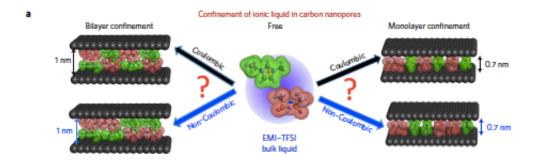


Just squeeze in—researchers discover when spaces are tight, nature loosens its laws

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When packed into pore channels as narrow as a nanometer or less, ions will forgo their typical positive-negative alternating charge ordering. They will form a single (right) or double-file (left) line, many times queuing up next to ions of the same charge. Credit: Drexel University

It turns out that when they're in a hurry and space is limited, ions, like people, will find a way to cram in—even if that means defying nature's norms. Recently published research from an international team of scientists, including Drexel University's Yury Gogotsi, PhD, shows that the charged particles will actually forgo their "opposites attract" behavior, called Coulombic ordering, when confined in the tiny pores of a nanomaterial. This discovery could be a pivotal development for energy storage, water treatment and alternative energy production technologies, which all involve ions packing into nanoporous materials.

In their paper, which was recently published in the journal Nature



Materials, the researchers explain how Coulombic ordering in liquid salts starts to break down when ions are confined in small spaces—specifically carbon pores less than a nanometer in diameter. And the narrower the <u>pore</u>, the less the ions adhere to Coulombic ordering.

"This is the first time breaking of the Coulombic ordering in subnanometer pores has been convincingly demonstrated," said Gogotsi, an author of the paper, who is the Distinguished University and Bach professor in Drexel's College of Engineering. "The breaking of symmetry principals, like Coulombic ordering, plays an essential role in nature. But many of these processes occur without us understanding them and knowing their mechanisms. Science can reveal those hidden processes. And if we understand them, we can eventually develop better technology by working at the same nanometer and subnanometer scales that nature does."

To make its discovery, the team—including researchers from Shinshu University in Japan; Loughborough University in the United Kingdom; The University of Adelaide in Australia; and Sorbonne University, the French Research Network on Electrochemical Energy Storage, and Paul Sabatier University in France—created two sets of carbon nanomaterials. One had pores at least a nanometer in diameter and one with pores less than a nanometer. They then used the materials to draw in ionic liquid as if they were a sponge sopping up water.

In <u>ionic liquids</u>, which are room-temperature liquid salts often used as solvents in the chemical industry, ions are layered in full compliance with the alternating positive-negative pattern of Coulombic ordering. But as the ionic liquid drew into the carbon nanopores it forced the ions to line up in single- and double-file lines. And, like a flock of elementary schoolers running for the bus, they didn't always end up in line next to their usual cohorts.



"In this state, the Coulombic ordering of the liquid is broken," the authors wrote. "Ions of the same charge neighbor each other due to a screening of their electrostatic interactions by the image charges induced in the pore walls."

The team observed this disruption in the natural order of ions through x-ray scattering and modeled the process to explain the experimental observations. They also reported that the non-Coulombic ordering became more pronounced when an electric charge was applied to the carbon material.

"Our results suggest the existence of a molecular-scale mechanism that reduces the Coulombic repulsion energy between co-ions that become closer to each other," they wrote. This mechanism, they theorize, is linked to the charge temporarily imposed on the walls of the carbon pores. This "image charge," they write, offsets the natural electrostatic repulsion of ions of the same charge, to allow the channels to fill with same-charged ions lined up next to each other.

Gogotsi suggests this discovery could make it more feasible to use ionic liquids in batteries and other <u>energy storage</u> devices, which has been examined as a method for making batteries safer but has yet to catch on because it limits their performance.

"We can get safer batteries and supercapacitors when using ionic liquid electrolytes because they are not flammable like the electrolyte solution currently used in these devices," Gogotsi said. "Also, since there is no solvent, the entire volume is occupied by ions and we may be able to store more energy compared to conventional electrolytes that use organic solvents."

He's also looking at this discovery as one that could have a significant impact on the push for water desalination technology. Membranes



currently being developed to turn salt water into drinking water could be improved with this knowledge about ion behavior within subnanometer pores.

More information: Ryusuke Futamura et al, Partial breaking of the Coulombic ordering of ionic liquids confined in carbon nanopores, *Nature Materials* (2017). DOI: 10.1038/nmat4974

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