

# Reactivity of Polar Organometallic Compounds in Unconventional Reaction Media: Challenges and Opportunities

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This paper is gratefully dedicated to the memory of Dr. Guy Lavigne

**Abstract:** Developing new green solvents in designing chemical products and processes or successfully employing the already existing ones is one of the key subjects in Green Chemistry and is especially important in Organometallic Chemistry, which is an interdisciplinary field. Can we advantageously use unconventional reaction media in place of current harsh organic solvents also for polar organometallic compounds? This Microreview critically analyses the state-of-the-art on this topic and showcases recent developments and breakthroughs which are becoming new research directions in this field. Because metals cover a vast swath of the periodic table, the content is organised into three Sections discussing the reactivity of organometallic compounds of s-, p- and d-block elements in unconventional solvents. The richness of bibliography reported witnesses the genuine, burning thirst for a deeper knowledge of this field, and forecasts an ever-bright future for Organometallics in Green Chemistry.

## 1. Introduction

“There are times when one can sense a sea change, a shift in the order of things that is profound and fundamental”.<sup>[1]</sup> A silent, but contagious revolution is taking place in the way of thinking and practising Organometallic Chemistry (OC) by academic and industrial groups worldwide, which is mainly driven by new insights, needs, and evidence on the horizon. OC plays an essential role across a wide spectrum of science, technology, medicine and industry, with a heavy impact on the environment, and still remains a core subject within the “grand challenges” or “big themes” (e.g., energy, materials, medicine) towards which priorities and policy often focus.

International strategies launched by institutions and organizations, such as The American Chemical Society’s (ACS) Green Chemistry Institute Pharmaceutical Roundtable (GCIPR), strive for the need to replace conventional hazardous volatile organic compounds (VOCs) in favour of safe, green and biorenewable reaction media that are not based on crude petroleum.<sup>[2]</sup> One large area of consumption of petroleum-based

chemicals in chemical transformations is, indeed, solvents used as reaction media, which account for 80–90% of mass utilization in a typical pharmaceutical/fine chemical operational process. Thus, the solvent itself is often a critical parameter especially in drug product manufacturing and is as well responsible for most waste generated in the chemical industries and laboratories.<sup>[3]</sup>

Following these considerations, some of the most critical and intriguing questions that arise are: Can we get traditional organic solvents out of organometallic reactions?<sup>[4]</sup> Can we use protic, recyclable, biodegradable, and cheap unconventional solvents also for highly reactive organometallic compounds? Answering these questions would not only mean to break new ground towards sustainable solutions to the aforementioned challenges, but it could also be rewarding from an intellectual point of view in order to investigate to what extent a certain organometallic compound does not react with the intended unconventional solvent and, if that is the case, to explain why this occurs.

In this Microreview, recent selected contributions published in the literature tackling the above timely topics have been highlighted, but seminal references have also been critically analysed. The contents are organized into three main sections with subheadings according to the nature of the Metal–Carbon (M–C) bond. The impact played by unconventional solvents (e.g., water, deep eutectic solvents, ionic liquids, and supercritical CO<sub>2</sub>) on the chemistry of compounds of s-, p- and d-block elements has been discussed. Topics that have recently been reviewed are not further detailed here.

## 2. Organometallic compounds of s-block elements

Within the periodic table, the s-block elements are the 14 elements contained in the first two columns (Groups 1A and 2A) plus helium. They are unified by the fact that their valence electrons are in an s orbital, and are very reactive due to highly polar M–C bonds. In this section, we will deal with the reactivity of organolithium and Grignard (organomagnesium) reagents in unconventional solvents.

### 2.1. Reactivity in protic reaction media (water and deep eutectic solvents)

Organolithium and Grignard reagents are among the most useful and versatile organometallic compounds in chemical synthesis, and functionalised organometallic species are very useful intermediates for the synthesis of many organic non-natural and natural products.<sup>[5]</sup> Opening chapters in classic organic textbooks, however, emphasise the need of the strict control of anhydrous conditions and the use of water-free reaction media for the successful handling of organometallic

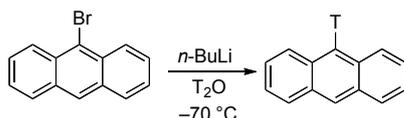
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compounds with highly polarised M–C bonds. Thus, at first sight, it might sound ridiculous to think to a protic solvent (e.g. water) as a potential “additive” (or, even worse, as a full component) for reaction media for carrying out s-block-metal-mediated organic transformations as these organometallics are extremely sensitive to traces of air and moisture.<sup>[6]</sup> Nevertheless, a perusal of the modern and present-day literature reveals, from time to time, some “perplexing” reactions strangely “accelerated” by water. These deserve consideration and still need an explanation.

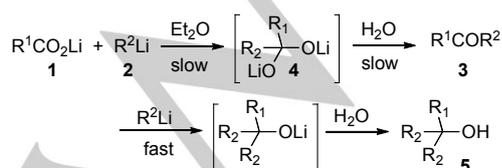
One of the first examples reported is the following. In order to label aromatics via hydrolysis of organometallics intermediates with tritiated water (T<sub>2</sub>O), Taylor made the observation that the most convenient way for the preparation of tritiated arenes was the addition of *n*-BuLi to a mixture of a bromoarene and *wetted* (T<sub>2</sub>O) sodium-dried Et<sub>2</sub>O. This result is consistent with a lithium-bromine exchange reaction surprisingly occurring “at least as fast and most probably faster” than the expected reaction of *n*-BuLi with T<sub>2</sub>O (Scheme 1).<sup>[7]</sup> The relative rates of deprotonation and of halogen-lithium exchange by organolithium compounds, however, have been a matter of controversy in the following years.<sup>[8]</sup>



**Scheme 1.** Preparation of 9-tritium-labelled anthracene.

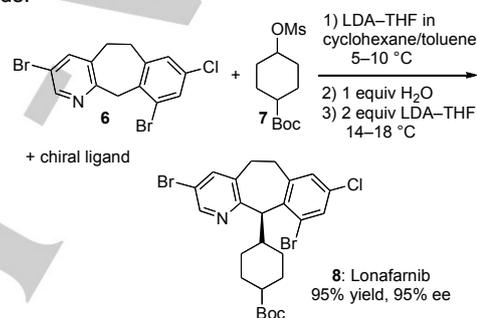
Lithium carboxylates **1** are known to react with organolithium compounds **2** in Et<sub>2</sub>O to give high yields of the corresponding ketones **3** after considerable time of reflux (at least 24 h). Under these conditions, tertiary alcohols **5** are not usually formed. However, if the reflux time is shorter (e.g. 30 min), mixtures of ketones **3** and tertiary alcohols **5** are alternatively produced upon quenching with H<sub>2</sub>O. This result implies that the excess organolithium **2** reacts *during the hydrolysis* with part of the ketone, which is formed by the hydrolysis of the intermediate adduct **4**, *faster* than it reacts with water (Scheme 2).<sup>[9]</sup> In highlighting these results,<sup>[10]</sup> Keith Smith finally commented: “Shall we await the day when reactions of organolithiums are routinely performed in aqueous solution?”

In general, Lewis basic solvents increase the reactivity of organolithiums as they become an integral part of the organolithium aggregate and, once used as additives or ligands, they sometimes proved to be effective in contributing to the optimization of organolithium reactions.<sup>[11]</sup>



**Scheme 2.** Preparation of ketones and tertiary alcohols by reacting organolithium compounds with lithium carboxylates.

For instance, in the asymmetric LDA-mediated synthesis of the anticancer Lonafarnib **8**, a unique *water effect* on the enantioselectivity was discovered. In the key alkylation step, an LDA–THF complex in cyclohexane was added to a toluene solution containing the tricyclic substrate **6**, the chiral norephedrine-based mediator, and the alkylating agent **7**. Counter-intuitively, the highest ee (95%) and the best yield (95%) in **8** were achieved once *one equivalent of water* was sequentially added to the above reaction mixture, then compensated by an additional equivalent of LDA. In the absence of water, both the ee and the yield in **8** dramatically dropped down to 50% (Scheme 3).<sup>[12]</sup> As an additional example, carboalumination of alkynes has been proved to occur within minutes at –23 °C (and is fast even at –70 °C) in the presence of *stoichiometric amounts of water* leading stereoselectively to alkenes.<sup>[13]</sup> To understand this *water effect* is extremely important to study the interaction of water with organometallic compounds.



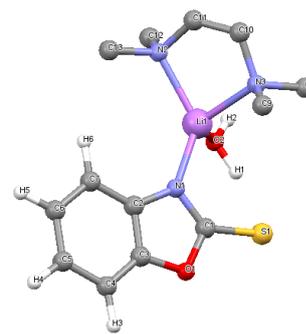
**Scheme 3.** Stereoselectivity achieved in the key alkylation step by adding water.

We have always been taught about the crucial role played by water in Life Sciences. Among its properties, it is worth mentioning its extraordinary capability to engage in strong intermolecular hydrogen bonding with a plethora of Lewis acid and basic sites, thereby promoting a self-organization in supramolecular ordered structures. Could the water play a similar role in organometallic compounds? Careful, systematic, and controlled hydrolysis studies performed by Roesky and co-workers of kinetically stabilised Group 13 trialkyl compounds, supported by spectroscopic and crystallographic evidences, have led to the isolation and the structural characterization of interesting intermediate oligomeric compounds that eventually result in final policondensed metalloxane clusters.<sup>[14]</sup> More recent work disclosed the synthesis of organoaluminum hydroxides<sup>[15]</sup> and of *three-in-one* clusters<sup>[16]</sup> comprising two tetranuclear aluminophosphate units and a tetrameric alumino hydroxide unit. These examples testify that water plays a major role also in organometallic chemistry contributing to the coordination sphere of the metal, and promoting intermolecular interactions and extensive self-assembly.

What about the “role of water” in the reactions of organometallic compounds of s-block metals? The hydrolysis of organolithium compounds by water or other proton donors is often assumed to be a very simple process yielding

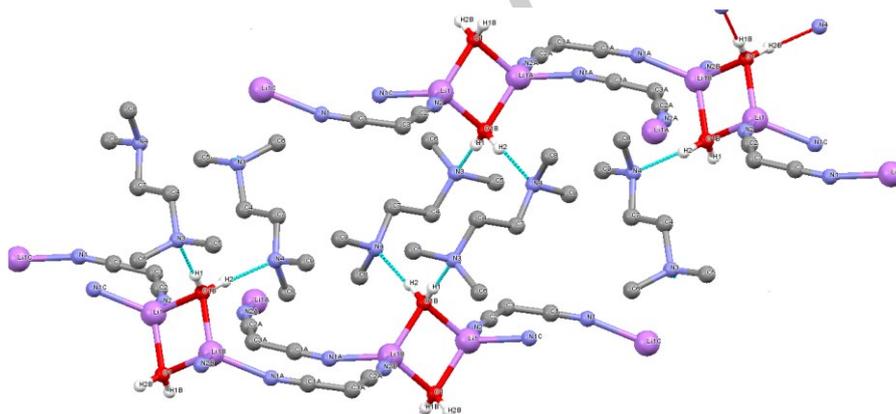
quantitatively the corresponding organic acid and LiOH.<sup>[17]</sup> As a matter of fact, it may not be as simple as is commonly believed. The rate of protonation of Et<sub>2</sub>O solution of PhLi and PhCH<sub>2</sub>Li and their O-deuterated analogues by water or alcohols, for instance, shows small isotope effects (1.0–1.5), which implies that the rate-determining step is the displacement of Et<sub>2</sub>O from the organolithium compound by the oxygen of the “acid”.<sup>[18]</sup> Could water act as a polar ligand towards Li<sup>+</sup> centres? The first lithiated organic compound containing water as a ligand was published by Wright and co-workers.<sup>[19]</sup> Lithiation of 2-mercaptobenzoxazole, in the presence of *N,N,N',N'*-tetramethylethylenediamine (TMEDA) and H<sub>2</sub>O (both adventitious and deliberately added to the reaction mixture), produced the monomeric complex **9** (Figure 1) exhibiting a strong hydrogen bonding between one proton of the coordinated H<sub>2</sub>O and the polarised sulfur centre of the organic anion, rather than a protonated Li–OH⋯SH species.

