



MUSiC: Reactivity Estimates Using Pulsed Neutron Source Methods

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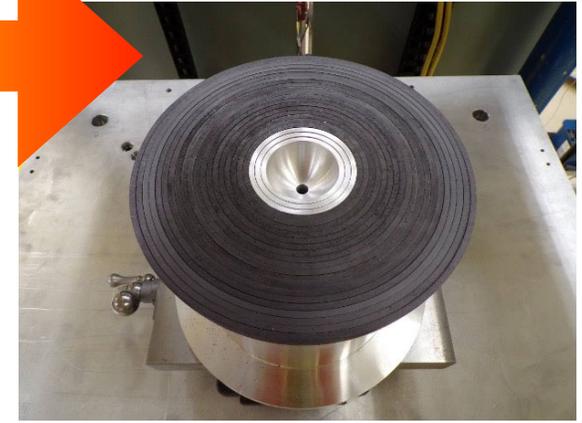
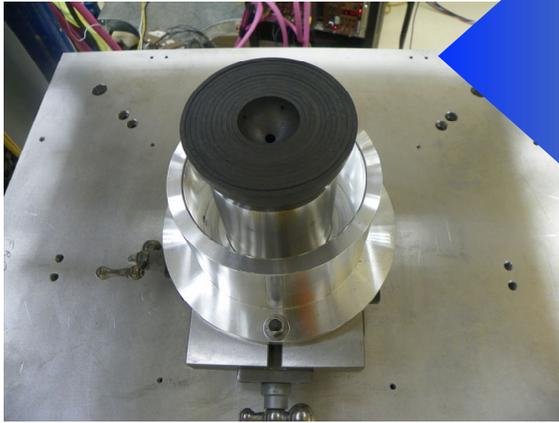
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The MUSiC experiment covers a broad range of subcritical to critical configurations

Deeply Subcritical
 k_{eff} 0.64 - 0.8

Nearly Delayed Critical
 k_{eff} 0.98 - 0.99

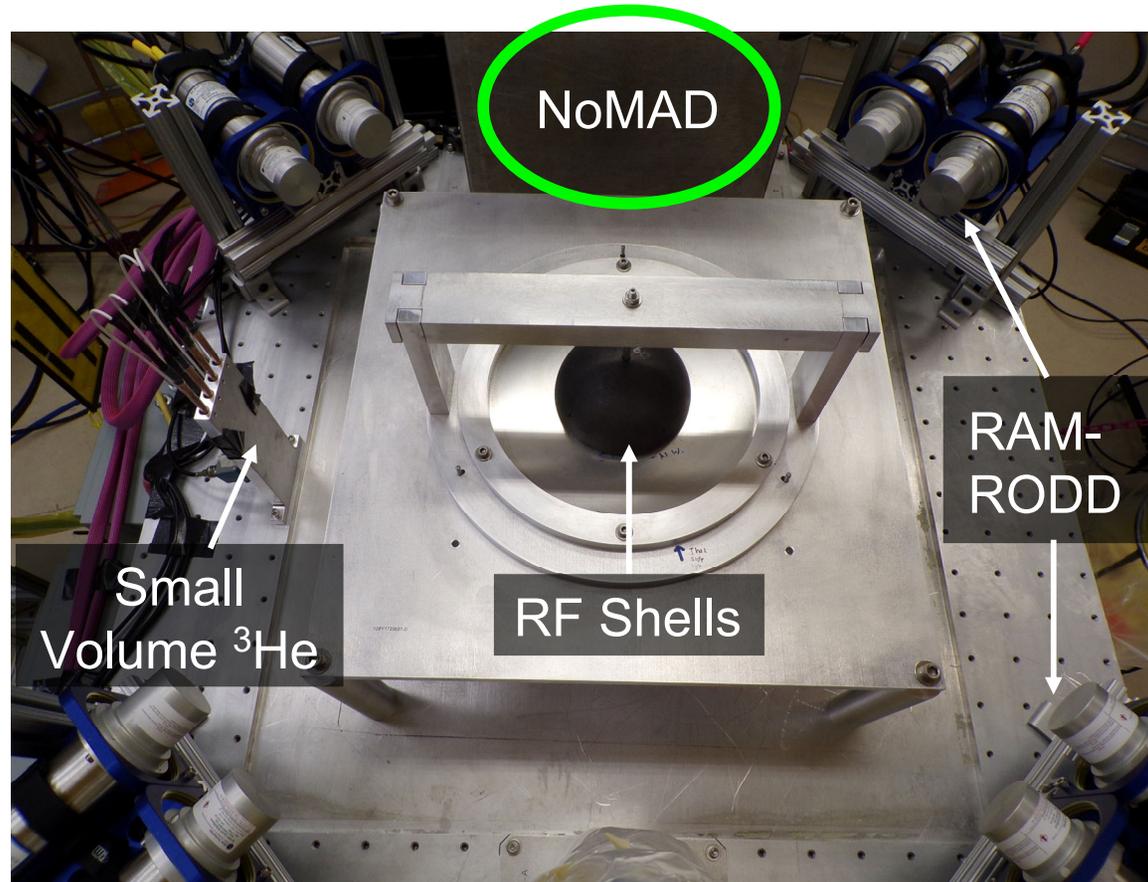


Many Possible Configurations

- Rocky Flats HEU shells can be configured for a variety of masses and reactivities
- Valuable test bed for measurement methods and instrumentation
- Platform for validated subcritical nuclear data benchmarks

