

ChemComm

Chemical Communications

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ISSN 1359-7345

COMMUNICATION

Christophe Werlé, Alexandre Vasseur *et al.*
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Cite this: *Chem. Commun.*, 2025, 61, 6296

Received 27th February 2025,
Accepted 21st March 2025

DOI: 10.1039/d5cc01079a

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Lithium zincate-enabled divergent one-pot dual C–C bond formation in thiophenes†

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We present a lithium zincate-enabled, divergent one-pot synthesis for regioselective dual C–C bond formation in thiophenes. By modifying the zinc coordination environment, a single set of reagents (ZnCl₂, R¹Li, and diethyl (5-halo)thienylphosphate) was found to generate two distinct products. This approach extends the versatility of lithium organozincates to regioselective C_{Ar}(sp²)–C_{thienyl}(sp²) and C_{thienyl}(sp³)–C_{Ar}(sp²) couplings without requiring transition metals and/or arenes pre-activated with a boronic acid.

Recent outbreaks of epidemic and pandemic diseases, such as those caused by Ebola, SARS-CoV-2, and the Mpox virus, emphasise the urgency of rapid drug discovery. Each crisis spurs a swift search for treatments, driving efforts to develop new therapies or improve existing ones. In this context, synthetic chemists, alongside biologists, are equally mindful of the importance of efficient methods for accessing diverse molecular libraries.¹ High-throughput screening of structurally related compounds often serves as a starting point for identifying potential treatments,² requiring synthetic methods that are both rapid and scalable. Synthetic strategies that quickly produce diverse, yet structurally related molecules are critical in this process. “One-pot synthesis” is particularly advantageous for such demands as it enables multiple bond-forming reactions within a single reaction vessel.³ This strategy encompasses domino, consecutive, and independent reaction sequences, streamlining processes by eliminating the need to isolate intermediates.⁴ Similarly, ligand-enabled metal-promoted divergent synthesis offers flexibility⁵ by allowing different products to form from the same starting material by adjusting the ligand environment around the

metal. However, these strategies are generally applied independently and have limited integration. A reagent capable of adapting to a wide range of reaction pathways could combine the principles of one-pot and divergent synthesis, offering increased versatility.

We hypothesised that lithium zincate (LiZn) of the PAIRiodic table,⁶—a synergistic complex combining group 1 and group 12 elements—could fulfil this role (Scheme 1a). When ZnCl₂ reacts with three equivalents of an organolithium compound (R¹Li), a 16-electron mono-anionic complex, (R¹)₃ZnLi, is formed. This complex features a dynamic Li–Zn bond (Scheme 1b), whose strength and adaptivity were recently demonstrated computationally for Et₃ZnLi·2THF (Scheme 1c).⁷ Lithium triorganozincates demonstrate significant structural flexibility in THF, existing as a network of dynamic equilibria between three distinct forms (Scheme 1d).⁷ This flexibility underpins their broad reactivity profile,⁸ which includes metallation *via* I/Zn exchange,⁹ deprotonation,¹⁰ and homologation through 1,2-ligand migration.¹¹ Additionally, they participate in 1,2- and 1,4-ligand additions to electrophiles,¹² enabling both ligand-enabled divergent synthesis¹³ and sequential carbon–carbon bond formation in one-pot synthesis.¹⁴ In addition to these capabilities, reacting ZnCl₂ with four equivalents of R¹Li produces an 18-electron di-anionic complex, (R¹)₄ZnLi₂ (Scheme 1b)¹⁵ which is thermodynamically more stable, kinetically more active and provides additional synthetic opportunities such as Br/Zn exchange metallations,¹⁵ cross-coupling reactions,¹⁶ conversions of carboxylic acid to ketones¹⁷ and transition metal catalyst-free conjugate addition to nitroolefins.¹⁸ With these characteristics established, we conceptualised two lithium zincate-mediated pathways as a framework for achieving regioselective functionalisation of thiophene derivatives. Using diethyl (5-halo)thienylphosphate **1** as a model substrate, we sought to explore their feasibility.

