



## Manuel F. Navedo, Ph.D., F.A.H.A.

### Research/Academic Interests

Research in the Navedo Lab aims to determine the cellular and molecular mechanisms regulating arterial smooth muscle excitability and vascular reactivity. Innovative techniques and approaches are used to specifically examine the mechanisms by which local and global calcium signals regulate excitation-contraction coupling and excitation-transcription coupling in arterial smooth muscle cells during physiological and pathological conditions such as hypertension and diabetes. The overarching objective is to uncover regulatory signaling pathways of vascular (dys)function that could help identify new targets for exploitation as novel therapeutic interventions to treat vascular complications during hypertension and diabetes.

**Title** Professor

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**Division** Pharmacology

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**Languages** Spanish

**Education** Ph.D., Neurobiology, University of Puerto Rico, San Juan PR 2003  
B.S., University of Puerto Rico, San Juan PR 1998

**Fellowships** PBio/Vascular Biology, University of Washington, Seattle WA 2004-2007

**Professional Memberships** American Heart Association  
American Physiological Society  
European Calcium Society

**Honors and Awards** Fellow of the American Heart Association, 2019  
Charter Member, National Institutes of Health Vascular Cell and Molecular Biology Study Section, 2016, 2017, 2018, 2019, 2020, 2021, 2022  
UCD Academic Senate Research Travel Award, 2016  
National Institutes of Health Early Career Reviewer Program Alumni, 2015

**Select Recent Publications** To view a complete list of Dr. Navedo's publications, please click [here](#).

Nieves-Cintrón M, Santana LF, Navedo MF. TRPML1ng on sparks. *Sci Signal*. 2020 Jun 23;13



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Syed AU, Reddy GR, Ghosh D, Prada MP, Nystoriak MA, Morotti S, Grandi E, Sirish P, Chiamvimonvat N, Hell JW, Santana LF, Xiang YK, Nieves-Cintrón M, Navedo MF. Adenylyl cyclase 5-generated cAMP controls cerebral vascular reactivity during diabetic hyperglycemia. *J Clin Invest*. 2019 Jun 4;129(8):3140-3152. doi:10.1172/JCI124705. PMID:31162142. Highlighted by several news outlets and commentary on *Journal of Diabetes Investigations*. Reviewed by F1000. PMC6668679.

Prada MP, Syed AU, Buonarati OR, Reddy GR, Nystoriak MA, Ghosh D, Simó S, Sato D, Sasse KC, Ward SM, Santana LF, Xiang YK, Hell JW, Nieves-Cintrón M, Navedo MF. A Gs-coupled purinergic receptor boosts Ca<sup>2+</sup> influx and vascular contractility during diabetic hyperglycemia. *Elife*. 2019 Mar 1;8:e42214. doi:10.7554/eLife.42214. PMID:30821687.

Nystoriak MA, Nieves-Cintrón M, Patriarchi T, Buonarati OR, Prada MP, Morotti S, Grandi E, Fernandes JD, Forbush K, Hofmann F, Sasse KC, Scott JD, Ward SM, Hell JW, Navedo MF. Ser1928 phosphorylation by PKA stimulates the L-type Ca<sup>2+</sup> channel Ca<sub>v</sub>1.2 and vasoconstriction during acute hyperglycemia and diabetes. *Sci Signal*. 2017 Jan 24;10(463):eaaf9647. doi:10.1126/scisignal.aaf9647. PMID:28119464. This article was highlighted in the cover and podcast of the journal and by Faculty of 1000. PMC5297430.

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