



porating (functionalized) substituents, since triazole precursors are conveniently prepared through Cu(I)-catalyzed azide-alkyne cycloaddition chemistry.<sup>33–38</sup> Here we report an efficient amine oxidation protocol based on a dual catalysis system consisting of triazolium cations and iodide as redox-active counteranion, with synergistic efficiencies that surpass by far the most active organic catalyst for this oxidative transformation,<sup>23</sup> even when carried out under air, and allowing for a shorter timescale for quantitative yields.

## Results and discussion

### Model reaction and controls

The catalytic oxidative dehydrogenation of benzyl amine was chosen as a model reaction for this study (Table 1). 1,2,3-Triazolium iodide salt **1**, with methyl and phenyl substituents, was first explored as a catalyst for this reaction, in refluxing 1,2-dichlorobenzene. These conditions have previously been used for this transformation when catalyzed by precious metal complexes.<sup>8–10</sup> With a loading of 5 mol% of **1**, selective formation of the imine product in quantitative spectroscopic yields was observed within 3 h (entry 1), with no other side-products detected by <sup>1</sup>H NMR analysis of the reaction mixture (Fig. S13†). No reaction was observed over the same timescale in the absence of triazolium salt (entry 2). Lowering the temperature to 115 °C or refluxing in lower-boiling solvents (toluene, 1,2-dichloroethane) resulted in no discernible reaction progress (Table S1†).

Decreasing catalyst loading to 2.5 mol% gave 94% yield after 4 h, with the rate of reaction approximately halved in line with the decrease in substrate/catalyst ratio. A low catalyst

loading of 0.5 mol% also gave quantitative yields, albeit after 6 h (entries 3 and 4; Fig. S1†). The robustness of the catalytic species derived from **1** was tested by performing the reaction with only 0.05 mol% triazolium salt. After 16 h a maximum yield of 36% was reached, indicating a catalyst turnover number of approximately 740 under these conditions.

No special precautions were taken to dry 1,2-dichlorobenzene for these reactions in order to achieve selective imine formation. Indeed, deliberate addition of 10 equivalents of H<sub>2</sub>O did not significantly slow the transformation (entry 5), although, notably, a drop in yield occurred towards the end of the reaction (at 4 h) and a corresponding amount of benzaldehyde was observed by <sup>1</sup>H NMR analysis, indicating that the presence of water leads to hydrolysis of the imine product. Increasing the water content tenfold completely inhibited catalysis (entry 6). Neither base nor acid additives were beneficial, as neither stoichiometric K<sup>t</sup>OBu nor TsOH accelerated conversion (entries 7 and 8). Notably, with either of the additives, quantitative yields of the imine product were still achieved after 4 h.

Different benzylamines with electronically active *para*-substituents were successfully oxidized with triazolium salt **1** (Table 2, entries 1–4). No specific trend correlating the Hammett parameter  $\sigma_p$  and catalytic activity was noted. Aliphatic *n*-butylamine was not oxidized to an imine under these conditions, demonstrating a rather limited substrate scope for dehydrogenation. Nonetheless the possibility to convert benzylic amines selectively offers attractive synthetic opportunities.

Catalysts for oxidative coupling of amines are often applied to alcohol oxidation,<sup>1,10,39</sup> however **1** was specific for amines as benzyl alcohol was not converted at all. Reaction of a 1 : 1 mixture of benzyl alcohol and benzylamine led to exclusive formation of the same imine product as is formed in the presence of benzylamine only (>98% yield after 4 h), while benzyl alcohol remained unreacted and was still present in the

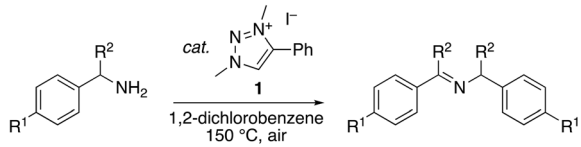
**Table 1** Optimization of reaction conditions for the triazolium-catalyzed homocoupling of benzylamine to benzylidene benzylimine<sup>a</sup>



