



## Philip Baran, Ph.D.

Professor  
Scripps Research  
La Jolla, CA

## Electrifying Chemistry

### ABSTRACT:

Electrochemistry offers new and sustainable means to accomplish enabling reactions for organic synthesis. This presentation will cover some of our latest developments in this area.

### BIO:

Phil Baran was born in 1977 in Denville, New Jersey. He received his B.S. in chemistry from NYU in 1997, his Ph.D. from Scripps Research Institute in 2001, and from 2001-2003 he was an NIH-postdoctoral fellow at Harvard. His independent career began at Scripps in the summer of 2003. Phil has published over 220 scientific articles, several patents, and has been the recipient of several ACS awards such as the Corey (2015), Pure Chemistry (2010), Fresenius (2006), and Nobel Laureate Signature (2003), and several international distinctions such as the Hirata Gold Medal and Mukaiyama Prize (Japan), the RSC award in Synthesis (UK), the Sackler Prize (Israel), and the Janssen Prize (Belgium). In 2013 he was named a MacArthur Foundation Fellow, in 2015 he was elected to the American Academy of Arts and Sciences, in 2016 he was awarded the Blavatnik National Award, and in 2017, he was elected to the National Academy of Sciences, USA. He has delivered hundreds of lectures around the world and consults for numerous companies such as Bristol-Myers Squibb, Gilead, Boehringer-Ingelheim, AstraZeneca, DuPont and TEVA, and is a scientific advisory board member for Eisai, Abide, Nutcracker, Quanta and AsymChem. In 2016 he was appointed as an Associate Editor for the *Journal of the American Chemical Society*. He co-founded Sirenas Marine Discovery (2012) and Vividion Therapeutics (2016) and in 2013 he co-authored *The Portable Chemist's Consultant*, an interactive book published on the iBooks store along with his graduate class in Heterocyclic Chemistry (viewable on YouTube). The Baran laboratory is committed to identifying areas of chemical synthesis that can have a dramatic impact on the rate of drug discovery and development. This is achieved both through the development of practical total syntheses of complex natural products (such as terpenes, alkaloids, peptides, and oligonucleotides) and by inventing reactions which can dramatically simplify retrosynthesis.