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Directed regioselective *ortho,ortho'*-magnesiations of aromatics and heterocycles using *s*Bu₂Mg in toluene†

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Aryl azoles are ubiquitous as bioactive compounds and their regioselective functionalization is of utmost synthetic importance. Here, we report the development of a toluene-soluble dialkylmagnesium base *s*Bu₂Mg. This new reagent allows mild and regioselective *ortho*-magnesiations of various *N*-arylated pyrazoles and 1,2,3-triazoles as well as arenes bearing oxazoline, phosphorodiamidate or amide directing groups. The resulting diarylmagnesium reagents were further functionalized either by Pd-catalyzed arylation or by trapping reactions with a broad range of electrophiles (aldehydes, ketones, allylic halides, acyl chlorides, Weinreb amides, aryl halides, hydroxylamine benzoates, terminal alkynes). Furthermore, several double *ortho,ortho'*-magnesiations were realized in the case of aryl oxazolines, *N*-aryl pyrazoles as well as 2-aryl-2*H*-1,2,3-triazoles by simply repeating the magnesiation/electrophile trapping sequence allowing the preparation of valuable 1,2,3-functionalized arenes.

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Introduction

The directed magnesiation of arenes and heteroarenes is an important synthetic tool for the preparation of polyfunctional aryl- and heteroaryl-magnesium organometallics.¹ Mixed magnesium and lithium amides R₂NMgX·LiCl are usually the most efficient reagents for such metalations.² Recently, we have examined the regioselective metalation of various pharmaceutically relevant aryl azoles such as **1**.³ We found that standard metal amides such as LDA or TMPLi (TMP = 2,2,6,6-tetramethylpiperidyl) gave the lithiated products **2** with poor regioselectivity, due to a competitive deprotonation at the 5-position of the triazole ring of **1**. The best result was achieved in toluene⁴ using the alkylmagnesium amide TMPMgBu⁵ prepared from commercial Bu₂Mg, which provided after cross-coupling with aryl bromides various products of type **3**. Although this base was highly regioselective in toluene, an excess of ArBr was required to compensate the formation of the Ar-*n*Bu side-product, originating from a faster cross-coupling of the *n*Bu moiety compared to the metalated azole **2** (Scheme 1).

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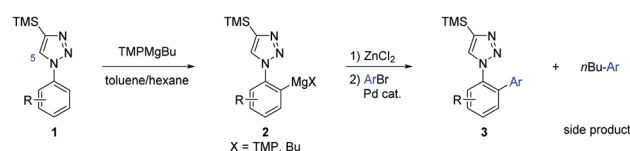
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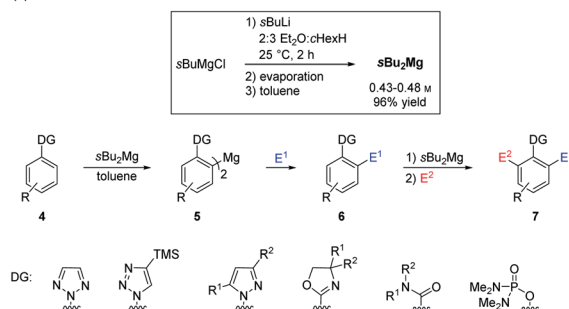
† Electronic supplementary information (ESI) available. CCDC 2061336 and 2061337. For ESI and crystallographic data in CIF or other electronic format see DOI: 10.1039/d1sc01777b

While commercially available Bu₂Mg contained a 60 : 40 mixture of *n*Bu₂Mg and *s*Bu₂Mg, we have found only small amounts of the branched coupling side-product Ar-*s*Bu, suggesting that the secondary alkyl moiety was reacting much slower than the primary one.

(a) Previous work:



(b) This work:



E¹, E²: aldehydes, ketones, allylic halides, acyl chlorides, Weinreb amides, aryl halides (Negishi cross-coupling).

Scheme 1 (a) Regioselective magnesiation and subsequent Negishi cross-coupling of aryl azoles (**1**) using TMPMgBu in toluene/hexane. (b) Regioselective magnesiation and *ortho,ortho'*-functionalization of arenes and heteroarenes using *s*Bu₂Mg in toluene.



Herein, we report the preparation of $s\text{Bu}_2\text{Mg}$,⁶ which avoided these side reactions and significantly increased the metalation scope. Thus, we showed that $s\text{Bu}_2\text{Mg}$ was an improved magnesiation reagent, which allowed a highly *ortho*-regioselective magnesiation of arenes **4** bearing various directing groups (DG), leading after trapping of the resulting diarylmagnesium species **5** with various electrophiles E^1 to products of type **6**. These polyfunctional arenes were in several cases magnesiated again using $s\text{Bu}_2\text{Mg}$ producing, after addition of a second different electrophile E^2 , valuable 1,2,3-polyfunctional arenes of type **7**.

