

PAPER

[View Article Online](#)
[View Journal](#) | [View Issue](#)Cite this: *Catal. Sci. Technol.*, 2025, 15, 4012

Influence of the supported ionic-liquid layer thickness on Z-selectivity in 1-alkyne hydrosilylation under continuous flow†

André Böth,^a Florian Kaltwasser,^a Christian Friedigkeit,^a Boshra Atwi,^b Wolfgang Frey,^c Michael R. Buchmeiser ^{*,b} and Ulrich Tallarek ^{*,a}

1-Butyl-3-methylimidazolium tetrafluoroborate containing different rhodium(I) N-heterocyclic carbene (NHC) complexes was immobilized as a supported ionic-liquid phase (SILP) inside the mesopores of a silica monolith to study the impact of SILP thickness (d_{SILP}) from the thin-SILP-limit ($d_{\text{SILP}} \approx 1$ nm) to complete mesopore filling ($d_{\text{SILP}} \approx 15$ nm) on *Z/E*-selectivity in the rhodium-catalyzed hydrosilylation of phenylacetylene with dimethylphenylsilane. A coupled analytical platform allowed monitoring of both yield and selectivity of the produced isomer pattern online in continuous-flow experiments of 600 minutes using methyl *tert*-butyl ether as mobile phase. The approach provided new insights into the mechanistic aspects of the reaction under liquid confinement conditions created by the varied SILP thickness. With decreasing d_{SILP} , the selectivity of a Rh-catalyst based on a chelating NHC is shifted towards the β (*Z*)-isomer, climaxing in a boost of the *Z/E*-ratio for $d_{\text{SILP}} = 1$ nm by a factor of >30 , while the selectivity is mostly unaffected for catalysts based on nonchelating NHCs. The spatial dimension of 1 nm reflects the rigid part of the SILP characterized by a quasi-frozen morphology of the ionic liquid. It shapes a local, spatially as well as molecularly confined catalytic environment, which, together with a tailored catalyst, facilitates the predominant formation of the β (*Z*)-isomer under kinetic control. Contrariwise, the random, mobile part of the adjoining bulk SILP, emerging with increasing d_{SILP} , generally favors the formation of the β (*E*)-isomer under thermodynamic control.

Received 9th April 2025,
Accepted 27th May 2025

DOI: 10.1039/d5cy00436e

rsc.li/catalysis

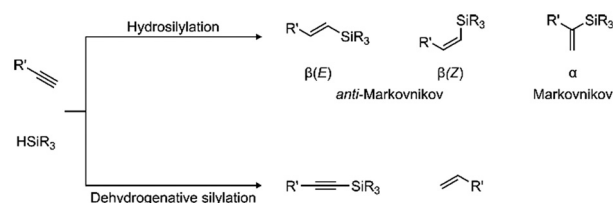
1. Introduction

The hydrosilylation of terminal alkynes is an important synthetic method for the preparation of vinylsilanes, which have a number of applications in organic synthesis^{1–5} and materials science,⁶ including the production of polymers⁷ and fine chemicals.⁸ By the addition of Si–H bonds to a terminal alkyne, the hydrosilylation reaction yields different vinylsilane isomers (Scheme 1).

Accordingly, the reaction can proceed in an *anti*-Markovnikov mode, leading to the β (*E*)- and β (*Z*)-vinylsilane isomers, or in a Markovnikov mode to give the α -vinylsilane. For some catalysts, dehydrogenative silylation products (the silyl-alkyne derivative and the corresponding

alkene) are received as by-products. Tailoring the synthesis towards an isomer is a challenging task, because the selectivity depends on a number of factors including the nature of the catalyst, the substrates, and the reaction conditions. Unsurprisingly, many catalysts have been designed with distinct electronic and steric properties to gain control over the regio- and stereoselectivity.⁹

Transition-metal N-heterocyclic carbene (NHC) complexes have received particular attention in recent years for alkyne hydrosilylation due to their versatility and wide range of applications.^{10–16} NHC–Ir(I), Rh(I), and Ru(I) catalysts demonstrate high *anti*-Markovnikov selectivity in the hydrosilylation of terminal alkynes, *i.e.*, they mainly produce



Scheme 1 Hydrosilylation and dehydrogenative silylation of terminal alkynes.

^a Department of Chemistry, Philipps-Universität Marburg, Hans-Meerwein-Strasse 4, D-35032 Marburg, Germany. E-mail: tallarek@staff.uni-marburg.de

^b Institute of Polymer Chemistry, Universität Stuttgart, Pfaffenwaldring 55, D-70569 Stuttgart, Germany. E-mail: michael.buchmeiser@ipoc.uni-stuttgart.de

^c Institute of Organic Chemistry, Universität Stuttgart, Pfaffenwaldring 55, D-70569 Stuttgart, Germany

† Electronic supplementary information (ESI) available: CCDC 2430795 and 2441044. For ESI and crystallographic data in CIF or other electronic format see DOI: <https://doi.org/10.1039/d5cy00436e>

mixtures of $\beta(E)$ - and $\beta(Z)$ -isomers.^{10,17,18} Nevertheless, the isomerization of the thermodynamically labile $\beta(Z)$ -isomer into the stable $\beta(E)$ -vinylsilane remains an issue.¹⁹ Further, cationic rhodium complexes were found to be highly $\beta(E)$ -selective,^{20,21} whereas, *e.g.*, zwitterionic Rh(III) bis(NHC) complexes demonstrate excellent selectivity for the thermodynamically less stable $\beta(Z)$ -isomer.²² Another strategy to control selectivity relies on the immobilization of the catalyst inside a confined space. Examples include Rh(I)/Rh(II) catalysts supported in mesoporous silica and rhodium nanoparticles in carbon reactors.^{23,24} Both studies report a change in selectivity with respect to the homogeneous conditions. It has been argued that the driving force originates in confinement effects, including preferred substrate orientation, increased local concentrations near active sites, the stabilization of intermediates, and enhanced steric congestion. Finally, Messerle and co-workers showed that Rh and Ir complexes covalently anchored to a carbon black-supported surface enable $\beta(Z)$ -selectivity, depending on the metal center, ligand environment, and tether length.^{25,26} Their results suggest that subtle changes in surface coverage and linker architecture can alter the product distribution, offering a modular, heterobimetallic approach to selectivity control. In a complementary approach, Sánchez-Page *et al.*²⁷ achieved high $\beta(Z)$ -selectivity using cyclometalated Rh(III)-NHC complexes, both in solution and supported on graphene, through a metal–ligand cooperative mechanism.

An intriguing approach that has become popular in recent years involves a supported ionic-liquid phase (SILP),^{28–31} where the homogeneous catalyst is dissolved in an ionic liquid (IL) that is physically adsorbed as a film on a high-surface area solid support. By providing the substrates in the gas phase, the reaction mixture can be easily separated from the SILP without catalyst leaching, so that the approach lends itself for the adaptation in continuous-flow experiments.^{32–38} We extended this technology to supported ionic liquid–liquid phase conditions^{32,39–41} by using a substrate- and, in course of the reaction, also product-containing liquid phase that does not mix with the IL. In this way, the SILP concept was successfully extended to more complex, high-boiling substrates.

In comparison to batch operation, the utilization of compact continuous-flow microreactors enables the effective use of large surface-to-volume ratios, facilitating heat and mass transfer. This, in turn, impacts yield and selectivity as well as the practicable reaction time.^{42,43} However, literature illustrating how the SILP and the SILP thickness (d_{SILP}), in particular, affect the selectivity of the dissolved catalyst in view of the spatial confinement engendered by the SILP is still very scarce.^{32,44} The aspect of the actual SILP thickness and catalyst condition is complemented by increased efforts in the rational design of task-specific ILs, which self-assemble on a surface (without covalent bonding) and offer a metal-binding site at a controlled distance from the solid support.⁴⁵

The main objective of the current work was to understand the predominant formation of the $\beta(Z)$ -isomer in the hydrosilylation of 1-alkynes catalyzed by SILP-embedded Rh-

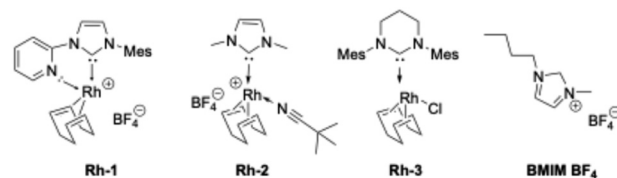
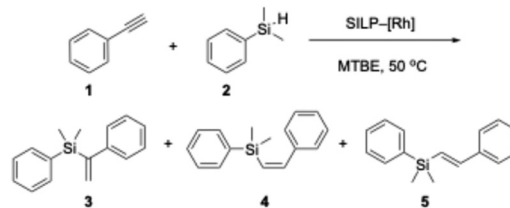


Fig. 1 The Rh-NHC complexes [(1-(pyrid-2-yl)-3-mesityl)-imidazol-2-ylidene](η^4 -1,5-cyclooctadiene) rhodium(I) tetrafluoroborate (**Rh-1**), [(1,3-dimethylimidazol-2-ylidene)(η^4 -1,5-cyclooctadiene) rhodium(I) tetrafluoroborate] (**Rh-2**), and [(1,3-dimesityltetrahydropyrimidin-2-ylidene)(η^4 -1,5-cyclooctadiene) rhodium(I) chloride] (**Rh-3**), as well as the room-temperature IL 1-butyl-3-methylimidazolium tetrafluoroborate, [BMIM][BF₄], used in this work. Mes = mesityl (2,4,6-trimethylphen-1-yl).

