

Actinides and Rare Earths Focus Topic

Room C124 - Session AC+MI+TH-TuM

Magnetism, Electron Correlation, and Superconductivity in the Actinides/Rare Earths

Moderators: Edgar Buck, PNNL, Tomasz Durakiewicz, Idaho National Laboratory, Krzysztof Gofryk, Idaho National Laboratory

8:00am AC+MI+TH-TuM-1 Uranium and Cerium Based Systems Probed with High-Pressure XANES and XMCD, *Fabrice Wilhelm*, A. Roagelv, ESRF, France

INVITED

This talk reviews recent advances in use of polarized x-rays to study local magnetic properties and electronic structure of uranium and cerium based compounds such as the ferromagnetic UGe_2 , the heavy-fermion paramagnet UTe_2 and the nanolamellar ferromagnetic Kondo lattice $Mo_4Ce_4Al_7C_3$ system.

8:40am AC+MI+TH-TuM-3 Searching for New Uranium-Based Arsenides, *Eteri Svanidze*, Max Planck Institute for Chemical Physics of Solids, Germany
As new classes of superconducting materials emerge, puzzles of high-temperature superconductivity continue to be one of the pressing issues in condensed matter physics and solid-state chemistry. In particular, iron pnictide superconductors still pose many open questions. Surprisingly, very few actinide-based analogues of iron pnictide high temperature superconductors have been reported so far, perhaps as a result of synthesis complications imposed by toxicity, reactivity, and high vapor pressure of constituent elements. In this work, we revisit the U-Fe-As ternary, in which only one compounds has been reported to exist so far – $UFeAs_2$ ($P4/nmm$ space group) [1]. By implementing flux synthesis, we were able to grow large single crystals of UFe_5As_3 compound, which adopts UCr_5P_3 structure type [2] ($P2_1/m$, $mP18$) with lattice parameters $a = 7.0501(17)$ Å, $b = 3.8582(9)$ Å and $c = 9.6342(13)$ Å, $\beta = 100.2(8)^\circ$ [3]. The refined composition agrees well with the elemental ratio established by the EDX analysis – $U_{9.9(2)}Fe_{58.7(2)}As_{31.4(2)}$. The magnetic behavior of UFe_5As_3 was studied by magnetic susceptibility measurements in the temperature range 1.8 - 600 K and an antiferromagnetic ordering below $T_N = 56$ K was established. The Sommerfeld coefficient $g = 138$ mJ/mol K^2 indicates enhanced effective electron mass. More in-depth investigations of this and other U-Fe-As ternary compounds are currently underway.

References:

- [1] D. Kaczorowski, J. Alloys. Compd. 186, 333-338 (1992).
- [2] W. Jeitschko, R. Brink, P.G. Pollmeier, Z. Naturforsch. (1993) 48b, p. 52-57.
- [3] N.Zaremba, M. Krnel, Yu. Prots, A. Leithe-Jasper, Yu. Grin, E. Svanidze, in preparation (2023)

9:00am AC+MI+TH-TuM-4 5f Magnetism at an Extreme, *Ladislav Havela*, Charles University, Faculty of Mathematics and Physics, Czechia; *V. Buturlim*, Idaho National Laboratory; *F. Honda*, Tohoku University, Japan; *D. Kaczorowski*, Institute of Low Temperature and Structure Research, Wrocław, Poland

Specific features of 5f magnetism in U-based systems, namely the strong spin-orbit coupling and very strong magnetic anisotropy or giant magnetoresistance, are degraded by low temperatures of magnetic ordering. The variability of T_C or T_N in intermetallics has been explored and it turns out that involvement of magnetic d -metals such as Fe or Co does not help, unless very high fraction of the transition metals turns a compound in fact a regular d -magnet, in which the unique 5f features are suppressed. The reason is the 5f- d hybridization with a destabilizing action on both d and 5f moments.

Seeking additional degrees of freedom we turned towards compounds with more polar character of bonding, which drives U to the verge of metallicity. Interesting alternative to intermetallics is provided by Zintl phases, typically ternary compounds with one cation (e.g. an f -metal) and a complex covalently bond anion, stabilized by a charge transfer from the cation. An excellent insight is provided by such compounds with layered crystal structures, which leads to strongly anisotropic electronic properties. Our research focuses on Zintl phases with the trigonal $CaAl_2Si_2$ type ($P-3m$), formed by alternating cationic (Ca) and anionic (Al-Si) layers. Those without f -metals or with 4f metals tend to be narrow-gap semiconductors. U-based phases such as UCu_2P_2 are semi-metallic. The prominence of this compound dwells in ferromagnetism with high Curie temperature $T_C = 216$ K, which is

further enhanced by pressure application to the vicinity of room temperature. Ab-initio calculations reveal it this material is actually a half-metal, with spin-split 5f band bringing spin-up only states to the Fermi level. The conduction electron states remain hybridized with the 5f states, which causes also the non-f electrons to be fully spin polarized. The magnetisms is of local moment type, with magnetic entropy exceeding $R \ln 2$, but arising from 5f bands. UCu_2P_2 can be compared with $EuZn_2P_2$. Here the Eu^{2+} moments are organized in an AF structure. The field alignment of the moments impacts the width of the gap seen in transport data, but the material remains semiconducting.

This work was supported by the Grant Agency of the Czech Republic under the grant No. 21-09766S.

9:20am AC+MI+TH-TuM-5 Valence-to-Core RIXS in Insulating Compounds with 4f and 5f Elements, *Jindrich Kolorenc*, Institute of Physics, Czech Academy of Sciences, Czechia

In the first part, we investigate the electronic structure of europium sulfide (EuS) with the aim to understand the valence-to-core resonant inelastic x-ray scattering (RIXS) spectra measured at the Eu L_3 edge. We show that the main part of the observed signal comes from the direct RIXS: an Eu 2p core electron is excited to an empty Eu 5d band above the Fermi level, and then another electron from an Eu 5d state hybridized with the S 3p bands (located below the Fermi level and hence occupied) fills back the core hole. Besides this straightforward channel, the measured RIXS spectra display two satellite features. After considering several candidates (such as excitations of the 4f shell excited by indirect RIXS processes, that is, by the interaction with the core hole in the intermediate state), we arrive at the conclusion that the satellites are excitons formed by a 4f hole and an 5d electron localized at the same Eu atom. Such excitons were suggested in the context of optical absorption a long time ago [1], but the concept is not widely accepted. It was argued that existence of these excitons is incompatible with photoconductivity measurements [2]. Our observations indicate that these arguments may need to be revisited.

In the second part, we simulate valence-to-core RIXS at the actinide M_5 edge in selected actinide oxides starting from the dynamical mean-field electronic structure obtained earlier [3] and using techniques similar to [4]. This investigation is motivated by the recent experimental study of UO_2 and UF_4 [5]. Our preliminary results suggest that the feature observed at an energy loss of roughly 10 eV above the white line is likely a charge-transfer excitation, the intensity of which is sensitive to the hybridization between the actinide 5f states and the ligand 2p states.

This work was supported by the Czech Science Foundation under the grant No. 21-09766S.

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- [3] J. Kolorenc, A. Shick, A. I. Lichtenstein, Phys. Rev. B 92 (2015) 085125, <https://doi.org/10.1103/PhysRevB.92.085125>
- [4] J. Kolorenc, Physica B: Condensed Matter 536 (2018) 695, <https://doi.org/10.1016/j.physb.2017.08.069>
- [5] J. G. Tobin et al., J. Phys.: Condens. Matter 34 (2022) 505601, <https://doi.org/10.1088/1361-648X/ac9bbd>

9:40am AC+MI+TH-TuM-6 Magnetism of Binary Actinide Oxides, *Binod Rai*, A. Bretaña, Savannah River National Laboratory; G. Morrison, University of South Carolina, Columbia; R. Greer, Savannah River National Laboratory; K. Gofryk, Idaho National Laboratory; H. zur Loye, University of South Carolina, Columbia

Historically, the bulk of actinide oxide research has focused on the properties of various uranium and plutonium compounds as they impact their use in the nuclear fuel industry. While the fundamental understanding of compounds such as UO_2 has improved, a fundamental grasp of the physical properties of other actinide oxides remains elusive. In actinide systems, the 5f electrons experience a uniquely delicate balance of effects and interactions having similar energy scales, which are often difficult to properly disentangle. The interplay of factors such as the dual character of 5f-states, competing interactions, and strong spin-orbit coupling results in magnetically unusual and intriguing behavior: multi-k antiferromagnetic ordering, multipolar ordering, mixed valence configurations, and more. In this presentation, I will provide an overview of the available synthesis techniques for selected binary actinide oxides. I will also show the current state of knowledge on their crystal structures and magnetic properties.

Moreover, I will talk about the future opportunities that are vital for a fundamental understanding of these systems.

11:00am **AC+MI+TH-TuM-10 N-Point Saddle-Band Model for the Hidden Order Phase of URu₂Si₂**, *J. D. Denlinger*, Lawrence Berkeley National Laboratory; *J. Kang*, The Catholic University of Korea; *L. Dudy*, SOLEIL, France; *J. Allen*, University of Michigan; *L. Wray*, New York University; *A. Gallagher*, *R. Baumbach*, National High Magnetic Field Laboratory; *N. Butch*, University of Maryland; *M. Maple*, University of California, San Diego

The 5f heavy fermion system of URu₂Si₂ is famous for an unconventional hidden order (HO) phase transition at T₀=17.5K and the various related doping phase diagrams explored to elucidate the nature of this unknown transition. A new model of an extended U 5f saddle-point dispersion at the N-point is proposed for the underlying physics of this 35-year old puzzle. With a 10X DFT energy renormalization suggested by angle resolved photoemission (ARPES) measurements in comparison to DMFT calculations [1], this feature lies only a few meV below E_F and thus is thermally active at T₀. ARPES reveals a distinct temperature-dependent evolution of incommensurate 5f nesting hotspots lying along Z-N-Z [1], whose origins are found in electronic susceptibility calculations of intraband q-scattering for thermal occupations in the temperature range of T₀. Tuning of the N-point region f-band energetics with chemical doping is thus naturally proposed to unify our understanding of the various phase diagrams. ARPES is presented which confirms the existence of N-point spectral and dichroic changes for ferromagnetic Re/Ru-substitution, for (001) antiferromagnetic Fe and Os/Ru-substitution, and for (1/2,1/2,1/2) antiferromagnetic P/Si-substitution, thus further demonstrating experimentally the key relevance of the N-point electronic structure to the HO phase. [1] J. D. Denlinger et al., *Electron. Struct.* 4, 013001 (2022).

