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A novel synthetic approach to poly(hydrosiloxane)s via hydrolytic oxidation of primary organosilanes with a AuNPs-stabilized Pickering interfacial catalyst†

Ravi Shankar,* Bhawana Jangir and Asmita Sharma

The triblock copolymer, $\text{PiBA}_{20}\text{-}b\text{-PDMS}_{75}\text{-}b\text{-PiBA}_{20}$ (PiBA = polyisobornylacrylate, PDMS = polydimethylsiloxane), **1** is employed as an amphiphilic scaffold for surface functionalization of AuNPs of 10–15 nm size in chloroform. In water–chloroform biphasic medium, the AuNPs form a randomly close packed assembly at the interface and remain stable for several months under ambient conditions. The interfacially stabilized AuNPs act as a selective and recyclable catalyst for the hydrolytic oxidation of primary organosilanes, RSiH_3 and offer a promising route for high yield synthesis of poly(alkyl/arylhydrosiloxane)s $\text{H}_2\text{RSiO}[\text{RHSiO}]_n\text{SiRH}_2$, **2–6** [$\text{R} = \text{Ph}$ (**2**), *n*-hexyl (**3**), *cyclo*-hexyl (**4**), $\text{Et}_3\text{SiCH}_2\text{CH}_2$ (**5**), $\text{PhMe}_2\text{SiCH}_2\text{CH}_2$ (**6**)].

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Introduction

Polysiloxanes, $[\text{RR}^1\text{SiO}]_n$ ($\text{R}, \text{R}^1 = \text{alkyl or aryl}$) are among the well-known synthetic polymers and differ in many respects from other inorganic polymers due to their unique properties associated with a flexible Si–O–Si backbone.¹ As a result, these polymers find widespread applications in industry and modern technology. The classical synthetic approach to polysiloxanes involves acid or base-catalyzed hydrolysis and condensation of chloro/alkoxy-substituted organosilane monomers.² However, a major limitation of this approach arises from the sensitivity of the Si–O–Si bond to pH conditions and formation of undesired side products due to cleavage or redistribution/disproportionation of the siloxane framework are often observed. The advent of anionic/cationic ring opening polymerization of *cyclo*-siloxanes has provided a great deal of impetus in synthetic chemistry of high molecular weight polysiloxanes and its scope to construct block copolymers of varying compositions has been well documented.^{1,3} In recent years, considerable attention has been devoted to develop catalytic routes to siloxane frameworks with well-defined molecular weight and control over dimensionality.⁴ In this context, Lewis acid catalysts such as $\text{B}(\text{C}_6\text{F}_5)_3$, BiCl_3 , $\text{Sc}(\text{OTf})_3$, *etc.* have shown promise in terms of their selectivity for various functional group transformation and formation of Si–O–Si bonds under non-hydrolytic conditions. These studies are primarily driven by an impetus to expand the

library of functional synthons for the development of new siloxane-based materials. Several metal-based heterogeneous and homogeneous catalysts have been investigated for hydrolytic oxidation of Si–H bonds in organosilanes.⁵ However, these catalysts often lack selectivity as the reactions are accompanied by the formation of a mixture of silanols, disiloxanes and siloxane oligomers. Recent studies have shown that gold nanoparticles on various solid supports are excellent heterogeneous catalysts for selective hydrolytic oxidation of tertiary organosilanes, R_3SiH to afford exclusive formation of organosilanols with the elimination of dihydrogen as the byproduct.⁶

Poly(hydromethylsiloxane), $[\text{Me}(\text{H})\text{SiO}]_n$ (abbreviated as PMHS) has attracted considerable attention in view of the presence of Si–H groups in the repeat unit. Hemery *et al.* have reported the synthesis of high molecular weight PMHS by cationic ring opening polymerization of 1,3,5,7-tetramethylcyclotetrasiloxane in aqueous emulsion using dodecylbenzenesulfonic acid and $\text{C}_{12}\text{H}_{25}(\text{OCH}_2\text{CH}_2)_{23}\text{OH}$ (Brij35) as emulsifying agents.⁷ The presence of Si–H groups in PMHS offers opportunity for chemical modification and incorporation of appended functional moieties on siloxane backbone by hydrosilylation, dehydrocoupling or click reactions.⁸ The functional groups endow these polymers with interesting physico-chemical properties and serve as scaffolds in the synthesis of Noble metal nanoparticles. The reducing property of PMHS has also been utilized for the reduction of a variety of organic substrates.⁹

The organization of colloidal metal/metal oxide particles at liquid–liquid interface, a phenomenon first described by Pickering, has emerged as an active area of research in catalysis.¹⁰ For example, the entrapment and stabilization of catalytically active nano/colloidal particles (*e.g.* Pd, SiO_2) at the interface has

Department of Chemistry, Indian Institute of Technology, Hauz Khas, New Delhi-110016, India. E-mail: shankar@chemistry.iitd.ac.in