In the first pathway (Scheme 1e), treating **1a** (X = I) with two equivalents of lithium triorganozincate (R¹)₃ZnLi (**2**) initiates an I/Zn exchange to produce intermediate **I**. The coexistence of a formally negatively charged Zn centre and a potentially cationic centre at the thienyl position could drive a homologation reaction *via* a 1,2-migration of an R¹ ligand, yielding intermediate **II** with temporary dearomatisation. A formally [1,5] metallotropic rearrangement

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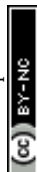
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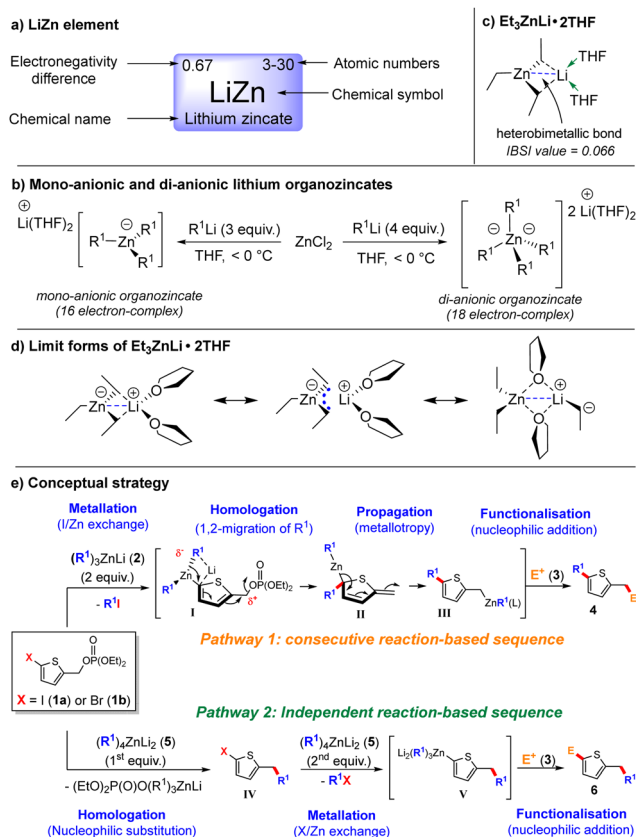
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† Electronic supplementary information (ESI) available: Experimental procedures, characterisation for new compounds including NMR spectra. See DOI: <https://doi.org/10.1039/d5cc01079a>





Scheme 1 Development of a lithium-zincate system for divergent bond formation. (a) Conceptual representation of the “LiZn element” as a synergistic assembly. (b) Structures of mono- and di-anionic zincates. (c) Representation of Li–Zn bonding. (d) Structural equilibria in THF. (e) Conceptual strategy.

would then restore aromaticity, producing the homologated thenyl-zinc intermediate **III**, which would react with an electrophile **3** to yield the final product (**4**).

In contrast, the second pathway (Scheme 1e) employs two equivalents of di-anionic tetraorganozincate (**5**), where homologation proceeds *via* nucleophilic substitution on **1a** or **1b** ($X = \text{Br}$). This generates an isomeric product (**6**) through an X/Zn exchange followed by electrophilic trapping. This pathway emphasises controlled sequential reactivity, requiring homologation to occur faster than metallation to maintain pathway fidelity. Directing each pathway enables access to a diverse array of heterocycles, highlighting the versatility of the lithium zincate-mediated approach. However, several challenges need to be addressed to ensure successful implementation. These include maintaining the stability of the thiophene framework under the polar organometallic conditions required for pathway 1,¹⁹ managing the competition between halogen/metal exchange and nucleophilic substitution that determines entry into either pathway 1 or 2, and achieving selective reactivity of **III** and **V** with electrophile **3** (transfer of the thenyl or thienyl ligand rather than the alkyl R^1 ligand).

