



# Doctoral College Metabolic & Cardiovascular Disease

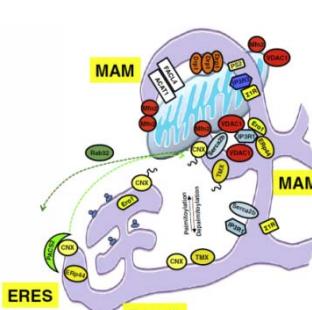
## THE MITOCHONDRIA-ASSOCIATED MEMBRANE (MAM): A CELLULAR STRUCTURE WHERE REDOX AND RABS DECIDE ON CANCER AND NEURODEGENERATION

GUEST LECTURE by

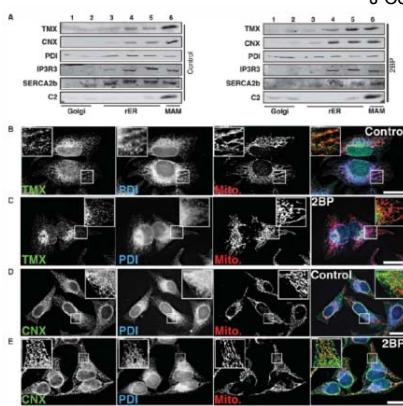


**Prof. Thomas Simmen, PhD**  
**Department of Cell Biology,**  
**University of Alberta, Edmonton, Canada**  
**Friday, 15.04.2016**  
**13:00**

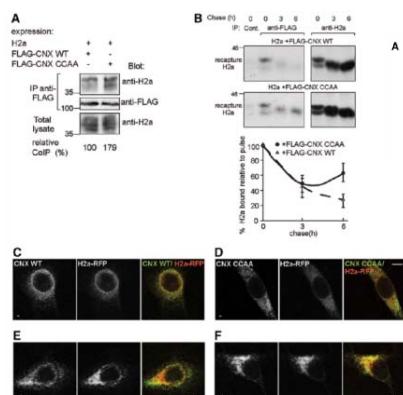
**SR 07.13, Preclinics, MUG  
(Harrachgasse 21, 1<sup>st</sup> floor)**



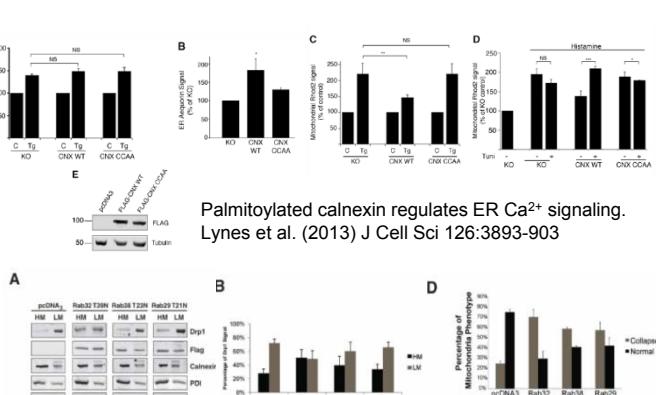
The MAM is a hub for lipid metabolism, mitochondrial fission and ER chaperones and oxidoreductases.  
Raturi & Simmen (2013) *Biochim Biophys Acta* 1833:213-24



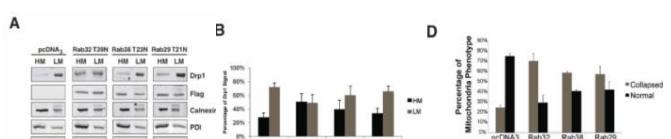
2-Bromopalmitate reduces the apposition of the TMX and calnexin signals with mitochondria. Lynes et al. (2012) *EMBO J* 31:457-70



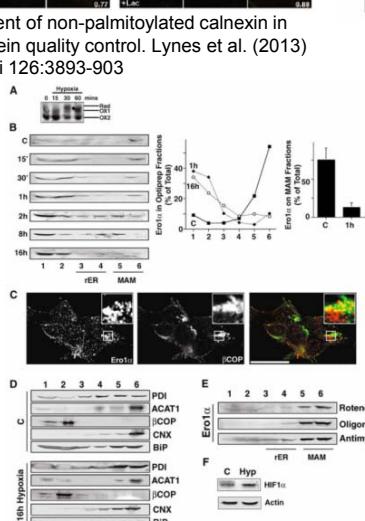
Involvement of non-palmitoylated calnexin in glycoprotein quality control. Lynes et al. (2013) *J Cell Sci* 126:3893-903



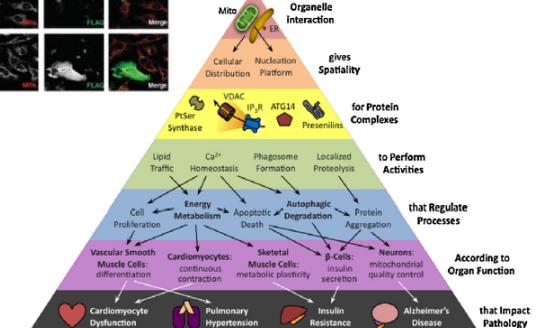
Palmitoylated calnexin regulates ER Ca<sup>2+</sup> signaling. Lynes et al. (2013) *J Cell Sci* 126:3893-903



Rab32 family proteins determine Drp1 localization and mitochondrial dynamics. Ortiz-Sandoval et al. (2014) *Cell Logist* 4(4):e986399



Ero1α MAM retention is oxygen-sensitive. Gilad et al. (2010) *Cell Stress Chaperones* 15:619-29



Physiopathological implications of organelle interaction. Bravo-Sagua et al. (2014) *Int J Biochem Cell Biol* 50:55-9