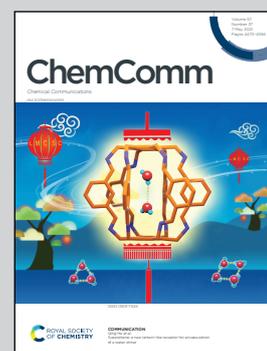


Showcasing research from the groups of Doctor Adrian Franczyk and Professor Jędrzej Walkowiak, Center for Advanced Technology, Adam Mickiewicz University in Poznań, Poland. Image used with the kind permission of Twinkl Educational Publishing. (c) Twinkl Ltd. All Rights Reserved.

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The first protocol for the synthesis of unsymmetrical bifunctional 1,1,3,3-tetramethyldisiloxane derivatives *via* subsequent hydrosilylation of alkenes and alkynes is presented. The methodology described has vast functional group tolerance and is extremely efficient towards the formation of novel disiloxane-based building blocks.

As featured in:



See Adrian Franczyk *et al.*,
Chem. Commun., 2021, **57**, 4504.


 Cite this: *Chem. Commun.*, 2021, 57, 4504

 Received 8th March 2021,
 Accepted 26th March 2021

DOI: 10.1039/d1cc01253c

rsc.li/chemcomm

Synthesis of bifunctional disiloxanes *via* subsequent hydrosilylation of alkenes and alkynes†‡

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The first protocol for the synthesis of unsymmetrical bifunctional 1,1,3,3-tetramethyldisiloxane derivatives *via* subsequent hydrosilylation of alkenes and alkynes is presented. The methodology described has vast functional group tolerance and is extremely efficient towards the formation of novel disiloxane-based building blocks.

The transition metal (TM) catalyzed addition of Si–H bonds to unsaturated bonds is the most powerful and commonly used laboratorial and industrial method for the preparation of silicon-based compounds.^{1–4} With practically unlimited potential for designing the organosilicon product structures and thus their chemical, physical, and biological properties, hydrosilylation is widely used in the synthesis of natural products,⁵ anti-corrosive coatings,⁶ hybrid materials,^{7,8} electrolytes,^{9,10} or polymer fire retardants.¹¹ Of particular interest regarding organosilicon compounds in terms of their modifiability, is 1,1,3,3-tetramethyldisiloxane (**1**). This inexpensive, low hazard, stable towards air and moisture, and commercially available hydrosiloxane derivative with low molecular weight, constitutes an attractive building block that has been applied in the reductive cleavage of the inert C–O bonds,¹² the reduction of tertiary carboxamides to tertiary amines,¹³ and the reduction of secondary and tertiary phosphine oxides.¹⁴ Moreover, functional disiloxanes have been widely applied as liquid crystals,¹⁵ solid polymeric electrolytes,¹⁶ or coupling agents.¹⁷

The functionalization of **1** can be achieved by hydrolysis/co-condensation of monochlorosilanes,¹⁸ condensation of silanols,¹⁹ or more preferably the hydrosilylation reaction, because of its simplicity, efficiency, and safety.^{20,21} This final method was recently

used for mono- or bisfunctionalization of **1** by γ -methacryloxypropyl,^{22–24} epoxy,^{22,25,26} aminopropyl,²⁷ or ethyleneoxide²⁸ groups or boryl, germyl,²⁹ or silyl moieties.^{30–33} The literature reports described above concern the hydrosilylation of unsaturated carbon–carbon double bonds (C=C) in vinyl or allyl groups. To the best of our knowledge, the hydrosilylation of carbon–carbon triple bonds (C≡C) by **1** or its monofunctionalized derivatives remains unexplored. The addition of the Si–H bond to the C≡C in terminal or internal alkynes can lead to a mixture of regio- and stereoisomers. Therefore, selective formation of unsaturated organosilicon compounds, which are extremely valuable building blocks in organic synthesis, is of importance.