Such a molecular structure, as suggested by the authors, can be viewed as a model for how hydrolysis of organolithium compounds might take place. Analogously to what has been ascertained in the case of organoalanes,<sup>[14]</sup> this process might proceed as well via preliminary organolithium–water complex formation followed by proton transfer to the carbanion.



**Figure 1.** Molecular structure of the complex (**9**) between lithiated 2-mercaptobenzoxazole and TMEDA.

The relative rate of these processes, however, might also be influenced by the nature of the aggregate involved because of the strong structure–reactivity relationship in organolithium compounds.<sup>[5b,c]</sup> These findings were then also extended to alkaline-earth metal complexes,<sup>[20]</sup> with the synthesis of aqua complexes carried out by “assembling” H<sub>2</sub>O ligands via solid metal hydroxides in a hydrocarbon solvent and in the presence of a Lewis base.



**Figure 2.** Molecular structure of the 1:1:1 complex (**10**) among lithiated malonodinitrile, TMEDA, and H<sub>2</sub>O.

The first structural characterisation of an H<sub>2</sub>O-containing complex of the lithium salt of an organic molecule containing an acidic C–H bond was also reported.<sup>[21]</sup> The three-dimensional polymeric structure **10** depicted in Figure 2 discloses an intriguing complex of lithiated malonodinitrile, TMEDA, and H<sub>2</sub>O in a 1:1:1 molar ratio. What is remarkable here is (a) the lack of TMEDA–Li interactions which is unprecedented in lithium chemistry especially considering that TMEDA is a bidentate ligand compared to water, which should favor complexation to lithium, and (b) the ability of *each* water molecule to engage simultaneously in hydrogen bonding to two TMEDA molecules (donor function) and to two lithium atoms (acceptor function) (Figure 2).

The intrinsic reactivity of a series of monomeric allylmetals with water and carbonyl compounds has been recently addressed by a theoretical study.<sup>[22]</sup> Interestingly, calculations

suggest that intrinsic kinetic preference of allylation over hydrolysis correlates quite well with the reactivity of hydrolysis. That is, a higher activation energy of hydrolysis corresponds to a higher kinetic preference toward allylation. However, the sole *polarity of the C–M bond* does not fully account for the reactivity of hydrolysis, but both *nucleophilicity* of the allylmetal and *thermodynamic driving forces* are likely significantly to contribute to the barrier of hydrolysis. Data relative to the organometallic compounds of the s-block elements also suggest that both  $\pi$ -complexes of Li and polarised  $\sigma$ -complexes of MgBr<sub>2</sub> may hydrolyse or allylate preferentially, *depending on the employed experimental conditions*. In particular, calculations for the reaction of allylMgBr with water and acetone show that both activation energies of hydrolysis (4.5 kcal/mol) and allylation (5.7 kcal/mol) are quite similar. This competition was later experimentally investigated by Madsen and Holm.<sup>[23]</sup> When

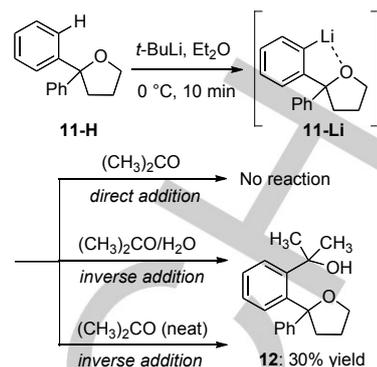
allylMgBr was reacted with either acetone or PhCHO in the presence of water (inverse addition), the yields of the corresponding addition products were found to be 91 and 75%, respectively. Grignard reagents are complex mixtures of the Schlenk components alkyl(aryl)magnesium halide, dialkyl(diaryl)magnesium, and magnesium halide salt, solvated by an ethereal solvent, and undergoing very fast chemical exchange in solution (Scheme 4).<sup>[59]</sup>



Scheme 4. The Schlenk equilibrium.

The position of the equilibrium is influenced by solvent, temperature, and the nature of the substituents. It is shifted left to right in strongly donating solvents (e.g., THF or dioxane) because the stabilization due to the interaction with solvent molecules decreases in the order:  $\text{MgX}_2 > \text{RMgX} > \text{R}_2\text{Mg}$ , that is according to the Lewis acidity of the various components.<sup>[24]</sup> Thus, the above results would seem to indicate that water preferentially coordinates magnesium, the complexation energy with one water molecule being  $-23.1$  kcal/mol.<sup>[22]</sup> It may be that “coordinated” water is less reactive in the protonation reaction, thereby allowing the addition reaction to take place more competitively. A possible “scavenging” effect toward water promoted by any electrophilic magnesium compound present in solution was supported by an experiment in which allylMgBr was added to a mixture of acetone–water in the presence of an extra  $\text{MgBr}_2$ . In this case, the yield of the addition product was quantitative. Benzylmagnesium chloride also proved to react sufficiently faster with acetone and benzaldehyde rather than being quenched with water, whereas butylmagnesium bromide yielded only traces of the expected addition products.

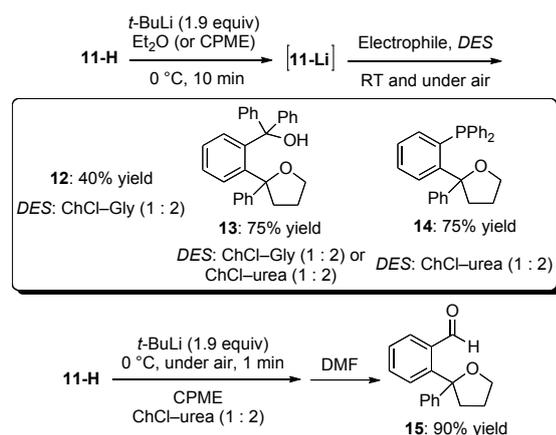
During the investigation of the directing ability of the tetrahydrofuranyl moiety in promoting regioselective *ortho*-lithiation/functionalisation of diaryltetrahydrofurans, an unexpected reactivity was observed by Capriati and co-workers in screening for electrophiles.<sup>[25]</sup> While no reaction was detected upon adding acetone to a dry- $\text{Et}_2\text{O}$  solution of the putative *ortho*-lithiated intermediate **11-Li** (generated via lithiation of the parent precursor **11-H** with *t*-BuLi at  $0^\circ\text{C}$  for 10 min (direct addition)), the expected hydroxyalkylated adduct **12** could instead be isolated in 30% yield once an  $\text{Et}_2\text{O}$  solution of **11-Li** was added over an acetone–water mixture (6 equiv each) at room temperature (inverse addition). It was ascertained, however, that water apparently did not boost any “rate acceleration” of the reaction as a similar yield was obtained in *neat conditions*, that is in the absence of water, which simply acts as a bystander (Scheme 5). The employment of cyclopentyl methyl ether (CPME) as an alternative environmentally friendly reaction medium considerably increased the yield of some adducts (e.g., EtI: 0% ( $\text{Et}_2\text{O}$ ), 80% (CPME);  $\text{Ph}_2\text{CO}$ : 40% ( $\text{Et}_2\text{O}$ ), 90% (CPME); 4- $\text{ClC}_6\text{H}_4\text{CHO}$ : 40% ( $\text{Et}_2\text{O}$ ), 80% (CPME);  $\text{Ph}_2\text{PCI}$ : 60% ( $\text{Et}_2\text{O}$ ), 85% (CPME)), but again **12** was isolated with a yield not higher than 30% (direct addition), enolization most probably still competing a lot with nucleophilic addition.



Scheme 5. *ortho*-Lithiation/functionalisation of diphenyltetrahydrofuran with acetone.

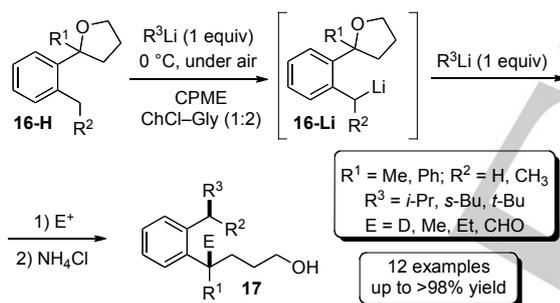
The potential impact of protic solvents in the above functionalisations was further investigated employing the so-called “deep eutectic solvents” (*DESs*). The concept of *DES* was firstly introduced by Abbott and co-workers to describe the formation of a liquid eutectic mixture (mp  $12^\circ\text{C}$ ) starting from two solid materials with high melting points: choline chloride ( $\text{ChCl}$ , mp  $133^\circ\text{C}$ ) and urea (mp  $302^\circ\text{C}$ ) in a ratio 1:2 (1 $\text{ChCl}$ /2 $\text{Urea}$ ).<sup>[26]</sup> *DESs* are today generally defined as combinations of two or three safe and inexpensive components which are able to engage in hydrogen bond interactions with each other to form an eutectic mixture with a melting point lower than either of the individual components.<sup>[27]</sup>  $\text{ChCl}$ , in particular, is nowadays one of the widespread ammonium salt used for the synthesis of *DESs*. The latter (also known as vitamin  $\text{B}_4$ ) is produced on the scale of million metric tons per year (ca. 2 €/Kg) as an additive for chicken feed and has many other applications. Thanks to their low ecological footprint and attractive low price, *DESs* have now become of growing interest both at academic and industrial levels especially for their unusual solvent properties. It is worth noting that the concept of *DESs* is quite different from that of traditional ionic liquids (*ILs*) (*vide infra*) because the former are not entirely composed of ionic species, and can also be obtained from non-ionic species.<sup>[27]</sup>

Both nucleophilic additions and substitutions proved to be effective in such eutectic mixtures providing the expected adducts in good yields and competitively with protonolysis.<sup>[25]</sup> Adduct **12**, for example, could now be recovered with a yield of 40% upon adding an  $\text{Et}_2\text{O}$  solution of **11-Li** to acetone (6 equiv) in a  $\text{ChCl}$ -Gly (1:2) eutectic mixture at room temperature and under air (Scheme 6). Similarly, the addition reaction of a CPME solution of **11-Li** to benzophenone, run either in a  $\text{ChCl}$ -Gly (1:2) or in a  $\text{ChCl}$ -urea (1:2) *DES* mixture, gave the hydroxyalkylated compound **13** in both cases in 75% yield. Chlorodiphenylphosphine also successfully underwent nucleophilic substitution in  $\text{ChCl}$ -urea (1:2) leading the phosphenyl derivative **14** in 75% yield. Remarkably, once a commercial pentane solution of *t*-BuLi (1.9 equiv) was rapidly spread out over a mixture of **11-H** (1 equiv) in CPME and  $\text{ChCl}$ -Gly (1:2), at  $0^\circ\text{C}$ , under air, and under vigorous stirring, and quenched after 1 min reaction time with neat DMF (2 equiv), the formylated adduct **15** could be isolated in 90% yield (Scheme 6).



**Scheme 6.** Regioselective preparation of adducts **12**–**15** via *ortho*-lithiation/electrophilic interception of **11-H** in DES mixtures.