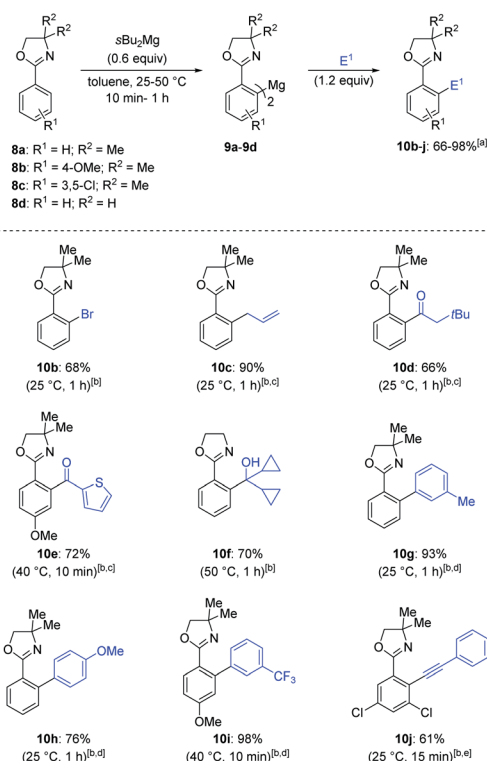
Results and discussion

The reaction of $s\text{BuMgCl}$ in diethyl ether with $s\text{BuLi}$ (1.0 equiv.) in cyclohexane at 25 °C (2 h) gave, after solvent evaporation under vacuum, redissolution in toluene and filtration, a 0.43–0.48 M solution of $s\text{Bu}_2\text{Mg}$ in 96% yield.^{7,8}

In preliminary experiments, we have observed a smooth magnesiation of oxazoline **8a** with a toluene solution of 0.6 equiv. of $s\text{Bu}_2\text{Mg}$ leading to the diarylmagnesium **9a** (Table 1). A full conversion to the diarylmagnesium species was achieved within 1 hour and the iodolyzed product **10a** was isolated in 80% yield (entry 1). $s\text{Bu}_2\text{Mg}$ gave also good results in cyclohexane or THF, albeit in lower yields (entries 2 and 3). $c\text{Hex}_2\text{Mg}$ in toluene delivered the desired product **10a** in only 46% yield and other bases such as Ph_2Mg , $(\text{TMSCH}_2)_2\text{Mg}$ and $t\text{Bu}_2\text{Mg}$ or $s\text{BuMgCl}^9$ did not give any conversion (entries 4–8).

Therefore, a range of oxazolines (**8a–d**) were magnesiated selectively on the aryl ring and the resulting diarylmagnesiums (**9a–d**) underwent Negishi cross-couplings,¹⁰ copper-catalyzed allylation or acylation,¹¹ *in situ* Sonogashira cross-coupling¹² or trapping reactions with tetrachlorodibromoethane or dicyclopropylketone, leading to the *mono-ortho* substituted oxazolines **10b–j** in 68–98% yield (Scheme 2).

Most of the C–H activation methods currently available for the arylation of aryl azoles were performed by using transition



Scheme 2 Regioselective magnesiation of oxazolines **8a–d** with $s\text{Bu}_2\text{Mg}$ leading, *via* diarylmagnesium species **9a–d**, to functionalized oxazolines **10b–j**. ^a All yields refer to isolated compounds. ^b Magnesiation conditions. ^c The reaction was catalyzed by $\text{CuCN}\cdot 2\text{LiCl}$ (20 mol%). ^d Obtained after transmetalation with ZnCl_2 (1.1 equiv.) and a palladium-catalyzed cross-coupling with $[\text{PdCl}_2(\text{dppf})]$ (5 mol%), dppe = diphenylphosphinoferrocene and an aryl halide (0.83 equiv.). ^e Obtained after transmetalation with ZnCl_2 (1.1 equiv.), subsequent iodine quench (1.1 equiv.) and Sonogashira cross-coupling with CuI (4 mol%), $\text{Pd}(\text{dba})_2$ (3 mol%), dba = dibenzylideneacetone, tri-(2-furyl)-phosphine (6 mol%) and phenylacetylene (1.3 equiv.).

Table 1 Magnesiation of oxazoline **8a** using various magnesium reagents in various solvents at 25 °C

Entry	Reagent	Solvent	Yield ^a
1	$s\text{Bu}_2\text{Mg}$	Toluene	91% (80) ^c
2	$s\text{Bu}_2\text{Mg}$	THF	73%
3	$s\text{Bu}_2\text{Mg}$	Cyclohexane	64%
4	$c\text{Hex}_2\text{Mg}$	Toluene	46%
5	$t\text{Bu}_2\text{Mg}$	Toluene	0%
6	$(\text{TMSCH}_2)_2\text{Mg}$	Toluene	0%
7	Ph_2Mg	Toluene	0%
8	$s\text{BuMgCl}^b$	Ether/toluene	0%