Herein, we report the highly efficient TM-catalyzed bisfunctionalization of **1** *via* the subsequent hydrosilylation of alkenes and alkynes. The main goal of our study was to develop a simple, straightforward, and easy to scale-up protocol that leads to the unique disiloxanes containing a broad gamut of reactive groups capable of further transformations.

In the first stage of our study, selective monofunctionalization of **1** with a series of olefins was carried out. The hydrosilylation occurred in the presence of commercially available chloro(1,5-cyclooctadiene) rhodium(i) dimer ([RhCl(cod)]₂). As shown in our previous studies,^{20,21,29} the application of this binuclear rhodium catalyst (unlike the commonly used Karstedt's (Pt₂(dvs)₃) or Wilkinson's catalysts) allowed a reduction of **1** over olefins with the retention of excellent selectivity towards β -isomers, and negligible formation of bisadducts if 4-fold excess of **1** was used. Thus, we examined six different allylic derivatives (**2a–f**) and one vinyl derivative (**2g**) (Scheme 1). For **2a** and **2c–e**, the earlier protection of hydroxyl or amine group(s) with chlorotrimethylsilane (TMSCl) was needed to avoid competitive O- or N-silylation. All products with TMS-protected functionalities were obtained with almost quantitative yields and fully characterized (see the ESI†).

Hydrosilylation of alkenes (**2a–g**) was performed under the optimized conditions, in toluene (1 M) at 60 °C under an air atmosphere, in the presence of [RhCl(cod)]₂ (10^{−4} mol per Rh).

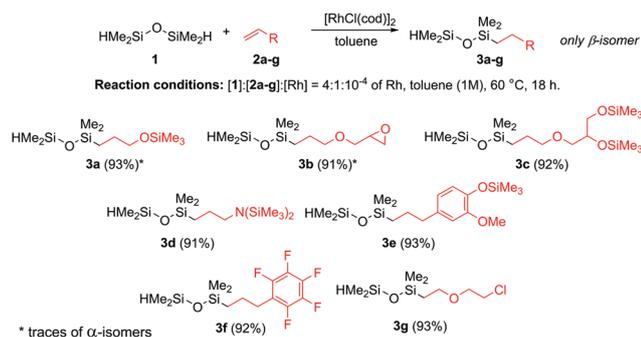
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‡ Electronic supplementary information (ESI) available. See DOI: 10.1039/d1cc01253c





Scheme 1 Hydrosilylation of allyl (**2a–f**) and vinyl (**2g**) derivatives with 1,1,3,3-tetramethyldisiloxane (**1**) in the presence of [RhCl(cod)]₂.

The reactions with (allyloxy)trimethylsilane (**2a**), allyl-glycidyl ether (**2b**), and *N*-allyl-*N,N*-bis(trimethylsilyl)amine (**2d**) resulted in the desired products **3a**, **3b**, and **3d** with very high yields. However, the formation of bisadducts or α -isomer in a very low quantity (<6%) was also determined by GC-MS analysis. Application of TMS-protected 3-(allyloxy)propane-1,2-diol (**2c**), eugenol-TMS (**2e**), allylpentafluorobenzene (**2f**) and 2-chloro(ethyl)vinyl ether (**2g**) smoothly led to β -isomers without formation of the by-products. It is worth emphasizing that the monofunctionalized compounds **3c** and **3f** are herein reported and fully characterized for the first time. In addition, **3a**, despite previous reports,³⁴ has been completely spectrally characterized for the first time. The presented above process has good tolerance towards various functional groups such as silyloxy, glycidyl, amine, aryl, and halogen, which can be easily modified in further transformations. Additionally, for the hydrosilylation of **3a** with **4b**, the gram-scale synthesis was performed leading to **5ab** with high efficiency.

In the next step with the series of diverse disiloxanes **3a–g** in hand, we examined the scope and limitations for the synthesis of their bifunctional, unsymmetrical derivatives *via* hydrosilylation of internal and terminal alkynes. The selective hydrosilylation of C \equiv C bonds allows the introduction of an alkenyl moiety into the compound structure, which makes them an attractive building block in *e.g.*, addition or coupling reactions. On the other hand, the selective hydrosilylation of the C \equiv C bonds, especially terminal ones, is a challenging task due to the possible formation of stereo- and regioisomers.

