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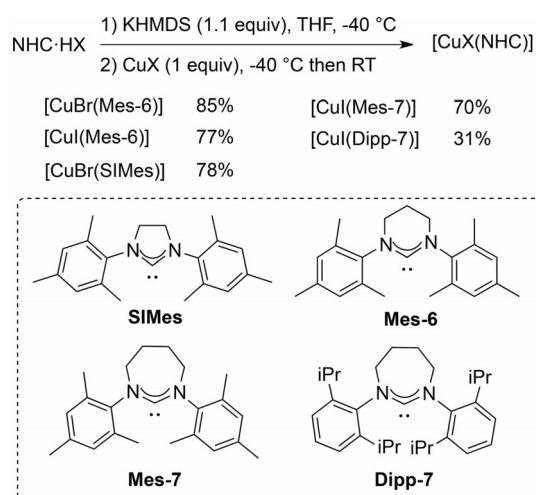
strongly favoured. Unfortunately, the scope and efficiency of [Cu(CAAC)] complexes have not been explored yet. Similarly, to the best of our knowledge, to date there is only one report on the activity of ring-expanded NHC ($^{\text{RE}}\text{NHC}$) ligands to this cycloaddition reaction.^[17] [Cu(*o*-Tol-7)₂]BF₄ was found to be the best catalyst of the series in the absence of solvent (0.5 mol% [Cu]). However, only 2 out of 10 cycloadditions gave conversions above 10% under the reported conditions.^[18]

Still, $^{\text{RE}}\text{NHC}$ s are known to improve the catalytic efficiency in important transformations such as cross-coupling,^[19] or allylic boronation^[20] reactions, as well as allowing the study of elusive species (that is, three-coordinated nickel(I) complexes,^[21] or copper(I)–hydrides^[22]). Herein we report the remarkable catalytic activity of several [CuX($^{\text{RE}}\text{NHC}$)] complexes in the formation of triazoles under click-suitable conditions.

Results and Discussion

Catalyst preparation and characterisation

The novel copper catalysts were prepared by the addition of the appropriate CuX salt to the NHC ligand formed in situ by prior deprotonation of the required NHC-HX in THF solution (Scheme 1). Previous reports relied on transmetallation reac-



Scheme 1. Synthesis of [CuX(NHC)] complexes.

tions from the corresponding silver complexes to prepare related complexes,^[23] but the approach adopted herein has the advantage of yielding the desired complexes in a single step and avoiding the formation of stoichiometric amounts of silver waste. For comparison purposes, benchmark [CuBr(SIMes)]^[12] was also prepared in a similar manner.

The desired complexes were isolated in good yields as cream coloured solids after work-up and characterised by NMR, HRMS, and elemental analysis. All spectroscopic data showed that the prepared complexes are monomeric in solution at detectable levels, with the notable exception of [CuBr(Mes-6)]. In this particular case, the isolated product contained 20% of [Cu(Mes-6)₂]⁺ in solution, according to the

¹H NMR spectrum.^[24] The counterion for this complex would presumably be CuBr₂[−].^[25,26] Such a ligand redistribution process led to the formation of an homoleptic complex as the only reaction product in a similar synthesis of a chloro derivative,^[25] but it could not be detected for the related iodo analogue. The reasons for this are not clear, but we had already observed improved bridging abilities for copper(I)–bromo complexes compared to iodo ones with triphosphorous macrocycles.^[27] It is conceivable that $\mu\text{-Br}$ species might facilitate the formation of bis-NHC complexes in this context.

Catalytic studies

With a series of [CuX(NHC)] complexes in hand, we next tested them on a model reaction under identical reaction conditions. A first set of experiments for the cyclisation of benzyl azide and phenylacetylene with 0.5 mol% [Cu] led to complete conversion with several of the tested catalysts, and in consequence, the metal loading was then lowered to 0.05 mol% to differentiate them (Table 1).

Table 1. Catalyst screening.

