



ADVANCED REACTOR SAFEGUARDS & SECURITY

# Process monitoring for MC&A: Optical spectroscopy

*FY26 Spring Program Review*

PRESENTED BY

**Shirmir D Branch, Sam A Bryan, Amanda M Lines**

PNNL-SA-222542

# On-line or In-line Measurement for Real-time, Continuous Salt Analysis



- MSR systems are receiving significant attention due to the potential benefits they offer in safety and efficiency
- MSR systems pose unique challenges to MC&A analysis
  - Due to complexity of the salt/reactor system
  - Particularly in the case of on-line conditioning or treatment of salts
- In situ and real-time analysis of salt chemistry is a potential game changer in terms of enabling operators to understand MC&A impacts

**Continuous (On-line):**  
lower accuracy but unparalleled insight into trends and notably improved relevance in statistical sampling

**Vs**

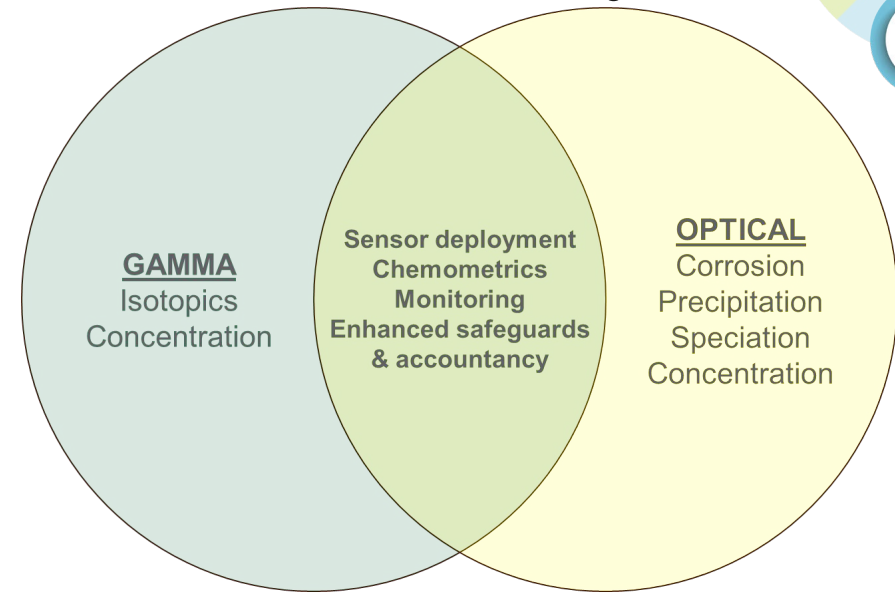
**Periodic (Off-line):**  
higher accuracy but limited to no insight into trends and greater sensitivity to heterogeneous sampling

Combining for a comprehensive and accurate analysis

# Selecting Impactful Tools for On-line Monitoring of MSRs



- Ultimately multiple on- or in-line monitoring tools will be needed to provide the full picture
- Two key tools for chemical/isotopic analysis include Raman and gamma spectroscopy
- **Goal: provide needed information and measurement uncertainty for actinides and other key targets without placing undue burden on the MSR system**

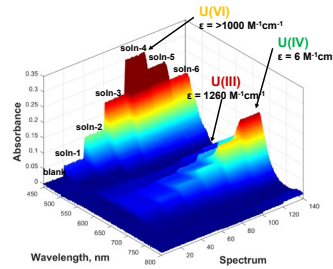


- Combination of multiple analytical techniques builds a more complete description of complex chemical information
  - **Optical (vis-NIR, Raman)** = oxidation state, precipitation, speciation, etc.
  - **Gamma** = isotopic information
- Application of chemometrics to data analysis to improve accuracy and flexibility in analyzing highly complex data exhibiting multiple interfering signals

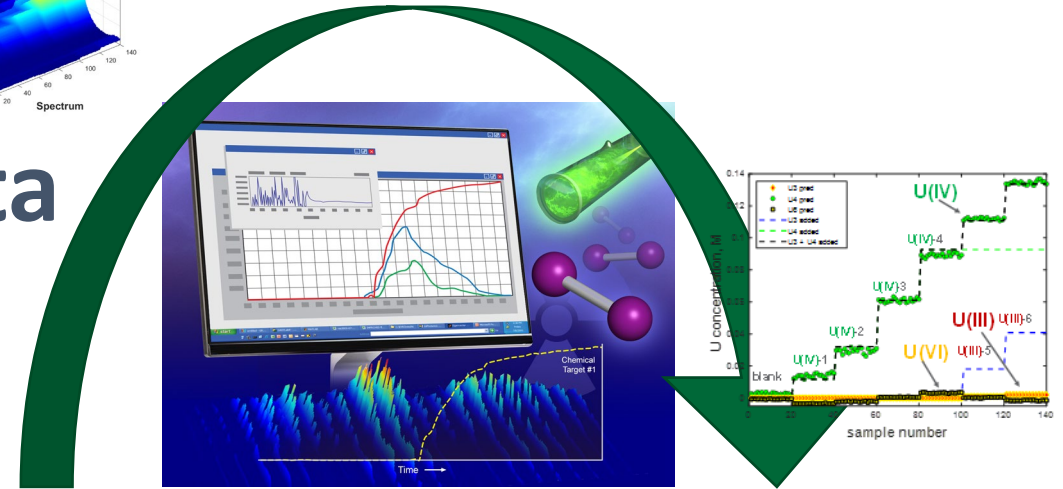
# Sensor Impact is Solidified by Partnering with Robust and Automated Data Analysis



- Sensor data will be complex. To realize benefits of on-line monitoring, automated data analysis is essential
- Furthermore, fusing or combining data from multiple sensors to support analysis builds a more robust tool that can handle salt variability and properly highlight potential heterogeneity
- **Real time insight into complex chemistry**



