BIOGRAPHICAL SKETCH

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NAME: Daniel Louis Minor, Jr., Ph.D.

eRA COMMONS USER NAME (credential, e.g., agency login): dminor

POSITION TITLE:

Professor, Departments of Biochemistry and Biophysics, & Cellular and Molecular Pharmacology Investigator, Cardiovascular Research Institute, University of California San Francisco Faculty Scientist, Molecular Biophysics & Integrated Imaging Division, Lawrence Berkeley National Laboratory, Berkeley

EDUCATION/TRAINING (Begin with baccalaureate or other initial professional education, such as nursing, include postdoctoral training and residency training if applicable. Add/delete rows as necessary.)

INSTITUTION AND LOCATION	DEGREE (if applicable)	Completion Date MM/YYYY	FIELD OF STUDY
University of Pennsylvania, Philadelphia, PA	B.A. magna cum laude	05/1989	Biochemistry (Honors) Biophysics (Honors)
Massachusetts Institute of Technology, Cambridge, MA	Ph.D.	02/1996	Chemistry
MRC Laboratory of Molecular Biology, Cambridge, UK	postdoc	09/1996	Ion channel structure
University of California, San Francisco, CA	postdoc	12/2000	Ion channel structure and function

A. Personal Statement

I have a broad background in ion channel structural biology and functional characterization. My interest in the physical chemistry of biological phenomena began with my undergraduate study in biophysics and biochemistry at the University of Pennsylvania. As a graduate student in the Department of Chemistry at MIT with Prof. Peter S. Kim, I focused on understanding the basic principles of protein folding and molecular interactions. While at MIT, I developed a keen interest in the proteins involved in electrical signaling. To pursue this interest, I worked as a postdoctoral fellow with Dr. Nigel Unwin at the LMB Cambridge and with Prof. Lily Y. Jan at UCSF where I was able to apply my background in structural biology to specific questions regarding ion channel structure and regulation. As a PI, I have focused my laboratory's efforts on structural and mechanistic understanding of ion channels and in the development of new pharmacological tools for orphaned channel classes. My lab is pursing a research program that combines structural biology, ion channel functional studies, and chemical biology approaches to develop new channel pharmacologies. I am a Professor of Biochemistry and Biophysics and Cellular and Molecular Pharmacology, an Investigator in the Cardiovascular Research Institute at UCSF, and a Faculty Scientist at LBNL.

My laboratory has made many contributions to the structural understanding of the function of various classes of ion channels and development of new channel modulators using a multidisciplinary approach employing genetic selections, biophysical approaches, chemical biology, and X-ray crystallography. Our work is exemplified by the following six key papers:

- 1. Van Petegem, F., Clark, K.A., Chatelain, F.C., and **Minor, D.L., Jr.**, "Structure of a complex between a voltage-gated calcium channel β-subunit and an α-subunit domain" *Nature* **429** 671-675 (2004) **PMID:15141227** (*Research Highlight Nature Rev. Neuroscience* 5:517, 2004; rated 'Exceptional' Faculty of 1000)
- 2. Bagriantsev, S., Clark, K.A., Peyronnet, R., Honoré, E., and **Minor, D.L., Jr.,** 'Multiple modalities act through a common gate to control K₂ channel function' function' The EMBO Journal **30** 3594-3606 (2011) **PMID: 21765396 PMCID: PMC3181481**
- 3. Bagriantsev, S. N., Ang, K.H., Gallardo-Godoy, A, Clark, K.A., Arkin, M.R., Renslo, A.R, and **Minor, D.L.**, **Jr.**, '*A high-throughput functional screen identifies small molecule regulators or temperature- and mechano-sensitive* K₂, *channels' ACS Chemical Biology* **8** 1841-1851 (2013) **PMID: 23738709PMCID: PMC3747594**
- 4. Lolicato, M., Riegelhaupt, P.M., Arrigoni, C., Clark, K.A., Minor, D.L., Jr.' *Transmembrane helix straightening and buckling underlies activation of mechanosensitive and thermosensitive K*₂, *channels' Neuron* 84 1198-1212 (2014) PMID: 25500157 PMCID: PMC4270892
- Arrigoni, C., Rohaim, A., Shaya, D., Findeisen ,F., Stein, R.A. Nurva, S.R., Mishra, S., Mchaourab, H.S., and Minor, D.L., Jr., 'Unfolding of a temperature-sensitive domain controls voltage-gated channel activation' Cell 164 922-936 (2016) PMID: 26919429 PMCID:PMC4769381
 Feb 18

