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Non-aggregated ruthenium naphthalocyanine enabling homogeneous carbene insertion into N–H bonds at low catalyst loading†

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The novel ruthenium octa-*n*-butoxy-naphthalocyanine complex was shown to retain an essentially monomeric state in dilute solutions. It was successfully applied as a homogeneous catalyst for carbene insertion into N–H bonds of amines with various substitution patterns, providing high yields of glycine derivatives.

A wide variety of complexes based on porphyrin,^{1–3} phthalocyanine,⁴ corrole,⁵ corrolazine⁶ and porphyrazine^{7,8} macrocyclic ligands are widely applied as catalysts for many reactions. In particular, these tetrapyrrolic complexes are robust and versatile catalysts for various carbene transfer reactions.^{9,10} For instance, carbene insertion into N–H bonds, considered as a challenging transformation owing to the high coordinating properties of amines leading to catalyst poisoning, can provide a wide range of bioactive compounds and pharmaceuticals. Compared to traditional Rh-based catalysts, which are usually required in amounts of 1 to 10 mol%,¹¹ tetrapyrrolic complexes efficiently catalyze carbene transfer reactions at a rather low loading. While ruthenium^{12–16} and iron^{15,17–19} porphyrins mediated N–H carbene insertion when using 1 mol% catalyst loading, iridium porphyrin afforded high product yields in the presence of only 0.06 mol%.²⁰ Furthermore, iron corroles were shown to be particularly efficient at 0.1 mol% loading providing a fast carbene insertion into N–H bonds of primary and secondary amines at high substrate concentration.^{21,22} In turn, iron and ruthenium phthalocyanines exhibited promising activity in cyclopropanation and carbene insertion into N–H bonds.^{23–29} Our groups

have shown that ruthenium octa-*n*-butoxyphthalocyanines in amounts as low as 0.05–0.2 mol% catalyze the insertion of carbene into N–H bonds of aromatic and aliphatic amines under mild conditions.^{27,28} The high catalytic efficiency of different tetrapyrrolic complexes suggests that other overlooked porphyrin-like complexes deserve more careful evaluation in carbene transfer reactions. Indeed, modification of the tetrapyrrolic structure strongly influences their spectroelectrochemical properties^{30–32} and hence should affect their catalytic properties.

Naphthalocyanine complexes can be regarded as the closest phthalocyanine analogues derived from the annulation of benzene moieties onto the isoindoline fragments of a phthalocyanine core. To date, their application in catalysis has been very rare and limited to heterogeneous processes. For example, unsubstituted cobalt naphthalocyaninate CoNc deposited on a rotating disc electrode electrocatalytically reduced oxygen and oxidized hydrazine.³³ Cathode-supported CoNc was superior to carbon black in the oxygen reduction reaction (ORR).³⁴ Lithium naphthalocyaninate Li₂Nc deposited on graphite catalyzed the electroreduction of CO₂ in KHCO₃ aqueous solution.³⁵ However, to the best of our knowledge, naphthalocyanine complexes have never been used as catalysts in carbene transfer and other homogeneous reactions. In fact, they tend to strongly aggregate in solution and their aggregation properties are much more pronounced compared to their phthalocyanine counterparts which can hinder access to the metal site and therefore prevent the catalytic reaction. Although several strategies can be employed to suppress naphthalocyanine aggregation—such as dilution,³⁶ heating,^{37,38} supramolecular interactions,^{38,39} attachment of appropriate groups to the macrocyclic core³⁹ or addition of axial bulky ligands⁴⁰—their homogeneous catalytic applications have yet to be demonstrated.

Herein, we report the first catalytic application of a naphthalocyanine complex in a carbene transfer reaction. The ruthenium complex $[(\gamma\text{-BuO})_8\text{NcRu}](\text{CO})$, bearing eight *n*-butoxy groups in the peripheral positions, was synthesized and characterized by UV-Vis, HR ESI MS and ¹H NMR techniques. Unexpectedly, $[(\gamma\text{-BuO})_8\text{NcRu}](\text{CO})$ is non-aggregated

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at 2–20 μM concentrations, which allows its application in homogeneous catalysis exemplified by the reaction of amines with ethyl diazoacetate (EDA).