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been employed as an alternative to classical methods for chemical transformation of organic substrates and biomass refining.¹¹ The energy (ΔE) of nanoparticles at the interface is governed by nature of the biphasic medium, wettability of the nanoparticle surface and effective radius of nanoparticles, as illustrated in the following equation:

$$\Delta E = -\pi(d/2)^2\gamma_{w/o}(1 \pm \cos \theta)^2 \quad (1)$$

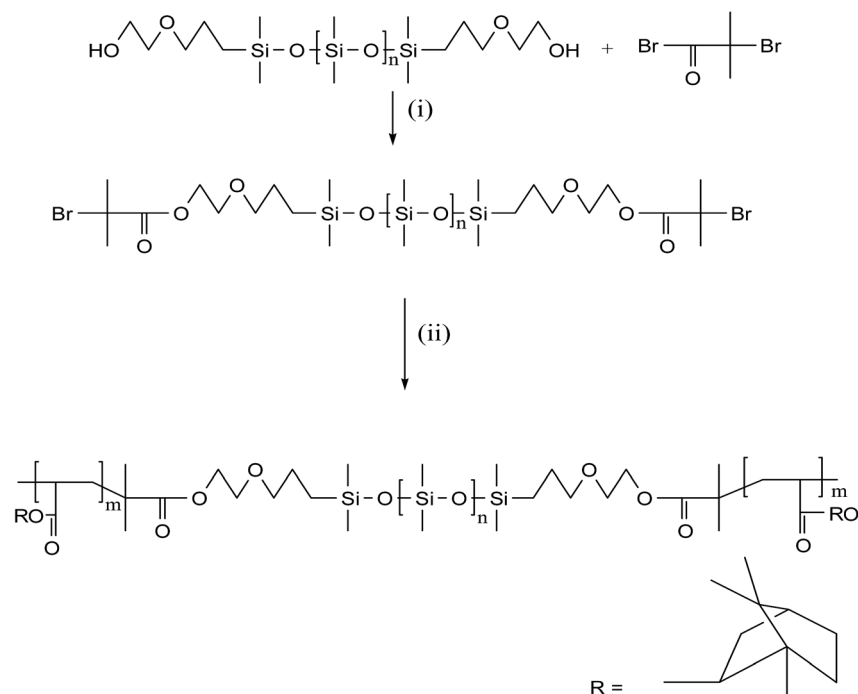
where $\gamma_{w/o}$ is the interfacial tension between water and oil, d the particle diameter, and θ the contact angle of particles at the interface.

The study described herein relates to the formation of a self-assembly of AuNPs at water–chloroform interface and its application as a selective Pickering catalyst for the synthesis of poly(alkyl/arylhydrosiloxane)s, $H_2RSiO[RHSiO]_nSiRH_2$ [$R = Ph$ (2), n -hexyl (3), *cyclo*-hexyl (4), $Et_3SiCH_2CH_2$ (5), $PhMe_2SiCH_2CH_2$ (6)]. The formation of the Pickering catalyst is achieved by surface decoration of AuNPs with amphiphilic block copolymer, $PiBA_{20}$ - b - $PDMS_{75}$ - b - $PiBA_{20}$ ($PiBA$ = polyisobornylacrylate, $PDMS$ = polydimethylsiloxane), **1**.

Results and discussion

The block copolymer, $PiBA_{20}$ - b - $PDMS_{75}$ - b - $PiBA_{20}$ (**1**) used for surface functionalization of AuNPs was synthesized by following atom transfer radical polymerization (ATRP)¹² of isobornylacrylate (iBA) monomer using difunctional macroinitiator, (2-bromoisobutyrate terminal-PDMS), copper bromide (CuBr) and N,N,N',N'',N''' -pentamethyldiethylenetriamine

(PMDETA) as complexing ligand (Scheme 1). The onset of polymerization was indicated by increase in viscosity of the solution. The overlaid SEC traces (Fig. S1†) of the aliquots at different time intervals reveal a progressive shift towards lower elution time (higher molar mass) compared to the PDMS-macroinitiator. An analysis of number average molecular weight (M_n) as a function of percentage conversion of the monomer (obtained from 1H NMR data) shows a linear plot (Fig. S2†), suggesting that the initiation rate is sufficiently fast relative to propagation and termination is maintained at low level. This is further supported by a decrease in the value of polydispersity index from 2.0 to 1.12 as the polymerization progressed. In the 1H NMR spectrum (Fig. S3†) the resonance at δ 1.9 due to terminal $(CH_3)_2CBr$ groups of the difunctional macroinitiator is completely absent, suggesting the efficacy of each individual initiating site during polymerization reaction. The detailed synthetic procedure and spectroscopic characterization of **1** is described in the experimental section. An examination of gel permeation chromatographic data reveals $M_n = 12\,570\text{ g mol}^{-1}$ and $PDI = 1.12$. The assignments of complex 1H NMR signals of **1** are made by comparison with those reported for poly(isobornylacrylate).¹³ The Me_2Si (δ 0.06) and CH -isobornyl (δ 4.25) protons associated with PDMS and $PiBA$ segments appear in the intensity ratio of 1.8 : 1 and is consistent with the composition of the polymer. The ^{29}Si NMR spectrum reveals a single resonance at δ -21.9 analogous to that observed in poly(dimethylsiloxane). The FT-IR spectrum in the solid state exhibits strong peaks at 1726, 1260 and 1089 cm^{-1} due to $C=O$ (ester), $Si-Me$ and $Si-O-Si$ groups respectively. Thermo gravimetric analysis (TGA) and differential scanning



