## **RSC** Advances

### PAPER

View Article Online View Journal | View Issue

### Oxazol-2-ylidenes. A new class of stable carbenes?†

Cite this: RSC Advances, 2013, 3, 7970

Received 10th January 2013, Accepted 15th March 2013

DOI: 10.1039/c3ra41177j

www.rsc.org/advances

#### Introduction

The chemistry of stable carbenes<sup>1</sup> has been flourishing in the past decades. Apart from their unusual structure, the interest is generated by their catalytic applicability in various organic and organometallic reactions. As ligands they can fine-tune the activity of transition metal catalysts,<sup>2</sup> furthermore, they can mimic<sup>3</sup> transition metals and can be powerful organoca-talysts.<sup>1b,4</sup>

Perhaps the most widely investigated carbenes are imidazol-2-ylidenes (1) – see Scheme 1, which exhibit remarkable stability: these compounds can be kept for years under inert atmosphere, and the first synthesized example could even be molten at *ca.* 240 °C without any decomposition.<sup>5</sup>

This stability can be attributed to the effective  $\pi$ -electron donating character of the nitrogen atoms, and also to the  $6\pi$ -aromaticity, which brings an additional stabilization of about 10 kcal mol<sup>-1</sup>,<sup>6</sup> contributing to their reduced reactivity,<sup>7</sup> and affecting their catalytic activity.<sup>8</sup> The significant stabiliz-

Zsolt Kelemen,<sup>a</sup> Oldamur Hollóczki,<sup>ab</sup> Julianna Oláh<sup>ac</sup> and László Nyulászi\*<sup>a</sup>

The investigation of the stability of several imidazol-2-ylidene analogue cyclic carbenes by an isodesmic reaction has revealed that the hitherto unknown oxazol-2-ylidene exhibits only slightly smaller stability than imidazol-2-ylidene, outperforming some of the already synthesized carbenes. Selenazol-2-ylidene also shows significant stability. The contribution of aromaticity to the stabilization has been analysed for the different five-membered ring carbenes, and was found to be relatively small for the oxygen containing systems. Investigation of possible reactivity/decomposition pathways reveals that properly substituted oxazol-2-ylidene is stable against dimerization. The thermodynamically feasible cycloreversion reaction yielding isocyanate and acetylene is prevented by a significant barrier, and furthermore with proper substitution (ring annellation) the ring can be stabilized thermodynamically as well. While in the presence of water a hydrolytic ring opening occurs; this reaction can be hindered if the water content of the reaction mixture is reduced to a few equivalents. This hydrolytic behaviour as well as the electrophilicity and nucleophilicity indices of several known nucleophilic carbenes were compared, revealing that oxazol-2-ylidene exhibits a reduced nucleophilicity with respect to imidazol-2-ylidene, while its electrophilicity is only slightly increased. This unique combination might result in unexpected (organo)catalytic activities, further expanding the colourful applications of NHCs.

ing effect of amino groups at the hypovalent center prompted the synthesis of further diaminocarbenes, such as imidazolidine-2-ylidene<sup>9</sup> (2), triazol-2-ylidene<sup>10</sup> (3) and also a stable noncyclic bisdiisopropylaminocarbene (4).<sup>11</sup> Despite the apparent dominance of nitrogen substitution in stabilizing the carbenes, some derivatives could also be synthesized with two significantly flattened<sup>12</sup> phosphorus<sup>13</sup> (5) which also benefit from aromaticity,<sup>14</sup> or most remarkably, derivatives could be synthesized with a nitrogen and a chalcogen atom, like thiazol-2-ylidene<sup>15</sup> (6) and non-cyclic aminothio-<sup>16</sup> and aminooxycarbenes<sup>16</sup> (7 and 8). Even cyclic (alkyl)(amino) carbenes (CAAC) are known.<sup>17</sup>



Scheme 1  $\pi$ -electron donor stabilized carbenes.

<sup>&</sup>lt;sup>a</sup>Department of Inorganic and Analytical Chemistry, Budapest University of Technology and Economics Szt, Gellért tér 4, Budapest, H-1111, Hungary. E-mail: nyulaszi@mail.bme.hu; Fax: +3614633408

<sup>&</sup>lt;sup>b</sup>Wilhelm-Ostwald-Institut für Physikalische und Theoretische Chemie Universität Leipzig, Linnéstr. 2, D-04103 Leipzig, Germany

<sup>&</sup>lt;sup>c</sup>Materials Structure and Modeling Research Group Hungarian Academy of Sciences, at Budapest University of Technology and Economics Szt, Gellért tér 4, Budapest, H-1111, Hungary

<sup>†</sup> Electronic supplementary information (ESI) available. See DOI: 10.1039/ c3ra41177j

# I. $R_1R_2C$ : + $CH_4 \longrightarrow R_1R_2CH_2$ + : $CH_2$

**Scheme 2** Isodesmic reaction to estimate the stability  $\Delta E_{(carbene)}$  of a carbene.

To estimate the (thermodynamic) stabilization of carbenes, an isodesmic reaction I (Scheme 2) has been applied,<sup>18</sup> exhibiting more than  $\Delta E_{\text{(carbene)}} = 90 \text{ kcal mol}^{-1}$  stabilization for each of the known synthesized carbenes. Furthermore  $\Delta E_{(\text{carbene})}$  has been shown to exhibit excellent correlation with the dimerization energy of carbenes.<sup>18</sup> Very recently, we have shown<sup>19</sup> that N-methyloxazol-2-ylidene exhibits 96.1 kcal  $mol^{-1}$  (B3LYP/6-311+G<sup>\*\*</sup>)  $\Delta E_{(carbene)}$  stabilization, exceeding the 92.4 kcal mol<sup>-1</sup> value for the analogous thiazol-2-ylidene (6) derivative. Similar conclusions were obtained in a different study, in which a singlet-triplet gap of 75.8 kcal mol<sup>-1</sup> (B3LYP/  $6-311+G^{**}$ ) was reported<sup>20</sup> for oxazol-2-ylidene (9) that is only slightly lower than the corresponding value for imidazol-2ylidene (83.0 kcal mol<sup>-1</sup> at the B3LYP/6-311+G\*\*//B3LYP/6-31G\* level).<sup>21</sup> It was also surmised that oxazol-2-ylidene (9) benefits from  $6\pi$ -aromatic stabilization, although this effect has not been quantified comprehensively.<sup>20</sup> Considering these indications of stability, and particularly in the light of the stability of 8, it is surprising that known chalcogen-substituted N-heterocyclic carbenes (NHCs) are limited to thiazol-2-ylidene (6),<sup>15,22</sup> and no free derivatives of oxazol-2-ylidene (9) have been prepared yet, although its complexes with transition metals<sup>23</sup> and BPh<sub>3</sub><sup>24</sup> are known. To assess the overall stability of a carbene, however, not only thermodynamical aspects,<sup>25</sup> but possible chemical decomposition reactions should also be considered. The computational investigation of the cycloreversion of cyclic dioxocarbenes<sup>26</sup> to ethyne and carbon dioxide (Scheme 3) revealed an exothermic reaction, with a low (8.8 kcal mol<sup>-1</sup> MP2/6-31G\*\*) barrier. This finding might indicate the tendency of cyclic chalcogen-substituted carbenes to decompose as a possible reason for the lack of success in synthesis of the derivatives.