The syntheses of all ruthenium naphthalocyanines described in the literature were performed by the template condensation of the corresponding phthalonitrile in the presence of a ruthenium precursor.^{41–44} However, the template condensation of 6,7-dibutoxynaphthalene-2,3-dicarbonitrile with $\text{Ru}_3(\text{CO})_{12}$ in refluxing *o*-dichlorobenzene (*o*-DCB) did not yield even trace amounts of the target ruthenium naphthalocyanine (Scheme 1). Next, we attempted the direct insertion of the ruthenium cation into the preformed macrocycle following a method previously used for the synthesis of ruthenium phthalocyanines (Scheme 1).^{25,29,45,46} The peripherally substituted octa-*n*-butoxy-naphthalocyanine $\text{H}_2[(\gamma\text{-BuO})_8\text{Nc}]$ was prepared from the corresponding phthalonitrile by Li-template condensation with subsequent acid treatment of the lithium complex.⁴⁷ The reaction of $\text{H}_2[(\gamma\text{-BuO})_8\text{Nc}]$ with $\text{Ru}_3(\text{CO})_{12}$ proceeded smoothly in refluxing *o*-DCB with the progressive disappearance of the broad Q-bands of $\text{H}_2[(\gamma\text{-BuO})_8\text{Nc}]$ at 713 nm and 783 nm and the appearance of the narrow Q band of the metal complex at 726 nm (Fig. S1†). The unreacted naphthalocyanine was separated by column chromatography on silica. Further purification by size-exclusion chromatography allowed for the isolation of the target ruthenium naphthalocyanine $[(\gamma\text{-BuO})_8\text{NcRu}](\text{CO})$ in a 25% yield. It should be noted that in a similar reaction of peripherally substituted phthalocyanines with $\text{Ru}_3(\text{CO})_{12}$ in *o*-DCB, in addition to the expected mononuclear complexes $[\text{PcRu}](\text{CO})$, μ -carbido binuclear complexes $[\text{PcRu}]_2(\mu\text{-C})$ were also formed.^{25,45} In contrast, we did not observe any other naphthalocyanine complexes in the MALDI TOF spectra of all isolated fractions, indicating more selective ruthenium insertion in the naphthalocyanine case.

The obtained ruthenium complex was characterized by UV-Vis, ^1H NMR, MALDI-TOF MS and HR ESI-MS techniques. The UV-Vis spectrum of naphthalocyanine $[(\gamma\text{-BuO})_8\text{NcRu}](\text{CO})$ in CH_2Cl_2 exhibits a narrow Q-band at 726 nm (Fig. S2†). The UV-vis spectra remained unchanged in CH_2Cl_2 and in $\text{C}_2\text{H}_4\text{Cl}_2$

in the concentration range of 2–20 μM (Fig. 1 and Fig. S7†), showing a linear dependence of absorbance intensity on concentration at five wavelengths, which suggests the absence of aggregation. A comparison of the normalized UV-vis spectra of 2 μM and 20 μM solutions indicates very small differences in the Soret region as well as in the shoulders at 697 nm and around 800 nm (Fig. S8†), which might be compatible with a minor aggregation phenomenon. Thus, $[(\gamma\text{-BuO})_8\text{NcRu}](\text{CO})$ maintains essentially a monomeric state in dilute solutions, although the presence of trace amounts of aggregated species cannot be excluded. Previous studies showed that the addition of eight butoxy substituents at either peripheral or non-peripheral positions of phthalocyanine complexes prevents their aggregation.^{25,28} Consequently, the introduction of eight butoxy groups into the naphthalocyanine platform, which is more prone to aggregation due to its extended aromatic plane, should reduce the aggregation. The presence of the CO axial ligand also hinders the aggregation. Thus, the metal-free