One primary goal of MUSiC is to evaluate and compare subcritical system measurement methods

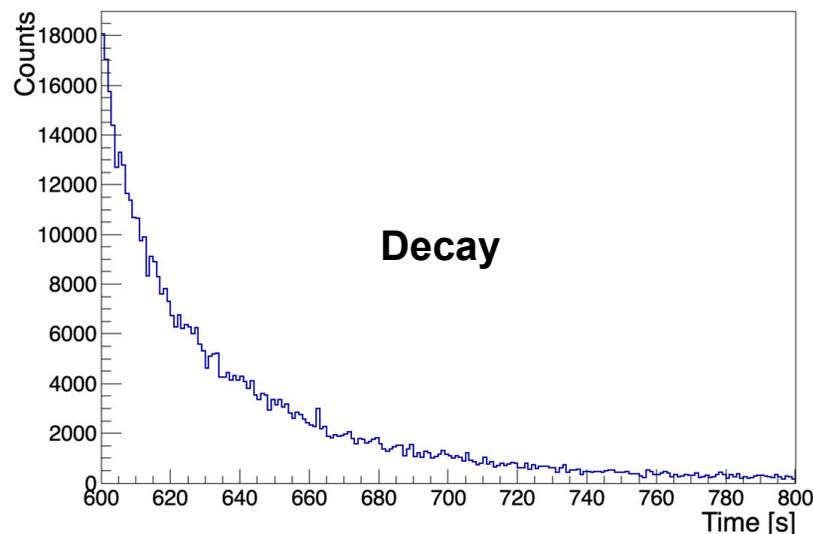
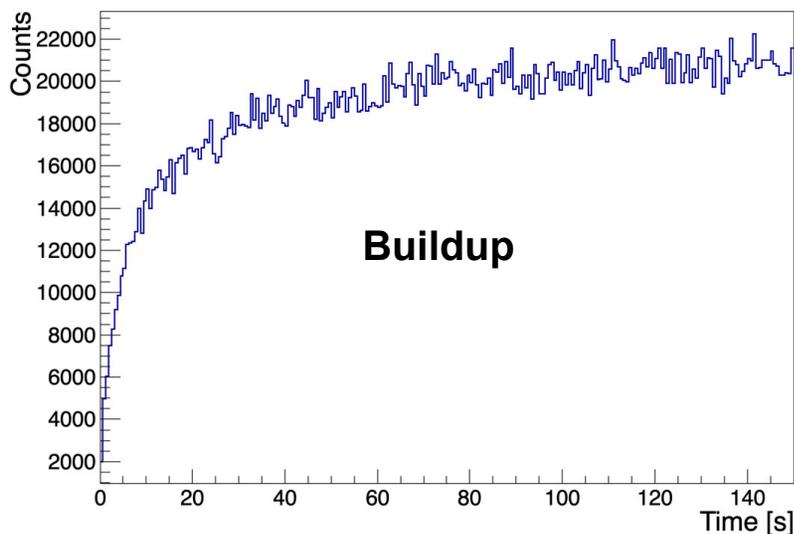
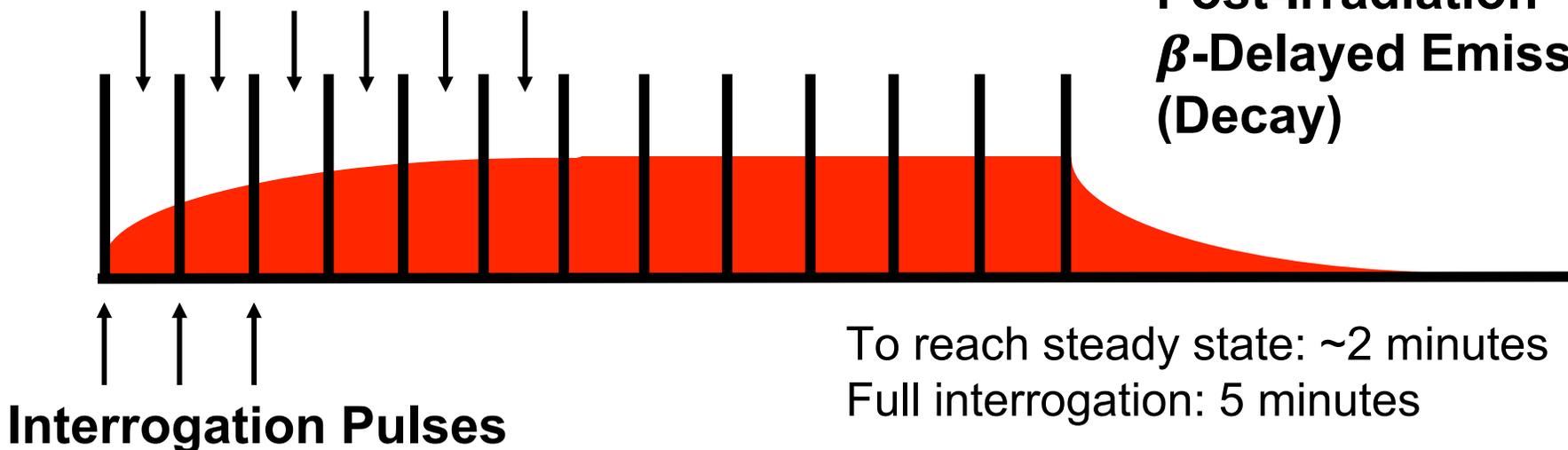
- Instrumentation:
 - ^3He detector systems
 - Scintillators
- Broad range of reactivities
- Multiple interrogation sources
- Rich data for analysis methods and parameter studies
- Consistent, repeatable configurations
- All methods are subject to bias → cross-compare and adjust experiments to account for it
- Preliminary results will guide future experimental improvements
- **Focus of this presentation: pulsed neutron method for reactivity analysis, using DT interrogation source and NoMAD detector**