On the other hand, carbon disulfide can react with strained<sup>27*a*</sup> or electron-deficient<sup>27*b*</sup> acetylenes at high temperature or under high pressure yielding 1,3-dithiolium carbenes, which can be trapped by the substrate, or dimerize to tetrathiafulvenes (Scheme 4).<sup>27</sup> The reaction of carbon disulfide with *t*-BuCP was also shown to proceed *via* the cycloaddition intermediate.<sup>28</sup> While for some 1,3-heterocumulenes the cycloreversion was investigated and the stability of



Scheme 3 Decomposition reactions of dioxol-2-ylidene and dioxolidin-2-ylidene.



Scheme 4 Reaction between acetylene and carbon disulfide (R: electron withdrawing group e.g.  $CF_3$ ).

nitrogen-containing rings was established, in that study<sup>29</sup> neither oxazol-2-ylidene (9) nor selenazol-2-ylidene (10) was considered. Changes in the carbenes' stability might trigger changes also in their catalytic activity. We have recently demonstrated<sup>19</sup> that the Breslow-intermediate – which is a key intermediate in many organocatalytic reactions (e.g. in the benzoin condensation) - possesses superior stabilization with thiazol-2-ylidene catalysts compared to other NHCs, affecting the catalytic activity. Also N,N'-diamidocarbenes (DAC) are engaged in reactions associated with electrophilic carbenes.<sup>30</sup> Furthermore, even the substituents on the nitrogen of the NHC were shown to affect the catalytic effect significantly.<sup>31</sup> In light of these results the design of new stable carbenes is of great importance in order to broaden the number of available catalysts, which might aid in choosing the most proper (and selective) one for a given reaction. As shown above, little is known about the stability of chalcogen-substituted N-heterocyclic carbenes, despite the apparent stability of the open-chain analogues. Thus, the aim of the present computational study is to explore the stability and reactivity of these species in detail, with a special emphasis on oxazol-2-ylidene.

#### **Computational methods**

Unless otherwise stated, all calculations were carried out with the Gaussian 03 program package,<sup>32</sup> using the B3LYP hybrid functional in conjunction with the 6-311+G\*\* basis. Full geometry optimizations were performed for all molecules, and harmonic vibrational frequencies were calculated at the same level of theory to establish the nature of the stationary points obtained, as characterized by none or a single negative eigenvalue of the Hessian for minima and transition structures, respectively. Intrinsic reaction coordinate (IRC) calculations were started to locate the corresponding minima. Nucleus-independent chemical shift (NICS) values<sup>33</sup> were calculated as the negative of the magnetic shielding at the geometric centre of the ring (NICS(0)) and also by 1 Å above the plane of the ring atoms (NICS(1)). Bird indices<sup>34</sup> (BI) were determined from the Gordy bond orders<sup>35</sup> of the ring using the bond distances of the simplest single- and double-bonded compounds (CH3-CH3, CH2=CH2, CH3-NH2, CH2=NH, CH3-OH, CH2=O, CH3-SH, CH2=S, CH3-SeH, and CH2=Se) calculated at the same level of theory as the references. Isomerisation stabilization energies (ISE, Scheme 5)<sup>36</sup> were calculated as an energetic measure of aromaticity.

Scheme 5 Isomerisation stabilization energy reaction.

In order to verify our results, certain calculations were also performed at the B3LYP/aug-cc-PVTZ and MP2/6-311+G\*\* levels of theory. To calculate (gas phase) Gibbs free energies the computed harmonic frequencies were used at 298 K and 1 atm. In order to get a deeper insight into the factors determining the reactivity of the investigated carbenes, we employed conceptual DFT indices,37 originating from the ground state electron density of the molecules, to estimate their hardness, electrophilic and nucleophilic characters. Pérez showed<sup>38</sup> that a good qualitative agreement existed between the theoretical and experimental philicity of carbenes by comparing the electrophilicity index of singlet carbenes to the experimental Moss scale,<sup>39</sup> which is based on a kinetic model of carbene addition to simple alkenes. Similarly, in a study of the divalent silicon and germanium analogues of carbenes, the electrophilicity index has been shown to be a valuable tool in assessing the reactivity of the compounds.<sup>40</sup> The electrophilicity index ( $\omega$ , eqn (1)) is calculated as

$$\omega = \frac{\mu^2}{2\eta} \tag{1}$$

where  $\mu$  is the chemical potential<sup>41</sup> (eqn (2)) and  $\eta$  is the chemical hardness<sup>41</sup> (eqn (3)) of the molecule.

$$\mu = \left(\frac{\partial E}{\partial N}\right) \approx \frac{\mathrm{IE} + \mathrm{EA}}{2} \approx \frac{\varepsilon_{\mathrm{HOMO}} + \varepsilon_{\mathrm{LUMO}}}{2} \tag{2}$$

$$\eta = \left(\frac{\partial^2 E}{\partial N^2}\right) \approx \frac{\mathrm{IE} - \mathrm{EA}}{2} \approx \frac{\varepsilon_{\mathrm{LUMO}} - \varepsilon_{\mathrm{HOMO}}}{2} \tag{3}$$

As direct evaluation of these indices is not straightforward we used two finite differences approaches,<sup>37</sup> commonly applied in the literature, based on the ionization energy (IE) and electron affinity (EA) of the molecules, and on their approximations based on Koopman's theorem.<sup>42</sup> As the trends obtained from the two formulas were identical, only results based on orbital energies are presented here, while the rest of the data are given in the SI. The nucleophilicity index (*N*) has been calculated as the energy difference between the HOMO energy of the compound and that of tetrachloretylene ( $E_{\text{HOMO(TCE)}}$ , eqn (4)) as was proposed by Domingo *et al.*<sup>43</sup> and was successfully used earlier for carbenes as well:<sup>44</sup>

$$N = \varepsilon_{\rm HOMO} - \varepsilon_{\rm HOMO(TCE)} \tag{4}$$



**Fig. 1** Stabilities of heterocyclic carbenes ( $\Delta E_{(carbene)}$  [kcal mol<sup>-1</sup>]) for different X and Y ring heteroatoms.

