

# Organic & Biomolecular Chemistry

Accepted Manuscript



This is an *Accepted Manuscript*, which has been through the Royal Society of Chemistry peer review process and has been accepted for publication.

*Accepted Manuscripts* are published online shortly after acceptance, before technical editing, formatting and proof reading. Using this free service, authors can make their results available to the community, in citable form, before we publish the edited article. We will replace this *Accepted Manuscript* with the edited and formatted *Advance Article* as soon as it is available.

You can find more information about *Accepted Manuscripts* in the [Information for Authors](#).

Please note that technical editing may introduce minor changes to the text and/or graphics, which may alter content. The journal's standard [Terms & Conditions](#) and the [Ethical guidelines](#) still apply. In no event shall the Royal Society of Chemistry be held responsible for any errors or omissions in this *Accepted Manuscript* or any consequences arising from the use of any information it contains.



Journal Name

ARTICLE

## Synthesis of Quinoxalines or Quinolin-8-amines from *N*-Propargyl Aniline Derivatives employing Tin and Indium Chlorides

Stefan Aichhorn,<sup>a</sup> Markus Himmelsbach<sup>b</sup> and Wolfgang Schöfberger\*<sup>a</sup>Received 00th January 20xx,  
Accepted 00th January 20xx

DOI: 10.1039/x0xx00000x

www.rsc.org/

Pyrazino compounds such as quinoxalines are 1,4-diazines with widespread occurrence in nature. Quinolin-8-amines are isomerically related and valuable scaffolds in organic synthesis. Herein, we present intramolecular main group metal Lewis acid catalyzed formal hydroamination as well as hydroarylation methodology using mono-propargylated aromatic *ortho*-diamines. The annulations can be conducted utilizing equal aerobic conditions with either stannic chloride or indium(III) chloride and represent primary examples for main group metal catalyzed 6-*exo*-dig and 6-*endo*-dig, respectively, cyclizations in such settings. Both types of reactions can also be utilized in a one-pot manner starting from *ortho*-nitro *N*-propargyl anilines using stoichiometric amounts SnCl<sub>2</sub>·2H<sub>2</sub>O or In powder. Mechanistic considerations are presented regarding the substituent-depending regioselectivity.

### Introduction

Pyrazine motifs<sup>1</sup> are widespread in biological organisms<sup>2</sup> and integrated into versatile chemical structures<sup>3</sup> also with pharmaceutical value.<sup>4</sup> 1,4-Diazines are commonly prepared from alpha-keto carbonyl moieties and 1,2-diamines through twofold imine condensation.<sup>5</sup> The second conventional route is to initiate self-condensation of alpha-amino ketones, often derived from phenacyl halides.<sup>6</sup> These syntheses appear to be quite straightforward cyclizations, but only a few supply a broader substituent scope.<sup>7</sup> Also, routes *via N*-oxides have been utilized to synthesize polycyclic quinoxalines.<sup>8</sup> On the other hand, propargylic amines and related structures found some use for the transition metal mediated construction of aromatic nitrogen heterocycles.<sup>9</sup> Furthermore, quinoline-8-amines, which act as valuable directing groups,<sup>10</sup> ligands for coordination chemistry<sup>11</sup> as well as agents for various diseases,<sup>12</sup> can be considered as structural isomers of methylquinoxalines. Functionalized quinoline platforms found broad interest with organic chemists over decades.<sup>13</sup> In the course of attempting to synthesize *N*<sup>4</sup>-propargylquinolin-3,4-diamine **2** from its 3-nitro precursor **1**, the reduction using stannous chloride dihydrate in ethanol gave a minor amount of **2**, but a considerable fraction of the product mixture was found to consist of an unknown compound. The seemingly mismatching simplicity of its NMR spectra encouraged us to elucidate the structure. It could be verified by HMBC

experiments as well as HR-MS that the resulting assembly appears to be 3-methylpyrazino[2,3-*c*]quinoline **3**. To the best of our knowledge, such 3-alkylated pyrazino[2,3-*c*]quinoline scaffolds and the preparation of pyrazine or quinoxaline derivatives in this fashion have not been reported so far.

### Results and discussion

In order to exploit this observation towards more general application for aromatic heterocycles, we investigated a one-pot procedure for six-membered nitrogen motifs,<sup>14</sup> starting from *ortho*-nitro *N*-propargyl aniline derivatives. Furthermore, we decided to examine whether next to tin also homologous indium species could be facilitated in this context, as indium powder is a reducing agent for nitro groups and In(III) has been demonstrated to activate alkynes.<sup>15</sup> Performing initial screening reactions revealed that the use of methanol as solvent did lead to high quantity of an unidentified side product. It is noteworthy that it was possible to conduct the six-membered nitrogen ring formations in aqueous solution at 85°C, although such reactions resulted in decreased yields. Farther, the same is true for employing toluene as solvent at 100°C. Finally, we have developed two pragmatic protocols for the construction of pyrazines (Fig. 1). Both standard procedures are based on the use of isopropanol as solvent while stirring the reaction mixtures at reflux under aerobic conditions. To obtain the different 2-methylquinoxalines, the best yields were obtained employing 5.5 equivalents of SnCl<sub>2</sub>·2H<sub>2</sub>O. Moving to elementary indium, 3 equivalents in combination with hydrochloric acid gave similar results. With these optimized protocols in hand, we conducted one-pot syntheses of quinoxalines using starting materials **1**, **4**, and **13**. With reaction times ranging from two to 20 hours, derivatives

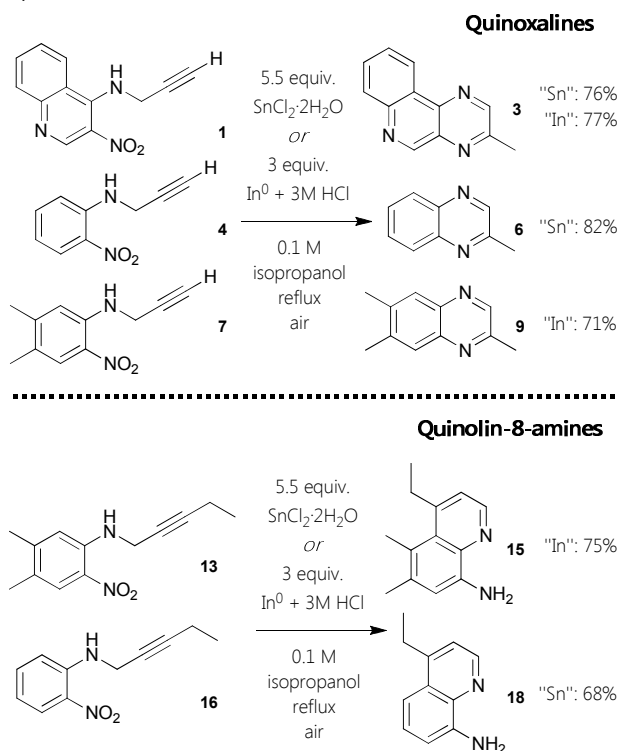
<sup>a</sup> Institute of Organic Chemistry, Johannes Kepler University Linz, Altenberger Straße 69, 4040 Linz, Austria.