$\text{Ph-CH}_2\text{-N}_3 + \text{Ph-C}\equiv\text{C-H} \xrightarrow[\text{neat, RT}]{\text{Cu}^{\text{I}} \text{ catalyst}} \text{Ph-CH}_2\text{-N=N-C(Ph)=N-Ph} \quad \mathbf{1a}$				
Entry	Catalyst	[Cu] [mol%]	<i>t</i> [h]	Conv [%] ^[a]
1	[CuBr(SIMes)]	0.5	1	> 95
		0.05	2	8
		0.05	24	9
2	[CuBr(Mes-6)]	0.5	1	> 95
		0.05	2	8
		0.05	4	17
		0.05	8	36
		0.05	24	> 95
3	[CuI(Mes-6)]	0.5	1	> 95
		0.05	2	> 95
4	[CuI(Mes-7)]	0.5	1	< 5
		0.5	4	36
		0.5	24	> 95
5	[CuI(Dipp-7)]	1.0	24	NR

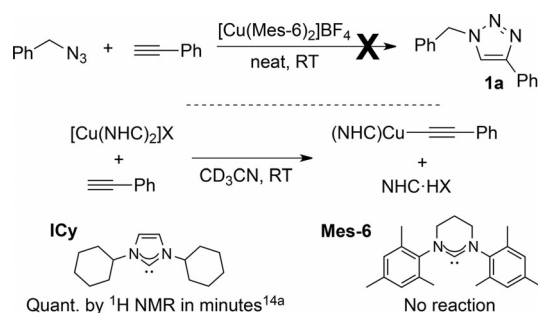
[a] ¹H NMR conversions are the average of at least two independent experiments. NR = No reaction.

Overall, [CuI(Mes-6)] displayed the best catalytic performance and **1a** was formed quantitatively in 2 h with only 0.05 mol% metal loading (Table 1, entry 3). In these reactions, it was clearly observed that with [CuX(NHC)] complexes, the activity could be ordered as NHC-6 > NHC-5 ≫ NHC-7. It is important to note that no catalyst decomposition was observed in these reactions, particularly with [CuI(Dipp-7)]. The disappointing catalytic activity of copper complexes bearing seven-membered NHC ligands is uncharacteristic of this family of ligands. Indeed, this trend is the reverse of that observed in

other catalytic reactions such as palladium-mediated C–C cross-couplings.^[19b,28] This might be due to the greater steric hindrance of NHC-7 ligands when compared to their smaller ring analogues,^[29] but general correlations in catalysis are hard to define owing to the complexity of the mechanisms involved and insufficient comparative data reported to date.

Results in Table 1 confirmed the previously established trend where $[\text{CuI}(\text{NHC})] > [\text{CuBr}(\text{NHC})] > [\text{CuCl}(\text{NHC})]$.^[30] Nevertheless, in this particular case, not only the halogen, but also the specification in solution needs to be taken into account. Hence, it appears that the specific behaviour of $[\text{CuBr}(\text{Mes-6})]$ in solution is linked to its diminished catalytic activity and therefore, $[\text{CuX}(\text{R}^{\text{E}}\text{NHC})]$ are privileged complexes for this cycloaddition reaction when compared to $[\text{Cu}(\text{R}^{\text{E}}\text{NHC})_2]\text{X}$ complexes. As previously mentioned,^[17] related $[\text{Cu}(\text{o-Tol-7})_2]\text{BF}_4$ only displayed poor catalytic activity in azide–alkyne cycloadditions and we hypothesised this might be due to an inefficient activation of the latter under the reaction conditions.

To be able to draw a direct comparison, $[\text{Cu}(\text{Mes-6})_2]\text{BF}_4$ was prepared following a similar procedure to that shown in Scheme 1 and it was then used as catalyst in our model reaction but no triazole formation was observed with either 0.5 or 1 mol% of the homoleptic complex after 24 h (Scheme 2). As



Scheme 2. Catalytic tests and proposed activation step for $[\text{Cu}(\text{NHC})_2]^+$ complexes.