#### **Results and discussion**

First, the thermodynamical stability has been evaluated for five-membered, chalcogen-substituted carbenes using  $\Delta E_{(carbene)}$ , to find possible target molecules for synthesis. In Fig. 1. the B3LYP/aug-cc-PVTZ stability of the investigated saturated and unsaturated carbenes are presented, while the MP2/6-311+G\*\* and B3LYP/6-311+G\*\* data are compiled in Table S1 in the ESI.<sup>†</sup>

These results show the decreasing stabilizing effect of the heteroatoms in the order of  $N > O > S \sim$  Se. Oxazol-2-ylidene (9) exhibits the second largest stabilization energy (98.5 and 100.7 kcal mol<sup>-1</sup> at the B3LYP/aug-cc-pVTZ and MP2/6-311+G\*\* levels, respectively), exceeding the stabilization of some already synthesized carbenes, such as 2 and 6 (96.2 and 95.5 kcal mol<sup>-1</sup> at B3LYP/aug-cc-pVTZ, respectively). It should also be mentioned that the stabilization of selenazol-2-ylidene 10 is similar to that of its sulphur analogue (6). All the unsaturated compounds exhibit higher  $\Delta E_{(carbene)}$  values than their saturated counterparts, showing the stabilizing contribution of aromaticity. The difference in  $\Delta E_{(\text{carbene})}$  values of the saturated/unsaturated pairs varies between 6 and 14 kcal mol<sup>-1</sup> (at B3LYP/aug-cc-pVTZ) and 6 and 18 kcal mol<sup>-1</sup> (at MP2/6-311+G\*\*). This indicates that the extent of aromaticity is influenced by the heteroatom, therefore we decided to investigate the aromaticity of the unsaturated systems in more detail. The largest<sup>45</sup> aromaticity is exhibited by imidazol-2ylidene (1) according to the different measures BI (60), ISE (-14.4 kcal mol<sup>-1</sup>) and NICS(0) (-13.0 ppm) NICS(1) (-10.1 ppm) (Table 1), indicating considerable cyclic delocalisation as noted before.<sup>6</sup> Gradual substitution of the nitrogen(s) by other heteroatoms results in a decrease of the aromaticity in the order of N > S > Se  $\sim$  O. Accordingly, the energetic measure of aromaticity also shows the rather small (ISE -9.6 kcal mol<sup>-1</sup>) but still non-negligible<sup>26b</sup> aromatic stabilization of oxazol-2-ylidene in line with the rather small (6.8 kcal  $mol^{-1}$ -Table S1, ESI<sup>†</sup>, Fig. 1) difference between the saturated and unsaturated  $\Delta E_{(carbene)}$  stabilization energies. It is worth noting that from the investigation of the magnetic susceptibilities of 1, 6 and 9, the smallest aromaticity was also found for **9**.<sup>46</sup>

**Table 1** Aromaticity measures: NICS(0) and NICS(1) (in ppm), Bird Index (BI), and ISE values (in kcal mol<sup>-1</sup>) for the unsaturated carbenes. Electrophilicity ( $\omega$  in au) and nucleophilicity (N in au) indices for all investigated carbenes

			<b>X</b> X						
Х	Y	NICS(0)	NICS(1)	BI	ISE	ω	Ν	ω	Ν
NH NH NH S S Se O S	NH S O S Se Se O	-13.0 -13.0 -12.0 -12.2 -16.6 -13.4 -12.7 -12.3 -11.6	-10.1 -10.6 -9.6 -9.8 -11.9 -11.3 -10.8 -9.6 -10.2	60 56 54 41 53 51 50 28 41	-14.4 -12.6 -11.1 -9.6 -13.3 -11.9 -11.3 -8.7 -11.1	0.0558 0.0838 0.0926 0.0686 0.1318 0.1417 0.1507 0.0963 0.1170	$\begin{array}{c} 0.0509\\ 0.0313\\ 0.0344\\ 0.0244\\ 0.0158\\ 0.0168\\ 0.0166\\ 0.0039\\ 0.0175\end{array}$	0.0555 0.0703 0.0751 0.0637 0.1333 0.1419 0.1524 0.0768 0.1020	0.0561 0.0408 0.0441 0.0395 0.0249 0.0265 0.0289 0.0152
Se	0	-10.2	-9.5	37	-10.6	0.1270	0.0235	0.1020	0.0322

From the analysis of the effect of the carbene substituents and the aromatic stabilization it can be concluded that the exceptional stability of imidazol-2-ylidenes is mainly due to the favourable stabilizing effect of the nitrogen substituent on the carbene, supported further by aromaticity.<sup>6</sup> Interestingly, although the oxygen substituent brings a significant stabilization in the case of oxazol-2-ylidene, this heteroatom contributes to the aromaticity to a smaller extent only. In this respect it is worth pointing out the analogy with the wellknown trend in the aromaticity of five-membered heterocycles, namely that the aromaticity of furan is also significantly reduced with respect to pyrrole and thiophene.<sup>47</sup> Clearly, the stabilization of the  $\pi$ -type lone pair ionization energy with respect to its saturated counterpart is the smallest for furan in the series of common five-membered heterocycles.<sup>48</sup> A further important factor from the point of view of the carbene stability is its propensity for dimerization. While the Gibbs free energy for the dimerization of 6 is -22.1 kcal mol<sup>-1</sup>, for 9 it amounts only -12.2 kcal mol<sup>-1</sup>. The reduced dimerization energy of the (unsubstituted) 9 with respect to 6 is in accordance with the increased  $\Delta E_{(carbene)}$  value, and also with our previous results on the good correlation of these quantities.<sup>18</sup> The N-methylderivative of oxazol-2-ylidene 9, however, has only -6.8 kcal mol<sup>-1</sup> dimerization Gibbs free energy, showing that the N-substitution, even by the small methyl group, significantly reduces the propensity of dimerisation. Since N-substituted derivatives of 6 (which are less stabilized than 9) are known to exist in monomeric form,15 among N-substituted oxazol-2ylidenes several derivatives (see below) should exist in monomeric form. A further important chemical transformation, the 1,2-hydrogen shift resulting in oxazol from 9, has been shown earlier to proceed via a high barrier, which has further increased for the synthetically most interesting alkyl

The fragmentation of oxazol-2-ylidene to acetylene and an isocyanate by a cycloreversion, however, may occur and this reaction is also entropically favoured. Indeed, some of the studied reactions are exergonic, but not exothermic (Table 2).

