



FUNCTIONAL MATERIALS  
& NANOTECHNOLOGIES



**Joint International Conference  
Functional Materials and Nanotechnologies and  
Nanotechnology and Innovation in the  
Baltic Sea Region**

**FM&NT – NIBS 2022**

Riga, Latvia  
July 3 – July 6, 2022

**BOOK OF ABSTRACTS**



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ISBN 978-9934-23-645-7 (for pdf files)  
ISBN 978-9934-23-644-0 (for USB files)

## THE USE OF TUNGSTATE NANOPARTICLES IN HYBRID X-RAY DETECTORS

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### Keywords

Tungstates, nanoparticles, XRD, SEM, X-ray absorption spectroscopy

### Actuality and aim

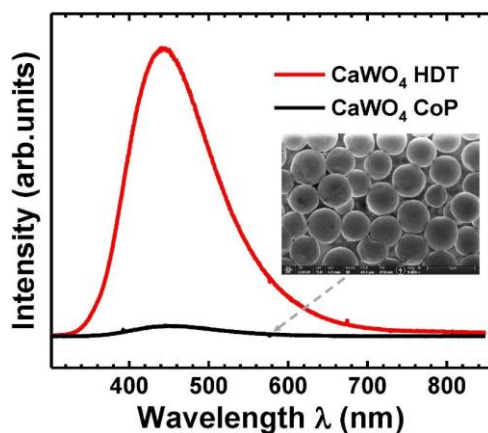
Tungstates of divalent metals form a large class of materials with various applications including but not limited to scintillators, photocatalysis, supercapacitors and sensors. The high value of the atomic number of tungsten ( $Z=74$ ) and the possibility to vary the atomic number of the second cation in a wide range ( $Z=12$  for Mg,  $Z=56$  for Ba) make tungstates attractive for the development of novel hybrid organic-inorganic X-ray detectors.

### Methods

In this study, scheelite ( $A=Ca, Sr$ ) and wolframite-type ( $A=Zn, Cd$ ) tungstates with different crystallinity were prepared using co-precipitation (CoP) and hydrothermal (HDT) synthesis. Nano- and polycrystalline powders were characterized by XRD, scanning electron microscopy (SEM), X-ray absorption spectroscopy (XAS), Raman spectroscopy and X-ray excited optical luminescence (XEOL).

### Results

Different synthesis parameters affect the size and morphology of nanoparticles (NPs). CoP at room temperature results in agglomerated NPs with an average size of 15-30 nm, but using citric acid as a capping agent the average size of NPs was smaller than 5 nm. However, XEOL is suppressed in NPs with low crystallinity because of high lattice defect concentration (see figure). In XAS spectra, size-induced local structure relaxations are observed which are more pronounced in NPs of wolframite-type. The HDT synthesis results in highly crystalline NPs with improved XEOL.



XEOL of two CaWO<sub>4</sub> samples.  
A SEM micrograph of CaWO<sub>4</sub> microspheres is shown in the inset.

### Conclusions

We show that the best sensitivity to X-rays is determined by a subtle interplay between particle size and their crystallinity.

### Acknowledgements

Financial support by the Latvian Council of Science project no. lzp-2019/1-0071 is acknowledged.