To overcome these challenges, we developed a LiZn element-promoted divergent process that regioselectively forms two carbon–carbon bonds five atoms apart in a single reaction vessel. By modifying the Zn coordination environment, we

achieved significant structural transformations through the distinct reaction pathways. Given the widespread applications of thiophenes in pharmaceuticals²⁰ and materials science,²¹ this investigation focuses on leveraging diethyl (5-halo)thenyl-phosphate **1** to explore regioselective functionalisation strategies.

As an initial step, we identified optimal conditions to carry out the consecutive reaction sequence in pathway 1 (Scheme 1e). Treating **1a** with $(n\text{-Bu})_3\text{ZnLi}$ at -85°C in 2-MeTHF for two hours, followed by the addition of 1 M HCl (**3a**), afforded the desired product **4aa** along with three minor Wurtz-type homocoupling by-products (**7**, **8**, and **9**), in a favourable **4aa**/(**7**+**8**+**9**) ratio of 90/10 (see table in Section 3a, S7 in the ESI† for optimisation details). These results align with prior lithium zincate-assisted functionalisation studies on 4-iodobenzyl derivatives, where 2-MeTHF served as a bio-solvent that promoted sequential steps and minimised Wurtz-type by-products.²² Notably, using THF instead of 2-MeTHF reversed the product ratio, giving a 20/80 distribution of **4aa** to by-products, underscoring the role of the solvent in selectively driving the transformation sequence. A key difference in this reaction sequence compared to previous reports is the temperature requirement for the 1,2-migration homologation step. With **1a**, the homologation was completed at -85°C , whereas a warmer environment (-40°C) was necessary for similar transformations with 4-iodobenzyl mesylate substrates. The synthetic scope of the reaction was next examined under the optimised conditions (Table 1).

Various electrophiles were tested to achieve a second C–C bond formation five atoms away from the homologation site in a single operation. We found that the intermediate **III** (Scheme 1e), formed after the metallotropic rearrangement, was generally less reactive than its counterpart derived from 4-iodobenzyl mesylate.²² Nevertheless, it reacted readily with secondary aldehyde **3b** at -40°C over three hours to afford homothienyl alcohol **4ab** in an isolated yield of 73%. Notably, this transformation proceeded with complete selectivity, as no transfer of the butyl ligand to **3b** was observed. Several homothienyl alcohols were synthesised by reacting intermediate **III** with a range of aldehydes, including tertiary (**4ac**), enolisable primary (**4ad**), aromatic (**4ae–4ah**), and heteroaromatic (**4ai–4al**) aldehydes, with isolated yields generally exceeding 60%, except for **4af** (41%). In some cases, warming the reaction mixture from -40°C to -20°C after adding the electrophile improved yields. We also observed efficient C–Si bond formation with electrophile **3m**, yielding **4am** in 83% yield. Our approach was further validated for synthesising sterically hindered (**4an**, 60%) and less hindered (**4ao**, 70%) homothienyl carbonyl compounds. Sterically hindered products were selectively synthesised using the corresponding acid chlorides (e.g., **3n** for **4an**), while Weinreb amides proved essential for less hindered compounds, as PhCOCl mainly led to double addition by-products (see Section 3c, S8 for details, ESI†). Beyond $(n\text{-Bu})_3\text{ZnLi}$, other lithium trialkylzincates with more nucleophilic (**4bb**, **4bn**) or bulkier ligands (**4cb**, **4cc**, **4db**, **4eb**) were also compatible, demonstrating the versatility of this methodology. Additionally, methyl ligands, traditionally considered less nucleophilic, could participate in the 1,2-migration, as seen with Me_3ZnLi (**2f**), yielding **4fb** in fair yield (37%). This contrasts with earlier findings where only the hybrid zincate Me_2PhZnLi facilitated similar migrations.²² A major challenge in expanding the substrate scope



2: **R**¹ = **n-Bu** (2a), **n-Hex** (2b), **t-Bu** (2c), **i-Bu** (2d), **i-Pr** (2e), **Me** (2f), **Ph** (2g)

R² = **i-Pr** (3b), **t-Bu** (3c)

R³ = **H** (3e), **2-Et** (3f), **4-t-Bu** (3g), **C₆H₄** (3h)