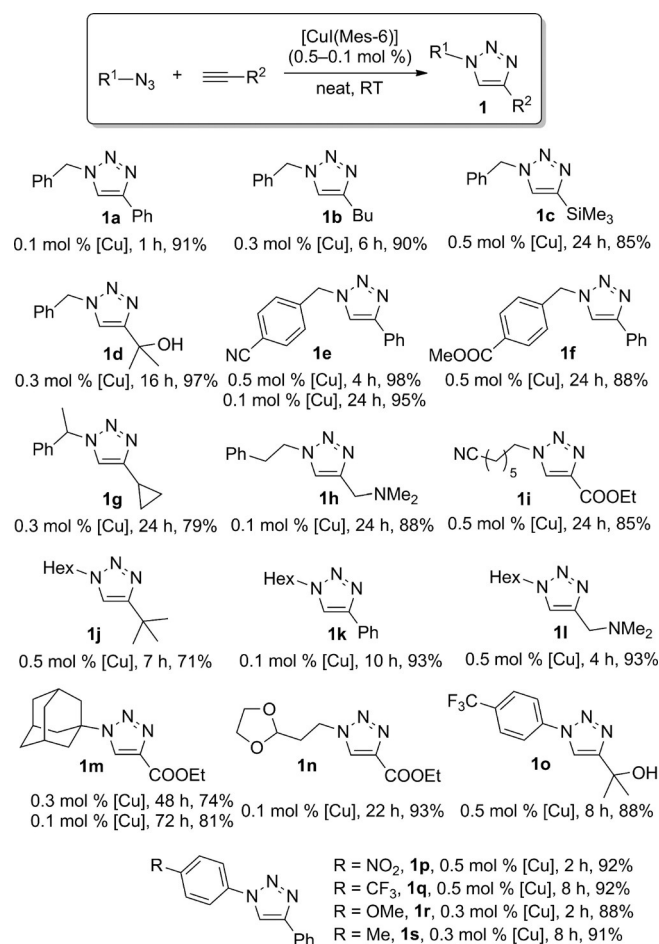
we had previously proposed that the first step in the catalytic cycle with homoleptic imidazol-2-ylidene complexes involves the displacement of one of the NHC ligand by the alkyne, with the generation of a copper acetylide intermediate,^[14a] we then carried out a similar reaction with $[\text{Cu}(\text{Mes-6})_2]\text{BF}_4$, and again no reaction was observed in CD_3CN , even in the presence of an excess of alkyne (Scheme 2).^[24]

While these experiments explain the low catalytic activity of homoleptic copper complexes bearing ring-expanded NHCs (as well as heteroleptic ones that might rearrange into homoleptic complexes, such as $[\text{CuBr}(\text{Mes-6})]$), the reasons behind this lack of reactivity are not obvious owing to the lack of data. While tetrahydropyrimidinyl-2-ylidenes have been found to be more basic than analogous imidazol- or imidazolin-2-ylidenes,^[31] and are better electron donor ligands,^[32] little is known about relative dissociation energies and strength of the newly formed NHC–H bonds.

In the light of these results, we also reassessed the activity of $[\text{CuBr}(\text{Mes-6})]$ in azide–alkyne cycloadditions. The model re-

action was carried out with 0.05 mol% of $[\text{CuBr}(\text{Mes-6})]$. In this case, the metal loading was calculated taking into account the presence of inactive $[\text{Cu}(\text{Mes-6})_2][\text{CuBr}_2]$ and it was assumed that the copper complexes did not interconvert under catalytic conditions. Higher conversions into triazole **1a** were then obtained, as expected.^[33] However, this complex still failed to match the performance of the iodo analogue.

Having established $[\text{CuI}(\text{Mes-6})]$ as the catalyst of choice, we next explored the scope of the reaction (Scheme 3). Isolated

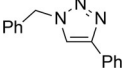
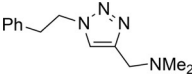
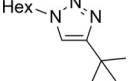
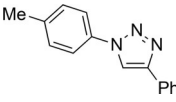


Scheme 3. $[\text{CuI}(\text{Mes-6})]$ -catalyzed azide–alkyne cycloaddition reaction.

yields in these reactions ranged from 71 to 98% with most triazoles obtained in 85% yield or higher. Benzyl, alkyl, and aryl azides were successfully employed under the optimised conditions. Also, a variety of functional groups (such as alcohol, nitrile, amine, silyl, nitro) were tolerated by the catalytic system. Sterically hindered substrates could also be employed, even if a substantially lower reaction rate was observed when adamantyl azide was used as a cycloaddition partner (**1m** in Scheme 3). Nevertheless, these reactions also show how robust the complex is, since $[\text{CuI}(\text{Mes-6})]$ remained an active catalyst even after three days.

This observation led us to test some of the reactions shown in Scheme 3 with lower metal loadings. The obtained results are summarised in Table 2. Gratifyingly, all four azides tested

Table 2. Low catalytic loading experiments.