Entry	Catalyst		Additive		Yield <sup>b</sup> (%)		
	mol%	mol%	mol%	mol%	1 h	3 h	4 h
1	<b>1</b>	5	—	—	60	100	99
2	—	0	—	—	0	<2	<2
3	<b>1</b>	2.5	—	—	33	79	94
4	<b>1</b>	0.5	—	—	24	60	75 <sup>c</sup>
5	<b>1</b>	5	H <sub>2</sub> O	1000	51	93	88 <sup>d</sup>
6	<b>1</b>	5	H <sub>2</sub> O	10 000	0	0	0
7	<b>1</b>	5	KO <sup>t</sup> Bu	5	47	93	98
8	<b>1</b>	5	TsOH	5	47	91	96

<sup>a</sup> General conditions: Benzyl amine (0.25 mmol), triazolium salt **1** (5 mol%) and hexamethylbenzene (internal standard), 1,2-dichlorobenzene (2 mL), 150 °C. <sup>b</sup> Spectroscopic yield determined by <sup>1</sup>H NMR analysis with respect to hexamethylbenzene. <sup>c</sup> 99% after 6 h. <sup>d</sup> 6% benzaldehyde formed in addition.

**Table 2** Reactivity of different amine substrates in the presence of catalyst **1**<sup>a</sup>



Entry	Substrate			Yield <sup>b</sup> (%)				TOF <sub>25</sub> (h <sup>-1</sup> )
	R <sup>1</sup>	R <sup>2</sup>	$\sigma_{para}$	1 h	3 h	4 h		
1	OMe	H	-0.27	32	97	92 <sup>c</sup>	6	
2	H	H	+0.00	60	100	99	16	
3	F	H	+0.06	19	71	91	4	
4	CF <sub>3</sub>	H	+0.54	65	98	100	21	
5	H	CH <sub>3</sub>	+0.00	26 <sup>d</sup>	35 <sup>d</sup>	33 <sup>d</sup>	5	

<sup>a</sup> General conditions: Benzyl amine (0.25 mmol), triazolium salt **1** (5 mol%), hexamethylbenzene (internal standard), 1,2-dichlorobenzene (2 mL), 150 °C. <sup>b</sup> Spectroscopic yield determined by <sup>1</sup>H NMR analysis with respect to hexamethylbenzene. <sup>c</sup> 8% aldehyde formed in addition. <sup>d</sup> Acetophenone formed in addition (9% after 1 h, 20% after 3 h, 26% after 4 h; total conversion after 4 h is 59%).



final reaction mixture after 6 h. Finally, branched primary amine  $\alpha$ -methylbenzylamine was also employed as a substrate for the homocoupling reaction (entry 5), but this transformation was less selective. While complete conversion occurred, only 33% yield of imine was seen within 4 h in addition to acetophenone (26%) and other unidentified products.

### Catalyst variation

In order to better understand this new catalytic system and identify its relevant components, a number of related species were tested for their catalytic activity. Variation of the wingtip groups of triazolium iodide catalyst **1** was investigated through use of salts **2–4** (Table 3, entries 2–4), which have been reported previously for applications including as ionic liquids<sup>32</sup> and precursors to 1,2,3-triazolyldene organometallic ligands.<sup>27–29</sup> All these triazolium iodide salts efficiently catalyze the oxidative homocoupling. Salts **2** and **4** displayed activities comparable to **1** ( $\text{TOF}_{50} = 9, 9$  and  $13 \text{ h}^{-1}$ , respectively) and give quantitative spectroscopic yields within 3 h, while **3** required 6 h to reach completion ( $\text{TOF}_{50} = 5 \text{ h}^{-1}$ ). Bearing in mind the similarities between imidazolium and triazolium salts, 1,3-dimethylimidazolium iodide **5** was also assessed as a catalyst (entry 5) and did demonstrate activity. However, the imidazolium salt was a poorer catalyst than any of the triazolium salts, reaching only 19% spectroscopic yield after 3 h ( $\text{TOF}_{50} < 2$ ). Considerably lower activity of **4** and, in particular, **5** strongly suggests that the azolium cation plays a relevant role in tuning the activity of catalysts for this dehydrogenative coupling.