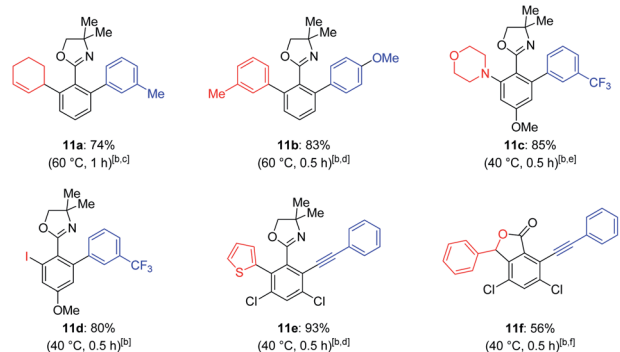
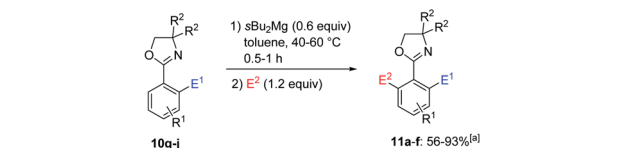
^a Calibrated GC-yield using undecane as internal standard. ^b 1.2 equiv. of $s\text{BuMgCl}$ were used. ^c Isolated yield.

metal catalysts and suffered from the unwanted formation of symmetrical bis-arylated products and the selective preparation of unsymmetrical *ortho-ortho'*-bis-functionalized¹³ aryl azoles remained challenging.¹⁴ We have found that various oxazolines **10g–j** were again magnesiated at 40–60 °C with $s\text{Bu}_2\text{Mg}$ in toluene (Scheme 3).¹⁵ The intermediate diarylmagnesium species were further functionalized by a copper-catalyzed allylation, Negishi cross-coupling, cobalt-catalyzed electrophilic amination¹⁶ and iodolysis furnishing the desired products **11a–e** in 74–93% yield. Interestingly, magnesiation of **10j** followed by trapping with benzaldehyde and subsequent treatment with 6 M HCl provided lactone **11f** in 56% yield.¹⁷

To demonstrate the versatility of the oxazoline directing group, the strongly sterical hindered *ortho,ortho'*-functionalized oxazoline **11b** was successfully converted to the corresponding nitrile **11g** using thionyl chloride and DMF¹⁸ in 92% yield (Scheme 4).¹⁹

We then turned our attention to the magnesiation of various *N*-aryl pyrazoles (**12a–c**). $s\text{Bu}_2\text{Mg}$ proved also to be an excellent base for the regioselective magnesiation of *N*-aryl pyrazole **12a**, affording the corresponding bis-arylmagnesium species **13a**



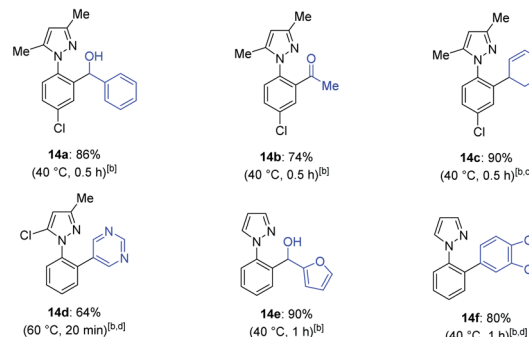
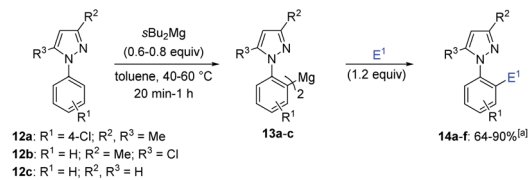


Scheme 3 Regioselective magnesiation of mono-functionalized oxazolines **10g–j**, leading to *ortho,ortho'*-functionalized oxazolines **11a–f**. ^a All yields refer to isolated compounds. ^b Magnesiation conditions. ^c The reaction was catalyzed by CuCN·2LiCl (20 mol%). ^d Obtained after transmetalation with ZnCl₂ (1.1 equiv.) and a palladium-catalyzed cross-coupling with [PdCl₂(dppf)] (5 mol%) and an aryl iodide (0.83 equiv.). ^e Obtained after transmetalation with ZnCl₂ (1.1 equiv.) and a cobalt-catalyzed electrophilic amination with CoCl₂ (5 mol%) and morpholino benzoate (1.2 equiv.). ^f Obtained after addition of benzaldehyde (1.2 equiv.) followed by treatment with 6 M HCl.



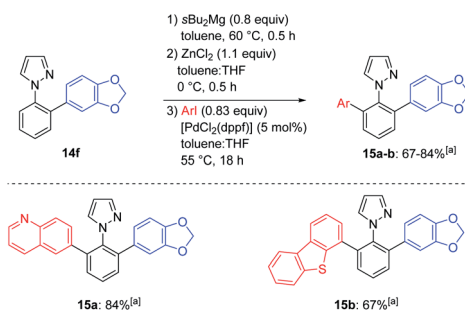
Scheme 4 Transformation of *ortho,ortho'*-functionalized oxazoline **11b** to the corresponding nitrile **11g**. ^a Isolated yield.