Pulsed neutron active interrogation can reveal characteristic signatures in the time domain

Interpulse β -Delayed Emission (Buildup)

Post-Irradiation β -Delayed Emission (Decay)



The reactivity of a subcritical system can be estimated using the pulsed neutron method

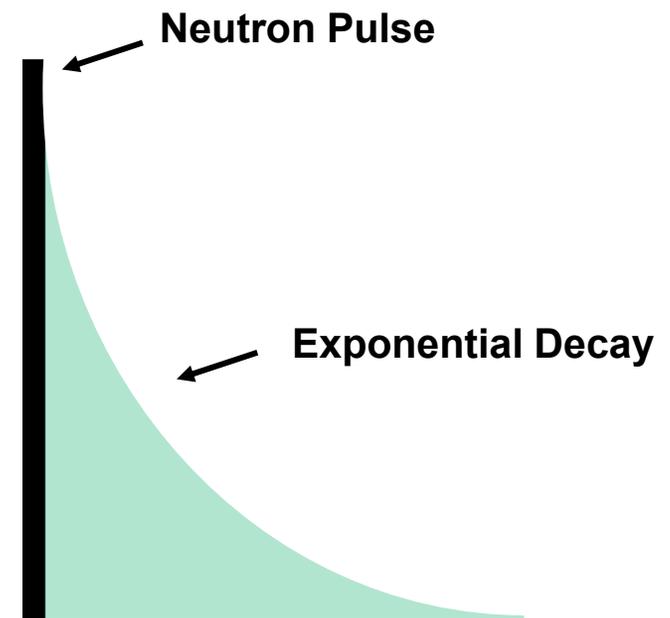
Basic method: a burst of neutrons is injected into a subcritical multiplying system and the decay of the neutron population is observed over time

The measured fundamental decay constant of the system, α_0 , and the decay constant at delayed critical, α_{DC} , are related to the reactivity (ρ) by

$$\alpha_0 = \alpha_{DC} (\rho - 1)$$

Thus, the reactivity is directly proportional to α_0 , and can be obtained by measurement of the decay constant at delayed critical.