To evaluate potential interaction between the triazolium cation and the anion, triazolium tetrafluoroborate **1'** was synthesized by treating **1** with  $\text{AgBF}_4$ .  $^1\text{H}$  NMR analysis (300 MHz,

$\text{CDCl}_3$ ) of **1'** showed only minor changes in the spectrum compared to **1**, most diagnostically a significantly upfield shifted resonance at 8.65 ppm attributed to the triazolium CH (*cf.*  $\delta_{\text{H}}$  9.36 for **1**).<sup>27</sup> Such a resonance shift in the spectra of these azolium salts is indicative of less hydrogen-bonding counterions in solution.<sup>40</sup>

Both **1** and **1'** were further characterised by single crystal X-ray diffraction analysis (Fig. 1 and Table S4†). The molecular structure of iodide salt **1** displays weak non-classical interaction between the triazolium CH and iodide counterion (Fig. 1a), with a  $\text{H}\cdots\text{I}$  distance of 2.857 Å. Similar interactions exist in other triazolium iodide structures,<sup>41,42</sup> and molecular modelling of iodide anion binding by tetra-triazolium receptors by Beer and co-workers predicted binding distances on the same scale.<sup>43</sup> There was also a potential anion- $\pi$  interaction with a centroid- $\cdots\text{I}$  distance of 3.710 Å. The structure of the tetrafluoroborate salt **1'** revealed the  $\text{BF}_4^-$  counterion disordered over two sites without a clear hydrogen bonding pattern (Fig. 1b), in agreement with solution analysis.

This tetrafluoroborate analogue allowed the role of the anion in the catalysis to be probed. In contrast to the triazolium iodide **1**, the tetrafluoroborate analogue **1'** was catalytically completely inactive under the same conditions (Table 3, entry 6). This inactivity was also observed with tetrafluoroborate analogues **2'**, **3'**<sup>30</sup> and **5'** (entries 7–9), pointing towards a critical role of the iodide anion in catalysis.

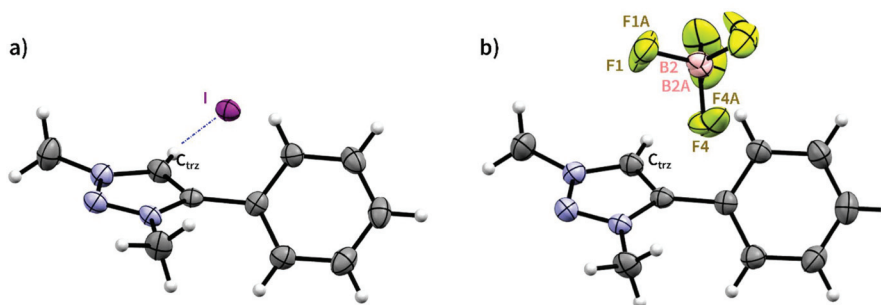
To further explore the role of cations and anions for catalysing this transformation, various other salts were evaluated (Table S2†). Ionic liquid  $(\text{Bu}_4\text{N})\text{Br}$ , previously reported to catalyze amine homocoupling of benzylamines when used as a solvent,<sup>26</sup> gave negligible conversion when added in catalytic amounts. Similarly  $(\text{H}_4\text{N})\text{Cl}$  was inactive, while  $(\text{Et}_4\text{N})\text{I}$  gave