NHC catalysts, which otherwise predominantly produce the $\beta(E)$ -isomer under homogeneous, that is, biphasic bulk IL-organic solvent conditions.^{32,44} For that purpose, we immobilized the IL 1-butyl-3-methylimidazolium tetrafluoroborate [BMIM][BF₄] on the surface of a macro-mesoporous silica monolith, suitable for continuous-flow operation. The employed catalysts (**Rh-1**, **Rh-2**, **Rh-3**) and the IL are shown in Fig. 1.

To better understand the confinement effect engendered by the SILP, we investigated the influence of its thickness on the *Z/E*-selectivity in the hydrosilylation reaction between phenylacetylene (PA) and dimethylphenylsilane (DMPS) under biphasic, liquid–liquid, continuous-flow conditions on the monolithic column using methyl *tert*-butyl ether (MTBE) as liquid mobile phase that does not mix with the IL (Scheme 2). Continuous-flow operation was established with high-precision, high-pressure pumps connected to the SILP/catalyst-loaded column, and the real-time monitoring of the reaction was achieved by hyphenating the reactor (first) dimension online with a concentration analytics (second) dimension for chromatographic separation as well as detection and quantification of all compounds of interest.

The two-dimensional setup facilitated the automation of the entire sampling process and enabled us to repeatedly run experiments with varied d_{SILP} over extended periods, typically for 600 minutes, during which the production of the targeted isomers, especially the *Z/E*-selectivity of the reaction, was continuously monitored. This would otherwise become very laborious as well as time-consuming with batch operation.



Scheme 2 Biphasic, continuous-flow hydrosilylation between PA (**1**) and DMPS (**2**) with **Rh-1** in the SILP formed by [BMIM][BF₄] on a macro-mesoporous silica monolith. Stationary phase: solid silica and SILP (containing **Rh-1**), mobile phase: MTBE (containing reactants and products).



Data acquired with the monolithic microreactor for SILP thicknesses between 1 and 15 nm reveal that the *Z/E*-selectivity of the reaction is generally low at the beginning and for $d_{\text{SILP}} \geq 3$ nm also increases only moderately during the 600 minutes of the experiments. However, for $d_{\text{SILP}} = 1$ nm the *Z/E*-selectivity demonstrates a real boost. These observations highlight that more attention should be paid to the morphology as well as the dynamics of the SILP in view of catalytic applications, in particular, to the rigid, low-mobility part of the SILP next to the surface, comprising 1–2 more ordered molecular layers of the IL, and the bulk (random) and correspondingly mobile region of the SILP emerging beyond the rigid part. Our data indicate that the catalytic environment in the ~ 1 nm-thick rigid part favors the formation of the $\beta(Z)$ -isomer. In contrast, the bulk SILP favors the formation of the $\beta(E)$ -isomer. It is obvious that the *Z/E*-selectivity can then be tuned in a simple way through d_{SILP} via the catalytic environments prevailing in rigid and mobile parts of the SILP.

This effect will be useful and tailored further in certain applications, as these characteristic regions in SILPs are a fundamental property of SILP interfacial morphology.

2. Results and discussion

To perform the continuous-flow hydrosilylation experiments outlined in Scheme 2, we adapted a silica monolith as support for the catalyst-containing IL. Silica-based monoliths with a hierarchical macro-mesopore space morphology, as used in this work, are attractive supports in liquid chromatography⁴⁶ and heterogeneous catalysis.^{47–49} Their bimodal pore size distribution is realized with a continuous block of silica perforated by intersecting networks of macropores and mesopores.^{50,51} While the macropores allow for fast (advection-dominated) transport through the material, the mesopores (accessible only by diffusion) provide sufficient surface area for adsorption and reaction. Fig. 2 highlights the morphology of the monolith⁵² implemented in the present work as support for the IL [BMIM][BF₄] containing the corresponding Rh-catalyst. The monolith was used in the form of an analytical-scale 4.6 mm inner diameter $\times$ 100 mm length column clad in polyether ether ketone (Chromolith® Si 100–4.6 mm).^{53,54} Relevant morphological properties of the monolith (provided

by the manufacturer) are a mean macropore size and a specific macropore volume of 1.84 μm and 2.58 mL g^{-1} , respectively, and a mean mesopore size, specific mesopore volume, and specific mesopore surface area of 32.9 nm, 0.91 mL g^{-1} , and 110 $\text{m}^2 \text{g}^{-1}$, respectively, as derived from mercury intrusion porosimetry (macropore space) and nitrogen physisorption analysis (mesopore space). For the preparation of the SILP, the monolith was flushed with a solution of [BMIM][BF₄] in CH_2Cl_2 containing the corresponding Rh-catalyst. Importantly, the IL does not interfere with the catalyst through anion metathesis and both, IL and catalyst, do not dissolve in MTBE. This enabled the continuous-flow experiments with MTBE as mobile phase in a biphasic SILP approach and prevented both IL and Rh-catalyst dissolved therein from leaching. Variation of the SILP thickness was achieved by adjusting the IL volume deposited inside the mesopores of the monolith (see Experimental section and ESI† for further details).

Four SILP thicknesses were realized in this work ($d_{\text{SILP}} = 1, 3, 5$, and 15 nm), ranging from the thin-SILP-limit ($d_{\text{SILP}} = 1$ nm), below which a uniform monolayer of [BMIM][BF₄] is difficult to establish on the silica surface considering the molecular structure of the supported IL, to the thick-SILP-limit ($d_{\text{SILP}} = 15$ nm), where the ~ 30 nm wide mesopores are completely filled with [BMIM][BF₄]. SILP thicknesses with applied IL volumes and amounts of **Rh-1** catalyst in the SILPs are summarized in Table S2 (ESI†). The hydrosilylation of PA with DMPS in the presence of **Rh-1** under continuous flow was accomplished by adapting the two-dimensional microreaction-separation setup shown in Fig. 3.⁵⁵ While the first dimension serves for the management of the reaction, the second one involves the chromatographic separation of the isomers as well as their subsequent quantification. In the first dimension, a binary pump (4, Fig. 3) was employed to deliver the substrate solution in MTBE to the microreactor fixed in a temperature-controlled oven compartment (8, Fig. 3) at a precisely adjustable and constant volumetric flow rate Q . The residence time of the substrate solution on the microreactor (and thus the reaction time) was therefore controlled by the pump flow rate and reflects the volume in the monolith available to the pumped solution.

To arrive at quantitative information about the reaction solution, the second dimension was coupled online to the first one to allow for the separation of the targeted isomers

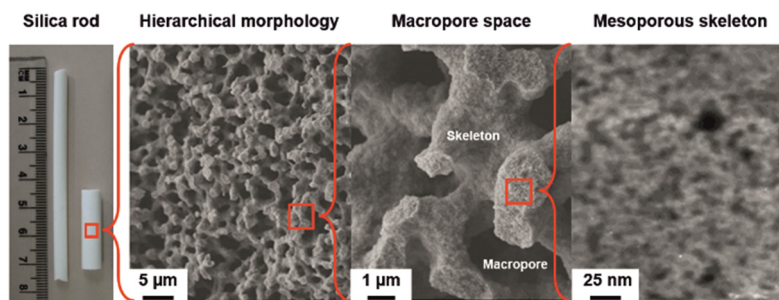


Fig. 2 Hierarchically structured silica monolith. The macropores form an interskeleton network of flow-through channels dedicated to advection-dominated transport, while transport in the intraskeleton mesopores remains diffusion-limited. Adapted with permission from Stoeckel *et al.*⁵²



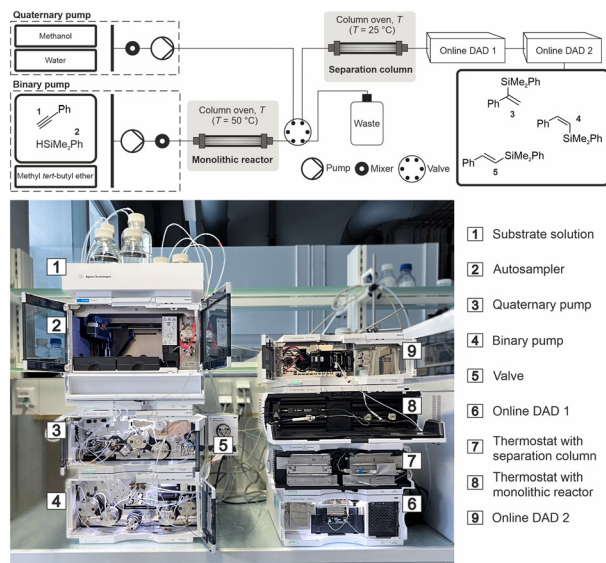


Fig. 3 Photograph of the two-dimensional continuous-flow platform used to study the hydrosilylation reaction between PA and DMPS with Rh-1 in the SILP formed by [BMIM][BF₄] on a macro-mesoporous silica monolith (cf. Fig. 2). An injection valve (5) behind the microreactor compartment (8) connects the first (reaction) dimension with the second (analytical) dimension, which enables chromatographic separation of the injected reaction solution on a phenyl-hexyl-modified UHPLC column (7) and online analysis of the separated compounds with a diode-array detector (9).

