

Bromine-Doped Pentacene Acts as a Phototransistor with Gain over a Large Spectral Range

Recent interest in the development of organic compounds for application as electronic devices has been stimulated in part by the capability of depositing the compounds over large areas on flexible substrates and at a low manufacturing cost. Doped-pentacene thin films have the potential to work as field-effect transistors (FETs) or photovoltaic diodes due to their high charge carrier mobility. Doping of pentacene enhances the benefits obtained from its electronic properties, and this represented an advantage for Jan Hendrik Schön and Christian Kloc from Bell Laboratories/Lucent Technologies when they fabricated an optically activated FET, as they reported in the May 28 issue of *Applied Physics Letters*. The researchers obtained such a device by using bromine-doped pentacene, which becomes a *p*-type semiconductor.

Single crystals of pentacene, 1- μm thick, were grown from vapor in a flow of hydrogen. The source and drain contacts were 300-nm-thick gold strips deposited by shadow masking, and the Schottky gate electrode was a 30-nm-thick aluminum strip. A Xe lamp and a monochromator or a He-Ne laser was used to illuminate the device. As the device is normally off, it develops a potential barrier at the Al-Schottky contact. In this case, the drain current increases with increasing gate bias and saturates with increasing drain voltage. The operative conditions of this device can be manipulated by changing the dopant amount, film thickness, or Schottky barrier height.

Illumination changes the behavior of the drain current and causes the device response to increase abruptly when the energy of the light quanta are equivalent

to that of the absorption edge of the semiconductor. This device seems to act in a similar manner to a photodiode which, however, incorporates the amplification of a FET. Changes in the illumination intensity affect the voltage at the gate and therefore affect the drain current, whose square root was found to vary logarithmically with illumination power. For this device, a constant gain of 8 can be achieved in the energy range between 1.8 eV and 2.5 eV. The authors define gain as the ratio of the total external current to the incident photon flux. In recent experiments, the researchers also observed a similar behavior in iodine-doped pentacene single crystals and thin films.

SIARI S. SOSA

Self-Assembled DNA Basis Serves as Semiconductor in MSM Photodetector

R. Rinaldi and colleagues at Università di Lecce (National Nanotechnology Laboratory of INFN) and Università di Bologna in Italy have fabricated metal-semiconductor-metal (MSM) photodetectors using a layer of a DNA basis. By varying the concentrations of a chloroform solution of this DNA basis—a deoxyguanosine (DG) derivative—the researchers were able to find a concentration that would self-assemble once deposited on a silicon dioxide substrate. The self-assembly took place in an approximately 120-nm-wide space between 5-mm-long gold electrodes. Continued pursuits with these materials may lead to extremely dense logic and memory circuits.

As reported in the May 21, 2001 issue of *Applied Physics Letters*, the researchers used electron-beam lithography to pattern a SiO₂ substrate. Next, they used lift-off processing and layers of chromium and gold to complete the 10- μm -wide electrodes. After completing the electrodes, they used a Hamilton syringe to drop a solution of deoxyguanosine derivative in chloroform between the electrodes on the substrate. When the solvent evaporated, a biological semiconductor material with an energy bandgap of approximately 3.4 eV self-assembled. By superimposing the chemical scheme onto atomic force microscopy images, the researchers demonstrated that the DG had self-assembled into aligned biological wires. However, based on *I-V* characteristics, the researchers believe that charge transport does not occur directly through these wires. The researchers said that, instead, “conductivity is a bandlike phenomenon involving the crystal of DG as a whole.” They said that the effect of the internal dipole induces the electron

donor properties of the DG crystal.

The researchers anticipate further development of molecular-scale electronics as a result of the high responsivity (1 A/W) of this biological material. The molecular scale of this photodetector tempts the researchers with the possibility of bio-molecular planar devices written by ink-jet printers.

PAMELA JOHNSON

Synthetic Clay Removes Radium from Water and Soil

Radium, a natural decay product of uranium, is found where large deposits of uranium are mined, but is also present in many other areas in rocks and soils. Coal and phosphate processing also produce tailings that contain radium. Having tested a variety of synthetic micas for radium removal, Sridhar Komarneni, professor of clay mineralogy at The Pennsylvania State University, has found that sodium-4 mica serves as a prime synthetic clay for this purpose.

Natural mica is a mineral containing a combination of aluminum, silica, magnesium, and potassium. The mineral is found in sheets that are bonded to each other to form a solid, layered mass. Komarneni said that natural mica has a closed structure with all the spaces between layers filled and is not a good ion-exchange medium.

Sodium-4 mica, like natural mica, contains aluminum, silica, and magnesium, but each potassium atom is replaced by two sodium ions, and fluorine is also added. The two sodium ions take up more space than the potassium ion and the layers of mica become offset, creating a space to capture water and radium.

Komarneni said, “Sodium-4 mica has an interlayer spacing of 2.6 Å, too small to capture ions of hydrated sodium, calcium, magnesium, or potassium. Radium, however, is less hydrated and therefore small enough to fit between the layers, as are barium, copper, nickel, and zinc.”

These other transition metals are not usually found in great abundance in radium-contaminated water or in tailings containing radium, so they would not compete for space between the layers. As Komarneni reported in the April 12 issue of *Nature*, when the mica is filled with radium, a shift in the layers occurs and the atoms of radium are trapped between the layered structures.

“Once the radium is trapped, it will not leave the mica,” Komarneni said. “Disposal and storage requirements would then depend only on the radioactivity of the material and not whether radium could

MgB₂ Superconductors Update Available

The discovery of 39 K superconductivity in crystallographically simple MgB₂ in January has launched a flurry of intense international research and development activities, as scientists search for higher transition temperatures (*T_c*) as well as better understanding and improvement of MgB₂'s materials characteristics. See the Materials Research Society Web site for an update on recent findings presented at the 2001 MRS Spring Meeting: www.mrs.org/meetings/spring2001/news/mgb2.html.