Table 3 Catalytic activity of triazolium and imidazolium salts<sup>a</sup>

Entry	Catalyst	Heterocycle	R	R'	X	Yield <sup>b</sup> (%)		$\text{TOF}_{50}$ ( $\text{h}^{-1}$ )
						1 h	3 h	
1	<b>1</b>	Triazolium	Me	Ph	I	60	100	13
2	<b>2</b>	Triazolium	<sup>n</sup> Bu	<sup>n</sup> Bu	I	45	99	9
3	<b>3</b>	Triazolium	<sup>n</sup> Bu	Ph	I	16	75 <sup>c</sup>	5
4	<b>4</b>	Triazolium	<sup>n</sup> Bu	Mes	I	47	100	9
5	<b>5</b>	Imidazolium	Me	Me	I	5	19 <sup>d</sup>	<2
6	<b>1'</b>	Triazolium	Me	Ph	$\text{BF}_4$	0	0	—
7	<b>2'</b>	Triazolium	<sup>n</sup> Bu	<sup>n</sup> Bu	$\text{BF}_4$	0	0	—
8	<b>3'</b>	Triazolium	<sup>n</sup> Bu	Ph	$\text{BF}_4$	0	0	—
9	<b>5'</b>	Imidazolium	Me	Me	$\text{BF}_4$	0	0	—

<sup>a</sup> General conditions as described in Table 1. <sup>b</sup> Spectroscopic yield determined by  $^1\text{H}$  NMR analysis with respect to hexamethylbenzene. <sup>c</sup> 96% after 6 h. <sup>d</sup> 48% after 6 h.





**Fig. 1** (a) Molecular structure of triazolium salt **1**, determined by single crystal X-ray diffraction analysis. Selected bond lengths (Å) and angles (°):  $|C_{\text{trz}}\cdots I| = 3.783(3)$ ,  $|H_{\text{trz}}\cdots I| = 2.857$ ,  $|\angle C_{\text{trz}}-H_{\text{trz}}\cdots I| = 173.69$ . (b) Molecular structure of triazolium salt **1'**, determined by single crystal X-ray diffraction analysis.  $BF_4$  anion is disordered over two sites B2 (65.2%) and B2A (34.8%).

modest yields and reached nearly 40% after 4 h, a level of activity similarly to imidazolium salt **5** and indicative for an essential role of iodide for inducing catalytic activity. In agreement with this model, KI gave comparable yields, which implies a notable 8 turnover numbers (TONs) for this simple salt. While active, these yields are significantly lower than those of the triazolium iodides, which suggests a prominent role of the cation for inducing high conversion. In line with this notion, experiments with KI in the presence of 5 mol% of 18-crown-6 to passivate the cation quenched activity completely.

### Mechanistic insight

From the catalytic reactions described above, it is possible to gain some insight into mechanistic aspects of this process, and rule out certain pathways. For instance, involvement of an N-heterocyclic carbene as catalytically active species is unlikely, since addition of stoichiometric  $K^t\text{O}Bu$  to triazolium salt **1** induced slower conversion (*cf.* Table 1, entry 7), even though this base promotes the formation of free carbenes.<sup>44</sup> Although NHCs generated *in situ* have been reported as organocatalysts for various transformations,<sup>45–48</sup> a similar reaction pathway is evidently not occurring here.

Instead, iodide plays an essential role, which may be related to its unique redox properties and the low potential to form molecular iodine,  $I_2$ .<sup>49</sup> To probe this hypothesis, reactions were performed under deliberately oxidizing or reducing conditions. Thus, when the reaction was performed in deaerated solvent and under inert atmosphere, significantly slower conversion rates were observed (Table 4, entry 1 *vs.* 2), while saturation with  $O_2$  accelerates amine oxidation when compared to standard conditions in air (entry 3,  $\text{TOF}_{50} \sim 30 \text{ h}^{-1}$  *vs.*  $13 \text{ h}^{-1}$ ). Dependence on dioxygen as an oxidant for amine homocoupling has previously been observed in ionic liquid solvents,<sup>26</sup> as well as with photocatalytic systems.<sup>24,25</sup> Carrying out the reaction in the absence of light had no notable impact on catalyst activity (entry 4), discounting the possibility of photocatalysis occurring under these conditions.

