

mine the average grain size after crystallization. This could not be done with conventional JMAK analysis because it gives an overall crystallization rate wherein the nucleation and growth rates are coupled. Armed with the ability to determine the rates of nucleation and growth independently, the researchers found them to be constant. By plugging these values into a mathematical expression from the JMAK theory, the researchers found that the experimentally measured grain sizes and the mathematical predictions agree over a broad range of temperatures. This method can be used in practical applications to control grain size and materials properties that are strongly related to microstructure.

TAO XU

Shock-Loading Strengthening Mechanism of Nanocrystalline Materials Revealed

The Hall–Petch effect, in which the strength or hardness of crystalline materials increases inversely with the square root of their grain size, makes clear that grain boundaries (GBs) are natural obstacles for dislocation motions. The limitations of the dislocation motion cause the increase in the hardness and strength of the crystal materials. When the grain size approaches the nanoscale, this strengthening mechanism is no longer applicable, because the grain boundary itself is susceptible to motion (e.g., sliding). Also, the dislocation tends to be unstable at such a small grain size. High-strain-rate deformation experiments, as in shock-loading conditions, have been found to exhibit increased strength in nanocrystalline Ni and Cu. However, the explanation of such a high strain rate (10^4 s⁻¹ to 10^6 s⁻¹) is still not clear, since the high strain rate is increased in conjunction with increasing pressure. Now, a team of researchers led by Eduardo M. Bringa at Lawrence Livermore National Laboratory (LLNL) and Helena Van Swygenhoven at the Paul Scherer Institute (Switzerland) has used molecular dynamics (MD) simulations as well as experimental methods to verify the underlying strengthening mechanism at such shock-loading conditions in nanocrystalline materials.

As reported in the September 16 issue of *Science* (p. 1838; DOI: 10.1126/science.1116723), the researchers used MD simulations to systematically model shock-loading behaviors during the plastic deformation processes of Cu nanocrystalline materials at different grain sizes (5–50 nm) and shockwave pressures (5–47 GPa). The MD simulations showed that with increasing hydrostatic pressure, a nanocrystalline metal Cu that deforms

plastically by GB accommodation (e.g., GB sliding) becomes harder, due to the reduction of the GB sliding and the initialization/emergence of dislocations along the GB. The researchers also performed high-resolution electron microscope (HREM) experiments on Ni nanocrystals after shock-loading at 40 GPa. Ni was used for the experiments because Ni and Cu are fcc materials with similar shock impedances, but Ni has a larger stacking fault energy. The dislocation activity is clearly seen by HREM inside the grains, in excellent agreement with the MD simulation results.

FENGTING XU

Bioelectronic Nanodevices Fabricated on Live Bacteria

Electrically percolating clusters of metal nanoparticles are ideal materials for electronic nanodevices because interparticle current occurs by single-electron transport. The negative surface charges of microorganisms such as viruses, yeasts, and bacteria, make them attractive scaffolds for templating metal nanoparticles. Now, researchers from the University of Nebraska–Lincoln have developed a method for building hybrid bioelectronic devices using gold nanoparticles and live bacteria. Furthermore, a biological response is used to control the electrical response of the devices.

As reported in an article published in the October 21 issue of *Angewandte Chemie, International Edition* (p. 6668; DOI: 10.1002/anie.200501711), V. Berry and R.F. Saraf deposited the gram-positive bacteria *Bacillus cereus* on a silica substrate containing linear gold electrodes 7 μ m apart, 10 mm long, and coated with poly(L-lysine) (PLL). Filtration and ultracentrifugation were used to extract from culture similar-sized bacteria—about 4–6 μ m in length and 0.8–1.0 μ m in diameter. Typically, about 10 bridges formed along a pair of electrodes, which were 10 mm long, with a typical bridge composed of two bacteria. The researchers then immersed the bacteria-deposited chip in a solution containing PLL-coated gold nanoparticles (diameter = 30 nm). Berry and Saraf said that the deposition of the gold nanoparticles, regulated simply by time, is highly selective with formation of a gold monolayer only on the negatively charged bacterial surface, because both the nanoparticles and the substrate are positively charged. The negatively charged teichoic acid (a polyelectrolyte on the bacterium surface) wraps the particle, screening the particle–particle repulsion and making the monolayer electrically percolating. A standard assay showed that the bacteria survived the fabrication process.

The researchers actuated the hydrophilic bacterial cell membranes by controlling the humidity, thereby modulating the separation between gold nanoparticles and concomitantly the interparticle electron tunneling current. Defining a device as a single bacterial bridge and measuring normalized current as a function of relative humidity, the researchers found that a decrease in humidity from 20% to ~0% resulted in a >40-fold increase in tunneling current, even though the interparticle separation decreased by only 0.2 nm. Berry and Saraf said, “We believe that such hybrids will be the key to conceptually new electronic devices that can be integrated with live microorganisms on flexible, plastic-like substrates by using simple chemistry.”