by ultrahigh-performance liquid chromatography (UHPLC)⁵⁶ as well as their detection with a diode-array detector. Both dimensions were connected by a 2-position/6-port valve (5, Fig. 3). Continuous-flow conditions were imposed by flushing the microreactor with the substrate solution at $Q = 0.066 \text{ mL min}^{-1}$, which translates into a reaction time of $t_{\text{rct}} = 20 \text{ min}$. This time was adjusted to enable a meaningful comparison between the different reactor loadings with SILP and catalyst (cf. Table S2†) rather than to maximize conversion. As a consequence, unreacted starting materials were generally present in these runs (as can be recognized in Fig. 4). Complementary to the reaction time t_{rct} , the total experiment time (t_{total}) is the overall time for which the microreactor was monitored under a given set of reaction (continuous-flow) conditions. For example, as both yield and selectivity (eqn (1) and (2) below) were repeatedly determined over a total period of 600 minutes, then $t_{\text{total}} = 600 \text{ min}$ for this experiment with the selected SILP thickness, catalyst and substrate concentrations, reaction time, and reaction temperature.

The peak areas of the baseline-separated compounds of interest, i.e., $\beta(\text{Z})$ - and $\beta(\text{E})$ -isomers, were converted into masses using calibration curves. Relative chromatographic yields of the individual isomers and Z/E -selectivities were quantified with these data according to

$$\text{Relative yield} = \frac{m_{\beta(i)}}{m_{\beta(\text{Z})} + m_{\beta(\text{E})}} \quad (1)$$

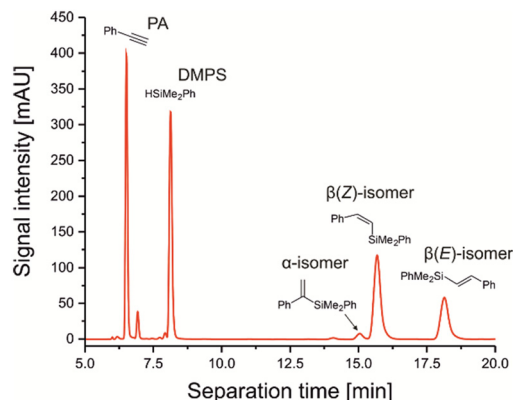


Fig. 4 Chromatogram of the reaction solution, recorded by UV/vis diode-array detection at 260 nm, highlighting the baseline separation of compounds 1–5. Column: 3.0 mm inner diameter $\times$ 100 mm length packed with phenyl-hexyl-modified, fully porous silica particles (1.8 μm particle size, 9.5 nm mesopore size), mobile phase: 75/25 (v/v) methanol/water, flow rate: 0.6 mL min^{-1} , column backpressure: 800 bar.

$$\text{Selectivity} = \frac{m_{\beta(\text{Z})}}{m_{\beta(\text{E})}} \quad (2)$$

where $m_{\beta(\text{Z})}$ and $m_{\beta(\text{E})}$ denote the respective masses and i stands for the $\beta(\text{Z})$ - or $\beta(\text{E})$ -isomer. Required calibration curves and details of their acquisition are provided in the ESI.†

One prerequisite for these continuous-flow studies is a fast baseline separation of the compounds of interest, because a short separation cycle allows to monitor yield and selectivity at high temporal resolution over an experiment (t_{total}) and address, for example, the influence of d_{SILP} in a series of experiments. For apolar molecules like compounds 1–5 (Scheme 2), a phenyl-hexyl silica column was found appropriate, as that surface chemistry reflects the presence of the aromatic phenyl groups in all these hydrosilylation compounds, offering selective interactions.⁵⁷ Fig. 4 highlights their baseline separation in less than 20 minutes using 75/25 (v/v) methanol/water as UHPLC eluent at a flow rate of 0.6 mL min^{-1} . This in turn enabled the quantitative analysis of all compounds *via* total numerical integration of their chromatographic peaks together with the acquired calibration curves. As can be seen in Fig. 4, the bottleneck in this analysis was particularly the baseline separation of the α -isomer from the $\beta(\text{Z})$ -isomer to allow for an unbiased analysis of the latter.

Accordingly, this optimized UHPLC method was implemented as default protocol in the second (separation) dimension of the setup shown in Fig. 3. With this implementation, we were prepared for a deeper investigation of mechanistic aspects of the hydrosilylation reaction occurring in the SILP, with the main focus on the determination of the yields of the $\beta(\text{Z})$ - and $\beta(\text{E})$ -isomers and, thus, of Z/E -selectivity for each SILP thickness over extended experiment times ($t_{\text{total}} = 600 \text{ min}$). Both reaction time ($t_{\text{rct}} = 20 \text{ min}$) and reaction temperature ($T = 50 \text{ }^\circ\text{C}$) were constant during all experiments in which the four SILP thicknesses (Table S2†) were analysed.



The basic data sets recorded with the four experiments are summarized in Fig. 5 as relative yield for each isomer and in Fig. 6 in the form of *Z/E*-selectivity (see Tables S3 and S4† for associated weights and moles of the two isomers). Three important observations can be made from the yields of the $\beta(Z)$ - and $\beta(E)$ -isomers in combination with the four different SILP thicknesses ($d_{\text{SILP}} = 15, 5, 3$, and 1 nm). First, compared to $d_{\text{SILP}} = 15$ nm, none of the yields for $d_{\text{SILP}} < 15$ nm reaches a steady-state even during the extended monitoring periods of 10 hours. While the yields mostly show only small changes and overall reveal a smooth progression, a true steady-state with constant yield is never attained. For $d_{\text{SILP}} = 15$ nm, however, a steady-state in the yields is recognizable after ~ 2 hours, characterized by a relatively low $m_{\beta(Z)}/m_{\beta(E)}$ -ratio of ~ 0.041 . That is, with a thick SILP layer (mesopores completely filled with IL), the reactor constantly produces almost exclusively the $\beta(E)$ -isomer. Second, for $d_{\text{SILP}} < 15$ nm the relative $\beta(E)$ -isomer content decreases over time (and with decreasing SILP thickness) while at the same time, the $\beta(Z)$ -isomer content rises gently with $d_{\text{SILP}} = 5$ and 3 nm. The strongest increase in the $\beta(Z)$ -isomer content was observed for the 1 nm-thick SILP after $t_{\text{exp}} \approx 130$ min and continued until the end of the experiment ($t_{\text{total}} = 600$ min). In fact, the $\beta(Z)$ -isomer content even exceeded the one of the $\beta(E)$ -isomer (for $t_{\text{exp}} > 400$ min), which remained unseen with the other SILP thicknesses. Summarizing, Fig. 5 shows that there is a systematic development in the relative yields of the two isomers with decreasing SILP thickness, which starts at the high and constant $\beta(E)$ -isomer level for $d_{\text{SILP}} = 15$ nm and

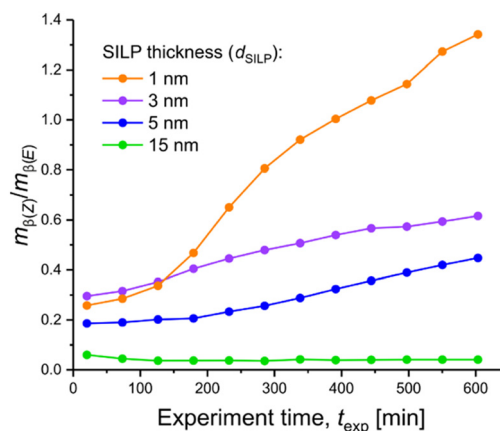


Fig. 6 Comparison of *Z/E*-selectivities via eqn (2) over the course of the four continuous-flow experiments with varied SILP thickness (cf. Fig. 5). Reaction time, $t_{\text{rct}} = 20$ min; reaction temperature, $T = 50$ °C.

climaxes in the boost of the $\beta(Z)$ -isomer yield with the thinnest SILP ($d_{\text{SILP}} = 1$ nm).