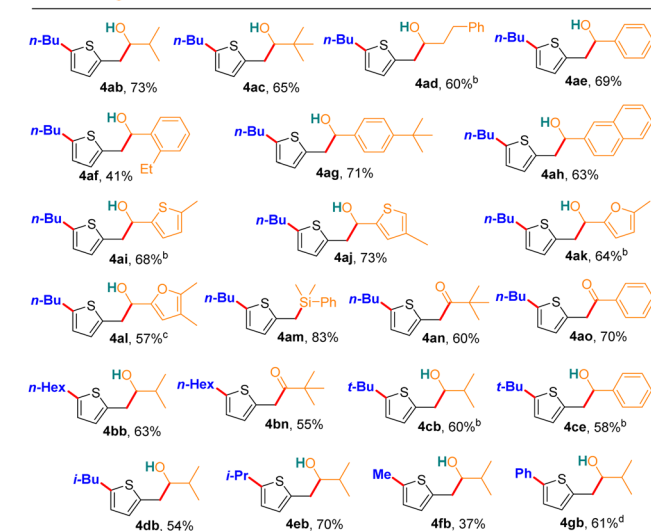
R⁴ = **5-Me** (3i), **4-Me** (3j)

R⁵ = **H** (3k), **Me** (3l)

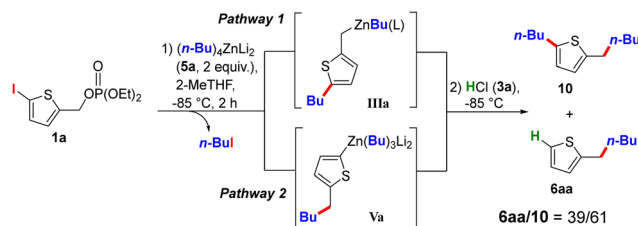
3m

t-BuCOCl (3n)

3d



The next objective was to develop a ligand-enabled divergent pathway relying on two independent reactions in a precise order: homologation *via* nucleophilic substitution of the phosphate by the zinc R¹ ligand, followed by Zn/X exchange metallation. Treating **1a** with 2 equivalents of the di-anionic reagent (*n*-Bu)₄ZnLi₂ (**5a**) in 2-MeTHF at –85 °C yielded a mixture of **6aa** and **10** in a 39/61 ratio after hydrolysis (Scheme 2), with pathway 1 favoured. The product **6aa** arises from hydrolysis of **Va** (pathway 2), while **10** results from the reaction of **IIIa** with *n*-BuI released during the I/Zn exchange (pathway 1). This result indicates that metallation proceeds faster than nucleophilic substitution, a finding further supported by solvent effects: replacing 2-MeTHF with THF reversed the **6aa/10** ratio, while using substrate **1b** (X = Br) delayed metallation, providing only **6aa** (see table in Section 3b, S7, for further details, ESI[†]).



Given the limited data on the reactivity of **Va**,²⁴ we investigated its reactions with aldehydes (Table 2). **Va** reacted successfully with secondary and tertiary aliphatic acyclic aldehydes (**6ab**, **6ac**, **6ap**) and cyclic aliphatic aldehydes (**6aq**, **6ar**, **6as**), yielding thienyl alcohols in 62% to 87%. This approach also enabled the synthesis of thienyl aldehyde **6at** (61%) and several ketones (**6an**, **6ao**, **6au**, **6av**), with Weinreb amides (**3o**, **3u**, **3v**) delivering higher yields than acid chlorides (**3n**). **Va** could also be successfully converted into ester compounds, as shown by the isolation of **6aw** and **6ax** in 65% and 76%, respectively.

