

Seminar announcement

Tuesday, October 24, 2023 4 pm WSI, Seminar room S 101

"Developing scalable routes to photocatalytic coatings; Exploring the chemical vapour deposition of metal oxidebased materials for NO_x remediation and solar water splitting"

<u>Abstract</u>: For any technology to be translated from the laboratory to industry, the fabrication process should be inherently scalable and its components should be economical. In the Solar Coatings Group, we are focussed on developing scalable synthetic routes to photocatalytic materials, which can enable their uptake by industry. We are working on a number of technologies, and the focus of this talk will be on two of them: (i) photocatalytic coatings for window glass that can remediate surrounding NOx pollution using ambient light and (ii) photocatalytic coatings for photoelectrochemical devices that can split water and produce clean H2 fuel from sunlight. In both cases we are producing materials on the 50 cm2 scale using chemical vapour deposition (CVD) (Figure 1a); a technique used widely by industry to grow coatings at scale with nanoscale control. These two aspects of my talk are summarised below:

NOx pollution

NOx (NO + NO2) pollution is a major cause of poor health, and has a range of detrimental environmental consequences [1]. NOx pollution can be remedied by photocatalysis using TiO2, which can use ambient light to oxidise this toxin into the comparatively benign nitrate (NO3-) [2]. Photocatalytic coatings of TiO2 are mass produced on window glass using chemical vapour deposition (CVD), but currently show nominal activities towards NOx remediation [3].

In this section of my talk, I will present how we used aerosol-assisted CVD (AA-CVD) to produce ~50 unique TiO2 coatings on window glass by systematically varying the process parameters. The coatings produced showed wide ranging physicochemical properties, including differences in phase composition (anatase/ anatase: rutile composite), optical bandgap energy, surface roughness, charge carrier generation and charge carrier lifetime. Their photocatalytic activities towards NO gas was measured using ISO protocol (ISO 22197-1:2016) [4], and the observed relationships between the processing parameters, the physicochemical properties and photocatalytic activity are discussed.

Water splitting

Current H2 demands are vast, with the industry valued in excess of \$150 billion. Today, most H2 is produced from the nonrenewable reformation of natural gas, and accounts for ~3% of total CO2 emissions. Various renewable methods of producing H2 are being developed, with solar-driven photoelectrochemical (PEC) water splitting seen as one of the most promising routes in terms of efficiency and potential cost [5].

Bismuth vanadate (BiVO4) has emerged as one of the most PEC materials in water splitting devices. State-of-the-art systems have achieved solar-to-hydrogen (STH) efficiencies above 8%; nearing benchmark efficiencies for commercial viability [6].

In this talk, I will present the work we have done in my group on the development of scalable AA-CVD routes to high performance BiVO4-based photoanodes (Figure 1b). I will also show how the intrinsic charge carrier behavior in these photoanodes was studied using time-resolved optical spectroscopies to reveal the kinetics of water oxidation, and provide insight on the reaction mechanism. And lastly, I will present our work on developing prototypes, incorporating these BiVO4-based PECs into custom made reactors (Figure 1c).

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