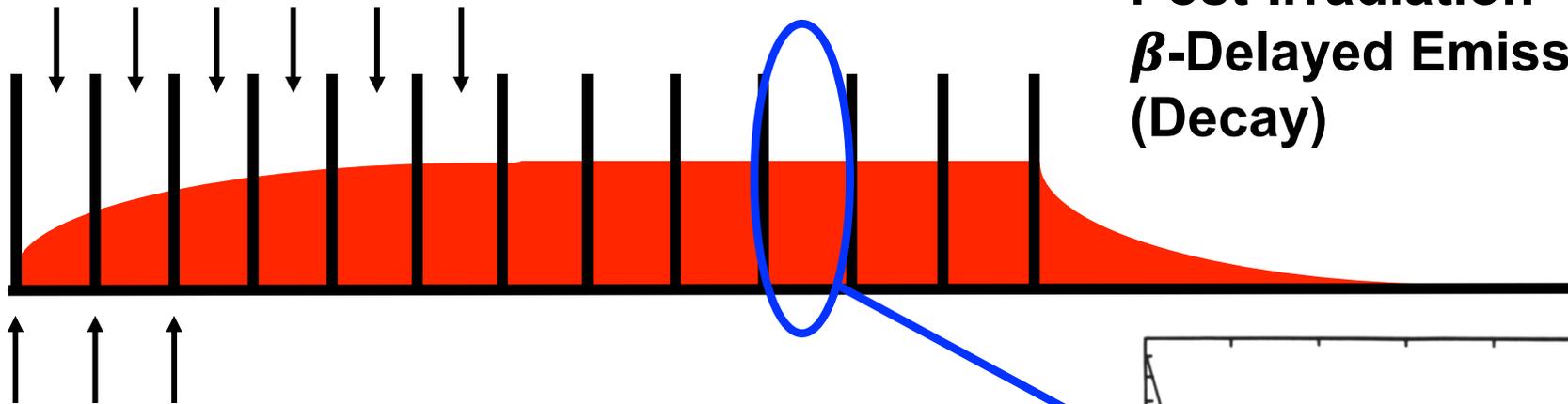
Measuring α_{DC} can be difficult for subcritical systems.



G. Keepin, *Physics of Nuclear Kinetics* (Addison-Wesley, 1965), 273-277

The reactivity of a subcritical system can be estimated using the pulsed neutron method

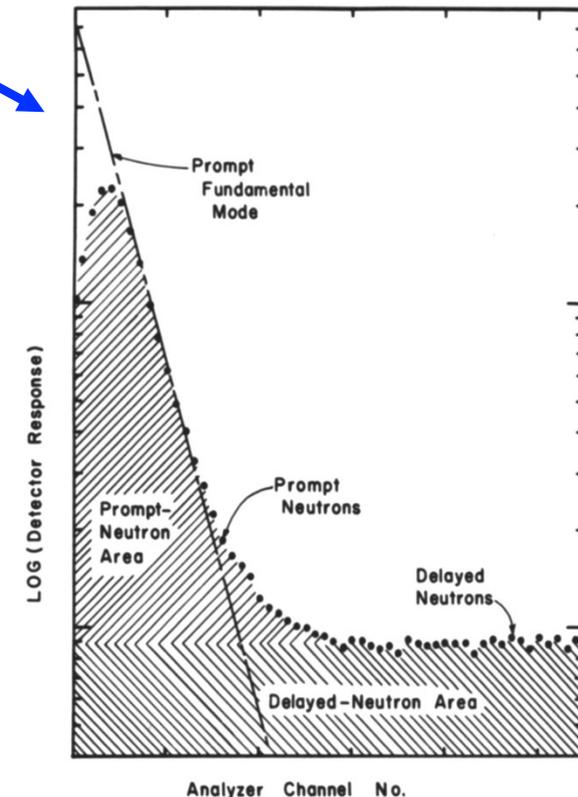
Interpulse β -Delayed Emission
(Buildup)



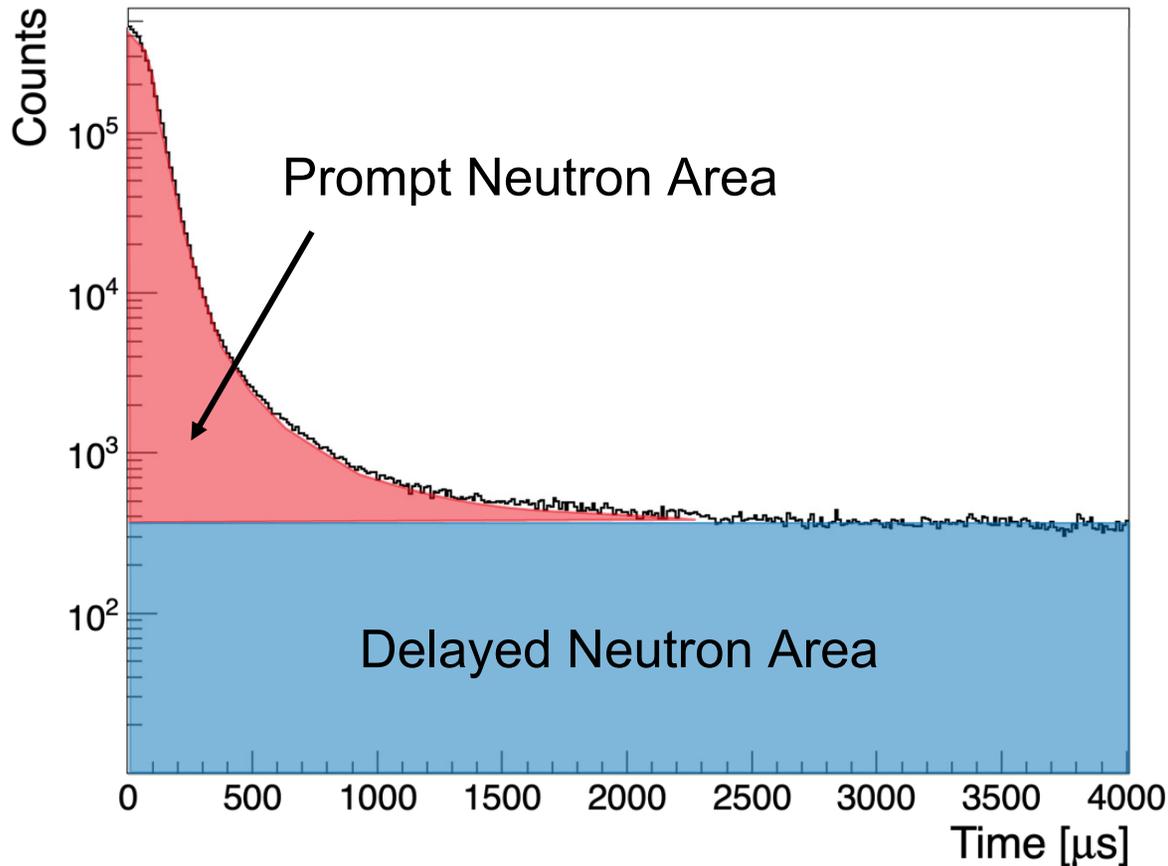
Post-Irradiation
 β -Delayed Emission
(Decay)

