

# STRUCTURAL CHANGES INDUCED BY FLASH IN A SINGLE CRYSTAL OF $\text{Pr}_2\text{CuO}_4$

Dmitry Reznik, University of Colorado-Boulder  
[dmitry.reznik@colorado.edu](mailto:dmitry.reznik@colorado.edu)

Susmita Roy, University of Colorado-Boulder

Yu Zhang, University of Colorado-Boulder

Gang Cao, University of Colorado-Boulder

Syed I. A. Jalali, University of Colorado-Boulder

Rishi Raj, University of Colorado-Boulder

Zachary Morgan, Martin Greven, University of Minnesota

Feng Ye, Zachary Morgan, Oak Ridge National Lab

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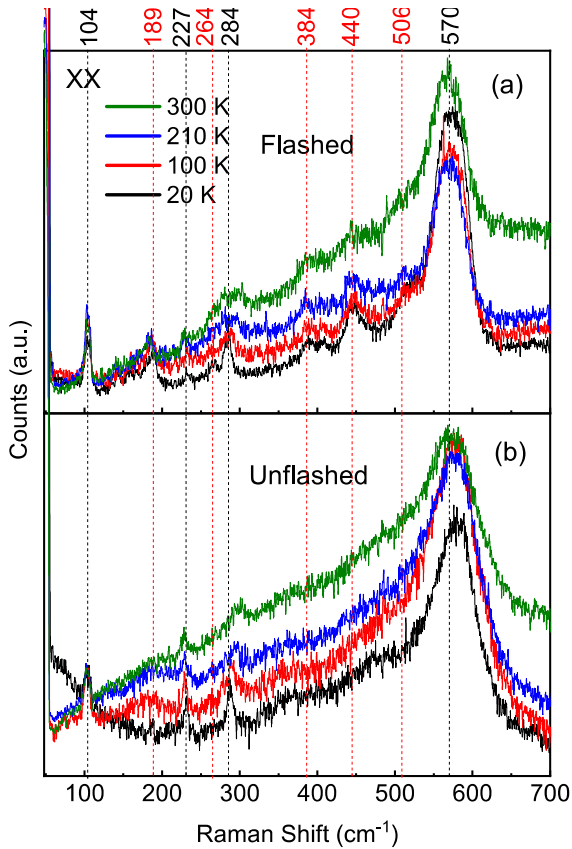


Figure 1 – Comparison of Raman spectra of unflashed and flashed single crystal of  $\text{Pr}_2\text{CuO}_4$

We demonstrate a novel approach to the structural and electronic property modification of perovskites, focusing on  $\text{Pr}_2\text{CuO}_4$ , an undoped parent compound of a class of electron-doped copper oxide superconductors. Currents were passed parallel or perpendicular to the copper oxygen layers with the voltage ramped up until a rapid drop in the resistivity was achieved, a process referred to as "Flash". The current was then further increased tenfold in current-control mode. This state was quenched by immersion into liquid nitrogen ( $\text{LN}_2$ ). Flash can drive many compounds into different atomic structures with new properties, whereas the quench freezes them into a long-lived state. Single-crystal neutron diffraction of as-grown and modified  $\text{Pr}_2\text{CuO}_4$  revealed a  $\sqrt{10} \times \sqrt{10}$  superlattice due to oxygen vacancy order. The diffraction peak intensities of the superlattice of the modified sample were significantly enhanced relative to the pristine sample. Raman-active phonons in the modified sample were considerably sharper. Measurements of electrical resistivity, magnetization and two-magnon Raman scattering indicate that the modification affected only the Pr-O layers, but not the Cu-O planes. These results point to enhanced oxygen vacancy order in the modified samples well beyond what can be achieved without passing electrical current. Our work opens a new avenue toward electric field/quench control of structure and properties of layered perovskite oxides.

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