

## RECENT ADVANCES IN PHOSPHORUS CHEMISTRY

Peruzzini M.

*Istituto di Chimica dei Composti Organo Metallici (CNR – ICCOM) and Dipartimento di Scienze Chimiche e Tecnologie dei Materiali, (CNR – DSCTM), Consiglio Nazionale delle Ricerche, Via dei Taurini, 19 – 00185 Rome – Italy; e-mail: maurizio.peruzzini@iccom.cnr.it*

In this lecture some of the most recent achievements in the field of the chemistry of elemental phosphorus, deriving from the author's own research in Florence (Italy) during the last two decades, will be presented and discussed.

The lecture will encompass the three allotropes of elemental phosphorus with particular emphasis to the extremely reactive and toxic white allotropic form and, on the opposite end, to the incredibly inert black allotrope. More in details, the lecture will include:

*i)* the ruthenium mediated activation of white phosphorus pointing out the unusual and fascinating hydrolytic degradation of the P<sub>4</sub> molecule which follows its η<sup>1</sup>-coordination to the metal. The stepwise hydrolysis results in a variety of unusual P<sub>x</sub> fragments (x ≤ 4), such as P-oxyacids, mono-, di-, and triphosphanes and hydroxyphosphanes, stabilised by coordination to one or two metal centers;<sup>1</sup>

*ii)* our most recent results in the chemistry of the less reactive allotrope of the element, i.e. black phosphorus, which may be easily exfoliated to form 2D-flakes of phosphorene (the all-P analogue of graphene).<sup>2</sup> This new material exhibits intriguing electronic properties and behaves as a perfectly tailored platform to host transition-metal nanoparticles<sup>3</sup> and, likely, to coordinate different transition metal fragments.<sup>4,5</sup>

*MP thanks all the coworkers listed in the references for their invaluable help in developing this chemistry. Thanks are also expressed to the European Research Council (ERC Grant Agreement No. 670173) for funding the project PHOSFUN “Phosphorene functionalization: a new platform for advanced multifunctional materials” through an ERC Advanced Grant. Finally, the Italian Ministry for Education and Research (MIUR) is kindly acknowledged for financial support through Project PRIN 2015 (grant number 20154X9ATP).*

## References

1. P. Barbaro, C. Bazzicalupi, M. Peruzzini, S. Seniori Costantini, P. Stoppioni *Angew. Chem. Int. Ed.* 2012, 51, 8628 - 8631 and references therein.
2. M. Peruzzini, R. Bini, M. Bolognesi, M. Caporali, M. Ceppatelli, F. Cicogna, S. Coiai, S. Heun, A. Ienco, I. Iglesias, A. Kumar, G. Manca, E. Passaglia, D. Scelta, M. Serrano-Ruiz, F. Telesio, S. Toffanin, M. Vanni *Eur. J. Inorg. Chem.* 2019, 41, 1476 – 1494.
3. M. Caporali, M. Serrano-Ruiz, F. Telesio, S. Heun, G. Nicotra, C. Spinella, S. Caporali, M. Peruzzini *Chem. Commun.* 2017, 53, 10946 – 10949.
4. M. Peruzzini et al. to be published.
5. For a theoretical analysis of the bonding capabilities of phosphorene see: A. Ienco, G. Manca, C. Mealli, M. Peruzzini *Dalton Trans.* 2018, 47, 17243 – 17256.