


 Cite this: *Chem. Commun.*, 2026, 62, 6931

 Received 12th December 2025,  
Accepted 25th February 2026

DOI: 10.1039/d5cc07035j

rsc.li/chemcomm

**We report the selective nucleophilic addition of aryllithium species to acyl cyanides—electrophiles bearing both carbonyl and cyano functional groups—under catalyst-free continuous flow conditions. Using benzoyl cyanide as a model substrate, we demonstrate that the flow microreactor system enables highly selective addition to the carbonyl moiety while eliminating the cyano group as a leaving group, affording polyfunctional ketones in significantly improved yields compared to traditional batch processes.**

The acylation of organometallic intermediates with acid chlorides represents a classical and widely utilized strategy for the synthesis of polyfunctional ketones, which are key structural motifs in numerous pharmaceuticals and materials science applications.<sup>1–3</sup> A broad spectrum of organometallic reagents—including organomanganese, organozinc, and organomagnesium compounds—have been developed and applied in acylation reactions.<sup>4–8</sup> While organomanganese reagents have demonstrated particular utility in such transformations, polyfunctional organozinc reagents, especially when combined with stoichiometric copper reagent<sup>9–11</sup> or palladium catalysis, have also enabled efficient access to functionalized ketone frameworks.<sup>12,13</sup> The preparation of functionalized arylzinc species remains synthetically challenging,<sup>14</sup> often necessitating cobalt catalysis<sup>15–17</sup> or the use of highly reactive activated zinc.<sup>18–20</sup> Arylmagnesium halides can be readily generated *via* halogen–magnesium exchange methods and have been successfully employed in the synthesis of multifunctional diarylketones through acylation reactions.<sup>21–23</sup> Previous studies have demonstrated moderate success in these transformations, with yields typically ranging from 50% to 58% when employing benzoyl chloride as the electrophile, despite optimization efforts and catalyst screening.<sup>24–39</sup> These results align with broader literature observations indicating that acylation of organometallics with acid chlorides often proceeds with only moderate efficiency.<sup>3</sup> A significant improvement in acylation efficiency was achieved by replacing acid chlorides with

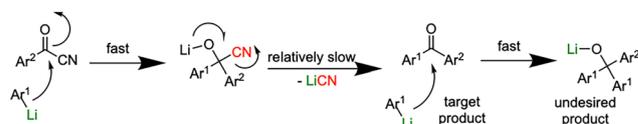
## Selective nucleophilic addition of aryllithium reagents to $\alpha$ -ketonitriles giving ketones under microflow conditions

 Mohmmad S. Qenawy,<sup>1</sup> Kazuhiro Okamoto<sup>2</sup> and Aiichiro Nagaki<sup>1\*</sup>

acyl cyanides.<sup>40–42</sup> Acyl cyanides exhibit enhanced electrophilicity due to the strong electron-withdrawing nature of the cyano group.

Unlike the chlorine substituent in acid chlorides, the cyano group withdraws electron density from the adjacent carbonyl, thereby increasing its susceptibility to nucleophilic attack (Scheme 1).<sup>43</sup> Given the high reactivity of organolithium species and their propensity for over-addition or decomposition under conventional batch conditions, we hypothesized that flow microreactor technology could overcome these limitations by offering rapid mixing, precise temperature control, and minimized residence times.<sup>44–65</sup> Here we report the unique reactivity of acyl cyanides towards aryllithium reagents under microflow conditions. While prior studies predominantly focused on organomagnesium species using batch processes, the chemoselectivity of highly reactive aryllithium intermediates toward bifunctional electrophiles bearing both carbonyl and cyano groups remains underexplored under flow microreactor systems.<sup>66</sup>

