

Special issue on “Fluorescent probes”

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Fluorescent probes are intelligently designed molecules that transform the act of binding/reacting with a target analyte into an optical signal [1,2]. Their high sensitivity and specificity coupled with high spatial and temporal resolution offers a non-invasive approach to observe a specific analyte within complex biological environments in real time with high precision [3]. These “molecular tools” have the capability to interrogate the role of specific biomarkers that are associated with a particular disease state or can be used to improve our understanding of the role or certain species in a biological pathway [4,5]. In addition to their biological applications, fluorescent probes have found use for the detection of environmental, agricultural, and industrial pollutants, making them critically important for public health and safety [6].

This special issue on “fluorescent probes” serves to highlight and illustrate various approaches towards fluorescent probes and their potential applications. Gunnlaugsson and co-workers have developed a series of 4-amino-1,8-naphthalimide fluorescent probes to evaluate the substitution effects on the extent of pH dependent photoinduced electron transfer (PeT) quenching [7]. These molecules were used as pH dependent lysosomal imaging agents. Biological processes are complex involving more than one biomarker, this has led to the use of fluorescence based molecular logic probes [8–14], an example includes the development of a dual enzyme activated fluorescent probe by Odyniec et al. [15]. Fluorescence imaging often suffers from poor light penetration and high background interference from autofluorescence of endogenous proteins. To overcome these limitations, researchers have begun to pursue near infrared (NIR) imaging agents [16,17]. O’Shea and co-workers report on a series of NIR aza-BODIPY fluorophores for the potential application of fluorescence guided surgery [18]. Peng and co-workers report on a NIR fluorescent probe for the subcellular imaging of the COX-2 enzyme in cancer cells [19].

Porphyrin-based systems have unique photophysical properties with the ability to form a range of metal complexes [20]. As such, Sessler and co-workers exploited the metal coordination properties of the expanded porphyrin Texaphyrin to detect heavy metals in drinking water [21]. Whereas Payne and co-workers have developed a diporphyrin tweezer to measure enantiomeric excess [22]. Azulene has unique photophysical properties [6], this prompted Lewis and co-workers to exploit the organic transformation of an aminoazulene to a diazo-azulene for the colorimetric detection of nitrite in drinking water [23]. Fluorescent probes are not just limited to small molecules, Minami and co-workers developed a “ligand-free” conjugated copolymer for the fluorescence detection of Copper(II) [24]. Giordani and co-workers synthesized *N*-heterocycle-based fluorophores attached to multilayer fullerenes—nano-onions for material-based applications [25]. Lastly, James and co-workers have developed a fructose responsive fluorescent hydrogel as a first step on the road to develop saccharide responsive molecular devices for clinical applications [26].

Upon reading this editorial, if you have found yourself intrigued and excited about the endless possibilities of fluorescent probes. Please join us at the 7th International Conference on Molecular Sensors & Molecular Logic Gates (MSMLG 2020) a celebration of the career of Seiji Shinkai in Reno, Nevada.

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Selfie of Tony D. James (Left) and Adam C. Sedgwick (Right) at the Weizmann Institute in Israel (January 2019)

Adam C. Sedgwick is a Postdoctoral research fellow working under the supervision of Prof. Jonathan L. Sessler at The University of Texas in Austin, Texas. He obtained his PhD with Prof. Tony D. James and Prof. Steven D. Bull at the University of Bath focusing on the development of new synthetic transformations for application in fluorescence sensing. This includes the development of a single molecule that can simultaneously detect multiple biologically relevant analytes at one time (molecular logic gates). His current research interests are in the realms of stimuli-responsive materials, molecular imaging agents, and theranostics agents. To date he has published > 40 papers in international peer reviewed journals and has a h-index of 15 (December 2019, google scholar).

Tony D. James is a Professor at the University of Bath and Fellow of the Royal Society of Chemistry, who currently holds a prestigious Royal Society Wolfson Research Merit Award (2017–2022). He has developed a broad interdisciplinary approach to research, with an underpinning focus on the development of modular sensors where he has pioneered a range of reporting regimes.

He has been a visiting professor at Tsukuba, Osaka, Kyushu and Sophia Universities, an AMADEus invited professor at the University of Bordeaux and is a guest Professor at East China University of Science and Technology, Xiamen University, Shandong Normal University, Nanjing University, Shaanxi University of Science and Technology, Changzhou University, Zhejiang University, Qufu Normal University, Henan Normal University and is a Hai-Tian (Sea-Sky) Scholar at Dalian University of Technology.

He has published over 314 publications, including two books, 9 book chapters and 303 papers in international peer reviewed journals. He has an h-index of 67 (December 2019, google scholar). In addition, he received the Daiwa-Adrian Prize for developing scientific networks with Japan in 2013, the inaugural CASE Prize for establishing scientific networks with China in 2015, and the MSMLG Czarnik Award in 2018. Since 2019, he has been a member of the editorial board of *Frontiers of Chemical Science and Engineering*.