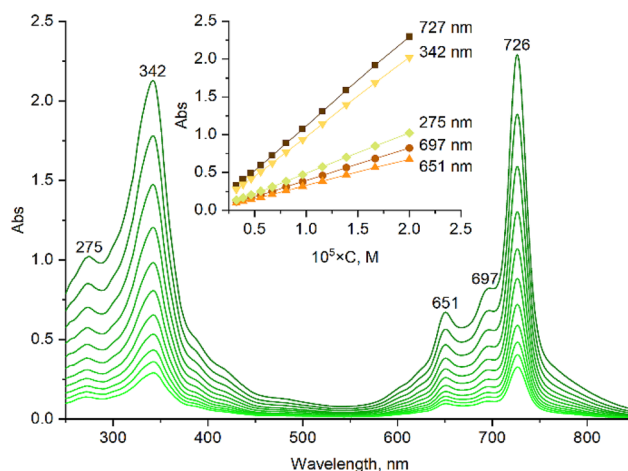
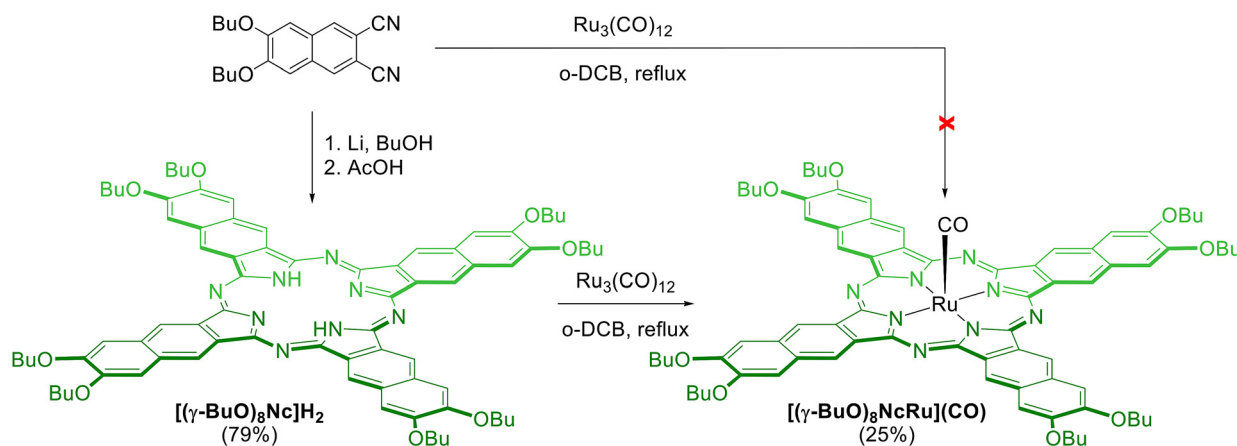


Fig. 1 The concentration dependence of the UV-Vis spectra of the ruthenium naphthalocyanine $[(\gamma\text{-BuO})_8\text{NcRu}](\text{CO})$ in CH_2Cl_2 .



Scheme 1 Synthesis of the ruthenium naphthalocyanine $[(\gamma\text{-BuO})_8\text{NcRu}](\text{CO})$.



naphthalocyanine $\text{H}_2[(\gamma\text{-BuO})_8\text{NcRu}]$ is strongly aggregated at 10^{-4} – 10^{-5} M concentrations, whereas $[(\gamma\text{-BuO})_8\text{NcRu}](\text{CO})$ is in the monomeric form (Fig. S1†).

This is in sharp contrast to the metal-free $\text{H}_2[(\beta\text{-BuO})_8\text{Pc}]$ and its ruthenium complex $[(\beta\text{-BuO})_8\text{PcRu}](\text{CO})$ found in the monomeric state in this concentration range in nonpolar solvents such as CH_2Cl_2 and CHCl_3 .²⁵

Compared to the peripherally substituted octa-*n*-butoxy phthalocyanine $[(\beta\text{-BuO})_8\text{PcRu}](\text{CO})$, the maximum of the Q band in the UV-vis spectrum of the naphthalocyanine complex bathochromically shifted to 71 nm (Fig. S2†). Since this absorption originates from the HOMO–LUMO transition, this shift can be explained by the destabilization of the HOMO upon annulation with the benzene rings.³⁰