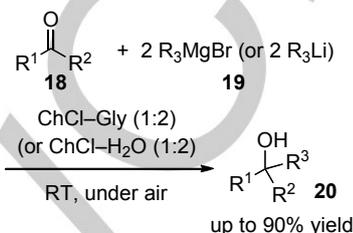
*o*-Tolyl-substituted tetrahydrofuran derivatives **16-H** have been recently found to undergo an unprecedented highly regioselective intramolecular C–O bond breaking reaction, triggered by the corresponding laterally lithiated intermediates **16-Li**, ending up with the formation of functionalised primary alcohols **17** showing incorporation in their skeletons of both a second equiv of base and of an electrophile (if any) at a tertiary carbon atom (Scheme 7).<sup>[28]</sup>



**Scheme 7.** Site-selected lateral lithiation/ring-opening of **16-H** and the regioselective preparation of functionalised alcohols **17** via intermediates **16-Li**.

This novel organic transformation can be also conveniently run directly in a glycerol-containing eutectic mixture, as a benign reaction medium, competitively with protonolysis. As a general reaction procedure, a commercial hydrocarbon solution of the organolithium (*s*-BuLi, *i*-PrLi, *t*-BuLi) was added by rapidly spreading it out over a mixture of **16-H** in CPME and ChCl–Gly (1:2), at 0 °C, under air, and under vigorous stirring, and quenched after 3 min reaction time with the electrophile to give alcohols **17** in yields up to >98%. The scope, limitation, and mechanistic aspects of this reaction, which pioneers “greener” alkylative THF ring-opening processes, have been discussed.<sup>[28]</sup> Interestingly, *s*-BuLi was found to promote a faster deprotonation compared to *t*-BuLi, and *ortho*-lithiation seriously competes with lateral lithiation only in the case of substrates possessing an ethyl group in an *ortho* position at one of the two aromatic rings.

The chemoselective nucleophilic addition of organolithium and Grignard reagents to ketones in ChCl-based eutectic mixtures was in depth investigated by Hevia, García-Álvarez and co-workers.<sup>[29]</sup> A range of the above reagents (**19**) could successfully be added, under air and at room temperature, to aromatic and aliphatic ketones **18** in both ChCl–Gly (1:2) and ChCl–H<sub>2</sub>O (1:2) mixtures, thereby affording the corresponding tertiary alcohols **20** in good yields (up to 90%) and competitively with protonolysis (Scheme 8).



**Scheme 8.** Chemoselective addition of Grignard and organolithium reagents to ketones in ChCl-based eutectic mixtures.

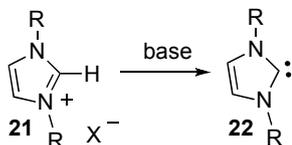
A comparison of the reactivity profiles of these organometallic reagents in DESs with those in pure water, suggest that a *kinetic activation* takes place in the former most probably due to the formation of more nucleophilic halide-rich magnesiate or lithiate species further to the reaction of the alkylating reagent with ChCl. Thus, ChCl may be playing a double role in these processes, that is as a component of both the DES mixtures and the new “ate” complexes. This conclusion was supported by X-ray crystallographic studies, multinuclear magnetic resonance investigations, and <sup>1</sup>H DOSY NMR experiments.

In a recent paper, Song showed as well that catalytic amounts of NBu<sub>4</sub>Cl in THF solutions of Grignard reagents enhanced the efficiency of addition reactions to carbonyl compounds producing tertiary alcohols in excellent yields, while minimizing the formation of enolization and reduction products.<sup>[30]</sup> The authors proposed that the presence of this ammonium salt should help to shift the Schlenk equilibrium of Grignard reagents in solution (Scheme 4) to the side of the dimeric species that would favour the addition reaction thanks to the 2:1 complex involved in the six-membered transition state.

## 2.2. Reactivity in ionic liquids

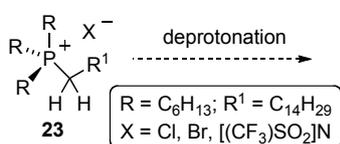
Generally, the term “ionic liquids” (*ILs*) stands for liquids composed of poorly coordinated ions with a melting point below 100 °C.<sup>[31]</sup> At least one ion has a delocalised charge, and one component is organic, which prevents the formation of a stable crystal lattice. They have recently attracted great interest as “greener” alternative to conventional organic solvents because of their thermal stability, non-flammability, easy of recycling, low vapour pressure, and catalytic properties.<sup>[32]</sup> Their use as solvents for reactions involving organometallic compounds of *s*-block elements, however, is still in its infancy. One of the most extensively studied class of *ILs* is based on imidazolium cations with an appropriate counter anion (*ImILs*), which are known to support many organic transformations.<sup>[32]</sup> Because of an acidic

hydrogen substituent at the C-2 position, *Im*ILs (**21**) have been shown to react under basic conditions to produce *N*-heterocyclic carbenes (NHCs) (**22**), which are neutral, highly reactive, six-electron species possessing a dicoordinate carbon atom with two nonbonding electrons, and are responsible of many side reactions (Scheme 9).



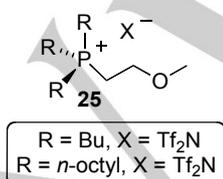
**Scheme 9.** Formation of a NHC (**22**) species from an imidazolium cation (**21**).

This problem was overcome by Clyburne and co-workers who showed that dried phosphonium *IL*s **23** are inert towards reactions with strong bases and are not reduced even by potassium metal, thereby representing the first suitable solvents for Grignard chemistry.<sup>[33]</sup> Commercially available THF solutions of PhMgBr, once dissolved in **23** (ratio THF : **23** = 1 : 3), proved to cleanly promote carbonyl additions, benzyne reactions, halogenation, hydroxy(alkylation)arylation, and coupling reactions. Most importantly, competitive deprotonation of **23** to produce the phosphorane **24** did not take place (Scheme 10). The inertness of phosphonium cations towards Grignard solutions appears to have primarily a kinetic basis and to be anion dependent: small bases are more prone to deprotonate **23**, whereas large bases are more reluctant. Several other causes, however, seem also to contribute to such an inertness; e.g., the bulkiness and the flexibility nature of the cation as well as its electrochemical robustness compared to unsaturated ions.<sup>[34]</sup>



**Scheme 10.** Possible formation of phosphorane **24** by deprotonation of **23**.

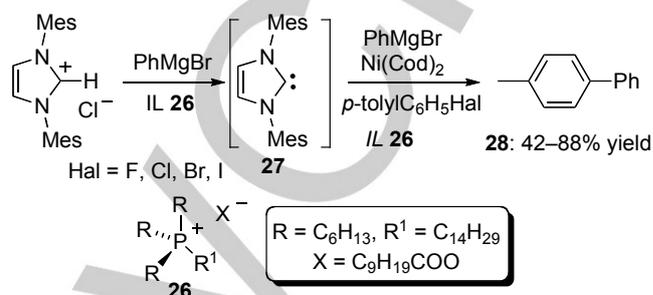
The introduction of an ether oxygen on the side arm of a phosphonium salt (**25**) contributes to stabilising the organomagnesium reagent, thereby improving the capability of the corresponding *IL* to act as a solvent even for aliphatic Grignard reagent-mediated reactions (Figure 3).<sup>[35]</sup>



**Figure 3.** Phosphonium *IL*s with an ether functionality.

Walsby and co-workers also demonstrated that while in molecular solvents Grignard reagents react according to nucleophilic pathways, *IL*s are ideal reaction media to promote electron-transfer processes.<sup>[36]</sup> The Kumada-Corriu reaction,

which involves the coupling of Grignard reagents with aryl halides mediated by transition metal catalysts (typically nickel or palladium), has been successfully carried out between PhMgBr and aryl halides in the phosphonium *IL* **26** in the presence of a Ni(0) complex of NHC **27** to afford biaryl derivatives **28** with yields up to 88%, thereby supporting the *in situ* generation of carbene species (Scheme 11). Remarkably, such a reaction even facilitates the activation of C–F bonds.



**Scheme 11.** Kumada–Corriu cross-coupling reaction in the phosphonium *IL* **27**.

Apart from phosphonium *IL*s, also some imidazolium-based *IL*s can withstand the strong basicity of the Grignard reagents. These include *Im*ILs with a phenyl substituent<sup>[37]</sup> and with an isopropyl group<sup>[38]</sup> at the vulnerable C–2 position. Both *IL*s **29** and **30** (Figure 4) have been successfully employed as suitable solvents in reactions involving the addition of aliphatic and aromatic Grignard reagents to aldehydes, ketones and esters affording the expected hydroxyalkylated adducts in good yields (68–83%). These *Im*ILs can be recycled and reused several times without appreciable loss of the same *IL*. All attempts, however, to generate Grignard reagents in *IL*s failed. In 2006, Chan and co-workers reported the first example of an organomagnesium species generated directly in the pyridinium-based *IL* **31** having a tetrafluoroborate as a counter ion (Figure 4).<sup>[39]</sup> The reactivity pattern showed towards carbonyl compounds for reactions run in **31**, however, was different from that exhibited by Grignard reagents in conventional organic solvents, the yields in the final adducts also being critically dependent on the molar ratio of the reagents and the presence of additives in the reaction mixture. New alkyipyridinium (**32**) and tetralkylphosphonium (**33**) *IL*s, possessing an ether functionality to provide stabilisation to the Grignard reagent, have been prepared by Scammells and co-workers and evaluated as solvents for Grignard reactions.<sup>[40]</sup> Interestingly, different outcomes have been observed according to the presence or not of an ethereal co-solvent. When the addition reaction to carbonyl compounds was run in *IL*s **32**, the expected adducts occurred only in the presence of Et<sub>2</sub>O, whereas in the absence of this ether an unusual reduction of aldehydes to the corresponding primary alcohols was favoured. On the other hand, aldehydes cleanly reacted with Grignard reagents in *IL*s **33** affording the corresponding addition products only in the presence of Et<sub>2</sub>O as a co-solvent.

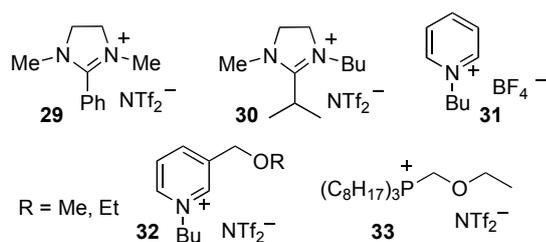


Figure 4. ILs employed as alternative reaction media for Grignard reagents.

To the best of our knowledge, analogous reactions of organolithium compounds run in *ILs* always produced decomposition and unidentified products. Because of the broad use of organolithium and Grignard reagents in the pharmaceutical and fine chemical industry, a judicious choice of the reaction solvent is crucial from both safety and environmentally standpoints. In this context, 2-MeTHF (derived from a renewal source) and CPME (directly obtainable from cyclopentene) are emerging “greener” alternatives for organometallic reactions to the common Et<sub>2</sub>O and THF, and have also proved to be more effective in improving product yield and in suppressing side reactions.<sup>[41]</sup> The challenge will be the use of commercially prepared Grignard and organolithium solutions directly in the above solvents.

