

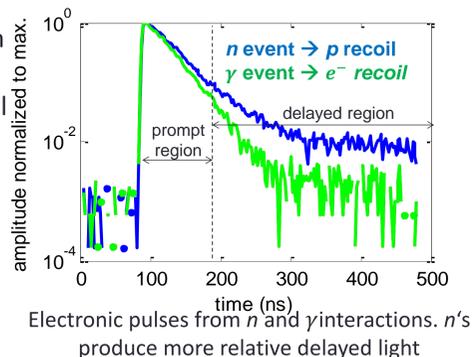
ABSTRACT

- Organic scintillator materials are becoming increasingly valuable as radiation detectors in nuclear safeguards and security. They offer simultaneous detection of fast neutrons and gamma-rays, provide information about the neutron energy, can be made in large sizes, and are relatively inexpensive.
- Organic crystal scintillators offer minimal thermal expansion compared to liquids and can be made in large sizes with excellent performance. However, a directional dependence that degrades the energy resolution has long been observed in heavy charged particle interactions in organic crystal scintillators.
- This directional dependence is generally seen as a problem, but may be an opportunity to learn more about the material properties in organic crystal scintillators or use them as directional neutron detectors.

ORGANIC SCINTILLATOR OUTPUT DEPENDS ON PARTICLE TYPE

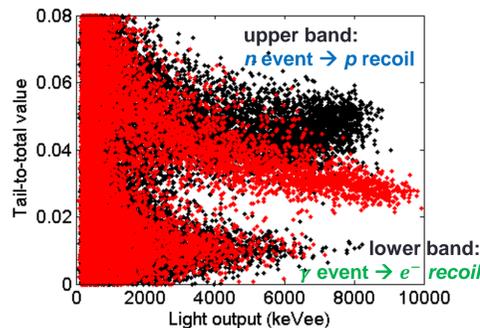
- Interactions by neutrons and gamma-rays in organic scintillators produce different distributions of light vs. time. Both the total light output per energy deposited and the shape of the time distribution differ.
- The pulse shape is quantified by measuring the fraction of delayed light, calculated as the tail-to-total (TTT) value:

$$TTT = \frac{\text{delayed light}}{\text{prompt} + \text{delayed light}}$$



CRYSTAL ORGANIC SCINTILLATOR OUTPUT ALSO DEPENDS ON HEAVY CHARGED PARTICLE DIRECTION

- In crystalline materials, the light output and pulse shape also depend on the direction of the heavy charged particle recoil with respect to the crystal axes.
- A measurement of events from a dT neutron generator at two different orientations (red and black data) produce different TTT vs. Light Output distributions as shown on right.

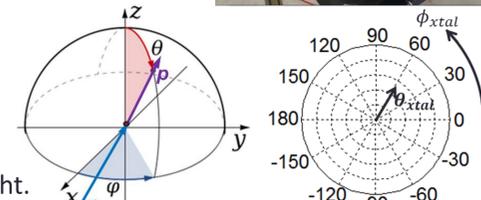


- The magnitude of change in the light output is greater at lower proton recoil energies, and the magnitude of change in the tail-to-total value is greater at higher proton recoil energies.
- While the light output changes by a similar factor in both materials, the tail-to-total value changes much more in anthracene than in stilbene.