11:20am **AC+MI+TH-TuM-11 Magnetoelastic Properties of 5f Ferromagnet UCu₂P₂**, *Volodymyr Buturlim*, Idaho National Laboratory; *P. Doležal*, *O. Koloskova*, *J. Prchal*, Charles University, Czechia; *I. Turek*, Charles university, Czechia; *M. Martinez Celis*, CRISMAT Laboratory, France; *F. Honda*, Kyushu University, Japan; *M. Divis*, Charles University, Czechia; *D. Kaczorowski*, Polish Academy of Sciences, Poland; *K. Gofryk*, Idaho National Laboratory; *L. Havela*, Charles University, Czechia

UCu₂P₂, a Zintl phase with the trigonal CaAl₂Si₂ structure type, is a 5f ferromagnet with a record-high Curie temperature among U compounds, T_c = 216 K [1]. While the size of the magnetic moment 2.0 μ_B/U is not surprising due to U-U spacing exceeding the Hill limit, the reasons for the high T_c are less understood. Ab-initio calculations reveal only very weak hybridization of the U-5f states with the 6d states as well as with electronic states of Cu and P. It seems that a transfer of U-6d states to the P-3p states is an important ingredient, which was highlighted by a rapid increase of T_c under hydrostatic pressure so that a room-temperature 5f ferromagnetism could be demonstrated. Besides magnetization, transport, and heat capacity studies on single crystals we performed also characterization of a polycrystalline material, which has T_c = 219 K. Thermal expansion study revealed a moderate increase of both lattice parameters just below T_c, so we can exclude that the pressure enhancement of T_c is driven simply by thermodynamics (via the Ehrenfest relation). Hence the reasons have to be attributed to the enhancement of specific U-U couplings upon compression. Indeed, ab initio calculations probing the energy enhancement upon moments reversal gave a semi-quantitative account of the observed tendency of T_c.

[1] D. Kaczorowski, R. Troc, Magnetic and transport properties of a strongly anisotropic ferromagnet UCu₂P₂, *J. Phys. Condens. Matter*, 1990, 2, 4185.

*This work was supported by the Czech Science Foundation under the grant No. 21-09766S.

11:40am **AC+MI+TH-TuM-12 Density Functional Theory Calculations of the Phonons in Gamma and Delta Phase Pu**, *Sven P. Rudin*, Los Alamos National Laboratory

The thermodynamics of plutonium challenges both theory and experiment. The challenge arises from the multiple structural phase transitions, the nature of the 5f electrons, and the effects due to the material's self-irradiation. Density functional theory calculations have made steady progress toward clarifying experimental measurements. Recently, for the delta phase a non-collinear 3Q spin structure was shown to make all structurally equivalent bonds equivalent in their bonding character. This results in elastic constants and phonons with the correct symmetry, and the calculated phonon dispersion agrees well with experiment. Extending the calculations into the gamma phase of Pu now provides a prediction for the

phonons of that phase.

12:00pm **AC+MI+TH-TuM-13 The 5f UDOS of the Actinide Dioxides: Why Pu is n = 5 in PuO₂**, *James Tobin*, University of Wisconsin-Oshkosh; *H. Ramanantoanina*, KIT, Germany; *C. Daul*, U. Fribourg, Switzerland; *S. Yu*, LLNL; *P. Roussel*, AWE, UK; *S. Nowak*, *R. Alonso-Mori*, *T. Kroll*, *D. Nordlund*, *T. Weng*, *D. Sokaras*, SSRL

The Unoccupied Density of States (UDOS) of ThO₂, UO₂ and PuO₂ have been investigated with a combined experimental and theoretical approach.^{1,2}Ligand field density functional theory calculations of the dioxides of thorium, uranium, and plutonium have been combined with high-energy-resolution fluorescence detection (HERFD) in x-ray absorption spectroscopy and inverse photoelectron spectroscopy (IPES) measurements to provide powerful insight into the underlying composition of the unoccupied 5f electronic structure in these 5f localized systems. Fine structure in the 5f_{5/2} transitions in HERFD can be directly correlated with the fine structure in the leading edge of the IPES. The shapes, intensities, and systematics in HERFD and IPES are explained in a consistent and rigorous fashion in terms of the j-specific 5f electronic structure. Additionally, an actinide N_{4,5} branching ratio analysis of PuO₂ and UO₂ has been performed, including measurements with a scanning transmission electron microscope at the Advanced Light Source and simulation with FEFF. It is shown that the 5f occupation of the Pu in plutonium dioxide is n = 5.

References

1. J. G. Tobin, H. Ramanantoanina, C. Daul, S.-W. Yu, P. Roussel, S. Nowak, R. Alonso-Mori, T. Kroll, D. Nordlund, T.-C. Weng, D. Sokaras, "The Unoccupied Electronic Structure of Actinide Dioxides," *Phys. Rev. B* 2022, **105**, 125129. <https://doi.org/10.1103/PhysRevB.105.125129>
2. J. G. Tobin and S.-W. Yu, "Pu 5f Occupation in Plutonium Dioxide," *Inorg. Chem.* 2023, **62**, 6, 2592–2598, <https://doi.org/10.1021/acs.inorgchem.2c03202>

Actinides and Rare Earths Focus Topic

Room C124 - Session AC+MI+TH-TuA

Chemistry and Physics of the Actinides/Rare Earths

Moderators: Ladislav Havela, Charles University, Czech Republic, Eteri Svanidze, Max Planck Institute for Chemical Physics of Solids, Gertrud Zwicknagl, Technical University Braunschweig

2:20pm **AC+MI+TH-TuA-1 Electronic Structure Methods for f-Block Elements: Are We There Yet?**, X. Li, Chad Hoyer, University of Washington
INVITED

The computational modeling of rare-earth and heavy-element complexes requires an accurate treatment of spin-orbit and electron correlation effects to fully understand the physical underpinnings of their chemical properties. The fully correlated frequency-independent Dirac-Coulomb-Breit Hamiltonian in a multireference framework provides the most accurate description of electron-electron interaction before going to a genuine relativistic quantum electrodynamics theory of many-electron systems. Our recent studies using many-body multireference methods suggest that there is a significant correlation effect of inner-valence electrons in rare-earth and heavy-element complexes. Ignoring this effect could lead to inaccurate descriptions of molecular properties, such as covalency, bonding, and spectroscopic response. In this talk, we will examine how the correlation effect of inner-valence electrons is manifested in molecular properties with a focus on the covalency in rare-earth and heavy-element complexes.

3:00pm **AC+MI+TH-TuA-3 Structures and Electronic States of Actinide and Lanthanide Complexes with Phenanthroline Derivatives**, Tsuyoshi Yaita, Japan Atomic Energy Agency, Japan
INVITED

Actinides and lanthanides are f-electron elements, and thus their chemical behavior is very similar if they have the same valence. Recently, the demand for lanthanides, especially specific lanthanides such as Nd, Tb, and Dy, is increasing with the realization of a carbon-neutral society and the shift to EVs for vehicle, and there is a growing need to find new resources and recycle them from urban mines. On the other hand, in the case of geological disposal of radioactive waste generated from nuclear power generation, separation of minor actinides such as Am, which are long-lived α -nuclides, is very important from the viewpoint of recycling in a geological repository in a small country like Japan, and the establishment of separation and transmutation technology is desired. This means that the need for separation of specific f elements from solutions such as high-level liquid waste containing a wide variety of lanthanides and actinides will, needless to say, increase more and more.

Here, we have been new synthesized phenanthroline amide derivatives (phenanthroline amide: PTA) with N and O as donor elements for these purposes. These ligands achieve separation of a special f-element by recognize slight size differences or utilize differences in their interactions with the donor elements.

In this talk, the chemical bonding properties of phenanthroline ligands to actinides and lanthanides by XAS/XES and RIXS using soft X-rays, the structural properties of tridentate and tetradentate PTA derivatives by single crystal structural analysis and EXAFS, respectively, will be presented. Taken together, the mechanisms governing ion recognition of PTA derivatives will be reported.

4:20pm **AC+MI+TH-TuA-7 Unraveling the Unique Properties of f-Element Terpyridyl Complexes**, Alyssa Gaiser, Michigan State University; C. Celis-Barras, Colorado School of Mines; F. White, Oak Ridge National Laboratory; T. Albrecht-Schoenert, Colorado School of Mines
INVITED

A terpyridine derivative was synthesized with the lanthanides and several trivalent actinides targeted at probing differences between americium and curium. In addition to investigating high quality structural data on both the americium and curium concluding in a slight increase in covalent character in the americium system, the cerium and berkelium analogs proved to be even more interesting, exhibiting enhanced covalent character in the system. This terpyridyl system continues to provide novel behavior throughout the f-elements exemplifying the inverse trans influence throughout the series with the interaction between the terpyridyl and the nitrate molecule in the same plane.