Data

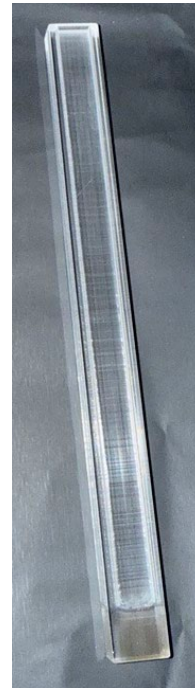
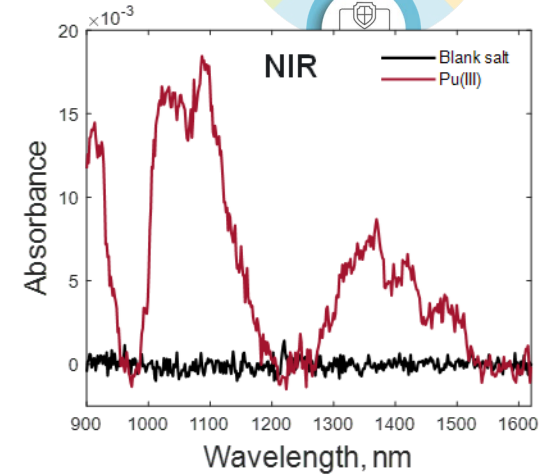
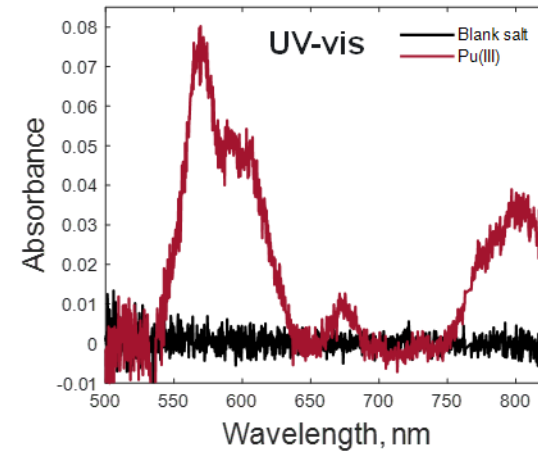


Enabling researchers and operators to understand complex processes with *in situ* and real-time feedback on process conditions

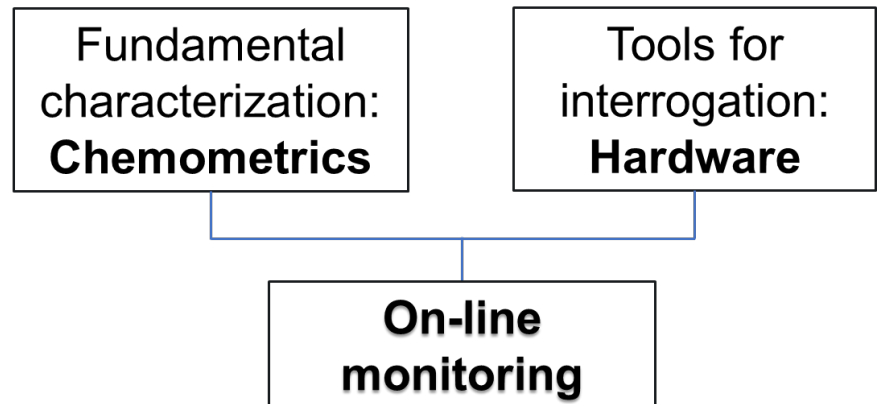
# Considerations for On-line Monitoring



- Target optimal quantification uncertainty for various analytical techniques
- Variable experimental scale
  - Calibration transfer
- Discernment between operation/safeguards effects
  - Temperature, corrosion, etc.
- Modeling techniques
- Monitoring hardware/tools for deployment



\*developed under other efforts



# Analyte accountancy



- Building widely applicable chemometric models and solidifying understanding of achievable uncertainties requires collection of a significant amount of data
- A select list of measurable chemical systems is shown here
- A sequential list of experiments and usability of data will be shown next

Analyte	Molar extinction coeff. (abs)	Chem. form(s) (experimental)	Observed challenges for optical measurement
U(IV)	6.46 M <sup>-1</sup> cm <sup>-1</sup> (670 nm) 13 M <sup>-1</sup> cm <sup>-1</sup> (1100 nm)	UCl <sub>4</sub>	Low molar absorptivity
U(III)	1260 M <sup>-1</sup> cm <sup>-1</sup> (480 nm) 17.5 M <sup>-1</sup> cm <sup>-1</sup> (1100 nm)	UCl <sub>3</sub>	Plating out in presence of some corrosion and FP surrogates
[UO <sub>2</sub> ] <sup>2+</sup>	57.2 M <sup>-1</sup> cm <sup>-1</sup> (450 nm)	UCl <sub>3</sub> (oxidation to U <sup>6+</sup> )	Indicates presence of oxygen in molten salt system
Pu(III)	20 M <sup>-1</sup> cm <sup>-1</sup> (563 nm) 0.05 M <sup>-1</sup> cm <sup>-1</sup> (1100 nm)	K <sub>2</sub> PuCl <sub>6</sub> Na <sub>2</sub> PuCl <sub>6</sub> PuCl <sub>3</sub>	Low solubility at lower concentrations; optical fingerprint can overlap other actinide species
Nd(III)	10.7 M <sup>-1</sup> cm <sup>-1</sup> (589 nm)	NdCl <sub>3</sub>	Optical fingerprint can overlap actinide species
Eu(III)	420 M <sup>-1</sup> cm <sup>-1</sup> (330 nm)	EuCl <sub>3</sub>	Precipitation with corrosion products
Er(III)	9.2 M <sup>-1</sup> cm <sup>-1</sup> (377 nm)	ErCl <sub>3</sub>	Low molar absorptivity
Ho(III)	10.10 M <sup>-1</sup> cm <sup>-1</sup> (450 nm)	HoCl <sub>3</sub>	Low molar absorptivity
Dy(III)	0.233 M <sup>-1</sup> cm <sup>-1</sup> (1111 nm)	DyCl <sub>3</sub>	Low molar absorptivity; optical fingerprint can overlap actinide species
Cr(III)	46 M <sup>-1</sup> cm <sup>-1</sup> (545 nm)	CrCl <sub>3</sub>	Plating out in presence of other analytes
Co(II)	204 M <sup>-1</sup> cm <sup>-1</sup> (609 nm)	CoCl <sub>2</sub>	Plating out in presence of other analytes
Fe(III)	398 M <sup>-1</sup> cm <sup>-1</sup> (295 nm)	FeCl <sub>3</sub>	Optical fingerprint overlaps with actinides of interest
Ni(II)	63.8 M <sup>-1</sup> cm <sup>-1</sup> (628 nm)	NiCl <sub>2</sub>	Plating out in presence of other analytes

# Accountancy in mixed analyte systems



- Several variants of mixed salt systems were studied
- Three examples of experimental schemes are shown here
- Experimental approaches to mixed U/Pu systems precipitated, meaning data could not be leveraged for model building efforts

