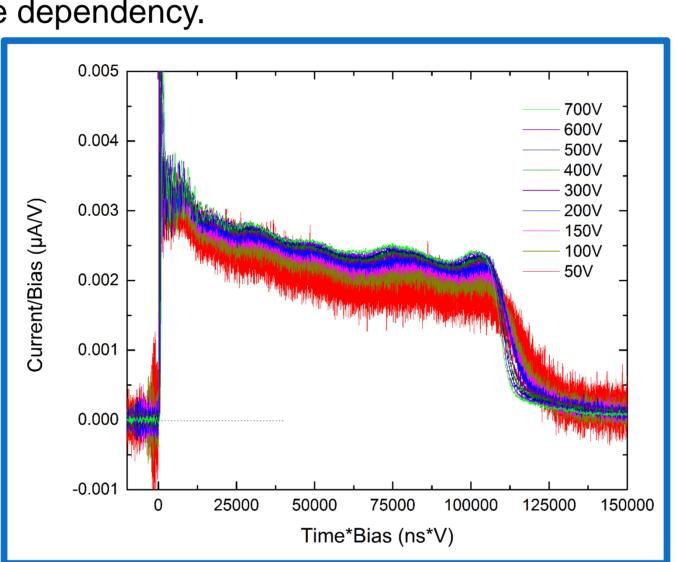
Modified Hecht equation (MH). Influence of different Vi produced by polarization field within the device.



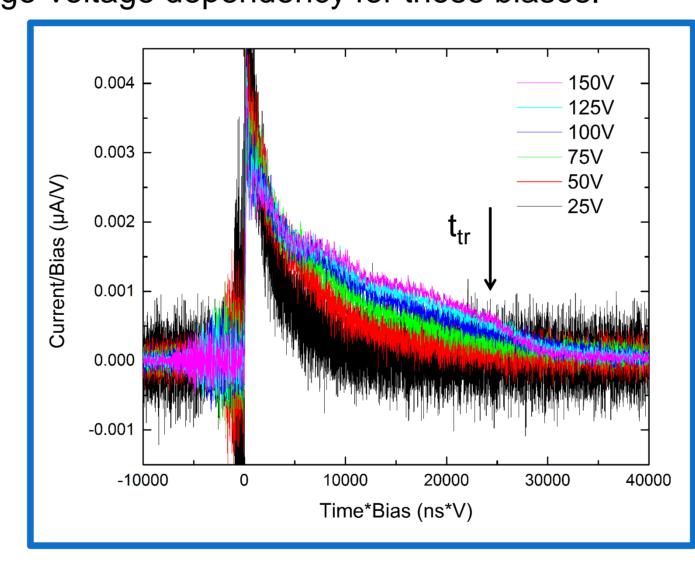
Surface Hecht equation (SH). Influence of different surface recombination velocity s.



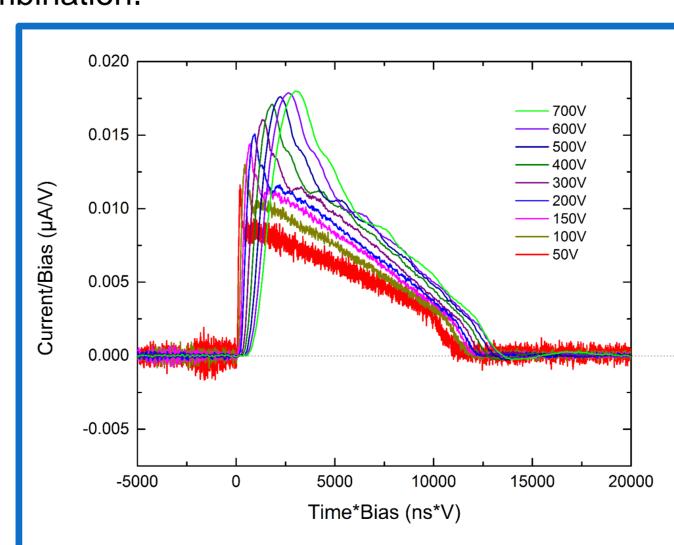
Normalized TCT waveforms of CdZnTe sample. Similar curves indicates a rapid saturation in collected chargevoltage dependency.