The peculiar rise in yield seen for the $\beta(Z)$ -isomer with the 1 nm-thick SILP manifests itself also in the *Z/E*-selectivities, which are shown in Fig. 6 for all four experiments from Fig. 5. Clearly $m_{\beta(Z)}/m_{\beta(E)}$ increases with decreasing SILP thickness. After 600 min, $m_{\beta(Z)}/m_{\beta(E)} = 0.041$ for $d_{\text{SILP}} = 15$ nm, 0.43 for $d_{\text{SILP}} = 5$ nm, 0.64 for $d_{\text{SILP}} = 3$ nm, and 1.35 for $d_{\text{SILP}} = 1$ nm. Overall, the transition from a thick to a thin SILP layer under continuous-flow conditions is accompanied by a remarkable increase in *Z/E*-selectivity, that is, by a factor of ~ 33 . This is in line with earlier findings using surface-

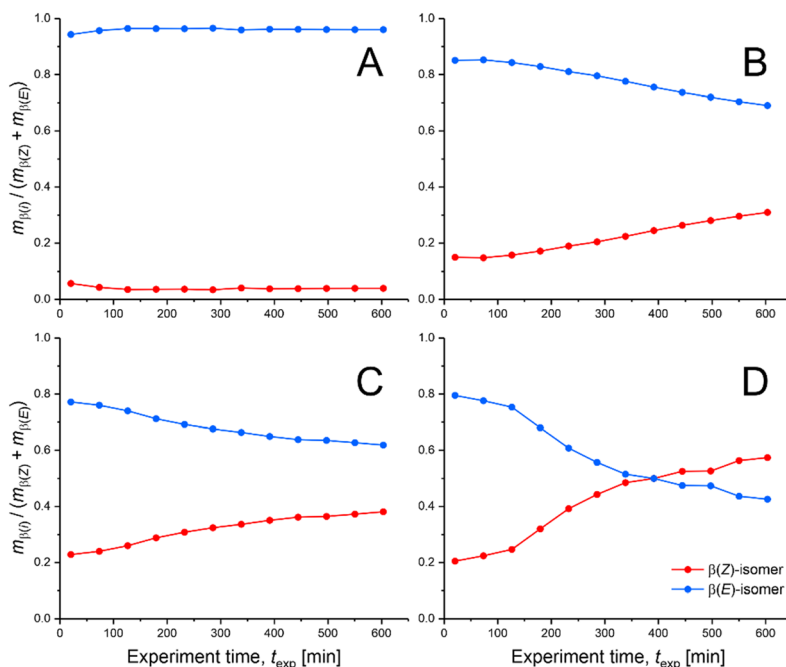


Fig. 5 Comparison of relative yields of the produced $\beta(Z)$ - and $\beta(E)$ -isomers via eqn (1) over the course of the four continuous-flow experiments ($t_{\text{total}} = 600$ min) with varied SILP thickness. The SILP thickness d_{SILP} was adjusted between (A) 15 nm (thick-SILP-limit), (B) 5 nm, (C) 3 nm, and (D) 1 nm (thin-SILP-limit). Reaction time, $t_{\text{rct}} = 20$ min; reaction temperature, $T = 50$ °C. Weights and moles of $\beta(Z)$ - and $\beta(E)$ -isomers underlying the reported yields can be found in Tables S3 and S4 (ESI†).



functionalized, polymeric, monolithic supports and clearly shows that this effect is independent of the support used.³²

Our findings are a result of different catalytic pathways that occur in a thin (1 nm wide) and in thicker SILPs, which explain the different *Z/E*-selectivities we observed. With increasing SILP layer thickness, the catalytic pathway responsible for the thin-SILP behaviour, *i.e.*, the formation of $\beta(Z)$ -isomer, becomes increasingly superimposed by those dominating in thicker SILPs, favouring the formation of the $\beta(E)$ -isomer. It is well known that the structural and dynamic properties of ILs can change dramatically at solid surfaces and under spatial confinement, in particular.⁵⁸ Indeed, their structure–transport relationships are modulated by interactions of the cations and anions with the support surface, *e.g.*, with silanol groups of the amorphous silica in the monolithic microreactor employed in the present work. As a consequence, structural ordering phenomena and a slow-down of the translational and rotational ion dynamics are often observed in ILs near solid surfaces.^{59–62}

For [BMIM][BF₄] used in this work, insightful results have been derived with molecular dynamics simulations conducted in a slit-pore confined by amorphous silica walls.^{62,63} Density oscillations were observed near the walls for both cations and anions indicating a layering effect, which is stronger for the [BF₄] anions than for the [BMIM] cations, as the former are more symmetric and smaller than the latter. In addition, the [BMIM] cations exhibit a preferred orientation in their first layer, where the methyl group and butyl tail point towards the silica wall and pore centre, respectively, and the imidazolium ring aligns with the surface. Furthermore, regarding the ion dynamics at a temperature of 300 K, the cation and anion motions are slowed down by roughly two orders of magnitude at the surface with respect to the centre of the pore. Here, the retardation of the ion dynamics rapidly declines with increasing distance from the surface and mainly affects the first two [BMIM][BF₄] layers.

These findings are highly relevant to our work, as distinctive density layers and strong mobility gradients exist near the surface. As a consequence, self-diffusion and structural relaxation are much slower there than in the bulk IL. The effects culminate in a highly dense and ordered (quasi-solid) first [BMIM][BF₄] layer at the silica surface, which is about 1 nm wide and characterized by a very low (quasi-frozen) mobility of the ionic species. It is reasonable to assume that in this quasi-frozen state the catalyst also experiences a slowdown of its translational and rotational dynamics and will most likely prefer the conformation that is found in the solid state, which we successfully determined by single-crystal X-ray analysis. **Rh-1** crystallizes in the triclinic space group *P* $\bar{1}$ with *a* = 975.6(5) pm, *b* = 1114.7(5) pm, *c* = 1352.7(7) pm, α = 75.446(16)°, β = 83.737(16)°, γ = 73.626(17)°, *Z* = 2. Relevant bond lengths and angles are summarized in Fig. 7. The most striking feature is that in the solid state, the pyridine group coordinates to the Rh-centre, thereby preventing any

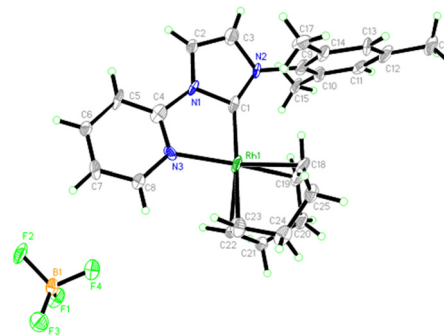


Fig. 7 Single-crystal X-ray structure of **Rh-1** (cf. Fig. 1). Relevant bond lengths (pm) and angles (°): Rh(1)–C(1) 197.4(10), Rh(1)–N(3) 213.4(8), Rh(1)–C(18) 213.7(9), Rh(1)–C(19) 214.4(10), Rh(1)–C(22) 220.6(9), Rh(1)–C(23) 222.3(10); C(1)–Rh(1)–N(3) 78.8(4), C(1)–Rh(1)–C(18) 96.1(4), N(3)–Rh(1)–C(18) 160.3(3), C(1)–Rh(1)–C(19) 100.0(4), N(3)–Rh(1)–C(19) 162.8(3), C(18)–Rh(1)–C(19) 36.6(3), C(1)–Rh(1)–C(22) 161.6(4), N(3)–Rh(1)–C(22) 94.6(3), C(18)–Rh(1)–C(22) 95.5(4), C(19)–Rh(1)–C(22) 81.1(4), C(1)–Rh(1)–C(23) 161.8(4), N(3)–Rh(1)–C(23) 98.0(3), C(18)–Rh(1)–C(23) 80.9(4), C(19)–Rh(1)–C(23) 88.3(4), C(22)–Rh(1)–C(23) 35.5(3).

rotation of the NHC, which is of utmost relevance for explaining our experimental findings (*vide infra*).

For comparison, the single-crystal X-ray structure of **Rh-2** together with relevant bond lengths and angles is shown in Fig. 8. **Rh-2** crystallises in the monoclinic space group *C*₂ with *a* = 2068.43(17) pm, *b* = 826.12(6) pm, *c* = 1516.11(13) pm, α = γ = 90°, β = 117.168(7)°, *Z* = 4.

Notably, the very low mobility will also affect the diffusion of solute molecules (that interact with the IL) in and through the density and mobility gradients of a SILP. For example, using fluorescence recovery after photobleaching (FRAP) in confocal microscopy,⁶⁴ it has been demonstrated that the

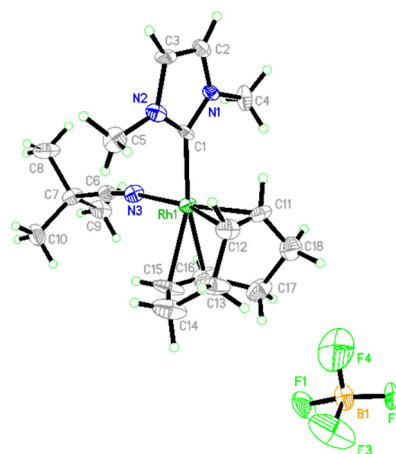


Fig. 8 Single-crystal X-ray structure of **Rh-2** (cf. Fig. 1). Relevant bond lengths (pm) and angles (°): Rh(1)–C(1) 202.4(3), Rh(1)–N(3) 205.5(3), Rh(1)–C(11) 207.7(10), Rh(1)–C(12) 217.0(11), Rh(1)–C(16) 217.8(8), Rh(1)–C(15) 225.7(7); C(1)–Rh(1)–N(3) 92.26(11), C(1)–Rh(1)–C(11) 89.8(3), N(3)–Rh(1)–C(11) 161.1(4), C(1)–Rh(1)–C(12) 90.9(3), N(3)–Rh(1)–C(12) 160.9(4), C(11)–Rh(1)–C(12) 37.73(15), C(1)–Rh(1)–C(16) 168.0(4), N(3)–Rh(1)–C(16) 92.0(2), C(11)–Rh(1)–C(16) 82.7(3), C(12)–Rh(1)–C(16) 88.8(3), C(1)–Rh(1)–C(15) 157.1(4), N(3)–Rh(1)–C(15) 88.6(2), C(11)–Rh(1)–C(15) 96.8(3), C(12)–Rh(1)–C(15) 81.3(3), C(16)–Rh(1)–C(15) 34.3(2).



diffusivity of a neutral dye in [BMIM][PF₆] confined in mesoporous silica is about two orders of magnitude lower than in the bulk IL. Similarly, the densified structure and slow dynamics of the SILP over a distance of 1–2 nm from the surface is expected to have an impact on the reactant molecules entering and moving within this region (in order to get in contact and arranged with the catalyst molecules residing there) and on the product molecules as they leave the local catalytic environment and the SILP after the reaction.