<b>Table 2</b> The energy ( $\Delta E$ ), Gibbs free energy ( $\Delta G$ ), activation energy ( $\Delta E^{\#}$ ) and
activation Gibbs free energy ( $\Delta G^{\#}$ ) (at the B3LYP/6-311+G** level) of the
cvcloreversion reactions in kcal mol <sup>-1</sup>

Carbene			x_	Y		ẍ́y				
X	Y	$\Delta E$	$\Delta E^{\#}$	$\Delta G$	$\Delta G^{\#}$	$\Delta E$	$\Delta E^{\#}$	$\Delta G$	$\Delta G^{\#}$	
NH	NH	-1.3	35.0	-18.2	30.7	32.1	70.7	14.0	63.1	
NH	S	-12.8	20.7	-28.7	17.6	16.2	49.6	-0.3	44.7	
NH	Se	-10.6	20.6	-26.5	17.6	16.7	48.2	0.3	43.4	
NH	0	-27.1	18.0	-42.5	14.9	-1.3	43.7	-17.8	38.3	
S	S	-27.1	6.7	-39.3	5.4	-1.7	30.3	-14.6	27.5	
S	Se	-24.2	6.6	-34.0	5.4	0.0	29.4	-10.5	26.7	
Se	Se	-21.4	6.6	-32.8	4.7	1.7	28.7	-10.4	26.1	
0	0	-54.7	7.7	-65.4	6.0	-35.2	26.6	-46.6	23.5	
S	0	-47.3	5.7	-57.4	4.1	-24.3	26.2	-34.9	23.3	
Se	0	-45.5	5.3	-55.2	3.6	-23.8	24.5	-33.8	21.7	

The stability against cycloreversion drops significantly for the saturated compounds. The reaction proceeds via a concerted pathway (we were not able to locate transition structures for a stepwise reaction for imidazol-2-ylidene, or for oxazol-2ylidene). Nevertheless, the barrier in the case of the cyclic saturated diaminocarbene 2 is large enough to prevent the decomposition reaction in agreement with its synthesizability. Again, the saturated compounds have much lower decomposition barriers than their unsaturated counterparts (Table 2), and alkenes and isocyanates have been obtained from transient oxazolidin-2-ylidenes in agreement with this.49 Apparently, the effect of the heteroatoms on the cycloreversion differs from that observed for  $\Delta E_{\text{(carbene)}}$ .<sup>50</sup> In the case of the cycloreversion reaction nitrogen containing compounds still exhibit the largest stability, however, sulfur (and selenium) containing systems are less prone to cyclorevert than their oxygen analogues. For oxazol-2-ylidene 9 the cycloreversion reaction is exergonic. Nevertheless, the calculated 38.3 kcal mol<sup>-1</sup> barrier of the concerted reaction is large, and thus we can conclude that this compound is synthesizable. It is also noteworthy that for selenazol-2-ylidene 10 the exothermicity and the significant barrier of the reaction are comparable to those of the known sulfur analogue.

Since the substituents on the nitrogen and the carbon atoms have some apparent effect on the stability<sup>6a</sup> and reactivity<sup>31</sup> of carbenes, we examined the effects of substituents on oxazol-2-ylidene **9** and on selenazol-2-ylidene **10**. For the substitution at the nitrogen the bulky adamantyl, mesityl and supermesityl groups have been chosen, which provide extra steric shielding for the carbene centre (Table 3). These nitrogen substituents result in slightly increased stabilization energies, similar to our previous observations in case of pyridine-2-ylidene.<sup>6a</sup> It should be emphasized again that the effect of the steric shielding – which also contributes to the prevention of dimerization – is not included in the  $\Delta E_{(carbene)}$ energies, thus, the real stabilization should be larger. At the ring carbon atoms (positions 4 and 5), however, the size of the substituent is likely to be less important; thus, groups with

substituted systems.<sup>20</sup>

**Table 3** Effect of substitution on the carbene stabilization ( $\Delta E_{(carbene)}$ ), reaction energy ( $\Delta E$ ), reaction Gibbs free energy ( $\Delta G$ ), activation energy ( $\Delta E^{\#}$ ) and activation Gibbs free energy ( $\Delta G^{\#}$ ) of the cycloreversion reactions of oxazol-2-ylidene and selenazol-2-ylidene in kcal mol<sup>-1</sup>. Me: methyl, Mes: mesityl, Mes\* supermesityl (2,4,6-tritert-butylphenyl), Ad: adamantyl

R₁∼Ŋ <sup>΅</sup> X			0					Se				
	$R_2 R_2$	2										
R <sub>1</sub>	$R_2$	$R_2$	$\Delta E_{ m carbene}$	$\Delta E$	$\Delta E^{\#}$	$\Delta G$	$\Delta G^{\#}$	$\Delta E_{ m caebene}$	$\Delta E$	$\Delta E^{\#}$	$\Delta G$	$\Delta G^{\#}$
Н	Н	Н	96.1	-1.3	43.7	-17.8	38.3	92.3	16.7	48.2	0.3	43.4
Ме	Н	Н	99.8	1.5	45.2	-15.5	39.4	93.1	14.1	46.8	-4.1	43.1
Mes	Н	Н	100.8	-4.5	41.1	-20.5	36.0	93.1	7.0	42.5	-9.2	38.5
Mes*	Н	Н	102.6	-10.3	40.5	-29.0	35.6	93.1	2.6	40.9	-15.2	36.8
Ad	Н	Н	101.2	-2.2	44.0	-19.4	39.2	94.8	9.1	44.1	-5.6	39.8
н	Cl	Cl	95.0	2.7	47.8	-13.2	42.8	89.9	22.1	54.6	6.1	50.2
н	$BH_2$	$BH_2$	93.4	-14.1	26.3	-31.5	23.0	87.0	1.6	26.6	-15.9	23.3
н	$CH_3$	$CH_3$	98.7	-4.6	44.3	-22.5	39.4	92.1	9.2	47.0	-8.6	43.0
н	$CF_3$	$CF_3$	92.6	-1.4	39.1	-18.5	34.8	88.9	17.5	42.7	-0.9	37.5
н	$NH_2$	$NH_2$	95.3	-1.1	46.0	-17.9	41.7	88.9	13.9	52.4	-3.0	47.4
Н	$\mathbf{V}$		91.7	43.6	56.9	25.9	50.7	87.7	61.2	67.3	43.4	60.8
н	$\searrow$		96.1	34.8	55.2	16.6	48.7	91.0	51.6	62.0	33.4	55.9

different electronic nature (boro, methyl, trifluoromethyl, chloro, amino) have been selected. Similarly to the minor influence of substitution at the nitrogen, substituents at the carbon atoms in positions 4 and 5 exerted also a small effect; only the strong  $\pi$ -electron withdrawing groups (-BH<sub>2</sub> and -CF<sub>3</sub>) resulted in a few kcal mol<sup>-1</sup> decrease of the stability of the carbene (Table 3). Thus, the effect of the substituents on the  $\Delta E_{(\text{carbene})}$  stabilization of oxazol-2-ylidene **9** (and its selenium analogue **10**) is minor.