\* E-mail: wolfgang.schoefberger@jku.at

<sup>b</sup> Institute of Analytical Chemistry, Johannes Kepler University Linz, Altenberger Straße 69, 4040 Linz, Austria.

Electronic Supplementary Information (ESI) available: [details of any supplementary information available should be included here]. See DOI: 10.1039/x0xx00000x

**3**, **6**, and **15** could be afforded in good yields from **71-82%** (Fig. 1).

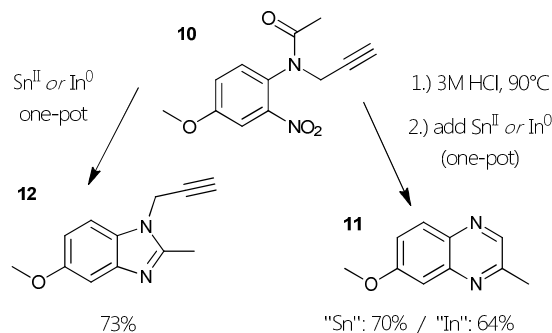


**Fig. 1** Substituent-dependent one-pot synthesis of either pyrazines or quinolin-8-amines.

To explore a variant of this approach with related substrates, readily available anilides as masked anilines were chosen. The monofunctionalization of an anilide appears to be less susceptible compared to primary *ortho*-nitro anilines. An acetyl group as in anisyl-compound **10** can be cleaved under acidic conditions like present in the one-pot procedures. By subjecting the *N*-propargyl anilide to hot hydrochloric acid solution, quantitative deacetylation can be established. When a standard reductive one-pot cyclization-aromatization reaction is conducted with **10**, almost exclusively *N*1-propargyl-C2-methyl-benzimidazole **12** is created. Under these circumstances, reduction of the nitro-group and subsequent condensation of the five-membered ring seems to be faster than the incorporation of the alkyne. In this fashion, functionalized benzimidazoles can be generated in good yield (Fig. 2). For the synthesis of quinoxalines, the protocol had to be adjusted by stirring the alkynyl-anilide in 3 M HCl at 90°C first. Next, by employing our developed standard protocol in the same pot (isopropanol and the respective reducing agent), gave the quinoxaline **11** in good yields.

Soon, it became inevitable to examine whether only terminal  $\beta$ - $\gamma$ -alkynylamines would serve as platform for such formal 6-*exo*-dig cyclizations.<sup>16</sup> To access 3'-alkylated derivatives, the primary *ortho*-nitro arylamines were treated with sodium hydride and 1-bromo-2-pentyne, which gave the internal alkynes **13**, **16**, and **19**. Those were subjected to reflux in

isopropanol under the established aerobic one-pot conditions. At this point, it was surprising to observe complete consumption of educts without creation of pyrazine products.



**Fig. 2** Alkynylanilides as substrates for either propargylic benzimidazoles or quinoxalines.

While the reduced primary amines in *ortho*-position remained not further functionalized, we realized that indeed six-membered rings were synthesized from **13** and **16** (Fig. 1), but in contrast to previous intramolecular annulations, quinoline-8-amines **15** (75%) and **18** (68%) were afforded, formally in a 6-*endo*-dig fashion, exclusively. In case of 3-nitro-*N*-(pent-2-yn-1-yl)quinolin-4-amine, only reduction giving the corresponding 3,4-diamine occurred. These tendencies allow for a distinction between two different types of cascade reactions, which can be conducted employing identical conditions whilst generating unlike heterocycles. When a *para*-diazine compound is generated, the initial cyclization of the reduced intermediate can be considered as intramolecular hydroamination reaction. In contrast, the formation of quinolinamine structures follows a hydroarylation pathway prior to aromatization.

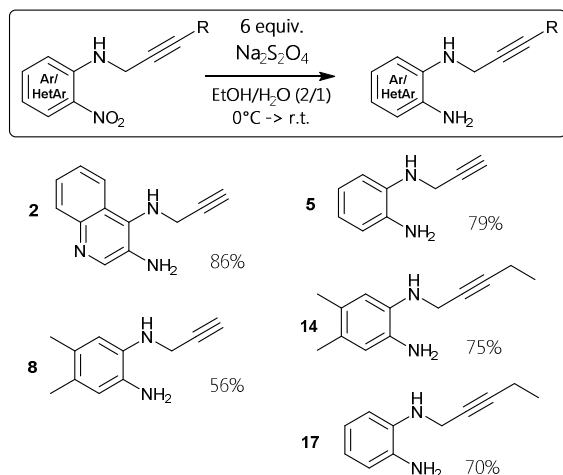
With protocols for specific six-membered rings in hand, we became interested in exploring the mechanistic aspects of this kind of annulation reactions. Preliminary experiments were conducted with  $\beta$ - $\gamma$ -alkynyl-diamines like **2** or **14** exposed to oxygen in isopropanol at 82°C. In the absence of any additives, no conversion was observed. The same was true for the addition of water or stannic chloride. By using less than one equivalent of  $\text{SnCl}_2 \cdot 2\text{H}_2\text{O}$  or  $\text{InCl}_3$ , product formation was achieved analogous to one-pot results. With these promising results, we were interested to demonstrate our synthesis procedures to obtain the six-membered rings in catalytic fashion with well-chosen substrates.

A study of literature reveals that there are a number of related syntheses starting from propargylamine derivatives towards either diaza-compounds (Fig. S1 – left column) or other six-membered rings (Fig. S1 – right column). Gold-catalysis sets the standards in the field due to the advantageous properties of Au-complexes concerning the interactions with C-C triple bonds.<sup>17</sup> Regarding the hydroamination type reactions, a few older reports cover intramolecular conversions attaining quinoxalines similarly to our approach. However, copper and mercury based mediating reagents have to be utilized in

stoichiometric fashion.<sup>18</sup> In this sense, the tin(II) as well as indium(III) promoted cyclizations described here are the first main group metal catalyzed 6-*exo*-dig hydroamination. The situation with electrophilic aromatic substitution reactions is different as there are no reports on vicinal monoalkynylated diamino-substrates, but several transition metal catalyzed annulations giving quinoline derivatives.<sup>19</sup> The best matching related ring formations are In(III) catalyzed syntheses of phenanthrenes and chromenes.<sup>20</sup> Considering this, the Sn(II) and In(III) mediated cyclizations described here are the first examples of main group metal catalyzed 6-*endo*-dig hydroarylation to afford quinoline derivatives.

To provide starting materials for several screening reactions, the nitro precursors had to be reduced to the corresponding diamines in a controlled manner. Therefore, chemoselective transformations of **1**, **4**, **7**, **13**, and **16**, respectively, were conducted employing sodium dithionite as reducing agent to prepare **2**, **5**, **8**, **14**, and **17** (Fig. 3). This proved to be a useful reaction procedure, since the reaction stopped at the *ortho*-diamine stages in reasonable yields.