(i) Et_3N , THF, RT, 20 h (ii) iBA, CuBr, PMDETA, Toluene, 80 °C, 4-5 h

Scheme 1 Synthesis of polymer **1**.



calorimetric (DSC) studies reveal that the polymer is thermally stable up to 280 °C and exhibits glass transition temperature (T_g) of PiBA segment at 85 °C. The PDMS segment could not be analysed by DSC due to temperature constraint of the available instrument. The glass transition temperature of PiBA segment in **1** is significantly lowered as compared to that of poly-(isobornylacrylate) homopolymer (94 °C)¹⁴ and the phenomenon likely arises from phase mixing of the PDMS and PiBA segments or configurational effects of the polymer chain.¹⁵

Synthesis of AuNPs-stabilized Pickering interfacial catalyst in water–chloroform biphasic medium

For the synthesis of Pickering interfacial catalyst, the initial step involves the reduction of hydrogen tetrachloroaurate(III) trihydrate in chloroform with triethylsilane¹⁶ under ultrasonication in presence of polymer **1** as the stabilizing matrix. The formation of AuNPs is evident by a gradual colour change of the solution from yellow to purple within 5–10 minutes. The solution was allowed to attain equilibrium for 24 h under ambient conditions and used for further analysis. The UV-Vis spectrum of the colloidal suspension of AuNPs exhibits surface plasmon resonance at 540 nm along with a shoulder at 650 nm (Fig. 1). The observed spectral profile is suggestive of polydispersed nature of AuNPs.¹⁷ The plasmonic band does not show perceptible change even after a month suggesting that AuNPs are well passivated by the polymer matrix. Transmission electron microscopic (TEM) study was performed by depositing one drop of the solution on a carbon-coated copper grid. The TEM image shown in Fig. 1 is typical of that found over the entire grid and reveals the formation of polydispersed AuNPs of 10–15 nm size domain. Under high magnification, the TEM image

identifies well resolved crystal lattice fringes featuring fcc gold (111) plane with a 0.236 nm d spacing between the adjacent lattice planes (Fig. S4†). The corresponding selected area electron diffraction (SAED) pattern confirms their polycrystalline nature (Fig. S5†). In subsequent step, addition of deionized water (1.0 mL) to a stirred colloidal suspension of AuNPs in chloroform (4.0 mL, 0.02 wt% Au) under ambient conditions results in entrapment of the AuNPs at the water–chloroform interface, while the organic layer becomes colourless within 30 minutes (Fig. 2). The interfacially stabilized polymer–AuNP assembly is hereafter referred as [Au]. It is noteworthy to mention that optimization of AuNP concentration (0.1 wt%) and water–chloroform volume ratio (1 : 1) affords the formation of a sheet like structure at the interface (Fig. 2). The AuNPs thus formed exhibit metallic lustre when light is passed into the sample. The HRTEM micrograph of [Au] is shown in Fig. 2. The result reveals an aggregation of individual nanoparticles that are not densely packed and do not show long-range order possibly because of the broad size and shape distribution. The plasmonic resonance appears at $\lambda_{\text{max}} = 540$ nm and extends in the near IR region. The stability of the nanoparticles is not affected by changing the organic medium such as CH_2Cl_2 , THF and toluene. Nevertheless, surface tethering of AuNPs by PiBA homopolymer does not favour the self-assembly process at water–chloroform interface and provides a basis to infer that organization of AuNPs at the interface is primarily driven by amphiphilic character of polymer **1**. The results find analogy with an earlier report by Wang *et al.* The authors have demonstrated that organization of AuNPs at water–toluene interface is primarily governed by surface functionalization of AuNPs with amphiphilic 2-bromo-2-methyl-propionate terminated long chain thiolate groups while the size of nanoparticles does not affect this phenomenon significantly.¹⁰

