Graphene-adsorbate van der Waals bonding memory inspires 'smart' graphene sensors

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Monolayer graphene, an atomic-layer thick sheet of carbon, has found immense applications in diverse fields including chemical sensors and detecting single molecule adsorption events electronically. Therefore, monitoring physisorbed molecule induced changes of the electrical response of graphene has become ubiquitous in graphene based sensors. Electric field tuning of the physisorbed molecule-graphene interaction results in enhanced gas sensing due to unique electric field dependent charge-transfer between the adsorbed gas and graphene. Molecular identification in graphene sensors was predicted based on this unique electrically tunable charge-transfer, which is a signature for different adsorbed molecules.

Nevertheless, to achieve molecular identification functionality in graphene sensors, an understanding of the gas adsorption/desorption events and retention of the graphene-gas molecule interaction after turning off the electric field is desired. Until now, the graphene-gas molecule bonding interactions were considered randomized by ambience thermal energy after the electric field is turned off, which is not surprising since these interactions are van der Waals (vdW) bonding and so inherently weak. Nevertheless, this assumed thermal randomization of the graphene-gas molecule vdW bonding was unverified experimentally and a major drawback towards electrically tunable charge-transfer based molecular identification in graphene gas sensors.

To clarify the bonding retention of adsorbed gas molecules on graphene with and without electric field tuning, Osazuwa Gabriel Agbonlahor (current doctoral student), Tomonori Imamura (graduated master's student), Dr. Manoharan Murugananthan (Senior Lecturer), and Professor Hiroshi Mizuta of Mizuta Laboratory at the Japan Advanced Institute of Science and Technology (JAIST) monitored the time-dependent vdW interaction decay of adsorbed CO$_2$ molecules on graphene at different electric fields. Using the electric field to tune the interaction between the adsorbed gas and graphene, the charge-transfer between the adsorbed CO$_2$ molecules and graphene was monitored while the tuning electric field was turned on and after it was turned off. Remarkably, the graphene-gas molecule van der Waals interactions were retained hours after the electric field was turned off, demonstrating both charge-transfer and carrier scattering retention characteristic of the previously applied electric field magnitude and direction i.e. the adsorbed CO$_2$ molecules demonstrated a 'vdW bonding memory'.

Due to this bonding memory, the charge-transfer and scattering properties of the adsorbed gas molecules on graphene can be studied hours after the electric field is turned off which is critical for identifying adsorbed molecules based on their signature charge-transfer response to an applied electric field. Furthermore, the long bonding retention time (over 2h) of these electrically tuned adsorbed molecules, sets graphene-based sensors apart as platforms for developing 'smart' sensors.
suitable for 'beyond-sensing' applications in memory devices and conformational switches.

**More information:** Osazuwa G. Agbonlahor et al, Adsorbed Molecules as Interchangeable Dopants and Scatterers with a Van der Waals Bonding Memory in Graphene Sensors, *ACS Sensors* (2020). [DOI: 10.1021/acssensors.0c00403](https://doi.org/10.1021/acssensors.0c00403)

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