Concerning the mediating metal species, tin(II) as well as indium(III) could be employed as catalysts (Tab. 1). In contrast to that, Sn(IV) salts and In(0) did not convert the substrates to the desired products. Soon we realized that no product is formed under strictly anaerobic conditions. On the other hand, atmospheric oxygen supply usually proves adequate and O<sub>2</sub>-bubbling is crucial only when it comes to syntheses in NMR tubes. It was observed that fast conversions are initiated above 70°C, hence refluxing at 82°C was chosen as general measure. Moreover, it turned out to be an important finding that a minimum amount of water is necessary for conversion of educts to proceed. With respect to one-pot procedures, which are not quenched in presence of an excess of water, it is noteworthy that water of crystallization from catalyst salts is sufficient for successful conversions.



**Fig. 3** Dithionite as reducing agent to receive *ortho*-diamine compounds for catalytic annulation.

Since a distinctly acidic regime is prevalent in a nitro-one-pot reaction mixture, intensely protic or Brønsted acidic conditions do not inhibit the intramolecular cyclization. In turn, our simple methodology demonstrated that product formation is promoted by Lewis acids, which result in a moderately acidic pH regime.

The overall tendency revealed that a well-working standard protocol consisted of refluxing the isopropanol solution of *N*<sup>1</sup>-β-γ-alkynyl *ortho*-diamine under air with stoichiometric water along with 25–50 mol% main group metal chloride. We reasoned that indium(III) is slightly more effective as catalyst than tin(II). In terms of catalyst loading, complete conversion could not be attained with less than 25 mol%. The best results were obtained with 0.5 equivalents loading, when syntheses could be completed within two to twelve hours. Decreasing the amount of SnCl<sub>2</sub> or InCl<sub>3</sub> would elongate reaction times up to two days. The highest yields with both catalyst systems were achieved by refluxing **2** overnight using 50 mol% metal chloride affording **3** in 91% (SnCl<sub>2</sub>·2H<sub>2</sub>O) and 93% (InCl<sub>3</sub>). Results of catalytic hydroamination and hydroarylation reactions are listed in Table 1 (also compare Figures 1, 3, 4).

**Tab. 1** Pyrazine/quinoline syntheses in catalytic fashion.

entry	catalyst	mol%	t / h	s.m. <sup>[a]</sup>	product	yield
1	SnCl <sub>2</sub>	50	12	<b>2</b>	<b>3</b>	91%
2	InCl <sub>3</sub>	50	12	<b>2</b>	<b>3</b>	93%
3	SnCl <sub>2</sub>	50	12	<b>5</b>	<b>6</b>	85%
4	InCl <sub>3</sub>	25	36	<b>8</b>	<b>9</b>	66%
5	SnCl <sub>2</sub>	25	36	<b>17</b>	<b>18</b>	76%
6	InCl <sub>3</sub>	25	36	<b>14</b>	<b>15</b>	80%

<sup>[a]</sup> conditions: 1 eq. starting material (0.1 M isopropanol soln), 82°C, cat., 2 eq. H<sub>2</sub>O, air.

A set of control experiments was conducted: we examined whether Brønsted acid could facilitate ring closure. Therefore, solutions of **5** and **8** were refluxed after the addition tosylic acid, which only lead to the recovery of starting material both with 0.5 and 5 equivalents of TsOH·H<sub>2</sub>O, respectively. To study the behavior of the nitro-compounds, precursor **4** was reacted in combination with a non-reducing metal chloride. During both cases (catalytic and stoichiometric amounts of InCl<sub>3</sub>), no conversion of starting material **4** was observed utilizing our standard reaction protocol. The same was true for the *N*<sup>1</sup>-Boc-protected derivative of **5**. Additionally, 2-(prop-2-yn-1-yloxy)aniline and 4-(pent-2-yn-1-yloxy)quinolin-3-amine were synthesized in order to study the probability of generating oxygen-heterocycles with our main group metal system. None

of the alkynyloxy-compounds was converted to a cyclized scaffold after treatment with catalytic amounts of  $\text{SnCl}_2 \cdot 2\text{H}_2\text{O}$ .

Further insight regarding the reactivity of the substrate-catalyst combinations could be achieved by subjecting simple *N*-(prop-2-yn-1-yl)-*p*-toluidine as well as *N*-(pent-2-yn-1-yl)-*p*-toluidine to standard catalytic conditions with both Sn(II) and In(III), respectively. After two days of stirring, the attempts with terminal alkynes just gave starting materials. On the other hand, reactions employing internal alkynes afforded small amounts of the hydroarylation product 4-ethyl-6-methylquinoline ("Sn": 7%; "In": 9%) besides educt.

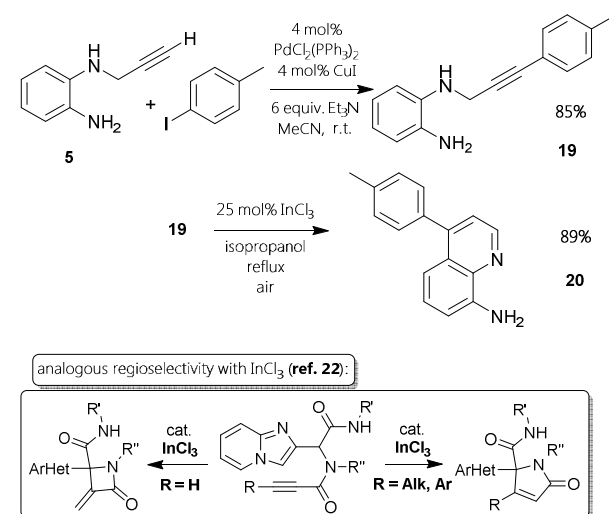
Considering those tendencies, we have performed quantum mechanics calculations to determine the atomic charges and total charge densities (electrostatic potentials, ESPs) of compounds **5** and **17**, respectively. By juxtaposing their relative charges concerning the propargylic moiety, there is a trend that the triple bond is much less polarized at the internal alkyne (Fig. S7, S8). In turn, the  $\beta$ -carbon of the terminal alkyne is clearly more electrophilic.

It is particularly remarkable that catalytic annulations of internal  $N^1$ - $\beta$ - $\gamma$ -alkynyl diamines produce quinoline-8-amines equally exclusive as with the corresponding one-pot attempts. This indicates that the nature of the alkyne moiety decides about the type of ring closure. There does not seem to be a competition between rates of conversion for reduction vs. aromatic substitution reaction, which would have had influenced the given selectivity.