after 0.5 h at 40 °C. After addition of benzaldehyde or Weinreb amide MeCON(OMe)Me, alcohol **14a** and ketone **14b** were obtained in 74–86% yield (Scheme 5). Copper-catalyzed allylation with 3-bromocyclohex-1-ene produced the pyrazole **14c** (90% yield). Interestingly, *N*-aryl pyrazoles **12b** and **12c** although bearing relatively acidic protons at the heterocyclic ring were selectively magnesiated at the *ortho*-position of the phenyl ring. In particular, unsubstituted pyrazole **12c** was metalated in 94% yield and >98 : 1 : 1 selectivity, as determined by deuterolysis of a reaction aliquot.²⁰ These results further confirm the key role of the coordination at the N(2)-atom of the pyrazole to direct the metalation selectively on the aryl ring in a non-polar solvent like toluene. Thus, the functionalized pyrazoles **14d–f** were obtained after Negishi cross-coupling with 5-bromopyrimidine, 5-bromobenzo[*d*][1,3]dioxole or addition of furfural in 64–90% yield. We also achieved an unsymmetrical *ortho,ortho'*-functionalization and mono-substituted pyrazole **14f** was selectively magnesiated at 60 °C (0.5 h) and trapped by Negishi cross-coupling with 6-iodoquinoline and 4-iododibenzo[*b,d*]thiophene providing the products **15a–b** in 67–84% yield (Scheme 6).



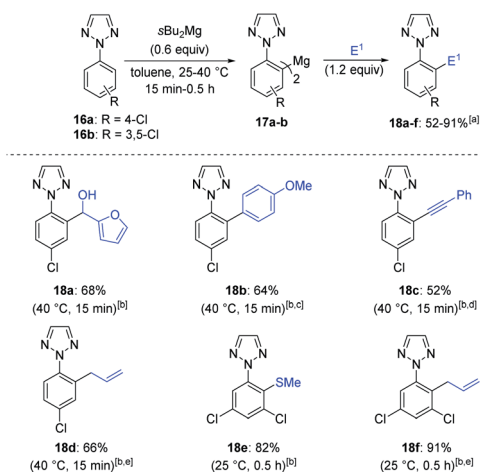
Scheme 5 Regioselective magnesiation of *N*-aryl pyrazoles **12a–c** with sBu₂Mg leading, via diarylmagnesium species **13a–c**, to functionalized *N*-aryl pyrazoles **14a–f**. ^a All yields refer to isolated compounds. ^b Magnesiation conditions. ^c The reaction was catalyzed by CuCN·2LiCl (20 mol%). ^d Obtained after transmetalation with ZnCl₂ (1.1 equiv.) and a palladium-catalyzed cross coupling with [PdCl₂(dppf)] (5 mol%) and an aryl bromide (0.83 equiv.).

The functionalization of less common heterocycles is of key importance for pharmaceutical applications.²¹ Thus, the metalation of symmetrical 2-aryl-2*H*-1,2,3-triazoles **16a–b** was then investigated (Scheme 7).^{22,23} After metalation of **16a** with 0.6 equiv. of sBu₂Mg for 15 min at 40 °C, the resulting bis-arylmagnesium species **17a** was then trapped with furfural, affording the functionalized 1,2,3-triazole **18a** in a 68% yield. Further trapping reactions such as Negishi cross-coupling, copper-catalyzed allylation and oxidative alkylation with (phenylethynyl)lithium²⁴ lead to 2-aryl-1,2,3-triazoles **18b–d** in 52–66% yield. Similarly, 1,2,3-triazole **16b** was readily magnesiated at 25 °C (0.5 h) as shown by the quantitative formation of a single regioisomer by NMR-analysis of a deuterolyzed reaction aliquot.²⁰ Further quenching reactions of **17b** like thiomethylation and allylation furnished triazoles **18e–f** in 82–91% yield. A second functionalization was performed on *N*-aryl



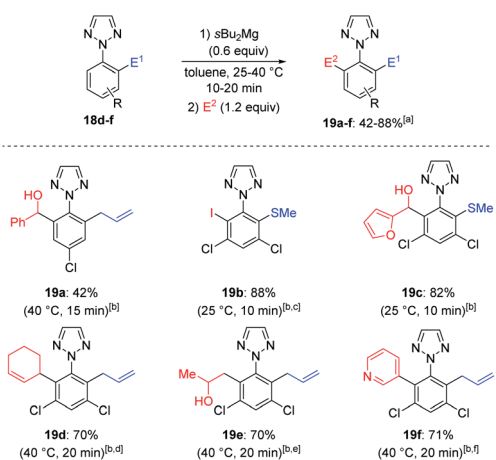
Scheme 6 Regioselective magnesiation of mono-functionalized *N*-aryl pyrazole **14f** with sBu₂Mg leading to *ortho,ortho'*-functionalized *N*-aryl pyrazoles **15a–b**. ^a All yields refer to isolated compounds.