These results point to a potential catalytic activity of molecular iodine for this transformation.  $I_2$ -Catalyzed and

**Table 4** Catalytic activity of various iodine-containing species, with variation of dioxygen content<sup>a</sup>

Entry	Catalyst	Atmosphere <sup>b</sup>	Yield <sup>c</sup> (%)			$\text{TOF}_{50} \text{ (h}^{-1}\text{)}$
			1 h	3 h	4 h	
1	<b>1</b>	Air	60	100	99	13
2	<b>1</b>	$N_2$	0	19	32	—
3	<b>1</b>	$O_2$	96	97 <sup>d</sup>	—	30
4	<b>1</b>	Air (darkness)	52	97	—	11
5	$I_2$	Air	42	91	98	7
6	$I_2$	$N_2$	10	15	28	—
7	<b>6</b>	Air	57	100	—	11
8	<b>6</b>	$N_2$	13	22	36	—
9	<b>1'</b> + $I_2$	Air	87	100 <sup>d</sup>	—	17
10	<b>1'</b> + $I_2$	$N_2$	11	20	33	—

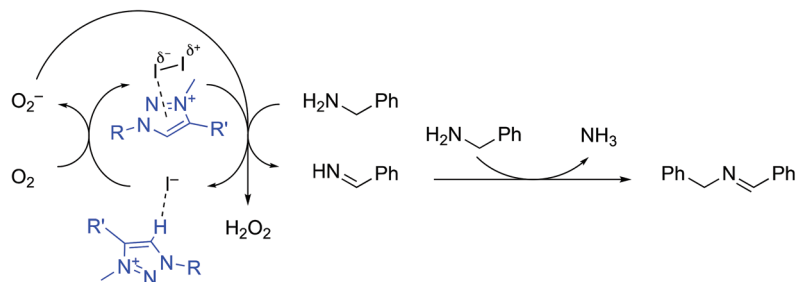
<sup>a</sup> General conditions: Benzyl amine (0.25 mmol), catalyst (5 mol%), hexamethylbenzene (internal standard), 1,2-dichlorobenzene (2 mL), 150 °C. <sup>b</sup> Solution saturated with the corresponding gas ( $N_2$  or  $O_2$ ) for 10 min before substrate addition. <sup>c</sup> Spectroscopic yield determined by  $^1\text{H}$  NMR analysis with respect to hexamethylbenzene. <sup>d</sup> Yield at 2 h.

-mediated reactions are of increasing interest for organic chemistry, including *e.g.* oxidative couplings of 2-aminopyridines and 1,3-diones, and for synthesis of other heterocycles.<sup>13,50–52</sup> Of relevance to this work, iodine was reported to produce imines as either proposed intermediates or low-yielding by-product upon oxidation of benzylamines to nitriles, indicating that high loadings of  $I_2$  (0.25–3 equivalents) induce transformations of these substrates.<sup>53,54</sup> However, no efficient and selective intermolecular imine synthesis by related catalytic methods has been reported.

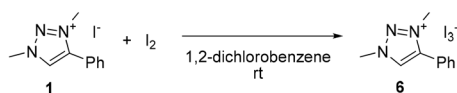
Indeed, when 5 mol% molecular iodine was reacted with benzyl amine under the conditions described above for triazolium iodide **1** (Table 4, entry 5), quantitative spectroscopic yields of the corresponding imine were obtained within 4 h. This activity is only slightly lower than with the triazolium iodide (7 *vs.*  $13 \text{ h}^{-1}$ , *cf.* entry 1, though the mol% in iodide is doubled). Upon addition of the substrate benzylamine, the initially purple  $I_2$  solution in 1,2-dichlorobenzene underwent a characteristic color change to yellow, which may be attributed to an amine-induced reduction of iodine to iodide.<sup>55</sup> The activity of  $I_2$  as a catalyst, as well as the key role of dioxygen







**Scheme 1** Proposed reaction pathway and role of the azolium salt and iodine in the amine oxidation.



**Scheme 2** Synthesis of triiodide salt **6**.

described above, suggests that molecular iodine may indeed be on the catalytic cycle of this reaction. Interestingly, when catalysis is carried out with  $I_2$  (5 mol%) under inert atmosphere (entry 6), 10% of imine was observed after 1 h, indicating approximately one turnover achieved in the absence of  $O_2$ . Notably, the triazolium iodide **1** is inactive under identical inert conditions.

