

TOPOLOGICALLY CONTROLLED GROWTH OF MAGNETIC-METAL-FUNCTIONALIZED SEMICONDUCTOR OXIDE NANORODS

Marianna Casavola,^{1,} Vincenzo Grillo,² Elvio Carlino,² Cinzia Giannini,³ Fabia Gozzo,⁴*

Enrique Fernandez Pinel,⁵ Miguel Angel Garcia,⁵ Liberato Manna,¹

Roberto Cingolani,¹ Pantaleo Davide Cozzoli¹

¹ National Nanotechnology Laboratory of CNR-INFM, Unità di Ricerca IIT, Distretto Tecnologico ISUFI, via per Arnesano km 5, 73100 Lecce, Italy

² TASC-INFM-CNR National Laboratory, Area Science Park - Basovizza, Bld MM, SS 14, Km 163.5, I-34012 Trieste, Italy

³ CNR-Istituto di Cristallografia (IC), via Amendola 122/O, I-70126 Bari, Italy

⁴ Swiss Light Source, Paul Scherrer Institute, 5232 Villigen PSI, Switzerland

⁵ Institute of Applied Magnetism and Department of Materials Physics, UCM, P. O. Box 155, 28230 Las Rozas, Madrid, Spain

* Presenting author: Phone:+39 0832 298231. Fax: +39 0832 298238.

E-mail: marianna.casavola@unile.it

The development of colloidal hybrid nanocrystals (HNCs), in which two or more material sections with peculiar chemical, optical, magnetic and/or catalytic properties are connected by epitaxial interfaces, represents an emerging field of great interest in nanoscience.¹ Nanocrystal heterostructures, characterized by a topologically controlled distribution of their chemical composition, are extremely attractive as advanced generations of nano-objects potentially able to perform multiple tasks in optoelectronic devices, biomedical engineering, diagnostics, sensing, and catalysis.⁽²⁻¹⁰⁾ To date, the chemical fabrication of HNCs remains a challenging task, as the ability to tailor the size and shape of the individual material domains has to be integrated with the control over additional parameters at the nanoscale, such as inter-compound miscibility, interfacial strain, and facet-specific chemical reactivity.

Here we present a seeded growth approach to synthesize a novel type of colloidal semiconductor/magnetic-metal HNCs, each made of spherical ε-Co domains epitaxially attached to one anatase TiO₂ rod-shaped portion.⁽¹⁰⁾ We have been able to control the heterogeneous nucleation of spherical ε-Co domains onto preformed TiO₂ nanorods in suitable surfactant mixture, achieving metal deposition on either the tips or on multiple locations along the longitudinal sidewalls of the oxide seeds. A detailed compositional, structural, and magnetic characterization of the as-prepared heterostructures has been carried out by combining powder X-ray Diffraction, high angle annular dark field (HAADF) imaging, and high-resolution transmission electron microscopy (HRTEM) analyses, superconducting quantum interference device (SQUID) magnetic measurements. Our results suggest that the possibility of switching between either TiO₂-Co HNC topologies arises from the facet-dependent chemical reactivity of the TiO₂ seeds, which is mainly governed by surface

selective adhesion of the surfactants, rather than by misfit-related interfacial strain at the relevant junction points. These TiO₂-Co HNCs could find relevant applications as bi-functional, magnetically recoverable (photo)catalysts and as active elements in novel magneto-optical applications.

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