Compounds **1–5** dissolve well in the bulk IL. In the rigid, 1 nm-thick part of the SILP the reaction proceeds slower and diffusion of the larger products is delayed, which must result in an enrichment of the produced isomers in a thin SILP. To get evidence for a slower reaction and especially a slow release of the hydrosilylation products from the quasi-frozen part of the SILP, we performed continuous-flow experiments for two SILP thicknesses, *i.e.*, for $d_{\text{SILP}} = 1$ nm and $d_{\text{SILP}} = 15$ nm, with a subsequent rest period of the microreactor in pure MTBE of 18 hours, followed by workup of the MTBE phase. The results are compared in Fig. 9 with respect to *Z/E*-selectivity.

The first data point recorded ~ 20 min after the rest period of 18 hours corresponds to an elution volume ($0.066 \text{ mL min}^{-1} \times 20 \text{ min} = 1.32 \text{ mL}$) of roughly one reactor void volume ($V_{\text{void}} \approx 1.46 \text{ mL}$, ESI†). This point therefore represents the (almost entire) MTBE solution that had contact with the SILP throughout the 18 hours. Remarkably, a *Z/E*-ratio of 4.72 was received for the thin SILP. Such a high value has never been indicated or even measured in our continuous-flow experiments. It demonstrates that predominantly the $\beta(\text{Z})$ -isomer was produced and then trapped in the thin (rigid) SILP. In contrast, a *Z/E*-ratio of only 0.15 was observed for the thick SILP after this long extraction period, which confirms our conjecture that mostly $\beta(\text{E})$ -isomer was produced and retained in the bulk SILP saturating the mesopores of the monolith. Control experiments were performed to ensure that **Rh-1** does not catalyse the isomerization of the less stable $\beta(\text{Z})$ -isomer into

the thermodynamically favoured $\beta(\text{E})$ -isomer during the rest period²¹ and thereby cause a bias in the subsequently analysed isomer pattern. For this purpose, microreactors with freshly prepared thin or thick SILP (containing **Rh-1**) were flushed with MTBE containing only a mixture of $\beta(\text{E})$ - and $\beta(\text{Z})$ -isomers that were beforehand synthesized and isolated. This isomer solution was then also allowed to rest in the microreactor for 18 hours, as in the extraction experiment, and then analysed. In fact, no changes in the original isomer composition could be detected after this period, implying that **Rh-1** does not catalyse *Z/E*-isomerization. The strong dependence of the $\beta(\text{Z})/\beta(\text{E})$ -ratio from the thickness of the SILP layer can thus be fully explained considering both, the structure of the catalyst (and of the resulting transition states during hydrosilylation) and the different morphologies of the SILP layers with different thickness.

Clearly, the SILP possesses a substantial gradient in terms of morphology and rigidity as we move from the pore walls to the centre of the pores. In the thin-SILP-limit ($d_{\text{SILP}} \approx d_1$, Fig. 10), the SILP predominantly consists of a quasi-frozen monolayer with a much lower diffusive mobility of the substrates and products than in the bulk IL. The cationic catalyst will also be firmly held in the rigid SILP, most likely supported by silanophilic interactions with the surface (as implied by its positioning in Fig. 10). As a consequence, the integration of the reactants into (as well as the release of the even larger products from) the rigid SILP, indicated by the phase transfer in Fig. 10, is slow, as confirmed by the extraction experiment (Fig. 9). On the other hand, in the thick-SILP-limit ($d_{\text{SILP}} \approx d_2$, Fig. 10), the SILP mostly consists of bulk SILP, which exhibits the properties (local density and diffusive mobility) of the bulk IL. There, the catalyst can move more freely as compared to the rigid SILP and also phase transfer of reactants and products between the mobile MTBE phase and the bulk region of the SILP is faster than between MTBE and rigid SILP.

The observed preferential formation of the $\beta(\text{Z})$ -isomer by **Rh-1** in the thin SILP can be accounted for by the quasi-frozen IL resulting in a preferred transition state with the pyridine group coordinating to the Rh-centre, as found in the solid-state structure of **Rh-1** (Scheme 3). The resulting $\beta(\text{Z})$ -selectivity can be explained by the modified Chalk–Harrod mechanism:^{9,65,66} Formation of a Rh-hydride is followed by the coordination of the alkyne. Alkyne insertion into the Rh–Si bond leads to the $\beta(\text{Z})$ -silylvinylidene intermediate. Subsequently, two reaction pathways can emerge, depending on whether the reaction proceeds in the bulk or in the quasi-frozen SILP. Thus, the formed $\beta(\text{Z})$ -silylvinylidene may lead to the $\beta(\text{E})$ -silylvinylidene and, consequently, to the $\beta(\text{Z})$ -vinylsilane (red pathway). The quasi-frozen SILP restricts the rotation of the NHC of the confined catalyst, which results in a strong steric interaction between the pyridine or mesityl moiety and the SiMe₂Ph group. Rotation of the HC–CPh bond eliminates this steric interaction making the $\beta(\text{E})$ -silylvinylidene the preferred

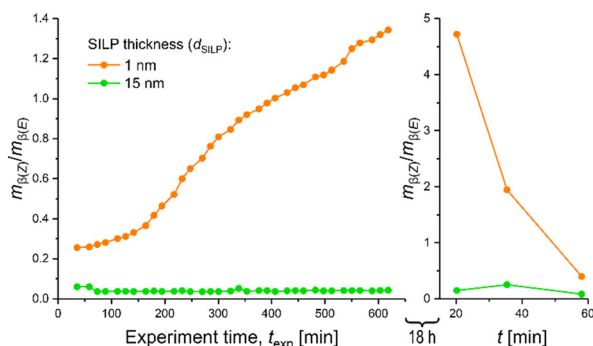


Fig. 9 Continuous-flow experiments with thin SILP ($d_{\text{SILP}} = 1$ nm) and thick SILP ($d_{\text{SILP}} = 15$ nm), followed by a flushing step of the microreactor with pure MTBE and a rest period (stopped flow) of 18 hours. Afterwards, the MTBE solution was flushed out and the effluent analysed for *Z/E*-selectivity using the second dimension of the setup depicted in Fig. 3.



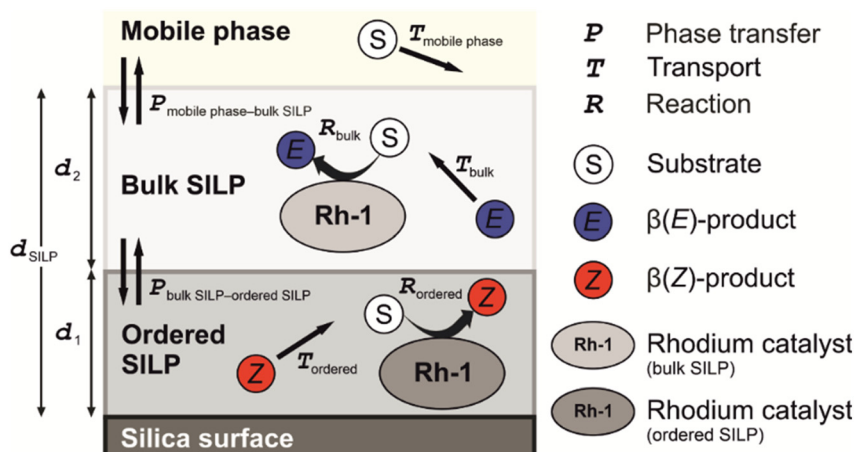
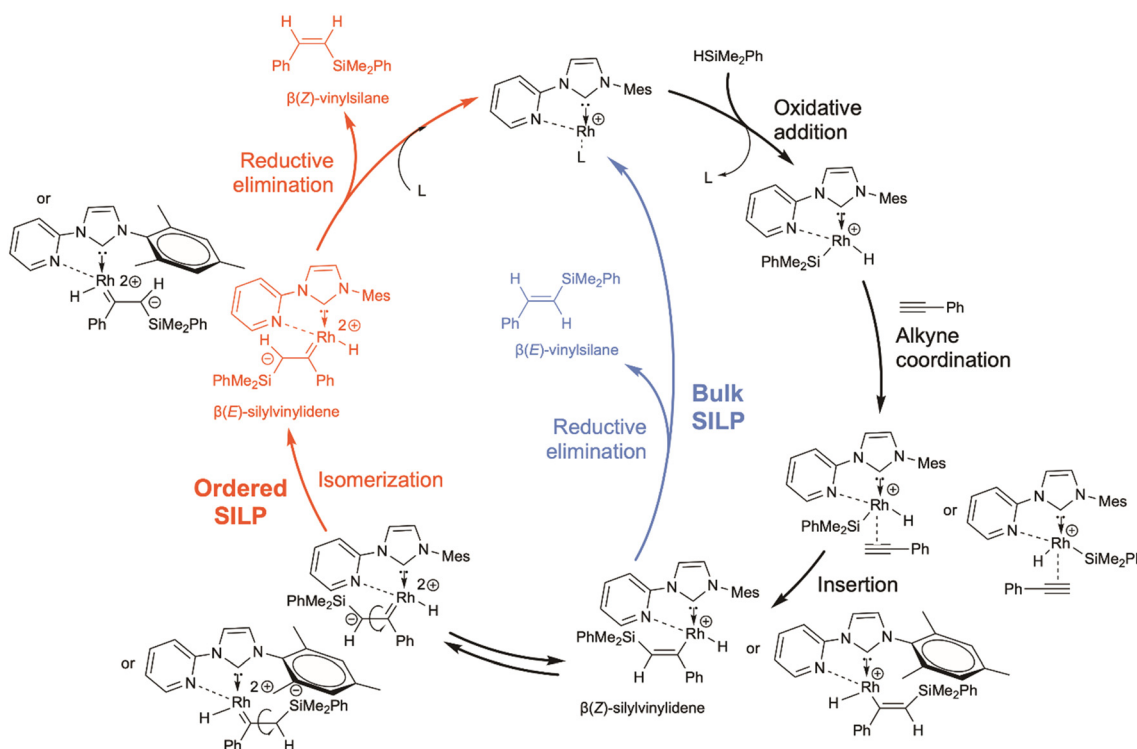