$\text{R}^1\text{-N}_3 + \text{R}^2\text{-C}\equiv\text{C} \xrightarrow[\text{neat, RT}]{[\text{CuI}(\text{Mes-6})] \text{ (100–500 ppm)}} \text{R}^1\text{-N}=\text{N}-\text{C}(\text{R}^2)=\text{N}$					
Triazole	1	[Cu] [ppm]	t [h]	Conv [%] ^[a]	TON
	1a	100	24	> 95	10000
	1h	500	24 48 72	57 64 71	1420
	1j	500	24 48	50 61	1220
	1s	500	24 48 72	30 60 95	1900

[a] ¹H NMR conversions are the average of at least two independent experiments.

led to the corresponding triazoles in good to excellent conversions with only 500 ppm of catalyst (100 ppm in the case of **1a**) and a TON ranging from 1220 and 10⁴. These experiments were not limited to activated substrates and triazoles with bulky substituents (that is, **1j**) or bearing an additional functional group (for example, amino group on **1h**) could be prepared under these conditions. In some reactions, though, the catalytic activity eventually seemed to level off, which might be due to the presence of a coordinating group in the substrates (**1h**) or a significant steric hindrance (**1j**).

Conclusions

The screening of different [CuX(NHC)] on a model azide–alkyne cycloaddition reaction has clearly established six-membered NHC ligands as promising scaffolds for the preparation of 1,4-disubstituted triazoles under click-suitable reaction conditions.

These seem to have an optimal stereoelectronic profile for this transformation and a range of triazoles could be efficiently prepared with metal loadings between 0.5 mol% and 100 ppm. We have also shown that the very low catalytic activity of related [Cu^{RE}(NHC)₂]X complexes is most probably due to an inefficient activation step. Further catalytic applications of copper(I) complexes bearing ring-expanded NHC ligands are currently ongoing in our laboratories and will be reported in due course.

Experimental Section

Catalytic reactions were carried out in air and using technical-grade solvents without any particular precautions to exclude moisture or oxygen. The reported isolated yields for the catalytic studies are the average of two independent reactions.

CAUTION: Although we did not experience any problems, the cycloaddition of azides and alkynes is highly exothermic and, as a consequence, adequate cooling should always be available when performing these reactions in the absence of solvent.

Model procedure for click cycloaddition reactions: In a vial fitted with a screw cap, [CuI(Mes-6)] (0.5 mol%–100 ppm), azide (0.5 mmol), and alkyne (0.5 mmol) were loaded. The reaction was allowed to proceed at room temperature until full (or no further) conversion was observed by ¹H NMR spectroscopy. Then, saturated aqueous ammonium chloride solution (10 mL) was added and the resulting mixture was stirred vigorously for 3 h. The resulting precipitate was filtered and washed with water and pentane and then dried under reduced pressure. In all examples, the crude products were estimated to be > 95 % pure by ¹H NMR.

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Conflict of interest

The authors declare no conflict of interest.

Keywords: alkynes • azides • click chemistry • copper • N-heterocyclic carbenes

- [1] H. C. Kolb, M. G. Finn, K. B. Sharpless, *Angew. Chem. Int. Ed.* **2001**, *40*, 2004–2021; *Angew. Chem.* **2001**, *113*, 2056–2075.
- [2] G. L'Abbé, *Bull. Soc. Chim. Belg.* **1984**, *93*, 579–592.
- [3] V. V. Rostovtsev, L. G. Green, V. V. Fokin, K. B. Sharpless, *Angew. Chem. Int. Ed.* **2002**, *41*, 2596–2599; *Angew. Chem.* **2002**, *114*, 2708–2711.
- [4] C. W. Tornøe, C. Christensen, M. Meldal, *J. Org. Chem.* **2002**, *67*, 3057–3064.
- [5] a) Special issue on click chemistry, *Chem. Soc. Rev.* **2010**, *39*, 1221–1408; b) M. Meldal, C. W. Tornøe, *Chem. Rev.* **2008**, *108*, 2952–3015.
- [6] a) E. Haldón, M. C. Nicasio, P. J. Pérez, *Org. Biomol. Chem.* **2015**, *13*, 9528–9550; b) S. Díez-González, *Catal. Sci. Technol.* **2011**, *1*, 166–178.
- [7] T. R. Chan, R. Hilgraf, K. B. Sharpless, V. V. Fokin, *Org. Lett.* **2004**, *6*, 2853–2855.
- [8] a) V. O. Rodionov, S. I. Presolski, S. Gardinier, Y. H. Lim, M. G. Finn, *J. Am. Chem. Soc.* **2007**, *129*, 12696–12704; b) V. O. Rodionov, S. I. Presolski, D. D. Diaz, V. V. Fokin, M. G. Finn, *J. Am. Chem. Soc.* **2007**, *129*, 12705–12712; c) V. Bevilacqua, M. King, M. Chaumonnet, M. Nothisen, S. Gabillet, D. Buisson, C. Puente, A. Wagner, F. Taran, *Angew. Chem. Int. Ed.* **2014**, *53*, 5872–5876; *Angew. Chem.* **2014**, *126*, 5982–5986.
- [9] I. Kalvet, J. Tammiku-Taul, U. Mäeorg, K. Tamm, P. Burk, L. Sikk, *ChemCatChem* **2016**, *8*, 1804–1808.
- [10] a) *N-Heterocyclic Carbenes: From Laboratory Curiosities to Efficient Synthetic Tools*, 2nd ed. (Ed.: S. Díez-González), RSC, Cambridge, **2017**; b) S. Díez-González, N. Marion, S. P. Nolan, *Chem. Rev.* **2009**, *109*, 3612–3676; c) *N-Heterocyclic Carbenes in Synthesis* (Ed.: S. P. Nolan), Wiley-VCH, Weinheim, **2006**.
- [11] a) C. Nolte, P. Mayer, B. F. Straub, *Angew. Chem. Int. Ed.* **2007**, *46*, 2101–2103; *Angew. Chem.* **2007**, *119*, 2147–2149; b) B. T. Worrell, J. A. Malik, V. V. Fokin, *Science* **2013**, *340*, 457–460; c) A. Makarem, R. Berg, F. Rominger, B. F. Straub, *Angew. Chem. Int. Ed.* **2015**, *54*, 7431–7435; *Angew. Chem.* **2015**, *127*, 7539–7543.
- [12] S. Díez-González, A. Correa, L. Cavallo, S. P. Nolan, *Chem. Eur. J.* **2006**, *12*, 7558–7564.
- [13] a) W. Wang, J. Wu, C. Xia, F. Li, *Green Chem.* **2011**, *13*, 3440–3445; b) J.-M. Collinson, J. D. E. T. Wilton-Ely, S. Díez-González, *Chem. Commun.* **2013**, *49*, 11358–11360.