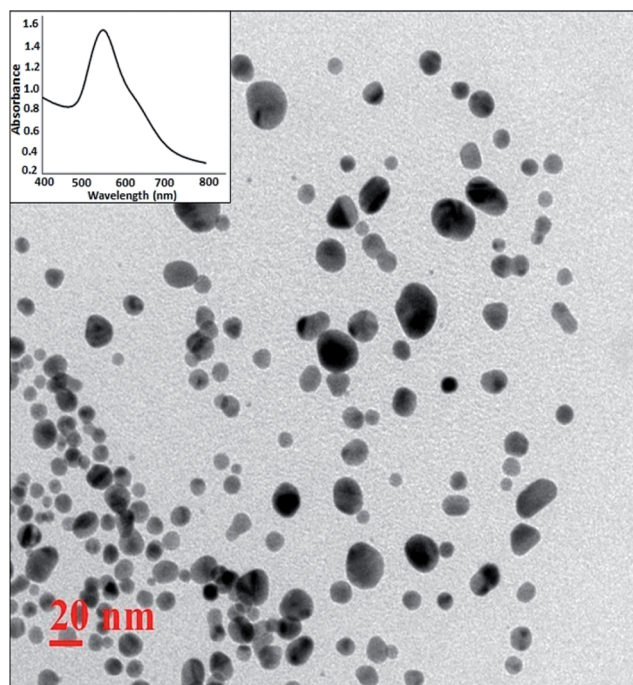


Fig. 1 HRTEM image of AuNPs in chloroform.

Synthesis and characterization of poly(hydrophenylsiloxane), **2** using Pickering interfacial catalyst, [Au]

As a case study, phenylsilane was taken as a model substrate to study the hydrolytic oxidation of the Si–H groups. The reaction was performed with 0.02 wt% of the [Au] catalyst at 80 °C and monitored at different time intervals by ¹H NMR spectroscopy (Fig. 3). As the reaction progresses, a new resonance at δ 5.1 appears *in lieu* of the signal at δ 4.1 associated with phenylsilane. The former value is typical of $\text{PhHSiO}/\text{H}_2\text{PhSiO}$ units¹⁸ suggesting the onset of hydrolytic oxidation of the precursor. The GC-MS spectrum (Fig. S6†) of the sample obtained after 2 h of the reaction identifies linear siloxanes of compositions, $\text{H}_2\text{PhSiOSiPh(H)(OH)}$ (A) $m/z = 245$; $[\text{M} - \text{H}]$, $\text{H}_2\text{PhSiO}[\text{PhHSiO}]_n\text{SiPhH}_2$ (B) $m/z = 351$; $[\text{M} - \text{H}]$ and $\text{H}_2\text{PhSiO}[\text{PhHSiO}]_2\text{SiPhH}_2$ (C) ($m/z = 474$) in 3 : 4 : 1 ratio. The signal due to the Si–H protons of phenylsilane disappears after 9–10 h. A usual work up (see Experimental section) of the reaction mixture affords the isolation of $\text{H}_2\text{PhSiO}[\text{PhHSiO}]_n\text{SiPhH}_2$, **2** as viscous oil with $M_w = 1.5 \times 10^3$ and PDI = 1.4 as evident from gel permeation chromatography. In the ¹H NMR spectrum, the integral intensity of Si–Ph and Si–H signals in 5 : 1 ratio is in conformity with the composition of the



