

# Direct Transformation of Esters into Arenes with 1,5-Bifunctional Organomagnesium Reagents

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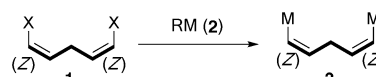
Dedicated to Professor Steven V. Ley on the occasion of his 70th birthday

**Abstract:** A direct transformation of carboxylic acid esters into arenes with 1,5-bifunctional organomagnesium reagents is described. This efficient and practical method enables the one-step defunctionalization of various carboxylic acid esters to prepare benzene, anthracene, tetracene, and pentacene derivatives. A double nucleophilic addition of the 1,5-organo-dimagnesium reagent to the ester is followed by an immediate 1,4-elimination reaction that leads to the direct [5+1] formation of a new aromatic ring.

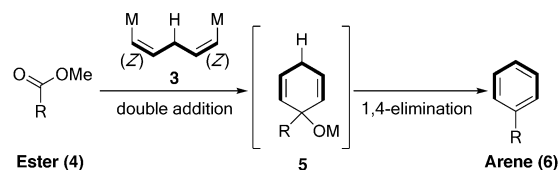
Organomagnesium reagents have an ideal reactivity for carbon–carbon bond forming reactions, as is evident from the enduring relevance of Grignard chemistry.<sup>[1]</sup> In recent years, we have experienced a dramatic improvement in the accessibility of organomagnesium compounds by the emergence of mild halogen–metal exchange methods. Seminal studies by Knochel and co-workers have shown that Grignard reagents bearing various reactive functional groups can be readily prepared and utilized.<sup>[2]</sup> Compared to the insertion of elemental magnesium into the C–X bond, these exchange reactions are characterized by an exceptional range of applications. For instance, Oshima et al. and researchers at Banyu Pharmaceuticals have reported on the highly efficient conversion of (Z)-alkenyl iodides without chelating groups to form the corresponding alkenylmagnesium compounds by using lithium trialkylmagnesates (R<sub>3</sub>MgLi).<sup>[3]</sup> These magnesiations proceed with complete retention of double-bond configuration and without undesired elimination reactions.

Considering this remarkable advancement, we anticipated the development of new synthetic methods based on bifunctional organomagnesium compounds resulting from a double halogen–metal exchange.<sup>[4]</sup> A stereospecific double halogen–metal exchange of (1Z,4Z)-1,5-dihalopenta-1,4-diene **1** with an exchange reagent **2** would lead to a 1,5-bifunctional organomagnesium reagent **3** (Scheme 1a). A subsequent double nucleophilic addition of reagent **3** to carboxylic acid ester **4** generates cyclohexa-2,5-dienolate **5**, which is directly transformed into an arene by means of a 1,4-elimination reaction (Scheme 1b).

## a) Double Halogen–Metal Exchange



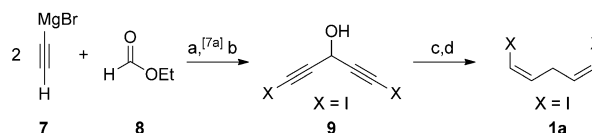
## b) Direct Ester to Arene Transformation



Scheme 1. a) Double halogen–metal exchange to form (1Z,4Z)-1,5-dimetalla-1,4-pentadiene **3**. b) Direct transformation of carboxylic acid ester **4** into arene **6** through the double addition of (1Z,4Z)-1,5-dimetalla-1,4-pentadiene **3** followed by a 1,4-elimination.

An analogous [5+1]-formation of an aromatic ring with known methods would require several stages and harsh reaction conditions.<sup>[5]</sup> Given the high availability of carboxylic acid esters, a mild one-step process would give expedient access to compounds typically prepared by transition-metal-catalyzed cross-coupling reactions from complementary sub-strates.<sup>[6]</sup> Intrigued by this prospect, we evaluated the feasibility of a direct conversion of esters into arenes with 1,5-bifunctional organomagnesium compounds.