Based on our experience in hydrometallation of the unsaturated C \equiv C bonds, for the Si–H addition to internal alkynes, the commercially available and commonly used Karstedt's catalyst was selected.³⁵ For terminal alkynes, *in situ* generated catalytic system based on PtO₂/Xphos was the most efficient.^{35,36} Due to the vast diversity of monofunctionalized **3a–g** derivatives and alkynes **4a–k**, hydrosilylation was performed at 100 °C for 18 hours with the equimolar ratio of the substrates, which were used as received from the suppliers, without any initial purification step.

To determine the influence of alkyne structure on the process efficiency, we chose derivative **3a** and used it in the hydrosilylation with internal (**4a–d**) and terminal alkynes (**4e–k**) (Table 1). The application of symmetrical and aromatic alkynes

Table 1 Hydrosilylation of internal and terminal alkynes (**4–k**) with (**3a**) in the presence of Pt catalysts

| Entry | Alkyne 4 | | Conv. of 3a ^a [%] | Selectivity 5 = 6/7 or 5/6/7 ^a [%] | Isolated yield of 5 [%] |
|-------|----------|---------------------------------------|-------------------------------------|---|--------------------------------|
| | R' | R'' | | | |
| 1 | a | Ph | 99 | 100/0 | aa 81 |
| 2 | b | 4-BrPh | 99 | 100/0 | ab 77 |
| 3 | c | C ₃ H ₇ | 100 | 100/0 | ac 96 ^b |
| 4 | d | 4-BpinPh | 99 | 100/0 | ad 73 |
| 5 | e | c-C ₆ H ₁₁ | 96 | 98/2/0 | ae 82 |
| 6 | f | CM ₂ (OSiMe ₃) | 100 | 85/15/0 | af — |
| 7 | g | Ph | 100 | 100/0/0 | af 94 ^c |
| 8 | h | 4-BrPh | 91 | 98/2/0 | ag 93 |
| 9 | i | 4-MeOPh | 99 | 96/4/0 | ah 69 |
| 10 | j | 4-MeOPh | 99 | 99/1/0 | ai 95 ^c |
| 11 | k | 9-Phenanthryl | 94 | 90/10/0 | aj 68 |
| 12 | k | 3-Thienyl | 91 | 99/1/0 | ak 84 |

Reaction conditions: for entries 1–4, 6: [**3a**]:[**4a–d**, **f**]:[Pt₂(dvs)₃] = 1:1:10⁻⁴ of Pt, 100 °C, 18 h, toluene (0.05–0.1 M); for entries 5, 7–12: [**3a**]:[**4e–k**]:[XPhos]:[PtO₂] = 1:1:2 × 10⁻²:10⁻² of PtO₂, 100 °C, 18 h THF (0.07 M). ^a Determined by GC-MS or NMR analyses. ^b Analysed without any purification step. ^c Filtration through a syringe filter (0.2 μ m) without further purification.

(**4a–b**) smoothly led to (*E*)-isomers (**5aa–ab**) *via cis*-addition of Si–H to the C \equiv C bond, with 99% conversion of reagents (Table 1, entries 1 and 2). In the same manner, aliphatic 4-octyne (**4c**) was transformed into **5ac** (Table 1, entry 3). The hydrosilylation of unsymmetrical internal 4-[(trimethylsilyl)ethynyl]-phenylboronic acid pinacol ester (**4d**) with **3a** resulted in the selective formation of the (*E*)-isomer. 1D NOESY analysis revealed that the proton from SiH group is attached to the alkenyl carbon with SiMe₃ group (Table 1, entry 4; ESI, ‡ Fig. S44).