We therefore propose the reaction mechanism shown in Scheme 1, consisting of a cascade redox process with as key process the presumably rate-limiting iodine-mediated oxidation of primary amine to the aldimine. Aldimine condensation with another equivalent of amine generates the final secondary imine, while the iodide by-product is regenerated under aerobic conditions back to iodine. The  $H_2O_2$  by-product was not directly observed in the reaction mixture, but it is proposed that the well-known ability of iodide to catalyze  $H_2O_2$  decomposition<sup>56</sup> makes detection unlikely. Moreover, iodide-mediated peroxide detection strategies are based on the development of a diagnostic yellow color of the solution,<sup>57</sup> which was observed to evolve when using **1** as catalyst (*vide supra*). The accelerating role of the triazolium salt has been tentatively attributed to potential interactions between the electron-poor azolium ring and molecular iodine, which induces polarisation of the I–I bond and facilitates the reduction to  $I^-$ .<sup>52,58</sup> The concomitant benzylamine C–H activation is postulated to be the rate-determining step and accordingly, activation of  $I_2$  through polarisation increases the observed reaction rate. A further beneficial role of the azolium cation may also be imparted by the stabilization of the iodide through hydrogen bonding or anion $\cdots\pi$  interactions.<sup>35,59,60</sup> This amine oxidation may therefore be considered as an anion–cation dual catalytic process, where the cation assists the redox process of the anion, which in turn induces substrate oxidation.

To evaluate the possible involvement of a triiodide anion due to the presence of both iodide and iodine,<sup>49,61</sup> the triazolium triiodide salt **6** was prepared by mixing an equimolar ratio of **1** and  $I_2$  in solution (Scheme 2). In  $CDCl_3$ , this reaction

led to an immediate color change from yellow to orange and the triazolyl proton resonance shifted about 1 ppm upfield to  $\delta_H = 8.55$  in the  $^1H$  NMR spectrum, indicative of weaker C–H $\cdots$ X hydrogen bonding upon transforming  $I^-$  to  $I_3^-$  (Fig. S8†). Compound **6** crystallized upon slow evaporation and the single crystals were analyzed by X-ray diffraction (Fig. S11†). The use of this triiodide salt **6** as precatalyst gave identical results, under air as the iodide salt **1** (Table 4, entry 7,  $TOF_{50}$  11 *vs.* 13  $h^{-1}$ ). Under inert conditions compound **6** achieved 13% yield after 1 h (entry 8), again suggesting one turnover in the absence of dioxygen. This behavior is reminiscent of that of  $I_2$ . Finally, when the inactive triazolium tetrafluoroborate **1'** was used in combination with  $I_2$ , excellent catalytic activity was observed that was far superior to  $I_2$  alone (entry 9,  $TOF_{50}$  17 *vs.* 7  $h^{-1}$  in air), and in fact even more active than **1** or **6** under aerobic conditions. Again, under inert atmosphere (entry 10), only a single turnover was observed in the first hour of reaction. These data support the mechanistic model with  $I_2$  as the catalytically active species for oxidation of the amine, and the triazolium cation playing a crucial role in accelerating turnover frequency.

## Conclusions

We report a novel metal-free approach to the oxidative homocoupling of benzylic amines catalyzed by triazolium iodide salts, selectively giving imine products in quantitative yields. Turnover frequencies of up to 13  $h^{-1}$  were achieved with **1** under aerobic conditions and up to 30  $h^{-1}$  under  $O_2$  atmosphere. Preliminary mechanistic investigations indicate critical roles of both the cation and the anion of the precatalyst, which has been attributed to an iodide/iodine couple for the actual oxidation, and a triazolium $\cdots I_2$  interaction for facile iodine reduction and concomitant amine oxidation. Moreover, the activity of the triazolium iodide catalysts is dependent on dioxygen as benign terminal oxidant. While the substrate scope is limited to benzylic amines, the simplicity of the system is appealing. This triazolium iodide/iodine catalysis constitutes a more cost-effective approach to amine oxidation than the more widely reported transition-metal based strategies. Notably, the synergistic effects of triazolium cations and iodide/iodine-mediated redox catalysis may be of broader rele-



vance and an advantageous new concept to other organocatalytic redox reactions such as the azapyrrole synthesis or Bayer-Villiger oxidation.