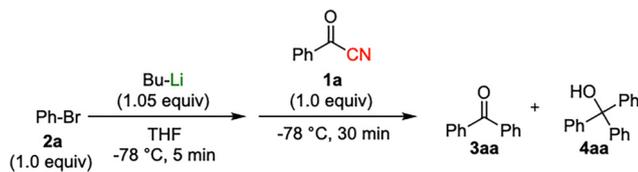
Initial study begun with the investigation of the chemoselectivity of aryllithium reagents toward bifunctional electrophiles bearing both carbonyl and cyano groups under flow conditions, employing benzoyl cyanide as a model electrophile (Scheme 2). In a batch reactor system, aryllithium intermediates were generated *in situ via* halogen–lithium exchange between bromobenzene and *n*-butyllithium (*n*-BuLi) at  $-78\text{ }^{\circ}\text{C}$ , followed by the reaction with 1 equiv. of benzoyl cyanide (Scheme 1). The desired benzophenone (**3aa**) was obtained in 15% yield because of nucleophilic addition of the aryllithium species to the carbonyl group of benzoyl cyanide, accompanied by the formation of triphenylmethanol (**4aa**) in



**Scheme 1** Reaction pathways for nucleophilic addition of  $\text{Ar}^1\text{Li}$  to acyl cyanide  $\text{Ar}^2\text{COCN}$ .

Department of Chemistry, Faculty of Science, Hokkaido University, Kita 10-jo, Nishi 8-chome, Kita-ku, Sapporo 060-0810, Japan



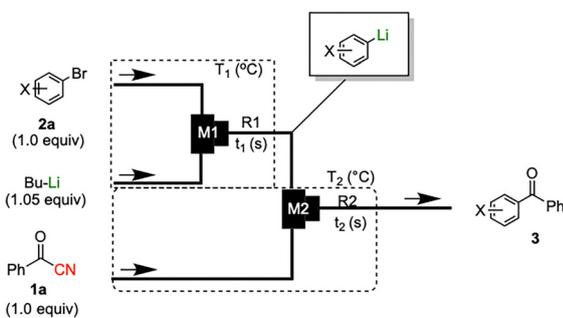


Scheme 2 Reaction of benzoyl cyanide (**1a**) and PhLi generated from bromobenzene (**2a**).

42% yield. Given the high reactivity of organolithium species, any unreacted aryllithium species could readily undergo secondary addition to the ketone intermediate.

The same transformation was subsequently examined under continuous flow conditions utilizing a microreactor system composed of two T-shaped micromixers and two microtube reactors (Scheme 3). A systematic investigation of the reaction temperature was conducted wherein the temperature at the second micromixer was varied between 0 °C and -78 °C. A pronounced temperature dependence was observed, whereby both the yield of the desired product **3aa** and the overall chemoselectivity were markedly enhanced under flow conditions relative to the corresponding batch reaction. The reaction yield progressively increased as the temperature decreased, reaching a maximum at -78 °C, at which the highest yield and selectivity were obtained (Fig. 1). These findings underscore the critical role of low-temperature operation in optimizing reaction efficiency under flow conditions. Importantly, to evaluate the practical applicability of the protocol, a gram-scale experiment was performed under the optimized flow conditions, affording product **3aa** in 87% isolated yield. This result demonstrates the scalability and synthetic utility of the methodology (see SI, p. S4, for details). A series of experiments were conducted wherein the total flow rate was maintained constant at 10.5 mL min<sup>-1</sup>, while the internal diameter of micromixer M2 was systematically varied. As the diameter increased, both the yield of product **3aa** and the overall selectivity decreased (Fig. 2), underscoring the critical importance of efficient mixing. Notably, the highest yield was obtained using a micromixer with an inner diameter of 250 μm at a flow rate of 10.5 mL min<sup>-1</sup>.

With the optimized conditions established, we subsequently explored the scope of the reaction by employing a range of



Scheme 3 A flow microreactor system for reaction of benzoyl cyanide **1a** and an aryllithium reagent generated from aryl halide **2a**. T<sub>1</sub> = 0 °C. When total flow rate is 10.5 mL min<sup>-1</sup>, t<sub>1</sub> = 1.7 s, t<sub>2</sub> = 2.2 s.

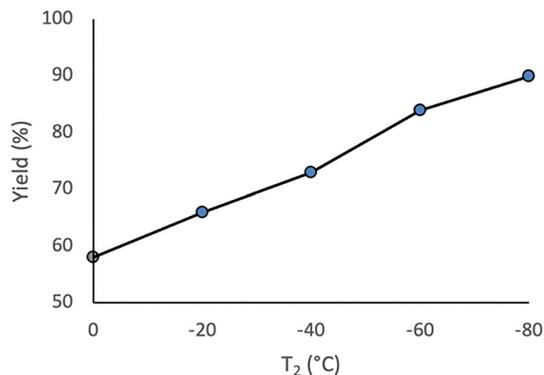


Fig. 1 A flow microreactor system for reaction of benzoyl cyanide (**1a**) and PhLi generated from bromobenzene (**2a**): the effect of the temperature.