- [14] For selected examples, see: a) S. Díez-González, S. P. Nolan, *Angew. Chem. Int. Ed.* **2008**, *47*, 8881–8884; *Angew. Chem.* **2008**, *120*, 9013–9016; b) R. Berg, J. Straub, E. Schreiner, S. Mader, F. Rominger, B. F. Straub, *Adv. Synth. Catal.* **2012**, *354*, 3445–3450; c) B. Liu, C. Chen, Y. Zhang, X. Liu, W. Chen, *Organometallics* **2013**, *32*, 5451–5460; d) H. Ibrahim, R. Guillot, F. Cisnetti, A. Gautier, *Chem. Commun.* **2014**, *50*, 7154–7156.
- [15] a) T. Nakamura, T. Terashima, K. Ogata, S.-i. Fukuzawa, *Org. Lett.* **2011**, *13*, 620–623; b) S. Hohloch, C.-Y. Su, B. Sarkar, *Eur. J. Inorg. Chem.* **2011**, 3067–3075; c) S. C. Sau, S. R. Toy, T. K. Sen, D. Mullangi, S. K. Mandal, *Adv. Synth. Catal.* **2013**, *355*, 2982–2991; d) S. Hohloch, D. Scheiffele, B. Sarkar, *Eur. J. Inorg. Chem.* **2013**, 3956–3965; e) S. Hohloch, B. Sarkar, L. Nauton, F. Cisnetti, A. Gautier, *Tetrahedron Lett.* **2013**, *54*, 1808–1812; f) Y. D. Vidal, M. Lesieur, M. Melaimi, F. Nahra, D. B. Cordes, K. S. A. Arachchige, A. M. Z. Slawin, G. Bertrand, C. S. J. Cazin, *Adv. Synth. Catal.* **2015**, *357*, 3155–3161; g) S. Hohloch, L. Suntrup, B. Sarkar, *Inorg. Chem. Front.* **2016**, *3*, 67–77.
- [16] a) L. Jin, D. R. Tolentino, M. Melaimi, G. Bertrand, *Sci. Adv.* **2015**, *1*, e1500304; b) L. Jin, E. A. Romero, M. Melaimi, G. Bertrand, *J. Am. Chem. Soc.* **2015**, *137*, 15696–15698.
- [17] L. R. Collins, T. M. Rookes, M. F. Mahon, I. M. Riddlestone, M. K. Whittlesey, *Organometallics* **2014**, *33*, 5882–5887.
- [18] During the preparation/revision of this manuscript a related article on eight-membered ring NHCs in this cycloaddition reaction was reported, see: G. A. Chesnokov, M. A. Topchiy, P. B. Dzhevakov, P. S. Gribanov, A. A. Tukov, V. N. Khrustalev, A. F. Asachenko, M. S. Nechaev, *Dalton Trans.* **2017**, *46*, 4331–4345.
- [19] a) T. Tu, J. Malineni, X. Bao, K. H. Dötz, *Adv. Synth. Catal.* **2009**, *351*, 1029–1034; b) J. J. Dunsford, K. J. Cavell, *Dalton Trans.* **2011**, *40*, 9131–9135.
- [20] a) J. K. Park, H. H. Lackey, B. A. Ondrusek, D. Tyler McQuade, *J. Am. Chem. Soc.* **2011**, *133*, 2410–2413; b) J. K. Park, D. T. McQuade, *Angew. Chem. Int. Ed.* **2012**, *51*, 2717–2721; *Angew. Chem.* **2012**, *124*, 2771–2775.