### 3. Organometallic compounds of d-block elements

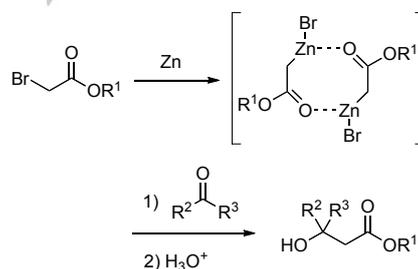
As has been assessed in the Introduction, OC has become a cornerstone of modern organic synthesis, and nowadays hardly any total synthesis endeavour can be envisioned without a key step involving the use of polarised organometallic compounds containing d-block elements, that is the *transition metals*.<sup>[42]</sup> These fundamental reagents (most commonly organozinc and organocopper compounds) are able to deliver carbon residues (M–C bonds) from zinc<sup>[43]</sup> or copper<sup>[44]</sup> to carbon halides or pseudohalides (C–X), to form new C–C bonds. However, the chemoselectivity of these processes can be seriously compromised by: i) the formation of undesired products, ii) the use of low temperatures (ranging from 0 to –78 °C), and iii) the employment of dry and hazardous ethereal solvents, and inert atmosphere protocols (to avoid fast degradation of the polar reagents). All these experimental restrictions hinder the synthetic application of these polarised species under environmentally-friendly reactions conditions (*i.e.* at room temperature and in the absence of protecting atmosphere),<sup>[1,3,45]</sup> and their use in the presence of unconventional solvents [*e.g.*, water, *ILs*, supercritical CO<sub>2</sub> (scCO<sub>2</sub>) or perfluorinated solvents] as reaction media.<sup>[46]</sup> Despite all these drawbacks, during the last decades the chemistry of polarised organometallic compounds containing d-block elements has crossed the frontiers between their application in modern synthetic organic chemistry and the growing area related to the employment of unconventional solvents. Since the synthesis of the first polarised organozinc compound (ZnEt<sub>2</sub>) by Wanklyn in 1848,<sup>[47]</sup>

it is well-known that these basic compounds are able to react with several unconventional solvents (like water or scCO<sub>2</sub>).

This Section covers the progress made in the application of the aforementioned unconventional solvents as reaction media in a variety of organozinc-, organocopper-, and organogold-mediated organic reactions. In particular, the following reactions will be surveyed: i) Reformatsky- and Barbier-type reactions, ii) addition of organozinc derivatives to  $\alpha,\beta$ -unsaturated carbonylic compounds, iii) cross-coupling reactions of the *in situ* generated organozinc reagents and organic halides (Negishi coupling), iv) polymerization reactions, v) organogold reactions, and vi) iridium-promoted C–H bond activation reactions.

#### 3.1 Reformatsky reaction in unconventional solvents

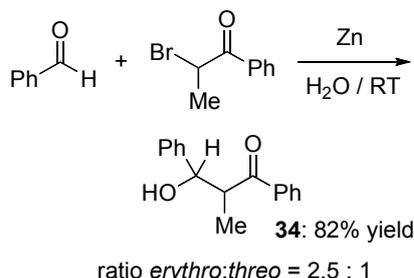
The Reformatsky reaction,<sup>[48]</sup> which involves the treatment of a haloester with a carbonyl compound (ketone or aldehydes) in the presence of Zn (Scheme 12), was the first example of addition of polarised organometallic reagents containing d-block elements to carbonyl compounds. Since its discovery in the 19<sup>th</sup> century, it was believed that this Zn-mediated addition reaction could only take place under protective atmosphere (using Schlenk techniques) and employing dry organic solvents.<sup>[49]</sup> However, several examples were then reported in the literature showing the reactions between carbonyl compounds and organic halides, mediated by reactive d-block metals (Zn, Cu), in both wet solvents and pure water.<sup>[50]</sup>



Scheme 12. Reformatsky reaction between haloesters and carbonyl compounds in the presence of Zn.

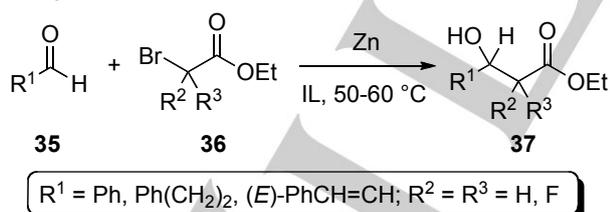
In 1990, Chan, Li and co-workers lighted the way by studying the direct Reformatsky-type conversion between benzaldehyde and  $\alpha$ -bromopropiophenone in pure water as the solvent, mediated by metallic zinc (Scheme 13).<sup>[51]</sup> Both the yield (82%) and the ratio of the two diastereomeric products (**34**, 2.5:1 *erythro:threo*) gathered in this aqueous reaction are comparable to those obtained with preformed organometallic reagents under anhydrous conditions. Latter, Bieber and co-workers demonstrated that this pioneering idea could be extended to a wide variety of carbonyl substrates (including aromatic and aliphatic aldehydes and ketones) and to different haloesters.<sup>[52]</sup> Again, yields achieved under these aqueous conditions proved to be comparable to the ones obtained in anhydrous organic solvents. Chan and co-workers proposed that this reaction may follow a radical mechanism (Single Electron Transfer), as no

formation of the desired product was observed in the presence of galvinoxyl or hydroquinone (radical scavengers).<sup>[51b]</sup>



**Scheme 13.** Reformatsky-type reaction between benzaldehyde and  $\alpha$ -bromopropiophenone in water.

Not only water, but also other unconventional solvents like *ILs* have been employed as reaction media for Reformatsky-type reactions. As previously discussed in Section 2.2 of this Microreview, *ILs* have received much attention as a new class of unconventional solvents during the last decades.<sup>[32]</sup> To the best of our knowledge, the first Reformatsky reaction in a variety of *ILs* was reported by Kitazume and co-workers (Scheme 14).<sup>[53]</sup> The authors studied the Reformatsky reaction: i) between aromatic, aliphatic or alkenyl aldehydes (**35**) with different haloesters (**36**), ii) mediated by metallic Zn, and iii) in the *ILs* [EtDBU][OTf] (8-ethy-1,8-diazabicyclo[5,4,0]-7-undecenium trifluoromethanesulfonate), [BMIM][BF<sub>4</sub>] (1-Butyl-3-methylimidazolium tetrafluoroborate), and [BMIM][PF<sub>6</sub>] (1-Butyl-3-methylimidazolium hexafluorophosphate). At room temperature, only a moderate yield (52%) of the desired compound **37** was achieved in the *IL* [EtDBU][OTf]. However, upon heating the reaction to 50 °C, almost quantitative conversion (93%) in **37** was reached. As has been pointed out in Section 2.2, one of the major advantages associated with the use of *ILs* as solvents is the possibility of reusing the *IL* by a simple extraction of the desired organic product (**37**) with organic solvents.<sup>[32,46]</sup> In this way, the *IL* could be recycled up to three consecutive cycles without any loss of activity or selectivity.



**Scheme 14.** Reformatsky reaction in *ILs*.

### 3.2 Barbier reaction in unconventional solvents

The Barbier reaction,<sup>[54]</sup> which involves the reaction of organic halides and carbonyl compounds in the presence of magnesium, aluminium, zinc, indium, tin or its salts is one of the most important methods for creating C–C bonds and has widespread synthetic applications in organic chemistry.<sup>[55]</sup> The reaction proceeds via the nucleophilic attack of the *in situ* generated

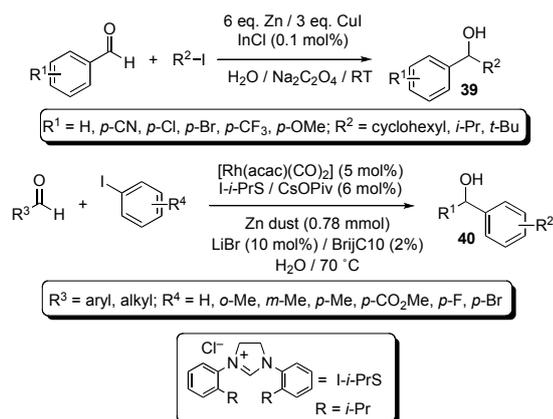
organometallic compound on the carbonyl electrophile (generally an aldehyde). Since its discovery, there has been considerable attention towards the development of this Zn-mediated reaction in water.<sup>[56]</sup> In fact, the allylation of aldehydes and ketones under the Barbier conditions usually occurs faster and gives rise to higher yields when water is used as the (co)solvent.<sup>[57]</sup> In this sense, Li and Chan reported one of the first and innovative allylation reactions of carbonyl compounds promoted by Zn in water (Scheme 15).<sup>[58]</sup> Again, the presence of water was critical to the success of the coupling step for the formation of **38**. Thus, when the reaction was performed in dry conventional ethereal solvents (e.g., Et<sub>2</sub>O or THF), poor formation of compound **38** was encountered.



**Scheme 15.** Zn-mediated Barbier reaction in water.

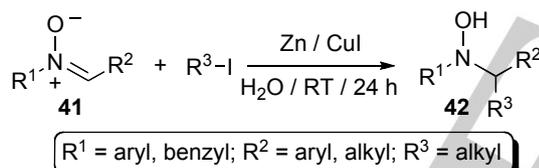
Nowadays, a plethora of Zn-mediated allylation of different electrophiles (e.g., aldehydes, ketones, acetals or dioxolanes) in aqueous conditions is known in the literature, allowing the direct synthesis of homoallylic alcohols under environmentally-friendly reaction conditions.<sup>[59]</sup> Recently, the spectrum of unconventional solvents available to accomplish this Zn-mediated allylation reaction under green conditions has been enlarged by Leeke and co-workers. These authors reported the employment of subcritical CO<sub>2</sub>/H<sub>2</sub>O (30 °C/80 bar) as a renewable solvent mixture to increase the desired allylation reaction with a variety of aryl aldehydes.<sup>[60,61]</sup>

Not only Zn-mediated allylations,<sup>[59]</sup> but also propargylations<sup>[62]</sup> and benzylations<sup>[63]</sup> of different carbonyl compounds can be conveniently performed in the presence of water. In recent years, Li and co-workers have expanded the scope of this Barbier-type reaction to the more challenging carbonyl alkylations<sup>[64]</sup> and arylations<sup>[65]</sup> with non-activated halides in water. In both the examples cited, the desired aromatic aldehyde suffers the corresponding alkylation or arylation in water, which is mediated by stoichiometric amounts of Zn dust and catalysed by InCl or [Rh(acac)(CO)<sub>2</sub>] (acac = acetylacetonate), respectively (Scheme 16). These processes allowed the mild and the straightforward synthesis of benzyl alcohols (**39**) and aryl methanols (**40**), thereby unlocking one of the last challenges in the field of Barbier-type reactions in water.



**Scheme 16.** Barbier-type alkylation and arylation of aldehydes in water.

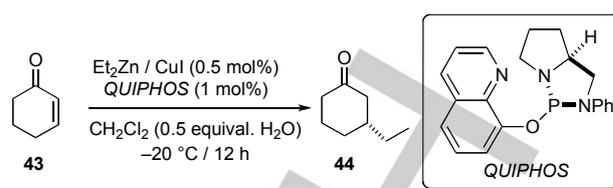
Amines could also be conveniently prepared by direct addition of zinc organometallic reagents to imines in the presence of water. Savoia, Umani-Ronchi and co-workers first developed the enantioselective synthesis of homoallylic amines by addition of *in situ* generated (allyl)ZnBr reagents to imines in a mixture of THF/H<sub>2</sub>O.<sup>[66]</sup> More recently, Naito and co-workers reported the Zn-mediated addition of alkyl iodides to imines in the absence of VOCs by using a saturated NH<sub>4</sub>Cl aqueous solution as the solvent.<sup>[67]</sup> Nitrones **41** can also suffer a Barbier-type alkylation in pure water as the solvent and at room temperature, yielding the corresponding hydroxylamines **42** (Scheme 17).<sup>[68]</sup>



**Scheme 17.** Synthesis of hydroxylamines *via* Barbier-type alkylation of nitrones in water.

### 3.3 Conjugate addition of organozinc derivatives to $\alpha,\beta$ -unsaturated carbonylic compounds in water

The conjugate 1,4-addition of organometallic compounds to electron-deficient olefins represents one of the most powerful tools currently exploited to create new C–C bonds. Among the various methods available, the most commonly employed strategies involve the use of organometallic species such as Grignards reagents (RMgX) or organolithium (RLi) compounds. However, the use of these highly reactive organometallic derivatives can lead to undesired side reactions (e.g., Wurtz coupling, reduction of the carbonyl compounds, hydrolysis, competitive 1,2-addition, etc.).<sup>[69]</sup> Thus, dialkylzinc reagents have dominated the field of copper-mediated enantioselective conjugate addition since their first application in the mid-1990s.<sup>[70]</sup> From that moment on, several reports have appeared in recent years demonstrating that the addition of stoichiometric or sub-stoichiometric quantities of water increases the rate and/or the enantioselectivity of this organic transformation.<sup>[71]</sup>



**Scheme 18.** Cu(I)-catalysed conjugate 1,4-addition of Et<sub>2</sub>Zn to cyclohexenone (**43**) accelerated in the presence of water.