An important remaining aspect was, if aryl-substituents<sup>21</sup> at the alkyne would favour an equal mode of ring closing as ethyl-alkynes. We did not succeed to functionalize *ortho*-nitro precursors like **4** employing Sonogashira cross-coupling methodology. However, we could generate that kind of internal alkynyl-amine by using the reduced compound **5**. It was possible to afford derivative **19** in very good yield (Fig. 4). Subjecting it to standard conditions with 25 mol% of indium(III) chloride for 36 h, 4-*p*-tolylquinolin-8-amine **20** was synthesized in 89% yield. This result evidently demonstrates the tendency that in settings with *ortho*-diamino-scaffolds substituted triple bonds follow a hydroarylation pathway, whereas terminal alkynes undergo hydroamination. This distinction at ring closing reactions with propargylic amine-type substrates catalyzed by  $\text{InCl}_3$  was also described in a recent publication.<sup>22</sup>

As it could be confirmed that the unexpected ring closing reactions, depending on the given substrates, indeed produce either methylpyrazino- or quinoline-8-amino-compounds, the explored set of reactions can therefore be differentiated between 6-*exo*-dig and 6-*endo*-dig annulations, respectively.<sup>23</sup> Under the given conditions, this would involve a Lewis acid (LA) complex to increase the electrophilic character of an alkyne. Metallic Lewis acids can bind weakly in a side-on manner to the triple bond or attach directly to an *sp*-carbon<sup>24</sup>. Besides the more frequent reports on transition<sup>25</sup> or main group<sup>26</sup> metals and iodine<sup>27</sup> to promote *exo*- and *endo*-dig-cyclizations,<sup>28</sup> indium(III) species<sup>29</sup> are able to promote such

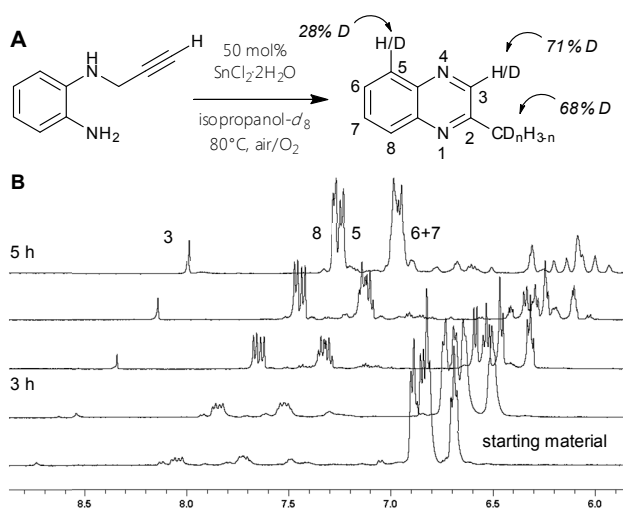
bond formations. The given reaction mixtures seem to be favouring intramolecular hydroaminations<sup>30</sup> or hydroarylations.<sup>31</sup> In order to examine mechanistic aspects of the annulation, syntheses were conducted in deuterated solvent and the mixtures analyzed by means of NMR and HR-MS.



**Fig. 4** Sonogashira coupling for aryl-alkynes, subsequent cyclization, and literature comparison concerning In(III).

The  $^1\text{H}$  NMR spectra of reactions in isopropanol- $d_8$  reveal which hydrogen positions are affected during the procedure. The most striking spectroscopic difference to standard protic condition spectra is the low intensity of two signals: the aryl-singlet of the pyrazine ring as well as the methyl-peak show integrals of less than a third compared to their supposed values. This is true for every successfully tested experiment in deuterated reaction mixture, only the relative amounts of decreased peak area vary specifically with the employed metal salt type. The corresponding  $^{13}\text{C}$  NMR spectra exhibit split signals for those carbons, which hold primarily deuterium instead of protons. Hence, heteronuclear coupling of carbons with  $^2\text{H}$  nuclei is observable ( $^1J_{\text{C-D}} = 27$  Hz). Furthermore, the aromatic peaks in the quinoxaline case appear to display differently overlapping multiplicities and to a minor extent reduced integrals, which can be explained by isotope exchange effects. ESI-mass spectra illustrated a range of product isomers with various deuteration grades. The influence of the alkyne-manipulation and cyclization on the initial aromatic body of the molecule can also be studied by a quinoxaline synthesis in deuterio-solvent<sup>32</sup> (Fig. 5):  $^1\text{H}$  and  $^{13}\text{C}$  NMR spectra indicate changes in peak patterns and intensities for the phenylene scaffold, mass spectra signify considerable amounts of isomers with more than four Ds (Figs. S2–5). Most importantly, this experiment proves that the *ortho*-H position, which gives rise to the electrophilic aromatic substitution pathway, is also activated at substrates with terminal alkynes, but does not afford quinoline product in such cases.

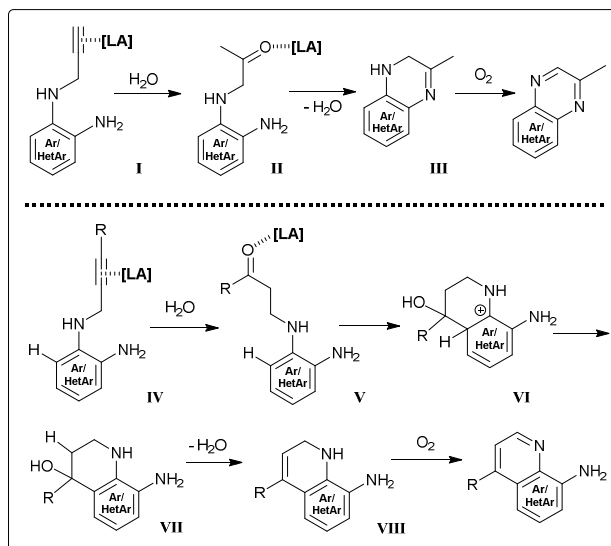
Comprehension of the ring closure might be facilitated by investigating certain screening reactions using **2** at reflux times of only 20 minutes along with catalytic amounts of Sn(II) and In(III) chlorides. After work-up, NMR spectra indicated approx. 50/50 mixtures of pyrazino-product and a species that was determined to be an intermediate. A similar observation was made during NMR-tube reactions of **14** with InCl<sub>3</sub> (resembling the experiment depicted in Fig. 5). Considerable amounts of compound holding a CH<sub>2</sub>-group positioned between a double bond and an NH-group could be detected. HRMS and NMR measurements suggested dihydropyrazine/dihydroquinoline species (scheme 1, structures III/VIII), which is likely to be a somehow stable structure prior to oxidation of the C–N bond. We were not able to isolate these intermediates by column chromatography through silica, as the whole reduced species were converted into final product. The observation of these intermediates allows better understanding of the isotope distribution effects in deuterated media. A range of tautomeric equilibria might foster H/D exchange. Indium(III) and tin(II) chlorides lower the signal intensity of the methylene group ( $HC=N$  in products) down to values between 15–30%.



**Fig. 5** A) Reaction of **5** to **6** with 50 mol% SnCl<sub>2</sub>·2H<sub>2</sub>O in isopropanol-*d*<sub>8</sub> to study the degree of activation on certain positions. B) Time-course <sup>1</sup>H NMR spectra of A).