Next, terminal alkyne screening was carried out. It was initially based on the reaction conditions determined for internal alkynes. Application of Pt₂(dvs)₃ resulted in a mixture of regioisomers **5** and **6**, even for sterically hindered [(1,1-dimethyl-2-propynyl)oxy]trimethylsilane (**4f**) (**5af/6af** = 85/15). Thus, relying on our previous studies and literature reports, the generated *in situ* PtO₂/Xphos system was applied. It is known as the selective catalyst for the addition of the Si–H bond to the C \equiv CH bond. Indeed, when it was used in the hydrosilylation of **4f** with **3a**, the exclusive formation of **5af** occurred (Table 1, entry 7). For the linear 1-heptyne (**4e**), a very high selectivity was noticed as well. However, traces of **6ae** were formed. The same observations were carried out for the aromatic terminal alkynes such as phenylacetylene (**4g**) (Table 1, entry 8). The incorporation of a weakly electron-withdrawing atom, *i.e.* Br, to the phenyl ring had a negative influence on the **3a** conversion and reaction selectivity (**5ah/6ah** = 96/4) (Table 1, entry 9).

In turn, a donating group (–OMe) ensures a very high conversion of **3a** and negligible formation of isomeric by-products (Table 1, entry 10). Polycyclic aromatic and heterocyclic alkynes such as 9-ethynylphenanthrene (**4j**) and 3-ethynylthiophene (**4k**) can also be effectively used for the hydrosilylation of **3a**, although



under the applied reaction conditions, a complete conversion of **3a** was not observed (Table 1, entries 11 and 12).

Encouraged by the results obtained for the hydrosilylation of **3a** with internal and terminal alkynes, we decided to expand the library of bifunctional unsymmetrical substituted derivatives. We applied **2b–g** in the hydrosilylation of selected alkynes. The substrate scope is summarized in Scheme 2. In general, mono-substituted 1,1,3,3-tetramethyldisiloxane bearing glycidyl ether (**3b**) selectively reacted with aromatic (**4a–b**) and alkyl (**4c**) internal alkynes, smoothly leading to (*E*)-isomers (**5ba–bc**), whereas with terminal **4f**, β -(*E*)-isomer (**5bf**) was formed. The same observation was made for the internal (**4a–c**) and terminal (**4f**) alkynes when **3c** containing two OSiMe₃ groups and **3d** bearing N(SiMe₃)₂ groups were used. Our protocol was also suitable for the derivatives with eugenol (**3e**) or pentafluorobenzene (**3f**) moieties. For both, excellent selectivity and high isolation yields ($\geq 71\%$) were observed (**5ea–ef**, **5fb**, **5fc**). The hydrosilylation of 1-(2-(2-chloroethoxy)ethyl)-1,1,3,3-tetramethyldisiloxane (**3g**) with internal and terminal alkynes, gave also the desired products (**5gb**, **5gf**) with a very high reaction efficiency.

By the *in situ* FT-IR method, the influence of the structure of **3a–g** on the rate of the hydrosilylation of diphenylacetylene (**4a**) was determined, by the following of the peak area at 907 cm⁻¹ which corresponds to the Si–H bond. It was found that the addition of the Si–H bond to the C \equiv C proceeded rapidly within 1 hour for most siloxanes. Afterward, the reaction rate decreased due to the lower concentration of the substrates. The fastest consumption of siloxane occurred for **3a** (2 h), while the