6. Lolicato, M., Arrigoni, C., Mori, T., Sekioka, Y., Bryant, C., Clark, K.A., **Minor, D.L., Jr.** '*K*₂2.1(*TREK*-1):activator complexes reveal a cryptic selectivity filter binding site' Nature **547** 364-368 (2017) **PMID: 28693035 PMCID: PMC5778891**

B. Positions and Honors:

Positions and Employment

1990-1996	Graduate Student, Department of Chemistry, Massachusetts Institute of Technology			
1770 1770	Advisor: Peter S. Kim, Ph.D.			
1996	Postdoctoral Fellow, MRC-Laboratory of Molecular Biology Cambridge, England			
1996-2000	Advisor: Nigel Unwin, Ph.D. Postdoctoral Fellow, Howard Hughes Medical Institute, Department of Physiology,			
1990-2000	University of California, San Francisco, Advisor: Lily Y. Jan, Ph.D.			
2000-2007	Assistant Professor, Department of Biochemistry and Biophysics, UCSF			
2002-2007	Assistant Professor, Department of Cellular and Molecular Pharmacology, UCSF			
2000-present	Investigator, Cardiovascular Research Institute, UCSF			
2007-2011	Associate Professor (w/ tenure), Departments of Biochemistry and Biophysics & Cellular and			
	Molecular Pharmacology, UCSF			
2009-present	Biochemist, Faculty Scientist, Physical Biosciences Division, Lawrence Berkeley National			
	Laboratory (as of 2016, renamed as Molecular Biophysics & Integrated Imaging Division)			
2011-present	Professor (w/ tenure), Cardiovascular Research Institute, Departments of Biochemistry and			
	Biophysics & Cellular and Molecular Pharmacology, UCSF			
	ence and Professional Memberships			
2000-present	Member, Graduate Programs in Biological Sciences: Biochemistry, Biophysics, Chemistry and			
2000 2007	Chemical Biology, Neuroscience, Program in Molecular Medicine, UCSF			
2000-2006	Member, Graduate program in Biomedical Science			
	Protein Society, Member Biophysical Society, Member			
	Society for Neuroscience, Member			
2007 present	NIH BST-Q Study Section (<i>ad hoc</i> member)			
2007	NIH NTRC Study Section (<i>ad hoc</i> member)			
2008	NIH BPNS Study Section (ad hoc member)			
2008- present	Member, Graduate program in Biomedical Science			
2008 - present	Society of General Physiologists, Member			
2009	NIH NIDA CEBRA Study Section (ad hoc member)			
2011-present	NIH BPNS Study Section (permanent member)			
2012-2015	Biophysical Society Council Member			
2013-2014	US-Israel Binational Science Foundation – Scientific Advisory Board Member			
2015-2016	Beckman Young Investigator, Beckman Foundation - Selection Committee Member			
2016 2016	NIH ZEY1 VSN Study Section (<i>ad hoc</i> member)			
2010	NSF CAREER Review Study Section, MCB Division, Molecular Biophysics Cluster NIH: Special Emphasis Panel 'Biophysics' ZRG1 MDCN-R(04) (<i>ad hoc</i> member)			
2017	NSF Biomolecular Dynamics and Function II Study Section, MCB Division, Molecular			
2017	Biophysics Cluster			
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	Neuron Editorial Board			
	Journal of Molecular Biology Editorial Board			
Honors:				
1985 ILGWU				
	88 Dean's list, University of Pennsylvania,			
1988-89 Penn Student Agencies Scholarship, University of Pennsylvania				
1989 Helix Prize in Biochemistry, University of Pennsylvania 1989 Phi Beta Kappa				
1999 Fill beta Kappa 1996 Burroughs Wellcome Hitchings-Elion Fellowship				
2001-2004 McKnight Scholar in Neuroscience				
2001-2005 Rita Allen Scholar				
2002-2004 Alfred P. Sloan Research Fellow				
	kman Young Investigator			
	2002-2004 March of Dimes, Basil O'Connor Scholar			