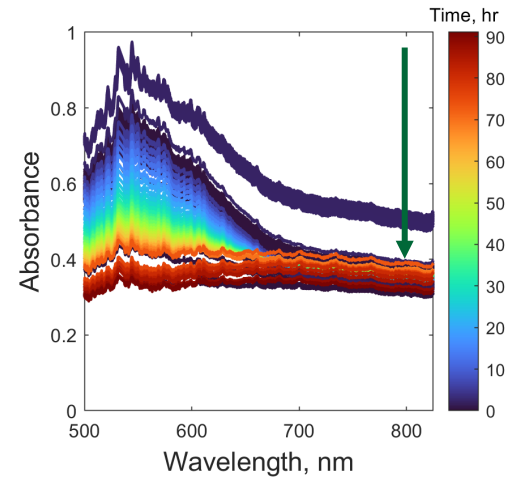
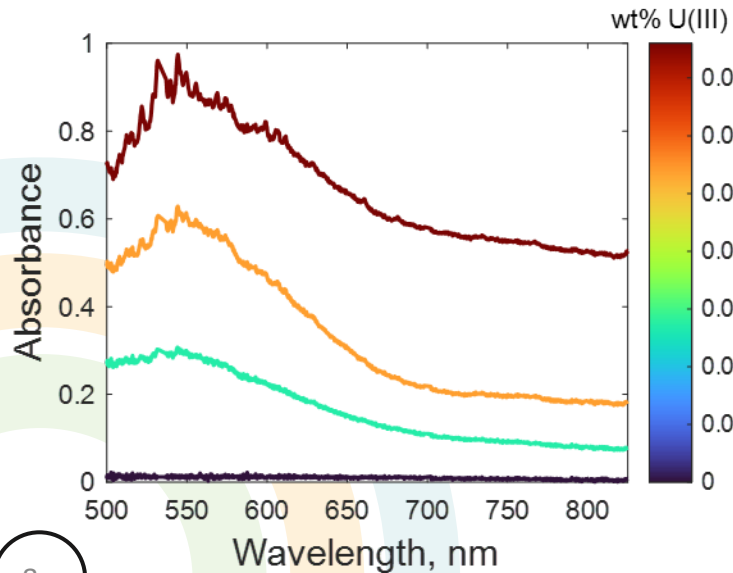
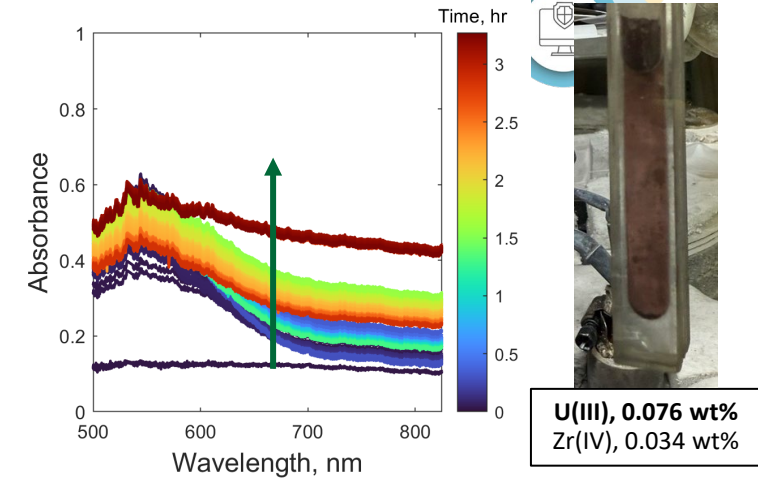
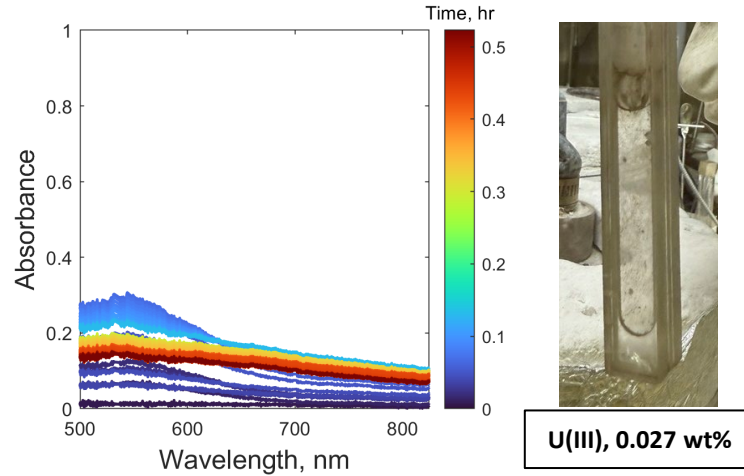
Experiment #1	Experiment #2	Experiment #3
<ol style="list-style-type: none"><li>1. Blank salt</li><li>2. Add U</li><li>3. Add Nd (precipitation)</li></ol>	<ol style="list-style-type: none"><li>1. Blank salt</li><li>2. Add Nd</li><li>3. Add Cr</li><li>4. Add U (precipitation)</li></ol>	<ol style="list-style-type: none"><li>1. Blank salt</li><li>2. Add Pu (precipitation)</li><li>3. Add Zr (dissolution)</li><li>4. Add Nd</li><li>5. Add U (precipitation)</li><li>6. Add Zr (no change)</li></ol>

**A review of the multiple mixed system attempts is detailed in PNNL-39142.**

# Precipitation can still be tracked



- Dissolved U species
- Oxidation and addition of Pu causes formation of precipitates
- Quantifiable U fingerprint still observable