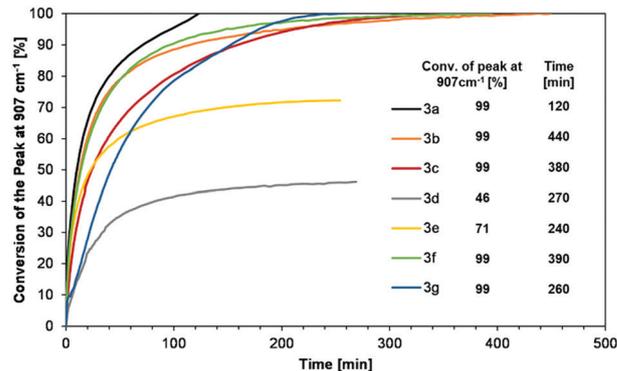
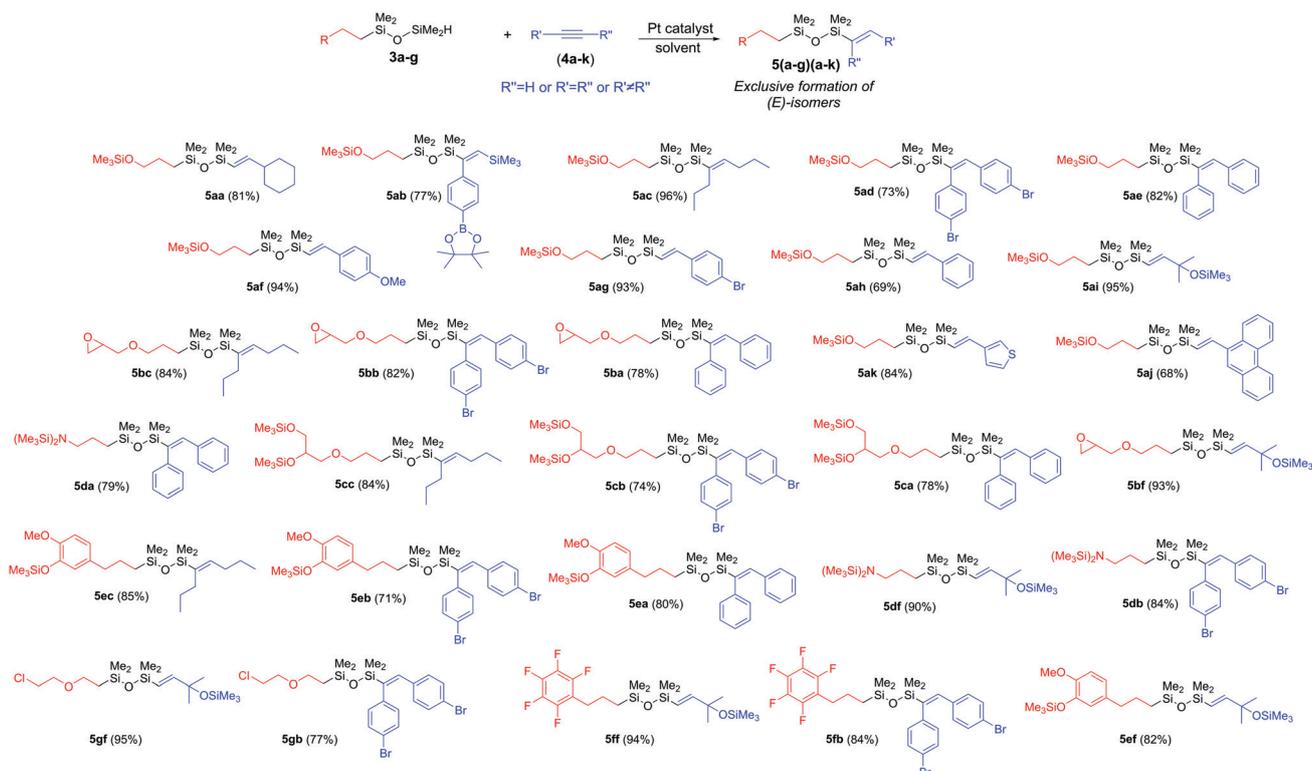


Fig. 1 The hydrosilylation of **4a** with **3a–g** monitored by *in situ* FT-IR.

slowest for **3b** (over 7 h). For the rest of the reagents, the reaction time was between 3 and 6 hours (Fig. 1). For **3d** and **3e**, the peak conversion was not complete due to partial overlapping of the band assigned to Si–N and ether bonds with the Si–H band. However, GC analysis performed after flattening of the conversion curves confirmed the 100% consumption of the siloxanes.

