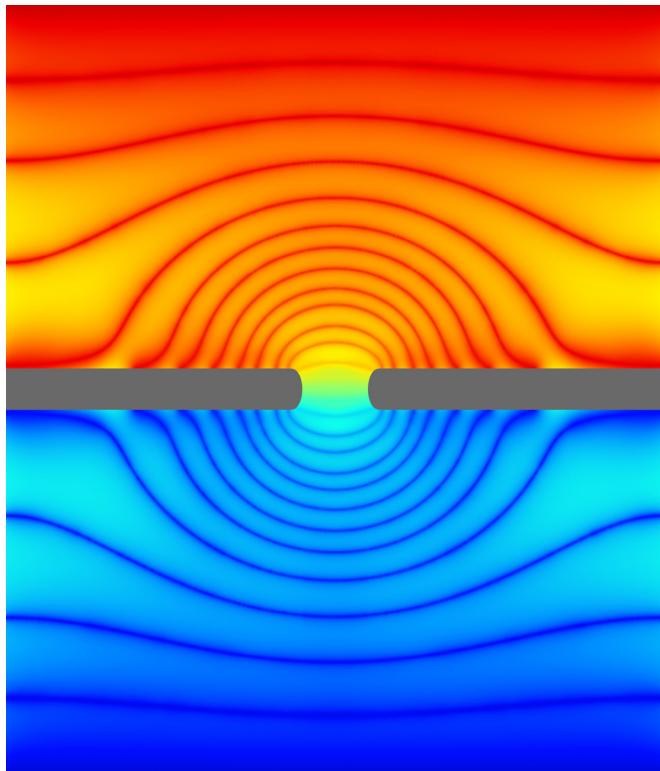


Scientists channel graphene to understand filtration and ion transport into cells

8 December 2017, by Ben Stein



In this simulation, a biological membrane (gray) with an ion channel (center) is immersed in a solution of water and ions. This cross section of a simulation "box" shows the electric potential, the externally supplied "force" that drives ions through the channel. A dazzling pattern emerges in this potential due to the presence of the channel -- the colors show the lines of equal potential. The slowly decaying nature of this pattern in space makes simulations difficult. The golden aspect ratio -- the chosen ratio of height to width of this box -- allows for small simulations to effectively capture the effect of the large spatial dimensions of the experiment. Credit: NIST

Tiny pores at a cell's entryway act as miniature bouncers, letting in some electrically charged atoms—ions—but blocking others. Operating as exquisitely sensitive filters, these "ion channels" play a critical role in biological functions such as

muscle contraction and the firing of brain cells. To rapidly transport the right ions through the cell membrane, the tiny channels rely on a complex interplay between the ions and surrounding molecules, particularly water, that have an affinity for the charged atoms. But these molecular processes have traditionally been difficult to model—and therefore to understand—using computers or artificial structures.

Now, researchers at the National Institute of Standards and Technology (NIST) and their colleagues have demonstrated that nanometer-scale pores etched into layers of graphene—atomically thin sheets of carbon renowned for their strength and conductivity—can provide a simple model for the complex operation of ion channels.

This model allows scientists to measure a host of properties related to [ion transport](#). In addition, graphene nanopores may ultimately provide scientists with efficient mechanical filters suitable for such processes as removing salt from ocean water and identifying defective DNA in genetic material.

NIST scientist Michael Zwolak, along with Subin Sahu (who is jointly affiliated with NIST, the University of Maryland NanoCenter and Oregon State University), has also discovered a way to simulate aspects of [ion channel](#) behavior while accounting for such computationally intensive details as molecular-scale variations in the size or shape of the [channel](#).

To squeeze through a cell's ion channel, which is an assemblage of proteins with a pore only a few atoms wide, ions must lose some or all of the water molecules bound to them. However, the amount of energy required to do so is often prohibitive, so ions need some extra help. They get that assistance from the ion channel itself, which is lined with molecules that have opposite charges to certain

ions, and thus helps to attract them. Moreover, the arrangement of these charged molecules provides a better fit for some ions versus others, creating a highly selective filter. For instance, certain ion channels are lined with negatively charged molecules that are distributed in such a way that they can easily accommodate [potassium ions](#) but not [sodium ions](#).

It's the selectivity of ion channels that scientists want to understand better, both to learn how biological systems function and because the operation of these channels may suggest a promising way to engineer non-biological filters for a host of industrial uses.

By turning to a simpler system—graphene nanopores—Zwolak, Sahu, and Massimiliano Di Ventra of the University of California, San Diego, simulated conditions that resemble the activity of actual ion channels. For example, the team's simulations demonstrated for the first time that nanopores could be made to permit only some ions to travel through them by changing the diameter of the nanopores etched in a single sheet of graphene or by adding additional sheets. Unlike biological ion channels, however, this selectivity comes from the removal of [water molecules](#) only, a process known as dehydration.

Graphene nanopores will allow this dehydration-only selectivity to be measured under a variety of conditions, another new feat. The researchers reported their findings in recent issues of *Nano Letters* and *Nanoscale*.

In two other preprints, Zwolak and Sahu address some of the complexity in simulating ions' [constriction](#) and [transport](#) through the nanopore channels. When theorists simulate a process, they choose a certain size "box" in which they perform those simulations. The box might be bigger or smaller, depending on the breadth and detail of the calculation. The researchers showed that if the dimensions of the simulation volume are chosen such that the ratio of the width of the volume to its height has a particular numerical value, then the simulation can simultaneously capture the influence of the surrounding ionic solution and such thorny details as nanoscale fluctuations in the diameter of

the pores or the presence of charged chemical groups. This discovery—which the team calls "the golden aspect ratio" for simulations—will greatly simplify calculations and lead to a better understanding of the operation of ion channels, Zwolak said.

More information: Subin Sahu et al. Dehydration as a Universal Mechanism for Ion Selectivity in Graphene and Other Atomically Thin Pores, *Nano Letters* (2017). [DOI: 10.1021/acs.nanolett.7b01399](https://doi.org/10.1021/acs.nanolett.7b01399)

Subin Sahu et al. Ionic selectivity and filtration from fragmented dehydration in multilayer graphene nanopores, *Nanoscale* (2017). [DOI: 10.1039/C7NR03838K](https://doi.org/10.1039/C7NR03838K)

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