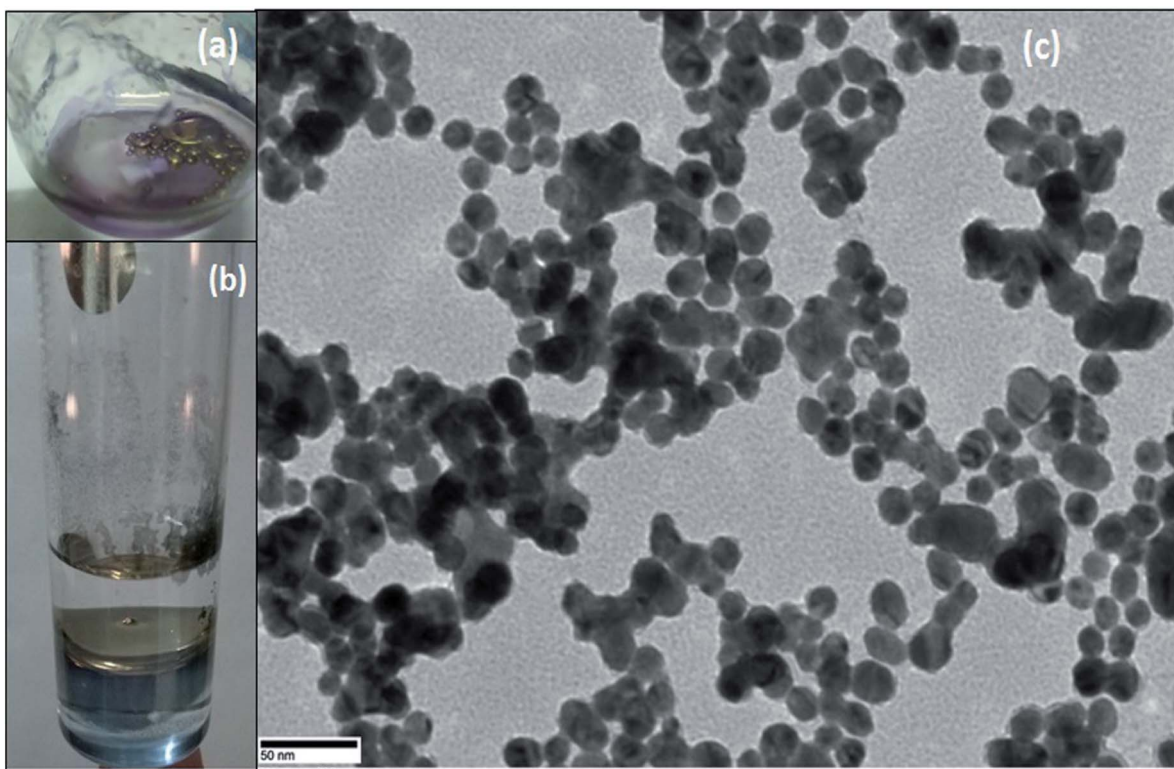


Fig. 2 Images of AuNPs (a) stabilized on water droplets (b) sheet like structure at water–chloroform interface, (c) TEM image of AuNPs obtained from (a).

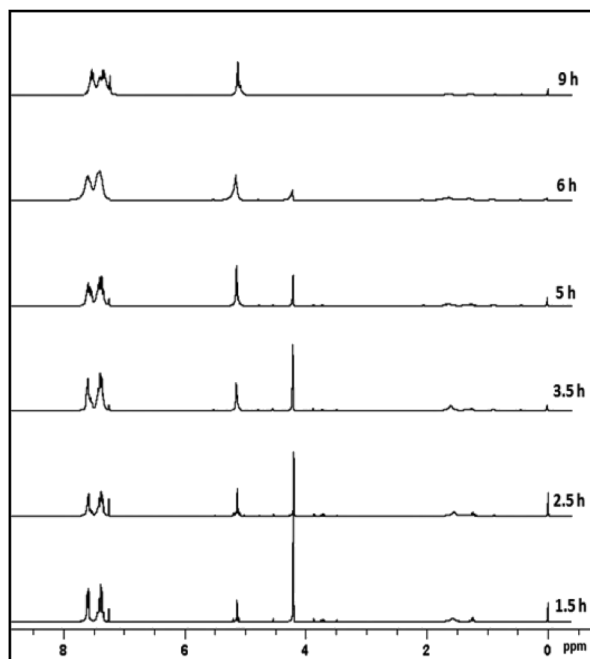


Fig. 3 ^1H NMR spectra at different time intervals during hydrolytic oxidation of phenylsilane.

polymer. The IR spectrum exhibits characteristic absorption at 2173 cm^{-1} due to Si–H groups. The ^{29}Si NMR spectrum (Fig. 4) identifies two distinct regions of chemical shifts at $\delta -44$ to

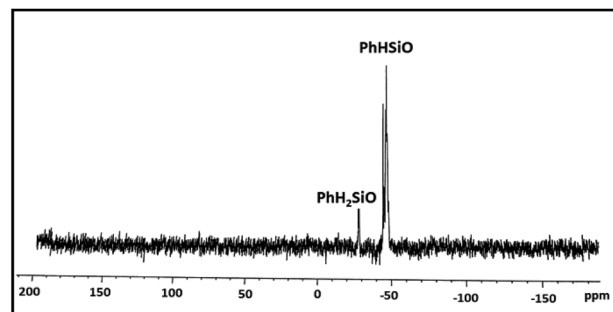


Fig. 4 $^{29}\text{Si}\{^1\text{H}\}$ NMR spectrum of poly(hydrophenylsiloxane), 2.

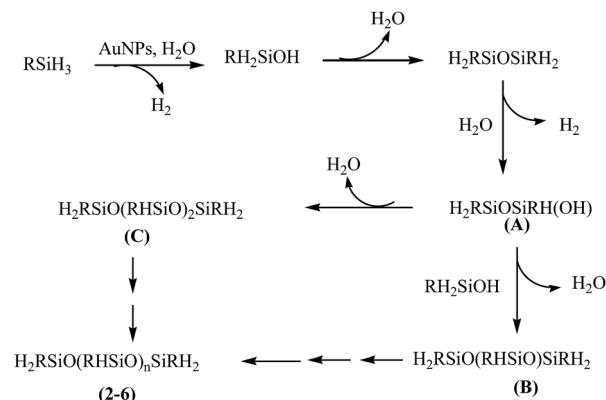
-47 and -27.0 . The former value is assigned to PhHSiO units while the latter to H_2PhSiO end groups. Similar chemical shifts have been reported earlier for $\text{H}_2\text{PhSiOSiPhH}_2$ ($\delta -27$, t , $^1\text{J}_{\text{Si-H}} = 224\text{ Hz}$) and $[\text{PhHSiO}]_n$ ($\delta -46.8$, d , $^1\text{J}_{\text{Si-H}} = 259\text{ Hz}$) which has been prepared by controlled hydrolysis of PhHSiCl_2 .¹⁸ Thermo gravimetric analysis (TGA) and differential scanning calorimetric (DSC) study of 2 reveal that the polymer is thermally stable up to $280\text{ }^\circ\text{C}$ and exhibits the glass transition temperature (T_g) at $-59\text{ }^\circ\text{C}$ (Fig. S7†). Interestingly, T_g is markedly higher than that of poly(hydromethylsiloxane) ($T_g = -138\text{ }^\circ\text{C}$) suggesting a reduced flexibility of the siloxane backbone.¹⁹ It is pertinent to mention that the blank experiments in the absence of $[\text{Au}]$ catalyst and water separately do not proceed at all and the substrate is recovered unchanged after 24 h.