Our method's versatility extends to zinc ligands beyond *n*-Bu. For instance, reagent **5b**, with *n*-hexyl ligands, yielded

5: $R^1 = n\text{-Bu}$ (5a), $n\text{-Hex}$ (5b), $t\text{-Bu}$ (5c), $i\text{-Bu}$ (5d), Me (5e), Ph (5f)

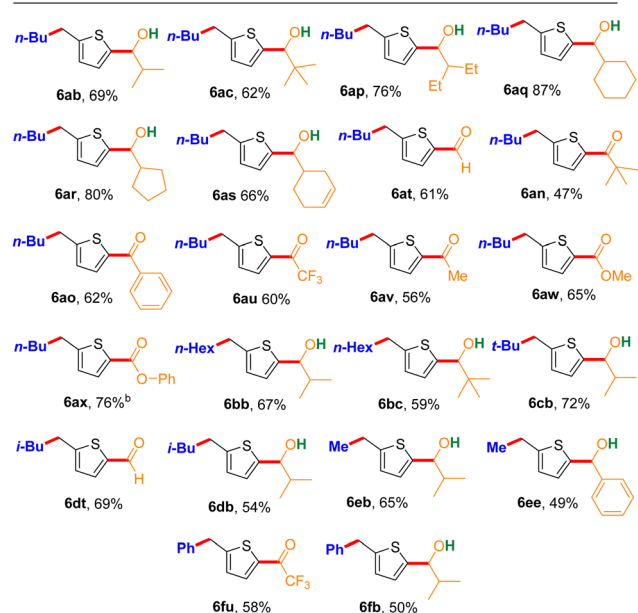
$R^2 =$

$i\text{-Pr}$ (3b), $t\text{-Bu}$ (3c) $n = 2$ (34), Et_2CH (3p), Ph (3e) $n = 1$ (3r)

3s , 3t , 3n

$R^3 = \text{Ph}$ (3o), CF_3 (3v), Me (3w), Ph (3x)

$R^4 = \text{Me}$ (3w), Ph (3x)



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products **6bb** and **6bc** in over 59% yield. Even bulkier ligands, such as those in **5c** (*t*-Bu) and **5d** (*i*-Bu), proved effective, as shown by products **6cb**, **6dt**, and **6db**, with yields between 54% and 72%. Notably, **5e**, containing less transferable ligands than *n*-Bu, afforded products **6eb** and **6ee** in fair yields (49–53%).

Finally, the presented one-pot strategy also supports C(sp³)–C(sp²) cross-coupling. This was demonstrated by the synthesis of **6fu** and **6fb**, with yields of 58% and 50%, respectively. To our knowledge, this achievement represents a rare C(sp³)–C(sp²) cross-coupling in the absence of a boronic acid and/or transition metal catalyst.²⁵ In total, nine constitutional isomers of the products shown in Table 2 were successfully synthesised, underscoring the broad applicability of this approach.

In summary, we developed a divergent, one-pot synthetic method for regioselective formation of two carbon–carbon bonds five atoms apart, leveraging the reactivity of lithium zincate (LiZn) element. By adjusting zinc's coordination environment, a single set of starting materials (ZnCl₂, organolithium R¹Li, and diethyl (5-halo)thienylphosphate) yielded distinct products: pathway 1 forms product **4** via a consecutive reaction sequence, while pathway 2 produces its constitutional isomer **6** through independent sequential reactions. This outcome underscores the versatility of lithium organozincates, with pathway 1 adapting our remote functionalisation approach²² to heterocycles for the first time and pathway 2 representing a new methodology. A critical factor was the solvent choice: bio-solvent 2-MeTHF promoted pathway 1, while THF favoured pathway 2. Across 44 examples, this method enabled heteroaryl-aryl couplings without transition metal and/or boronic acid activation, highlighting its synthetic versatility. Other heterocycles are currently under study, and the role of the bio-solvent is also being examined. The results will be disclosed in due course.

The authors acknowledge financial support from the French Agence Nationale de la Recherche (ANR-19-CE07-0032), the French Ministry of Higher Education and Research (MESR), and the Max Planck Society. They also thank F. Dupire (MAS-SLOR), P. Lemi re (SynBioN), D. Dodin for designing custom devices to enhance chiller functionality & the ‘‘Plateforme de RMN’’ (P le CPM/UL).

Open Access funding provided by the Max Planck Society.

Data availability

The data supporting this article are included in the ESI.†

Conflicts of interest

There are no conflicts to declare.

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