

## Thesis abstract

# Functional magnetic interface phenomena in nano-architectures

Grace L. Causer

Abstract of a thesis for a Doctorate of Philosophy submitted to University of Wollongong

The work embodied in this thesis aims to investigate the occurrence of magnetic interface phenomena in low-dimensional thin-film systems which have conceivable utility in future condensed-matter technologies. Namely, the magnetic interface quality of an FePt<sub>3</sub> nano-magnet formed via ion-induced chemical disorder will be critically analysed, in addition to a Co/Pd bilayer which features modifiable magnetic surface anisotropy upon exposure to hydrogen gas. The studies are enabled chiefly through advanced X-ray and neutron scattering techniques specifically chosen to probe interface structure as well as chemical and magnetic orders, and supplemented by traditional lab-based characterisation tools.

To begin, a much-anticipated experimental confirmation of the intrinsic sharpness of magnetic interfaces formed by locally driving magnetic phase transitions in materials using ion beams is presented. This is achieved through a unique experimental design whereby a room-temperature ferromagnetic nano-layer is encoded with depth-control onto a paramagnetic FePt<sub>3</sub> film by inducing chemical disorder using energy-specific He<sup>+</sup> ions. The magnetic transition is investigated through theoretical modelling, whereby the first density functional theory results for the entire suite of potential long-

range magnetically ordered states of FePt<sub>3</sub> are presented. In doing so, the energetically favourable ground-state spin structure is identified. By analysing several localised defect structures which may form in FePt<sub>3</sub> under ion irradiation, the fundamental mechanism of the disorder-driven magnetic transition is revealed and shown to be caused by an intermixing of Fe and Pt atoms in anti-site defects above a threshold density.

In a second study, hydrogen-induced modifications to the layer-averaged static magnetisation and macroscopic magnetodynamic behaviours of a Co/Pd heterostructure are investigated. The modifications are observed and examined in detail through simultaneously probing the magnetic anisotropy energy and studying the changing chemical and magnetic depth-profiles across the entire bilayer during primary hydrogen gas absorption. It is revealed that the in-plane interfacial magnetisation of the Co/Pd bilayer irreversibly increases after primary hydrogen-gas absorption, indicating a weakening of the perpendicular magnetic anisotropy energy. To aid in conducting this analysis, an original experimental method is first developed which innovatively combines neutron scattering and microwave spectroscopy; equipment is then commissioned, and feasibility studies are performed.

JOURNAL & PROCEEDINGS OF THE ROYAL SOCIETY OF NEW SOUTH WALES  
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Dr Grace L. Causer,  
Institute for Superconducting and Elec-  
tronic Materials and School of Physics,  
University of Wollongong,  
Wollongong NSW 2018

E-mail: [grace.causer@tum.de](mailto:grace.causer@tum.de)  
URL: <https://ro.uow.edu.au/theses1/411>