5:00pm **AC+MI+TH-TuA-9 Electronic Properties of Plutonium Oxycarbide**, Paul Roussel, AWE plc, UK

Plutonium is both electropositive and highly reactive, such that an oxide film of varying thickness is always present on metal samples. It is of interest from a safety point of view (reduced handling/processing) to investigate methods that either prevent or slow down the rate of the corrosion reaction of the metal. This can be achieved by alloying with a suitable quantity of gallium [1] or via the formation of a surface film of plutonium oxycarbide, PuO_xC_y (where $x+y \leq 1$) [2]. Plutonium oxycarbide films have the NaCl structure and were initially called plutonium monoxide, PuO . Plutonium oxycarbide films are formed from the bulk to surface diffusion of the ubiquitous carbon impurity in the metal [3]. However, to date, there is little understanding of the electronic properties of plutonium oxycarbide films. The aim of this work is to investigate the electronic structure of plutonium oxycarbide surface films using X-ray photoelectron spectroscopy (XPS), ultraviolet photoelectron spectroscopy (UPS) and inverse photoemission spectroscopy (IPES). XPS measurements have been acquired at elevated temperature following the Pu 4f, O 1s and C 1s / Pu 5p_{1/2} spectral regions on both alloyed and pure plutonium samples following sputter cleaning and oxidation at room temperature. UPS and IPES were acquired at room temperature following the formation of the surface plutonium oxycarbide film. All three spectroscopic measurements display a two peak structure consistent with a partially localized and itinerant 5f electronic structure for plutonium oxycarbide. Finally, the Auger parameter (Pu N₇O₅V - Pu 4f_{7/2}) for plutonium oxycarbide films of varying carbon and oxygen stoichiometries are compared to the recently reported values for plutonium metal and the homoleptic plutonium oxides [1].

[1] XPS characterization of a PuGa-7 at. % alloy, P. Roussel, S. C. Hernandez, J. J. Joyce, K. S. Graham, T. Venhaus, J. Vac. Sci. Technol. A 41, 023204 (2023).

[2] Retardation of plutonium oxidation by a PuO surface film, D. T. Larson, D. L. Cash, J. Vac. Sci. Technol. 9, 800 (1972).

[3] Initial studies of plutonium corrosion, P. Roussel, D. S. Shaw, D. A. Geeson, J. Nucl. Sci. Technol. 39:sup3, 78 (2002).

5:20pm **AC+MI+TH-TuA-10 Exploring the Surface and Subsurface Behavior of Hydrogen in δ -Pu(100) and Bulk δ -Pu Through Density Functional Theory**, Charles Fricke, S. Hernandez, Los Alamos National Laboratory

The corrosion of plutonium to form its hydride creates significant challenges in handling, researching, and storing of the metal. Plutonium hydrides rapidly, much faster than most metals, and the kinetics are less well understood. Significant questions remain, including the energetics of hydrogen formation in different lattice sites, such as the tetrahedral, octahedral, and vacancy sites, as well as its reactive behavior on and in the surface of metallic plutonium. As such, this work aims to explore how hydrogen behaves within face-centered-cubic δ -plutonium from the surface into the subsurface and into the bulk through density functional theory. We find that hydrogen is energetically favored to remain on the surface, but that the energetics of hydrogen in the first subsurface remains thermodynamically favorable for both the subsurface tetrahedral and octahedral sites when compared to the referenced gas-phase hydrogen molecule. We also find that the octahedral site remains thermodynamically favorable for hydrogen species to fill from the 2nd subsurface to the bulk in δ -Pu(100). In addition, we find we have subsurface to bulk formation energy convergence at the 3rd sublayer of δ -Pu(100). Finally, we will discuss the energetics differences of hydrogen between the surface, subsurface, and bulk within δ -Pu.

5:40pm **AC+MI+TH-TuA-11 Nanoscale Nuclear Materials: Synthesis and Advanced X-Ray Characterization of Uranium Oxide Nanoparticles**, Liane Moreau, Washington State University, US

Nanostructures (particularly with sizes below 10 nm) are inherently challenging to characterize on the atomic scale, due to broadening which occurs in diffraction-based characterization methods, and the high concentration of surface defects and energy-minimization effects. Characterization challenges compound when investigating nanoscale actinide oxides, such as uranium oxide, due to radioactive sample constraints and rich electronic structure which can potentially stabilize a wide range of crystallographic arrangements. To address these challenges, x-ray spectroscopic and scattering based methods are used to probe challenging nanoscale systems in detail over multiple length scales. In particular, synthetic developments are paired with robust characterization, in order to improve knowledge of nanoparticle growth and transformation,

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as well as develop ways to make advanced x-ray characterization analysis more informed and accessible, in tandem.

Nanoscale uranium oxides are of interest towards the development of accident-tolerant nuclear fuels and are also relevant to the migration of actinides in environmental systems. Synthetic methods will be presented, which have enabled preliminary investigation into the size and morphology-dependent properties of uranium oxides. Specifically, the synthesis of UO₂ anisotropic nanoparticles and the deposition of uranium onto iron nanoparticles will be highlighted. These synthetic pursuits have been paired with x-ray spectroscopy and x-ray scattering to create new analysis strategies for decoupling surface vs. interior chemistry of nanoparticles and to interrogate the in-depth structural and electronic properties of uranium oxide as a function of particle size and morphology. The studies presented give a glimpse into the interesting fundamental and behavioral differences between actinide nanomaterials and their counterparts elsewhere on the periodic table.

6:00pm **AC+MI+TH-TuA-12 A Statistical Mechanics Treatment of Multiconfigurational Ground States in Isolated Neutral Atoms, Miles Beaux**, Los Alamos National Laboratory

Janoschek, et al. described the ground state of plutonium (Pu) as being, “governed by valence fluctuations, that is, a quantum mechanical superposition of localized and itinerant electronic configurations.” [*Science Advances*, 1, 6 (2015)]. However, a casual internet search of the electronic structure of Pu gives [Rn]7s²5f⁶ which is wholly insufficient to describe the complex nature of its multiconfigurational ground state as described by Janoschek, et al. A better description might be given by

$$[\text{Rn}]7s^{2+\gamma_{s,\text{Pu}}} 5f^{6+\gamma_{f,\text{Pu}}} 6d^{0+\gamma_{d,\text{Pu}}} 7p^{0+\gamma_{p,\text{Pu}}}$$

where $\gamma_{s,\text{Pu}}$, $\gamma_{f,\text{Pu}}$, $\gamma_{d,\text{Pu}}$, $\gamma_{p,\text{Pu}}$ represent potentially non-integer deviations from the integer electron occupancies of their respective orbitals. By judiciously selecting bounding conditions for how the eight valence electrons can distribute themselves within the given orbitals, a limited number of occupancy configurations can be identified with an identifiable number of cumulative allowable electron permutations. For example, the bounding conditions

$$-2 \leq \gamma_{s,\text{Pu}} \leq 0,$$

$$-2 \leq \gamma_{f,\text{Pu}} \leq 2,$$

$$0 \leq \gamma_{d,\text{Pu}} \leq 2,$$

$$\gamma_{p,\text{Pu}} = 0, \text{ and}$$

$$\gamma_{s,\text{Pu}} + \gamma_{f,\text{Pu}} + \gamma_{d,\text{Pu}} + \gamma_{p,\text{Pu}} = 0$$

result in nine possible occupancy configurations

$$7s^2 5f^6 6d^0 7p^0, 7s^2 5f^5 6d^1 7p^0, 7s^2 5f^4 6d^2 7p^0, 7s^2 5f^3 6d^3 7p^0, 7s^2 5f^2 6d^4 7p^0, 7s^2 5f^1 6d^5 7p^0, 7s^2 5f^0 6d^6 7p^0, 7s^1 5f^6 6d^0 7p^0, 7s^0 5f^6 6d^0 7p^0$$

with 487,630 cumulative possible electron permutations. With the specific number of permutations identified, principles of statistical mechanics can be applied to multiconfigurational ground states.

Similar generalized representations of ground state electronic structures will be described for each element in the periodic table. Classical Boltzmann’s entropies for the respective electronic structures will be calculated by considering the possible electron permutations as statistical mechanics microstates. These entropies will then be compared to known standard molar entropies.

LA-UR-23-24803

Wednesday Afternoon, November 8, 2023

Actinides and Rare Earths Focus Topic Room C124 - Session AC+AS+MI+TH-WeA

Emerging Topics and Methods in Actinide/Rare Earth Sciences

Moderators: Edgar Buck, PNNL, Krzysztof Gofryk, Idaho National Laboratory, Liane Moreau, Washington State University

2:20pm **AC+AS+MI+TH-WeA-1 Chemical Imaging and Applications Using High Energy Resolution Fluorescence Detection for the Actinides, Samuel Webb, N. Edwards, V. Noel**, SLAC National Accelerator Laboratory **INVITED**
Microscale synchrotron radiation-based x-ray fluorescence (SR-XRF) chemical analyses can provide a unique capability for chemical signature recognition and classification capabilities for actinide micro-particle analysis. SR-XRF is well suited to forensic type analyses of small particles because it is rapid, non-destructive, highly sensitive, has good spatial resolution, and can provide chemical information on the elements that are present when combined with x-ray absorption spectroscopy (XAS). The combination of spatially resolved distribution and chemical information, often known as chemical imaging, effectively provides a “chemical morphology” of the sample of interest and can show how chemical states are distributed within and among a series of particles. This type measurement is critical for understanding particle origin and history, as the spectroscopy, and its spatial distribution, can provide unique and complementary chemical signatures that may not be elucidated with other forms of measurement.

However, the conventional XAS capability in the near edge region as commonly implemented is often inadequate for systems that require high sensitivity or require a higher detail of spectroscopic information. This can be overcome with the combination of traditional micro SR-XRF and XAS, integrated with a high energy resolution fluorescence detector (HERFD) crystal analyzer. This has been recently implemented at BL 6-2b at SSRL and applied in the determination of the micron-scale oxidation state of uranium in particles. A discussion of the image and data processing techniques that can be applied using spatially resolved HERFD to obtain chemical and structural information, as well as the distribution of phases across different particles at the micro-scale, will also be presented.