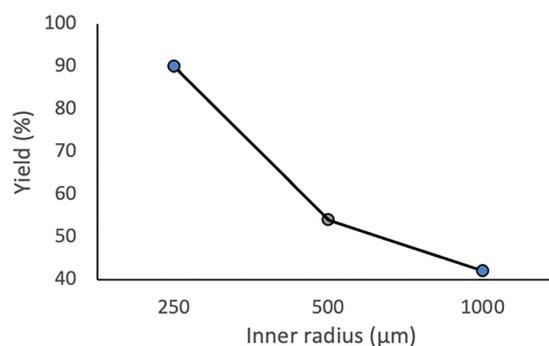


Fig. 2 A flow microreactor system for reaction of benzoyl cyanide (**1a**) and PhLi generated from bromobenzene (**2a**): the effect of the mixing. Flow rate at M2: 10.5 mL min<sup>-1</sup>. Reaction's temperature: T<sub>2</sub> = -78 °C.

aryllithium reagents bearing various electrophilic functional groups, with particular emphasis on evaluating the influence of steric hindrance (Table 1). Reactions involving benzoyl cyanide and aryllithium reagents substituted with either electron-donating or electron-withdrawing groups furnished the desired products in yields ranging from 71% to 90%. Aryllithium species bearing electron-donating substituents such as methyl, methoxy and isobutyl groups were well tolerated under the optimized flow conditions and consistently delivered the corresponding diaryl ketones in high yields, indicating that increased electron density on the aromatic ring does not adversely affect the nucleophilic acylation process. For example, the reaction of (4-cyanophenyl)lithium with benzoyl cyanide afforded product **3ae** in 71% yield. These high yields are attributed to the advantages of the flash microflow system, which facilitates the efficient generation and immediate consumption of short-lived aryllithium intermediates through rapid mixing and precise control over temperature and residence time, thereby minimizing byproduct formation. Furthermore, the applicability of this methodology was confirmed by extending the protocol to other α-keto nitriles, such as acetyl cyanide and ethyl cyanofornate, which similarly afforded high yields under flow conditions.



Table 1 Selective reaction of  $\alpha$ -Keto nitriles **1** with organolithium reagents generated from aryl halides **2** using flow microreactors