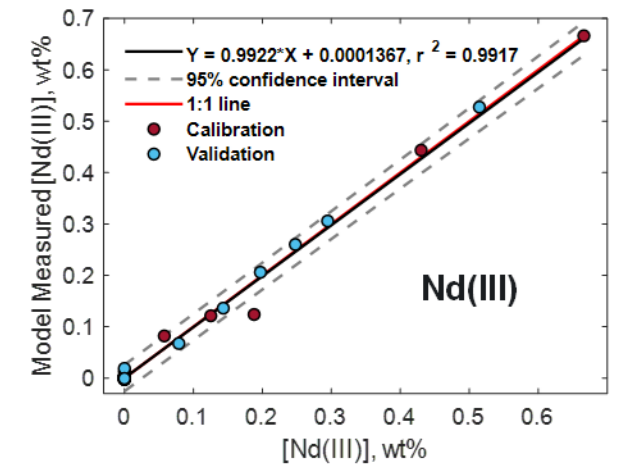
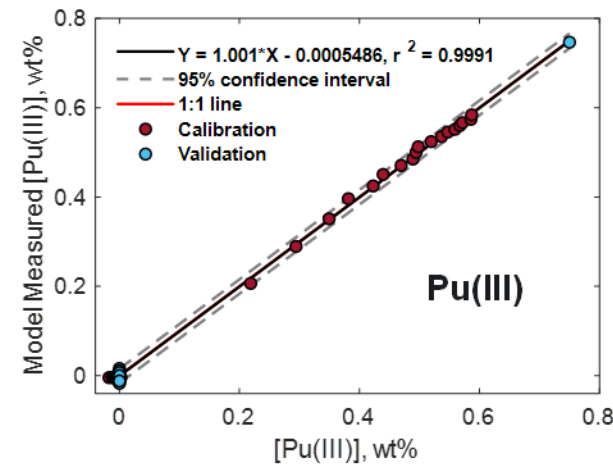
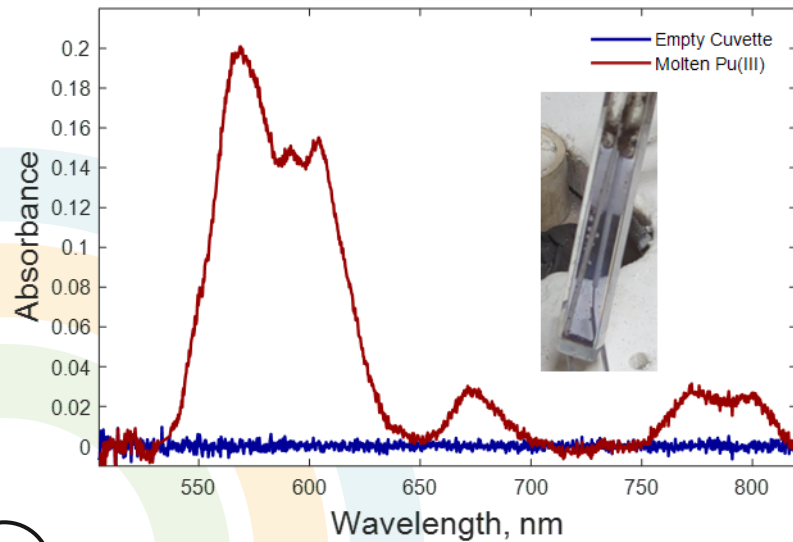
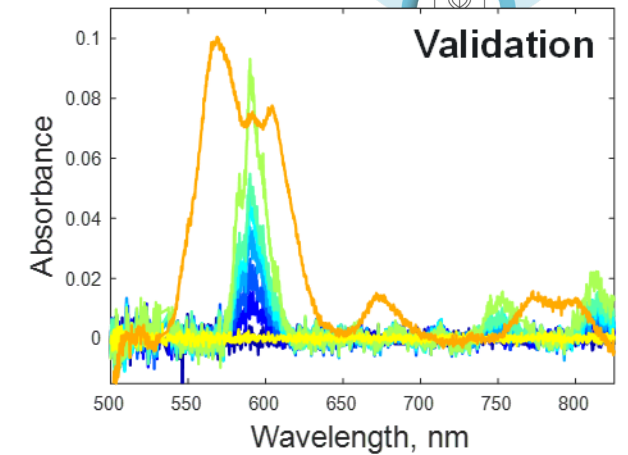
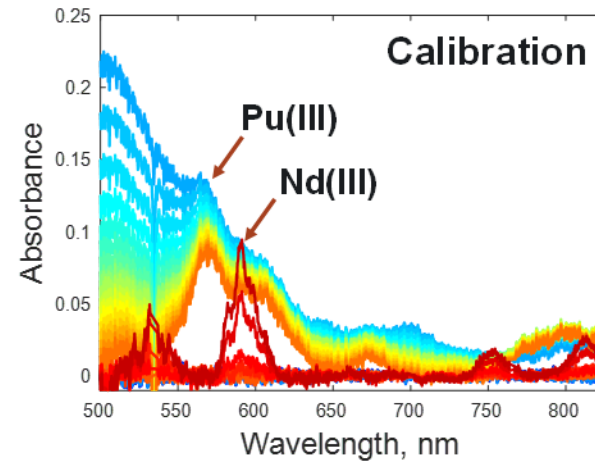


**“Bad” data is not bad data**

# Accountancy of Pu in mixed analytes in molten chloride

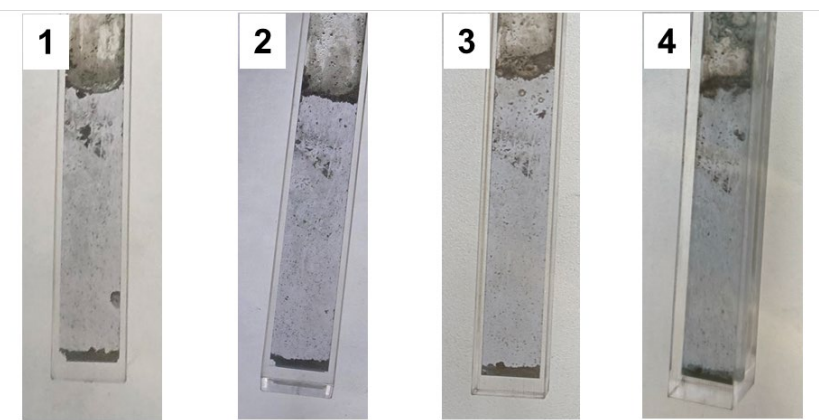


- Pu in single and multi-component salts was studied to gain an understanding of modeling uncertainties
- Individual calibration (model building) and validation (model testing) data sets were completed/collected
- Calibration data is shown here