The ^1H NMR spectrum of $[(\gamma\text{-BuO})_8\text{NcRu}](\text{CO})$ recorded in CDCl_3 in the presence of 5 μL pyridine-*d*₅ exhibits two singlet signals at 7.70 and 9.51 ppm attributed to the aromatic protons of the naphthalocyanine macrocycle and four signals at 4.35, 2.03, 1.67 and 1.11 ppm corresponding to the aliphatic chains of the *n*-butoxy groups (Fig. S3†). Upfield shifted shoulder signals of low intensity near the signals of H_{Ar2} (7.70 ppm), $\alpha\text{-CH}_2$ (4.35 ppm), $\gamma\text{-CH}_2$ (1.67 ppm) and CH_3 (1.11 ppm) were also observed (Fig. S3†). Given the much higher $[(\gamma\text{-BuO})_8\text{NcRu}](\text{CO})$ concentration ($\sim 10^{-3}$ M) used in ^1H NMR measurements compared to UV-vis measurements and catalytic experiments (2–20 μM), this observation can be explained by minor aggregation occurring even in the presence of 5 μL of pyridine-*d*₅ which was added to prevent the aggregation due to the axial coordination of pyridine. We have recorded the ^1H NMR spectrum of $[(\gamma\text{-BuO})_8\text{NcRu}](\text{CO})$ in pure pyridine-*d*₅ and observed no additional minor signals, thus confirming the purity of $[(\gamma\text{-BuO})_8\text{NcRu}](\text{CO})$ and its low tendency for aggregation.

The MALDI TOF mass spectrum shows two molecular clusters corresponding to mononuclear $\{[(\gamma\text{-BuO})_8\text{NcRu}] + \text{H}\}^+$ and binuclear $\{[(\gamma\text{-BuO})_8\text{NcRu}]_2 + \text{H}\}^+$ species without axial ligands (Fig. S6†).

However, the presence of the coordinated CO was evidenced by the HR ESI-MS technique, which showed the molecular

cluster of $[(\gamma\text{-BuO})_8\text{NcRu}](\text{CO})$ (Fig. S9†). Previously, the CO axial ligand was also found in ruthenium phthalocyanine complexes.^{29,45,48–50} Although a range of ruthenium naphthalocyanines have been prepared,^{41–44,51–53} complexes containing a coordinated CO ligand have not yet been published.

The lack of $[(\gamma\text{-BuO})_8\text{NcRu}](\text{CO})$ aggregation at 2–20 μM concentrations prompted us to probe its catalytic activity in the benchmark carbene transfer reaction between ethyl diazoacetate (EDA) and aniline using low catalyst loadings (Table 1).

We have initially used 2 equiv. of EDA in order to determine whether $[(\gamma\text{-BuO})_8\text{NcRu}](\text{CO})$ is capable of performing single and double N–H insertions. The complete aniline conversion and a 96% yield of the tertiary derivative were obtained using 0.1 mol% $[(\gamma\text{-BuO})_8\text{NcRu}](\text{CO})$ (Table 1, entry 1). A decrease in the catalyst loading to 0.025 mol% slightly reduced the yield of the double N–H insertion product (Table 1, entry 3). Further reduction to 0.01 mol% resulted in comparable yields of mono- and disubstituted glycine derivatives (59% and 41% yields, respectively) (Table 1, entry 4). It is noteworthy that even with a 0.002 mol% catalyst loading, a 39% yield of the single N–H insertion product was obtained, although the reaction required 16 h (Table 1, entry 5). The very high turnover number (TON) of 20 050 obtained under these conditions indicates the high efficiency of $[(\gamma\text{-BuO})_8\text{NcRu}](\text{CO})$ as a catalyst for carbene insertion reactions. Interestingly, when the previously reported phthalocyanine counterpart $[(\gamma\text{-BuO})_8\text{PcRu}](\text{CO})$ ²⁸ was used under the same reaction conditions, only a 26% yield of the single insertion product and less than 1% yield of the double insertion product were obtained (TON = 13 250) (Table 1, entry 6). Previously, iron porphyrin and iron corrole complexes were reported as efficient catalysts for N–H insertion, affording a TON of up to 970 with 0.1 mol% catalyst loading.²² Iridium porphyrin catalyzed both single and double EDA insertion to aromatic and aliphatic amines with a total TON of up to 3180 in the presence of 0.06 mol% catalyst.²⁰ In turn, engineered variants of myoglobin mediated asymmetric N–H carbene insertion of aromatic amines with 2-diazopropanoate to achieve a TON of up to 2470 with 0.02 mol%.⁵⁴

Table 1 Interaction of aniline and EDA catalyzed by the ruthenium naphthalocyanine $[(\gamma\text{-BuO})_8\text{NcRu}](\text{CO})$ at different concentrations^a