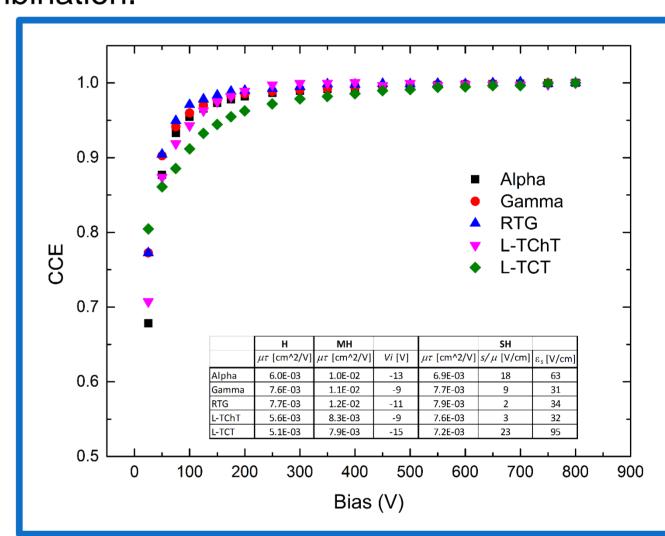
Normalized TCT waveforms of CdTe:In sample. Low voltages show the absence of transit time, thus we have inactive layer under the anode = low values in collected charge-voltage dependency for those biases.



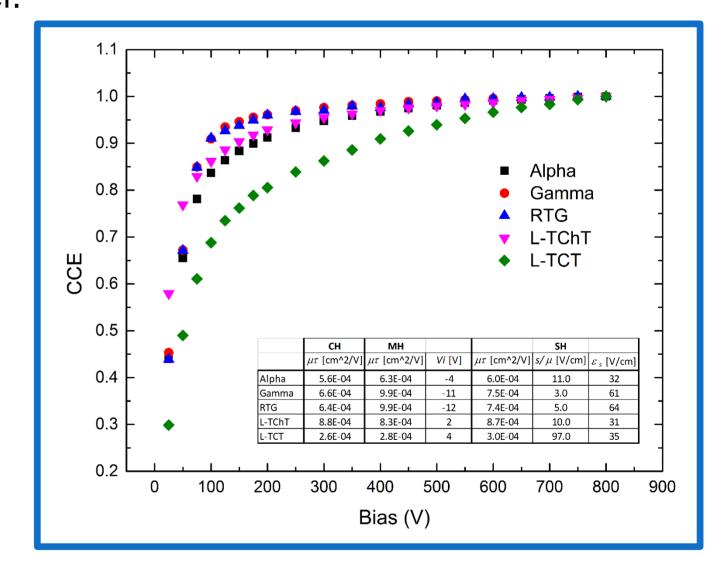
Normalized TCT waveforms of CdTe:Cl sample. The increasing character correspond to a large surface recombination.



Collected charge-voltage dependency of CdZnTe **sample.** High Vi in (MH) indicates wrong values of $\mu\tau$. We obtained good match in (CH) and (SH) for small surface recombination.

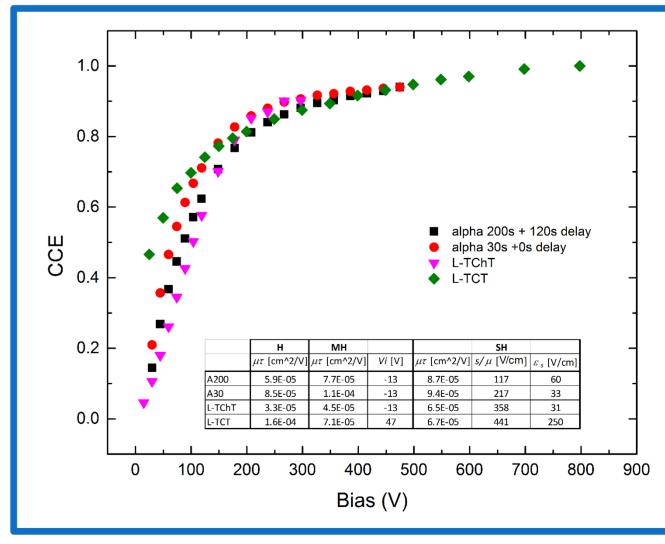


Collected charge-voltage dependency of CdTe:In sample. Low obtained values are distored by inactive layer.



Collected charge-voltage dependency of CdTe:Cl sample. Due to a similar resistance of the sample as the value of R_B, it is not possible to measure spectral analysis

to higher voltages.



CdTe:Cl L = 1 mm

CdZnTe

L = 3,15 mm

CdTe:In

L = 1,65 mm

Conclusion

- \Longrightarrow Modified surface recombination Hecht equation (SH) gives good values of $\mu\tau$ parameter, if we counted the influence of characteristic contribution of band banding to a mean field at the surface – $\varepsilon_{\rm s}$.
- ⇒ It is always good to complete spectral analysis with TCT measurements, because the samples with inactiv layer under the low voltages distors the obtained values of $\mu\tau$ parameter due to this effect.
- Normalized TCT waveforms show the influence of surface recombination.
- Simple modification of Hecht equation (MH) is not sufficient and the high values of *Vi* do not correspond the real state.

References

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