The need of molecular oxygen for the progress of the reaction can clearly be related to the oxidation of the 1,2-dihydro compounds (Scheme 1) to the final product. The gain of stability by reaching aromaticity explains the fast conversion of the cyclized skeleton. No evidence was found for the formation of an allene prior to ring closure.<sup>33</sup> The prerequisite of water for the reactions to proceed might favour a hydration step prior to ring closure. Both tin and indium are known to be able to facilitate hydration reactions of alkynes to generate ketones.<sup>34</sup> Those carbonyl intermediates could additionally be activated by Lewis acids. In a quinoxaline case, a structure like

**II** may give rise to the condensation resulting in dihydro-compound **III** (Scheme 1, Fig. S2).



**Scheme 1** Proposed mechanisms of the annulation reactions.

Applying this concept to a hydroarylation reaction, the hydration step (**IV** to **V**) would create the ketone at  $\gamma$ -position. This would then be attacked in an S<sub>E</sub>Ar manner (**V** to **VI**), followed by rearomatization and water elimination (**VI** to **VII** to **VIII**). However, there is no evidence for such reactivity, although it would serve as credible explanation for the extensive deuteration also (Figs. S4–S5). For all those reasons mentioned, the course of the reactions might be covered by the sequence of steps depicted in scheme 1.

What appears to be of special importance for the conception of a mechanism and the composition of the Lewis acid catalyst is the lack of reactivity in cases of nitro-precursors like **4** in combination with metal chlorides. The aromatic *ortho*-diamines are well suitable to act as bidentate ligand. As a consequence, we wish to emphasize that there could be deciding influence of a metal cation centred complex, involving an *ortho*-phenylene diamine type ligand,<sup>35</sup> on mediating the cyclization steps.

## Conclusions

We have demonstrated the utility of stannous chloride as well as indium powder under acidic conditions for intramolecular one-pot synthesis of either methylquinoxaline or quinolin-8-amine derivatives. Starting from *ortho*-nitro-*N*-propargylaniline compounds, selectivity is dependent on the substitution of the alkyne. It was shown that such aerobic annulation reactions are suitable to be conducted in catalytic fashion employing tin(II) and indium(III) chlorides as Lewis acids starting from *ortho*-diamino compounds. Those are the first examples of 6-

*exo*-dig hydroaminations and 6-*endo*-dig hydroarylations with alkylnylanilines, respectively, catalyzed by main group metals.

## Experimental part

**General procedure for one-pot syntheses of quinoxalines or quinolin-8-amines:** SnCl<sub>2</sub>·2H<sub>2</sub>O (5.5 equiv.) or indium powder (3 equiv. - in combination with hydrochloric acid solution (3 M; 15 equiv.)) were added to a stirred 0.15 M solution of *ortho*-nitro *N*-propargylaniline compound (1 equiv.) in 2-propanol at room temperature. The mixture was heated at reflux under air until judged completed as indicated by thin layer chromatography. The mixture was brought to room temperature, filtrated and the filtrate concentrated under reduced pressure. The residue was quenched with saturated NaHCO<sub>3</sub> solution until pH 10. The aqueous solution was extracted with EtOAc three times, the combined organic layers were dried over anhydrous Na<sub>2</sub>SO<sub>4</sub> and evaporated under reduced pressure. The residue was purified by column chromatography (silica gel, heptanes/EtOAc = 3 / 1).

**General procedure for main group metal chloride catalyzed formation of quinoxalines or quinolin-8-amines:** 25–50 mol% of stannous chloride dihydrate or indium(III) chloride were added to a stirred 0.1 M solution of *ortho*-amino *N*-propargylaniline compound (1 equiv.) in 2-propanol at room temperature. Unless water of crystallization was present, 1 equiv. H<sub>2</sub>O was added. The mixture was heated at reflux under air until judged completed as indicated by thin layer chromatography. The mixture was concentrated under reduced pressure and the residue quenched with saturated NaHCO<sub>3</sub> solution until pH 10. The aqueous solution was extracted with EtOAc three times, the combined organic layers were dried over anhydrous Na<sub>2</sub>SO<sub>4</sub> and evaporated under reduced pressure. The residue was purified by column chromatography (silica gel, heptanes/EtOAc = 3 / 1).

**General procedure for alkylation of *ortho*-nitro anilines using NaH:** sodium hydride (95%, 1.1 equiv.) was added to a 0.2 M acetonitrile solution of 2-nitroaniline compound (1 equiv.). The mixture was stirred at 0°C for 15 min. Then, 1-bromoalk-2-yne (1.1 equiv.) was added and the mixture was brought to room temperature. It was thoroughly stirred under N<sub>2</sub> atmosphere for 40 h. The suspension was quenched with H<sub>2</sub>O and saturated NaHCO<sub>3</sub> solution was added until pH 10. The resulting aqueous layer was extracted with DCM three times, the combined organic layers were dried over anhydrous Na<sub>2</sub>SO<sub>4</sub> and evaporated under reduced pressure. The residue was purified by column chromatography (silica gel, heptanes/EtOAc or MeOH/DCM).

**General procedure for the reduction of *ortho*-nitro *N*-alkynylanilines to diamines using Na<sub>2</sub>S<sub>2</sub>O<sub>4</sub>:** to a stirred 0.1 M EtOH solution of 2-nitro-*N*-alk-2'-ynylaniline compound (1 equiv.) at 0°C were added portions of sodium dithionite (85%, 6 equiv.) and H<sub>2</sub>O (half volume of EtOH) alternately. Then the mixture was allowed to warm to room temperature and stirred for 1 h. Then it was filtrated and the bulk of aqueous EtOH of the filtrate was evaporated *in vacuo*. Under reduced pressure, quick column chromatographic purification

of the residue over a short pad of silica in a sintered glass filter was performed using a solvent gradient from DCM/MeOH (10 / 1) to pure MeOH. The polar fractions were collected and concentrated *in vacuo*.

**3-methylpyrazino[2,3-*c*]quinoline (3):** pale orange solid, m.p. 120–122°C; <sup>1</sup>H NMR (700 MHz, CDCl<sub>3</sub>, 298 K): δ = 2.88 (s, 3H), 7.81 (t, *J* = 7.5 Hz, 1H), 7.90 (t, *J* = 7.6 Hz, 1H), 8.27 (d, *J* = 8.1 Hz, 1H), 9.01 (s, 1H), 9.07 (d, *J* = 8.1 Hz, 1H), 9.52 (s, 1H) ppm; <sup>13</sup>C NMR (176 MHz, CDCl<sub>3</sub>, 298 K): δ = 22.4, 123.5, 124.7, 128.2, 129.6, 130.6, 135.4, 141.4, 146.0, 148.6, 153.9, 155.2 ppm; IR (film):  $\bar{\nu}$  = 2921, 2853, 1458, 1376, 951, 773 cm<sup>-1</sup>; HRMS (ESI): *m/z* calcd for C<sub>12</sub>H<sub>9</sub>N<sub>3</sub>: 196.0869 [M+H]<sup>+</sup>; found: 196.0870.

