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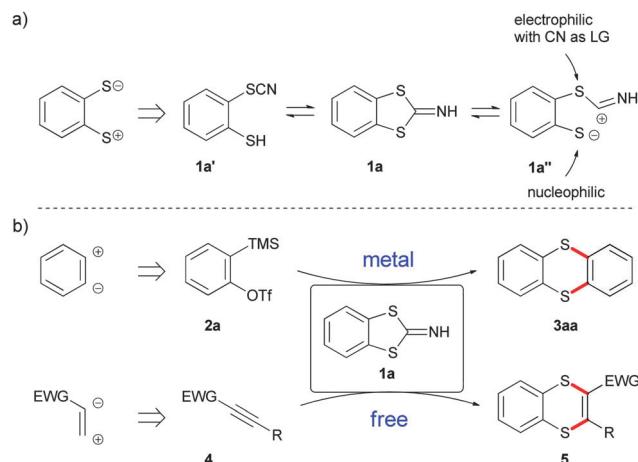
Benzodithioloimines are reacted with arynes or alkynes substituted with electron-withdrawing groups to afford the corresponding thianthrene or benzo[b][1,4]dithiine derivatives. The transformation takes place under mild reaction conditions without any transition metal. Furthermore, the reaction mode could be expanded to 2-thiocyanatopyrroles yielding pyrrolothiazoles.

In the last decades sulphur-containing molecules have received great attention because of their interesting biological properties¹ and their unique application in material science.² For that purpose, the development of new methodologies dealing with the formation of carbon–sulphur bonds has become a major topic in modern organic synthesis.³ Traditional methods which take advantage of the high nucleophilicity of sulphur require harsh reaction conditions to form arene–sulphur bonds. Therefore, elevated temperatures, strong bases or a specific substitution pattern at the arene are needed making them unattractive for numerous target molecules.⁴ These drawbacks could be bypassed utilising metal catalysed cross-coupling reactions for C–S bond formation.⁵ Nevertheless, applications in material science require highly pure compounds without any extraneous metal impurities. Thus, transition metal free protocols for the formation of C–S bonds under mild reaction conditions are of considerable interest.⁶

Regarding the reaction properties of thiophenols, sulphur reacts as a nucleophile. However, its philicity can be reversed by converting thiophenol into the corresponding sulphenyl chloride or thiocyanate. In this shape, sulphur acts as electrophile suitable for substitution by nucleophiles (e.g. lithium or Grignard reagents).⁷ For the facile synthesis of heterocycles containing two sulphur atoms, 1,2-thiophenol derivatives equipped with both abilities are of utmost importance (Scheme 1a). This means one

Exploiting amphiphilicity: facile metal free access to thianthrenes and related sulphur heterocycles†

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Scheme 1 (a) Reaction mode of mononitrile-substituted 1,2-dithiophenol. (b) Possible reaction partners and desired products.

sulphur atom should act as nucleophile, the other one as electrophile. 2-Thiocyanatobenzenethiol (**1a'**) which exists in its closed form **1a** was chosen as target substrate.⁸ Providing the required abilities easily visualised by its open form **1a''**, compound **1a** should be able to react with other easily polarisable or polarised substrates such as arynes⁹ or alkynes substituted with electron-withdrawing (EWG) groups.¹⁰ As products, the corresponding dithiahetocycles **3aa** and **5**, respectively, should be formed (Scheme 1b).

