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## Calixarene alpha-ketoacetylenes: versatile platforms for reaction with hydrazine nucleophile†

Anton A. Muravev,<sup>a</sup> Svetlana E. Solovieva,<sup>\*ab</sup> Farida B. Galieva,<sup>b</sup> Olga B. Bazanova,<sup>a</sup> Ildar Kh. Rizvanov,<sup>a</sup> Kamil A. Ivshin,<sup>b</sup> Olga N. Kataeva,<sup>ab</sup> Susan E. Matthews<sup>c</sup> and Igor S. Antipin<sup>ab</sup>

Late stage diversification of calix[4]arenes and thiacalix[4]arenes with heterocycles remains a significant synthetic challenge and hampers further exploitation of the scaffolds. Here we describe the development of a short and facile synthetic route to conformationally diverse novel calix[4]arene and thiacalix[4]arene ynones using a palladium cross coupling approach (5% Pd(II) + 10% Cu(I)) with benzoyl chloride. Their successful conversion to heterocycles to afford pyrazoles was demonstrated through treatment with hydrazine. Functionalisation is calixarene conformation and linker independent enabling access to a library of structures.

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### Introduction

(Thia)calix[4]arenes are a widely investigated platform for molecular recognition due to the ease of their preparation and functionalisation through regio-, stereo- and iterative modification<sup>1–5</sup> and the ability to immobilise in four distinct conformations.<sup>6</sup> These macrocycles not only provide a cavity for inclusion of various guest species (such as fullerenes<sup>7</sup> and dyes<sup>8</sup>), but also assist coordination (small ions<sup>9</sup> to large biomolecules<sup>10</sup>) by means of receptor groups, which form a binding site for a specific substrate.<sup>11–14</sup> The latter often leads to a synergetic effect, or multivalency,<sup>15</sup> in such applications<sup>16</sup> as biological recognition, sensing, luminescence, magnetism, catalysis, and membrane separation.

Heterocycles, including triazoles,<sup>17</sup> imidazoles,<sup>18</sup> pyridines<sup>19</sup> and pyrroles<sup>20</sup> are some of the most extensively investigated moieties on a calixarene framework,<sup>21</sup> however extension to more diverse rings remains challenging. Two fundamental approaches are available for inclusion of such functionality; introduction of a pre-assembled moiety (Fig. 1a) or *in situ* modification of an activated calixarene (Fig. 1b and c). The former route is convenient only when the heterocycle is stable under the employed reaction conditions or does not require the use of protecting groups for example terpyridines<sup>22,23</sup> or thiophenes.<sup>24</sup>

The most extensively studied example of the second approach is the functionalisation of calixarenes with alkynes. This route has offered up a number of derivatisation possibilities through participation in pericyclic reactions (Fig. 1b) with nitrile oxides<sup>25</sup> and azides<sup>26–28</sup> yielding isoxazoles and triazoles. Additional examples include the reaction of calixarene  $\beta$ -amidoamines with triethylamine to afford imidazolines<sup>29</sup> and the three-component Biginelli reaction (Fig. 1c) of calixarene aldehydes with urea and  $\beta$ -ketoesters to give dihydropyrimidines.<sup>30</sup> However, these methods offer limited diversity of heterocycle structure and a particular challenge is the avoidance of competitive inter and intramolecular reactions leading to mixed, bridged<sup>31–34</sup> or multimeric products.

To-date conversion to ynones and subsequent nucleophilic addition–elimination reactions have not been employed on macrocycles. This approach offers a rapid entry to a variety of nitrogen- and oxygen-containing heterocycles<sup>35</sup> through diverse cyclization pathways from a single precursor. In contrast to small molecules, a challenge to developing this approach on (thia)calixarenes is the avoidance of inter- or intramolecular crosslinking on reaction with nucleophiles; this is particularly important, when polytopic platforms are considered.

Here we describe the successful functionalisation of (thia)calixarenes with multiple ynones and demonstrate their potential for heterocycle formation through subsequent controlled reaction with hydrazine to produce pyrazoles without crosslinking.