2002-2005 Searle Scholar 2004-2006 McKnight Technological Innovations in Neuroscience Award 2007-2011 Established Investigator, American Heart Association 2010-2012 Fellow of the American Asthma Foundation 2011 Weizmann Institute of Science, Feinberg Visiting Faculty Fellowship

C. Contributions to science:

1) Protein folding: I established the experimental scale for β-sheet formation (*Minor and Kim, 1994a*) and uncovered that this property is context dependent (*Minor and Kim, 1994b*). Previously, there were no experimental measures of β-sheet formation. The observation of the effect of context led us to design an 11-residue sequence, the 'chameleon' sequence whose folding was entirely context dependent (*Minor and Kim, 1996*). *This work established that context could drive the formation of entire secondary structures, a demonstration that had implications for understanding structural transitions in amyloid and other fibril-forming proteins*.

Minor, D. L., Jr. and Kim P. S. "Measurement of the β -sheet forming propensities of amino acids" *Nature* **367** 660-663 (1994a) **PMID: 810785**

Minor, D.L., Jr. and Kim P.S. "Context is a major determinant of β-sheet propensity" *Nature* **371** 264-267 (1994b) **PMID: 8078589**

Minor, D.L., Jr. and Kim P.S. "Context-dependent secondary structure formation of a designed protein sequence" *Nature* **380** 730-734 (1996) **PMID: 8614471**

2) Voltage-gated calcium channels (Ca_xs): Voltage-gated calcium channels (Ca_xs) are central components of excitable tissues in the brain and heart. <u>When my laboratory started working on this channel family there were no high-resolution data for Ca_xs.</u> My laboratory determined the first high-resolution structure of part of a Ca_x, the β -subunit, alone and in complex with its interaction site from the channel (*Van Petegem et al., 2004*). <u>We subsequently have determined structures for all of the components for which there are presently high-resolution data</u> including calcium-calmodulin complexes of Ca_x1 (*Van Petegem et al., 2005; Kim et al., 2010*) and Ca_x2 IQ domains (*Kim et al., 2008*). These studies have established a structural foundation for investigating Ca_x function and have demonstrated the conformational complexities that underlie Ca_x feedback modulation by calcium-calmodulin.

Van Petegem, F., Clark, K.A., Chatelain, F.C., and **Minor**, **D.L.**, **Jr.**, "Structure of a complex between a voltage-gated calcium channel β -subunit and an α -subunit domain" *Nature* **429** 671-675 (2004) **PMID:15141227 PMCID: PMC3076333** (*Research Highlight Nature Rev. Neuroscience* **5:**517, 2004; rated 'Exceptional' Faculty of 1000)

Van Petegem, F, Chatelain, F.C., **Minor, D.L., Jr.,** "Insights into voltage-gated calcium channel regulation from the structure of the Ca.1.2 IQ domain-Ca²⁻/calmodulin complex" *Nature Structural & Molecular Biology* **12** 1108-1115 (2005) **PMID: 16299511 PMCID: PMC3020901**

Kim, E.Y., Rumpf, C.H., Fujiwara, Y., Cooley, E.S., Van Petegem, F., and Minor, D.L., Jr., "Structures of Ca₂ Ca₂/CaM-IQ domain complexes reveal binding modes that underlie calcium-dependent inactivation and facilitation' *Structure* **16** 1455-1467 (2008) **PMID: 18940602; PMCID: PMC2701236** (Rated 'Must Read' by Faculty of 1000)