3:00pm **AC+AS+MI+TH-WeA-3 High Energy X-Ray Characterization of Microstructure at Macroscopic Depths in Pu Alloys, Donald Brown**, Los Alamos National Laboratory; *T. Carver, R. Pokharel*, Los Alamos National Laboratory; *A. Smith, J. Park*, Argonne National Laboratory **INVITED**

High energy (95keV) X-rays were utilized to characterize the microstructure of six PuGa alloy samples. The samples had different gallium content, age, and history (i.e. thermos-mechanical processing). The crystallographic textures, phase fractions, lattice parameters and dislocation densities of each sample were evaluated through distinct analysis techniques. The textures of the samples were modest. In each case, the dislocation densities were relatively high, comparable to cold worked metals, with some attributable to mechanical work and some likely due to self-irradiation damage (aging). The lattice parameters determined in the predominantly single-phase samples were larger than expected based on the nominal Ga concentration. The lattice parameter increase is attributed to self-irradiation induced Ga segregation leaving the matrix Ga lean. Finally, measurements of the lattice parameter were completed as a function of temperature to determine the crystallographic thermal expansion of the material.

4:20pm **AC+AS+MI+TH-WeA-7 Native Oxide Growth of Pu—Ga Alloys, Kasey Hanson, S. Hickam, D. Olive, A. Pugmire**, Los Alamos National Laboratory

Plutonium corrosion has proven to be an intricate area of research that garners considerable attention. In particular, understanding plutonium oxidation is significant to the safe handling and storage of plutonium metal, which is known to readily oxidize under ambient conditions. In order to mitigate this, plutonium metal is commonly stored in inert atmospheres. These include gloveboxes as well as ultra-high vacuum (UHV) chambers used for materials characterization.

The objective of this work is to measure oxidation rates of plutonium metal in various storage environments. To achieve this, spectroscopic ellipsometry was selected due to its prevalence in a variety of industries to measure thin-film thickness. A non-destructive technique, ellipsometry exposes the sample to polarized light, and through changes in the amplitude ratio and phase difference of the reflected vs. incident light, one can generate a material's optical properties that can be used to determine oxide thicknesses.

Building upon previous work, this presentation will present the optical constants of plutonium metal, achieved through the use of ion sputtering under UHV, and plutonium oxide formed on plutonium metal in various storage and exposure conditions. These results will then be used to model oxide thickness values directly from ellipsometry measurements as well as provide additional insights into the nature of the oxide films relevant to plutonium storage and handling.

4:40pm **AC+AS+MI+TH-WeA-8 Molecular Beam Epitaxy of Ternary Nitrides: From Transition Metals to Actinoids, Kevin Vallejo**, Idaho National Laboratory; *S. Gutierrez Ojeda*, Universidad Nacional Autonoma de Mexico; *G. Hernandez Cocoltzi*, Benemerita Universidad Autonoma de Puebla, Mexico; *S. Zhou, K. Gofryk, B. May*, Idaho National Laboratory

The exceptional properties of transition metal nitrides for coatings, as well as photo- and electrochemical applications have opened the doors for more detailed studies of their structure and synthesis. Mn- and Cr- nitrides are strong candidates to showcase the application of spintronic and magnetic sensing applications partially because of their ability to form several different metastable phases. Similarly, rare-earth nitrides have been studied for over 50 years due to their optical properties and strong electron correlations, leading to behaviors that range from insulating to metallic. This study uses molecular beam epitaxy to synthesize epitaxial thin films of different Mn-, Cr-, Ga-, and Ce-nitride phases. The electrical and magnetic properties of these films are investigated with the rocksalt MnN and CrN both showing metallic behavior, with the latter showing a magnetic transition ~280K. However, when combining these materials at similar growth conditions, instead of maintaining the rocksalt structure, a new ternary cubic phase of Mn_xCr_yN is obtained which shows narrow-gap semiconducting behavior. Additionally, density functional theory calculations show potential new phases where lanthanoid ternary phases are experimentally realizable. This work presents an avenue for the epitaxial integration of metallic, magnetic, and semiconductor materials via transition metal and lanthanoid compounds, en route to the synthesis of actinoid compounds.

5:00pm **AC+AS+MI+TH-WeA-9 Formation and Electronic Properties of Uranium Tellurides: A Thin Films Study, Evgenia Tereshina-Chitrova, S. Alex**, Institute of Physics CAS, Prague, Czechia; *O. Koloskova, L. Horak*, Charles University, Prague, Czechia; *O. Romanyuk, Z. Soban*, Institute of Physics CAS, Prague, Czechia; *T. Gouder, F. Huber*, JRC Karlsruhe, Germany

Understanding the interaction of 5f electrons with other electrons in actinide compounds presents a significant challenge. Factors such as relativistic effects, strong electron correlations, and hybridization with ligands contribute to the complexity of observed effects. This complexity is exemplified in the case of the recently discovered unconventional superconductor UTe_2 [1]. The ongoing debate surrounding UTe_2 revolves around understanding its unconventional superconductivity, the nature of its electronic correlations [2,3], and the character of the 5f electrons. While it's evident that UTe_2 exhibits intermediate occupancy of the 5f states at ambient pressure, there are two opposing interpretations of experimental findings regarding the degree of localization of the 5f electrons: the model based on $5f^26d^1$ states with singlet crystal field states [4,5], and a $5f^3$ -based model with atomic Kramers doublet and greater 5f itinerancy [6,7]. Interestingly enough, ab-initio calculations considering the $4f^2$ U-ion ground state multiplet reproduce bulk experimental observations, such as the Schottky anomaly and magnetic anisotropy at low temperatures [8].

To contribute to this subject, we provide a comprehensive and detailed investigation of thin films of uranium tellurides with various compositions U_xTe_y by means of X-ray Photoelectron Spectroscopy (XPS) and Ultra-violet Photoelectron Spectroscopy (UPS). We also explore the structural and bulk

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properties of the films, including their magnetic and transport characteristics. This comprehensive approach allows us to correlate the changes in XPS spectra across different compositions and crystal structures with resultant bulk properties, contributing to our understanding of the nature of the 5f electrons within uranium tellurides.

The work by E.A. T.-Ch. is supported by the Czech Science Foundation (GAČR) under the grant number 22-19416S. E.A. Ch.-T. is a recipient of the L'Oreal-UNESCO for Women in Science prize.

- [1] S. Ran et al., *Science* 365, 684 (2019).
- [2] I. M. Hayes et al., *Science*, 373, 797 (2021).
- [3] C. Duan et al., *Phys. Rev. Lett.* 125, 237003 (2020).
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- [5] S. Liu et al., *Phys. Rev. B* 106, L241111 (2022).
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- [8] S. Khmelevsyi et al., *Phys. Rev. B* 107, 214501 (2023).

5:20pm AC+AS+MI+TH-WeA-10 Nanoscale Uranium and Thorium Oxides for Applications in Advanced Nuclear Fuels, *Matthew Heaney, L. Moreau, X. Guo*, Washington State University

Nanomaterials of uranium and thorium oxides and their mixtures are of significance to the development of advanced nuclear fuels with properties such as self-healing and greater fission gas retention. They also serve as a means for furthering our understanding of spent nuclear fuel behaviors. Furthermore, thorium incorporation into the uranium sublattice induces changes in oxidation behavior, which would be a promising means to provide additional redox resistance. Coupling this with the possible application in fast neutron reactors presents a potential fuel for the Th fuel cycle, which is inherently proliferation resistant. However, these possible applications are purely hypothetical given that, compared to nanomaterials from the rest of the periodic table, actinide nanomaterials are heavily understudied. Therefore, it will be imperative to interrogate the properties of these nanomaterials. This work aims to explore three key areas. (1) How careful kinetic control through variations in synthetic methods can be used to produce particles with different sizes, shapes, and heterogeneities of mixing. (2) Elucidation of the structural and thermodynamic properties of these nanomaterials and how characteristic size, shape, and heterogeneity of mixing affect them through a variety of X-ray characterization and calorimetry techniques. X-ray characterization techniques such as X-ray absorption spectroscopy (XAS) is used to evaluate the heterogeneity of mixing, local structure, and oxidation state of particles while small-angle X-ray scattering (SAXS) is used to determine statistical values on particle size, shape, and polydispersity. Calorimetry techniques like high temperature drop calorimetry are used to determine enthalpies of formation for nanoparticles and the contribution surface effects have on the thermodynamics or thermogravimetric analysis-differential scanning calorimetry (TGA-DSC), which can be used to evaluate oxidation behaviors. (3) How sintered nanoparticles behave in *in-situ* reactor conditions as nuclear fuels and how particle characteristics can be chosen to achieve more desirable radiation-resistance and thermal properties for nuclear fuel, through irradiation studies performed at the 1-MW TRIGA reactor on the campus of Washington State University. Overall, the work presented provides initial evidence for the benefits of nanomaterials in the production of advanced reactor fuels and outlines important fundamental properties of actinides on the nanoscale.

Magnetic Interfaces and Nanostructures Division

Room B110-112 - Session MI+2D+TF-WeA

Special Symposium on Coupling Phenomena in Magnetism

Moderator: Hendrik Ohldag, Lawrence Berkeley National Laboratory

2:20pm MI+2D+TF-WeA-1 Coupling Spin-Orbit and Exchange Interaction in a Low-Dimensional Magnet, *Pascal Jona Grenz¹, M. Donath, P. Krüger*, University of Münster, Germany

Coupling exchange interaction (SOI) and spin-orbit interaction (XCI) provides the foundation for many prospective spin-based information technology applications. For example, it was suggested that the strength of SOI at a ferromagnet/heavy-metal (FM/HM) interface is decisive for the efficiency of writing magnetic bits in spin-orbit-torque MRAM devices [1].

Exploring the electronic structure is the key to access the factors underlying the coupling of SOI and XCI. We use spin- and angle- resolved inverse photoemission to study the interplay of SOI and XCI in the unoccupied electronic structure at the interface of a low-dimensional FM on a HM substrate. The prototypical FM/HM hybrid system Ni/W(110) exhibits exchange-split Ni-related states that become strongly influenced by SOI. A balanced ratio of SOI and XCI results in a magnetization- and k-dependent quenching or enhancement of the spin splitting. This remarkably large interplay of SOI reflected in the adlayer states contrasts previous studies of the occupied electronic structure of the same system, where the observations were attributed solely to either SOI or XCI [2].