We started our investigations with the development of a concise synthesis of (1Z,4Z)-1,5-diiodopenta-1,4-diene (**1a**) from readily available starting materials (Scheme 2).<sup>[7]</sup> Two equivalents of ethynylmagnesium bromide (**7**) were added to ethyl formate (**8**), followed by diiodination with N-iodosuccinimide. The subsequent (Z)-selective double alkyne reduction with diimide and a trifluoroacetic acid mediated dehydroxylation with triethylsilane provided (1Z,4Z)-1,5-diiodopenta-1,4-diene (**1a**) in 41% yield over four steps.<sup>[8]</sup>



Scheme 2. Synthesis of (1Z,4Z)-1,5-diiodopenta-1,4-diene **1a**. a) THF, RT; b) N-Iodosuccinimide, AgNO<sub>3</sub>, acetone, RT; c) KO<sub>2</sub>CN=NCO<sub>2</sub>K, pyridine, AcOH, MeOH, RT, then aqueous HCl; d) Et<sub>3</sub>SiH, CF<sub>3</sub>CO<sub>2</sub>H, CH<sub>2</sub>Cl<sub>2</sub>, 0 °C; 41% over four steps.

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**Table 1:** In situ generation of reagent **3**<sup>[a]</sup> and optimization of the reaction parameters for the direct ester to arene transformation.<sup>[b]</sup>

**1a**  $\xrightarrow[\text{THF, } T]{\mathbf{2}}$  **3**

**4a**  $\xrightarrow[\text{THF, } T, t]{\mathbf{3}}$  **6a**

Entry	X–M Exchange Reagent ( <b>2</b> )	T [°C]	t	Yield [%] <sup>[c]</sup>
1	<i>n</i> BuLi <sup>[d]</sup>	–40	2 h	46
2	<i>i</i> PrnBu <sub>2</sub> MgLi <sup>[e]</sup>	0	30 min	70
3	<i>i</i> PrnBu <sub>2</sub> MgLi	0	30 min	70
4	<i>n</i> Bu <sub>3</sub> MgLi	0	30 min	64
5	<i>s</i> BunBu <sub>2</sub> MgLi	0	30 min	56
6	<i>i</i> PrnBu <sub>2</sub> MgLi	–20	30 min	69
7	<i>i</i> PrnBu <sub>2</sub> MgLi	–20	2 h	<b>82</b>
8	<i>i</i> PrnBu <sub>2</sub> MgLi	–40	2 h	61
9	<i>i</i> PrnBu <sub>2</sub> MgLi <sup>[f]</sup>	–20	2 h	70
10	<i>i</i> PrnBu <sub>2</sub> MgLi <sup>[g]</sup>	–20	2 h	73

[a] Reactions performed with 200 μmol **1a** and 200 μmol **2** for 5 min at *T*. [b] 100 μmol **4a** and in situ generated **3** for *t* at *T* followed by aqueous work-up (HCl 1.0 mol L<sup>–1</sup>). [c] Yield of isolated product **6a**. [d] Reaction was performed in Et<sub>2</sub>O. [e] Purchased from Sigma–Aldrich, No. 683418. [f] Reagent **3a** was kept 2 h at –20 °C prior the reaction with **4a**. [g] 160 μmol **1a** and 160 μmol **2**.