### Results and discussion

There are numerous approaches to the preparation of ketoacetylenes: the addition of lithium acetylenides to aromatic

<sup>a</sup>Arbuzov Institute of Organic and Physical Chemistry, FRC Kazan Scientific Center of RAS, Kazan, 420088 Russia. E-mail: antonm@iopc.ru; svsol@iopc.ru

<sup>b</sup>A.M. Butlerov Institute of Chemistry, Kazan Federal University, Kazan, 420008 Russia

<sup>c</sup>School of Pharmacy, University of East Anglia, Norwich, NR4 7TJ UK

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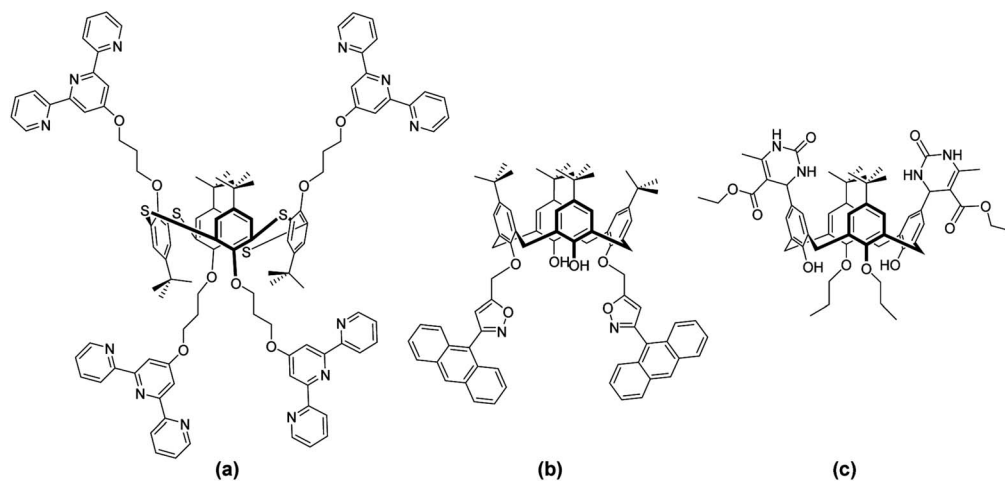


Fig. 1 Heterocycle derivatised (thia)calixarenes.

aldehydes with subsequent oxidation of benzyl alcohol by  $\text{MnO}_2$ ,<sup>36</sup> alkylation of benzaldehydes by phenylalkynyl iodides in the presence of iridium,<sup>37</sup> and cross-coupling of carboxylic acid chloroanhydrides with terminal aryl and heteroalkynes,<sup>38</sup> to name just a few. Despite this to-date, only a single example of a calixarene ynone has been reported, where upper-rim functionalisation was achieved through coupling of an aldehyde with a Grignard salt in the presence of Jones reagent.<sup>39</sup> Functionalisation of the more synthetically useful lower-rim of calixarenes has not been reported.

In this study, the per functionalisation of the lower-rim of calix- and thiacalix[4]arenes with ynones has been investigated through derivatisation of alkyne precursors. Alkynyl derivatives of (thia)calixarenes have been extensively exploited for click chemistry and are readily accessible in a range of conformations.<sup>40</sup>

The presence of sulphide bridges, in thiacalixarenes, creates an additional challenge to functionalisation and necessitates the use of milder conditions. In particular, the sulphur bridge in thiacalixarenes excludes the possibility of general ynone synthesis using organolithium reagents and  $\text{MnO}_2$  oxidation. Additionally, the sulphur atoms may coordinate palladium and copper and require use of higher quantities in the cross-coupling reactions as catalysts along with possible steric hindrance towards the attack on the carbon-carbon triple bond.

We employed various routes to increase the reactivity of the functional groups during ynone formation (Table 1) using **1c** in the 1,3-alternate stereoisomeric form as a model calix[4]arene, due to the larger distance between alkynyl groups compared to cone and partial cone stereoisomers (Scheme S1†). Interestingly, reaction of **1c**<sup>40a</sup> with benzoyl chloride and cyanuric chloride (TCT) in the presence of magnesium bromide as a Lewis acid,<sup>41</sup> to enhance the activity of carbonyl component, was unsuccessful (Table 1). Equally activation of the C-C triple bond with trimethylaluminum<sup>42</sup> proved ineffective. Mixed results were seen with copper complexes,<sup>43,44</sup> with the reaction either not proceeding with the DABCO complex of  $\text{Cu}(\text{I})$ <sup>43</sup> or yielding a hardly separable mixture of the products of cross-

Table 1 Variation of the reaction conditions of **1c** (solvent, catalyst system, temperature of reaction mixture) with carbonyl reagent

Reagent	Catalyst	<i>t</i> , °C	Solvent	Yield
PhCOONa	2 eq. TCT/MgBr <sub>2</sub>	80	MeCN	No reaction
PhC(O)Cl	Me <sub>3</sub> Al	80	TEA/PhCH <sub>3</sub>	No reaction
	Cu(I)	80	TEA/PhCH <sub>3</sub>	— <sup>a</sup>
	Cu(I) + DABCO	80	TEA/PhCH <sub>3</sub>	No reaction
	5% Pd(0) + 10% Cu(I)	20	TEA/THF	Mixture
	Pd(0)	20	TEA/THF	Mixture
	5% Pd(II) + 10% Cu(I)	20	TEA/THF	91% of <b>2c</b>

<sup>a</sup> Mixture of the products of cross-coupling and acetylide Glaser homocoupling.