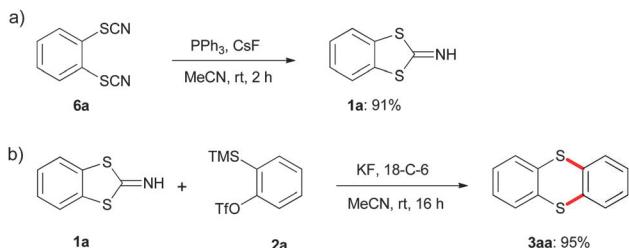
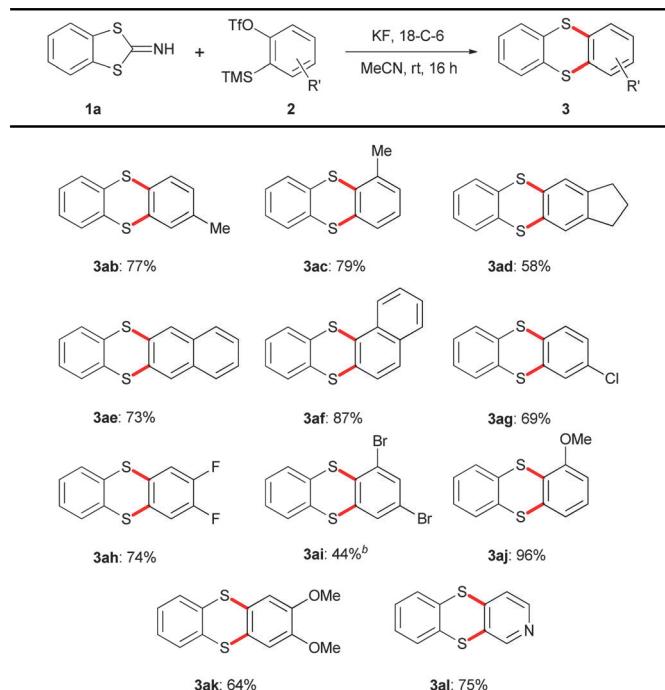
At the beginning of our investigations, we intended to establish a fast and efficient access to key compound **1a** since the methods described in the literature proved to be tedious.⁸ We discovered, when treating aromatic 1,2-dithiocyanatobenzene **6a** with PPh_3 in the presence of base in acetonitrile, **1a** is formed in up to 91% yield within 1 h at room temperature (Scheme 2a). With compound **1a** in hand, we proved the amphiphilic behaviour of the two sulphur atoms. Based on our previous considerations, we chose a common aryne precursor for the desired metal free transformation which would provide a novel entry to the thianthrene scaffold.¹¹ To our delight, common

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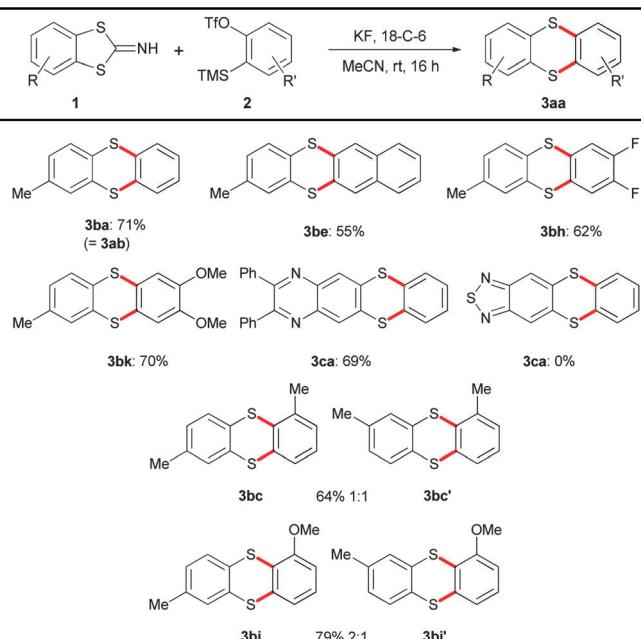
Scheme 2 (a) Synthesis of **1a**. (b) Reaction of **1a** with aryne precursor **2a**.Table 1 Scope of thianthrenes with respect to aryne^a

^a Reaction conditions: **1a** (50 μmol), **2** (75 μmol), KF (150 μmol), 18-C-6 (150 μmol), MeCN (2 mL), rt, 16 h. ^b 200 μmol of **2** were used.

conditions using KF and 18-crown-6 in acetonitrile at ambient temperature yielded the desired thianthrene **3aa** in excellent 95% yield (Scheme 2b).

Exposition of **1a** to various aryne precursors under the above-mentioned reaction conditions yielded the corresponding thianthrenes (Table 1). Alkyl-substituted arynes and such with extended π -systems were successfully converted to their thianthrene counterparts **3ab**–**3af** in 58–87% yield. Chloro-, fluoro-, and bromo-substituted thianthrenes **3ag**, **3ah**, and **3ai** were obtained in yields ranging from 44 to 74%. In the case of the dibromo derivative **3ai** higher loadings of aryne precursor were required. Thianthrenes equipped with one or two methoxy substituents **3aj** and **3ak** were accessed in 64% and 96% yield, respectively. Furthermore, also a pyridine-based aryne precursor was able to undergo the transformation to azathianthrene **3al** in 75% yield.

Similarly, variations with respect to the dithioloimine bearing core are possible (Table 2). Methyl-substituted **1b** in combination with the unfunctionalised aryne precursor was exposed to the

Table 2 Scope of thianthrenes with respect to dithioloimine and aryne^a

^a Reaction conditions: **1** (50 μmol), **2** (75 μmol), KF (150 μmol), 18-C-6 (150 μmol), MeCN (2 mL), rt, 16 h.

reaction conditions yielding **3ba** in 71%. Naphthyl-based **3be**, difluoro- **3bh** and dimethoxy-substituted systems **3bk** were obtained in yields ranging from 55 to 70%. Because of their high electronegativity the latter two substituents render the important in-plane π orbital of the aryne highly electron-deficient. Further, benzodithioloimines containing pyrazine or thiadiazole heterocyclic cores were also applied. Pyrazine derivative **3ca** could be obtained in 69% yield, whereas the formation of **3ca** stayed out.