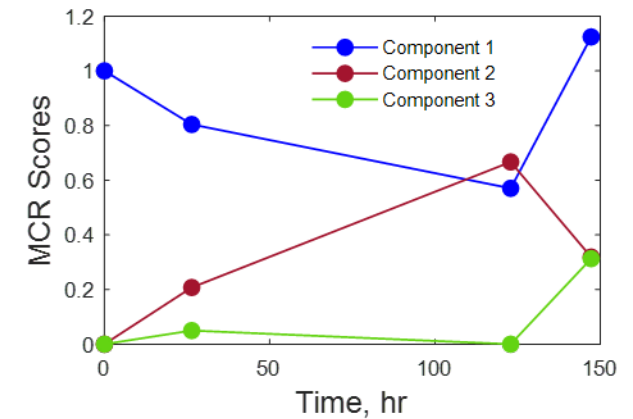
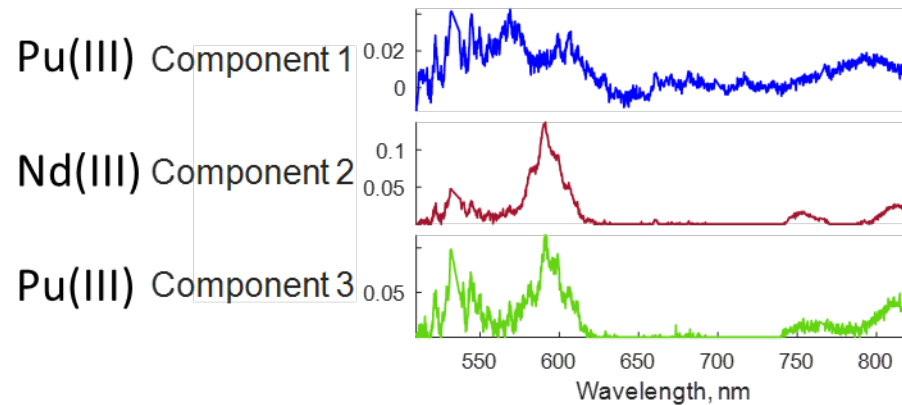
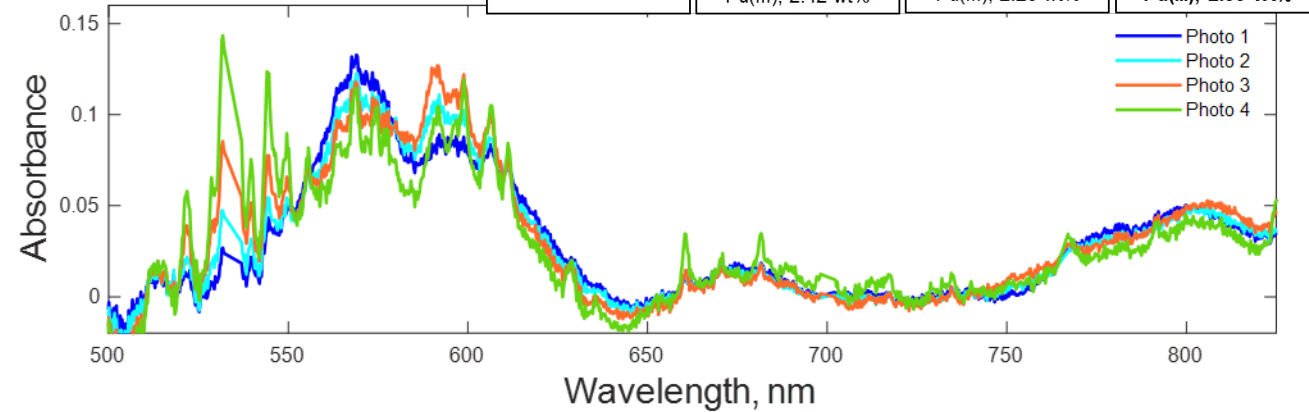


# Accountancy of Pu in mixed analytes in molten chloride

- Pu ( $\lambda_{\max} = 568 \text{ nm}$ ) and Nd ( $\lambda_{\max} = 589 \text{ nm}$ ) have broad, overlapping peaks
- Precipitation impacts use of quantification tools such as Partial Least Squares (PLS) Regression
- Multivariate Curve Resolution (MCR) loadings discern impact of interfering species (Nd) on target analyte (Pu)