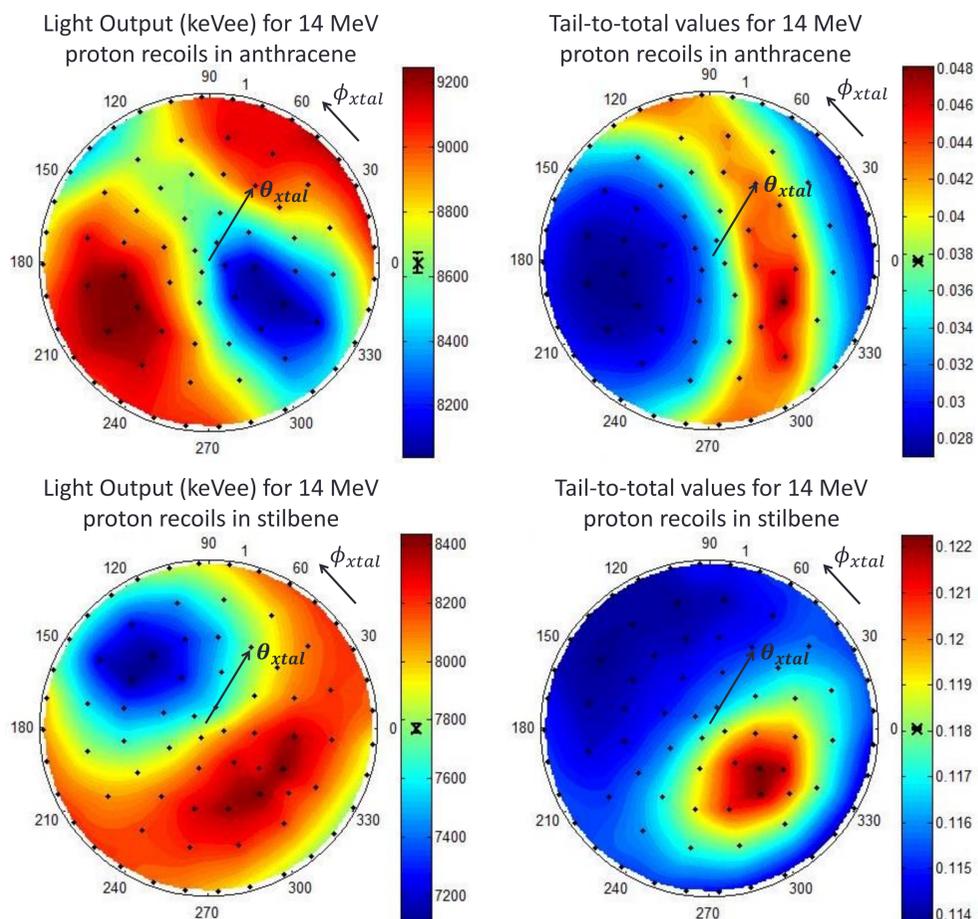
Anthracene		
E_p	L.O. Max/Min	TTT Max/Min
14 MeV	1.155 ± 0.006	1.798 ± 0.006
2.4 MeV	1.396 ± 0.036	1.317 ± 0.007
Stilbene		
	L.O. Max/Min	TTT Max/Min
14 MeV	1.191 ± 0.004	1.076 ± 0.001
2.4 MeV	1.320 ± 0.020	1.031 ± 0.003

CHARACTERIZING LIGHT OUTPUT AND PULSE SHAPE VS. PROTON RECOIL DIRECTION

- To control the proton recoil direction and energy, full energy interactions from monoenergetic neutrons are used in which the proton travels in a single known direction. The proton recoil direction is adjusted by rotating the detector with respect to the neutron source.
- A motor-driven rotational stage can position four detectors at all possible proton recoil directions in the crystal axes.
- Proton recoil directions are expressed in spherical coordinates as (ϕ, θ) in an arbitrary axes system, with a 2D representation for plotting purposes shown on right.



- The following figures show the light output and tail-to-total value as a function of proton recoil direction at 14 MeV in anthracene and stilbene.
- In anthracene, the direction of maximum light output corresponds to that of minimum tail-to-total value. In stilbene, that relationship is reversed.



CONCLUSION AND FUTURE WORK

- Measurements of proton recoils from neutron interactions have demonstrated that the light output and pulse shape depend on the direction of the proton recoil. The observed directional dependence is significant and complex in anthracene and stilbene.
- Future work aims to characterize additional materials of different crystal quality, $n-g$ separation, and with known crystal axes to better understand the mechanism that produces the effect.
- A deeper understanding may make it possible to correct for the effect in order to improve energy resolution, or produce new materials that eliminate or enhance the effect.
- These materials offer the potential for a compact directional neutron detector. A hand-carryable, relatively robust system could be constructed with a single detector whose material properties provide internal directionality.

ACKNOWLEDGEMENTS

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