Delapierre and co-workers reported the dramatic beneficial effect of addition of water (0.5 equiv.) in the asymmetric addition of diethylzinc to cyclohexenone (**43**) catalysed by CuI in the presence of chiral ligands (Scheme 18).<sup>[72]</sup> Thus, when the reaction was performed in dry CH<sub>2</sub>Cl<sub>2</sub>, only 55% yield of the desired cyclohexanone (**44**) was achieved in 45% ee. However, formation of the desired carbonyl compound in higher yield (76%) and enantiomeric excess (ee 61%) was observed upon the addition of sub-stoichiometric amounts of water. The authors proposed that the *in situ* formation of Zn(OH)<sub>2</sub> (which is a stronger and more effective Lewis acid), activates the carbonyl moiety. This suggestion was confirmed as the direct addition of Zn(OH)<sub>2</sub> to the reaction provided analogous results to those observed with water.<sup>[73]</sup> Similarly, the addition of sub-stoichiometric amounts of water (0.3–0.33 equiv.) to lithium dimethylcuprate (LiCuMe<sub>2</sub>), generated a more reactive and stereoselective reagent for the conjugate addition to linear  $\alpha,\beta$ -enones.<sup>[74]</sup> More recently, Lipshutz and co-workers reported the conjugate addition of *in situ* generated organocopper reagents to enones<sup>[75]</sup> in water and at room temperature by using small amounts of commercially available amphiphiles (TPGS-750-M, polyoxyethanyl- $\alpha$ -tocopheryl succinate) that are able to form nanomicelles in water.



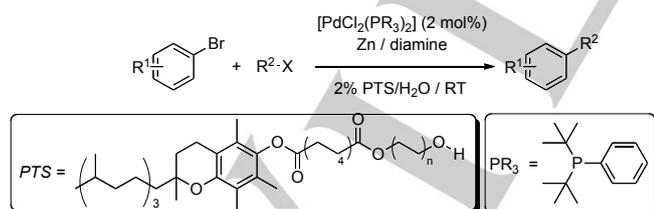
**Scheme 19.** Synthesis of lupidine analogue **45** mediated by Zn/CuI under sonication conditions in the presence of water.

Conjugated 1,4-addition of alkyl halides (R-X) to  $\alpha,\beta$ -unsaturated aldehydes, ketones, esters, amides,<sup>[76]</sup> or nitriles<sup>[77]</sup> can be mediated in EtOH–H<sub>2</sub>O or THF–H<sub>2</sub>O mixtures by the combination of Zn and Cu, under sonication conditions. It is worth noting that this methodology has been fruitfully applied to the synthesis of: i) a variety of vitamin D<sub>3</sub> derivatives,<sup>[78]</sup> ii) dioxolanes,<sup>[78d]</sup> iii) oxazolidinones,<sup>[78h]</sup> and iv) sinfungin analogues.<sup>[79]</sup> Finally, the intramolecular version of this 1,4-addition reaction mediated by Zn/CuI allowed the straightforward synthesis of the lupinine analogues **45** (Scheme 19).<sup>[80,81]</sup> Thus, the 1,4-conjugated addition of alkyl halides to  $\alpha,\beta$ -unsaturated carbonyl compounds mediated by Zn–Cu mixtures in aqueous media proved to be the key step in the total synthesis of a diversity of natural products.

### 3.4 Cross-coupling of *in-situ* generated organozinc reagents with organic halides (Negishi coupling) in unconventional solvents.

Metal-catalysed cross-coupling reactions between an organic electrophile (typically an organic halide) and an organic nucleophile have developed into a standard component of the armamentarium synthetic chemist's toolbox for the formation of C–C and C–heteroatom bonds.<sup>[82]</sup> Palladium-catalysed reactions, which can be generally carried out under milder conditions and with a wider range of substrates than reactions promoted by other metals, clearly dominate the field. The organic halide can be a  $sp$ -,  $sp^2$ -, or  $sp^3$ -hybridised carbon with any halogen or pseudo-halogen leaving group. Different organometallic nucleophiles (e.g., organoboron, organotin, organozinc, organomagnesium) and organic nucleophilic reagents (such as amines, alkenes or alkynes) are routinely used in different cross-coupling reactions. In this Section, the attention will be mainly focused on the Pd-catalysed Negishi coupling (with reference to the cross-coupling reactions of polarised organozinc reagents with organic halides),<sup>[83]</sup> in different unconventional solvents, like water, *ILs* and perfluorinated solvents.

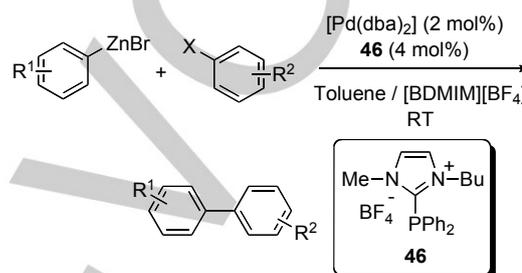
Lipshutz and co-workers have almost dominated the field of aqueous Negishi-type cross-coupling reactions by describing a new technology that allows the Pd-catalysed Zn-mediated cross-couplings to be conducted in water and at room temperature, without the need to preform the corresponding organozinc reagent ( $RZnX$ ).<sup>[84,85]</sup> Lipshutz's approach uses homogeneous micellar catalysis within catalytic nanoreactors formed spontaneously upon dissolution in water of different surfactants (PTS, TPGS, Brij 30, Solutol, SPGS). The scope of this process has been studied in the Pd-catalysed coupling between: i) alkyl halides and aryl or heteroaryl halides (Scheme 20),<sup>[84a,f-1]</sup> ii) alkyl or benzylic halides and alkenyl halides,<sup>[84b,e,h]</sup> and iii) benzyl halides and aryl or heteroaryl halides.<sup>[84c,d]</sup> More recently, Lipshutz and co-workers reported the reduction of alkyl halides,<sup>[86]</sup> and nitroaromatics<sup>[87]</sup> in water and at room temperature by using Zn dust in the presence of nanomicelles composed of the aforementioned surfactants.



**Scheme 20.** Representative example of Negishi-type coupling reaction in water at room temperature employing the Lipshutz conditions.

Pd-catalysed Negishi-type reaction has also been reported in *ILs*. In this regard, Knochel and co-workers described at the beginning of this millennium the cross-coupling reaction between preformed aryl- or benzylzinc halides ( $RZnX$ ) and various aryl iodides in the *IL*  $[BDMIM][BF_4]$  ( $BDMIM$  = 1-butyl-2,3-dimethylimidazolium), and using as catalytic system the mixture

formed by  $[Pd(dba)_2]$  an ionic phosphine **46** (Scheme 21).<sup>[88a]</sup> In most cases, the reaction proceeded at room temperature within minutes leading to the desired product in almost quantitative yields. The work up of this reaction is remarkably simple, as the *IL* phase containing the palladium catalyst can be separated from the organic product simply by extraction with toluene. Attempts to reuse the palladium catalyst showed that after the third cycle, a significant decrease in the yield was observed. The same authors enlarged the scope of unconventional solvents that could be used in the Negishi reaction by describing the Pd-catalysed cross-coupling of organozinc bromides with aryl iodides in perfluorinated solvents.<sup>[88b]</sup>

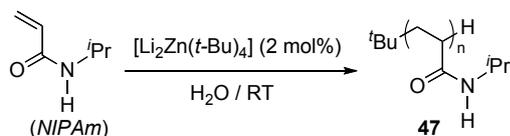


**Scheme 21.** Pd-catalysed Negishi-type coupling between preformed organozinc reagents and aryl halides in the ionic liquid  $[BDMIM][BF_4]$ .

### 3.5 Application of lithium organozincates for chemoselective anionic polymerization

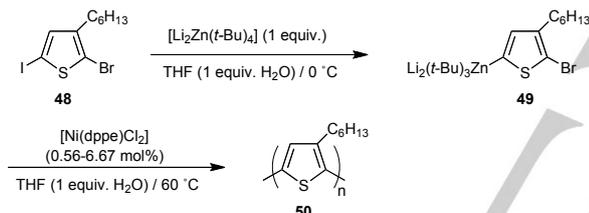
Highly coordinated dianion-type zincates ( $Li_2ZnR_4$ ) were reported in the mid-1990s by Uchiyama and co-workers as a new type of zincate complexes, and added a new dimension to organozincate reagents because they were able to promote bromine-zinc exchange and carbozincation reactions.<sup>[89]</sup> In this regard, the dilithium tetra-*tert*-butylzincate  $[Li_2Zn(t-Bu)_4]$  turned out to be a highly crowded and bulky zincate with an excellent anionic polymerization ability, even in the presence of acidic protons.<sup>[90]</sup> Uchiyama and co-workers studied the anionic polymerization of *N*-isopropylacrylamide (*NIPAm*) using  $Li_2Zn(t-Bu)_4$  as initiator in both organic solvents and water (Scheme 22).<sup>[91]</sup> Surprisingly, an interesting solvent effect was found in this polymerization reaction: with THF as the reaction medium, only 8% of the desired polymeric material **47** was obtained after 24 h, whereas the polymer **47** could be isolated in high yields (92–76%) after 3 hours only when protic solvents (like  $H_2O$  or MeOH) were alternatively used. Upon monitoring the time/yields profile of this polymerization reaction in water, the authors noticed that polymer **47** could be obtained in 92% yield after 15 minutes. The nature of the organometallic compound proved to be crucial as no polymerization reaction took place when  $Li_2Zn(t-Bu)_4$  was replaced by  $Li^+t-Bu^-$ ,  $ZnCl_2$ ,  $LiCl$  or  $LiOH$ . This Zn-mediated anionic polymerization in water could also be extended to other acryl acid derivatives, such as *N,N*-dimethylacrylamide (DMA, 74% yield), acrylamide (AM, 84% yield) and 2-hydroxyethylmethacrylate (HEMA, 92% yield). One of the main drawbacks of this  $Li_2Zn(t-Bu)_4$ -mediated polymerization in aqueous media is the impossibility of induce the polymerization of styrene (one of the most important synthetic polymers). In this

case, deprotonation of the solvent took place before polymerization.<sup>[92]</sup>



**Scheme 22.**  $\text{Li}_2\text{Zn}(\text{t-Bu})_4$  catalysed anionic polymerization of *N*-isopropylacrylamide (*NIPAm*) in water.

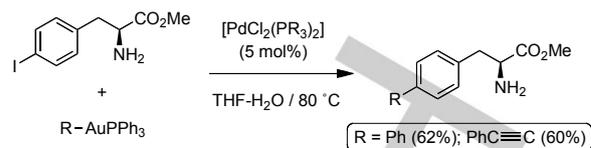
Recently, Higashihara and co-workers reported another possibility for application of zincate  $\text{Li}_2\text{Zn}(\text{t-Bu})_4$  in polymerization reactions, that is through an exchange-cross-coupling process.<sup>[93]</sup> Thus, when 2-bromo-3-hexyl-5-iodothiophene **48** was treated with  $\text{Li}_2\text{Zn}(\text{t-Bu})_4$ , the iodine-zinc exchange reaction took place selectively. Upon heating to 60 °C the resultant zincate **49** with the nickel catalyst  $[\text{Ni}(\text{dppe})\text{Cl}_2]$  ( $\text{dppe}$  = 1,2-Bis(diphenylphosphino)ethane), polymerization proceeded in a controlled manner affording poly(3-*n*-hexylthiophene) **50** in high yield (80-85%) and low polydispersities (PDIs < 1.2). As analogously observed in the anionic polymerization of *N*-isopropylacrylamide (*NIPAm*) in water,<sup>[91]</sup> the high-molecular-weight-polymer **50** could be obtained in a THF solution containing a small amount of water (Scheme 23).