The position of the substituent has a significant effect on the cycloreversion reaction. While substitution at the nitrogen has a small impact (except for the bulky supermesityl group), having the substituent at positions 4 and 5 significantly effects both the reaction energy and the barrier (Table 3). For the boro-substituted derivates of oxazol-2-ylidene **9** and selenazol-2-ylidene **10** the cycloreversion reaction is more exothermic than for the parent carbenes, and also the reaction barrier is reduced. Ring anellation and condensation have an important effect on the reaction. First, the kinetic hindrance of cycloreversion increases significantly; furthermore, the reaction becomes endothermic due to the formation of the ringstrained cyclohexyne or benzyne, while the carbene stabilization energy exhibits only a small decrease.<sup>27a,29,51</sup> Thus the most stable synthetic targets as free carbenes are **11** and **12** (Scheme 6).

In the assessment of the stability an important aspect is the reactivity against common reactants such as water, acids and bases. Oxazol-2-ylidenes were already considered in early works<sup>52</sup> as alternative catalysts in the benzoin condensation to thiazol-2-ylidenes or imidazol-2-ylidenes, but the oxazolium salts did not show catalytic activity in the presence of a base.<sup>52</sup> Haake has shown that although oxazolium salts are more acidic than their thiazolium counterparts,<sup>53</sup> (*i.e.* they should produce the carbene easily) they exhibit a facile irreversible ring opening resulting in **13** (X = O, see Scheme 7) at pH values being sufficiently basic for the deprotonation.<sup>54</sup> In contrast, thiazolium exhibits a reversible ring opening, while to open up the imidazolium ring extremely basic conditions are needed.<sup>54</sup>

To describe the effect of water as solvent, recently we used a microsolvatation approach by embedding the starting carbene, intermediates and product in a water cluster.<sup>7,19</sup> For imidazol-2-ylidene, product **13** (X = NMe) was less stable than imidazolium-hydroxide by 4.7 kcal mol<sup>-1</sup>,<sup>7</sup> while thiazol-2-ylidene **13** (X = S) was more stable than the hydroxide by 7.4 kcal mol<sup>-1</sup>, using a cluster made from 31 water molecules.



Scheme 6 The most suitable targets for synthesis (R: bulky substituent e.g.: adamantyl or mesityl.



Scheme 7 Ring opened products from the hydrolysis of azolium salts.



**Fig. 2** B3LYP/6-311+G<sup>\*\*</sup>//B3LYP/6-31+G<sup>\*</sup> level optimized structures and relative energies (in kcal  $mol^{-1}$  units) of oxazol-2-ylidene with 31 water, oxazolium-hydroxide, **13** and **14** with 30–30 water molecules.

Using the same approach for oxazol-2-ylidene the water microsolvated structures for the oxazol-2-ylidene, oxazolium hydroxide, **13** (X = O) and its H-shifted isomer **14** were optimized as shown in Fig. 2. For the oxazol case, **13** (X = O) is significantly more stable than oxazolium-hydroxide. All these results are in full agreement with the earlier observations<sup>54</sup> of the irreversible ring opening in the case of the oxygen compound, and the intact nature of the imidazolium moiety in the presence of excess water.<sup>7</sup>

For the sake of completeness we have made similar calculations for triazol-2-ylidene and selanazole-2-ylidene. While the results (shown as Fig. S1 and S2 in the ESI†) for the former one are close to those for imidazol-2-ylidene, the selenazol-2-ylidene results are similar to those for thiazol-2-ylidene. It is worth noting that the energy of oxazol-2-ylidene in the 31 water cluster is nearly the same as for oxazolium-hydroxide, while imidazolium-hydroxide was more stable than the corresponding carbene by 6.1 kcal mol<sup>-1</sup>, in agreement with the relative basicities of these carbenes.<sup>19</sup> This shows that the microsolvation approach applied provides a reasonably good description for the behaviour in water as solvent.

Recently, however, we have shown in a combined computational-experimental study that the mechanism of imidazol-2ylidene hydrolysis depends on the number of participating water molecules.<sup>7</sup> While with excess water a fast protonation takes place – as was shown by the computations using the microsolvatation approach – with a small amount of water molecules the ring-opening reaction is a kinetically-hindered slow reaction. The computational studies indicate that the



**Fig. 3** The relative energies of the products of the hydrolysis of different carbenes at B3LYP/6-311+G<sup>\*\*</sup> level (in kcal mol<sup>-1</sup> units).

open chain products (Scheme 7) are more stable than the imidazol-2-ylidene-water complex, but a significant reaction barrier hinders the reaction with one water molecule, which is only slightly reduced in the presence of two or three reacting water molecules.<sup>7</sup>

A similar conclusion was drawn for thiazol-2-ylidene,<sup>19</sup> although the relative stability of the open chain 13 (X = S) was somewhat increased. We have carried out analogous calculations for the hydrolysis of oxazol-2-ylidene and for the sake of completeness we considered selenazol-2-ylidene and triazol-2ylidene as well. The details of the reaction pathways are given as Fig. S3-S5 of the ESI<sup>†</sup>, while the energetically most important points of the reactions are compiled in Fig. 3 together with the previously published data for imidazol-2ylidene<sup>7</sup> and thiazol-2-ylidene.<sup>19</sup> The comparison of the energetics for the different hydrolysis reactions reveals that while the activation energy for the reaction is reduced only slightly with respect to imidazol-2-ylidene and thiazol-2ylidene, the product 13 is significantly stabilized in case of X = O. For the calculation with two and three water molecules (the reaction pathways are presented in the ESI as Fig. S4-5, S7-8, S10-11<sup>†</sup>) again similar transition structures and intermediates were obtained as in case of imidazol-2-ylidene<sup>7</sup> and thiazol-2-ylidene.<sup>19</sup> Since in all these calculations the barriers are similar to the imidazol-2-ylidene case, it seems reasonable to assume that the hydrolysis reaction is likewise hindered for oxazol-2-ylidene, if the water concentration is small (and the clustering is reduced).55