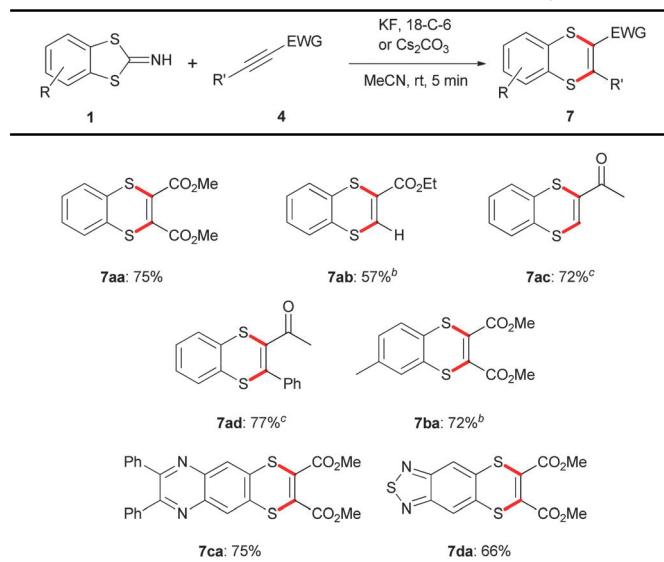
Finally, **1b** was exposed to unsymmetrical aryne precursors substituted with a methyl or a methoxy group. As result, inseparable mixtures of **3bc**/**3bc'** (64%, ratio of 1:1) and **3bj**/**3bj'** (79%, ratio of 2:1) were obtained, respectively. This observation demonstrates that even the methoxy-substituted aryne, known for its high distortion and hence its high regioselectivity in aryne chemistry, does not achieve a satisfying ratio of regioisomers.¹²

In addition, we also tried a one-step procedure applying 1,2-dithiocyanatobenzene **6a** by addition of PPh_3 to the aryne reaction. Unfortunately, the resulting yields were lower with 65% as the best result. A further drawback was the separation of PPh_3 from the product by column chromatography wherefore the two-step procedure proved to be the method of choice.

With this procedure for thianthrenes in hand, we tried to expand the scope to other heterocyclic scaffolds. Therefore, alkynes **4** bearing an electron-withdrawing group were utilised resulting in benzo[*b*][1,4]dithiines.¹³ Slight adjustments of the reaction conditions were necessary to obtain the desired products, but in all cases much faster transformations were observed using these polarised alkynes.¹⁴

Consequently, dimethylester **7aa** and mono-methylester **7ab** were reached in 75% and 57% yields, respectively (Table 3).



Table 3 Scope of dithioloimines with EWG-substituted alkynes^a

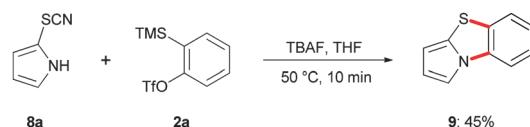
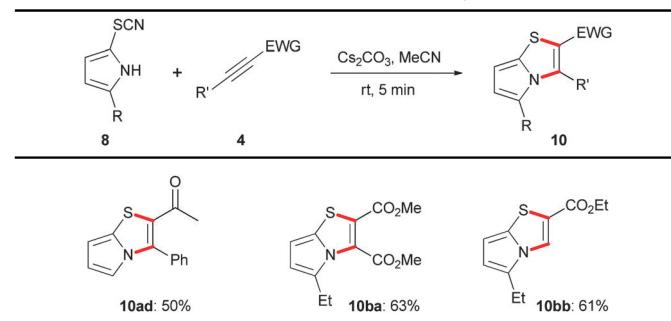
^a Reaction conditions: **1a** (200 μ mol), **4** (300 μ mol), KF (600 μ mol), 18-C-6 (600 μ mol), MeCN (2 mL), rt, 5 h. ^b Reaction conditions: **1** (500 μ mol), **4** (750 μ mol), KF (1.50 mmol), 18-C-6 (1.50 mmol), MeCN (5 mL), rt, 5 h. ^c Cs_2CO_3 (600 μ mol) instead of KF and 18-C-6.