Scheme 7 Regioselective magnesiation of 2-aryl-2H-1,2,3-triazoles **16a–b** with $s\text{Bu}_2\text{Mg}$ leading, via diarylmagnesium species **17a–b**, to functionalized 2-aryl-2H-1,2,3-triazoles **18a–f**. ^a All yields refer to isolated compounds. ^b Magnesiation conditions. ^c Obtained after transmetalation with ZnCl_2 (1.1 equiv.) and a palladium-catalyzed cross-coupling with $[\text{PdCl}_2(\text{dppf})]$ (5 mol%) and an aryl halide (0.83 equiv.). ^d Obtained after transmetalation with $\text{CuCN}\cdot 2\text{LiCl}$ (1.2 equiv.) and subsequent addition of (phenylethynyl)lithium (2.0 equiv.), followed by addition of chloranil (1.3 equiv.). ^e The reaction was catalyzed by $\text{CuCN}\cdot 2\text{LiCl}$ (20 mol%).

triazoles **18d–f** using again $s\text{Bu}_2\text{Mg}$ in toluene, followed by quench with a different electrophile (E^2) (Scheme 8). We observed a complete magnesiation of **18d** with $s\text{Bu}_2\text{Mg}$ within 15 min at 40 °C and a subsequent reaction with benzaldehyde produced the mixed bis-functionalized 1,2,3-triazole **19a** in 42% yield. Similarly, **18e** and **18f** were magnesiated under the



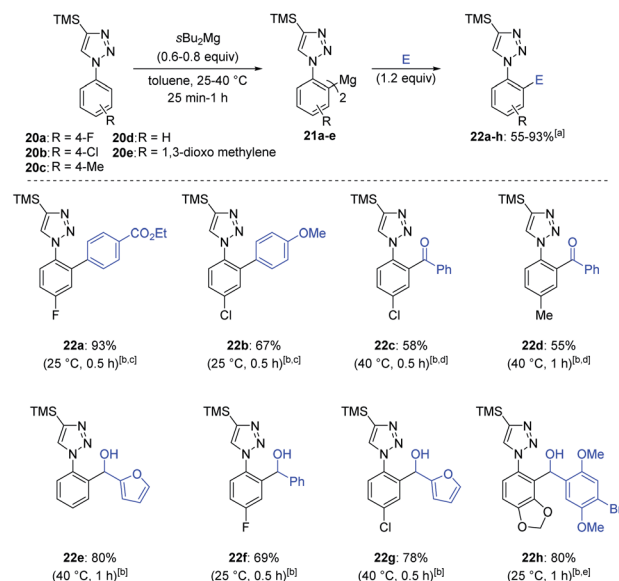
Scheme 8 Regioselective magnesiation of mono-functionalized 2-aryl-2H-1,2,3-triazoles **18d–f** with $s\text{Bu}_2\text{Mg}$ leading to *ortho,ortho'*-functionalized 2-aryl-2H-1,2,3-triazoles **19a–f**. ^a All yields refer to isolated compounds. ^b Magnesiation conditions. ^c The regioselectivity was determined by crystal structure analysis, see ESI.† ^d The reaction was catalyzed by $\text{CuCN}\cdot 2\text{LiCl}$ (20 mol%). ^e The reaction was catalyzed by CuI (10 mol%). ^f Obtained after transmetalation with ZnCl_2 (1.1 equiv.) and a palladium-catalyzed cross-coupling with $[\text{PdCl}_2(\text{dppf})]$ (5 mol%) and an aryl bromide (0.83 equiv.).

standard conditions and the resulting bis-arylmagnesium species were trapped with a different electrophile (E^2) leading to a range of unsymmetrical functionalized 1,2,3-triazoles **19b–f** in 70–88% yield.

Finally, we examined the metalation of 1-aryl-1H-1,2,3-triazoles such as **20a–e** and found that $s\text{Bu}_2\text{Mg}$ led to a highly regioselective magnesiation at the *ortho*-position of the aryl ring in toluene (25–40 °C, 0.5–1 h), affording the bis-aryl-magnesium species **21a** in 75% yield and 97 : 3 regioselectivity (Scheme 9).²⁰

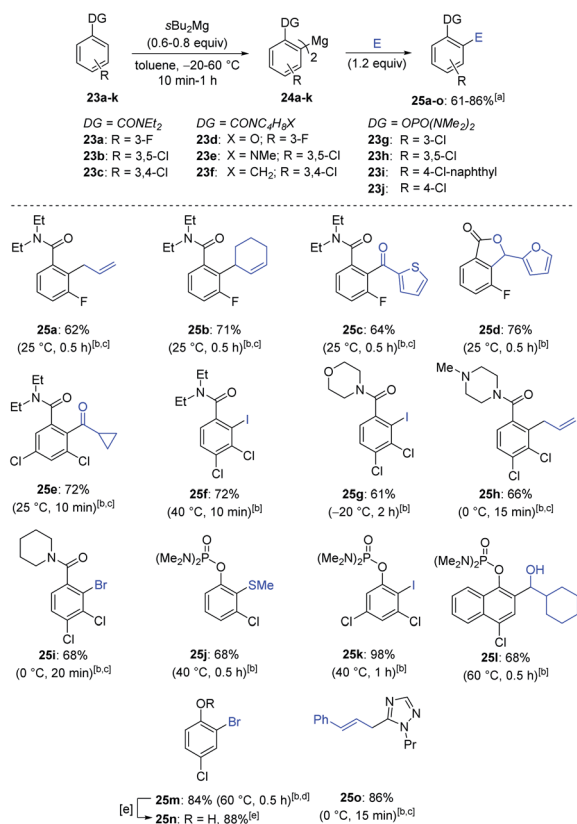
This new metalation procedure occurred twice as fast as the previously reported TMPMgBu base.³ 1,2,3-Triazoles **22a** and **22b** were isolated in 93% and 67% yields respectively after Negishi cross-couplings with only 0.83 equiv. of aryl bromide.³ Copper-catalyzed acylation¹¹ with benzoyl chloride lead to products **22c** and **22d** in 55–58% yield and quenching with various aldehydes afforded compounds **22e–h** in 69–80% yield.