Kim, E.Y., Rumpf, C.H., Fujiwara, Y., Van Petegem, F., Arant, R., Findeisen, F., Cooley, E.S., Isacoff, E.Y. and **Minor**, **D.L.**, **Jr.**, 'Multiple C-terminal Tail Ca²/CaMs regulate Ca₂1.2 function but do not mediate channel dimerization' *The EMBO Journal* **29** 3924-3938 (2010) **PMID: 20953164 PMCID: PMC3020648**

3) Bacterial voltage gated sodium channels (BacNa,s): BacNa,s are model systems for understanding the fundamental principles of voltage gated channel function. Using a protein dissection approach, we demonstrated that BacNa,s are modular within the membrane and that the pore domain can be excised from the voltage sensors to produce functional sodium or calcium selective channels. These 'pore-only' proteins establish a general design principle of modularity within voltage-gated ion channel membrane domains and show that the pore domain and voltage sensor domains are separable units (*Shaya et al. 2011*). We determined the structure of a bacterial 'pore-only' sodium channel. *This work uncovered the complete structure of a pore-only BacNa*, and revealed the structure of a crucial part, the cytoplasmic tail that had evaded other <u>structural studies (Shaya et al. 2014)</u>. This finding together with functional tests resolved controversy over the location of the intracellular gate that controls channel opening and established a role for the cytoplasmic domain in controlling channel opening. Structure determination of a 'pore-only' bacterial sodium channel also identified an ion binding site having common to human voltage-gated calcium channel pores and

discovered a previously unknown determinant of ion selectivity in human voltage-gated calcium channels (Shaya et al. 2014).

There has been much debate about the origins of thermosensitivity in ion channels and whether such responses rely on a dedicated sensing to domain or a more distributed property of the channel. <u>Our recent</u> <u>studies of BacNa_v mechanisms have defined the first authentic temperature-sensitive module in an ion channel and</u> <u>show that it is possible for a single, defined domain to control channel thermal responses.</u> (Arrigoni *et al.* 2016).

Shaya, D., Kreir, M., Robbins, R.A., Wong, S., Hammon, J., Brüggemann, A., and **Minor, D.L., Jr.,** 'Voltage-gated sodium channel (Na_v) protein dissection creates a set of functional 'pore-only' proteins' Proc Natl Acad Sci USA **108** 12313-12318 (2011) **PMID: 21746903: PMCID: PMC3145705**

Shaya, D., Findeisen, F., Abderemane-Ali, F., Arrigoni, C., Wong, S., Reddy Nurva, S., Loussouarn, G., and Minor, D.L., Jr., 'Structure of a prokaryotic sodium channel pore reveals essential gating elements and an outer ion binding site common to eukaryotic channels' Journal of Molecular Biology **426** 476-483 (2014) **PMID: 24120938 PMCID: PMC3947372**

Arrigoni, C., Rohaim, A., Shaya, D., Findeisen ,F., Stein, R.A. Nurva, S.R., Mishra, S., Mchaourab, H.S., and **Minor**, **D.L.**, **Jr.**, 'Unfolding of a temperature-sensitive domain controls voltage-gated channel activation' Cell 164 922-936 (2016) **PMID: 26919429 PMCID:PMC4769381**