Having concluded that properly substituted oxazol-2ylidenes (11) and selenazol-2-ylidenes (12) (Scheme 6) are viable synthetic targets, it is worth investigating their expected reactivity. It is well known that the divalent carbon centre in singlet carbenes is characterized by an empty 2p orbital and an in-plane lone pair-like orbital, which may give rise to its electrophilic and/or nucleophilic character. In general,  $\pi$ -electron donation from the substituents to the empty p orbital of the divalent centre is expected to reduce its electrophilic character. Furthermore, it has been shown that the nucleophilicity remains virtually unchanged upon aromatization (comparing 1 and 2),<sup>56</sup> likewise the donating ability as Paper

measured by the Tolman electronic parameter.<sup>57</sup> In the case of silvlenes (the silicon analogues of carbenes) the relationship between the  $\pi$ -electron donating ability, nucleophilic and electrophilic character has been studied in detail,40,58 and a clear relationship between the  $\pi$ -electron donating ability of the substituent, the decrease in the electrophilicity of the silylene, and the emergence of its nucleophilic character was demonstrated. Here in this study (see data in Table 1) we did not find such a straightforward relationship. The most important aspect from the point of view of our target molecule oxazol-2-ylidine 9 is, however, that its electrophilicity is only slightly increased compared to imidazol-2-vlidene 1, while its nucleophilicity<sup>59</sup> (and also its basicity)<sup>19,26b</sup> is much smaller, similar to that of the also widely applied organocatalysts thiazol-2-ylidene 6 which has significantly higher electrophilicity compared to both 1 and 9. Thus one may expect that oxazol-2-ylidene will show different reactivity, and organocatalytic reactivity than imidazol-2-ylidine and thiazol-2-ylidene. Selenazol-2-ylidenes (10), however, should have similar electronic properties to their sulfur analogues 6.

#### Conclusions

Combining the stabilizing effect of the (reasonably large) substituents (R) on the nitrogen and annellated rings on the carbon atoms, the most promising synthetic targets **11** and **12** are shown in Scheme 6. These compounds should be stable in dimerization and in cycloreversion reactions, provided that the water content of the reaction mixture is significantly reduced. The electrophilicity index of **9** indicates a comparable reactivity to the known imidazol-2-ylidene (**1**), while its lower nucleophilicity and basicity will result in changes in complex formation, and organocatalytic activities, opening new perspectives in the fine-tuning of catalytic reactions.

#### Acknowledgements

Financial support from New Szechenyi Plan TAMOP-4.2.2/B-10/1-2010-0009 and OTKA K 170356 is gratefully acknowledged. J. O. acknowledges receipt of an EU Marie Curie Fellowship (Project "Oestrometab") and O. H. the Alexander von Humboldt fellowship. Support from Sumitomo LTD for Zs. K. is also gratefully acknowledged.

#### Notes and references

 (a) D. Bourissou, O. Guerret, F. P. Gabbai and G. Bertrand, *Chem. Rev.*, 2000, **100**, 39–92; (b) D. Enders, O. Niemeier and A. Henseler, *Chem. Rev.*, 2007, **107**, 5606–5655; (c) F. E. Hahn and M. C. Jahnke, *Angew. Chem., Int. Ed.*, 2008, **47**, 3122–3172; (d) A. J. Arduengo III and G. Bertrand, *Chem. Rev.*, 2009, **109**, 3209; (e) J. Vignolle, X. Cattoen and D. Bourissou, *Chem. Rev.*, 2009, **109**, 3333; (f) M. Melaimi, M. Soleilhavoup and G. Bertrand, *Angew. Chem., Int. Ed.*, 2010, **49**, 8810–8849; (g) T. Droge and F. Glorius, *Angew.* *Chem., Int. Ed.*, 2010, **49**, 6940–6952; (*h*) D. Martin, M. Melaimi, M. Soleilhavoup and G. Bertrand, *Organometallics*, 2011, **30**, 5304–5313.

- 2 (a) D. R. Anderson, V. Lavallo, D. J. O'Leary, G. Bertrand and R. H. Grubbs, Angew. Chem., Int. Ed., 2007, 46, 7262-7265; (b) R. H. Crabtree, J. Organomet. Chem., 2005, 690, 5451-5457; (c) M. C. Perry and K. Burgess, Tetrahedron: Asymmetry, 2003, 14, 951-961; (d) S. Leuthaöusser, D. Schwarz and H. Plenio, Chem.-Eur. J., 2007, 13, 7195-7203; (e) S. Diez-Gonzalez, N. Marion and S. P. Nolan, Chem. Rev., 2009, 109, 3612-3676.
- 3 D. Martin, M. Soleilhavoup and G. Bertrand, *Chem. Sci.*, 2011, 2, 389–399.
- 4 (a) D. Enders and T. Balensiefer, Acc. Chem. Res., 2004, 37, 534–541; (b) N. Marion, S. Díez- González and S. P. Nolan, Angew. Chem., Int. Ed., 2007, 46, 2988; (c) P.-C. Chiang and J. W. Bode, N-Heterocyclic Carbene Catalyzed Reactions of alpha-Functionalized Aldehydes, Science of Synthesis: Asymmetric Organocatalysis, vol. 1, 2012, Georg Thieme Verlag, pp. 639–672.
- 5 A. J. Arduengo, R. L. Harlow and M. Kline, *J. Am. Chem. Soc.*, 1991, **113**, 361–363.
- 6 (a) O. Hollóczki and L. Nyulászi, J. Org. Chem., 2008, 73, 4794–4799; (b) C. Boehme and G. Frenking, J. Am. Chem. Soc., 1996, 118, 2039–2046; (c) C. Heinemann, T. Müller, Y. Apeloig and H. Schwartz, J. Am. Chem. Soc., 1996, 118, 2023–2038; (d) M.-J. Cheng and C.-H. Hu, Chem. Phys. Lett., 2001, 349, 477–482; (e) S. P. Nolan, Coord. Chem. Rev., 2007, 251, 874–883; (f) A. K. Guha, S. Sarmah and A. K. Phukan, Dalton Trans., 2010, 39, 7374–7383; (g) S. Gronert, J. R. Keeffe and R. A. More O'Ferrall, J. Am. Chem. Soc., 2011, 133, 3381–3389; (h) S. Gronert, J. R. Keeffe and R. A. More O'Ferrall, J. Am. Chem. Soc., 2011, 133, 11817–11818.
- 7 The hydrolysis of imidazol-2-ylidene was studied by O. Hollóczki, P. Terleczky, D. Szieberth, G. Mourgas, D. Gudat and L. Nyulászi, *J. Am. Chem. Soc.*, 2011, 133, 780–789. Much higher activation barriers were found than for the saturated counterpart imidazolidine-2-ylidene.
- 8 B. Maji, M. Horn and H. Mayr, *Angew. Chem., Int. Ed.*, 2012, 51, 6231–6235.
- 9 A. J. Arduengo, J. R. Goerlich and W. Marshall, *J. Am. Chem. Soc.*, 1995, **117**, 11027.
- D. Enders, K. Breuer, G. Raabe, J. Runsink, J. H. Teles, J. P. Melder, K. Ebel and S. Brode, *Angew. Chem., Int. Ed. Engl.*, 1995, 34, 1021–1023.
- 11 P. R. Allen, M. Murray and G. Orpen, *Angew. Chem., Int. Ed.* Engl., 1996, **35**, 1121–1123.
- 12 (a) A. Fekete and L. Nyulászi, J. Organomet. Chem., 2002, 643-644, 278-284; (b) L. Nyulászi, Tetrahedron, 2000, 56, 79-84.
- 13 D. Martin, A. Baceiredo, H. Gornitzka, W. W. Schoeller and G. Bertrand, *Angew. Chem., Int. Ed.*, 2005, 44, 1700–1703.
- 14 L. Nyulászi, Chem. Rev., 2001, 101, 1229-1246.
- 15 A. J. Arduengo, J. R. Goerlich and W. Marshall, *Liebigs Ann.*/ *Recl.*, 1997, 365–369.
- 16 (a) R. W. Alder, C. P. Butts and A. G. Orpen, J. Am. Chem. Soc., 1998, 120, 11526–11527; (b) N. Merceron-Saffon, A. Baceiredo, H. Gornitzka and G. Bertrand, Science, 2003, 301, 1223.