Using density-functional-theory (DFT) calculations, we investigate the underlying drivers responsible for the experimentally observed coupling of SOI and XCI. We find that hybridization between adsorbate and substrate states, along with the strongly localized wave functions at the heavy W nuclei, cause the strong influence of SOI within the Ni-related exchange-split states.

[1] I.M. Miron *et al.*, *Nature*, **476** 189 (2011)

[2] A. Nuber, PhD Thesis (University of Würzburg, Germany, 2011)

2:40pm MI+2D+TF-WeA-2 AVS Graduate Research Awardee Talk: Temperature Dependent Magnetic and Electronic Properties of NiCo₂O₄ Thin Film Surfaces, *Arjun Subedi^{1,3}*, University of Nebraska-Lincoln; *D. Yang, C. Mellinger, X. Xu*, University of Nebraska-Lincoln; *P. Dowben*, University of Nebraska-Lincoln

Although NiCo₂O₄ thin film is shown to possess perpendicular magnetic anisotropy [1], we have observed in-plane spin polarization of NiCo₂O₄ thin film in spin polarized inverse photoemission spectroscopy (SPIES). The unoccupied states of NiCo₂O₄, above Fermi level, were observed to have unequal density of states for spin majority and spin minority carriers in SPIES, and the spectra obtained from the SPIES have spectral features that can be compared to the XMCD spectra. The in-plane spin polarization of NiCo₂O₄ is found to decrease with increasing temperature, as expected. In addition to the temperature dependent change in spin polarization, we observed that there is change in the surface electronic properties of NiCo₂O₄ from conducting to insulating when the temperature is increased. X-ray photoemission spectroscopy (XPS) studies show that there exist no appreciable binding energy changes of Ni 2p_{3/2} and Co 2p_{3/2} core levels with change in temperature (T) when the NiCo₂O₄ film exists in conducting phase. However, when the NiCo₂O₄ films became insulating, the core level binding energies changed reversibly with change in temperature during annealing and cooling cycles. The core level binding energy (BE) change with temperature (T) is found to closely follow a modified Arrhenius type model. The proposed model is also followed by Co 2p_{3/2} and Fe 2p_{3/2} core levels in temperature dependent XPS of insulating CoFe₂O₄ thin films. Our studies indicate that thermal effects and oxygen defects should play the roles in changing both magnetic and electronic properties of NiCo₂O₄ thin films with temperature.

[1] C. Mellinger et al., *Phys. Rev. B* 101, 014413 (2020).

¹ Falicov Student Award Finalist

² AVS Graduate Research Awardee

³ Falicov Student Award Finalist

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3:00pm **MI+2D+TF-WeA-3 Antiferromagnetic Real-Space Configuration Probed by Dichroism in Scattered X-Ray Beams with Orbital Angular Momentum**, *Sophie Morley*, *M. McCarter*, *A. U. Saleheen*, *A. Singh*, Lawrence Berkeley Lab; *R. Tumbleson*, University of California Santa Cruz; *J. Woods*, Argonne National Laboratory; *A. Tremsin*, UC Berkeley; *A. Scholl*, Lawrence Berkeley Lab; *L. de Long*, *J. Hastings*, University of Kentucky; *S. Roy*, Lawrence Berkeley Lab

INVITED

X-ray beams with orbital angular momentum (OAM) are a promising tool for x-ray characterization

techniques. Beams with OAM have an azimuthally varying phase, and new material properties can

potentially be probed by utilizing this unique phase structure. Here, we show how OAM beams are

created through resonant diffraction from an artificial antiferromagnet with a topological defect. The

scattered OAM beams have circular dichroism whose sign is coupled to the phase of the beam [1]. Using

magnetic scattering calculations, we show that this dichroism is related to the real-space configuration

of the antiferromagnetic ground state. Thermal cycling of the artificial antiferromagnet can change the

ground state, as indicated by the changing dichroism. These results exemplify the potential of OAM

beams to probe matter in a way that is inaccessible using currently available x-ray techniques.

[1] M. R. McCarter et al., Phys. Rev. B **107**, L060407 (2023)

5:00pm **MI+2D+TF-WeA-9 Spin-dependent Hybridization of Image-potential States with TI States in TI/Ag(111)**, *Sven Schemmelmann*¹, Universität Münster, Germany; *P. Härtl*, Universität Würzburg, Germany; *P. Krüger*, Universität Münster, Germany; *M. Bode*, Universität Würzburg, Germany; *M. Donath*, Universität Münster, Germany

The BiAg₂ and PbAg₂ surface alloys exhibit giant Rashba splittings [1-4]. The related TI/Ag₂ surface alloy is expected to show states with smaller but still large Rashba splittings due to the lower atomic number of TI. However, this alloy forms only small patches with long-range structural order [5]. For higher coverages of TI on Ag(111), a smooth and stable wetting layer forms with a moiré superstructure due to the lattice mismatch between TI and Ag. We present a study of the unoccupied electronic structure of this superstructure by spin- and angle-resolved inverse photoemission. The experimental results are accompanied by DFT calculations. We observe surface states and an image-potential resonance located several Å in front of the surface. Surprisingly, one surface state exhibits almost no Rashba splitting even though it is located around the TI atoms. This result is explained by the orbital symmetry of the respective state. For the image resonance, we find hybridization with a downward dispersing TI state leading to spin-dependent spectral intensities that vary strongly close to the hybridization point. This observation, both in experiment and bandstructure calculations, is supported by charge distribution calculations showing an expeditious change from the image resonance to the TI-induced surface state depending on k_{\parallel} .

[1] C. R. Ast et al., Phys. Rev. Lett. **98**, 186807 (2007)

[2] G. Bihlmayer, S. Blügel, and E. V. Chulkov, Phys. Rev. B **75**, 195414 (2007)

[3] S. Wissing et al., Phys. Rev. Lett. **113**, 116402 (2014)

[4] L. El-Kareh et al., New J. Phys. **16**, 045017 (2014)

[5] P. Härtl, S. Schemmelmann, P. Krüger, M. Donath, and M. Bode, submitted to Phys. Rev. B

5:20pm **MI+2D+TF-WeA-10 Distinct Tamm and Shockley Surface States on Re(0001) Mixed by Spin-Orbit Interaction – A Photoemission Study**, *Marcel Holtmann*, *P. Krüger*, University of Münster, Germany; *K. Miyamoto*, *T. Okuda*, HiSOR, Japan; *P. Grenz*, University of Münster, Germany; *K. Shimada*, HiSOR, Germany; *M. Donath*, University of Münster, Germany

Tamm and Shockley states, these two paradigmatic concepts are used to describe surface states not only in electronic systems but also in photonic and phononic crystals. The Re(0001) surface was found to host both types of electronic surface states in neighboring, but qualitatively different gaps [1]. Interestingly, spin-orbit interaction generates a double W-shaped energy vs k_{\parallel} dispersion by mixing both types of states and lifting their spin degeneracy. We employ a combination of spin- and angle-resolved photoemission, tight-binding model calculations, and density functional theory that accounts for the photoemission process to establish reliable criteria for distinguishing between the two types of surface states. Our approach leads to a coherent understanding of the mechanism of spin-orbit interaction in such a situation.

From a detailed investigation of the Re(0001) surface along the ΓM and ΓK high-symmetry directions [2], we obtain Rashba parameters of 0.32 and 0.34 eVÅ along the two respective directions. This indicates a slight warping of the surface state. Regarding the aforementioned state's spin polarization: at variance with theoretical predictions for a perfect hcp(0001) of rhenium [3], we do not find any out-of-plane spin polarization. This is attributed to monatomic steps of a real Re(0001) surface with alternating terminations, leading on average to an effective sixfold surface symmetry and vanishing net out-of-plane spin polarization.

[1] M. Holtmann et al., Phys. Rev. B **105**, L241412 (2022)

[2] M. Holtmann et al., Phys. Rev. B **107**, 165420 (2023)

[3] A. Urru and A. Dal Corso, Surf. Sci. **686**, 22 (2019)

5:40pm **MI+2D+TF-WeA-11 Coupling between Spin Order and Orbital Order in 2D-Superlattice Perovskite Film**, *Bin Hu*, University of Tennessee Knoxville

INVITED

The coupling between spin order and orbital order presents a fundamental request to develop advanced multifunctional materials. 2D-superlattice perovskite films, known as solution-processing semiconductors, possess strong orbital order within non-degenerate Rashba band structures under the concurrent influence of spin-orbital coupling and symmetry breaking. This provides a fundamental condition to dynamically couple spin order and orbital order through multiferroic interface design. Here, we combine ferroelectric 2D-superalattice perovskite (4,4-DFPD₂PbI₄) film and ferromagnetic cobalt (Co) film to form multiferroic perovskite/Co interface. By using this multiferroic interface design, we found that the circularly polarized orbitals with right and left handedness (s^+ and s^-) in Rashba band structures can selectively interact with spin-up and spin-down spin dipoles on the Co surface, leading to a mutually selectivity between spin order and orbital order. Particularly, this selective interaction between spin order and orbital order can enable spin-switchable phenomena towards developing emerging functionalities in these solution-processing hybrid metal halide perovskites. When the ferromagnetic spins on the Co surface are altered between positive and negative magnetic field directions (+**B** and -**B**), the circularly polarized luminescence (CPL) in 2D-superlattice perovskite can be switched between s^+ and s^- polarizations, leading to spin-switchable phenomena at room temperature. More interestingly, our polarized neutron reflectometry (PNR) studies found that circularly polarized photoexcitation generates a static magnetization within 2D-superlattice perovskite film prepared on the Co surface. This presents an optically induced magnetization phenomenon. Essentially, this optically induced magnetization reveals a long-range coupling between the spin order on the Co surface and the orbital order within Rashba band structures in 2D-superlattice perovskite film. This presentation will discuss the fundamental coupling between spin order and orbital order through Rashba band structures in 2D-superlattice perovskite film.