Interrogation Pulses

- For subcritical systems, it may not be feasible to directly measure α_{DC}
- The need to measure α_{DC} can be circumvented by measuring pulsed response once delayed neutron equilibrium is reached
- This is possible when the pulse rate R satisfies $\alpha_0 \gg R \gg \lambda$



The Sjöstrand Method uses a simple area ratio to estimate reactivity

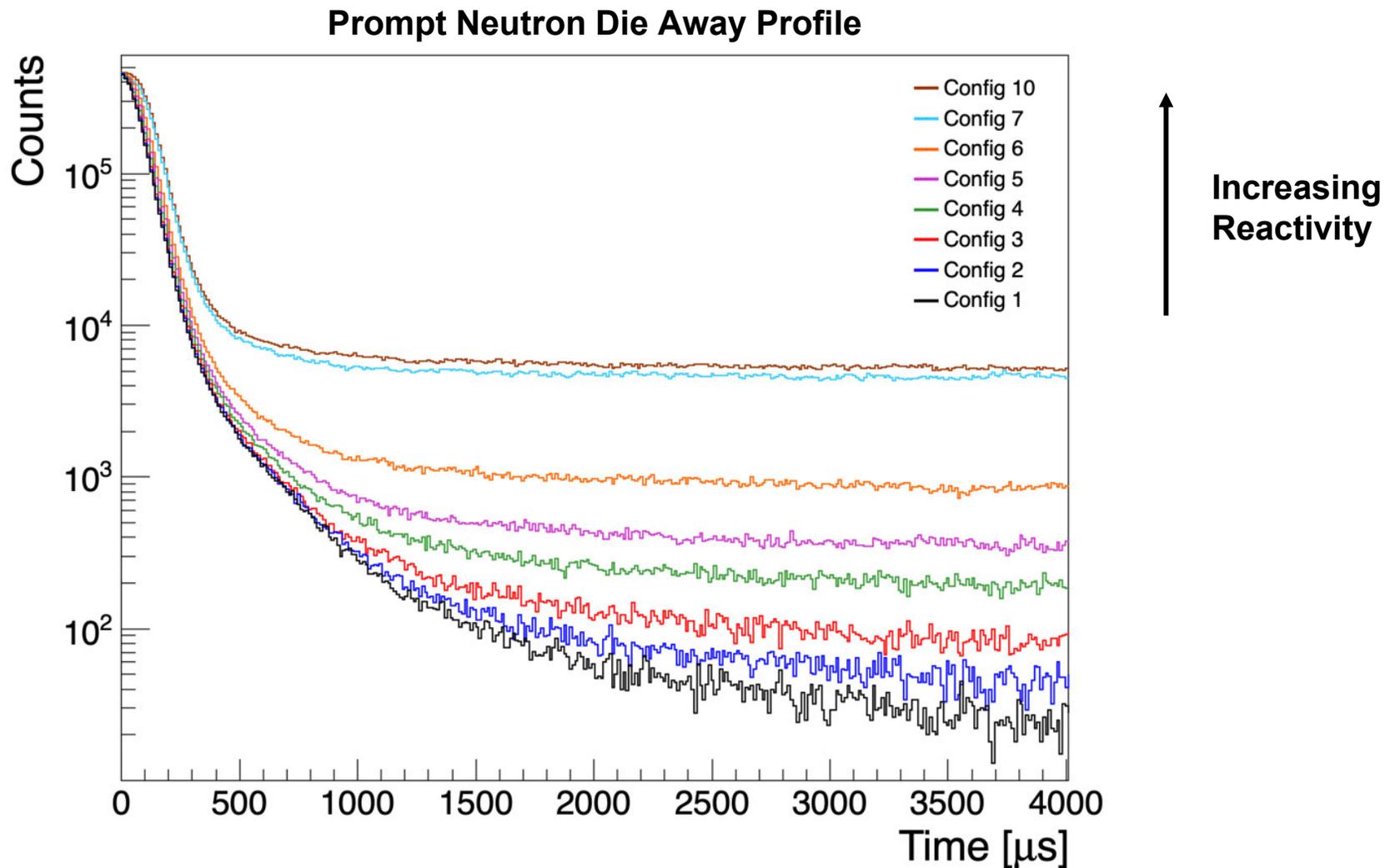


$$\rho_{SJ} \equiv \frac{\text{Prompt Neutron Area}}{\text{Delayed Neutron Area}}$$

- **Sjöstrand method¹**
 - Simplest method to apply
 - Reactivity estimated by ratio of prompt/delayed neutron areas
 - Assumes point kinetic model (ignores spatial harmonics)
- **More complex analysis methods are also being pursued**
 - Account for some spatial effects
 - Involve fitting prompt decay modes, alternative calculations of characteristic constants