4) K_x channels K_x channels are a diverse set of potassium channels that produce 'leak' currents that control cellular excitability. K_x2.1 (TREK-1) is a classic polymodal channel that is gated by a wide range of stimuli including pH, temperature, and pressure. When we began studying this channel family, it was unclear whether the various inputs controlled the channel by separate or common mechanisms. Using a gain-of-function selection that identified K₂2.1 (TREK-1) mutants that rescued a potassium transport deficient yeast strain, we identified a key element of the channel gate and established that pH, temperature, and pressure act through a common mechanism to control the gate, which is the channel selectivity filter (Bagriantsev et al., 2011). We established that signals from the intracellular sensors for temperature and phosphorylation are relayed to the selectivity filter via transmembrane helix M4 (*Bagriantsev et al. 2012*). Structure determination of two gain-of-function mutants in K₂4.1 (TRAAK) revealed an unexpected mechanism that is opposite from prior expectations based on studies of other potassium channel classes (Lolicato et al. 2014). Upon activation M4 straightens at a conserved glycine and causes the M2 transmembrane helix to buckle at a conserved 'GXG' sequence. Structure-based functional tests demonstrate that this mechanism operates in the thermosensitive and mechanosensitive K_{x} subfamily. Our studies have shown that K_{x} channels gate at the selectivity filter rather than an intracellular gate. Our latest work has defined the structure of K₂2.1 (TREK-1) and uncovered a previously unknown small molecule modulatory site and demonstrates that small molecules can directly activate the selectivity filter gate (Lolicato et al., 2017).

Bagriantsev, S., Clark, K.A., Peyronnet, R., Honoré, E., and **Minor, D.L., Jr.,** 'Multiple modalities act through a common gate to control K₂ channel function' function' The EMBO Journal **30** 3594-3606 (2011) **PMID: 21765396 PMCID: PMC3181481**

Bagriantsev, S., Clark, K.A., and **Minor, D.L., Jr.,** '*Metabolic and thermal stimuli control* K₂2.1 (TREK-1) through modular sensory and gating domains' The EMBO Journal **31** 3297-3308 (2012) **PMID: 22728824 PMCID: PMC3411076**

Lolicato, M., Riegelhaupt, P.M., Arrigoni, C., Clark, K.A., and **Minor, D.L., Jr.** '*Transmembrane helix* straightening and buckling underlies activation of mechanosensitive and thermosensitive K₂, channels' Neuron **84** 1198-1212 (2014) **PMID: 25500157 PMCID: PMC4270892'**

Lolicato, M., Arrigoni, C., Mori, T., Sekioka, Y., Bryant, C., Clark, K.A., **Minor, D.L., Jr.** *'K*₂2.1(*TREK*-1):activator complexes reveal a cryptic selectivity filter binding site' Nature **547** 364-368 (2017) **PMID: 28693035 PMCID: PMC5778891**

5) Ion channel-modulator interactions: We established a yeast genetic selection system for investigating ion channel-modulator interactions. By selecting for barium resistant mutants of the inward rectifier Kir2.1, we identified a T \rightarrow K mutant that demonstrated that the pore helix dipole, thought by many to be important for function, is not involved in ion permeation (*Chatelain et al.* 2005). We showed that the yeast system could be used generally to identify residues important for ion channel-small molecule interactions. Recently, we have established this system as a means to identify small molecule modulators of K_w channel function and have developed a novel, selective activator of the K_w subclass of mechanosensitive and thermosensitive channels (*Bagriantsev et al.*, 2013). <u>Our has defined the first set selective K_w activators and resulted in a patent (*Bagriantsev et al.*, 2013). We have also shown that the yeast system can be used to discover novel protein-based regulators of ion channel function (*Bagriantsev et al.*, 2014). <u>We have recently defined a novel class</u></u>

of K_x activators that bind to a previously unknown site supporting the channel selectivity filter, the ' K_x modulator <u>pocket'</u>. These studies set the stage for the development of a new class of ion channel modulators (Lolicato et al., 2017).

Chatelain, F.C., Alagem, N., Xu, Q., Pancaroglu, R., Reuveny, E., and **Minor, D.L., Jr.**, "The pore helix dipole has a minor role in inward rectifier channel function" *Neuron* **47** 833-843 (2005) **PMID: 16157278 PMCID: PMC3017504 (Preview** *Neuron* **47** 777-778, 2005)

Chatelain, F.C., Gazzarrini, S., Fujiwara, Y., Arrigoni, C., Domigan, C., Ferrara, G., Pantoja, C., Thiel, G., Moroni, A., and **Minor, D.L., Jr**., 'Selection of inhibitor-resistant viral potassium channels identifies a selectivity filter site that affects barium and amantadine block' *PLoS ONE* 4 (10) e7496. doi:10.1371/journal.pone.0007496 (2009) **PMID: 19834614; PMCID: PMC2759520**