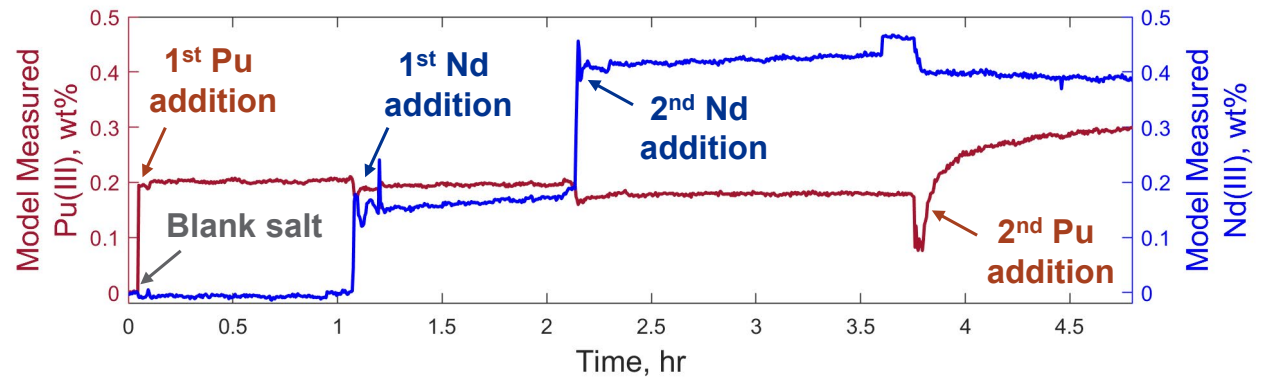
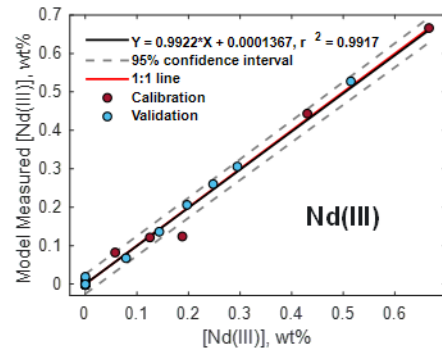
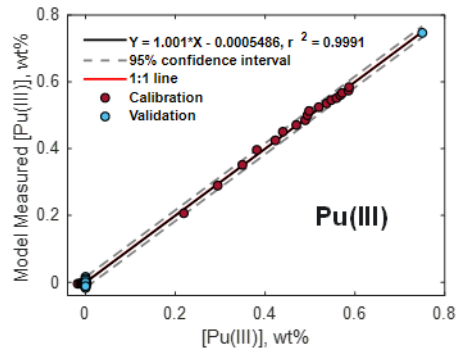
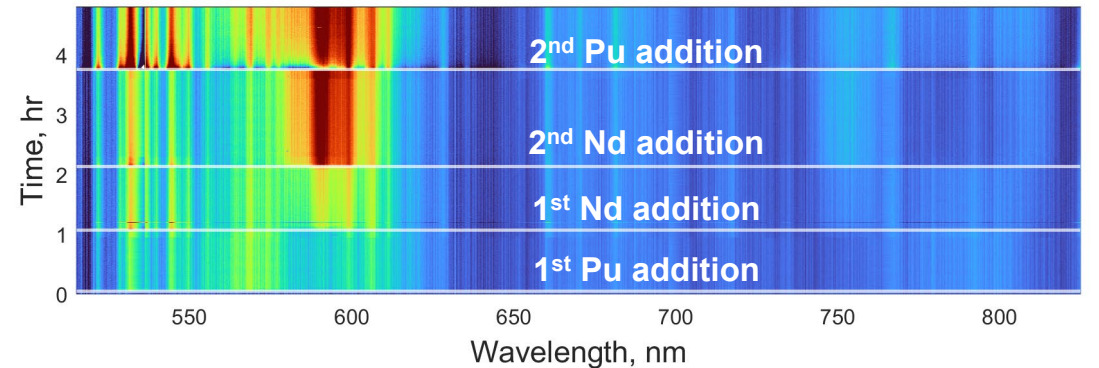
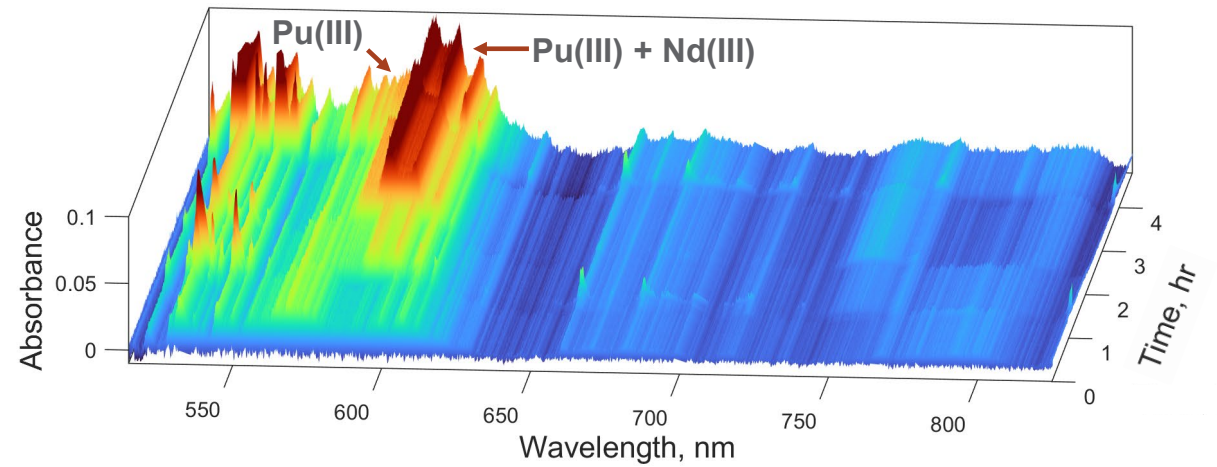


Pu(III), 2.58 wt%      Nd(III), 0.33 wt%  
Pu(III), 2.42 wt%      Nd(III), 0.61 wt%  
Pu(III), 2.28 wt%      Nd(III), 0.58 wt%  
Pu(III), 2.99 wt%



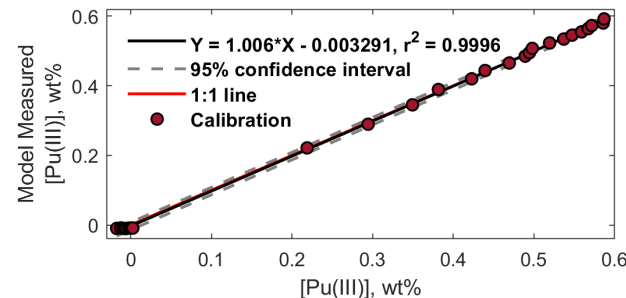
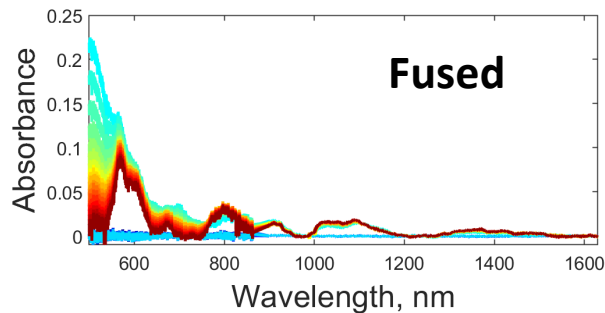
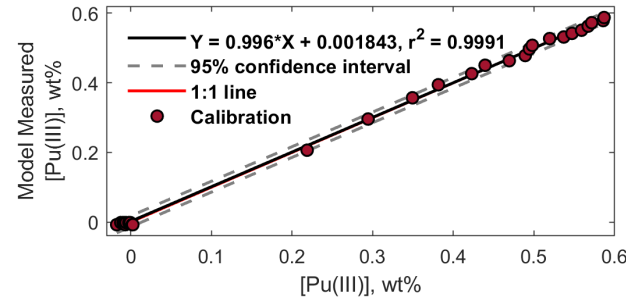
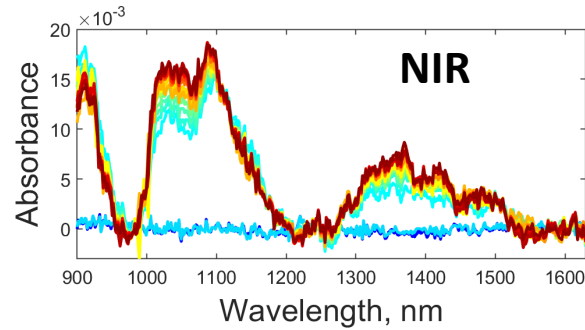
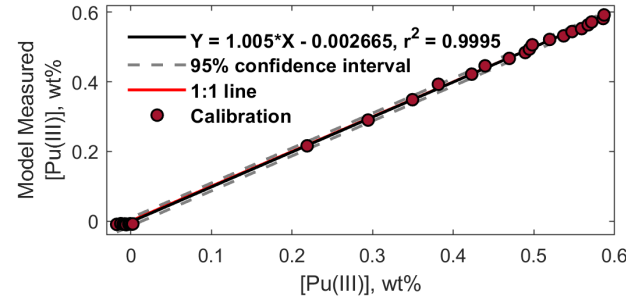
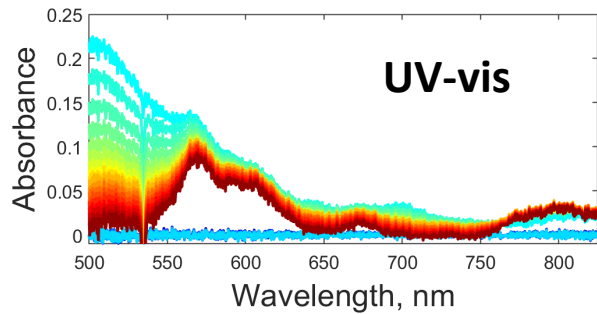
# Accountancy of Pu in mixed analytes