Remarkably, $s\text{Bu}_2\text{Mg}$ was also an excellent base for the magnesiation of various arenes bearing directing groups such as a tertiary amide or phosphorodiamidate (**23a–j**; Scheme 10).²⁵ The addition of $s\text{Bu}_2\text{Mg}$ to the aromatic amide **23a** in toluene led to a clean magnesiation within 0.5 h at room temperature. The resulting diarylmagnesium species **24a** was then further allylated with allyl and cyclohexenyl bromides, leading to **25a** and **25b** in 62% and 71% yield respectively. Copper-catalyzed acylation of **23a** with thiophene-2-carbonyl chloride or trapping with furfural furnished the ketone **25c** (64% yield) and the lactone **25d** (76% yield).²⁶ Similarly, the amides **23b–f** afforded with the same magnesiation/trapping sequence the polyfunctional amides (**25e–i**) in 61–72% yield. Various

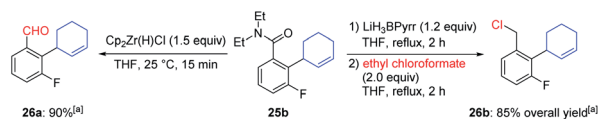


Scheme 9 Regioselective magnesiation of 1-aryl-2H-1,2,3-triazoles **20a–e** with $s\text{Bu}_2\text{Mg}$ leading, via diarylmagnesium species **21a–e**, to functionalized 1-aryl-2H-1,2,3-triazoles **22a–h**. ^a All yields refer to isolated compounds. ^b Magnesiation conditions. ^c Obtained after transmetalation with ZnCl_2 (1.1 equiv.) and a palladium-catalyzed cross-coupling with $[\text{PdCl}_2(\text{dppf})]$ (5 mol%) and an aryl halide (0.83 equiv.). ^d The reaction was catalyzed by $\text{CuCN}\cdot 2\text{LiCl}$ (20 mol%). ^e The regioselectivity was determined by crystal structure analysis, see ESI.†





Scheme 10 Regioselective magnesiation of various arenes bearing an amide or a phosphorodiamidate directing group as well as 1-propyl-1,2,4-triazole **23a–k** with $s\text{Bu}_2\text{Mg}$ leading, *via* diarylmagnesium species **24a–k**, to functionalized arenes **25a–o**. ^a All yields refer to isolated compounds. ^b Magnesiation conditions. ^c The reaction was catalyzed by $\text{CuCN} \cdot 2\text{LiCl}$ (20 mol%). ^d R = $\text{P}(\text{O})(\text{NMe}_2)_2$. ^e Obtained after treating **25m** with 2 M HCl in dioxane (105 °C, 1 h).



Scheme 11 Synthetic transformations of magnesiated product **25b**. ^a All yields refer to isolated compounds.

phosphorodiamidates (**23g–j**) were also metalated with $s\text{Bu}_2\text{Mg}$ at 40–60 °C (0.5–1 h) providing the diarylmagnesiums **24g–j**, which were trapped with a range of electrophiles (MeSSO_2Me , I_2 , $c\text{HexCHO}$ and $(\text{BrCCl}_2)_2$) furnishing the phenol derivatives **25j–m** in 68–98% yield. Removal of the phosphorodiamidate group²⁷ in **25m** was achieved with a 2 M HCl treatment in dioxane (105 °C, 1 h) leading to phenol **25n** in 88% yield. Interestingly, 1-propyl-1,2,4-triazole (**23k**) was magnesiated with $s\text{Bu}_2\text{Mg}$ and allylated with cinnamyl bromide providing the *N*-heterocycle **25o** in 86% yield.²⁸

We performed some further transformations leading to polyfunctionalized 1,2,3-trisubstituted arenes to show the utility of these magnesiations. The newly prepared amide **25b** was thus selectively reduced with $\text{Cp}_2\text{Zr}(\text{H})\text{Cl}$ ²⁹ (25 °C, 15 min) to

the aldehyde **26a** in 90% yield. A two-step transformation consisting of a reduction with the complex borohydride $\text{LiH}_3\text{-BPyrr}$ ($\text{Pyrr} = \text{pyrrolidino}$)³⁰ followed by a treatment with ethyl chloroformate³¹ provided the benzylic chloride **26b** in 85% overall yield (Scheme 11).