Bagriantsev, S. N., Ang, K.H., Gallardo-Godoy, A, Clark, K.A., Arkin, M.R., Renslo, A.R, and **Minor, D.L.**, **Jr.**, '*A high-throughput functional screen identifies small molecule regulators or temperature- and mechano-sensitive* K₂, *channels' ACS Chemical Biology* 8 1841-1851 (2013) **PMID: 23738709PMCID: PMC3747594**

US patent application number 61/785,155 'Modulation of K₂ channels' Bagriantsev, S.N., Renslo, A.R., and **Minor, D. L., Jr.**

Bagriantsev, S.N., Chatelain, F.C., Clark, K.A., Alagem, N., Reuveny, E., Minor, D.L., Jr. '*Tethered protein display identifies a novel Kir3.2 (GIRK2) regulator from protein scaffold libraries' ACS Chemical Neuroscience* **5** 812-822 (2014) **PMID: 25028803 PMCID: PMC4176385**

Lolicato, M., Arrigoni, C., Mori, T., Sekioka, Y., Bryant, C., Clark, K.A., **Minor, D.L., Jr.** '*K*₂2.1(*TREK*-1):activator complexes reveal a cryptic selectivity filter binding site' Nature **547** 364-368 (2017) **PMID**: **28693035 PMCID: PMC5778891**

URL to myNCBI a full list of published work

http://www.ncbi.nlm.nih.gov/sites/myncbi/daniel.minor.1/bibliography/41458029/public/?sort=date&dir ection=ascending.

Complete list of Peer-reviewed publications (in chronological order):

- 1. **Minor, D. L., Jr.** and Kim P. S. "Measurement of the β-sheet forming propensities of amino acids" *Nature* **367** 660-663 (1994) **PMID: 810785**
- 2. **Minor**, **D.L.**, **Jr.** and Kim P.S. "Context is a major determinant of β-sheet propensity" *Nature* **371** 264-267 (1994) **PMID: 8078589**
- 3. Schumacher, T.N.M., Mayr, L.M., **Minor, D.L., Jr.,** Milhollen, M.A., Burgess, M.W. and Kim, P.S. "Identification of (D)-peptide ligands through Mirror-Image phage display" *Science* **271** 1854-1857 (1996) **PMID: 8596952**
- 4. Minor, D.L., Jr. and Kim P.S. "Context-dependent secondary structure formation of a designed protein sequence" *Nature* **380** 730-734 (1996) **PMID: 8614471**
- 5. Minor, D.L., Jr., Masseling, S.J., Jan, Y.N. and Jan, L.Y. "Transmembrane structure of an inwardly rectifying potassium channel" *Cell* **96** 879-891 (1999) **PMID: 10102275**
- 6. **Minor, D.L., Jr.,** Lin, Y.F, Mobley, B.C., Avelar, A., Jan, Y.N., Jan, L.Y. and Berger, J.M. "The polar T1 interface is linked to conformational changes that open the voltage-gated potassium channel" *Cell* **102** 657-670 (2000) **PMID: 11007484**
- 7. Mosavi, L. K., Minor, D.L., Jr., and Peng, Z.-y., "Consensus-derived structural determinants of the ankyrin repeat motif" *Proceedings of the National Academy of Sciences*, USA **99** 16029-16034 (2002) **PMID:12461176**; **PMCID: PMC138559**
- 8. Walden, H., Podgorski, M.S., Huang, D.T., Miller, D.W., Howard, R.J., **Minor, D.L., Jr.,** Holton, J.M., and Schulman, B.A., "The structure of APPBP-1UBA3-NEDD8-ATP complex reveals the basis for selective ubiquitin-like protein activation by an E1" *Molecular Cell* **12** 1427-1437 (2003) **PMID: 14690597**
- 9. Van Petegem, F., Clark, K.A., Chatelain, F.C., and Minor, D.L., Jr., "Structure of a complex between a voltage-gated calcium channel β-subunit and an α-subunit domain" *Nature* **429** 671-675 (2004) PMID:15141227 (*Research Highlight Nature Rev. Neuroscience* **5**:517, 2004; rated 'Exceptional' Faculty of 1000)
- 10. Chatelain, F.C., Alagem, N., Xu, Q., Pancaroglu, R., Reuveny, E., and **Minor, D.L., Jr**., "The pore helix dipole has a minor role in inward rectifier channel function" *Neuron* **47** 833-843 (2005) **PMID: 16157278** (Preview *Neuron* **47** 777-778, 2005)