- 17 V. Lavallo, Y. Canac, C. Prasang, B. Donnadieu and G. Bertrand, *Angew. Chem., Int. Ed.*, 2005, 44, 5705–5709.
- 18 L. Nyulászi, T. Veszprémi and A. Forró, *Phys. Chem. Chem. Phys.*, 2000, **2**, 3127–3129.
- 19 O. Hollóczki, Zs. Kelemen and L. Nyulászi, *J. Org. Chem.*, 2012, 77, 6014–6022.
- 20 F. Freeman, D. J. Lau, A. R. Patel, P. R. Pavia and J. D. Willey, *J. Phys. Chem. A*, 2008, 12, 8775–8784.
- 21 M.-J. Cheng and C.-H. Hu, Chem. Phys. Lett., 2000, 322, 83-90.
- 22 The synthesis of a stable isothiazol carbene has been claimed (J. Wolf, W. Bohlmann, M. Findeisen, T. Gelbrich, H.-J. Hofmann and B. Schulze, *Angew. Chem., Int. Ed. Engl.*, 2007, 46, 3118–21), but was subsequently shown to be a rearranged product (A. DeHope, V. Lavallo, B. Donnadieu, W. W. Schoeller and G. Bertrand, *Angew. Chem., Int. Ed. Engl.*, 2007, 46, 6922–6925), formed from the intermediate isothiazol carbene (J. Wolf, W. Bohlmann, M. Findeisen, T. Gelbrich, H.-J. Hofmann and B. Schulze, *Angew. Chem., Int. Ed. Engl.*, 2007, 46, 6926.
- 23 (a) K. R. Rundy and W. R. Roper, J. Organomet. Chem., 1975,
  91, C61–C64; (b) A. C. Tubaro, A. Biffs, M. Basato,
  F. Benetollo, K. J. Cavell and L. Ooi, Organometallics,
  2005, 24, 4153–4158; (c) J. A. Cabeza, I. del Río, D. Miguel,
  E. Pérez-Carreño and M. G. Sánchez-Vega, Organometallics,
  2008, 27, 211–217; (d) J. Zhang, J. Fu, X. Su, X. Wang,
  S. Song and M. Shi, Chem.-Asian J., 2013, 3, 552–555.
- 24 (a) C. Lambert, I. Lopez-Solera and P. R. Raithby, Organometallics, 1996, 15, 452–455; (b) F. E. Hahn, T. Lugger and M. Z. Beinhoff, Z. Naturforsch., 2004, 59b, 196–201.
- 25 R. Hoffmann, P. V. R. Schleyer and H. F. Schaefer III, *Angew. Chem., Int. Ed.*, 2008, 47, 7164–7167.
- 26 (a) R. R. Sauers, *Tetrahedron Lett.*, 1994, 35, 7213–7216; (b)
  R. R. Sauers, *Tetrahedron Lett.*, 1996, 37, 149–152.
- 27 (a) A. Krebs and H. Kimling, Angew. Chem., Int. Ed. Engl., 1971, 10, 509-510; (b) H. D. Hartzler, J. Am. Chem. Soc., 1973, 95, 4379-4387; (c) C. U. Pittman and M. Narita, Bull. Chem. Soc. Jpn., 1976, 49, 1996-1999 (the adduct of trialkylphosphine and carbon disulfide was reacted with electon deficient acetylene).
- 28 S. E. d'Arbeloff-Wilson, P. B. Hitchcock, S. Krill, J. F. Nixon, L. Nyulászi and M. Regitz, J. Am. Chem. Soc., 2000, 122, 4557–4562.
- 29 J. Fabian, A. Krebs, D. Schoönemann and W. Schaefer, J. Org. Chem., 2000, 65, 8940–8947.
- 30 (a) T. W. Hudnall and C. W. Bielawski, J. Am. Chem. Soc., 2009, 131, 16039–16041; (b) G. A. Blake, J. P. Moerdyk and C. W. Bielawski, Organometallics, 2012, 31, 3373–3378.
- 31 J. Mahatthananchai and J. W. Bode, *Chem. Sci.*, 2012, 3, 192–197.
- 32 M. J. Frisch, G. W. Trucks, H. B. Schlegel, G. E. Scuseria, M. A. Robb, J. R. Cheeseman, J. A. Montgomery, Jr., T. Vreven, K. N. Kudin, J. C. Burant, J. M. Millam, S. S. Iyengar, J. Tomasi, V. Barone, B. Mennucci, M. Cossi, G. Scalmani, N. Rega, G. A. Petersson, H. Nakatsuji, M. Hada, M. Ehara, K. Toyota, R. Fukuda, J. Hasegawa, M. Ishida, T. Nakajima, Y. Honda, O. Kitao, H. Nakai, M. Klene, X. Li, J. E. Knox, H. P. Hratchian, J. B. Cross, V. Bakken, C. Adamo, J. Jaramillo, R. Gomperts, R. E. Stratmann, O. Yazyev, A. J. Austin, R. Cammi, C. Pomelli, J. Ochterski, P. Y. Ayala,