**Scheme 23.** Halogen-exchange reactions and catalyst-transfer polycondensation for the synthesis of polymer **50** using  $\text{Li}_2\text{Zn}(\text{t-Bu})_4$  in the presence of water.

### 3.6 Organogold(I) compounds in Palladium-catalysed cross-coupling reactions in aqueous media

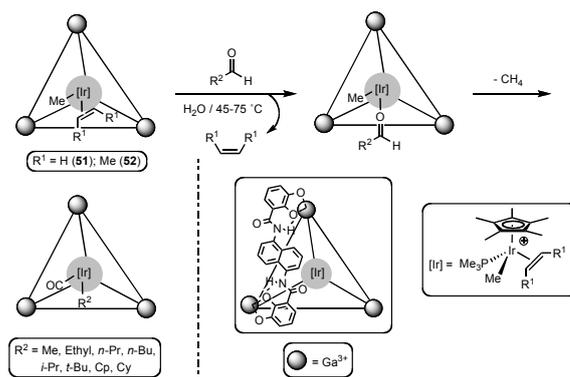
As previously discussed in Section 3.4, Pd-catalysed cross-coupling reactions are usually run with polarised organometallic nucleophiles (e.g., organoboron, organotin, organozinc, organomagnesium) in VOCs solvents. However, Sarandeses and co-workers have recently expanded the scope of this transformation first employing organogold(I)-phosphane derivatives ( $\text{RAuPPH}_3$ ) as organometallic nucleophiles in water. Under these conditions, reactions between isolated aryl-, alkenyl-, or alkynylgold(I)-phosphanes and aryl halides or triflates were shown to proceed at room temperature (or at 80 °C) in water/THF mixtures.<sup>[94]</sup> These Pd-catalysed reactions delivered the corresponding coupling products in good yield and with high chemoselectivity being compatible with free amino or hydroxyl groups present in the electrophile. As a proof of concept, this methodology was then successfully applied also to the preparation of substituted phenylalanine esters under protic conditions (Scheme 24).



**Scheme 24.** Synthesis of 4-substituted phenylalanines in a mixture THF-water.

### 3.7 Iridium-promoted C–H bond activation in water

Encapsulation of a variety of organometallic complexes into the internal cavity of hydrophilic supramolecular structures constitutes an innovative way to solubilise organometallic derivatives in aqueous media.<sup>[95]</sup> In this context, Raymond and co-workers have incorporated the cationic iridium complexes  $[(\text{Cp}^*)(\text{PMe}_3)\text{Ir}(\text{Me})(\eta^2\text{-olefin})[\text{OTf})]$  ( $\eta^2\text{-olefin}$  = ethylene or *cis*-2-butene) into a supramolecular  $[\text{Ga}_4\text{L}_6]$  tetrahedral assembly ( $\text{L}$  = 1,5-bis(2,3-dihydroxybenzoylamino)naphthalene) (Scheme 25). These species formed the host-guest complexes **51** and **52**, stabilised by hydrophobic effects as well as by  $\pi$ - $\pi$  interactions between the coordinated olefin and the  $\pi$ -basic naphthalene walls of the host.<sup>[96]</sup> The resulting water soluble host-guest systems (**51,52**) were then tested in the C–H activation of aldehydes in aqueous media. In order to generate the active iridium species, decoordination of the olefin was preliminary required. The simple heating of the host-guest complexes (45 °C for **51** and 75 °C for **52**) facilitated olefin dissociation, thereby allowing the C–H bond activation of the desired aldehyde. Interestingly, evidence for both size and shape selectivity was observed. Small aldehydes (e.g., acetaldehyde) are readily activated, whereas large aldehydes (e.g., benzaldehyde) are too large to fit inside the cavity. Also, the shape of the aldehyde proved to influence the reactivity of the encapsulated host-guest complex. For example, the host-guest complex reacted with isobutyraldehyde with a lower diastereoselectivity than with butyraldehyde. This experimental evidence was attributed to the more spherical shape of the isobutyraldehyde complex when compared to the butyraldehyde one.



**Scheme 25.** C–H activation of aldehydes in aqueous media promoted by the water-soluble host-guest complexes **51,52**.

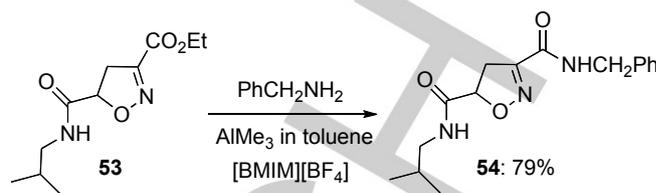
## 4. Organometallic compounds of p-block elements

Amongst the vast family of organometallic compounds, and within the subgroup of the p-block metals (elements whose valence electrons are in the p orbital), organoaluminium<sup>[97]</sup> and organotin<sup>[98]</sup> members play a pivotal role in organic synthesis, finding widespread applications in a myriad of C–C bond-forming processes. Although most of these reactivity studies have been performed using conventional VOCs, the potential of using these commodity organometallic reagents also in unconventional media (including *ILs*, *DESSs*, *scCO*<sub>2</sub>, and recently also neat water) has already been hinted at by several intriguing studies that will be discussed in this Section. In addition to these two important families of p-block metal reagents, the chemistry and applications of organoindium compounds are receiving increasing attention from the synthetic community, and are thus being rapidly developed. Bearers of an exceptional functional group tolerance and distinctive mild reactivity profiles, these compounds can render unique chemoselectivities in several C–C bond forming reactions which are difficult to achieve using more polar reagents such as organolithium or organomagnesium reagents. Intriguingly and contrasting with the typical extreme moisture sensitivity of these polar organometallics, organoindium reagents can be utilised in aqueous media which allows the functionalisation of water-soluble substrates such as carbohydrates, as well as the development of greener synthetic methodologies. This unique behaviour was first reported by Li and Chan in 1991 through a seminal study assessing the allylation reactions of aldehydes and ketones under Barbier conditions in water,<sup>[99]</sup> and nowadays it constitutes a signature attraction of organoindium chemistry. Of note, the chemistry of these reagents and their applications for the functionalisation of organic molecules (some of them employing aqueous media) have been recently summarised in a comprehensive review by Loh and co-workers.<sup>[100]</sup> Thus, they will be not covered in this overview.

### 4.1 Applications of Group 13 organometallic reagents

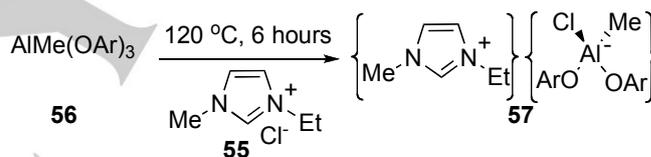
Organoaluminium reagents have received considerable attention in recent years not only due to their high chemoselective reactivity and exceptional functional group tolerance, but also because of their relative cheapness, ready availability and comparative low toxicity. The polarity of their Al–C bonds makes these commodity reagents extremely air and moisture sensitive, and they usually have to be manipulated under strict inert-atmosphere techniques. Therefore, although at present most of their applications require the use of dry organic solvents, some promising studies have already glimpsed the potential of applying these compounds in *ILs* as an alternative reaction media. In 2006, Taddei and co-workers have reported the multistep synthesis of isoxazolines using the *IL* [BMIM][BF<sub>4</sub>], where one of the key synthetic steps involves the transformation of an ester into an amide via an aluminium amide. This species is generated *in situ* by adding a solution of AlMe<sub>3</sub> in toluene to a

solution of the ester **53** and benzylamine furnishing **54** in 79% yield (Scheme 26).<sup>[101]</sup>



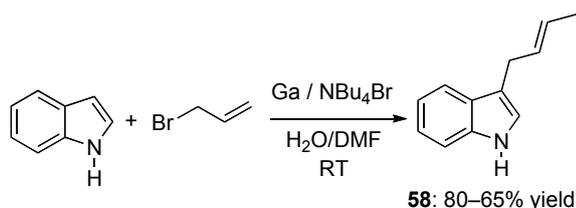
**Scheme 26.** Ester amidination of isoxazoline **53** with AlMe<sub>3</sub> in the *IL* [BMIM][BF<sub>4</sub>].

More recently, Chen and Liu have shown that aluminium alkyl and aryloxy compounds, used widely in polymerization processes, can effectively catalyse the conversion of glucose to HMF (5-hydroxymethylfurfural) using the *IL* 1-ethyl-3-methylimidazolium chloride, [EMIM]Cl **55**.<sup>[102]</sup> Trying to shed some light on the constitution of the active Al species involved in this transformation, alkylaryloxy aluminium MeAl(OAr)<sub>2</sub> **56** (OAr = 2,6-di-tert-butyl-4-methylphenoxide) was mixed with **55**, under the same glucose conversion conditions, and this enabled the isolation and subsequent structural elucidation of the new mixed imidazolium aluminate {ENIM}<sup>+</sup>{Me(Cl)Al(OAr)<sub>2</sub>}<sup>-</sup> **57**, where the chloride is now attached to Al (Scheme 27).



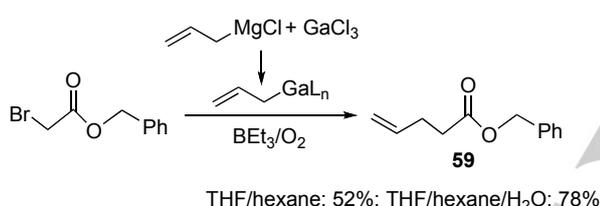
**Scheme 27.** Formation of imidazolium aluminate **57** by combining aluminium complex **56** with *IL* **55**.

As already alluded to, indium can mediate Barbier-type reactions in water.<sup>[99,100]</sup> Interestingly, metallic gallium can also be used to promote the allylation of aldehydes and ketones with allyl bromide in water, affording the relevant homoallyl alcohols in high yields.<sup>[103]</sup> Similarly, the coupling of indoles and pyrroles with allyl halides can be accomplished in a mixture of water and DMF in the presence of Ga metal using NBu<sub>4</sub>Br as an additive.<sup>[104]</sup> This method in granting access to C3-allylated indole species **58** represents a main-group metal-mediated alternative to other approaches employing Pd catalysts. Interestingly, the choice of solvent is crucial for the success of this transformation as a mixture of products results on using acetonitrile or THF. The effect of NBu<sub>4</sub>Br is also remarkable; in fact, the employment of other metal bromides such as MgBr<sub>2</sub> or KBr inhibits the coupling process (Scheme 28).



**Scheme 28.** Gallium-mediated allylation of indoles using a H<sub>2</sub>O/DMF mixture.

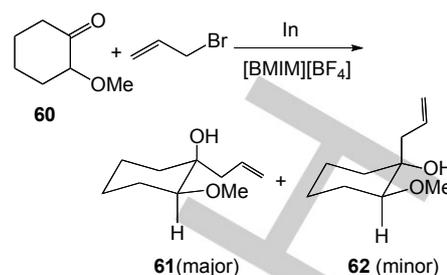
Oshima has shown that allylgallium reagents, generated *in situ* via salt metathesis of GaCl<sub>3</sub> and allylmagnesium bromide, promote radical allylation of  $\alpha$ -iodo and  $\alpha$ -bromo carbonyl compounds in the presence of BEt<sub>3</sub> and under air, using a mixture of THF/hexane and water (Scheme 29).<sup>[105]</sup> Interestingly, when assessing solvent effects, it was found that without using water as a co-solvent the yields in the formation of **59** were lowered significantly. Although the exact nature of this favourable solvent effect is unclear, the authors suggest the possible involvement of allylgallium hydroxide intermediates, which may be more reactive towards the radical allylation process.