¹N. G. Sjöstrand, *Arkiv Fysik*, 11, 233, 1956

Pulsed neutron method preliminary results

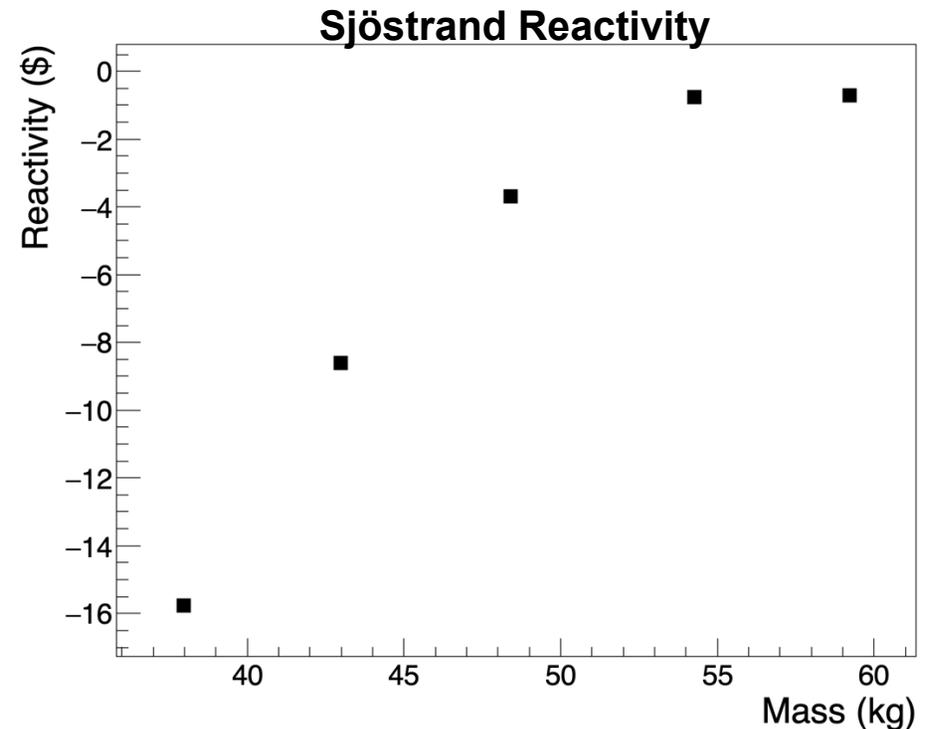


- Proportion of delayed neutron area increases with increasing reactivity
- Smaller prompt/delayed ratio \rightarrow closer to critical

Pulsed neutron method preliminary results

Configuration	Mass (kg)	Multiplication	Feynman- α Reactivity (\$)	Sjöstrand Reactivity (\$)	Sjöstrand k_{eff}
4	37.9617	8.3	-22.71	-15.76	0.907
5	42.9722	11.4	-16.90	-8.60	0.947
6	48.4099	18.4	-10.26	-3.68	0.977
7	54.2785	46.6	-4.81	-0.764	0.995
10	59.2075	>150	N/A	-0.713	0.995

- Pulsed (Sjöstrand) results higher than expected
- Feynman- α results somewhat lower than expected
- Sources of significant systematic error exist (dead time, integration windows), yet to be fully quantified