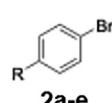
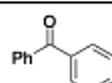
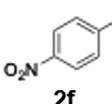
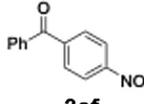
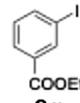
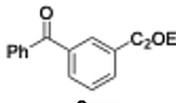
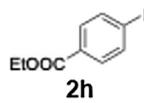
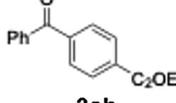
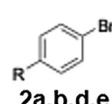
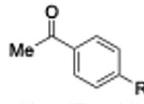
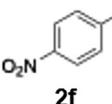
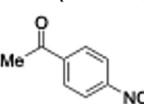
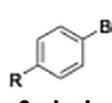
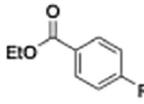
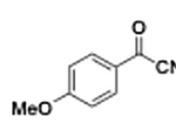
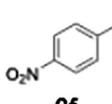
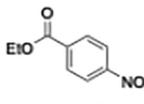
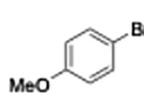
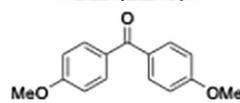
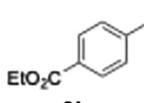
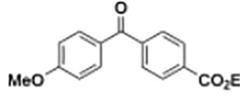
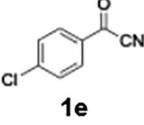
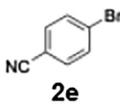
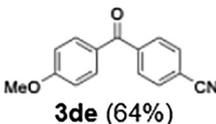
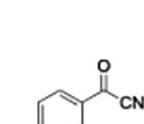
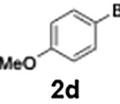
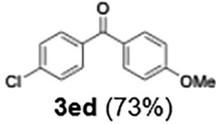
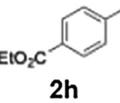
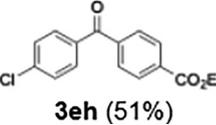
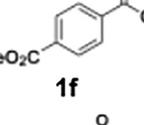
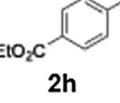
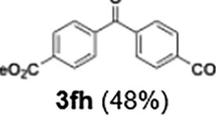
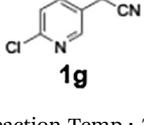
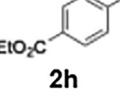
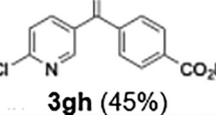
$\alpha$ -Keto nitriles	Aryl Halide	Product	Yield (%)			
 <b>1a</b>	 <b>2a-e</b>	 <b>3a</b> (R = H)	90			
		<b>3ab</b> (R = Me)	74			
		<b>3ac</b> (R = <i>i</i> Bu)	80			
		<b>3ad</b> (R = OMe)	75			
		<b>3ae</b> (R = CN)	71			
 <b>1b</b>	 <b>2f</b>	 <b>3af</b>	77			
		 <b>2g</b>	 <b>3ag</b>	53		
			 <b>2h</b>	 <b>3ah</b>	54	
				 <b>2a,b,d,e</b>	 <b>3ba</b> (R = H)	76
					<b>3bb</b> (R = Me)	72
<b>3bd</b> (R = OMe)	70					
 <b>1c</b>	 <b>2f</b>	 <b>3bf</b>	61			
		 <b>2a,b,d,e</b>	 <b>3ca</b> (R = H) (65%)	65		
			<b>3cb</b> (R = Me) (64%)	64		
			<b>3cd</b> (R = OMe) (64%)	64		
			<b>3ce</b> (R = CN) (74%)	74		
 <b>1d</b>	 <b>2f</b>	 <b>3cf</b> (62%)	62			
		 <b>2d</b>	 <b>3dd</b> (77%)	77		
			 <b>2h</b>	 <b>3dh</b> (56%)	56	



Table 1 (continued)

$\alpha$ -Keto nitriles	Aryl Halide	Product	Yield (%)
			64
			73
			51
			48
			45

Reaction Temp.:  $T_1 = 0\text{ }^\circ\text{C}$ . When total flow rate is  $10.5\text{ mL min}^{-1}$ ,  $t_1 = 1.7\text{ s}$ ,  $t_2 = 2.2\text{ s}$ .

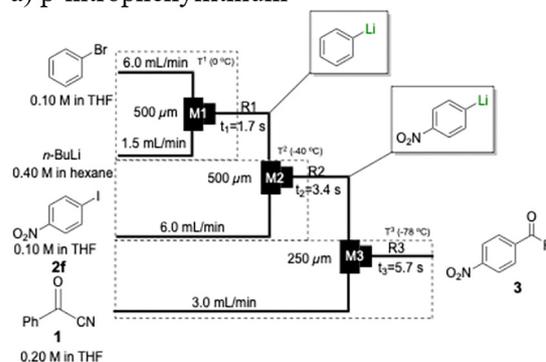
The generation of highly short-lived aryllithium species involving nitro and alkoxy carbonyl moieties presented considerable challenges due to the well-documented instability of the corresponding nucleophilic intermediates, which are prone to rapid decomposition under conventional batch conditions.<sup>65</sup> For the generation of nitrophenyllithium species, freshly generated PhLi with microflow mixing was immediately transferred to micromixer M2, maintained at  $-40\text{ }^\circ\text{C}$ , where it underwent halogen-lithium exchange with *p*-nitrophenyl iodide (**2f**), affording the desired aryllithium intermediate (Scheme 4a).

The acylation of *meta*- and *para*-substituted ethyl iodobenzoates presented significant synthetic challenges, largely due to their reduced reactivity under conventional halogen-lithium exchange conditions.