## Experimental section

### General considerations

All compounds, unless otherwise stated, were obtained from commercial sources and used as provided. Triazolium salts **1**,<sup>27</sup> **2**,<sup>32</sup> **2'**,<sup>28</sup> **3**,<sup>29</sup> **3'**,<sup>30</sup> and **4**<sup>28</sup> were prepared according to literature procedures, as were imidazolium salts **5**<sup>62</sup> and **5'**.<sup>63</sup> NMR spectra were recorded on a Bruker spectrometer operating at room temperature. Chemical shifts ( $\delta$  in ppm,  $J$  in Hz) were referenced to residual solvent resonances and are reported downfield from SiMe<sub>4</sub>. High resolution mass spectrometry was performed by the Analytical Research Services at University of Bern.

### General catalytic procedure

Benzyl amine (27  $\mu$ L, 0.25 mmol) and triazolium salt **1** (3.6 mg, 0.012 mmol, 5 mol%) were dissolved in 1,2-dichlorobenzene (2 mL) and heated at 150 °C in round-bottomed flasks fitted with water-cooled condensers. Hexamethylbenzene was added as internal standard and reaction progress monitored by <sup>1</sup>H NMR analysis (Fig. S1–S5 and S13–S18†). For reactions under N<sub>2</sub> atmosphere or O<sub>2</sub> atmosphere, reaction was carried out in a sealed Schlenk tube, and the reaction mixture was saturated with the relevant gas for 10 minutes before addition of benzyl amine, and the mixture was kept under a balloon of gas.

### Synthesis of **1'**

Triazolium salt **1** (0.100 g, 0.33 mmol) was dissolved in CH<sub>2</sub>Cl<sub>2</sub> (20 mL). AgBF<sub>4</sub> (0.070 g, 0.34 mmol) was added and the reaction mixture stirred at room temperature for 2 h. This was filtered through Celite to remove insoluble silver salts and concentrated under reduced pressure to yield **1'** as an off-white solid (0.080 g, 0.31 mmol, 92%). HRMS (ESI+): Calculated for C<sub>10</sub>H<sub>12</sub>N<sub>3</sub>  $m/z$  = 174.1026 [M – BF<sub>4</sub>]<sup>+</sup>. Found  $m/z$  = 174.1022; <sup>1</sup>H NMR (300 MHz, CDCl<sub>3</sub>):  $\delta$  = 8.65 (s, 1H, C<sub>trz</sub>H), 7.51–7.65 (m, 5H, C<sub>Ph</sub>H), 4.43 (s, 3H, N–CH<sub>3</sub>), 4.26 (s, 3H, C–CH<sub>3</sub>); <sup>13</sup>C NMR (75 MHz, CDCl<sub>3</sub>):  $\delta$  = 132.0, 129.9, 129.8, 129.5 (4  $\times$  C<sub>Ph</sub>), 128.9, 125.8 (2  $\times$  C<sub>trz</sub>), 40.5, 38.3 (2  $\times$  NCH<sub>3</sub>).

### Synthesis of **6 in situ**

Triazolium salt **1** (0.006 g, 0.02 mmol) and I<sub>2</sub> (0.005 g, 0.02 mmol) were dissolved in CDCl<sub>3</sub> (or 1,2-dichlorobenzene). <sup>1</sup>H NMR (300 MHz, CDCl<sub>3</sub>):  $\delta$  = 8.55 (s, 1H, triazolium C<sub>trz</sub>H), 7.69–7.75 (m, 5H, C<sub>Ph</sub>H), 4.57 (s, 3H, N–CH<sub>3</sub>), 4.37 (s, 3H, C–CH<sub>3</sub>). Single crystals suitable for X-ray diffraction analysis grew upon slow evaporation of this sample.