¹ Falicov Student Award Finalist

Light Sources Science Mini-Symposium Room C124 - Session LS+AC+LX+MI+TH-ThM

Tender X-ray Science and Time Resolved Studies

Moderators: Alison Pugmire, LANL, David Shuh, Lawrence Berkeley National Laboratory, James G. Tobin, University of Wisconsin-Oshkosh

8:00am **LS+AC+LX+MI+TH-ThM-1 Developments of High Resolution X-Ray Spectroscopic Tools for Probing Structural Properties of Actinide System from the Metal and Ligand Perspective, Tonya Vitova**, Karlsruhe Institute of Technology, Institute for Nuclear Waste Disposal, Germany **INVITED**

High energy resolution X-ray absorption and emission spectroscopic techniques became indispensable methods in actinide and radionuclide research.¹⁻⁵ One important motivation is studies concerning the mobilization and retention of long-lived actinides and fission products in geochemical processes relevant for safety studies of a potential deep geological nuclear waste repository.³⁻⁴ In-depth insights into the actinide-ligand binding properties is a main application of these novel experimental techniques too.⁶ Development at the ACT experimental station of the CAT-ACT wiggler beamline at the Karlsruhe Institute of Technology (KIT) Light Source will be discussed. One experimental technique especially powerful to differentiate oxidation states of actinides (An) is the An M_{4,5}-edge high-energy resolution X-ray absorption near-edge structure (HR-XANES).⁴ This presentation highlights the latest technological developments at the ACT station enabling the HR-XANES spectroscopic technique for samples with low radionuclide loading down to 1 ppm in combination with a cryogenic sample environment reducing beam-induced sample alterations.⁷⁻⁸ It paves the way for the examination of coupled redox/solid-liquid interface reactions.⁸ Examples of applications of An M_{4,5} edge core-to-core and valence band resonant inelastic X-ray scattering (CC-RIXS and VB-RIXS) for probing the electronic structure and binding properties of the actinide elements will be illustrated.⁶ First results obtained using a newly developed versatile chamber for soft X-ray spectroscopy at the X-SPEC beamline at the KIT Light Source will be discussed.

This work has received funding from the European Union's Horizon 2020 research and innovation program under grant agreement No. 847593. We also acknowledge funding from the ERC Consolidator Grant 2020 under the European Union's Horizon 2020 research and innovation program (grant agreement No. 101003292).

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4. Pidchenko, I. et al., *Environ Sci Technol* **2017**,51 (4), 2217-2225.
5. Bagus, P. S. et al., *Inorganic Chemistry* **2021**,60 (21), 16090-16102.
6. Vitova, T. et al., *Chem Sci* **2022**,13 (37), 11038-11047.
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8. Schacherl, B. et al., *Anal Chim Acta* **2022**,1202, 339636.

8:40am **LS+AC+LX+MI+TH-ThM-3 High-Energy-Resolution X-Ray Spectroscopy and Actinides Research at SLAC, Dimosthenis Sokaras**, SLAC National Accelerator Laboratory **INVITED**

Nowadays, high-energy-resolution x-ray spectroscopy is a well-established and powerful tool available in state-of-the-art synchrotron facilities. The suppression of the core-hole lifetime contribution within the conventionally broad spectroscopic features of actinide series has revitalized the role of x-ray spectroscopy in the study of actinide complexes and intermetallics. Numerous studies have leveraged the fine structure of M or L absorption edge resonances to sensitively probe and quantify the oxidation state, 5f delocalization, and ligation of the actinides species. The increasing availability of large solid angle instruments coupled with high flux beamlines is quickly enabling such advanced studies for dilute samples or samples under special sample environments. In this presentation we will summarize the high-resolution tender and hard x-ray spectroscopy advances at SLAC and the actinides research program that these capabilities have enabled during the last decade.

9:20am **LS+AC+LX+MI+TH-ThM-5 New Insight Into Excited-State Chemical Dynamics Using Ultrafast X-Rays:Recent Highlights, Future Opportunities & Development Plans at LCLS, Robert Schoenlein**, Linac Coherent Light Source - SLAC National Accelerator Laboratory **INVITED**

Ultrafast X-rays from free-electron lasers (XFELs) are driving a qualitative advance in our understanding of condensed-phase chemical dynamics and catalysis. Ultrafast soft X-rays provide element-specific mapping of chemical bonds, charge distributions, oxidation states and frontier orbitals. Ultrafast hard X-ray pulses reveal the atomic scale structural dynamics of excited-state dynamics – revealing relaxation pathways, and the coupling of atomic structure, electronic structure, and solvent dynamics. This talk will highlight recent results from the Linac Coherent Light source (LCLS) using advanced ultrafast X-ray methods to track excited-state charge-transfer and relaxation pathways, and reveal the influence of molecular structural dynamics, and solvent coupling. Notably, multi-modal methods combining time-resolved X-ray scattering and spectroscopy represent a powerful approach for linking X-ray experimental observables with theory to achieve a deeper understanding of excited-state dynamics to advance the development of design principles for creating molecules, complexes, and assemblies with desired functions.

In addition, new science opportunities enabled by the nearly-completed upgrade of LCLS (LCLS-II) coupled with advanced instrumentation and methods will be discussed. LCLS-II will provide tunable soft X-ray pulses (0.25 to 5.0 keV) at high repetition rate (up to 1 MHz) and hard X-rays up to 25 keV (at 120 Hz). This unprecedented capability will support powerful new methods such as time-resolved resonant inelastic X-ray scattering (RIXS). The new ChemRIXS instrument is optimized for studying solvated complexes with C, N, O (K-edges), 3d transition metals (L-edges), and rare-earth elements (M-edges) – where 2D RIXS maps of excited-state dynamics coupled with quantum chemical calculations will reveal the evolution of frontier orbitals. The Tender X-ray Instrument (TXI, 2.1-5.0 keV), now under development for LCLS-II, will support time-resolved tender X-ray spectroscopy (spanning the 4d transition metal L-edges and key functional ligands including P, S, and Cl), coherent scattering, and novel nonlinear X-ray pump / X-ray probe methods - combining X-rays from two independently tunable XFEL sources.

11:00am **LS+AC+LX+MI+TH-ThM-10 Attosecond Studies of Radiolysis at XFELs, Linda Young**, Argonne National Laboratory **INVITED**

We report the first attosecond x-ray pump/x-ray probe transient absorption study in condensed phases using a pure liquid water target. With tunable two-color attosecond x-ray pulses, the pump ionizes the valence band of water and the probe scans the oxygen K-edge absorption region. Theory establishes the nature of the detected transient absorption and models the observed signal for sub-femtosecond delay times.

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critical importance for fields ranging from cancer therapy to the longevity of nuclear reactors to space travel. In these applications, radiolysis is initiated by a high-energy particle that leads to the ejection of energetic primary electrons followed by inelastic and non-adiabatic processes that produce damaging low energy electrons and reactive radical species. A microscopic understanding of reaction mechanisms, especially in complex systems, is missing as typical techniques used to detect prominent species, EPR and UV spectroscopies, lack either time resolution or spectral clarity. Tunable ultrafast x rays can dissect the radiolysis process. That is, x-ray pump/x-ray probe studies can systematically either peel electrons from valence, or eject them from core orbitals and follow the ensuing dynamics on a site-specific basis.

11:40am **LS+AC+LX+MI+TH-ThM-12 First Real-Time Tracking of Oxidation States During Fast Redox of UO₂ Using a Microfluidic Electrochemical Cell and HR-XANES**, *Jennifer Yao*, Pacific Northwest National Laboratory; *B. Schacherl*, Karlsruhe Institute of Technology (KIT), Germany; *B. McNamara*, Pacific Northwest National Laboratory; *C. Vollmer*, Karlsruhe Institute of Technology (KIT), Germany; *N. Lahiri*, *E. Ilton*, *E. Buck*, Pacific Northwest National Laboratory; *T. Vitova*, Karlsruhe Institute of Technology (KIT), Germany

Real-time tracking of the oxidation states of a UO₂ electrode during electrochemical oxidation and reduction was achieved using operando high-resolution X-ray absorption near-edge structure (HR-XANES) spectroscopy at the ACT station of the CAT-ACT beamline at the KIT Light Source, Karlsruhe, Germany. This was made possible by utilizing a particle-attached microfluidic electrochemical cell (PAMEC) developed at PNNL, and employing KIT's advanced actinide M-edge HR-XANES technique.¹⁻² The PAMEC is a three-electrode system consisting of a working electrode (WE) made of the materials of interest a platinum (Pt) reference electrode, and a Pt counter electrode.³ The electrochemical analyzer connected to the PAMEC device controlled the redox process, e.g., applying constant potential on the UO₂ WE to reduce (-1.1 V vs Pt) or oxidize it (0.5 V vs Pt), while HR-XANES simultaneously scanned its surface chemistry. The U M₄-edge HR-XANES spectra revealed the evolution of U from U(IV) to U(V) and finally to U(VI) during the oxidation process. We were able to demonstrate the reversibility of this process by reducing the same electrode back to pure U(IV), as confirmed by HR-XANES. To our knowledge, this study reports the first in-situ and operando measurement of real-time oxidation state changes of UO₂. The spectra obtained also provided insight into the electronic structure of U(VI) in the UO₂ alteration process. This successful international scientific collaboration showcases the potential of a PAMEC for in-situ and operando experiments with UO₂ and highlights its promising broad application for characterization of spent nuclear fuel systems.