- Validation runs were completed on mixed Pu/Nd systems looking at different ratios of Pu to Nd for each experiment



Model Statistics	Pu Uncertainty, wt%	Pu % Uncertainty	Nd Uncertainty, wt%	Nd % Uncertainty
RMSEC	0.00753	1.3	0.0123	1.8
RMSECV	0.00835	1.4	0.0148	2.2
RMSEP	0.0100	1.7	0.0109	1.6

# Data fusion: Robust monitoring

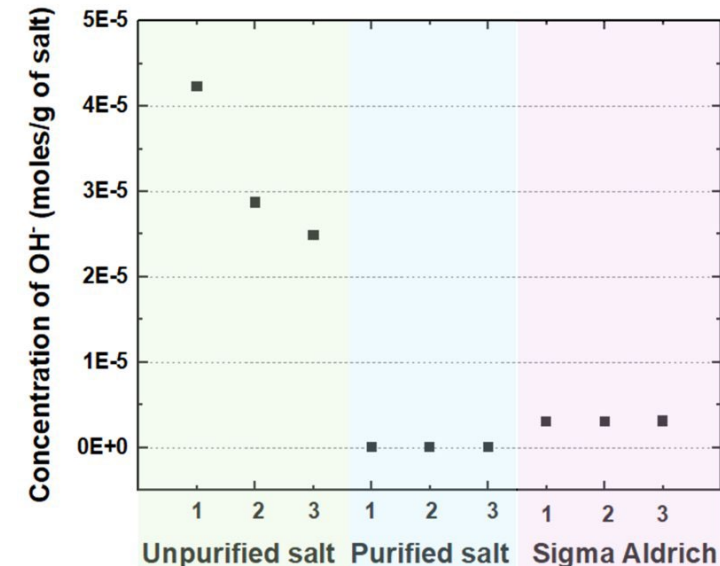
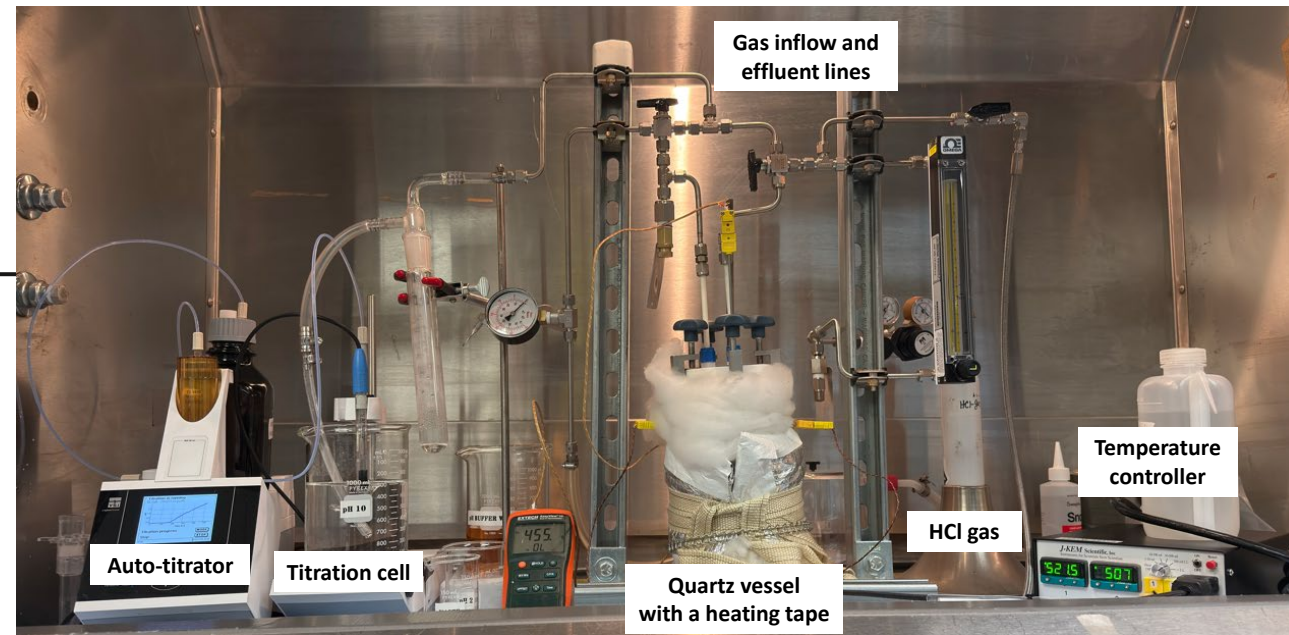


System	RMSEC, wt%	RMSECV, wt%	% Uncertainty
UV-vis	0.005577	0.006985	1.2
NIR	0.007369	0.008596	1.5
Fused	0.005481	0.006600	1.1

- Optical spectroscopy can leverage the entire spectrum for robust monitoring
- The model performance for one analytical technique can be used to validate another technique
- These model techniques are not limited to optical spectroscopy

