APR17-2016-000114

Abstract for an Invited Paper for the APR17 Meeting of the American Physical Society

## Nuclear Excitation by Electronic Transition of U-235<sup>1</sup> PERRY CHODASH, Lawrence Livermore National Laboratory

Nuclear excitation by electronic transition (NEET) is a rare nuclear excitation that is theorized to exist in numerous isotopes. NEET is the inverse of bound internal conversion and occurs when an electronic transition couples to a nuclear transition causing the nucleus to enter an excited state. This process can only occur for isotopes with low-lying nuclear levels due to the requirement that the electronic and nuclear transitions have similar energies. One of the candidate isotopes for NEET,  $^{235}$ U, has been studied several times over the past 40 years and NEET of  $^{235}$ U has never been conclusively observed. These past experiments generated conflicting results with some experiments claiming to observe NEET of  $^{235}$ U and others setting limits for the NEET rate. If NEET of  $^{235}$ U were to occur, the uranium would be excited to its first excited nuclear state. The first excited nuclear state in  $^{235}$ U is only 76 eV, the second lowest known nuclear state. Additionally, the 76 eV state is a nuclear isomer that decays by internal conversion with a half-life of 26 minutes. In order to measure whether NEET occurs in  $^{235}$ U and at what rate, a uranium plasma was required. The plasma was generated using a Q-switched Nd:YAG laser outputting 789 mJ pulses of 1064 nm light. The laser light was focused onto uranium targets generating an intensity on target of order  $10^{12}$  W/cm<sup>2</sup>. The resulting plasma was captured on a catcher plate and electrons emitted from the catcher plate were accelerated and focused onto a microchannel plate detector. Measurements performed using a variety of uranium targets spanning depleted uranium up to 99.4% enriched uranium did not observe a 26 minute decay. An upper limit for the NEET rate of  $^{235}$ U was determined.

<sup>1</sup>This work was performed under the auspices of the U.S. Department of Energy by Lawrence Livermore National Laboratory under Contract DE-AC52-07NA27344. The U.S. DHS, UC Berkeley, the NNIS fellowship and the NSSC further supported this work.