Conclusions

In summary, we have developed a new preparation of $s\text{Bu}_2\text{Mg}$ in toluene and showed its utility for the directed magnesiation of various aromatic and heterocyclic systems including pharmaceutically relevant *N*-arylated pyrazoles as well as *N*-arylated 1,2,3-triazoles. This method provides a unique access to various diarylmagnesium reagents in toluene. Furthermore, a range of arenes bearing various directing groups such as an oxazoline, phosphorodiamidate or an amide were magnesiated with $s\text{Bu}_2\text{Mg}$. Remarkably, a second unsymmetrical *ortho,ortho'*-functionalization was achieved in the case of aryl oxazolines, *N*-aryl pyrazoles as well as *N*-aryl triazoles, leading to valuable synthetic intermediates of potential pharmaceutical relevance. Further investigations of the use of $s\text{Bu}_2\text{Mg}$ as metalating agent are currently underway in our laboratory.

Author contributions

A. H., J. P. P., S. B. D., F. T. and K. K. performed and analyzed the experiments. A. H., F. H. L., S. L., S. W. and P. K. designed the experiments. A. H., S. L., S. W., and P. K. prepared the manuscript with contributions of all authors.

Conflicts of interest

There are no conflicts to declare.

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Notes and references

- (a) R. E. Mulvey, F. Mongin, M. Uchiyama and Y. Kondo, *Angew. Chem., Int. Ed.*, 2007, **46**, 3803; (b) K. W. Henderson and W. J. Kerr, *Chem.-Eur. J.*, 2001, **7**, 3430; (c) C. R. Hauser and H. G. Walker, *J. Am. Chem. Soc.*, 1947, **69**, 295; (d) J. Wei, W.-X. Zhang and Z. Xi, *Org. Chem. Front.*, 2014, **1**, 983.
- (a) A. Krasovskiy, V. Krasovskaya and P. Knochel, *Angew. Chem., Int. Ed.*, 2006, **45**, 2958; (b) G. C. Clososki, C. J. Rohbogner and P. Knochel, *Angew. Chem., Int. Ed.*, 2007, **46**, 7681; (c) B. Haag, M. Mosrin, H. Ila, V. Malakhov and P. Knochel, *Angew. Chem., Int. Ed.*, 2011, **50**, 9794; (d) P. E. Eaton, C. H. Lee and Y. Xiong, *J. Am. Chem. Soc.*, 1989, **111**, 8016; (e) P. E. Eaton and K. A. Lukin, *J. Am. Chem. Soc.*, 1993, **115**, 11370.