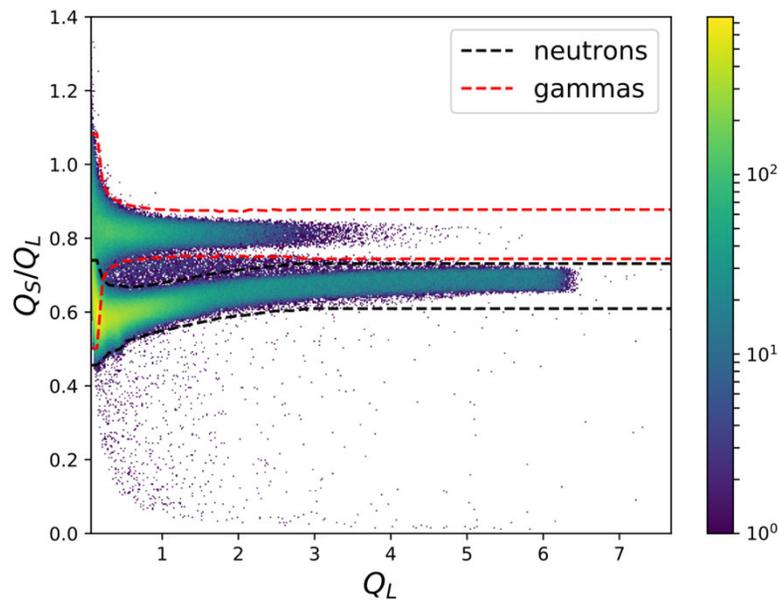
Separate Effort:

Organic scintillators for time correlated measurements

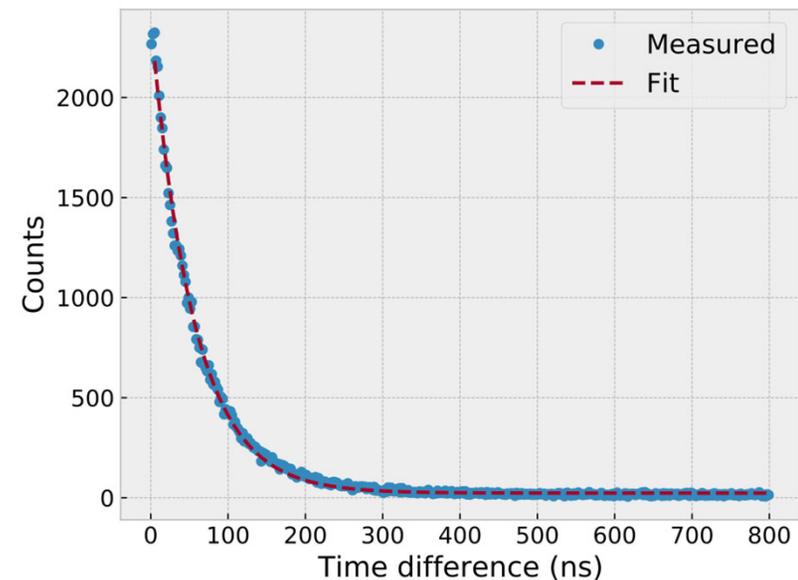
- We are also interested in fast timing applications
- ^3He systems like NoMAD have inherent time resolution limits
- R&D of organic scintillator applications led by Rob Weldon

Organic scintillators for time correlated measurements

- Organic liquid scintillators
- Direct detection of fast neutrons (recoil based)
- No moderation, fast response → ideal for bare systems
- Time resolution (~ 900 ps)
- Excellent for neutron noise measurements of fast systems
- Measurements: Rossi- α , Feynman- α
- Pulse shape discrimination – eliminate gamma signal
- Reactivity range: all configurations
- Neutron energy spectroscopy possible



Rossi- α Measurement



Conclusion & Future Work

Accomplishments

- Analysis of pulsed active interrogation data has begun
- Development of reactivity analysis based on pulsed neutron methods
- Preliminary results show expected trends
- Fast-timing analysis using organic scintillators

Next Steps

- Refinement of pulsed neutron method algorithms
- Analysis of sources of systemic bias
- Comparison with other reactivity estimation methods

Questions?

Calculating Reactivity of a Subcritical System Using the Pulsed Neutron Method

Basic method: a burst of neutrons is injected into a subcritical multiplying system and the decay of the neutron population is observed over time

The fundamental-mode decay constant for prompt neutrons, α_0 , is given by

$$\alpha_0 \equiv \frac{1}{n} \frac{dn}{dt} = \frac{k(1 - \beta_{\text{eff}}) - 1}{l}$$

where l is the prompt neutron lifetime and β_{eff} is the effective delayed neutron factor.