- 11. Van Petegem, F, Chatelain, F.C., **Minor, D.L., Jr.**, "Insights into voltage-gated calcium channel regulation from the structure of the Ca. 1.2 IQ domain-Ca²/calmodulin complex" *Nature Structural & Molecular Biology* **12** 1108-1115 (2005) **PMID: 16299511 PMCID: PMC3020901**
- 12. Tsuruda, P., Julius, D., and **Minor, D.L., Jr**., "Identification and characterization of a domain required for assembly of a cold-activated TRP channel" *Neuron* **51** 201-212 (2006) **PMID: 16846855 PMCID: PMC3014052**
- 13. Michelsen, K., Mrowiec, T., Duderstadt, K.E., Frey, S., **Minor, D.L., Jr**., Mayer, M.P., Schwappach, B., "A multimeric membrane protein reveals 14-3-3 isoform specificity in forward transport in yeast' *Traffic* 7 903-916 (2006) **PMID: 16734667**
- 14. Pioletti, M., Findeisen, F., Hura, G.L., and Minor, D.L., Jr., "Three-dimensional structure of the KChIP1/Kv4.3 T1 domain complex reveals a cross-shaped octamer" *Nature Structural & Molecular Biology* 13 987-995 (2006) PMID: 17057713 PMCID: PMC3018330
- 15. Howard, R.J., Clark, K.A., Holton, J.M., and Minor, D.L., Jr., "Structural insight into KCNQ (Kv7) channel assembly and channelopathy" *Neuron* 53 663-675 (2007) PMID: 17329207 PMCID: PMC3011230
- 16. Balss, J., Paptheodorou, P., Mehmel, M., Baumeister, D., Hertel, B., Delaroque, N., Chatelain, F. C., Minor, D.L., Jr., Van Etten, J.L., Rassaw, J., Moroni, A., and Thiel, G. "Transmembrane Domain Length of Viral Potassium Ion Channels is a Signal for Mitochondria Targeting" PNAS 105 12313-12318 (2008) PMID: 18719119; PMCID: PMC2518832
- 17. Van Petegem, F, Duderstadt, K.E., Clark, K.A., Wang, M., **Minor, D.L., Jr.**, "Alanine-scanning mutagenesis defines a conserved energetic hotspot in the Ca_va_v AID-Ca_vβ interaction site that is critical for channel modulation" *Structure* **14** 280-294 (2008) **PMID: 18275819 PMCID: PMC3018278**
- 18. Fujiwara, Y. and Minor, D.L., Jr., 'X-ray crystal structure of a TRPM assembly domain reveals an antiparallel four-stranded coiled-coil' *Journal of Molecular Biology* **383** 854-870 (2008) PMID: **18782578**; PMCID: PMC2630241
- 19. Kim, E.Y., Rumpf, C.H., Fujiwara, Y., Cooley, E.S., Van Petegem, F., and **Minor, D.L., Jr.**, "Structures of Ca₂2 Ca²⁻/CaM-IQ domain complexes reveal binding modes that underlie calcium-dependent inactivation and facilitation' *Structure* **16** 1455-1467 (2008) **PMID: 18940602; PMCID: PMC2701236** (Rated 'Must Read' by Faculty of 1000)
- 20. Hammon, J., Palanivelu, D.V., Chen, J., Patel, C., and Minor, D.L., Jr., 'A green fluorescent protein screen for identification of well-expressed membrane proteins' *Protein Science* **18** 121-133 (2009) **PMID: 19177357**; **PMCID : PMC2708023** (Rated 'Recommended' by Faculty of 1000)
- 21. Findeisen, F. and Minor, D.L., Jr., 'Disruption of the IS6-AID linker affects voltage-gated calcium channel inactivation and facilitation' *Journal of General Physiology* 133 327-343 (2009) PMID: 19237593: PMCID: PMC2654080
- 22. Xu, Q. and **Minor**, **D.L.**, **Jr.**, 'Crystal structure of a trimeric form of the Kv7.1 (KCNQ1) A domain Tail coiled coil reveals structural plasticity and context dependent changes in a putative coiled-coil trimerization motif' *Protein Science* **18** 2100-2114 (2009) **PMID 19693805 PMCID: PMC2786974**
- 23. Chatelain, F.C., Gazzarrini, S., Fujiwara, Y., Arrigoni, C., Domigan, C., Ferrara, G., Pantoja, C., Thiel, G., Moroni, A., and **Minor, D.L., Jr.,** 'Selection of inhibitor-resistant viral potassium channels identifies a selectivity filter site that affects barium and amantadine block' *PLoS ONE* **4** (10) e7496. doi:10.1371/journal.pone.0007496 (2009) PMID: 19834614; PMCID: PMC2759520
- 24. Kohout S.C., Bell S.C., Liu L., Xu Q., **Minor**, **D.L.**, **Jr.**, and Isacoff, E.Y., "Electrochemical coupling in the voltage-dependent phosphatase Ci-VSP *Nature Chemical Biology* **6** 369-375 (2010) **PMID: 20364128; PMCID: PMC2857593** (Preview *Nature Chemical Biology* **6** 315-316, 2010)
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- 26. Findeisen, F. and **Minor**, **D.L.**, **Jr.**, 'Structural basis for the differential effects of CaBP1 and calmodulin on Ca_v1.2 calcium-dependent inactivation' *Structure* **18** 1617-1631(2010) **PMID: 21134641: PMCID: In Progress** (Rated 'Recommended' by Faculty of 1000)
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Patents issued or pending