Fig. 10 Schematic representation of relevant structural and dynamic aspects in the continuous-flow hydrosilylation reaction between PA and DMPS with Rh-1 in the SILP formed by [BMIM][BF₄] in the ~30 nm wide mesopores of a (macro-mesoporous) silica monolith.



Scheme 3 Modified Chalk-Harrod mechanism for the preferential formation of $\beta(Z)$ - or $\beta(E)$ -vinylsilane isomer in the continuous-flow hydrosilylation reaction between PA and DMPS with Rh-1 in the SILP formed by [BMIM][BF₄] on a macro-mesoporous silica monolith.

configuration, which finally results in the formation of the $\beta(Z)$ -vinylsilane. *Vice versa*, the bulk SILP allows for a decooordination of the pyridine group and rotation of the NHC, which places less steric constraints on the initial $\beta(Z)$ -silylvinylidene, resulting in predominant formation of the $\beta(E)$ -vinylsilane. The joint role of a coordinating pyridine group and the restricted mobility within a thin SILP was confirmed by running the same hydrosilylation reaction in a thin SILP ($d_{\text{SILP}} = 1$ nm) at 50 °C by the action of Rh(i) NHC catalysts that did not possess any chelating groups, *i.e.*, **Rh-2** and **Rh-3** (Fig. 1). With these Rh-catalysts, $m_{\beta(Z)}/m_{\beta(E)}$

did not exceed 0.65 (**Rh-2**) and 0.12 (**Rh-3**) over a reaction time of 140 min and 540 min, respectively (*cf.* Fig. S7 and Table S5 in the ESI†).

All this fits to our experimental data, where starting in the thin-SILP-limit ($d_{\text{SILP}} = 1$ nm), the dominating *Z*-selectivity in the rigid SILP becomes increasingly superimposed by *E*-selectivity for larger SILP thickness ($d_{\text{SILP}} = 3$ and 5 nm) in the bulk SILP, until the $\beta(E)$ -isomer dominates and is correspondingly produced at a constantly high level, without sign of increasing *Z/E*-selectivity over time, in the thick-SILP-limit ($d_{\text{SILP}} = 15$ nm, *cf.* Fig. 5).



Conclusions

Continuous-flow studies of the hydrosilylation reaction between PA (**1**) and DMPS (**2**) in the presence of the cationic Rh-NHC complex **Rh-1**, taking place within the SILP formed by [BMIM][BF₄] on the mesopore surface of a macro-mesoporous silica monolith, have revealed a clear dependence of *Z/E*-selectivity on the SILP thickness varied from $d_{\text{SILP}} = 1$ nm to 15 nm. In the thick-SILP-limit ($d_2 \gg d_1$ and $d_{\text{SILP}} \approx d_2$, Fig. 10), the SILP mostly consists of bulk IL, in which hydrosilylation leads to the predominant formation of the thermodynamically more stable $\beta(E)$ -vinylsilane isomer (**5**). The thick-SILP-limit is approached in our experiments as the mesopores are completely filled with IL ($d_{\text{SILP}} = 15$ nm), while the thin-SILP-limit is reached for $d_{\text{SILP}} = 1$ nm ($d_2 \rightarrow 0$ and $d_{\text{SILP}} \approx d_1$, Fig. 10). In the latter situation, the SILP mostly consists of a quasi-frozen monolayer of IL cations and anions, in which the catalyst is firmly embedded experiencing restricted mobility. In this thin-SILP-limit, hydrosilylation results in the predominant formation of the kinetically favoured $\beta(Z)$ -vinylsilane isomer (**4**) via generation of the $\beta(E)$ -silylvinylidene intermediate through isomerization of the $\beta(Z)$ -silylvinylidene (Scheme 3).

From a more general point of view, our work highlights the importance of SILP morphology and, in particular, of the adjusted SILP thickness, on the structural and transport dynamics as well as the selectivity of reactions occurring inside a SILP. Implementation as continuous-flow experiment with UHPLC-separation based online analytics has been essential in recognizing the evolution of *Z/E*-selectivity over hours and thereby tracing the slow release of the *Z*-isomer from the rigid part of the SILP as a function of d_{SILP} . To the best of our knowledge, this is the first example for a confinement jointly generated by a tailored structural motif in a catalyst (chelating ligand) and the morphology of a very thin IL layer. We therefore anticipate similar effects of rigid vs. bulk SILP microstructure, which become convoluted as the SILP thickness increases from thin to thick, to be manifested also in other reactions, e.g., in the closely related hydroboration and hydroamination reactions.

Experimental

Chemicals and columns

Methanol (HPLC grade) came from VWR Chemicals (Darmstadt, Germany) and water was used as received from a Milli-Q gradient purification system (Millipore, Bedford, MA). HPLC grade MTBE and CH₂Cl₂ as well as PA (98%), DMPS ($\geq 98\%$), and [BMIM][BF₄] ($\geq 97\%$) were obtained from Sigma Aldrich (St. Louis, MO). DMPS was stored at 4 °C, while MTBE was dried over sodium to remove residual water and stored under inert gas with a molecular sieve (3 Å) until use. Catalysts **Rh-1**,²⁴ **Rh-2**,⁶⁷ and **Rh-3** (ref. 68) were prepared according to established protocols. For hydrosilylation, we used a macro-mesoporous silica monolith received from Merck KGaA (Darmstadt, Germany)

as 4.6 mm inner diameter $\times$ 100 mm length column clad in polyether ether ketone. The silica monolith had mean macropore and mesopore sizes of 1.84 μm and 32.9 nm, respectively. The reaction mixture (compounds **1–5**, Scheme 2) was separated on a 3.0 mm inner diameter $\times$ 100 mm length UHPLC column packed with phenyl-hexyl-modified, fully porous silica particles with nominal mean particle and mesopore sizes of 1.8 μm and 9.5 nm, respectively (Agilent Technologies, Santa Clara, CA).

Microreactor operation

Continuous-flow hydrosilylation was realized by implementation of commercially available HPLC instrumentation, as depicted in Fig. 3. Substrate solution (25 mM PA and 32.5 mM DMPS in dry, absolute MTBE) was connected to a binary pump with a septum and metal capillary. This solution was pumped at a flow rate of $Q = 0.066 \text{ mL min}^{-1}$ through the reactor placed in a thermostatted compartment (50 °C). The reaction solution from the microreactor was transferred by a valve to the second (analytical) dimension, where the reaction mixture was separated using UHPLC (cf. Fig. 4) and analyzed with a diode-array detector (260 nm).

Hardware components (as shown in Fig. 3)

1, substrate solution; 2, autosampler (Agilent 1290 Infinity II Series, G7129B); 3, quaternary pump for second-dimension online analytics (Agilent 1290 Infinity II Series, G7104A); 4, binary pump for first-dimension reaction control (Agilent 1290 Infinity II Series, G7120A); 5, 2-position/6-port valve for connecting first and second dimensions (Agilent 1290 Infinity Series, G1170A); 6, online diode-array detector (Agilent 1260 Infinity II Series, G7115A); 7, thermostatted compartment with UHPLC column (Agilent 1290 Infinity II Series, G7116B); 8, thermostatted compartment with monolithic microreactor (Agilent 1290 Infinity II Series, G7116B); 9, online diode-array detector (Agilent 1290 Series, G4212A).

