## <b>FBXO25-ASSOCIATED NUCLEAR DOMAINS (FANDS): A NOVEL SUBNUCLEAR STRUCTURE</b>

<b><u>Adriana O. Manfiolli</u><sub>1</sub></b>; Ana Leticia G. C. Maragno<sub>1</sub>; Munira M. A. Baqui<sub>2</sub>; Sami Yokoo<sub>1</sub>; Eduardo B. Oliveira<sub>1</sub>; Odete A. B. Cunha<sub>1</sub> and Marcelo D. Gomes<sub>1</sub>

<sub>1</sub>Departamento de Bioquímica e Imunologia, FMRP-USP; <sub>2</sub>Departamento de Biologia Celular e Molecular e Bioagentes Patogênicos, FMRP-USP.

SCF<sub>FBXO25</sub> is a Skp1/Cul1/E-box protein (SCF) productive ubiquitin-ligase complex containing the F-box protein FBXO25. To help define its cellular function, we have developed an affinity-purified antibody raised against to a recombinant FBXO25 NH<sub>2</sub>-terminal fragment (aa 2 to 62). In western blot, FBXO25 protein was expressed in all major mouse tissues but was not detected in the striate muscle. In confocal analysis, the endogenous FBXO25 was found accumulated in a novel dot-like nuclear domain that is distinct from clastosomes and other well-characterized structures. Additionally, these compartments also contain at least two other ubiquitin-proteasome system components: 20S proteasome and Skp1. We propose to name these compartments "FANDs," for FBXO25-associated nuclear domains. Remarkably, the inhibition of transcription with actinomycin D drastically affects FANDs nuclear organization indicating that they are dynamic compartments influenced by the transcriptional activity of the cell. Supported by FAPESP and FAEPA.