It should be emphasized, that the presented protocol has excellent functional group tolerance for monosubstituted 1,1,3,3-tetramethyldisiloxanes (siloxy, glycidyl, amine, aryl, halogen groups) and alkynes (*n*-alkyl, *c*-alkyl, aryl, halogen, alkoxy, heterocyclic groups). Moreover, our approach is extremely selective towards the formation of (*E*)-isomers. The presence of the



Scheme 2 The scope of the synthesized bifunctional unsymmetrical substituted disiloxane derivatives.



unsaturated C=C bond on the one side and the reactive moiety on the other makes them very appealing building blocks with potential usage as coupling agents or in materials science. It is worth noting that all bifunctional 1,1,3,3-tetramethyldisiloxanes are new compounds and were comprehensively characterized by NMR, FT-IR, and MS methods. Moreover, a gram-scale reaction of **2a-c** with **1** was carried out. Very high yields and selectivities of the monofunctionalized derivatives: **3a** (93%), **3b** (92%), and **3c** (89%) were obtained.

In summary, selective and highly efficient methods for the synthesis of 29 unsymmetrical bifunctional derivatives *via* simple hydrosilylation of structurally different alkynes and alkenes with 1,1,3,3-tetramethyldisiloxane have been successfully developed. The process was highly selective when the hydrosilylation of alkenes was carried out in the presence of [RhCl(cod)]₂ and for internal or terminal alkynes Pt₂(dvs)₃ or PtO₂/XPhos were respectively used. The protocol is effective towards a broad spectrum of alkenes as well as internal and terminal alkynes with various functionalities, leading to products with high isolation yields. At the same time it is easy to scale up, and therefore the synthesis can be performed on the gram scale.

This work was supported by the National Centre for Research and Development in Poland, Lider Programme No. LIDER/6/0017/L-9/17/NCBR/2018 and the National Science Centre in Poland No. UMO-2019/32/C/ST4/00235, UMO-2018/31/G/ST4/04012.

Conflicts of interest

There are no conflicts to declare.