**Scheme 29.** Triethylborane-induced radical allylation of  $\alpha$ -halo carbonyl compounds with allylgallium reagent in aqueous media.

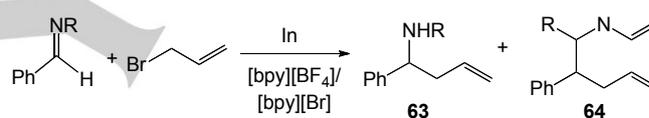
Significantly, indium-mediated allylation reactions cannot only be accomplished using water as the solvent.<sup>[100]</sup> Alternative reaction media such as *ILs*<sup>[106]</sup> and *scCO*<sub>2</sub><sup>[107]</sup> can be employed too. Indeed, Gordon and Ritchey have reported the use of indium metal and allyl bromide for the allylation of a wide range of aldehydes and ketones using *IL* [BMIM][BF<sub>4</sub>]. These reactions can be carried out at room temperature using stoichiometric amounts of In to afford the relevant homoallylic alcohols in yields ranging from 37 to 92%, which, in general, are comparable to those reported using organic solvents or water.<sup>[106a]</sup> Interestingly, this study reveals that, at the end of the reaction, addition of water to quench the putative indium alkoxide intermediate is essential in order to achieve the above yields for the relevant homoallylic alcohols. Using this approach for the allylation of 2-methoxycyclohexanone (**60**), the level of diastereoselectivity towards the *syn* product **61** was greater (**61** : **62** ratio = 18.6:1) than using pure water or a THF–water mixture (Scheme 30).

Related to these studies is the work of Chan and co-workers who compared the ability of In, Sn or Zn to mediate the allylation of carbonyl compounds using *ILs* [BMIM][BF<sub>4</sub>] and [EMIM][BF<sub>4</sub>]. This study reveals that under ambient temperature conditions each metal can effectively promote the formation of the relevant homoallylic alcohols in the above solvent systems, although the best conversions are observed for Sn.<sup>[106b]</sup>



**Scheme 30.** Indium-mediated allylation of 2-methoxycyclohexanone **60** using *IL* [BMIM][BF<sub>4</sub>].

The same group has also shown that aldimines can undergo nucleophilic addition with allylindium reagents, generated *in situ* from In and allylbromide in a mixture of *ILs* [bpy][BF<sub>4</sub>]/[bpy][Br] (bpy = *N*-butylpyridine), affording homoallylic amines **63** in good yields (66–99%) (Scheme 31). Mechanistic studies have revealed that in these reaction media allylindium(I) and allylindium(III) dibromide are in equilibrium, with the former being the most reactive towards imines in the formation of **63**, whereas the latter accounts for the formation of the bis(allylated) amine **64**. Interestingly, the use of bromide ion as an additive (in the form of *IL* [bpy][Br]) shifts the position of this equilibrium towards an In(I) species, which promotes the selective formation of **63**.<sup>[106c]</sup>



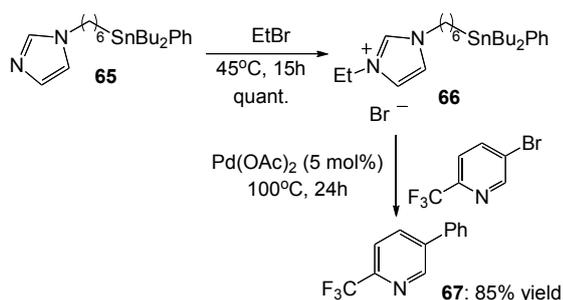
**Scheme 31.** Indium-mediated allylation of imines in [bpy][BF<sub>4</sub>]/[bpy][Br].

From a more cost-effective perspective, Hirashita has reported the efficient allylation of carbonyl compounds in *ILs* using catalytic amounts of In, which can be generated *in situ* by reduction of InCl<sub>3</sub> (10 mol%) with stoichiometric aluminium. Notably, these reactions occur faster when water is added to the *IL* [BMIM][PF<sub>6</sub>], although it should be noted that when the same approach was employed using neat water as the solvent, the allylation process was completely shutdown.<sup>[106d]</sup> In addition, once In-mediated aldehyde allylation processes are carried out using liquid CO<sub>2</sub> as the solvent, the relevant homoallylic alcohols can be isolated in 38 to 82% yield.<sup>[107]</sup> This method represents a cleaner and efficient alternative to conventional organic solvents, where the excess of CO<sub>2</sub> can be separated by depressurisation and subsequently be reused.

## 4.2. Applications of organotin reagents

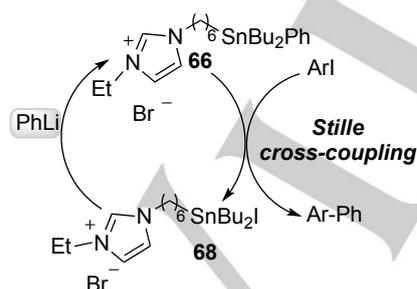
Finding widespread applications in cornerstone synthetic methods (Stille coupling, radical reactions, allylations, etc.), organotin compounds are a family of versatile organometallic reagents. Part of their popularity stems from their thermal stability and relatively straightforward preparations combined with their robustness to hydrolysis and oxidation. Furthermore, these reagents are also compatible with a myriad of organic

functional groups, showing an excellent balance between stability and reactivity. However, despite such an impressive synthetic background, one of their main drawbacks is related to their toxicity and the difficulties associated with the removal of residues from the final products. Some of the strategies developed to try to overcome this limitation include the use of organotin reagents supported by *ILs*.<sup>[108]</sup> These reagents can be easily prepared by treating imidazole derivative **65** with MeI or EtBr forming the *IL* supported tin reagent **66** that, in turn, can be used in Stille cross-coupling reactions, under solvent-free conditions and without the addition of additives or ligands, thereby affording a range of bis(aryl) compounds of the kind of **67** in good yields (Scheme 32).



**Scheme 32.** Synthesis of *IL* **66** and its application in a Stille cross-coupling reaction to give the biaryl derivative **67**.<sup>[108b]</sup>

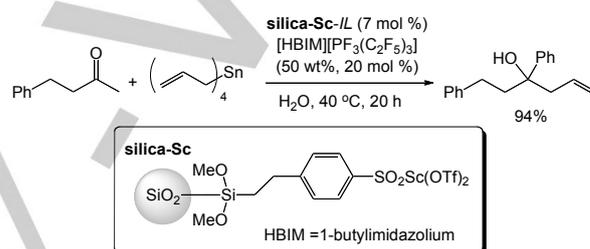
Furthermore, it is possible to recycle the tin compound/catalyst system at the end of the reaction by extracting the organic products with an organic solvent. By treating the *IL* phase containing the halogenotin-supported ionic liquid **68** with PhLi, it is possible to generate the arylating starting material **66** (Figure 5). These organotin reagents can also be used as effective catalysts for the reductive amination of aldehydes and ketones using PhSiH<sub>3</sub>. Reactions can also be carried out under solvent-free conditions, which facilitate the purification of the final products and minimise problems caused by tin contamination.<sup>[108c]</sup>



**Figure 5.** Recycling of *IL*-supported tin reagents **66**.

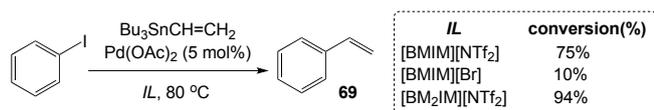
It should also be noted that allylation reactions and Stille couplings, two of the most powerful synthetic applications of organotin reagents, have also been investigated using *ILs* as alternative reaction media to organic solvents. Under these conditions, successful allylation methods of aldehydes and

ketones have been reported using several tin reagents, including Sn metal, SnCl<sub>2</sub> or tetra(allyl)tin. These reactions offer an excellent substrate scope and can be carried out at room temperature. Even more importantly, in many cases, the *IL* can be efficiently recycled without any further purification, thus making these protocols more environmentally benign.<sup>[109]</sup> Moreover, using tetra(allyl)tin it is possible to activate all-four allyl groups towards their transfer to the carbonyl substrates, thereby maximizing the atom-economy of the process.<sup>[109a]</sup> Related to this work is that by Kobayashi and co-workers who described a silica-gel-supported scandium system with an *IL* acting as a heterogeneous catalyst to efficiently promote several C–C bond-forming processes, including the allylation of ketones using tetra(allyl)tin (Scheme 33).<sup>[110]</sup> Intriguingly, this study shows that the key to success of this novel approach is the combination of a silica-gel-supported metal catalysts with an *IL*, which creates a hydrophobic reaction environment in water.



**Scheme 33** Organotin-mediated allylation reaction catalysed by silica-Sc-*IL* in water.

Stille couplings of organostannanes catalysed by Pd complexes have also been studied in *ILs*.<sup>[111]</sup> Pioneering work by Handy and Zhang,<sup>[112]</sup> indeed, showed that Stille coupling reactions can be successfully carried out using [BMIM][BF<sub>4</sub>] as the reaction medium, thus allowing the effective recycling of the solvent and the catalyst without significant activity loss. Notably, these processes are particularly sensitive to the structure of the *IL* employed.<sup>[113]</sup> For example, as illustrated in Scheme 34, once nucleophilic *ILs* such as [BMIM][Br] are used for the coupling of iodobenzene and tributylvinyl stannane under Pd(OAc)<sub>2</sub> catalysis, compound **69** is isolated in very low yields. Conversely, the employment of *ILs* with *N*-containing anions such as [NTf<sub>2</sub>]<sup>−</sup> (NTf<sub>2</sub><sup>−</sup> = bis(trifluoromethylsulfonyl)imide) allowed higher conversions (up to 94%) under ligand-free reaction conditions (Scheme 34). This dramatic difference in the *IL* performance has been attributed by Chiappe and co-workers to the nucleophilic assistance of the anion NTf<sub>2</sub><sup>−</sup> in the transmetallation step of the coupling reaction, which allows the coordination expansion of tin.<sup>[114]</sup> On the contrary, a similar type of activation would be less likely to occur in [BMIM][Br], due to the stronger cation–anion interactions present in this *IL*. Notably, and despite the higher reactivity observed in these NTf<sub>2</sub><sup>−</sup>-based *ILs*, the stability of the catalyst is very low, which precludes the effective recycling of the system.



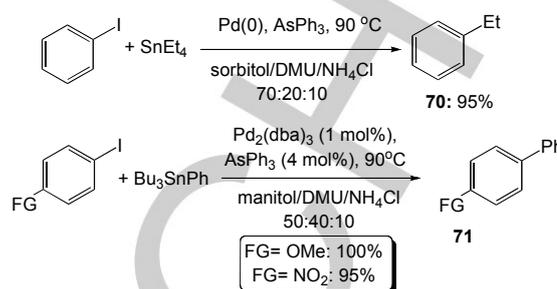
**Scheme 34.** Ligand-free Stille cross-coupling of iodobenzene with tributylvinyl stannane in *ILs*.