- [21] a) C. J. E. Davies, M. J. Page, C. E. Ellul, M. F. Mahon, M. K. Whittlesey, *Chem. Commun.* **2010**, *46*, 5151–5153; b) M. J. Page, W. Y. Lu, R. C. Poulten, E. Carter, A. G. Algarra, B. M. Kariuki, S. A. Macgregor, M. F. Mahon, K. J. Cavell, D. M. Murphy, M. K. Whittlesey, *Chem. Eur. J.* **2013**, *19*, 2158–2167.
- [22] a) L. R. Collins, I. M. Riddlestone, M. F. Mahon, M. K. Whittlesey, *Chem. Eur. J.* **2015**, *21*, 14075–14084; b) A. J. Jordan, C. M. Wyss, J. Bacsá, J. P. Sadighi, *Organometallics* **2016**, *35*, 613–616; c) L. R. Collins, N. A. Rajabi, S. A. Macgregor, M. F. Mahon, M. K. Whittlesey, *Angew. Chem. Int. Ed.* **2016**, *55*, 15539–15543; *Angew. Chem.* **2016**, *128*, 15768–15772.
- [23] a) E. L. Kolychev, I. A. Portnyagin, V. V. Shuntikov, V. N. Khrustalev, M. S. Nechaev, *J. Organomet. Chem.* **2009**, *694*, 2454–2462; b) O. S. Morozov, A. V. Lunchev, A. A. Bush, A. A. Tukov, A. F. Asachenko, V. N. Krustalev, S. S. Zaleskiy, V. P. Ananikov, M. S. Nechaev, *Chem. Eur. J.* **2014**, *20*, 6162–6170.
- [24] See the Supporting Information for full details.
- [25] B. Bantu, D. Wang, K. Wurst, M. R. Buchmeiser, *Tetrahedron* **2005**, *61*, 12145–12152.
- [26] G. M. Pawar, B. Bantu, J. Weckesser, S. Blechert, K. Wurst, M. R. Buchmeiser, *Dalton Trans.* **2009**, 9043–9051.
- [27] L. V. Wickramatunga, *Co-ordination & organometallic chemistry of facially capping tridentate triphosphorus macrocycles and the applications*, PhD Thesis, Cardiff University, **2014**.
- [28] J. J. Dunsford, K. J. Cavell, *Organometallics* **2014**, *33*, 2902–2905.
- [29] a) J. J. Dunsford, K. J. Cavell, B. M. Kariuki, *Organometallics* **2012**, *31*, 4118–4121; b) N. Phillips, T. Dodson, R. Tirfoin, J. I. Bates, S. Aldridge, *Chem. Eur. J.* **2014**, *20*, 16721–16731.
- [30] S. Díez-González, E. C. Escudero-Adán, J. Benet-Buchholtz, E. D. Stevens, A. M. Z. Slawin, S. P. Nolan, *Dalton Trans.* **2010**, *39*, 7595–7606.
- [31] a) A. M. Magill, K. J. Cavell, B. F. Yates, *J. Am. Chem. Soc.* **2004**, *126*, 8717–8724; b) E. M. Higgins, J. A. Sherwood, A. G. Lindsay, J. Armstrong, R. S. Massey, R. W. Alder, A. M. C. O'Donoghue, *Chem. Commun.* **2011**, *47*, 1559–1561.
- [32] D. J. Nelson, S. P. Nolan, *Chem. Soc. Rev.* **2013**, *42*, 6723–6753.
- [33] Conversions into triazole **1a**: 2 h, 26%; 4 h, 56%; 8 h, > 95%.

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