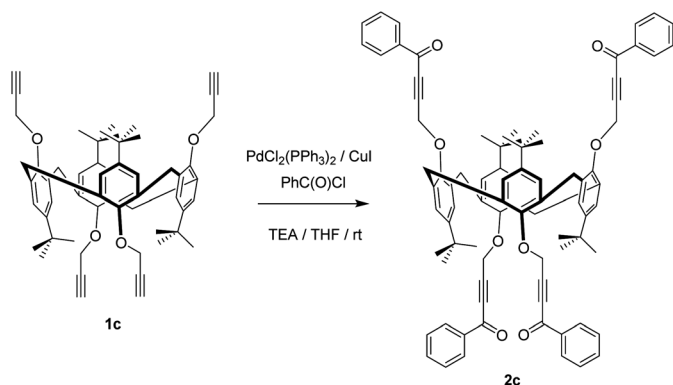
coupling and acetylide Glaser homocoupling in the presence of copper(I) iodide.<sup>44</sup>

In contrast, employment of palladium catalysis successfully yielded the desired ynones. Low reproducibility of the yields was noted when using air- and moisture-sensitive tetrakis(triphenylphosphine)palladium(0) as a catalyst. However, the use of 5% Pd(II) and 10% Cu(I) as the catalytically system in tetrahydrofuran<sup>45</sup> gave reproducibly high yield of the target ynone **2c**. Variation of the reaction time under these conditions showed that full conversion of **1c** is achieved within 1.5 h, while an increase in the time of reaction does not increase the yield.

In summary, Pd(II)/Cu(I) catalyst system is the most reactive among the conditions under study, enabling full conversion of all the alkynyl groups of calixarene **1c** to ynones with a reduced quantity and number of side products compared to copper(I) or Pd(0) catalysis conditions.

Thus, we carried out palladium/copper-catalysed cross-coupling of calix[4]arene in the cone, partial cone, and 1,3-alternate conformations of **1** with benzoyl chloride (an example synthesis is given in Scheme 1) giving **2a**, **b** and **c** in 79%, 88% and 91% yields, respectively. The reaction was extended to the thiacalix[4]arene derivatives **3** from **1d–1g**<sup>46</sup> using the same methodology to give an extended family of derivatives in which

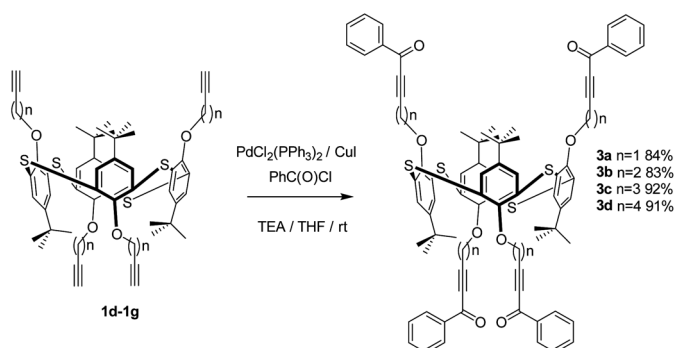




Scheme 1 Example synthesis of ketoacetylene **2c** from alkynyl precursor **1c**.

the distance between the macrocyclic core and the acetylenic function can be varied (Scheme 2).

Successful cross-coupling was indicated by the presence of the characteristic aromatic multiplets of the benzoyl substituents, the loss of the triplet associated with the terminal acetylenic proton and a simplification of the splitting pattern of oxymethylene groups (Fig. 4 and ESI†) in the  $^1\text{H}$  NMR spectra of



Scheme 2 Synthesis of thiocalix[4]arene ketoacetylenes **3**.