SILP preparation

For impregnation, the monolith was flushed with a solution of [BMIM][BF₄] in CH₂Cl₂ (also containing **Rh-1**). After removal of CH₂Cl₂ by overnight vacuum-drying, a layer of the catalyst-containing IL remained on the monolith surface, almost exclusively ($>99\%$) intraskelton mesopore surface. Variation of the SILP thickness was achieved by adjusting the IL volume deposited inside the mesopores (V_{IL})

$$d_{\text{SILP}} = \frac{V_{\text{IL}}}{m_{\text{silica}} A_{\text{meso}}} \quad (3)$$

where m_{silica} is the mass of solid silica present in the column and A_{meso} is the specific mesopore surface area of the monolith (110 m² g⁻¹). In practice, a SILP thickness is first specified and the corresponding V_{IL} calculated using eqn (3). m_{silica} required for calculation of d_{SILP} can be accessed by



$$m_{\text{silica}} = (V_{\text{column}} - V_{\text{void}})\rho_{\text{silica}} \quad (4)$$

where V_{column} is the volume of the empty column and V_{void} is the void volume of the column in the presence of the silica monolith; ρ_{silica} is the density of the solid, amorphous silica ($\sim 2.2 \text{ g cm}^{-3}$).^{69,70}

With V_{void} determined as the elution volume of a small, non-adsorbing tracer on the monolithic column, verified by the elution of PA in pure acetonitrile, we found $m_{\text{silica}} = 0.447 \text{ g}$ (additional details behind the determination of m_{silica} can be found in the ESI†). For a targeted SILP thickness of 1 nm, for example, V_{IL} is then given by

$$V_{\text{IL}} = 1 \times 10^{-7} \text{ cm} \times 0.447 \text{ g} \times 110 \times 10^4 \text{ cm}^2 \text{ g}^{-1} = 0.05 \text{ mL}$$

Accordingly, the dry monolithic column was saturated up to V_{void} with IL in CH_2Cl_2 containing the corresponding Rh-catalyst. The solution used for preparation of the 1 nm-thick SILP in the above example thus consisted of the $V_{\text{IL}} = 0.05 \text{ mL}$ [BMIM][BF_4] and a CH_2Cl_2 volume of $V_{\text{void}} - V_{\text{IL}}$. After SILP formation in the mesopores of the monolith, CH_2Cl_2 was evaporated and the column flushed with MTBE. Four SILP thicknesses were realized, ranging from the thin-SILP-limit ($d_{\text{SILP}} = 1 \text{ nm}$), below which a uniform monolayer of [BMIM][BF_4] will be difficult to establish along the mesopore surface, to the thick-SILP-limit ($d_{\text{SILP}} = 15 \text{ nm}$), when the $\sim 30 \text{ nm}$ wide mesopores become fully saturated with IL. SILP thicknesses and associated values of V_{IL} and $m_{\text{Rh-1}}$ (mass of Rh-catalyst in each SILP) are summarized in Table S2 of the ESI.†

Stopped-flow experiments

Experiments were run for $d_{\text{SILP}} = 1 \text{ nm}$ and 15 nm. After the (standard) continuous-flow experiment, in which the reactor was fed with reaction solution (25 mM PA and 32.5 mM DMPS in MTBE) for 600 min, the pumping system was switched to pure MTBE and the reactor flushed until no substrate and product molecules could be detected any longer in the effluent. Then, the flow was stopped and the reactor left at rest in pure MTBE for 18 hours. This long extraction period guaranteed a sufficient diffusive release of the trapped $\beta(\text{Z})$ - and $\beta(\text{E})$ -isomers from the SILP into the MTBE phase. After this period the flow was restarted (pure MTBE, $Q = 0.066 \text{ mL min}^{-1}$) and the reactor effluent analysed for the amount of the two isomers released from the corresponding SILP phase.

Data availability

- Data for this article, including primary data for the NMR spectra, kinetic measurements, and chromatograms are available at the Data Repository of the University of Stuttgart (DARUS) [<https://darus.uni-stuttgart.de>] in connection with the DOI of the published paper.

- Further data (experimental details, chromatograms, NMR spectra, crystallographic details for **Rh-1** and **Rh-2**) supporting this article have been included as part of the ESI.†

- Deposition numbers CCDC 2430795 (**Rh-1**) and 2441044 (**Rh-2**) contain the supplementary crystallographic data for this paper. These data are provided free of charge by the joint Cambridge Crystallographic Data Centre and Fachinformationszentrum Karlsruhe Access Structures Service.

Author contributions

André Böth: formal analysis, investigation, validation, visualization, writing – original draft. Florian Kaltwasser: formal analysis, investigation, validation, visualization, writing – review & editing. Christian Friedigkeit: formal analysis, investigation. Boshra Atwi: investigation: resources. Wolfgang Frey: formal analysis. Michael R. Buchmeiser: conceptualization, funding acquisition, resources, supervision, writing – original draft. Ulrich Tallarek: conceptualization, funding acquisition, methodology, resources, supervision, writing – original draft. All authors have approved the final version of the manuscript.

Conflicts of interest

The authors declare no competing financial interests.

Acknowledgements

Financial support by the Deutsche Forschungsgemeinschaft DFG (German Research Foundation, project ID 358283783 – SFB 1333/2 2022) is gratefully acknowledged. We further thank Agilent Technologies (Waldbronn, Germany) for support with analytical equipment and Benjamin Peters (Instrumental Analytics R&D, Merck KGaA, Darmstadt, Germany) for the gift of monolithic silica columns.

Notes and references

- 1 L. N. Lewis, J. Stein, Y. Gao, R. E. Colborn and G. Hutchins, *Platinum Met. Rev.*, 1997, **41**, 66–75.
- 2 L. Cornelissen, V. Cirriez, S. Vercrussey and O. Riant, *Chem. Commun.*, 2014, **50**, 8018–8020.
- 3 J. Szudkowska-Frątczak, G. Hreczycho and P. Pawluć, *Org. Chem. Front.*, 2015, **2**, 730–738.
- 4 T. Komiyama, Y. Minami and T. Hiyama, *ACS Catal.*, 2017, **7**, 631–651.
- 5 A. Vignesh, J. Liu, Z. Wang, Y. Liu and Z. Ke, *Org. Chem. Front.*, 2024, **11**, 576–596.
- 6 B. Marciniec, H. Maciejewski and P. Pawluć, Hydrosilylation of Carbon–Carbon Multiple Bonds—Application in Synthesis and Materials Science, in: *Organosilicon Compounds*, ed. V. Y. Lee, Academic Press, New York, 2017, vol. 2, pp. 169–217.
- 7 M. A. Brook, *Silicon in Organic, Organometallic and Polymer Chemistry*, John Wiley & Sons, New York, 2000.
- 8 A. K. Franz and S. O. Wilson, *J. Med. Chem.*, 2013, **56**, 388–405.
- 9 B. Marciniec, Hydrosilylation of Alkynes and Their Derivatives, in: *Hydrosilylation. Advances in Silicon Science*,