STEVEN TROHALAKI

Covalently Cross-linked Block Copolymer Micelles Encapsulate Magnetic Nanoparticles for Use in Biomedical Applications

Magnetic nanoparticles are currently used in numerous biomedical applications in which stability and surface functionality are crucially important. In the October 5 issue of *Nano Letters* (p. 1987; DOI: 10.1021/nl0513939), B.-S. Kim, J.-M. Qiu, J.-P. Wang, and T.A. Taton from the University of Minnesota reported the encapsulation of magnetic nanoparticles (γ -Fe₂O₃) within amphiphilic block copolymer micelles of poly(styrene₂₅₀-*block*-acrylic acid₁₃), which were subsequently cross-linked to secure the nanoparticles topologically. These “magnetomicelles” are stable to aggregation and have carboxylate groups on their surfaces that are readily modified.

The micelles, formed as a copolymer solution in *N,N*-dimethylformamide (DMF), were added to a second solution of nanoparticles in tetrahydrofuran (THF) in a defined ratio, forming a stable solution. Water was then slowly added to desolvate the copolymer core and nanoparticles simultaneously. Once cross-linked using a diamine cross-linker and a carbodiimide activator, the micelles offer a formidable defense against aggregation, changes in pH, and chemical etching from HCl (a known etchant of γ -Fe₂O₃ nanoparticles). No degradation of the nanoparticle cargo occurred during cross-linking. In addition, Kim and co-workers were able to alter the surface chemistry of the micelles by adding *N*-ethyl-*N'*-(3-dimethylaminopropyl)carbodiimide methiodide (EDC), *N*-hydroxylsulfosuccinimide sodium salt, and the fluorescent compound α -amino- γ -[5(6)carboxyamido fluorescein]-pentaethyleneglycol to the particle suspension. After this

reaction, the nanoparticles showed no structural damage, which was attributed to the protective polymer shell. The extent of functionalization was dictated by stoichiometric addition of the amine reagent. Magnetic susceptibility of the magnetomicelles was probed using superconducting quantum interference device magnetometry. Kim and co-workers showed that micelles containing few nanoparticles ($N_{ave} < 4$) have less interparticle coupling than micelles with many nanoparticles ($N_{ave} > 4$). This change was most likely due to increased first-neighbor distance for the micelles with fewer nanoparticles.

KEVIN P. HERLIHY

Novel Saccharide–Peptide Hybrid Polymers Show Potential for Biomedical Applications

Although a few biopolymers synthesized from natural building blocks exhibit good biocompatibility and have found clinical application, their structural diversity and functionality are limited. Recently, however, researchers from the University of California, Irvine, polymerized saccharide and amino acid monomers to form versatile biomaterials that display properties desirable for biomedical applications.

As reported in the October 14 issue of *Angewandte Chemie, International Edition* (p. 6529; DOI: 10.1002/anie.200501944), Z. Guan and co-workers synthesized three hybrid copolymers from a galactose-derived monomer and one of three different L-lysine-derived monomers. Gel permeation chromatography showed that each copolymer—poly(galactaro dilysine), poly(galactaro trilysine), and poly(galactaro tetralysine)—attained a high molecular weight. Enzymatic degradation studies showed that the polymers were almost completely degraded after 5–7 days. The researchers used a standard assay to demonstrate that their polymers exhibited minimal cytotoxicity, that is, toxicity at the cellular level. In addition, immunogenicity responses measured *in vivo* using rats as animal models showed no evidence of antibody response.

The researchers then evaluated their polymers as a vector for gene delivery—a biomedical application for which the polymers are particularly suited because of the cationic charges they possess at physiological pH. Current synthetic cationic polymers, such as poly(L-lysine) (PLL), condense DNA into particles that can enter cells through endocytosis, but

these polymers are also cytotoxic. Guan and co-researchers used electrophoretic mobility-shift assays to show that their polymers efficiently complexed DNA under physiological conditions. Atomic force microscopy showed that the condensed polymer–DNA particles are spherical, with diameters (50–200 nm) within the range typical for cellular internalization. Using a standard assay, the researchers found that two of their polymers transferred DNA into cells much more efficiently than PLL. The researchers said that “a diverse family of saccharide–peptide hybrid polymers is currently under development in our laboratory for various biomedical applications including gene/drug delivery and tissue engineering.”

STEVEN TROHALAKI

Targeted Delivery of Amphotericin B to Cells Accomplished with Functionalized CNTs

Carbon nanotubes (CNTs) can easily cross cell membranes without damaging them. Recent studies have shown that functionalized carbon nanotubes (f-CNTs) can carry specific drugs to the cells and they are known to be less toxic than existing mechanisms. W. Wu of Institut de



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