Notes and references

- B. Marciniec, *Hydrosilylation: a comprehensive review on recent advances*, Springer Science & Business Media, 2009.
- B. Marciniec, H. Maciejewski, C. Pietraszuk and P. Pawluć, *Applied Homogeneous Catalysis with Organometallic Compounds: A Comprehensive Handbook in Four Volumes*, 2017, pp. 569–620.
- A. K. Roy, in *Advances in Organometallic Chemistry*, ed. R. West, A. F. Hill and M. J. Fink, Academic Press, 2007, vol. 55, pp. 1–59.
- Y. Naganawa, K. Inomata, K. Sato and Y. Nakajima, *Tetrahedron Lett.*, 2020, **61**, 151513.
- E. Langkopf and D. Schinzer, *Chem. Rev.*, 1995, **95**, 1375–1408.
- J. Wojciechowski, K. Szubert, R. Peipmann, M. Fritz, U. Schmidt, A. Bund and G. Lota, *Electrochim. Acta*, 2016, **220**, 1–10.
- U. Díaz, T. García, A. Vely and A. Corma, *Chem. – Eur. J.*, 2012, **18**, 8659–8672.
- Y. Du and H. Liu, *Dalton Trans.*, 2020, **49**, 5396–5405.
- S. L. Guillot, A. Peña-Hueso, M. L. Usrey and R. J. Hamers, *J. Electrochem. Soc.*, 2017, **164**, A1907–A1917.
- T. Yong, J. Wang, Y. Mai, X. Zhao, H. Luo and L. Zhang, *J. Power Sources*, 2014, **254**, 29–32.
- Z. Han, A. Fina and G. Camino, *Polymer Green Flame Retardants*, Elsevier, 2014, pp. 389–418.
- P. Alvarez-Bercedo and R. Martin, *J. Am. Chem. Soc.*, 2010, **132**, 17352–17353.
- Y. Sunada, H. Kawakami, T. Imaoka, Y. Motoyama and H. Nagashima, *Angew. Chem., Int. Ed.*, 2009, **48**, 9511–9514.
- M. Berthod, A. Favre-Réguillon, J. Mohamad, G. Mignani, G. Docherty and M. Lemaire, *Synlett*, 2007, 1545–1548.
- G. Shanker, M. Prehm and C. Tschierske, *J. Mater. Chem.*, 2012, **22**, 168–174.
- I. J. Lee, G. S. Song, W. S. Lee and D. H. Suh, *J. Power Sources*, 2003, **114**, 320–329.
- F. J. LaRonde, A. M. Ragheb and M. A. Brook, *Colloid Polym. Sci.*, 2003, **281**, 391–400.
- Y. Isoda and K. Ayama, EP0657486B1, 1998.
- H. Friedrich, I. Jansen and K. Rühlmann, *Polym. Degrad. Stab.*, 1994, **46**, 9–18.
- R. Januszewski, I. Kownacki, H. Maciejewski and B. Marciniec, *J. Organomet. Chem.*, 2017, **846**, 263–268.
- R. Januszewski, I. Kownacki, H. Maciejewski, B. Marciniec and A. Szymańska, *Eur. J. Inorg. Chem.*, 2017, 851–856.
- R. Acosta Ortiz, M. Sangermano, R. Bongiovanni, A. E. Garcia Valdez, L. B. Duarte, I. P. Saucedo and A. Priola, *Prog. Org. Coat.*, 2006, **57**, 159–164.
- G. Erdodi and J. P. Kennedy, *J. Polym. Sci., Part A-1: Polym. Chem.*, 2007, **45**, 295–307.
- G. Guzman, T. Nugay, I. Nugay, N. Nugay, J. Kennedy and M. Cakmak, *Macromolecules*, 2015, **48**, 6251–6262.
- R. Malik and J. V. Crivello, *J. Macromol. Sci., Pure Appl. Chem.*, 1997, **34**, 247–263.
- D. Zhang, E. Liang, T. Li, S. Chen, J. Zhang, X. Cheng, J. Zhou and A. Zhang, *RSC Adv.*, 2013, **3**, 3095–3102.
- Y.-Z. Niu, L. Zhang, S.-J. Liang, D.-X. Wang and S.-Y. Feng, *Chin. Chem. Lett.*, 2014, **25**, 1419–1422.
- S. S. Sologubov, A. V. Markin, N. N. Smirnova, N. A. Novozhilova, E. A. Tatarinova and A. M. Muzafarov, *J. Phys. Chem. B*, 2015, **119**, 14527–14535.
- R. Januszewski, M. Grzelak, B. Orwat, M. Dutkiewicz and I. Kownacki, *J. Catal.*, 2020, **390**, 103–108.
- D. A. de Vekki, V. Ol'Sheev, V. Spevak and N. Skvortsov, *Russ. J. Gen. Chem.*, 2001, **71**, 1912–1923.
- D. A. de Vekki and N. Skvortsov, *Russ. J. Gen. Chem.*, 2004, **74**, 197–206.
- D. A. de Vekki, V. Uvarov, A. Reznikov and N. Skvortsov, *Russ. Chem. Bull.*, 2008, **57**, 349–357.
- R. Murthy, C. D. Cox, M. S. Hahn and M. A. Grunlan, *Biomacromolecules*, 2007, **8**, 3244–3252.
- G. D. Khatuntsev, V. D. Sheludyakov and V. F. Mironov, *Zh. Obshch. Khim.*, 1974, **43**, 2150–2155.
- K. Stefanowska, A. Franczyk, J. Szyling, K. Salamon, B. Marciniec and J. Walkowiak, *J. Catal.*, 2017, **356**, 206–213.
- A. Hamze, O. Provot, J.-D. Brion and M. Alami, *J. Organomet. Chem.*, 2008, **693**, 2789–2797.