**7-methoxy-2-methylquinoxaline (11):** to *N*-alkynyl-*ortho*-nitro-anilide (1 equiv.) was added 3 M hydrochloric acid solution (30 equiv.). The mixture was stirred at 90°C under N<sub>2</sub> atmosphere for 7 h. Then, the mixture was directly subjected to one-pot reaction conditions as described in the General procedure. Physical and spectroscopic properties are in accordance with published data.<sup>36</sup> <sup>1</sup>H NMR (300 MHz, CDCl<sub>3</sub>, 298 K): δ = 2.74 (s, 3H), 3.96 (s, 3H), 7.31 (d, *J* = 2.7 Hz, 1H), 7.35 (dd, *J* = 2.7 Hz, 9.1 Hz, 1H), 7.94 (d, *J* = 9.1 Hz, 1H), 8.59 (s, 1H) ppm; HRMS (ESI): *m/z* calcd for C<sub>10</sub>H<sub>10</sub>N<sub>2</sub>O: 175.0866 [M+H]<sup>+</sup>; found: 175.0862.

**5-methoxy-2-methyl-1-(prop-2-yn-1-yl)-1H-benzo[*d*]imidazole (12):** brown solid, m.p. 110–116°C; eluent DCM/MeOH = 7 / 1; <sup>1</sup>H NMR (300 MHz, CDCl<sub>3</sub>, 298 K): δ = 2.34 (t, *J* = 2.5 Hz, 1H), 2.62 (s, 3H), 3.84 (s, 3H), 4.80 (d, *J* = 2.5 Hz, 2H), 6.90 (dd, *J* = 2.4 Hz, 8.8 Hz, 1H), 7.18 (d, *J* = 2.4 Hz, 1H), 7.25 (d, *J* = 8.8 Hz, 1H) ppm; <sup>13</sup>C NMR (75 MHz, CDCl<sub>3</sub>, 298 K): δ = 13.9, 33.3, 56.0, 73.6, 76.7 (overlaid by solvent signal), 102.1, 109.5, 112.0, 129.3, 143.4, 151.5, 156.3 ppm; IR (film):  $\bar{\nu}$  = 3194, 2112, 1519, 1489, 1439, 1276, 1198, 1151, 1034, 787 cm<sup>-1</sup>; HRMS (ESI): *m/z* calcd for C<sub>12</sub>H<sub>12</sub>N<sub>2</sub>O: 201.1022 [M+H]<sup>+</sup>; found: 201.1027.

**4-ethyl-5,6-dimethylquinolin-8-amine (15):** brownish orange oil; <sup>1</sup>H NMR (700 MHz, CDCl<sub>3</sub>, 298 K): δ = 1.33 (t, *J* = 7.5 Hz, 3H), 2.40 (s, 3H), 2.62 (s, 3H), 3.25 (q, *J* = 7.5 Hz, 2H), 6.82 (s, 1H), 7.18 (d, *J* = 4.4 Hz, 1H), 8.55 (d, *J* = 4.4 Hz, 1H) ppm; <sup>13</sup>C NMR (176 MHz, CDCl<sub>3</sub>, 298 K): δ = 16.2, 18.6, 22.1, 29.8, 113.3, 120.0, 122.7, 128.7, 135.7, 138.4, 142.1, 145.7, 150.9 ppm; IR (film):  $\bar{\nu}$  = 2922, 2853, 1519, 1459, 1325, 1260, 1013, 797 cm<sup>-1</sup>; HRMS (ESI): *m/z* calcd for C<sub>13</sub>H<sub>16</sub>N<sub>2</sub>: 201.1386 [M+H]<sup>+</sup>; found: 201.1386.

**N<sup>1</sup>-(pent-2-yn-1-yl)benzene-1,2-diamine (17):** brown oil; <sup>1</sup>H NMR (300 MHz, CDCl<sub>3</sub>, 298 K): δ = 1.14 (t, *J* = 7.5 Hz, 3H), 2.21 (qt, *J* = 2.2 Hz, 7.5 Hz, 2H), 3.44 (br s, 3H), 3.89 (t, *J* = 2.2 Hz, 2H), 6.70–6.79 (m, 3H), 6.80–6.88 (m, 1H) ppm; <sup>13</sup>C NMR (75 MHz, CDCl<sub>3</sub>, 298 K): δ = 12.5, 14.0, 34.5, 76.5, 85.4, 113.0, 116.5, 119.8, 120.6, 135.1, 136.7 ppm; IR (film):  $\bar{\nu}$  = 3334, 3042, 1573, 1506, 1455, 1315, 1269, 738 cm<sup>-1</sup>; HRMS (ESI): *m/z* calcd for C<sub>11</sub>H<sub>14</sub>N<sub>2</sub>: 175.1230 [M+H]<sup>+</sup>; found: 175.1232.

**N<sup>1</sup>-(3-(*p*-tolyl)prop-2-yn-1-yl)benzene-1,2-diamine (20):** prepared in analogy to literature.<sup>37</sup> To a solution of 5 (0.68 mmol, 99 mg, 1 equiv.) in 4 ml acetonitrile were added 1-iodo-4-methylbenzene (0.75 mmol, 165 mg, 1.1 equiv.),

triethylamine (4.1 mmol, 566  $\mu$ l, 6 equiv.), copper(I) iodide (0.026 mmol, 5 mg, 0.04 equiv.), and bis(triphenylphosphine)palladium(II) dichloride (0.026 mmol, 19 mg, 0.04 equiv.). The mixture was stirred under N<sub>2</sub> atmosphere at room temperature for 20 h. Then it was filtrated and washed with EtOAc. The filtrate was quenched with water and saturated NaHCO<sub>3</sub> solution was added until pH 10. It was extracted with EtOAc three times, the combined organic layers were dried over anhydrous Na<sub>2</sub>SO<sub>4</sub> and evaporated under reduced pressure. The residue was purified by column chromatography (silica gel, heptanes/EtOAc = 3 / 1) and gave **20** as a light brown oil (0.578 mmol, 137 mg, 85%). <sup>1</sup>H NMR (300 MHz, CDCl<sub>3</sub>, 298 K):  $\delta$  = 2.34 (s, 3H), 3.49 (br s, 3H), 4.14 (s, 2H), 6.73–6.89 (m, 4H), 7.10 (d, *J* = 8.0 Hz, 2H), 7.31 (d, *J* = 8.0 Hz, 2H) ppm; <sup>13</sup>C NMR (75 MHz, CDCl<sub>3</sub>, 298 K):  $\delta$  = 21.6, 35.1, 83.8, 85.9, 113.4, 116.6, 120.0, 120.1, 120.6, 129.1 (2C), 131.7 (2C), 135.3, 136.6, 138.4 ppm; IR (film):  $\bar{\nu}$  = 3334, 2920, 1597, 1508, 1450, 1271, 816, 740 cm<sup>-1</sup>; HRMS (ESI): *m/z* calcd for C<sub>16</sub>H<sub>16</sub>N<sub>2</sub>: 237.1386 [M+H]<sup>+</sup>; found: 237.1384.