Entry	Catalyst ^b (mol%)	Reaction time, h	Aniline conversion %	Yield of double insertion product, %	Yield of single insertion product, %	TON
1	A (0.1)	1	100	96	4	1960
2	A (0.05)	3	100	92	8	3840
3	A (0.025)	5	96	91	5	7480
4	A (0.01)	6	100	41	59	14 100
5	A (0.002)	16	40	1	39	20 050
6	B (0.002)	16	26	<1	26	13 250

^a Reaction conditions: aniline (0.5 mmol), EDA (1.05 mmol), catalyst, $\text{C}_2\text{H}_4\text{Cl}_2$ (0.5 mL), 40 °C, and Ar. Yields and aniline conversions based on aniline were determined by the ^1H NMR method. ^b A = $[(\gamma\text{-BuO})_8\text{NcRu}](\text{CO})$ and B = $[(\beta\text{-BuO})_8\text{PcRu}](\text{CO})$.



Thus, the high efficiency of $[(\gamma\text{-BuO})_8\text{NcRu}](\text{CO})$ in carbene insertion into amines places this complex among the most efficient catalysts for this reaction.

A broad range of aromatic and aliphatic amines were then evaluated for N–H insertion with 1.1 equiv. of EDA with respect to the substrate using a catalyst loading of 0.0125 mol%.

Table 2 Reaction of amines with EDA catalyzed by the ruthenium naphthalocyanine $[(\gamma\text{-BuO})_8\text{NcRu}](\text{CO})^a$

Entry	Substrate	Reaction time	Yield of single N–H insertion product, %	Yield of double N–H insertion product, %	TON
1		1 h	95	5	8400
2		1 h	94	6	8480
3		2 h	91	9	8720
4		1 h	90	10	8800
5		1 h	98	2	8160
6		4 h	96	4	8320
7		1 h 10 min	94	Traces	7520
8		30 min	88	2	7360
9		30 min	78	20	9440
10		3 h 40 min	84	2	7040
11		3 h 40 min	93	3	7920
12		4 h 30 min	73	0	5840
13		4 h	52	Traces	4160
14		21 h	>99	—	7920
15		19 h	16	0	1280
16		21 h	69	—	5520

^a Reaction conditions: amine (0.5 mmol), EDA (0.55 mmol), $[(\gamma\text{-BuO})_8\text{NcRu}](\text{CO})$ (0.0125 mol%), $\text{C}_2\text{H}_4\text{Cl}_2$ (0.5 mL), Ar, and 40 °C.

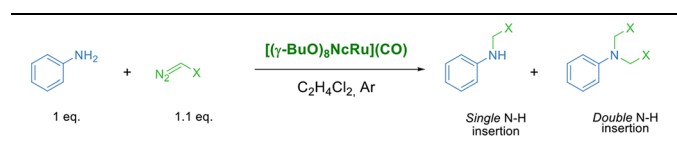


and its derivatives bearing electron-donating groups (*p*-Me and *p*-MeO) reacted smoothly, affording single N–H insertion products with 91–95% yields (Table 2, entries 1–3). Similarly, high product yields were obtained with halogen-substituted substrates (90, 98, 96 and 94% for *p*-F, *p*-Cl, *o*-F and *o*-Br derivatives) (Table 2, entries 4–7).

The introduction of strong electron-withdrawing groups, such as *p*-nitro and 3,5-bis-(trifluoromethyl), was tolerated, giving product yields of 84 and 93%, respectively, after longer reaction times (Table 2, entries 10 and 11). Even such an electron-deficient amine as pentafluoroaniline furnished the corresponding glycine derivative with a 73% yield (Table 2, entry 12). The presence of bulky substituents in the *ortho*-positions of 2,6-diisopropylamine did not prevent the reaction, though it led to a decrease in the product yield to 52% (Table 2, entry 13). Aliphatic amines were also suitable substrates for carbene insertion but required longer reaction times to achieve satisfactory yields. For instance, cyclopropylamine reacted slowly to furnish a 16% yield after 19 h (Table 2, entry 15). This decrease in the reactivity can be explained by the coordination of the amine to the ruthenium center, thus inhibiting the formation of active metal–carbene species. However, diisopropylamine underwent N–H insertion, giving a 69% product yield, most probably due to the bulky substituents hindering the coordination of this amine to the metal site (Table 2, entry 16). Finally, secondary amines can also be amenable to N–H carbene insertion. For example, *N*-methylaniline provided a quantitative yield of the target tertiary amine after 21 h (Table 2, entry 14).