- 3 F. H. Lutter, L. Grokenberger, L. A. Perego, D. Broggini, S. Lemaire, S. Wagschal and P. Knochel, *Nat. Commun.*, 2020, **11**, 4443.
- 4 D. S. Ziegler, K. Karaghiosoff and P. Knochel, *Angew. Chem., Int. Ed.*, 2018, **57**, 6701.
- 5 (a) E. Hevia, A. R. Kennedy, R. E. Mulvey and S. Weatherstone, *Angew. Chem., Int. Ed.*, 2004, **43**, 1709; (b) B. Conway, E. Hevia, A. R. Kennedy, R. E. Mulvey and S. Weatherstone, *Dalton Trans.*, 2005, 1532.
- 6 (a) L. J. Bole, N. R. Judge and E. Hevia, *Angew. Chem., Int. Ed.*, 2021, **60**, 7626; (b) C. W. Kamienski and J. F. Eastham, *J. Organomet. Chem.*, 1967, **8**, 542; (c) N. D. R. Barnett, W. Clegg, R. E. Mulvey, P. A. O'Neil and D. Reed, *J. Organomet. Chem.*, 1996, **510**, 297.
- 7 This reagent still contained 0.5 equiv. of Et₂O, having the formula sBu₂Mg·0.5Et₂O (determined by ¹H-NMR analysis) that we abbreviated sBu₂Mg for the sake of clarity.
- 8 The toluene solution of sBu₂Mg can be stored at –30 °C for at least 1 week.
- 9 A. Marxer and M. Siegrist, *Helv. Chim. Acta*, 1974, **57**, 1988.
- 10 A. O. King, N. Okukado and E.-I. Negishi, *J. Chem. Soc., Chem. Commun.*, 1977, **19**, 683.
- 11 P. Knochel, M. C. P. Yeh, S. C. Berk and J. Talbert, *J. Org. Chem.*, 1988, **53**, 2390.
- 12 (a) K. Sonogashira, Y. Tohda and N. Hagihara, *Tetrahedron Lett.*, 1975, **50**, 4467; (b) M. Mosrin, T. Bresser and P. Knochel, *Org. Lett.*, 2009, **11**, 3406.
- 13 (a) A. J. Martinez-Martinez, A. R. Kennedy, R. E. Mulvey and C. T. O'Hara, *Science*, 2014, **346**, 834; (b) A. J. Martinez-Martinez, S. Justice, B. J. Fleming, A. R. Kennedy, I. D. H. Oswald and C. T. O'Hara, *Sci. Adv.*, 2017, **3**, e1700832.
- 14 (a) M. Simonetti, D. M. Cannas, X. Just-Baringo, I. J. Vitorica-Yrezabal and I. Larrosa, *Nat. Chem.*, 2018, **10**, 724; (b) L. Ackermann, A. Althammer and R. Born, *Tetrahedron*, 2008, **64**, 6115; (c) S. Oi, H. Sato, S. Sugawara and Y. Inoue, *Org. Lett.*, 2008, **10**, 1823; (d) C. J. Teskey, S. M. A. Sohel, D. L. Bunting, S. G. Modha and M. F. Greaney, *Angew. Chem., Int. Ed.*, 2017, **56**, 5263; (e) O. Daugulis and V. G. Zaitsev, *Angew. Chem., Int. Ed.*, 2005, **44**, 4046.
- 15 A. I. Meyers and W. B. Avila, *J. Org. Chem.*, 1981, **46**, 3881.
- 16 (a) A. M. Berman and J. S. Johnson, *J. Am. Chem. Soc.*, 2004, **126**, 5680; (b) A. M. Berman and J. S. Johnson, *J. Org. Chem.*, 2005, **70**, 364; (c) M. Campbell and J. S. Johnson, *Org. Lett.*, 2007, **9**, 1521; (d) Y.-H. Chen, S. Graßl and P. Knochel, *Angew. Chem., Int. Ed.*, 2018, **57**, 1108; (e) S. Graßl, Y.-H. Chen, C. Hamze, C. P. Tüllmann and P. Knochel, *Org. Lett.*, 2019, **21**, 494.
- 17 C. A. Boulet and G. A. Poulton, *Heterocycles*, 1989, **28**, 405.
- 18 (a) O. Fischer, A. Muller and A. Vilsmeier, *J. Prakt. Chem.*, 1925, **109**, 69; (b) G. Jones and S. P. Stanforth, *Org. React.*, 2000, **56**, 355.
- 19 (a) I. M. Dordor and J. M. Mellor, *Tetrahedron Lett.*, 1983, **24**, 1437; (b) H. Petersen, *US Pat.* US2003/13895, A1, 2003.
- 20 See Supporting Information for further details.†
- 21 (a) V. M. Ahrens, K. Bellmann-Sickert and A. G. Beck-Sickinger, *Future Med. Chem.*, 2012, **4**, 1567; (b) C. R. Dass and P. F. M. Choong, *Peptides*, 2006, **27**, 3020.
- 22 (a) J. L. Riebsommer, *J. Org. Chem.*, 1948, **13**, 815; (b) G. F. Myachina, T. G. Ermakova, N. P. Kuznetsova, R. G. Sultangareev, L. I. Larina, L. V. Klyba, G. T. Suchanov and B. A. Trofimov, *Chem. Heterocycl. Compd.*, 2010, **46**, 79.
- 23 S. Shi, W. Liu, P. He and C. Kuang, *Org. Biomol. Chem.*, 2014, **12**, 3576.
- 24 S. R. Dubbaka, M. Kienle, H. Mayr and P. Knochel, *Angew. Chem., Int. Ed.*, 2007, **46**, 9093.
- 25 (a) P. Beak, G. R. Brubaker and R. J. Farney, *J. Am. Chem. Soc.*, 1976, **98**, 3621; (b) P. Beak and R. A. Brown, *J. Org. Chem.*, 1977, **42**, 1823; (c) P. Bowles, J. Clayden, M. Helliwell, C. McCarthy, M. Tomkinson and N. Westlund, *J. Chem. Soc., Perkin Trans. 1*, 1997, 2607; (d) J.-C. Cuevas, P. Patil and V. Snieckus, *Tetrahedron Lett.*, 1989, **30**, 5841; (e) W. H. Puterbaugh and C. R. Hauser, *J. Org. Chem.*, 1964, **29**, 853; (f) M. Watanabe and V. Snieckus, *J. Am. Chem. Soc.*, 1980, **102**, 1457; (g) F. Hintze and D. Hoppe, *J. Org. Chem.*, 1990, **55**, 1375; (h) H. Gilman and R. L. Bebb, *J. Am. Chem. Soc.*, 1939, **61**, 109; (i) G. Wittig and G. Fuhrman, *Chem. Ber.*, 1940, **73**, 1197; (j) V. Snieckus, *Chem. Rev.*, 1990, **90**, 879.
- 26 W. Lin, O. Baron and P. Knochel, *Org. Lett.*, 2006, **24**, 5673.
- 27 (a) M. Watanabe, M. Date, K. Kawanishi, M. Tsukazaki and S. Furukawa, *Chem. Pharm. Bull.*, 1989, **37**, 2564; (b) C. J. Rohbogner, S. Wirth and P. Knochel, *Org. Lett.*, 2010, **9**, 1984.
- 28 K. Schwärzer, C. P. Tüllmann, S. Graßl, B. Górski, C. E. Brocklehurst and P. Knochel, *Org. Lett.*, 2020, **5**, 1899.
- 29 (a) J. M. White, A. R. Tunoori and G. I. Georg, *J. Am. Chem. Soc.*, 2000, **122**, 11995; (b) Y. Zhao and V. Snieckus, *Org. Lett.*, 2014, **2**, 390.
- 30 G. B. Fisher, J. C. Fuller, J. Harrison, C. T. Goralski and B. Singaram, *Tetrahedron Lett.*, 1993, **34**, 1091.
- 31 H. Kapnang and G. Charles, *Tetrahedron Lett.*, 1983, **24**, 3233.