**4-(*p*-tolyl)quinolin-8-amine (21):** pale yellow oil; <sup>1</sup>H NMR (700 MHz, CDCl<sub>3</sub>, 298 K):  $\delta$  = 2.46 (s, 3H), 5.05 (br s, 2H), 6.94 (dd, *J* = 1.9 Hz, 6.7 Hz, 1H), 7.24–7.30 (m, 3H), 7.32 (d, *J* = 7.9 Hz, 2H), 7.41 (d, *J* = 7.9 Hz, 2H), 8.77 (d, *J* = 4.4 Hz, 1H) ppm; <sup>13</sup>C NMR (176 MHz, CDCl<sub>3</sub>, 298 K):  $\delta$  = 21.3, 109.9, 114.3, 121.7, 127.2, 127.4, 129.1 (2C), 129.4 (2C), 135.7, 138.1, 138.8, 144.2, 146.9, 148.4 ppm; IR (film):  $\bar{\nu}$  = 3447, 3319, 2360, 1618, 1502, 1358, 1282, 818, 761 cm<sup>-1</sup>; HRMS (ESI): *m/z* calcd for C<sub>16</sub>H<sub>14</sub>N<sub>2</sub>: 235.1230 [M+H]<sup>+</sup>; found: 235.1230.

## Acknowledgements

WS kindly acknowledges the financial support of the project 836532 ("Prä-klinische Entwicklung einer Off-the-Shelf individualisierten Krebsimmuntherapie") by the Austrian Research Promotion Agency (FFG). The NMR spectrometers were acquired in collaboration with the University of South Bohemia (CZ) with financial support from the European Union through the EFRE INTERREG IV ETC-AT-CZ program (project M00146, "RERI-uasb").

## Notes and references

- a) I. J. Krems and P. E. Spoerri, *Chem. Rev.*, 1947, **40**, 279–358; b) W. Kaim, *Angew. Chem.*, 1983, **95**, 201–221;
- a) R. Müller and S. Rappert, *Appl. Microbiol. Biotechnol.*, 2010, **85**, 1315–1320; b) P. Sjödin, H. Wallin, J. Alexander and M. Jägerstad, *Carcinogenesis*, 1989, **10**, 1269–1275;
- a) V. P. Mehta, A. Sharma, K. van Hecke, L. van Meervelt and Van der Eycken, Erik V., *J. Org. Chem.*, 2008, **73**, 2382–2388; b) S. G. Modha, J. C. Trivedi, V. P. Mehta, D. S. Ermolat'ev and Van der Eycken, Erik V., *J. Org. Chem.*, 2011, **76**, 846–856; c) A. Zaid, J.-S. Sun, C.-H. Nguyen, E. Bisagni, T. Garestier, D. S. Grierson and R. Zain, *ChemBioChem*, 2004, **5**, 1550–1557;
- a) S. B. Ferreira and C. R. Kaiser, *Expert Opin. Ther. Patents*, 2012, **22**, 1033–1051; b) J. Guillon, I. Forfar, M. Mamani-

- Matsuda, V. Desplat, M. Saliège, D. Thiolat, S. Massip, A. Tabourier, J.-M. Léger, B. Dufaure, G. Haumont, C. Jarry and D. Mossalayi, *Bioorg. Med. Chem.*, 2007, **15**, 194–210;
- a) F. Zhang, Y. Xi, Y. Lu, L. Wang, L. Liu, J. Li and Y. Zhao, *Chem. Commun.*, 2014, **50**, 5771–5773; b) H. A. Oskooie, M. M. Heravi, K. Bakhtiari and S. Taheri, *Monatsh. Chem. Chem. Mon.*, 2007, **138**, 875–877; c) N. Xekoukoulotakis, C. Hadjiantoniou-Maroulis and A. Maroulis, *Tetrahedron Lett.*, 2000, **41**, 10299–10302; d) C. Qi, H. Jiang, L. Huang, Z. Chen and H. Chen, *Synthesis*, 2011, 387–396; e) P. Ghosh and A. Mandal, *Green Chem. Lett. Rev.*, 2012, **5**, 127–134;
  - a) F. Buron, A. Turck, N. Plé, L. Bischoff and F. Marsais, *Tetrahedron Lett.*, 2007, **48**, 4327–4330; b) G. Liu, G. Xu, J. Li, D. Ding and J. Sun, *Org. Biomol. Chem.*, 2014, **12**, 1387–1390; c) Z. Chen, D. Ye, G. Xu, M. Ye and L. Liu, *Org. Biomol. Chem.*, 2013, **11**, 6699–6702;
  - a) C. Venkatesh, B. Singh, P. K. Mahata, H. Ila and H. Junjappa, *Org. Lett.*, 2005, **7**, 2169–2172; b) Z.-J. Yang, C.-Z. Liu, B.-L. Hu, C.-L. Deng and X.-G. Zhang, *Chem. Commun.*, 2014, **50**, 14554–14557;
  - a) F. Chen, X. Huang, X. Li, T. Shen, M. Zou and N. Jiao, *Angew. Chem.*, 2014, **126**, 10663–10667; b) Viswanadham, K K Durga Rao, M. Prathap Reddy, P. Sathyanarayana, O. Ravi, R. Kant and S. R. Bathula, *Chem. Commun.*, 2014, **50**, 13517–13520;
  - a) D. D. Vachhani, V. P. Mehta, S. G. Modha, K. van Hecke, L. VanMeervelt and Van der Eycken, Erik V., *Adv. Synth. Catal.*, 2012, **354**, 1593–1599; b) Q. Yang, T. Xu and Z. Yu, *Org. Lett.*, 2014, **16**, 6310–6313; c) D. D. Vachhani, S. G. Modha, A. Sharma and Van der Eycken, Erik V., *Tetrahedron*, 2013, **69**, 359–365; d) S. Jalal, K. Bera, S. Sarkar, K. Paul and U. Jana, *Org. Biomol. Chem.*, 2014, **12**, 1759–1770;
  - a) B. M. Monks, E. R. Fruchey and S. P. Cook, *Angew. Chem. Int. Ed.*, 2014, **53**, 11065–11069; b) V. G. Zaitsev, D. Shabashov and O. Daugulis, *J. Am. Chem. Soc.*, 2005, **127**, 13154–13155;
  - C. E. Meyet and C. H. Larsen, *J. Org. Chem.*, 2014, **79**, 9835–9841.
  - A. Shi, T. A. Nguyen, S. K. Battina, S. Rana, D. J. Takemoto, P. K. Chiang and D. H. Hua, *Bioorg. Med. Chem. Lett.*, 2008, **18**, 3364–3368.
  - a) M. Rehan, G. Hazra and P. Ghorai, *Org. Lett.*, 2015; b) X. Zhang, M. A. Campo, T. Yao and R. C. Larock, *Org. Lett.*, 2005, **7**, 763–766; c) G. A. Salman, R. U. Nisa, V. O. Iaroshenko, J. Iqbal, K. Ayub and P. Langer, *Org. Biomol. Chem.*, 2012, **10**, 9464–9473;
  - a) G. S. Welmaker, J. A. Nelson, J. E. Sabalski, A. L. Sabb, J. R. Potoski, D. Graziano, M. Kagan, J. Coupet, J. Dunlop, H. Mazandarani, S. Rosenzweig-Lipson, S. Sukoff and Y. Zhang, *Bioorg. Med. Chem. Lett.*, 2000, **10**, 1991–1994; b) Pereira, Maria de Fatima and V. Thiéry, *Org. Lett.*, 2012, **14**, 4754–4757;
  - a) N. Rajesh, R. Sarma and D. Prajapati, *Synlett*, 2014, **25**, 1448–1452; b) K. Surendra, W. Qiu and E. J. Corey, *J. Am. Chem. Soc.*, 2011, **133**, 9724–9726; c) Y. Zhang, P. Li, M. Wang and L. Wang, *J. Org. Chem.*, 2009, **74**, 4364–4367;