# Some Notes Regarding Precipitation

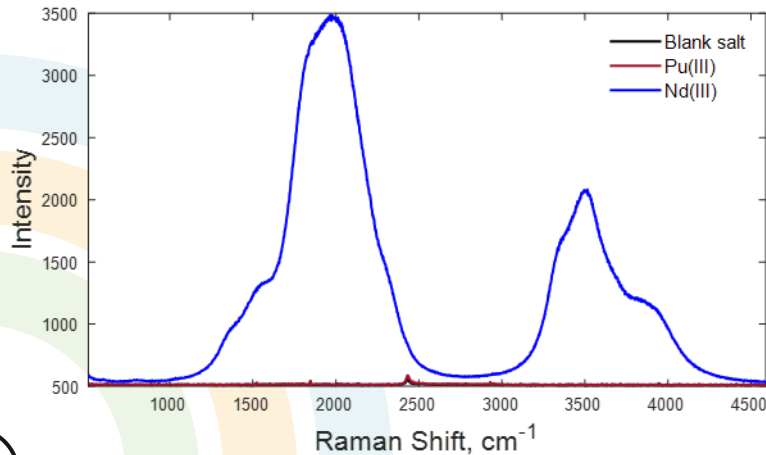
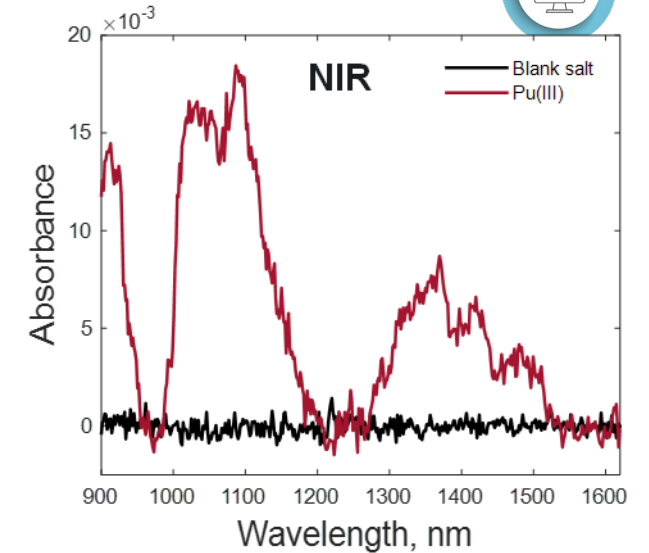
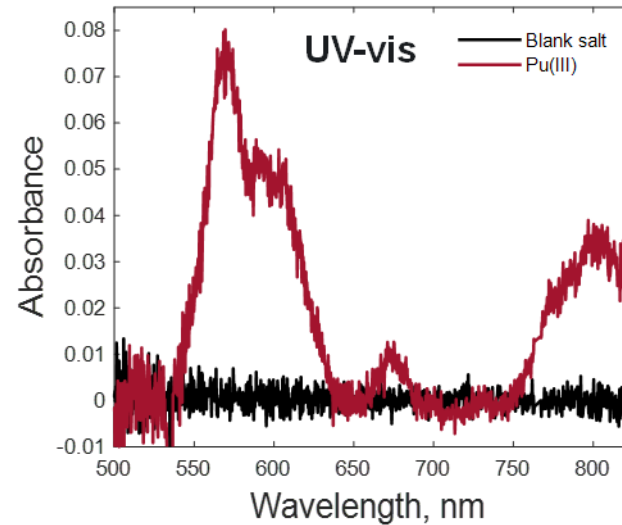
- Given the goal to assess accuracy/uncertainty of models, data of precipitated systems cannot be used in modeling
  - Model performance is assessed based on “known” values, precipitation adds significant uncertainty to the “known” concentration of dissolved metal species
  - U/Pu systems are a key target of study
- Salt purification systems underway
  - Non-rad purification system demonstrated in FY25
  - Continuing with rad purification system in FY26 to allow for a Pu/U study in the purified salt



# Expanding Sensors to Provide More System Flexibility



- Expansion of absorbance testing
  - Adding NIR to chemometric library
  - Potential “quiet” spots in high U systems



- Analytes of interest (e.g. U and Pu) have optical fingerprints in the NIR where some interferent species (e.g. Nd, Cr) have overlapping fingerprints

# Paths forward/Acknowledgements



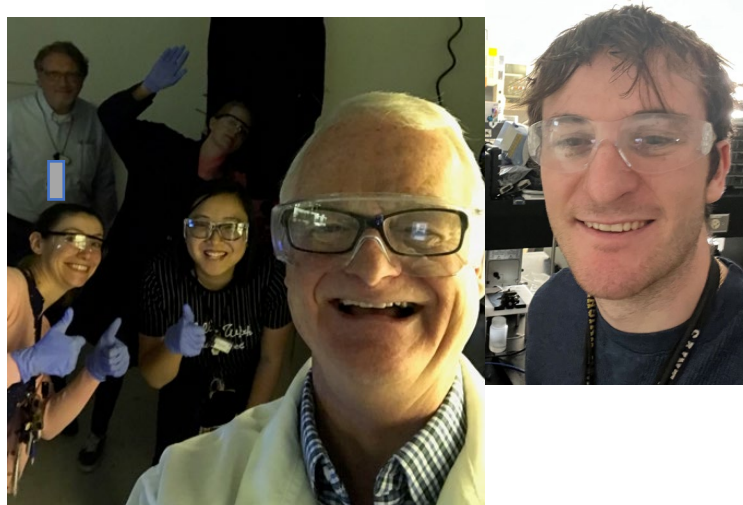
- Chemometric data fusion demonstrating with optical and gamma spectroscopy
- Technical evaluation on the complexity of molten salt chemistry for online monitoring
- Testing with robust monitoring tools

## • PNNL OLM team

- Shirmir Branch ([shirmir.branch@pnnl.gov](mailto:shirmir.branch@pnnl.gov))
- Amanda Lines ([amanda.lines@pnnl.gov](mailto:amanda.lines@pnnl.gov))
- Sam Bryan ([sam.bryan@pnnl.gov](mailto:sam.bryan@pnnl.gov))
- Heather Felmy
- Adan Schafer Medina
- Suhee Choi
- Jason Rakos
- Forrest Heller
- Dana Arbova
- Grey Batie
- Erin Good

Milestone	Description	Due date
M3RS-26PN0401031	Completion of technical report on salt chemistry complexity for role of online monitoring	31 Mar 2026 COMPLETE
M3RS-26PN0401032	Combined optical and gamma measurement on molten salt system	30 Sept 2026

# Questions?



Thank you!