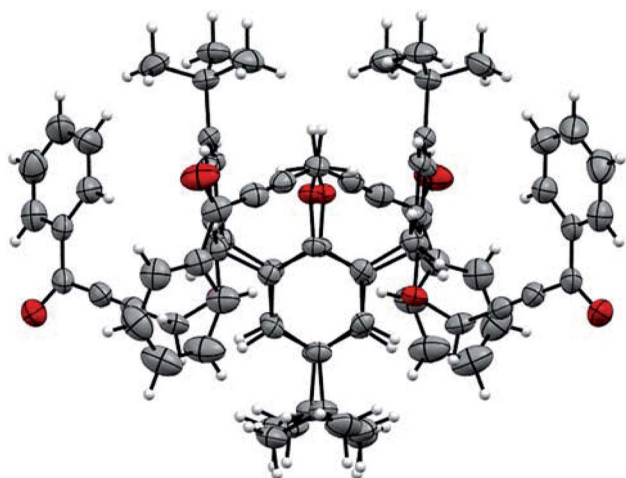


Fig. 2 The molecular geometry of compound **2c**.

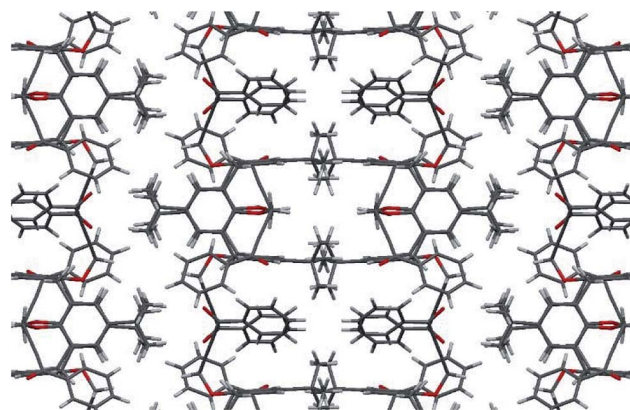


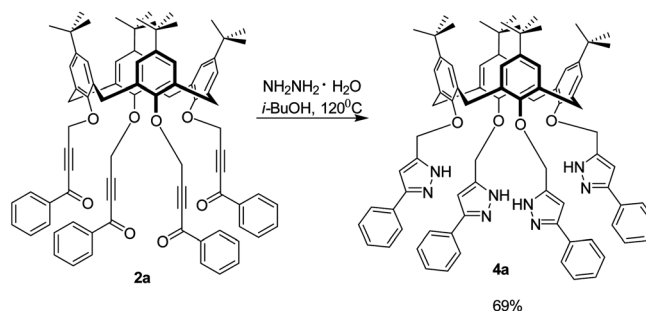
Fig. 3 A fragment of crystal packing of compound **2c**, view along *b* axis.

products **2** and **3**. Full derivatization was confirmed by HRMS data in the presence of CsCl.

The structure of compound **2c** was also determined by X-ray single diffraction (Fig. 2). Compound **2c** crystallizes in space group  $I4_1/a$ , with the asymmetric part of the unit cell containing 1/4 of the molecule. The whole molecule is obtained by a series of symmetry operations, including 4-fold rotoinversion and 2-fold axes, thus **2c** adopts symmetrical 1,3-alternate conformation. The crystal packing is determined by multiple weak  $\text{CH}\cdots\text{O}$  and  $\text{CH}\cdots\pi$  interactions (Fig. 3).

Ketoacetylenes offer diverse cyclization pathways to a range of nitrogen- and oxygen-containing heterocycles<sup>35</sup> with acidic protons including pyrazoles,<sup>47</sup> pyrroles,<sup>48</sup> pyrimidines.<sup>49</sup> Treatment with the highly reactive hydrazine<sup>45</sup> in refluxing isobutanol for 4 h (Scheme 3) to form pyrazoles was investigated on **2**. Surprisingly, in spite of the close arrangement of ynone fragments on the calixarene platform and the possibility of their intramolecular crosslinking by hydrazine molecules, the reaction afforded only the pyrazole derivatives **4**. These could be isolated by simple filtration, in 69–87% yields. Interestingly there is a direct dependence of the yields of pyrazoles with the distance between the ynone with the yield being lowest for the cone and highest for the 1,3-alternate.

Pyrazole ring formation instead of crosslinking could be related to at least one of three factors.



Scheme 3 Example synthesis of pyrazoles **4**.