- ed. B. Marciniak, Springer, Dordrecht, 2009, vol. 1, pp. 53–86.
- 10 A. Zanardi, E. Peris and J. A. Mata, *New J. Chem.*, 2008, **32**, 120–126.
 - 11 S. Demir, Y. Gökçe, N. Kaloğlu, J. B. Sortais, C. Darcel and I. Özdemir, *Appl. Organomet. Chem.*, 2013, **27**, 459–464.
 - 12 J. Sun and L. Deng, *ACS Catal.*, 2016, **6**, 290–300.
 - 13 Y. Mutoh, Y. Mohara and S. Saito, *Org. Lett.*, 2017, **19**, 5204–5207.
 - 14 P. Wang, Y. Lai, P. Wang, X. Leng, J. Xiao and L. Deng, *J. Am. Chem. Soc.*, 2021, **143**, 12847–12856.
 - 15 R. R. Behera, R. Saha, A. A. Kumar, S. Sethi, N. C. Jana and B. Bagh, *J. Org. Chem.*, 2023, **88**, 8133–8149.
 - 16 M. González-Lainez, M. V. Jiménez, V. Passarelli and J. J. Pérez-Torrente, *Dalton Trans.*, 2023, **52**, 11503–11517.
 - 17 A. Tyagi, S. Yadav, P. Daw, C. Ravi and J. K. Bera, *Polyhedron*, 2019, **172**, 167–174.
 - 18 A. Mumcu, M. O. Karataş, N. Özdemir, A. Erdoğan and H. Küçükbay, *J. Organomet. Chem.*, 2023, **997**, 122791.
 - 19 J. J. Pérez-Torrente, D. H. Nguyen, M. V. Jiménez, F. J. Modrego, R. Puerta-Oteo, D. Gómez-Bautista, M. Iglesias and L. A. Oro, *Organometallics*, 2016, **35**, 2410–2422.
 - 20 R. Takeuchi and I. Ebata, *Organometallics*, 1997, **16**, 3707–3710.
 - 21 L. P. Morales-Cerón, P. Lara, J. López-Serrano, L. L. Santos, V. Salazar, E. Álvarez and A. Suárez, *Organometallics*, 2017, **36**, 2460–2469.
 - 22 R. Puerta-Oteo, J. Munárriz, V. Polo, M. V. Jiménez and J. J. Pérez-Torrente, *ACS Catal.*, 2020, **10**, 7367–7380.
 - 23 W. A. Solomonsz, G. A. Rance, M. Suyetin, A. La Torre, E. Bichoutskaia and A. N. Khlobystov, *Chem. – Eur. J.*, 2012, **18**, 13180–13187.
 - 24 P. K. R. Panyam, B. Atwi, F. Ziegler, W. Frey, M. Nowakowski, M. Bauer and M. R. Buchmeiser, *Chem. – Eur. J.*, 2021, **27**, 17220–17229.
 - 25 M. Roemer, V. R. Gonçalves, S. T. Keaveney, I. Pernik, J. Lian, J. Downes, J. J. Gooding and B. A. Messerle, *Catal. Sci. Technol.*, 2021, **11**, 1888–1898.
 - 26 M. Roemer, S. T. Keaveney, J. E. Downes, S. Gautam, J. J. Gooding and B. A. Messerle, *Catal. Sci. Technol.*, 2022, **12**, 226–236.
 - 27 B. Sánchez-Page, J. Munárriz, M. V. Jiménez, J. J. Pérez-Torrente, J. Blasco, G. Subias, V. Passarelli and P. Patricia Álvarez, *ACS Catal.*, 2020, **10**, 13334–13351.
 - 28 A. Riisager and R. Fehrmann, Supported Ionic Liquid Phase Catalysts, in: *Ionic Liquids in Synthesis*, ed. P. Wasserscheid and T. Welton, Wiley-VCH, Weinheim, 2008, vol. 2, pp. 527–558.
 - 29 C. van Doorslaer, J. Wahlen, P. Mertens, K. Binnemans and D. de Vos, *Dalton Trans.*, 2010, **39**, 8377–8390.
 - 30 R. Kukawka, A. Pawłowska-Zygarowicz, J. Działkowska, M. Pietrowski, H. Maciejewski, K. Bica and M. Smiglak, *ACS Sustainable Chem. Eng.*, 2019, **7**, 4699–4706.
 - 31 J. Dupont, B. C. Leal, P. Lozano, A. L. Monteiro, P. Migowski and J. D. Scholten, *Chem. Rev.*, 2024, **124**, 5227–5420.
 - 32 P. K. R. Panyam and M. R. Buchmeiser, *Faraday Discuss.*, 2023, **244**, 39–50.
 - 33 G. Franciò, U. Hintermair and W. Leitner, *Philos. Trans. R. Soc., A*, 2015, **373**, 20150005.
 - 34 E. García-Verdugo, B. Altava, M. I. Burguete, P. Lozano and S. V. Luis, *Green Chem.*, 2015, **17**, 2693–2713.
 - 35 J. M. Marinkovic, A. Riisager, R. Franke, P. Wasserscheid and M. Haumann, *Ind. Eng. Chem. Res.*, 2019, **58**, 2409–2420.
 - 36 M. Aubermann, M. Haumann, D. Wisser and P. Wasserscheid, *ACS Sustainable Chem. Eng.*, 2022, **10**, 6973–6980.
 - 37 A. Billion, A. Vogel, J. Schulte, H. Scherer and I. Krossing, *ChemCatChem*, 2023, **15**, e202300844.
 - 38 P. Latos, A. Wolny and A. Chrobok, *Materials*, 2023, **16**, 2106.
 - 39 B. Autenrieth, W. Frey and M. R. Buchmeiser, *Chem. – Eur. J.*, 2012, **18**, 14069–14078.
 - 40 B. Sandig and M. R. Buchmeiser, *ChemSusChem*, 2016, **9**, 2917–2921.
 - 41 B. Sandig, L. Michalek, S. Vlahovic, M. Antonovici, B. Hauer and M. R. Buchmeiser, *Chem. – Eur. J.*, 2015, **21**, 15835–15842.
 - 42 A. Odedra, K. Geyer, T. Gustafsson, R. Gilmour and P. Seeberger, *Chem. Commun.*, 2008, 3025–3027.
 - 43 M. Jankowska-Wajda, R. Kukawka, M. Smiglak and H. Maciejewski, *New J. Chem.*, 2018, **42**, 5229–5236.
 - 44 H. Acikalin, P. K. R. Panyam, A. W. Shaikh, D. Wang, S. R. Kousik, P. Atanasova and M. R. Buchmeiser, *Macromol. Chem. Phys.*, 2023, **224**, 2200234.
 - 45 C.-L. Tavera-Méndez, A. Bergen, S. Trzeciak, F. W. Heinemann, R. Graf, D. Zahn, K. Meyer, M. Hartmann and D. Wisser, *Chem. – Eur. J.*, 2024, **30**, e202303673.
 - 46 G. Guiochon, *J. Chromatogr. A*, 2007, **1168**, 101–168.
 - 47 D. Enke, R. Gläser and U. Tallarek, *Chem. Ing. Tech.*, 2016, **88**, 1561–1585.
 - 48 A. Galarneau, A. Sachse, B. Said, C.-H. Péllisson, P. Boscaro, N. Brun, L. Courtheoux, N. Olivi-Tran, B. Coasne and F. Fajula, *C. R. Chim.*, 2016, **19**, 231–247.
 - 49 C. P. Haas, T. Müllner, R. Kohns, D. Enke and U. Tallarek, *React. Chem. Eng.*, 2017, **2**, 498–511.
 - 50 A. Feinle, M. S. Elsaesser and N. Huesing, *Chem. Soc. Rev.*, 2016, **45**, 3377–3399.
 - 51 A. Inayat, B. Reinhardt, J. Herwig, C. Küster, H. Uhlig, S. Krenkel, E. Raedlein and D. Enke, *New J. Chem.*, 2016, **40**, 4095–4101.
 - 52 D. Stoeckel, C. Kübel, K. Hormann, A. Hölzel, B. M. Smarsly and U. Tallarek, *Langmuir*, 2014, **30**, 9022–9027.
 - 53 S. Altmaier and K. Cabrera, *J. Sep. Sci.*, 2008, **31**, 2551–2559.
 - 54 K. Hormann, T. Müllner, S. Bruns, A. Hölzel and U. Tallarek, *J. Chromatogr. A*, 2012, **1222**, 46–58.
 - 55 A. Böth, T. Roider, F. Ziegler, X. Xie, M. R. Buchmeiser and U. Tallarek, *ChemCatChem*, 2023, **15**, e202201268.
 - 56 T. H. Walter and R. W. Andrews, *TrAC, Trends Anal. Chem.*, 2014, **63**, 14–20.
 - 57 A. Böth, D. Foshag, C. Schulz, B. Atwi, S. E. Maier, D. P. Estes, M. R. Buchmeiser, T. van de Goor and U. Tallarek, *J. Chromatogr. A*, 2024, **1730**, 465165.



- 58 S. Zhang, J. Zhang, Y. Zhang and Y. Deng, *Chem. Rev.*, 2017, **117**, 6755–6833.
- 59 S. Perkin, *Phys. Chem. Chem. Phys.*, 2012, **14**, 5052–5062.
- 60 M. Foroutan, S. M. Fatemi and F. Esmaeilian, *Eur. Phys. J. E: Soft Matter Biol. Phys.*, 2017, **40**, 19.
- 61 F. Borghi and A. Podestà, *Adv. Phys.: X*, 2020, **5**, 1736949.
- 62 R. Köster and M. Vogel, *J. Chem. Phys.*, 2022, **156**, 074501.
- 63 T. Pal, C. Beck, D. Lessnich and M. Vogel, *J. Phys. Chem. C*, 2018, **122**, 624–634.
- 64 A. D. Drake, Y. He, F. Ladipo, B. L. Knutson and S. E. Rankin, *J. Phys. Chem. B*, 2024, **128**, 3046–3060.
- 65 A. J. Chalk and J. F. Harrod, *J. Am. Chem. Soc.*, 1965, **87**, 16–21.
- 66 L.-W. Chung, Y.-D. Wu, B. M. Trost and Z. T. Ball, *J. Am. Chem. Soc.*, 2003, **125**, 11578–11582.
- 67 I. M. Daubitz, M. P. Sullivan, M. John, D. C. Goldstone, C. G. Hartinger and N. Metzler-Nolte, *Inorg. Chem.*, 2020, **59**, 17191–17199.
- 68 M. Mayr, K. Wurst, K.-H. Ongania and M. R. Buchmeiser, *Chem. – Eur. J.*, 2004, **10**, 1256–1266.
- 69 H. U. K. Jatoti, M. Goepel, D. Poppitz, R. Kohns, D. Enke, M. Hartmann and R. Gläser, *Front. Chem. Eng.*, 2021, **3**, 789416.
- 70 A. Galarneau, Z. Abid, B. Said, Y. Didi, K. Szymanska, A. Jarzębski, F. Tancrét, H. Hamaizi, A. Bengueddach, F. Di Renzo and F. Fajula, *Inorganics*, 2016, **4**, 9.