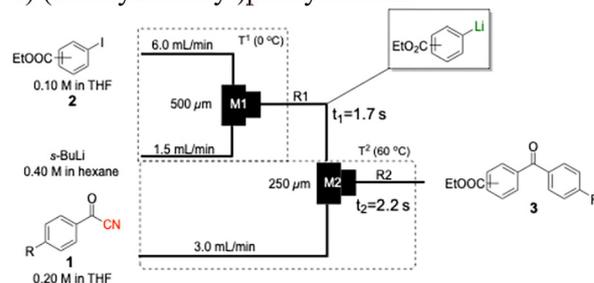
Efficient activation of these substrates could not be achieved using standard protocols. Subsequent nucleophilic addition to  $\alpha$ -ketonitriles (**1**) was performed at micromixer M3 under a range of temperatures, with optimal results obtained at  $-78\text{ }^\circ\text{C}$ , affording the desired product in excellent yields of 61–77% with high chemoselectivity. Specifically, ethyl iodobenzoates required activation with <sup>s</sup>BuLi instead of BuLi, and the exchange needed to be performed at  $-60\text{ }^\circ\text{C}$  rather than  $0\text{ }^\circ\text{C}$  to ensure efficient generation of the corresponding aryllithium intermediates (Scheme 4b). Furthermore, we extended the methodology to reactions between ethyl 4-iodobenzoate (**2h**) and  $\alpha$ -keto nitriles bearing electronically diverse substituents, including 4-methoxybenzoyl cyanide (**1d**) as an electron-donating group and 4-chlorobenzoyl cyanide (**1e**) as an electron-withdrawing group. In both cases, the corresponding products were obtained in good

yields without the use of any catalyst, demonstrating the generality and robustness of the catalyst-free flow protocol across electronically distinct substrates.

#### a) *p*-nitrophenyllithium



#### b) (ethoxycarbonyl)phenyllithium



Scheme 4 Flash-flow generation of short-lived aryllithium intermediates. (a) *p*-nitrophenyllithium; (b) (ethoxycarbonyl)phenyllithium.



In summary, we have developed a highly efficient and chemo-selective method for the synthesis of polyfunctional ketones through the nucleophilic addition of aryllithium reagents to acyl cyanides under continuous flow conditions. By utilizing a flow microreactor system without the need for any external catalysts, we achieved significant improvements in reaction yield and selectivity compared to conventional batch processes. This study demonstrates the utility of flow microreactor technology as a powerful tool for controlling highly reactive organolithium intermediates, enabling catalyst-free access to polyfunctional ketones with high efficiency and selectivity, and offers a valuable approach for future applications in complex molecule synthesis.

## Conflicts of interest

There are no conflicts to declare.

## Data availability

The data supporting this article have been included as part of the supplementary information (SI). Supplementary information is available. See DOI: <https://doi.org/10.1039/d5cc07035j>.

## Acknowledgements

This work was supported by MEXT KAKENHI number JP24H01051 (Digi-TOS), and JSPS KAKENHI numbers, JP23K04744, JP20KK0121, and JP21H01936. This work was also partially supported by AMED PJ24126027 (GAP-FREE), PJ22230021 (BINDS), NEDO PJ22240036, JKA Foundation, and Takahashi Industrial and Economic Foundation.