Michael acceptors based on ketones could be converted in yields of 72% for **7ac** and 77% for **7ad**. Modifications regarding the dithioloimine core were performed as well. Methyl-substituted dimethylester derivative **7ba** could be achieved in 72% yield. Also, other heterocyclic systems could be introduced resulting in pyrazine **7ca** and thiadiazol **7da** and in 75% and 66% yield. In addition, the application of Michael acceptors based on double bonds was unsuccessful. We assume that in such a case the retro-Michael reaction in contrast to the ring closure is the preferred mode of action.¹⁵

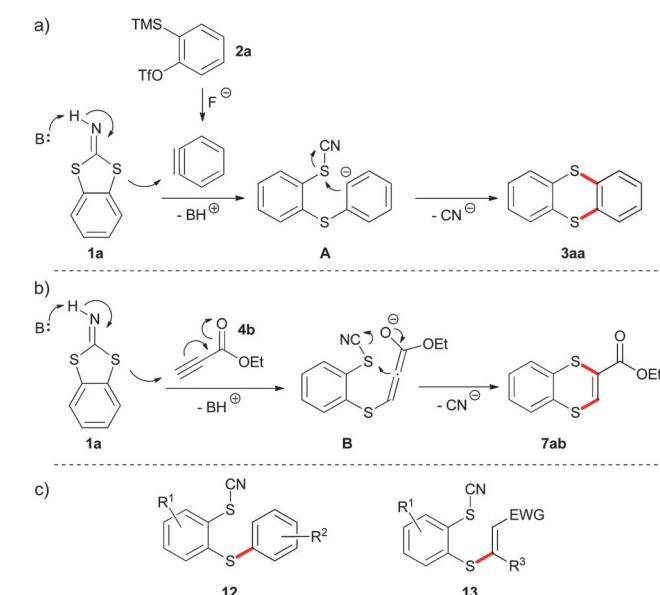
Encouraged by these results, we investigated whether also nitrogen might play the nucleophilic role. Changing the aromatic core to 2-thiocyanatopyrrole the endocyclic nitrogen should be prone to act as nucleophile. As result, the analogous transformation under modified reaction conditions yielded the desired benzo[*d*]pyrrolo[2,1-*b*]thiazole¹⁶ (**9**) in 45% yield (Scheme 3).

Further, also Michael acceptors were shown to be suitable substrates to afford the corresponding pyrrolo[2,1-*b*]thiazole¹⁷ derivatives **10ad**, **10ba** and **10bb** in 50–63% yield (Table 4). Because of the high reactivity and amphiphilic character of **8**, it rather tends to undergo a homocoupling than a 1,4-addition. This problem was partially counteracted by applying higher loadings of Michael acceptor.

We propose the following plausible reaction mechanisms being responsible for product formation (Scheme 4). Precursor **2a** is transformed into the reactive aryne. Deprotonation of

Scheme 3 Reaction of **8a** with aryne precursor **2a**.Table 4 Reaction of **8** with EWG-substituted alkynes **4** to afford **10**^a

^a Reaction conditions: **8** (200 μ mol), **4** (4.00 mmol), Cs_2CO_3 (300 μ mol), MeCN (3 mL), rt, 5 min.



Scheme 4 Proposed mechanistic scenario: (a) for the reaction with arynes; (b) for the reaction with Michael acceptors; (c) observed side products.

benzodithioloimine favours the nucleophilic attack on the aryne leading to intermediate **A** (Scheme 4a). This species undergoes a concomitant nucleophilic substitution on sulphur, whereby nitrile acts as leaving group to form product **3aa**. The pivotal reaction steps for alkynes substituted with electron-withdrawing groups are similar (Scheme 4b). Nevertheless, in this case the formation of the reactive triple bond is not necessary resulting in shorter reaction times in comparison with the previous transformation. The proposed reaction mechanisms are supported by side products **12** and **13** which are formed if ring-closure stays out because of protonation of intermediate **A** or **B** (Scheme 4c).

In conclusion, we developed a broadly applicable transition metal free method for the synthesis of thianthrenes and benzodithiines. For this purpose, amphiphilic dithioloimines containing negatively and positively polarised sulphur were reacted with arynes and alkynes substituted with electron-withdrawing groups. Additionally, the concept of two differently polarised heteroatoms was transferred to a nitrogen–sulphur combination.

The use of 2-thiocyanatopyrroles paves the way for a novel, fast and efficient entry to benzo[*d*]pyrrolo[2,1-*b*]thiazoles and pyrrolo[2,1-*b*]thiazoles, respectively. Further studies to expand this methodology to other heterocyclic scaffolds are ongoing.

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