Synthesis and characterization of poly(alkylhydrosiloxane)s, 3–6

To expand the substrate scope, the Pickering catalyst, [Au] has been further examined for the hydrolytic oxidation of primary alkylsilanes, RSiH_3 [$\text{R} = n\text{-hexyl, cyclo-hexyl, Et}_3\text{SiCH}_2\text{CH}_2, \text{PhMe}_2\text{SiCH}_2\text{CH}_2$] under conditions similar to those described above for phenylsilane. An illustrative ^1H NMR spectra showing the progress of the reaction for $n\text{-hexylsilane}$ as the precursor is given in ESI (Fig. S8).† A complete consumption of the organosilane in each case is observed within 10–12 h. The corresponding poly(hydrosiloxane)s, $\text{H}_2\text{RSiO}[\text{RHSiO}]_n\text{SiRH}_2$ 3–6 were isolated as viscous oils and characterized by GPC, IR, ^1H and ^{29}Si NMR spectroscopy. These are thermally stable upto 200 °C (Fig. S9†) and thermal stability decreases in the order: $\text{cyclo-hexyl} > n\text{-hexyl} > \text{Et}_3\text{SiCH}_2\text{CH}_2 \approx \text{PhMe}_2\text{SiCH}_2\text{CH}_2$. The GPC profiles of 3–6 shown in Fig. 5 reveal molecular weight (M_w) in the range of $1.1\text{--}1.8 \times 10^3$ ($\text{PDI} = 1.2\text{--}1.4$) suggesting that the polymerization rate is comparable to that of phenylsilane. The ^1H NMR spectra exhibit characteristic signals at δ 4.4–4.6 and 0.6–1.9 due to Si–H and alkyl groups respectively in 1 : 1 integral ratio suggesting the presence of RHSiO repeat units. Additional evidence with regard to the composition of these polymers comes from characteristic ^{29}Si NMR signals at δ –32 to –37 and –17 to –19 (Fig. S10†) which are attributed to RHSiO and H_2RSiO units respectively by comparison with those reported earlier for $[\text{RHSiO}]_n$ and $\text{H}_2\text{RSiOSiRH}_2$ ($\text{R} = \text{octadecyl}$).²⁰ The IR absorptions due to Si–H, Si–O and aliphatic groups appear at 2140–2160, 1050–1070 and 2900–3000 cm^{-1} respectively.

The efficiency of the Pickering catalyst after the first cycle of the reaction decreases as evident from a longer reaction time (18–24 h) for complete hydrolytic oxidation of organosilanes. To understand this phenomenon, the AuNPs were examined after the first cycle by HRTEM (Fig. S11†). The micrograph reveals that structural reorganization of the primitive nanoparticles occurs during the catalytic event at elevated temperature and



Scheme 2 Proposed catalytic pathway for the synthesis of 2–6.

aggregation of individual nanoparticles is discernable in the TEM image. The plasmonic band appears at 570 nm and undergoes a bathochromic shift of the order of 30 nm as compared to primitive AuNPs. Such modulations in the catalytic cycles have been previously observed in PdNPs during Suzuki coupling reaction between phenyl boronic acid and iodobenzene.²¹

There has been ample precedence in literature which provides a mechanistic rationale for the oxidation of organosilanes using AuNP catalyst. It has been suggested that the reaction proceeds *via* the formation of Au–H and Au–Si bonds and subsequent reaction of these species with water affords the formation of silanol with the elimination of dihydrogen.²² A similar pathway (Scheme 2) has been proposed in the present study to explain the catalytic activity of [Au] and formation of poly(alkyl/arylhydrosiloxane)s 2–6. The identity of A to C formed as intermediates is authenticated by GC–MS spectral studies (*vide supra*). An added evidence in favour of the selectivity of the [Au] catalyst comes from the catalytic oxidation of diphenylsilane which affords $\text{HPh}_2\text{SiOSiPh}_2\text{H}$ (7) within 2 h in >99% yield.

The results provide a basis to infer that the Si–H groups directly bonded to a siloxy unit is far less prone to oxidation under the reaction conditions employed. To our knowledge, there have been only a few reports on the AuNP-catalyzed hydrolytic oxidation of primary organosilanes. In a related study, the alkylsilane, $\text{C}_{18}\text{H}_{37}\text{SiH}_3$ is known to undergo complete oxidation of the Si–H groups and affords the formation of nanowire-like structures build up from $\text{RSiO}_{1.5}$ units.²³

Conclusions

We have demonstrated that surface decoration of AuNPs with amphiphilic block copolymer 1 assists the organization of the nanoparticles at water–chloroform interface. The interfacial stabilization of AuNPs provides a platform to study their catalytic activity and selectivity for hydrolytic oxidation of primary alkyl/aryl silanes. This has resulted in the development of a versatile approach for the synthesis of poly(alkyl/arylhydrosiloxane)s, 2–6 bearing [RHSiO] repeat units. Further studies are in progress to expand the scope of these poly(hydrosiloxane)s to incorporate new functional groups and explore their possible applications.