This drawback can be overcome by using Pd nanoparticles as catalysts rather than molecular palladium species.<sup>[115]</sup> The versatility and tuneability of *ILs* allows the stabilisation of nanoparticles, protecting them from agglomeration, while increasing the robustness to oxidation and hydrolysis of the nanoscale catalyst surface thanks to the formation of a protective shell. In these reactions, Pd nanoparticles act as catalyst reservoirs, while the active catalytic species are molecular Pd complexes, which can be leached out from the surface of the nanoparticle.<sup>[116]</sup> A wide range of *ILs* have been assessed for this type of Stille coupling, containing a variety of cations (e.g., pyridinium, imidazolium, tetraalkyl ammonium). Notably, Dyson has shown that the use of nitrile functionalised *ILs* improved significantly the stability of the catalytic systems allowing their efficient recycling and minimising catalyst leaching.<sup>[115b]</sup> Employing Pd nanoparticles stabilised by tetraalkyl ammonium salts bearing long alkyl chains as catalyst, Nacci and co-workers have reported efficient Suzuki and Stille cross-couplings involving a wide range of halide aryls, including aryl chlorides, whose applications in these type of processes can be particularly challenging due to their reduced reactivity.<sup>[115c]</sup>

*DESs* have also been successfully employed in Stille alkylations and biaryl synthesis. Köning and co-workers have reported that using low-melting mixtures of sugar, urea and inorganic salts as solvents it is possible to promote the fast and efficient Pd-catalysed alkyl transfer of tetraalkyltin reagents (Scheme 35).<sup>[117]</sup> Using conventional organic solvents, the transfer of simple alkyl groups requires instead special conditions including the use of toxic solvents such as HMPA or DMF. The smooth formation of coupling products **70** (Scheme 35) using this alternative method has been attributed to the high polarity and nucleophilic character of the *DES* combinations employed. Biaryls **71** can also be prepared in almost quantitative yields using this approach, which works well with both electron-poor and electron-rich aryl bromides (Scheme 35). Interestingly, the catalyst load can be reduced to 0.001 mol% and the catalyst-solvent mixture can be recycled up to three times. An added advantage to this approach is the simple work up and the ease at which products are isolated employing these melt mixtures; in fact, upon adding water, the organic products precipitate as amorphous solids and can be separated by filtration.

Although traditionally Stille couplings are carried out in organic solvents, the stability that aryl stannanes exhibit to air and moisture has allowed the development of alternative methods using water as the solvent.<sup>[118]</sup> For example, Wolf has reported an effective Stille cross-coupling methodology using a wide range of aryl chlorides and aryl bromides. Using an air-stable and water-soluble Pd-phosphinous acid catalyst, the synthesis of several bis(aryl) compounds can be accomplished

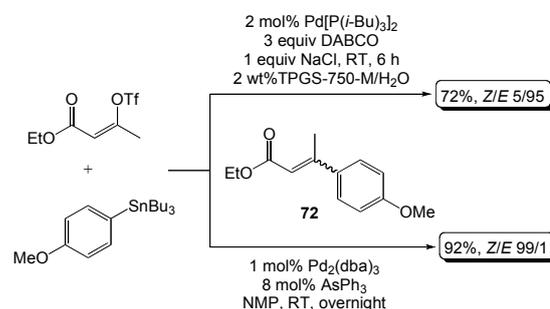
in good to high yields, using neat water as the solvent and without the need of an organic co-solvent.<sup>[118d]</sup>



**Scheme 35.** Stille couplings in sugar-urea-salt melts (DMU = dimethylurea).

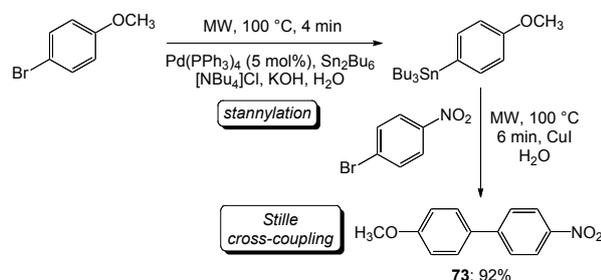
Interestingly, the catalyst can be recycled up to four runs with just a slight decrease in the observed yields (from 96% to 84% yield). Furthermore, these recycling studies demonstrate that under these conditions, the coupling products can be easily separated by extraction from the water-soluble catalyst. Notwithstanding, it should be noted that these reactions need to be carried out at high temperatures (135–140 °C) and over long periods of time (up to 24 hours). Milder reaction conditions for these couplings have been reported using dendrimer encapsulated Pd nanoparticles, which can catalyse Stille reactions in water at room temperature.<sup>[119]</sup> However, the substrate scope of these approaches is relatively limited.

In this regard, Lipshutz and Lu have developed an alternative strategy which builds on their previous work on Pd-catalysed Negishi cross-couplings in water (Scheme 20).<sup>[86]</sup> The use of TPGS-750M as a surfactant (able to undergo self-assembly in water to form nanomicelles) and a Pd[P(*t*Bu)<sub>3</sub>]<sub>2</sub>/DABCO (DABCO = 1,4-diazabicyclo[2.2.2]octane) combination as a catalyst, enables the efficient coupling of an impressive range of aryl and alkenyl halides to be conducted at room temperature with water as the only reaction medium.<sup>[120]</sup> These reactions not only take place in high yields but, in some cases, they also offer greater and different stereoselectivities than when using conventional solvents, as shown in Scheme 36 for the formation of **72**.



**Scheme 36.** Stille couplings with a *Z*-alkenyl triflate using NMP as the solvent and under aqueous micellar conditions.

Using water as the solvent, a simple and efficient one-pot methodology has been developed by Duan and co-workers: with Pd(PPh<sub>3</sub>)<sub>4</sub> as a catalyst, effective cross-coupling of a variety of aryl and heteroaryl bromides/iodides could be accomplished.<sup>[121]</sup> This method requires the use of microwave irradiation and involves the sequential stannylation followed by a Stille cross-coupling process, as depicted in Scheme 37 for the synthesis of **73**.



**Scheme 37.** Open-pot stannylation/Stille cross-coupling sequential reactions.

## 5. Conclusions

Overall, it is clear from the examples discussed in this Microreview that the employment of unconventional reaction media in the chemistry of s-, p-, and d-block elements is taking the stage today transforming everything and everyone. In a world with dwindling petroleum resources, organometallic scientists are slowly but firmly nurturing their own ability to improve the environmentally friendliness of chemical processes so as to produce the best and original “green organometallic chemistry”. The reticence to carry out the reactions of polar organometallic reagents under air, at room temperature, and in protic reaction media is also destined to fade away. The water itself, which is the most abundant and the only natural solvent on Earth, as well as the main traditional allied of biochemical processes, should no longer be considered a “foe” for organometallic compounds, but a “friend” to win.

The change in thinking will be the key to shift definitively our own paradigms and convictions. This will require a complete redesign of the experimental conditions, a deeper understanding of organometallic mechanisms and reactivities, and a close collaboration between theoreticians and experimental inorganic and organic researchers. In addition, fostering stronger industry/academic partnerships will expedite the transfer of new knowledge and emerging “greener technologies” to the marketplace to finally see realized in topic areas of science what they were simply considered not long ago as “fragile and visionary ideas”. Now that the first seeds are being planted, it would be expected and desirable that this infant field would grow up and develop quickly beyond our present concepts, and that the lion's share of organometallic transformations (stoichiometric, catalytic, and asymmetric applications) be designed and carried out mainly according to the “Green Chemistry” regulations. Until today, we have developed an extraordinary ability to adapt the word to our expectations. It is time now to adapt our expectations to fit the environment needs, hoping this may also lead to new scientific findings and to the disclosure of novel and

unpredictable aspects of organometallic reactivity in the years to come!

**Joaquín García Álvarez** studied chemistry at the University of Oviedo and received his PhD degree in 2005 under the supervision of Prof. José Gimeno and Dr. Victorio Cadierno studying the coordination of iminophosphorane-phosphine and bis(iminophosphorane)methane ligands in arene-Ru(II) fragments. Then, he joined the group of Prof. Robert E. Mulvey at the University of Strathclyde in Glasgow (Scotland, United Kingdom) working for two years and a half in the field of main group chemistry (alkali-metal-mediated-metallation). He returned to the University of Oviedo in 2008 where he is currently a “Ramón y Cajal” postdoctoral researcher. The current focus of his research are: (i) the study of synthetic organic reactions mediated by highly-polarized organometallic compounds in green and biorenewable reaction media (e.g., Deep Eutectic Solvents, DESs), and (ii) the application of transition metal complexes (e.g., Ru, Re, Pd, Au, Cu, Ag) containing iminophosphorane ligands in homogeneous catalysis using environmentally friendly solvents (water, ionic liquids, glycerol and DESs) as reaction media. He is co-author of 45 publications and eight book chapters in the field of *Organometallic Chemistry* and *Green Chemistry*.

**Eva Hevia** received her MSci degree in Chemistry from the Universidad de Oviedo (Spain) in 1998. She obtained her PhD degree from the same institution in 2002 under the supervision of Victor Riera and Julio Pérez. Next she was awarded a Marie Curie Fellowship held at the University of Strathclyde under the direction of Prof. Robert Mulvey. In 2006 she took up a Royal Society University Research Fellowship at the University of Strathclyde where she is currently a Professor of Inorganic Chemistry. Her research interests include s-block metal-mediated transformations with a particular emphasis on metal-metal cooperativity and synergistic effects.

**Vito Capriati** obtained his MSci degree in Chemistry and Pharmaceutical Technology (summa cum laude) from the University of Bari “Aldo Moro” (Italy) in 1990. After working as a forensic chemist officer within the Carabinieri's RIS (Scientific Investigation Department) of Rome and earning a two-year graduate fellowship within the Italian National Research Council (CNR Centre MISO, then merged into ICCOM-CNR), in 1993 he became Assistant Professor before taking up his present appointment as Associate Professor of Organic Chemistry in the University of Bari in 2002, where he leads the Bioorganic and Organometallic group. He has been Visiting Scientist at The Ohio State University (USA) (Prof. Fraenkel's group) (2001), and Visiting Professor at the Göthenburg University (Sweden) (2003). He is co-founder of the academic spin-off SYNCHIMIA srl and departmental coordinator of three Erasmus Programmes. His current research interests revolve around: functionalised organolithiums (structure elucidation, dynamic stereochemistry of chiral compounds, and the discovery of new reactions), synthesis and reactivity of new heterocycles, novel sustainable chemical processes using unconventional reaction media, organofluorine and organoboron chemistry, and the development of new drugs for rare diseases. He has published over 90 peer-reviewed journal articles, 5 book chapters and 7 reviews. He was the recipient of the CINMPIS Prize for

"Innovation in Organic Synthesis" (2009), of the Award of the Italian Chemical Society (Organic Division) for "Mechanistic and Theoretical Aspects of Organic Chemistry" (2014), and the Italian coordinator of an international Italian-German bilateral Vigoni Project (2012-2013). Co-Editor of "Lithium Compounds in Organic Synthesis – From Fundamentals to Applications"; Publisher: Wiley-VCH (2014).

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**Keywords:** organometallic compounds • water • deep eutectic solvents • ionic liquids • reactivity

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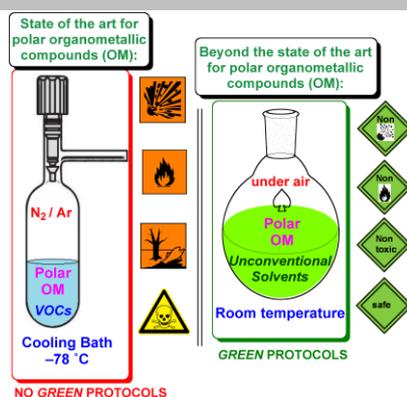
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## MICROREVIEW

**Stepping towards ideal polar organometallic processes!**

This Microreview provides a reliable evidence of the unstoppable transition of polar organometallic chemistry towards greener protocols, mainly represented by unconventional solvents. This contributes to make organometallic chemistry even more sustainable and prepares the ground for new, stimulating research.

**Green Chemistry**

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and Vito Capriati\*

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**Reactivity of Polar Organometallic  
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Opportunities**