K. Morokuma, G. A. Voth, P. Salvador, J. J. Dannenberg,
V. G. Zakrzewski, S. Dapprich, A. D. Daniels, M. C. Strain,
O. Farkas, D. K. Malick, A. D. Rabuck, K. Raghavachari, J.
B. Foresman, J. V. Ortiz, Q. Cui, A. G. Baboul, S. Clifford,
J. Cioslowski, B. B. Stefanov, G. Liu, A. Liashenko,
P. Piskorz, I. Komaromi, R. L. Martin, D. J. Fox, T. Keith,
M. A. Al-Laham, C. Y. Peng, A. Nanayakkara,
M. Challacombe, P. M. W. Gill, B. G. Johnson, W. Chen,
M. W. Wong, C. Gonzalez and J. A. Pople, *GAUSSIAN 03* (*Revision C.02*), Gaussian, Inc., Wallingford, CT, 2004.

- 33 (a) NICS(0) is the magnetic shielding in the geometrical center of the ring. P. V. R. Schleyer, C. Maerker, A. Dransfeld, H. Jiao and N. J. R. van. Eikema Hommes, J. Am. Chem. Soc., 1996, 118, 6317–6318; (b) NICS(1) is the magnetic shielding 1 Å above the geometrical center of the ring. P. V. R. Schleyer, H. Jiao, N. J. R. van Eikema Hommes, V. G. Malkin and O. L. Malkina, J. Am. Chem. Soc., 1997, 119, 12669–12670.
- 34 C. W. Bird, Tetrahedron, 1985, 41, 1409.
- 35 W. Gordy, J. Chem. Phys., 1947, 15, 305-310.
- 36 (a) P. V. R. Schleyer and F. Pühlhofer, Org. Lett., 2002, 4, 2873–2876 introduced ISE for six membered rings; (b) For five membered rings: L. Nyulászi, O. Hollóczki, C. Lescop, M. Hissler and R. Réau, Org. Biomol. Chem., 2006, 4, 996–998; (c) O. Hollóczki and L. Nyulászi, Struct. Chem., 2011, 22, 1385–1392. ISE was calculated for some NCH-s in ref. 6a.
- 37 P. Geerlings, F. De Proft and W. Langenaeker, *Chem. Rev.*, 2003, **103**, 1793–1874.
- 38 P. Pérez, J. Phys. Chem. A, 2003, 107, 522-525.
- 39 (a) R. A. Moss, Acc. Chem. Res., 1989, 22, 15–21; (b) R.
   A. Moss, Acc. Chem. Res., 1980, 13, 58–64.
- 40 J. Oláh, F. D. Proft, T. Veszprémi and P. Geerlings, *J. Phys. Chem. A*, 2005, **109**, 1608–1615.
- 41 (a) R. G. Parr and R. G. Pearson, J. Am. Chem. Soc., 1983, 105, 7512; (b) R. G. Parr, W. Yang, Density Functional Theory of Atoms and Molecules, Oxford University Press, New York, 1989.
- 42 T. Koopmans, Physica, 1934, 1, 104.
- 43 L. R. Domingo, E. Chamorro and P. J. Perez, *J. Org. Chem.*, 2008, **73**, 4615–4624.
- 44 M. Z. Kassaee, F. A. Shakib, M. R. Momeni, M. Ghambarian and S. M. Musavi, *J. Org. Chem.*, 2010, **75**, 2539–2545.
- 45 With the exception of the NICS values of some sulfur containing carbenes.
- 46 Yu. B. Vysotsky, V. S. Bryantsev and O. A. Gorban, *Chem. Heterocycl. Compd.*, 2002, 38, 1451–1468.
- 47 The aromaticity of the five membered heterocycles has been compared many times. For a work using several aromaticity measures see *e.g.* L. Nyulászi and P. V. R. Schleyer, *J. Am. Chem. Soc.*, 1999, 121, 6872–75. For evaluation of the ISE see ref. 36c.
- 48 L. Nyulászi, G. Keglevich and L. D. Quin, *J. Org. Chem.*, 1996, **61**, 7808–7812.
- 49 P. Couture, J. K. Terlouw and J. Warkentin, *J. Am. Chem. Soc.*, 1996, **118**, 4214–4215.
- 50 This difference is likely to be attributable to the different stability of the reaction product isocyanate analogues.
- 51 A. Krebs and J. Wilke, *Top. Curr. Chem.*, 1983, 190–233.
- 52 R. Breslow, J. Am. Chem. Soc., 1958, 40, 3719-3726.

Paper

- 53 P. Haake and W. B. Miller, J. Am. Chem. Soc., 1963, 85, 4044–4045.
- 54 J. M. Duclos and P. Haake, *Biochemistry*, 1974, 13, 5358–5362.
- 55 It is known that clustering of water is significantly reduced in ionic liquids. See *e.g.* C. G. Hanke and R. M. Lynden-Bell, *J. Phys. Chem. B*, 2003, **107**, 10873–10878.
- 56 B. Maji, M. Breugst and H. Mayr, *Angew. Chem., Int. Ed.*, 2011, **50**, 6915–6919.
- 57 R. A. Kelly III, H. Clavier, S. Giudice, N. M. Scott, E. D. Stevens, J. Bordner, I. Samardjiev, C. D. Hoff, L. Cavallo and S. P. Nolan, *Organometallics*, 2008, 27, 202–210.
- 58 (a) J. Oláh, F. De Proft, T. Veszpremi and P. Geerlings, J. Phys. Chem. A, 2007, 111, 10815–10823; (b) J. V. Correa, P. Jaque, J. Oláh, A. Toro-Labbe and P. Geerlings, Chem. Phys. Lett., 2009, 470, 180–186.
- 59 In a study of the related thiazol-2-ylidene the nucleophilic character of oxazol-2-ylidene was also considered. (*a*) D.
  B. DuPre and J. L. Wong, *J. Phys. Chem. A*, 2005, 109, 7606–7612; (*b*) D. B. DuPre and J. L. Wong, *J. Phys. Chem. A*, 2007, 111, 2172–2181.

Paper