We next explored different methods for the double metalation to form 1,5-organodimetallic reagents **3** (Table 1, M = Li or Mg). The two-fold iodine–lithium exchange at 40 °C described by Cutzi led to a highly reactive dilithium species<sup>[9]</sup> capable of converting methyl 4-phenylbenzoate (**4a**) into p-terphenyl (**6a**) in a promising yield of 46% (Table 1, entry 1). Since no product formation was observed after a transmetalation reaction with MgX<sub>2</sub> and since the direct reaction of **1a** with elemental magnesium was ineffective to give the corresponding organomagnesium compound, we focused on (Z)-alkenyliodide–magnesium exchange reactions.<sup>[10]</sup> However, even after prolonged reaction times, the reaction of **1a** with *i*PrMgCl·LiCl resulted mainly in monometalation. In contrast, *s*Bu<sub>2</sub>Mg·LiCl enabled complete halogen–metal exchange at both the 1- and 5-position. Nevertheless, the resulting reagent was unproductive in the conversion of **4a** to the desired p-terphenyl (**6a**).<sup>[11]</sup> Given the extraordinary modulation aptitude of magnesate reagents, we treated **1a** with equimolar amounts of commercially available heteroleptic *i*PrnBu<sub>2</sub>MgLi.<sup>[12]</sup> Even at 0 °C, (Z,Z)-1,4-pentadiene-1,5-diyl **3a**<sup>[13]</sup> was efficiently formed within 5 min and a significant increase in yield for the direct carboxylic acid ester to phenyl conversion was achieved (Table 1, entry 2, 70%). Alternatively, *i*PrnBu<sub>2</sub>MgLi can effortlessly be prepared from *i*PrMgCl and two equivalents of *n*BuLi (Table 1, entry 3). We further examined *n*Bu<sub>3</sub>MgLi and *s*Bu-*n*Bu<sub>2</sub>MgLi to identify *i*PrnBu<sub>2</sub>MgLi as the ideal halogen–metal exchange reagent (Table 1, entries 4

and 5 versus entry 3). A marked improvement in yield was obtained after optimizing the reaction time and temperature, and subsequent reactions were carried out for 2 h at 20 °C. The best results were obtained by the immediate use of the in situ prepared reagent **3a** and a compensation of decomposition pathways with an excess of the reagent (Table 1, entries 9 and 10 versus entry 7).

With the optimized reaction conditions in hand, we explored the substrate scope of the direct transformation of carboxylic acid esters into benzenes (Table 2). A comparable outcome was obtained with substrates bearing electron-withdrawing groups, whereas the lower reactivity of ester substrates with electron-donating groups was evident from a lower yield (Table 2, entry 1 versus entries 2–4). Nonetheless, phenyl-substituted heterocyclic compounds could be prepared in synthetically meaningful yields (Table 2, entries 5 and 6). Even alkyl and alkenyl esters were amenable to form the corresponding benzene derivatives **6g** and **6h**, thus highlighting the broad applicability of the method (Table 2, entries 7 and 8). Furthermore, when using an additional equivalent of reagent **3a**, even a protic substrate was tolerated and efficiently converted into the corresponding benzene derivative **6i** (Table 2, entry 9, 80%).

Having developed a method for the direct conversion of carboxylic acid esters into benzene derivatives, we turned our attention to 1,5-bifunctional diarylmagnesium reagents **3b–3d** for the transformation of esters into anthracene, tetracene, and pentacene derivatives. The direct addition of metallic

**Table 2:** Scope of the direct ester to benzene transformation using 1,5-bifunctional organomagnesium reagent **3a**.<sup>[a]</sup>

**4**  $\xrightarrow[\text{THF, } -20\text{ }^\circ\text{C, } 2\text{ h}]{\mathbf{3a}}$  **6**

Entry	Product <sup>[b]</sup>	Entry	Product <sup>[b]</sup>	Entry	Product <sup>[b]</sup>
1	 <b>6a</b> , 82%	4	 <b>6d</b> , 57% <sup>[c]</sup>	7	 <b>6g</b> , 68%
2	 <b>6b</b> , 80%	5	 <b>6e</b> , 72%	8	 <b>6h</b> , 60%
3	 <b>6c</b> , 59%	6	 <b>6f</b> , 68%	9	 <b>6i</b> , 80% <sup>[c]</sup>

[a] Reactions performed with 100 μmol **4** and in situ generated **3a** (from 200 μmol **1a** and 200 μmol **2**) for 2 h at –20 °C followed by aqueous work-up (HCl 1.0 mol L<sup>–1</sup>). [b] Yield of isolated product. [c] From 300 μmol **1a** and 300 μmol **2**.