Patent US9862684B2 'Modulation of K_a channels' Bagriantsev, S.N., Renslo, A.R., and Minor, D. L., Jr.

D. Research Support. **Ongoing Research Support:** R01HL080050 NIH/NĤLBI 5/1/05-3/31/18 Minor (PI) Structure and function of voltage-gated calcium channels \$388,174 The major goals of this project are to investigate the molecular origins of calcium channel function. Role: PÍ R01 MH093603-01 NIH/NIMH Minor (PI) 03/01/11-02/28/21 Genetic and chemical biological studies of K_a structure, function, and modulation \$400,000 The major goals are to develop genetic selection-based, approaches to define and characterize essential elements of K₂, channel gating and to discover and characterize small molecule K₂, modulators. Role: PI R01DC007664 NIH/NIDCD Minor (PI) 07/01/05-01/31/22 Structure and function of ion channel assembly and signaling complexes \$360.648 The major goal of this project is to study the structural biology of potassium channel regulation. Role: PÍ U.S.-Israel Binational Science Foundation Grant 2011124 Minor/Reuveny (PI) 10/1/16 – 09/30/20 'Molecular Mechanisms of the regulation of SOCE by SARAF' \$13,376 The major goals of this project are to develop an understanding the interactions between SOCE channels and calcium regulation Role: Co-PI with E. Reuveny, Weizmann Institute

Pending Research Support:

R01 MH116278-01 NIH/NIMHRenslo (PI)4/1/18 - 3/31/21'Expanding the chemical biology of K_x channels with selective cellular and in vivo probes'The major goals of this project are to advance new cellular and *in vivo* compatible probes of K_x channels.Role: Co-PI with A. Renslo, UCSF