Using the definition of reactivity $\$ \equiv \frac{\rho}{\beta_{\text{eff}}} = \frac{k - 1}{k\beta_{\text{eff}}}$, α_0 is related to reactivity by $\alpha_0 = \frac{k\beta_{\text{eff}}}{l} (\$ - 1)$

G. Keepin, *Physics of Nuclear Kinetics* (Addison-Wesley, 1965), 273-277

Calculating Reactivity of a Subcritical System Using the Pulsed Neutron Method

At delayed critical ($\beta = 0$), the decay constant α becomes

$$\alpha_{DC} = -\frac{k\beta_{\text{eff}}}{l}$$

and α_0 can be expressed as

$$\alpha_0 = \alpha_{DC} (\beta - 1)$$

Thus, the reactivity is directly proportional to the fundamental-mode decay constant α_0 , and the proportionality constant can be obtained by measurement of the decay constant at delayed critical.

G. Keepin, *Physics of Nuclear Kinetics* (Addison-Wesley, 1965), 273-277

Calculating Reactivity of a Subcritical System Using the Pulsed Neutron Method

- **Gozani method¹**

- $\left(\frac{\rho}{\beta}\right)_{GO} \equiv \frac{\text{Extrapolated Prompt Fundamental Mode Area}}{\text{Delayed Neutron Area}}$

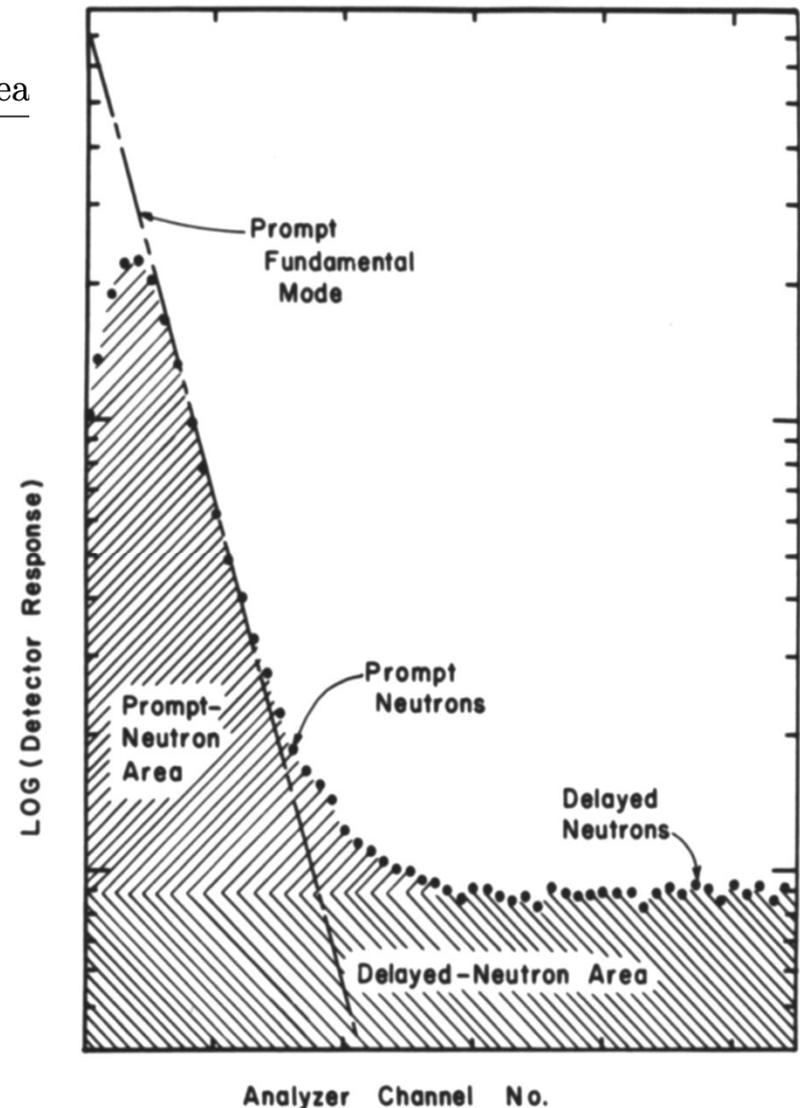
- Eliminates non-fundamental harmonics by extrapolating the fundamental mode back to the time of the pulse

- **Garelis-Russell method²**

- $\left(\frac{\rho}{\beta}\right)_{GR} \equiv \frac{\alpha_0}{\left(\frac{k\beta}{l}\right)} - 1$, where $\frac{k\beta}{l}$ is obtained from

$$\int_0^{\infty} N_p e^{\frac{k\beta}{l}t} dt - \int_0^{\infty} N_p dt = \frac{N_d}{R}$$

- Prompt neutron response is weighted by an exponential factor. More difficult to calculate and implement.



C. F. Masters and K. B. Cady, *Nuclear Science & Engineering*, 29:2, 272-282, 1967

¹T. Gozani, *Nukleonik*, 4, 348, 1962

²E. Garelis and J. Russell, *Nuclear Science & Engineering*, 16:3, 263-270, 1963

MUSiC: Pulsed Neutron Method Preliminary Results