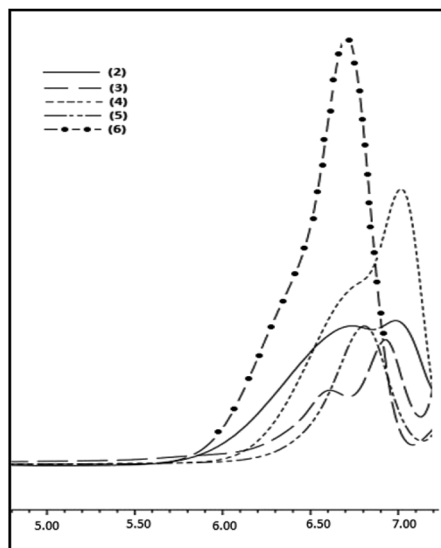


Fig. 5 GPC profiles of the polymers 2–6.



Experimental

Materials and methods

Solvents were freshly distilled over sodium wire/benzophenone (tetrahydrofuran, toluene) under inert atmosphere. Chloroform (Fischer, HPLC grade) was used as received. Glassware were dried in an oven at 110–120 °C and further flame dried under vacuum prior to use. Triethylamine was dried over potassium hydroxide and distilled before use. Copper(i) bromide (Sigma Aldrich) was stirred in glacial acetic acid overnight, filtered and washed with absolute ethanol under nitrogen atmosphere. Isobornyl acrylate (iBA) was passed through alumina column to remove the inhibitor. The polymer, bis(hydroxyethoxypropyl)-terminated PDMS ($M_n = 5600 \text{ g mol}^{-1}$), 2-bromoisobutyl bromide and N,N,N',N'',N'' -pentamethyldiethylenetriamine (PMDETA), phenylsilane, n -hexylsilane, *cyclo*-hexylsilane, triethylsilane and lithium aluminium hydride were purchased from Sigma Aldrich and used as procured. ^1H , ^{13}C and ^{29}Si NMR spectra were recorded in CDCl_3 on Bruker AVANCE II 400 NMR spectrometer and the chemical shifts are quoted relative to Me_4Si . IR spectra were recorded on Nicolet protege 460 ESP spectrometer using KBr optics. UV-Vis spectral studies were performed on Perkin-Elmer (Lambda Bio 20) spectrophotometer. Transmission electron microscopic (TEM) studies were carried out using carbon coated copper grid on a Philips CM 20 electron microscope operated at 100 kV. Molecular weight (M_w) and polydispersity index (PDI) of the polymers were estimated using Waters gel permeation chromatography (GPC) equipped with L-2414 refractive index detector and Waters styragel HR3 and HR4 columns in series using THF as eluent (flow rate 1 mL min^{-1} ; polystyrene standards). Thermo gravimetric analysis (TGA) and differential scanning calorimetry (DSC) measurements were carried out in nitrogen atmosphere at a heating rate $10 \text{ }^\circ\text{C min}^{-1}$ on a Perkin Elmer Thermal analysis system.

Synthetic methods

The organosilanes, RSiH_3 ($\text{R} = \text{Et}_3\text{SiCH}_2\text{CH}_2$, $\text{PhMe}_2\text{SiCH}_2\text{CH}_2$) were synthesized according to the procedure reported earlier.²⁴ The difunctional macroinitiator (Br-PDMS-Br) was prepared by following the procedure reported earlier,²⁵ using bis(hydroxyethoxypropyl) terminated PDMS (5.0 g, 1.0 mmol), triethylamine (0.63 g, 6.0 mmol) and 2-bromoisobutyl bromide (0.69 g, 3.0 mmol) as the starting materials. The IR, ^1H , ^{13}C and ^{29}Si NMR data of the macroinitiator are summarized below.

^1H NMR (CDCl_3): δ 0.07 (br, SiCH_3), 0.57 (br, SiCH_2), 1.57 (br, SiCH_2CH_2), 1.95 [br, $\text{BrC}(\text{CH}_3)_2$], 3.45 (br, $\text{SiCH}_2\text{CH}_2\text{CH}_2$), 3.67 (br, OCH_2), 4.32 (br, $\text{CH}_2\text{OC}=\text{O}$); $^{13}\text{C}\{^1\text{H}\}$ NMR (CDCl_3): δ 1.0 (SiCH_3), 14.1 (SiCH_2), 23.4 (SiCH_2CH_2), 30.7 [$\text{BrC}(\text{CH}_3)_2$], 65.2 ($\text{SiCH}_2\text{CH}_2\text{CH}_2$), 68.1 (OCH_2), 74.1 ($\text{CH}_2\text{OC}=\text{O}$), 175.2 ($\text{C}=\text{O}$); ^{29}Si NMR (CDCl_3 , δ): -21.9. IR (KBr, cm^{-1}): 1089 ($\nu_{\text{Si-O-Si}}$), 1260 ($\nu_{\text{Si-CH}_3}$), 1742 ($\nu_{\text{C=O}}$).