Table 3: Scope of the direct transformation of esters into anthracenes, tetracenes, and pentacenes using 1,5-bifunctional organomagnesium reagents 3b–3d.<sup>[a]</sup>

Entry	Product <sup>[b]</sup>	Entry	Product <sup>[b]</sup>	Entry	Product <sup>[b]</sup>
1		7		11	
	6j, 99%		6n, 87%		6r, 61%
2–4		8		12	
	6k, 97% 99%, <sup>[c]</sup> 94% <sup>[d]</sup>		6o, 55% <sup>[e]</sup>		6s, 89%
5		9		13	
	6l, 99%		6p, 88% <sup>[f]</sup>		6t, 97%
6		10		14	
	6m, 97%		6q, 99% <sup>[g]</sup>		6u, 82%

[a] Reactions performed with 100 mmol 4 in 1.0 mL THF and 140 mmol 3 for 4 h at RT followed by aqueous workup (HCl 1.0 mol L<sup>-1</sup>). [b] Yield of isolated product. [c] 1.00 mmol scale. [d] 10.0 mmol scale. [e] Threefold ester to anthracene transformation with 420 mmol 3b; the third aromatization was induced in a separate step through treatment with conc. HCl [f] 200 mmol 3b. [g] 240 mmol 3b.

magnesium to *o,o'*-dibromodiarlylmethanes led to the notably more stable reagents 3b–3d.<sup>[14]</sup> To our delight, the union of 3b and ester 4a resulted in the formation of anthracene derivative 6j with extraordinary efficiency (Table 3, entry 1, 99%). A more electron-rich aryl ester could also be converted into the biaryl product in excellent yield (Table 3, entry 2). The scalability of the reaction was verified by increasing the reaction scale (Table 3, entries 3 and 4, 1.00 mmol and 10.0 mmol). Moreover, the phenylanthracene with a bromo substituent and the naphthyl derivative with increased steric demand could also be obtained in remarkable yields (Table 3, entries 5 and 6, 99% and 97%). The reaction of an alkenylester underlines the wide substrate scope (Table 3, entry 7, 87%) and a triple carboxylic acid ester to arene transformation highlights the robustness of the reaction

(Table 3, entry 8, 55% for three anthracene formations). With an excess of reagent 3b, esters with protic functional groups could also be efficiently converted (Table 3, entries 9 and 10). Particularly gratifying was that tetracene derivatives substituted at the 5-position were also obtained in high yields (Table 3, entries 11 and 12). We furthermore found that by working in the dark with degassed THF, even monosubstituted pentacene derivatives were accessible after isolation by crystallization (Table 3, entries 13 and 14). These synthetically challenging monosubstituted pentacenes emphasize the advantages of the direct ester to arene transformation. The mild reaction conditions of the method also allow the preparation of sensitive compounds in high yields.

In summary, we have developed a method for the direct transformation of carboxylic acid esters into the corresponding benzene and acene derivatives by the use of 1,5-bifunctional organomagnesium reagents. An *in situ* iodine–magnesium exchange with lithium trialkylmagnesates enabled the generation of dialkenyldimagnesium reagents for the formation phenyl derivatives. In contrast, the diaryldimagnesium reagents were accessible when using metallic magnesium from *o,o'*-dibromodiarlylmethane precursors, which allowed the efficient synthesis of anthracenes, tetracenes, and pentacenes. Treatment of carboxylic acid esters with the 1,5-bifunctional organomagnesium reagents enabled the single-step preparation of various arenes under mild reaction conditions and in yields of up to 99%. We expect that this method will find a wide range of applications in discovery chemistry and sustainable process research, particularly in the synthesis of organic functional materials. Future studies will evaluate other selected substrate and reagent classes, the structure of the 1,5-bifunctional organomagnesium reagents, and the mechanism of this practical preparative method.

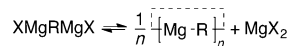
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Keywords: acenes · arenes · carboxylic acid esters · Grignard reactions · organomagnesium reagents

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