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- (3) Yao, J.; Lahiri, N.; Tripathi, S.; Riechers, S. L.; Ilton, E. S.; Chatterjee, S.; Buck, E. C., A microfluidic electrochemical cell for studying the corrosion of uranium dioxide (UO₂). *RSC Advances* **2022**,*12*, 19350-19358. 10.1039/D2RA02501A

12:00pm **LS+AC+LX+MI+TH-ThM-13 Use of Artificial Intelligence Techniques To Analyze Materials Characterization Data From Actinide Containing Materials**, *Jeff Terry*, Illinois Institute of Technology

We have developed artificial intelligence (AI) based methodology that can be utilized to reliably analyze experimental results from Extended X-ray Absorption Fine Structure (EXAFS), Nanoindentation, and core level photoemission. Specifically, we use a genetic algorithm to extract the relevant structural parameters through fitting of the measured spectra. The current approach relies on a human analyst to suggest a potential set of chemical compounds in the form of feff.inp input files that may be present. The algorithm then attempts to determine the best structural paths from these compounds that are present in the experimental measurement. The automated analysis looks for the primary EXAFS path contributors from the potential compounds. It calculates a goodness of fit value that can be used to identify the chemical moieties present. The analysis package is called EXAFS Neo and is open source written in Python. I will illustrate the use of this package with fits of actinide species in the barrier layer of Tristructural-isotropic (TRISO) encapsulated nuclear fuel particles. The current particle design consists of a two-phase uranium-oxide/uranium-carbide kernel of 19.74% ²³⁵U enrichment, a porous carbon buffer layer, and consecutive layers of pyrolytic carbon, silicon carbide (SiC) and pyrolytic carbon. The SiC layer provides the main barrier to fission product release. Much work has gone towards studying metallic fission product interaction in the SiC containment layer due to the propensity of metallic fission product release as a function of high temperature (safety) testing. Here, I will show how the interaction dynamics of plutonium and uranium within this layer have been determined through EXAFS measurements that have been fit with AI. One of the major benefits of using this technology is that actinide containing

materials often have edges from higher Z-elements that limit the usable range of the spectrum. Our method fits momentum space data which does not suffer from transformation artifacts of real space over a small momentum range.

**Magnetic Interfaces and Nanostructures Division
Room B110-112 - Session MI+2D+TF-ThM**

2D Magnetism and Superconductivity

Moderators: Markus Donath, Muenster University, Germany, Valeria Lauter, Oak Ridge National Laboratory

8:00am **MI+2D+TF-ThM-1 Heterostructures for Tunneling and Point-Contact Spectroscopy of Two-Dimensional Superconductors**, *Benjamin Hunt*, *Q. Cao*, Carnegie Mellon University; *E. Telford*, *C. Dean*, Columbia University

INVITED
Tunneling spectroscopy is an indispensable experimental tool of modern condensed matter physics. Vertical planar tunneling, which uses a fixed-width tunnel barrier, offers advantages over other spectroscopic tools such as scanning tunneling microscopy (STM). One such advantage is the ability to tunnel in reorientable and very large (≥ 40 T) magnetic fields at dilution refrigerator temperatures (≤ 30 mK), a capability that has application in, for example, determining the order parameter symmetry of novel two-dimensional (2D) superconductors. We demonstrate a novel vertical planar tunneling architecture for van der Waals heterostructures based on via contacts, namely, metallic contacts embedded into through-holes in hexagonal boron nitride (hBN). This via-based architecture overcomes limitations of other planar tunneling designs and produces high-quality, ultra-clean tunneling structures from a variety of 2D materials. The physical area of our via-based tunnel contacts is limited only by nanofabrication techniques, and we demonstrate a crossover from diffusive to point contacts in the small-contact-area limit by studying the spectrum of a 2D superconductor, NbSe₂. We show that our tunneling technique may enable highly-sought measurements of newly-discovered 2D superconductors such as monolayer 1T'-WTe₂, rhombohedral trilayer graphene, twisted trilayer graphene, and twisted bilayer BSCCO.

8:40am **MI+2D+TF-ThM-3 Ghost States and Topography Inversion in 2D Materials**, *Mina Yoon*, Oak Ridge National Laboratory, USA

INVITED
In this talk, I will discuss the challenges associated with characterizing the surface structures of single-atom thick materials, such as graphene and boron nitride, on metallic substrates or the surface of bulk systems, including quantum topological Kagome systems, using scanning tunneling microscopy (STM). The understanding of fundamental properties of two-dimensional (2D) materials and surface properties depends critically on the presence of "ghost" states, which arise due to different decay lengths in the wave function of the underlying layers and surfaces.

The existence of these ghost states, in conjunction with long-lived substrate states or underlying layers, plays a crucial role in interpreting and understanding the surface properties of 2D materials. These ghost states can originate from various sources, such as the bulk or the substrate, and can even arise from the boundary on the opposite side. The appearance of ghost states due to different decay lengths leads to unexpected results in surface structure measurements, including the intriguing phenomenon of topography inversion. Topography inversion refers to the counterintuitive result where the observed topography in STM images is opposite to the expected atomic geometry, as discussed in our recent study [1,2]. This inversion occurs as a consequence of the pervasive substrate states overshadowing the intrinsic states of the 2D materials. As a result, the measurement of the intrinsic properties of 2D materials becomes complicated, with the ghost and substrate states dominating the observed topography. To address these challenges, we employ a combination of first-principles density functional theory calculations and analytical modeling. Through our investigations, we demonstrate the critical role played by these ubiquitous substrate and ghost states in the observed topography inversion in STM images. By unraveling the influence of these states on STM measurements, we provide crucial insights for the accurate interpretation of STM topographies of atomically thin materials.

Our findings not only shed light on the phenomenon of topography inversion, but also contribute to the further development of 2D materials in (opto)electronic and quantum applications. Understanding and characterizing the ghost and substrate states is essential to unlock the full potential of 2D materials and enable their use in various technological advancements.

Thursday Morning, November 9, 2023

[1] "Spatially resolved on-dimensional boundary states in graphene-hexagonal boron nitride planar heterostructures", J. Park et al., Nat.Com. 5, 5403 (2014).

[2] "Topography inversion in scanning tunneling microscopy of single-atom-thick materials from penetrating substrate states", C. Park and M. Yoon, Sci. Reports 12, 7321 (2022).

11:00am **MI+2D+TF-ThM-10 Spatially-Resolved Photoemission Studies of Magnetic Weyl Semimetals**, S. Sreedhar, University of California, Davis; M. Staab, R. Prater, University of California at Davis; A. Rossi, Italian Institute of Technology, Italy; V. Ivanov, Lawrence Berkeley Lab; Z. Shen, University of California at Davis; G. Conti, Lawrence Berkeley Lab; V. Taufour, S. Savrasov, University of California at Davis; S. Nemsak, Lawrence Berkeley Lab; **Inna Vishik**, University of California-Davis **INVITED**

$\text{Co}_3\text{Sn}_2\text{S}_2$ is a magnetic Weyl semimetal below its Curie temperature (T_c) of 177K. I will discuss spatial and temperature-dependent angle-resolved photoemission spectroscopy (ARPES) and x-ray photoelectron spectroscopy (XPS) studies in this system. Across T_c , we observe signatures of a topological phase transition, but also observe changes in bulk bands which are inconsistent with a simple lifting of exchange interactions, suggesting enhanced electronic correlations in the regime without long-range magnetic order. I will also discuss spatial-dependent ARPES and XPS data which quantify the characteristic differences between Sn- and S-terminated surfaces, with relevance for interpreting surface-dominated phenomena.

11:40am **MI+2D+TF-ThM-12 High-Temperature Superconductor FeSe Films Enabled Through Temperature and Flux Ratio Control**, Maria Hilde, H. Yi, C. Chang, N. Samarth, The Pennsylvania State University; R. Engel-Herbert, Paul-Drude-Institut für Festkörperelektronik, Germany

FeSe, a bulk superconductor with a T_c of 9 K has attracted a high level of attention since a skyrocketing boost in TC was reported for a single unit cell (UC) layer of FeSe grown on $\text{SrTiO}_3(001)$ by molecular beam epitaxy (MBE) to as high as 100 K. FeSe- SrTiO_3 heterostructures have since been fabricated by many groups but the record TC proved difficult to reproduce and thus the mechanism behind it remains concealed. After extensive work in the past, the field appears to agree on certain key "ingredients" in the heterostructure sample preparation that are believed essential for the boost in TC. Those are; 1. an ultra-clean substrate surface of a double TiO_2 termination realized by a chemical and thermal *ex-situ* and/or thermal *in-situ* substrate preparation; 2. ultra-thin – one UC thickness – limit of FeSe; 3. a high number of Se vacancies in the FeSe film ensured through post-growth annealing steps in ultra-high vacuum (UHV) for several hours; 4. followed by a capping layer growth protecting FeSe against oxidation during *ex-situ* characterization.

We present our findings on FeSe thin film growth by MBE and present a roadmap for high- T_c – 222 % higher than the reported bulk value in *ex-situ* transport measurements – circumventing above mentioned steps 1, 2, and 3 by simple *in-situ* Se/Fe flux ratio and temperature control during FeSe growth. FeSe films of 20-UC-thickness grown at varying temperatures and Se/Fe flux ratios and the structural and morphological properties of the obtained uncapped FeSe films were analyzed. The morphology of the films showed a sensitive dependence on the growth temperature and flux ratio spanning from perfectly smooth and continuous films with atomic terraces at 450 °C growth temperature and a low flux ratio of 2.5 to exclusively disconnected island growth of large height but smooth top surfaces at lower temperatures and/or higher flux ratios. Surprisingly, the tetragonal $P4/nmm$ crystal structure of beta-FeSe was maintained for all investigated films and the *in-situ* observed diffraction pattern in reflection high energy diffraction also maintained the streaky pattern characteristic for smooth FeSe films even for the samples with the most pronounced island growth resulting in a root mean square atomic force microscopy roughness of more than 18 nm. Smaller flux ratios than 2.5 resulted in mixed – beta-FeSe/elemental Fe – phase samples. FeSe films grown under optimized conditions at 450 °C and a flux ratio of 2.5 (but without any post-growth UHV anneal) and capped with the commonly used FeTe (300 °C) and elemental Te (room temperature) layers yielded superconducting onset temperatures of about 30 K and a T_c of 20 K.