- 16 K. Gilmore and I. V. Alabugin, *Chem. Rev.*, 2011, **111**, 6513–6556.
- 17 a) T. S. Symeonidis, I. N. Lykakis and K. E. Litinas, *Tetrahedron*, 2013, **69**, 4612–4616; b) S. Basceken and M. Balci, *J. Org. Chem.*, 2015, **80**, 3806–3814; c) B. Alcaide, P. Almendros, J. M. Alonso, I. Fernández, G. Gómez-Campillos and M. R. Torres, *Chem. Commun.*, 2014, **50**, 4567–4570;
- 18 a) J. Barluenga, F. Aznar, R. Liz and M.-P. Cabal, *Synthesis*, 1985, 313–314; b) R. Mukhopadhyay and N. G. Kundu, *Tetrahedron Lett.*, 2000, **41**, 9927–9930; c) S. Knapp, J. Ziv and J. D. Rosen, *Tetrahedron*, 1989, **45**, 1293–1298;
- 19 a) H. Murase, K. Senda, M. Senoo, T. Hata and H. Urabe, *Chem. Eur. J.*, 2014, **20**, 317–322; b) K. C. Majumdar, R. Kumar Nandi, S. Ganai and A. Taher, *Synlett*, 2011, 116–120; c) B. Roy, I. Ansary, S. Samanta and K. C. Majumdar, *Tetrahedron Lett.*, 2012, **53**, 5119–5122;
- 20 a) Y. Kwon, H. Cho and S. Kim, *Org. Lett.*, 2013, **15**, 920–923; b) L. Alonso-Marañón, M. M. Martínez, L. A. Sarandeses and J. P. Sestelo, *Org. Biomol. Chem.*, 2015, **13**, 379–387;
- 21 X. Zhang, T. Yao, M. A. Campo and R. C. Larock, *Tetrahedron*, 2010, **66**, 1177.
- 22 Z. Li, U. K. Sharma, Z. Liu, N. Sharma, J. N. Harvey and Van der Eycken, Erik V., *Eur. J. Org. Chem.*, 2015, **2015**, 3957–3962.
- 23 a) M. J. Gainer, N. R. Bennett, Y. Takahashi and R. E. Looper, *Angew. Chem.*, 2011, **123**, 710–713; b) C. Shu, L. Li, C.-B. Chen, H.-C. Shen and L.-W. Ye, *Chem. Asian J.*, 2014, **9**, 1525–1529; c) N. T. Patil, A. Nijamudheen and A. Datta, *J. Org. Chem.*, 2012, **77**, 6179–6185;
- 24 L. Li, G. Huang, Z. Chen, W. Liu, X. Wang, Y. Chen, L. Yang, W. Li and Y. Li, *Eur. J. Org. Chem.*, 2012, 5564–5572.
- 25 a) T. E. Müller and M. Beller, *Chem. Rev.*, 1998, **98**, 675–704; b) D. Y. Li, H. J. Chen and P. N. Liu, *Org. Lett.*, 2014;
- 26 F. Pohlki and S. Doye, *Chem. Soc. Rev.*, 2003, **32**, 104–114.
- 27 N. M. Mishra, D. D. Vachhani, S. G. Modha and Van der Eycken, Erik V., *Eur. J. Org. Chem.*, 2013, 693–700.
- 28 a) A. Kumar, Z. Li, S. K. Sharma, V. S. Parmar and Van der Eycken, Erik V., *Chem. Commun.*, 2013, **49**, 6803–6805; b) J. M. French and S. T. Diver, *J. Org. Chem.*, 2014, **79**, 5569–5585; c) A.-L. Girard, T. Enomoto, S. Yokouchi, C. Tsukano and Y. Takemoto, *Chem. Asian J.*, 2011, **6**, 1321–1324;
- 29 a) R. Sarma and D. Prajapati, *Chem. Commun.*, 2011, **47**, 9525–9527; b) J. Augé, N. Lubin-Germain and J. Uziel, *Synthesis*, 2007, 1739–1764;
- 30 a) K. Okuma, J.-i. Seto, K.-i. Sakaguchi, S. Ozaki, N. Nagahora and K. Shioji, *Tetrahedron Lett.*, 2009, **50**, 2943–2945; b) R. Severin and S. Doye, *Chem. Soc. Rev.*, 2007, **36**, 1407–1420; c) A. Ranjan, R. Yerande, P. B. Wakchaure, S. G. Yerande and D. H. Dethe, *Org. Lett.*, 2014, **16**, 5788–5791;
- 31 a) X.-Y. Liu, P. Ding, J.-S. Huang and C.-M. Che, *Org. Lett.*, 2007, **9**, 2645–2648; b) B. Michelet, J.-R. Colard-Itté, G. Thiery, R. Guillot, C. Bour and V. Gandon, *Chem. Commun.*, 2015, **51**, 7401–7404;
- 32 a) N. Yoshikai, S. Asako, T. Yamakawa, L. Ilies and E. Nakamura, *Chem. Asian J.*, 2011, **6**, 3059–3065; b) W. Xie, J. Yang, B. Wang and B. Li, *J. Org. Chem.*, 2014, **79**, 8278–8287;
- 33 a) A. Saito, S. Oda, H. Fukaya and Y. Hanzawa, *J. Org. Chem.*, 2009, **74**, 1517–1524; b) J. K. Vandavasi, W.-P. Hu, G. C. Senadi, Boominathan, Siva Senthil Kumar, H.-Y. Chen and J.-J. Wang, *Eur. J. Org. Chem.*, 2014, 6219–6226;
- 34 a) X. Jin, T. Oishi, K. Yamaguchi and N. Mizuno, *Chem. Eur. J.*, 2011, **17**, 1261–1267; b) E. Bosch and L. Jeffries, *Tetrahedron Lett.*, 2001, **42**, 8141–8142; c) Thuong, Mathieu Bui The, A. Mann and A. Wagner, *Chem. Commun.*, 2012, **48**, 434–436; d) L. Hintermann and A. Labonne, *Synthesis*, 2007, 1121–1150; e) S. Liang, G. B. Hammond and B. Xu, *Chem. Commun.*, 2015, **51**, 903–906;
- 35 M. Normand, V. Dorcet, E. Kirillov and J.-F. Carpentier, *Organometallics*, 2013, **32**, 1694–1709.
- 36 S. Grivas, W. Tian, S. V. Meshkov, H. Lönnberg, M. Dahlqvist, M. M. Kady and S. B. Christensen, *Acta Chem. Scand.*, 1992, **46**, 1109–1113.
- 37 Y. Ichikawa, T. Nishimura and T. Hayashi, *Organometallics*, 2011, **30**, 2342–2348.