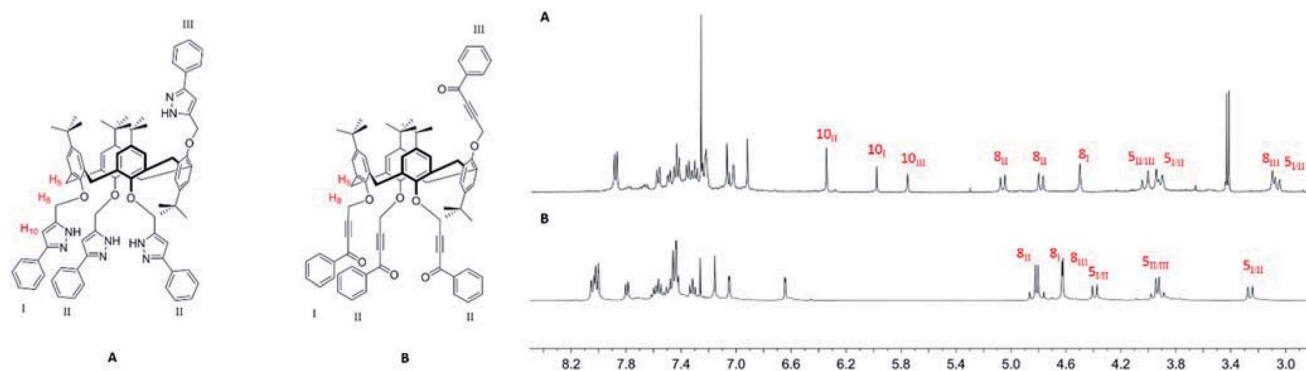


Fig. 4 Partial  $^1\text{H}$  NMR spectra of compounds **4b** (A) and **2b** (B) ( $\text{CDCl}_3$ , 400 MHz, 298 K).

(1) Kinetic effect, where the attack of the  $\text{NH}_2$  fragment on the terminal carbon of the carbon–carbon triple bond is faster than rearrangement of the neighboring carbonyl group from the same or another calixarene molecule.

(2) Thermodynamic effect, in which pyrazole ring formation is entropically more favorable and has a lower energy than any of crosslinked products.

(3) That the amino group in hydrazine is more nucleophilic than the amino group near the imine nitrogen formed after nucleophilic attack of hydrazine on carbonyl group of ynone fragment and dehydration. Assuming a large excess of hydrazine with respect to ynone fragments, this would result in a two-stage mechanism, namely, attack of four hydrazine molecules on the four carbonyl groups of the ynone derivative of calixarene followed by ring closures into four pyrazole fragments.

We postulate that the third factor is the most important as it is less dependent on the temperature of the reaction and concentrations of the reagents than the former two factors.

The structure assignment of pyrazoles **4a** and **4c**, fixed in the cone and 1,3-alternate conformations is straightforward due to high symmetry resulting in equivalency of the phenol rings. (ESI, Fig. S30 and S41 $^\dagger$ ). The key signals for formation of the pyrazole being found at 6.33 and 6.26 ppm, respectively. IR spectra of all the pyrazoles **4** display absorption bands at  $\approx 3350$  and  $\approx 1480\text{ cm}^{-1}$  corresponding to stretching vibrations of N–H and C=N bonds of the pyrazole ring (Fig. S29, S34 and S40 $^\dagger$ ). High-resolution MALDI mass spectra of the pyrazoles **4** mixed with CsCl salt (Fig. S33, S39 and S43 $^\dagger$ ) show the only peak of the product at  $m/z$  1405.5927 (**4a**), 1405.5915 (**4b**), and 1405.6020 (**4c**), while no peaks of the compounds with terminal alkyne groups were observed, which indicates full conversion of the alkyne precursors **2**.

In the case of partial cone stereoisomer **4b**, there is a significantly more complex splitting pattern in the  $^1\text{H}$  NMR spectrum due to the presence of three different phenolic rings (I, II and III) (Fig. 4). The conversion to the pyrazole is accompanied by large shifts in the oxymethylene ( $\text{H}_8$ ) protons and the appearance of three peaks for the pyrazole proton ( $\text{H}_{10}$ ) at 6.34, 5.98 and 5.75 ppm in a 2 : 1 : 1 ratio. Retention of the partial cone conformation is clearly indicated by the presence of two partially overlapped AB quadruplets in the region 4.02–3.90 and at 3.03 ppm. Full characterization was achieved through  $^{13}\text{C}/^1\text{H}$

HMBC, HSQC and described in Suppl. Information (Fig. S37 and S38 $^\dagger$ ).

## Conclusions

An optimal palladium catalysed cross-coupling method has been developed for the preparation of ynones on (thia)calixarenes from the common alkyne precursors. Treatment with hydrazine, as a model nucleophile, yielded the desired pyrazoles in high yield, without inter or intra molecular cross-linking, demonstrating the suitability of the ynone approach for heterocycle formation on these platforms.

## Conflicts of interest

There are no conflicts to declare.

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