To demonstrate the synthetic value of this protocol, we have carried out the cyclopropanation of *p*-chlorostyrene on a 10 mmol scale with 0.0125 mol% $[(\gamma\text{-BuO})_8\text{NcRu}](\text{CO})$ loading. After a 3 h reaction, 1.76 g (83% yield) of the pure target cyclopropanation product was isolated (Fig. S10†), thus confirming the scalability of this method.

Table 3 Reaction of aniline with diazo compounds catalyzed by $[(\gamma\text{-BuO})_8\text{NcRu}](\text{CO})^a$



Entry	Diazo compound	Reaction time, h	Yield of single N–H insertion product, %	Yield of double N–H insertion product, %	TON
1 ^b		17 h	86	13	8960
2 ^b		1 h	29	0	2320
3 ^c		4 h	88	7	4750
4 ^c		4 h	84	7	4550

^a Reaction conditions: aniline (0.5 mmol), diazo compound (0.55 mmol), $[(\gamma\text{-BuO})_8\text{NcRu}](\text{CO})$, and $\text{C}_2\text{H}_4\text{Cl}_2$ (0.5 mL). ^b 0.0125 mol% catalyst and 40 °C. ^c 0.02 mol% catalyst and 60 °C.

The versatility of $[(\gamma\text{-BuO})_8\text{NcRu}](\text{CO})$ was further explored using different diazo compounds in the cyclopropanation of styrene (Table 3).

When we used *t*-butyl diazoacetate and benzyl diazoacetate as carbene precursors at 40 °C, we observed lower activity compared to EDA, most probably because of the bulkiness of *t*-butyl and benzyl groups. However, a smooth carbene N–H insertion occurred at 60 °C, affording single insertion products with 88 and 84% yields, respectively (Table 3, entries 3 and 4). Diazoacetonitrile was also effectively activated by $[(\gamma\text{-BuO})_8\text{NcRu}](\text{CO})$ providing a single insertion product containing the cyano group with an 86% yield (Table 3, entry 1). The introduction of a fluorinated fragment was achieved using 1,1,1-trifluorodiazoethane, albeit with an inferior product yield (Table 3, entry 2). After a 1 h reaction, the reaction mixture was bleached, indicating degradation of $[(\gamma\text{-BuO})_8\text{NcRu}](\text{CO})$, leading to a 29% yield of the secondary amine bearing a CF_3 group.

Conclusions

Ruthenium naphthalocyanine $[(\gamma\text{-BuO})_8\text{NcRu}](\text{CO})$ containing peripheral butoxy substituents was prepared by the direct reaction of the metal-free ligand with $\text{Ru}_3(\text{CO})_{12}$ for the first time. The strong tendency for aggregation in solution is a significant drawback of naphthalocyanine complexes, preventing their catalytic applications. In contrast to published naphthalocyanine complexes, $[(\gamma\text{-BuO})_8\text{NcRu}](\text{CO})$ shows no appreciable aggregation at 2–20 μM concentrations. Taking advantage of this property, we have demonstrated the first catalytic application of the naphthalocyanine complex exemplified by the carbene transfer reaction. The complex $[(\gamma\text{-BuO})_8\text{NcRu}](\text{CO})$ was shown to be a very efficient catalyst for the insertion of ethyl diazoacetate into N–H bonds of sixteen aromatic and aliphatic amines at low catalyst loading (0.0125 mol%) and a quasi-equimolar EDA/substrate ratio. A wide range of monosubstituted glycine derivatives was obtained with up to quantitative yields and up to 9440 turnover numbers. Importantly, a catalyst loading as low as 0.002 mol% can be successfully used in the reaction between aniline and EDA to afford the single N–H insertion product with a TON of 20 050. The generality of this methodology was further demonstrated using several diazo compounds as carbene precursors. The promising results obtained in this work on the carbene insertion into N–H bonds suggest that the yet overlooked naphthalocyanine complexes might be efficient catalysts for other types of reactions.

Data availability

The data supporting this article have been included as part of the ESI.†

Conflicts of interest

There are no conflicts to declare.



Acknowledgements

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