

the floating gate makes the structure more tolerant to defects in the tunnel oxide and therefore reduces leakage. This permits the use of thinner tunnel oxides, which in turn reduces the operating voltage, improves endurance and retention, and increases the write/erase rate.

To fabricate their devices, the researchers took advantage of the fact that tungsten silicide on SiO₂ forms elemental tungsten when thermally oxidized. As described in the March issue of *Electrochemical and Solid-State Letters* (p. G71), the team began by growing a 4.5 nm tunnel oxide on silicon substrates, followed by sputter deposition of 8 nm of W₅Si₃ for the floating gate and 10 nm of amorphous silicon. A thermal oxidation step formed the control oxide and the elemental W in the SiO₂ matrix. The W particles had an average diameter of 4.5 nm and an areal density of $3.7 \times 10^{11} \text{ cm}^{-2}$. Fourier transform infrared spectroscopy analysis showed that elemental tungsten was clearly present, with little or no tungsten oxide formation.

The researchers used capacitance–voltage measurements to demonstrate the electron charging effect of their W particles. A bidirectional voltage sweep from 3 V to –4 V and back to 3 V yielded a shift in the threshold voltage ΔV_t of 0.95 V, which is sufficient for binary switching. The researchers found that there was no appreciable decrease in ΔV_t until after 10⁶ cycles and in fact, after 10⁹ cycles, ΔV_t had only dropped to 0.71 V.

The operating characteristics of the W nanoparticle EEPROMs represent an improvement over both the high 7 V operating voltage of conventional floating gate devices and the rapid drop-off seen in the endurance of other nanoparticle devices. In addition, this silicide technique is easily integrated with existing semiconductor technology, making the W nanocrystal devices viable competitors to conventional EEPROMs.

AMANDA GIERMANN

Transparent Hydroxyapatite with High Crystal Orientation Produced by Pulsed Electric Current Sintering

Hydroxyapatite [Ca₁₀(PO₄)₆(OH), HAp] is a major mineral component found in bones and teeth. As a biomaterial, it aids the growth and development of bones and teeth by providing attachment sites for new cells. HAp has a hexagonal crystal structure and there is a large difference in cell attachment between the *a* plane and the *c* plane. Sintered dense HAp bodies with a high degree of crystal orientation are useful for culturing cells and as scaffolds for the regeneration of bones and teeth.

In the January issue of the *Journal of American Ceramic Society* (p. 243; DOI: 10.1111/j.1551-2916.2004.00041.x), researchers Yujiro Watanabe of Hosei University, Japan, and Toshiyuki Ikoma of the National Institute for Materials Science, Japan, and colleagues reported a sintering process for producing dense, transparent, and highly oriented crystalline HAp bodies.

The researchers first prepared the HAp powder by reacting Ca(OH)₂ with H₃PO₄ solution, followed by spray drying and then calcining at 800°C for 3 h. Sintering of the HAp sample was carried out by pulsed electric current sintering (PECS) in a spark plasma sintering (SPS) system. In the PECS process, powder particles are charged with electrical energy and a high pressure is applied on the sample.

During the sintering of HAp, the sample was pressed uniaxially at 50 MPa in vacuum. The sintering temperature was elevated at a rate of 50°C/min to 1200°C. After maintaining the temperature for 10 min, the sample was slowly cooled to 600°C at a rate of 5°C/min. The electric current was then stopped, the pressure was released, and the sample was cooled to room temperature.

The resulting bulk sample has high optical transmittance at wavelengths above 700 nm to at least 1100 nm (i.e., >70% for a 1-mm-thick sample), and a density >99.7% of the theoretical value. As for the crystal orientation, the researchers concluded from x-ray diffraction measurements taken perpendicular and parallel to the direction of the pressure that there is a very significant crystal orientation on the *a*, *b*, and *c* planes. The *c* plane is aligned parallel to the pressure direction, while the *a* and *b* planes are aligned perpendicular to the pressure direction. Orientation indexes of the *a* and *b* planes are much higher than those of samples obtained with other processes.

SHIMING WU

LaPO₄:Eu³⁺ Nanowires Luminesce More Efficiently than Dots

One-dimensional LaPO₄:Eu³⁺ nanowires luminesce more efficiently than zero-dimensional, spherical LaPO₄:Eu³⁺ nanoparticles, said Hongwei Song and co-workers from the Chinese Academy of Sciences. As reported in the March 1 issue of *Optics Letters* (p. 483), the researchers studied the luminescent properties of low-dimensional LaPO₄ doped with rare-earth Eu in order to better understand the fundamental physics of this new condensed-matter system. With the technological drive toward smaller and novel devices, LaPO₄ is

appropriately studied as it is used in fluorescent lamps, cathode ray tubes, and plasma display panels.

The research team reported that the Eu atoms partially filled *f* orbitals, which carry magnetic moments. In addition, Eu³⁺ ions are sensitive activators for use in the study of local symmetry. The researchers used this property of Eu³⁺ to study the structural differences between nanowires, nanoparticles, micron-sized powders, and micron-sized rods.

The LaPO₄:Eu nanowires, nanoparticles, micropowders, and microrods were prepared by a wet chemical method. The colloidal nanoparticles and micropowders were basic and the nanowires and microrods were acid. The diameter of the nanoparticles and nanowires ranged from 10–20 nm, whereas the length of nanowires ranged by several hundred nanometers. The diameters of micropowders ranged over 1–2 μm. The diameters and lengths of the microrods were ~200 nm and 1–2 μm, respectively. X-ray diffraction patterns revealed that the crystal structures of all of the samples belong to the monoclinic monazite type. Time-resolved emission spectra and high-resolution transmission electron micrograph images show that the Eu³⁺ ions occupy two sites in the nanowires as well as the microrods, but not in the nanoparticles or the micropowders. The researchers attributed this result to a degeneration of the crystal field in the micropowders. They also concluded that the observed higher radiative rate for nanowires, as compared with the nanoparticles, micropowders, and microrods, was due to the “variation of the electronic–magnetic dipole field caused by shape anisotropy.”

VIVEK RANJAN

Raman Measurements in Silicon Nanowires Reveal the Diameter Dependence of Phonon Confinement

Phonon confinement in a small nanosystem leads to an asymmetric broadening of the Raman bands. Recently, P.C. Eklund from the Pennsylvania State University, G.U. Sumanasekera from the University of Louisville, and their co-workers determined the evolution of phonon confinement with wire diameter of long, crystalline Si nanowires using the Raman microprobe method.

As reported in the March 1 issue of *Nano-Letters* (p. 409; DOI: 10.1021/nl0486259), the researchers prepared Si nanowires by pulsed Nd:YAG laser vaporization of a Si target containing ~10 at.% Fe. The wire growth process occurs through the vapor–liquid–solid mechanism in which a