12:00pm **MI+2D+TF-ThM-13 Unraveling Picosecond Dynamic Material Processes on the Mesoscale by X-Ray Microscopy**, Thomas Feggeler, University of California, Berkeley; J. Lill, D. Guenzing, R. Meckenstock, D. Spoddig, B. Zingsem, University of Duisburg-Essen, Germany; M. Efremova, Eindhoven University of Technology, Netherlands; S. Pile, T. Schaffers, Johannes Kepler University, Austria; S. Wintz, Max Planck Institute for Intelligent Systems, Germany; M. Weigand, Helmholtz Center Berlin, Germany; A. Ney, Johannes Kepler University, Austria; M. Farle, H. Wende, K. Ollefs, University of Duisburg-Essen, Germany; D. Shapiro, Lawrence Berkeley National Laboratory; R. Falcone, University of California, Berkeley; H. Ohldag, Lawrence Berkeley National Laboratory

Dynamic processes govern a multitude of phenomena in physical, chemical and material sciences. Time- and spatially resolved element-specific monitoring of such processes is crucial in the understanding of phenomena like magnetization dynamics, battery charging and discharging, and phase transitions of several kinds. Time-Resolved Scanning Transmission X-ray Microscopy (TR-STXM) [1] is a versatile tool fulfilling these demands on the mesoscopic scale, offering element-specific observations with sub 50 nm spatial resolution and picosecond time sampling. By introducing a phased-locked-loop excitation synchronization scheme, TR-STXM also allows to sample dynamics originating from continuous wave excitations. This presentation introduces the TR-STXM technique and its principle of operation, and the setup developed at the Advanced Light Source at Lawrence Berkeley National Laboratory. The presentation is complemented by examples of dynamic magnetic measurements, which allow for local monitoring of magnetization dynamics in fields such as spintronics, magnonics, biomedical and energy related applications. Here we demonstrate TR-STXM results on Py/Co microstructures [2], Py stripe ensembles [3] and magnetite nanoparticle chains inside magnetotactic bacteria *Magnetospirillum Magnetotacticum* [4,5], showcasing localized uniform and non-uniform resonant magnetic responses, supplemented by micromagnetic simulations in good agreement.

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[1] T. Feggeler, A. Levitan, et al. J. Electron Spectrosc. Relat. Phenom. 2023. **267**: 147381.

[2] T. Feggeler, R. Meckenstock, et al. *Sci. Rep.* 2022, **12**:18724.

[3] S. Pile, T. Feggeler, et al. *Appl. Phys. Lett.* 2020, **116**(7): 072401.

[4] T. Feggeler, R. Meckenstock, et al. *Phys. Rev. Res.* 2021, **3**(3): 033036.

[5] T. Feggeler, J. Lill, et al. *New J. Phys.* 2023, **25**(4): 043010.

Thursday Afternoon, November 9, 2023

Light Sources Science Mini-Symposium

Room C124 - Session LS+AC+AS+LX+MI+TH-ThA

Facility Upgrades and Recent Capability Development

Moderators: David Shuh, Lawrence Berkeley National Laboratory, James G. Tobin, University of Wisconsin-Oshkosh, Gertrud Zwicknagl, Technical University Braunschweig

2:20pm **LS+AC+AS+LX+MI+TH-ThA-1 The Impact of Upgraded High-Brightness Synchrotron Lightsources on the Chemical Speciation of Nanoscale Heterogeneous Aggregates and Transformations, Andreas Scholl**, Advanced Light Source, Lawrence Berkeley National Laboratory

INVITED

ALS-U is an ongoing upgrade of the Advanced Light Source (ALS) at Berkeley Lab that will endow the ALS with revolutionary x-ray capabilities. The new storage ring will enable the production of highly focused beams of soft x-ray light that are at least 100 times brighter than those of the existing ALS. Applying this technology at the ALS will help us better understand and develop new materials and chemical systems needed to advance our research needs in energy science, environmental systems research, and biosciences in the 21st century. This will create a world-leading platform for next-generation soft x-ray and tender x-ray instrumentation.

Four beamlines with new and upgraded capabilities will become available after the upgrade. The FLEXON beamline (FLuctuation and EXcitation of Orders in the Nanoscale), a high-brightness coherent soft x-ray beamline, will provide x-ray photon correlation spectroscopy and diffraction imaging techniques to study electronic, chemical, and magnetic fluctuations in quantum materials with nanosecond temporal and nanometer spatial resolution. A new ALS-U developed tender x-ray beamline is designed to address challenges at the frontiers of diverse scientific areas, ranging from soft condensed matter and biomaterials to energy science and Earth and environmental sciences. It will offer state-of-the-art nanometer-resolved chemical imaging and resonant scattering nanoprobe, enabling operando and in situ studies of materials of K-edges of elements (Na through Ca) and the M and L edges of lanthanides and actinides. These two beamlines will be complemented by two upgraded beamlines for nanometer 3D chemical tomography based on ptychography of light elements and first-row transition metals and for high-resolution ARPES.

The high coherent flux of the upgraded ALS will drastically improve the speed, sensitivity, and spatial resolution of nanoprobe, enabling the speciation and forensic study of nanoscale constituents and contaminants via x-ray absorption spectroscopy and fluorescence detection. Chemical signatures can be correlated with morphology and compared with chemical standards. Operando experiments, for example, of liquid phase systems and studies under ambient conditions, will be enabled by specially designed sample holders and liquid cells using the high penetration of tender x-rays.

3:00pm **LS+AC+AS+LX+MI+TH-ThA-3 The Advanced Photon Source Upgrade: A transformative tool for understanding material structure., Jonathan Lang, J. Lang**, Argonne National Laboratory

INVITED

The APS is currently undergoing a major upgrade of the facility that will increase the brightness of the x-ray beams by factors of up to 500. This upgrade will provide transformational capabilities for examining the nanoscale structure and electronic configuration of materials and their evolution with external stimuli. This presentation will provide an update on the current status of the APS-U project, and discuss the new opportunities for imaging actinide and rare-earth compounds with this new source

3:40pm **LS+AC+AS+LX+MI+TH-ThA-5 Combining Focused Ion Beam Sectioning, Soft X-ray Spectromicroscopy, and Non-Negative Matrix Factorization to Reveal Actinide Chemical Speciation at the Nanoscale, Alexander Ditter, D. Smiles, J. Pacold, D. Lussier**, Lawrence Berkeley National Laboratory; Z. Dai, Lawrence Livermore National Laboratory; A. Altman, Lawrence Berkeley National Laboratory; M. Bachhav, Idaho National Laboratory; B. Chung, Lawrence Livermore National Laboratory; C. Degueldre, Lancaster University, UK; S. Donald, Lawrence Livermore National Laboratory; L. He, Idaho National Laboratory; M. Mara, S. Minasian, D. Shuh, Lawrence Berkeley National Laboratory

INVITED

Spectromicroscopy methods, combining the chemical insight of spectroscopy with microscopy imaging, can give a unique and informative view of a sample of interest. Scanning Transmission X-ray Microscope (STXM) spectromicroscopy is one such method, utilizing synchrotron

radiation to probe electronic structure with a spatial resolution in the tens of nanometers. Specialized methods like ptychography can push the spatial resolution even lower into the single nanometer range.

Discussed here, STXM spectromicroscopy is applied to two actinide samples: spent nuclear fuel, which offers a unique insight into the complex environment of nuclear fuel undergoing burnup, and uranium oxide allowed to age in a humid environment, which serves as a demonstration of the power of this technique for nuclear forensics investigations. These samples are created by focused ion beam (FIB) sectioning to generate cross-sections of ideal thickness for soft x-ray measurements (100-200 nm). The FIB method of sample preparation also allows for the measurement of highly radioactive spent fuel without containment due to the extremely small amount of material present.

Data analysis is a key component to the understanding of spectromicroscopy results for varied samples like these. Non-negative matrix factorization (NMF) is employed to identify key components and recent efforts to improve NMF to work with noisy individual STXM spectra are outlined here. Reproducibility of the analysis is a concern (as with similar methods like multivariate curve regression) and methods to enhance both reproducibility and interpretability of the results are discussed.

Combining STXM spectromicroscopy, FIB sectioning, and NMF analysis has allowed for unique insights into actinide materials. Potential future developments utilizing this method for other samples and with advanced techniques like ptychography are also discussed.

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 Tremsin, A.: MI+2D+TF-ThM-3, 7
 Tumbleson, R.: MI+2D+TF-WeA-3, 7
 Turek, I.: AC+MI+TH-TuM-11, 2
 — U —
 U. Saleheen, A.: MI+2D+TF-WeA-3, 7
 — V —
 Vallejo, K.: AC+AS+MI+TH-WeA-8, 5
 Vishik, I.: MI+2D+TF-ThM-10, 10
 Vitova, T.: LS+AC+LX+MI+TH-ThM-1, 8;
 LS+AC+LX+MI+TH-ThM-12, 9
 Vollmer, C.: LS+AC+LX+MI+TH-ThM-12, 9
 — W —
 Webb, S.: AC+AS+MI+TH-WeA-1, 5
 Weigand, M.: MI+2D+TF-ThM-13, 10
 Wende, H.: MI+2D+TF-ThM-13, 10
 Weng, T.: AC+MI+TH-TuM-13, 2
 White, F.: AC+MI+TH-TuA-7, 3
 Wilhelm, F.: AC+MI+TH-TuM-1, 1
 Wintz, S.: MI+2D+TF-ThM-13, 10
 Woods, J.: MI+2D+TF-WeA-3, 7
 Wray, L.: AC+MI+TH-TuM-10, 2
 — X —
 Xu, X.: MI+2D+TF-WeA-2, 6
 — Y —
 Yaita, T.: AC+MI+TH-TuA-3, 3
 Yang, D.: MI+2D+TF-WeA-2, 6
 Yao, J.: LS+AC+LX+MI+TH-ThM-12, 9
 Yi, H.: MI+2D+TF-ThM-12, 10
 Yoon, M.: MI+2D+TF-ThM-3, 9
 Young, L.: LS+AC+LX+MI+TH-ThM-10, 8
 Yu, S.: AC+MI+TH-TuM-13, 2
 — Z —
 Zhou, S.: AC+AS+MI+TH-WeA-8, 5
 Zingsem, B.: MI+2D+TF-ThM-13, 10
 zur Loye, H.: AC+MI+TH-TuM-6, 1