## References

- 1 *Modern Organocopper Chemistry*, ed. N. Krause, Wiley, 2002.
- 2 N. J. Lawrence, *J. Chem. Soc., Perkin 1*, 1998, 1739–1750.
- 3 R. K. Dieter, *Tetrahedron*, 1999, 55, 4177–4236.
- 4 G. Cahiez, *Tetrahedron Lett.*, 1981, 22, 1239–1242.
- 5 G. Cahiez and B. Laboue, *Tetrahedron Lett.*, 1989, 30, 3545–3546.
- 6 G. Cahiez and B. Laboue, *Tetrahedron Lett.*, 1989, 30, 7369–7372.
- 7 G. Cahiez and B. Laboue, *Tetrahedron Lett.*, 1992, 33, 4439–4442.
- 8 G. Cahiez, P.-Y. Chavand and E. Metais, *Tetrahedron Lett.*, 1992, 33, 5245–5248.
- 9 P. Knochel, N. Millot, A. L. Rodriguez and C. E. Tucker, *Organic Reactions*, Wiley, 2001, pp. 417–759.
- 10 M. J. Rozema, A. Sidduri and P. Knochel, *J. Org. Chem.*, 1992, 57, 1956–1958.
- 11 P. Knochel, M. C. P. Yeh, S. C. Berk and J. Talbert, *J. Org. Chem.*, 1988, 53, 2390–2392.
- 12 D. Wang and Z. Zhang, *Org. Lett.*, 2003, 5, 4645–4648.
- 13 E. Negishi, V. Bagheri, S. Chatterjee, F.-T. Luo, J. A. Miller and A. T. Stoll, *Tetrahedron Lett.*, 1983, 24, 5181–5184.
- 14 T. N. Majid and P. Knochel, *Tetrahedron Lett.*, 1990, 31, 4413–4416.
- 15 H. Fillon, C. Gosmini and J. Périchon, *Tetrahedron*, 2003, 59, 8199–8202.
- 16 I. Kazmierski, C. Gosmini, J.-M. Paris and J. Périchon, *Tetrahedron Lett.*, 2003, 44, 6417–6420.
- 17 H. Fillon, C. Gosmini and J. Périchon, *J. Am. Chem. Soc.*, 2003, 125, 3867–3870.
- 18 R. M. Wehmeyer and R. D. Rieke, *Tetrahedron Lett.*, 1988, 29, 4513–4516.
- 19 R. D. Rieke, *Science*, 1989, 246, 1260–1264.