### Crystal-structure determination

Triazolium iodide **1** crystallized upon slow diffusion of diethyl ether into a saturated methanol solution of the compound.

Triazolium tetrafluoroborate **1'** and triazolium triiodide **6** crystallized by slow evaporation of CDCl<sub>3</sub> solutions. All crystals were mounted in air at ambient conditions. The measurements for **1** were made on an Oxford Diffraction SuperNova area-detector diffractometer<sup>64</sup> using mirror optics monochromated Mo K $\alpha$  radiation ( $\lambda$  = 0.71073 Å) and Al filtered.<sup>65</sup> The unit cell constants and an orientation matrix for data collection were obtained from a least-squares refinement of the setting angles of reflections in the range 2.6° <  $\theta$  < 27.7°. A total of 1402 frames were collected using  $\omega$  scans, with 1.0 + 1.0 seconds exposure time, a rotation angle of 1.0° per frame, a crystal-detector distance of 65.0 mm, at  $T$  = 173(2) K. All measurements for **1'** and **6** were made on a RIGAKU Synergy S area-detector diffractometer<sup>64</sup> using mirror optics monochromated Cu K $\alpha$  radiation ( $\lambda$  = 1.54184 Å). The unit cell constants and an orientation matrix for data collection were obtained from a least-squares refinement of the setting angles of reflections in the range 5.6° <  $\theta$  < 76.9° (for **1'**) or 4.4° <  $\theta$  < 76.4° (for **6**). A total of 2578 frames (**1'**) or 3102 frames (**6**) were collected using  $\omega$  scans, with exposure times of 0.05 and 0.18 s (**1'**) or 5 and 20 s (**6**), a rotation angle of 0.5° per frame, a crystal-detector distance of 34.0 mm, at  $T$  = 173(2) K. Data reduction for all three data sets was performed using the CrysAlisPro<sup>64</sup> program. The intensities were corrected for Lorentz and polarization effects. For **1**, a numerical absorption correction based on Gaussian integration over a multifaceted crystal model was applied. For **1'** and **6**, the intensities were corrected for Lorentz and polarization effects, and an absorption correction based on the multi-scan method using SCALE3 ABSPACK in CrysAlisPro<sup>1</sup> was applied. Data collection and refinement parameters are given in Table S3.† The structure was solved by direct methods using SHELXT,<sup>66</sup> which revealed the positions of all non-hydrogen atoms, which were refined anisotropically. All H-atoms were placed in geometrically calculated positions and refined using a riding model where each H-atom was assigned a fixed isotropic displacement parameter; for **1** and **1'**, this value was equal to 1.2 $U_{eq}$  of its parent atom (1.5 $U_{eq}$  for the methyl groups), while for **6** the value was equal to 1.5 $U_{eq}$  of its parent atom. In the structure of **1'**, the BF<sub>4</sub> anion is disordered about two orientations. The geometries of the two moieties were restrained to be similar. Its ADP's were restrained by the SHELX SIMU and RIGU instructions. Refinement of the structure was carried out on  $F^2$  using full-matrix least-squares procedures, which minimized the function  $\sum w(F_o^2 - F_c^2)^2$ . The weighting scheme was based on counting statistics and included a factor to down-weight the intense reflections. All calculations were performed using the SHELXL-2014/7<sup>67</sup> program in OLEX2.<sup>68</sup> The crystal of **6** used for structure determination was twinned, and thus the integration of just the major domain was used. The quality of this single crystal structure determination is lowered by this fact. The results are sufficient for a discussion of the crystal packing but do not allow a discussion of bonding geometry. Further details are given in Table S4.† Crystallographic data for all structures have been deposited with the Cambridge Crystallographic Data Centre (CCDC) as



supplementary publication numbers 2016764 (1), 2016765 (1'), and 2016766 (6).†

## Conflicts of interest

The authors declare no conflict of interest.

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