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**Symposium: Probing Ion Channel Structure/Function Using Novel Tools**

**889-Symp**

**Conformational Changes in Voltage-Sensing Domains: Concerted Simulation and Scattering Studies**

*Douglas J. Tobias*

Chemistry, University California, Irvine, Irvine, CA, USA.

Electrical signals in excitable tissues such as nerves and muscles are generated by the synchronized opening and closing of ion channels in cell membranes, which generate transient transmembrane gradients in the concentrations of sodium and potassium ions. These channels open and close in response to local changes in transmembrane potential as the signal propagates, and it is well established that their voltage sensitivity is conferred by charged structural elements, referred to as voltage-sensing domains. The details of how conformational changes in the voltage-sensing domains lead to opening/closing of the ion-conducting pores of the channels are still being worked out. Crystal structures of the open states of voltage-gated potassium channels are available, but the structure of the closed/resting state has not been determined to high resolution experimentally. In this talk I will report our efforts to generate a model of the resting state of the archael KvAP potassium channel based on atomistic molecular dynamics (MD) simulations in explicit membrane environments, with restraints derived from experimental functional data. I will also present results from simulation studies of an isolated voltage-sensing domain in a hydrated membrane, which include validation by neutron diffraction measurements, and direct observations of elementary gating charge displacement events during a 30-microsecond MD simulation under applied transmembrane potential. Finally, I will discuss prospects for and present preliminary results on using x-ray and neutron interferometry measurements, carried out by our collaborators, to validate and refine simulation-based models of the resting state and the voltage-sensing mechanism.

**890-Symp**

**Tricking Out the Toolbox: Use of Genetic Code Expansion for the Study of Ion Channels**

*Chris Ahern*

Molecular Physiology and Biophysics, University of Iowa, Iowa City, IA, USA.

The application of nonsense suppression strategies to membrane proteins has great potential for the study and manipulation of ion channels and receptors. The technique relies on the incorporation of a stop codon that is subsequently ‘suppressed’ by an orthogonal misacylated tRNA. Ion channels in particular are especially amenable to these approaches due to the built-in signal gain from the high conductance of recorded ions. On one end of the scale of possibility, ‘small’ suppression provides the means for single atom replacement within an amino acid or sub-atomic redistribution of pi electrons of aromatic side-chains. Such approaches provide an atomic resolution functional correlates to match high-resolution structural data, and have been key in decoding the interaction energetics of ligands, therapeutics, backbone contributions, salt-bridges and H-bonds in gating of voltage- and ligand-gated channels and receptors. More unconventional side-chains, such as light activated cross-linkers, hold promise for their ability to characterize transient noncovalent protein contacts. Lastly, encoded amino acid fluorophores, while highly valued, have been technically resistant to ion channel application and largely remain on the experimental fringe. Data will be presented on the use of genetic code expansion as a tool to reveal the basis for inactivation mechanisms employed by voltage-gated sodium and potassium channels.

**891-Symp**

**Small Molecule Modulation of Voltage-Gated Ion Channels**

*Heike Wulf*, Vladimir Yarov-Yarovoy.

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Pharmacological modulation of voltage-gated-like ion channels offers tremendous opportunities for the discovery of new drugs for neurological, cardiovascular and metabolic disorders, as well as for immunosuppression and cancer. However, in contrast to other therapeutic targets such as protein kinases, voltage-gated ion channels have not been popular with medicinal chemists in the pharmaceutical industry. One reason for this unpopularity is the absence of co-crystals of medically relevant channels with drug molecules bound, a prerequisite for structure based drug design. To address this challenge, molecular modeling, which recently made significant advances due to improved computational methods, and the availability of more high-resolution ion channel structures, is increasingly being used to interpret the results of experimental and computational studies. As illustrated with the example of CaV3.1 and CaV1.2 channels, small molecules can modulate their activity. Here I will present recent findings on the modulation of voltage-gated ion channels by small molecules using computational and experimental approaches.

**Platform: Calcium Signaling**

**893-Plat**

**Decreased Polycystin 2 Expression Alters Calcium-Contraction Coupling and Changes Beta-Adrenergic Signaling Pathways**

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Cardiac disorders are the main cause of mortality in autosomal dominant polycystic kidney disease (ADPKD). However, how mutated polycystins predispose ADPKD patients to cardiac pathologies before development of renal dysfunction is unknown. The effect of decreased polycystin 2 (PC2) levels on myocardial function. We hypothesize that heterozygous PC2 mice (Pkd2+/−) undergo cardiac remodeling due to changes in calcium handling, separate to renal complications. We found that Pkd2+/− mice underwent cardiac remodeling due to changes in calcium handling, separate to renal complications. We found that Pkd2+/− mice have increased left ventricular ejection fraction after stimulation with isoproterenol (ISO), a β-adrenergic receptor (βAR) agonist. Blockers of βAR-1 and 2 inhibited the ISO response in Pkd2+/− mice, suggesting that the dephosphorylated state of phospholamban is primarily by βAR-2 signaling. Importantly, the Pkd2+/− mice were norepinephrine sensitive and had no evidence of renal cysts. Our results showed that decreased PC2 levels shifted the βAR pathway balance and changed expression of calcium handling proteins, which resulted in altered cardiac contractility. We propose that PC2 levels in the heart directly contribute to cardiac remodeling in ADPKD patients in the absence of renal dysfunction.

**894-Plat**

**In vivo Reconstitution of the Mitochondrial Unipporter**


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The mitochondrial unipporter is a highly selective calcium channel present broadly in eukaryotes, but absent in Saccharomyces cerevisiae. Therefore,