- 20 A. Guijarro, D. M. Rosenberg and R. D. Rieke, *J. Am. Chem. Soc.*, 1999, 121, 4155–4167.
- 21 A. E. Jensen, W. Dohle, I. Sapountzis, D. M. Lindsay, V. A. Vu and P. Knochel, *Synthesis*, 2002, 565–569.
- 22 P. Knochel, W. Dohle, N. Gommermann, F. F. Kneisel, F. Kopp, T. Korn, I. Sapountzis and V. A. Vu, *Angew. Chem., Int. Ed.*, 2003, 42, 4302–4320.
- 23 H. Liu, Y.-X. Chai, J.-J. Yang, D. Miao, M.-Y. Wang, T. Jiang, Y.-Y. Sun, X. Lu, Q. Sun, J.-H. Li and Y.-P. Zhu, *Org. Lett.*, 2024, 26(2), 477–482.
- 24 W. Dohle, F. Kopp, G. Cahiez and P. Knochel, *Synlett*, 2001, 1901–1904.
- 25 C. K. Reddy and P. Knochel, *Angew. Chem., Int. Ed. Engl.*, 1996, 35, 1700–1701.
- 26 A. Fürstner, D. De Souza, L. Parra-Rapado and J. T. Jensen, *Angew. Chem., Int. Ed.*, 2003, 42, 5358–5360.
- 27 A. Fürstner and M. Méndez, *Angew. Chem., Int. Ed.*, 2003, 42, 5355–5357.
- 28 A. Fürstner and A. Leitner, *Angew. Chem., Int. Ed.*, 2002, 41, 609–612.
- 29 A. Fürstner, A. Leitner, M. Méndez and H. Krause, *J. Am. Chem. Soc.*, 2002, 124, 13856–13863.
- 30 G. Cahiez and H. Avedissian, *Synthesis*, 1998, 1199–1205.
- 31 G. Cahiez and S. Marquais, *Tetrahedron Lett.*, 1996, 37, 1773–1776.
- 32 A. J. Birch, *Pure Appl. Chem.*, 1996, 68, 553–556.
- 33 R. S. Smith and J. K. Kochi, *J. Org. Chem.*, 1976, 41, 502–509.
- 34 M. Tamura and J. Kochi, *J. Organomet. Chem.*, 1971, 31, 289–309.
- 35 M. Tamura and J. Kochi, *Synthesis*, 1971, 303–305.
- 36 J. K. Kochi and M. Tamura, *J. Am. Chem. Soc.*, 1971, 93, 1485–1487.
- 37 V. Fiandanese, G. Marchese and L. Ronzini, *Tetrahedron Lett.*, 1983, 24, 3677–3680.
- 38 V. Fiandanese, G. Marchese, V. Martina and L. Ronzini, *Tetrahedron Lett.*, 1984, 25, 4805–4808.
- 39 C. Cardellicchio, V. Fiandanese, G. Marchese and L. Ronzini, *Tetrahedron Lett.*, 1987, 28, 2053–2056.
- 40 S. Hünig and R. Schaller, *Angew. Chem., Int. Ed. Engl.*, 1982, 21, 36–49.
- 41 A. J. Bellamy and L. Maclean, *J. Chem. Soc., Perkin 1*, 1979, 204.
- 42 J. Thesing, D. Witzel and A. Brehm, *Angew. Chem.*, 1956, 68, 425–435.
- 43 C. Duplais, F. Bures, I. Sapountzis, T. J. Korn, G. Cahiez and P. Knochel, *Angew. Chem., Int. Ed.*, 2004, 43, 2968–2970.
- 44 J. Yoshida, H. Kim and A. Nagaki, *ChemSusChem*, 2011, 4, 331–340.
- 45 D. Webb and T. F. Jamison, *Chem. Sci.*, 2010, 1, 675.
- 46 S. Marre and K. F. Jensen, *Chem. Soc. Rev.*, 2010, 39, 1183.
- 47 W.-Y. Lin, Y. Wang, S. Wang and H.-R. Tseng, *Nano Today*, 2009, 4, 470–481.
- 48 P. Watts and C. Wiles, *Chem. Commun.*, 2007, 443–467.
- 49 B. Ahmed-Omer, J. C. Brandt and T. Wirth, *Org. Biomol. Chem.*, 2007, 5, 733–740.
- 50 B. P. Mason, K. E. Price, J. L. Steinbacher, A. R. Bogdan and D. T. McQuade, *Chem. Rev.*, 2007, 107, 2300–2318.
- 51 M. Brivio, W. Verboom and D. N. Reinhoudt, *Lab Chip*, 2006, 6, 329.
- 52 J. Kobayashi, Y. Mori and S. Kobayashi, *Chem. – Asian J.*, 2006, 1, 22–35.
- 53 A. J. deMello, *Nature*, 2006, 442, 394–402.
- 54 K. Geyer, J. D. C. Codée and P. H. Seeberger, *Chem. – Eur. J.*, 2006, 12, 8434–8442.
- 55 P. Watts and S. J. Haswell, *Chem. Soc. Rev.*, 2005, 34, 235.
- 56 P. Natho and R. Luisi, *Tetrahedron, Green Chem.*, 2023, 2, 100015.
- 57 T. Zhao, L. Micouin and R. Piccardi, *Helv. Chim. Acta*, 2019, 102, DOI: [10.1002/hlca.201900172](https://doi.org/10.1002/hlca.201900172).
- 58 F. Lima, M. Meisenbach, B. Schenkel and J. Sedelmeier, *Org. Biomol. Chem.*, 2021, 19, 2420–2424.
- 59 L. Brassart, M. Manneveau, T. Poisson, J. Legros, P. Huez, M. Forcato, P. Jubault and L. Chausset-Boissarie, *Adv. Synth. Catal.*, 2025, 367(10), e20241438.
- 60 P. Natho, M. Colella, M. Andresini, L. Degennaro and R. Luisi, *Org. Lett.*, 2024, 26, 3032–3036.
- 61 P. He, P. Watts, F. Marken and S. J. Haswell, *Angew. Chem., Int. Ed.*, 2006, 45, 4146–4149.
- 62 K. Tanaka, S. Motomatsu, K. Koyama, S. Tanaka and K. Fukase, *Org. Lett.*, 2007, 9, 299–302.
- 63 H. R. Sahoo, J. G. Kralj and K. F. Jensen, *Angew. Chem., Int. Ed.*, 2007, 46, 5704–5708.
- 64 C. H. Hornung, M. R. Mackley, I. R. Baxendale and S. V. Ley, *Org. Process Res. Dev.*, 2007, 11, 399–405.
- 65 M. S. Qenawy, K. Okamoto and A. Nagaki, *Chem. – Eur. J.*, 2025, 31, e202500299.
- 66 C. Wiles, P. Watts, S. J. Haswell and E. Pombo-Villar, *Chem. Commun.*, 2002, 1034–1035.

