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Phase Transformation of U<sub>3</sub>O<sub>8</sub> and Enhanced Structural Stability at Extreme Conditions<sup>1</sup> FUXIANG ZHANG, MAIK LANG, RODNEY EWING, University of Michigan — A powder sample of  $\beta$ -U<sub>3</sub>O<sub>8</sub> was pressurized at room temperature up to 37.5 GPa with a symmetric diamond anvil cell. XRD patterns clearly indicated that a phase transition occurred between 3-11 GPa. The highpressure phase is a fluorite-like structure. The fluorite-like structure is stable up to 37.5 GPa. The high-pressure phase was then laser heated to over 1700 K in the diamond any il cell at high pressure conditions. No phase transition was found at high pressure/ temperature conditions, and the fluorite-like structure of  $U_3O_8$  is even fully quenchable. The lattice parameter of the fluorite-like high-pressure phase is 5.425 Å at ambient conditions, which is smaller than that of the stoichiometric  $UO_2$ . Previous experiments have shown that the stoichiometric uranium dioxide  $(UO_2)$  is not stable at high pressure conditions and starts to transform to a cotunnite structure at  $\sim 30$  GPa. When heating the sample at high pressure, the critical transition pressure is greatly reduced. However, the fluorite-like high-pressure phase of  $U_3O_8$ is very stable at high pressure/high temperature conditions. The enhanced phase stability is believed to be related to the presence of extra oxygen (or U vacancies) in the structure.

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