Synthesis of PiBA_{20} -*b*-PDMS₇₅-*b*-PiBA₂₀, 1

In a typical procedure, the macroinitiator, Br-PDMS-Br (1.0 g, 0.17 mmol), iBA (3.4 g, 17.0 mmol) and CuBr (0.048 g, 0.34 mmol) in dry toluene (2.0 mL) were introduced into a two-neck

round bottom flask equipped with a magnetic stirrer. The contents were subjected to three freeze-pump-thaw cycles to remove oxygen and N,N,N',N'',N'' -pentamethyldiethylenetriamine (PMDETA) (0.12 g, 0.68 mmol) was added while keeping the reaction mixture at 80 °C. The reaction was allowed to continue for 4–5 h and the content thereafter was dissolved in THF and passed through neutral alumina column to remove the copper catalyst. The polymer 1 was isolated as a white solid by repeated precipitation in methanol. The numerical notations for the assignment of various protons are shown in Scheme 1.

^1H NMR (CDCl_3): δ 0.06 (H1 , Si-CH_3), 0.57 (H_2 , SiCH_2), 0.80–1.70 [H10-H11 , H13-H20 (*isobornyl* ring) + H3 , SiCH_2CH_2], 2.27 [H9 , $\text{C}(\text{O})\text{CH}$], 3.45 (H4 , $\text{SiCH}_2\text{CH}_2\text{CH}_2$), 3.67 (H5 , OCH_2), 4.25 (H6 , $\text{CH}_2\text{OC}=\text{O}$), 4.32 (H12 , CH-isobornyl). IR (KBr, cm^{-1}) 1089 ($\nu_{\text{Si-O-Si}}$), 1260 ($\nu_{\text{Si-CH}_3}$), 1727 ($\nu_{\text{C=O}}$). ^{29}Si NMR (CDCl_3): δ -21.9.

Synthesis of Pickering interfacial catalyst (PIC)

To a solution containing hydrogen tetrachloroaurate(III) trihydrate (10.0 mg, 0.025 mmol) in chloroform (20.0 mL, HPLC), a solution of polymer 1, (10.0 mg, 0.04 mmol with respect to polymer repeat unit) in the same solvent (5.0 mL) was added under sonication. Thereafter, triethylsilane (10 μL , 0.1 mmol) was added and the resulting solution was kept for 24 h at room temperature and subsequently used for analysis. The above solution of AuNPs (4.0 mL) was taken separately and mixed with 1.0 mL of deionized water with constant stirring to obtain AuNP-stabilized water droplets in chloroform medium and used for catalytic reactions.

Synthesis of poly(aryl/alkyl)hydrosiloxanes, 2–6

In a typical procedure, hydrolytic oxidation of primary organosilanes, RSiH_3 (2.0 mL, 0.02 mmol) was performed in the presence of Au-stabilized PIC under aerobic conditions at 80 °C. After complete consumption of the monomer, the organic layer was separated and dried over sodium sulphate. The solvent was evaporated under vacuum and poly(hydrosiloxane)s, 2–6 were isolated as viscous oils.

$\text{H}_2\text{PhSiO}[\text{PhHSiO}]_n\text{SiPhH}_2$, 2. ^1H NMR (CDCl_3): δ 7.24 (br, 5H, SiPh), 5.14 (br, 1H, Si-H); $^{29}\text{Si}\{^1\text{H}\}$ NMR (CDCl_3): δ -27.0 [H_2PhSiO], -44 to -47 (HPhSiO); IR (KBr, cm^{-1}): 3070, 3052 ($\nu_{\text{C-H}}$, aromatic), 2173 ($\nu_{\text{Si-H}}$), 1429 ($\delta_{\text{C-H}}$), 1085 ($\nu_{\text{Si-O-Si}}$). $M_w = 1.5 \times 10^3$, PDI = 1.4.

$\text{H}_2(n\text{-hexyl})\text{SiO}[(n\text{-hexyl})\text{HSiO}]_n\text{Si}(n\text{-hexyl})\text{H}_2$, 3. ^1H NMR (CDCl_3): δ 0.54 (br, 2H, Si-CH_2), 0.67–0.79 (br, 4H, CH_2), 1.2 (br, 9H), 4.56 (br, 1H, Si-H); $^{29}\text{Si}\{^1\text{H}\}$ NMR (CDCl_3): δ -20.0 (H_2SiO), -33.2, -34.71, -35.89 (HSiO); IR (KBr cm^{-1}): 2957, 2924 ($\nu_{\text{C-H}}$, aliphatic), 2160 ($\nu_{\text{Si-H}}$), 1461 ($\delta_{\text{C-H}}$), 1089 ($\nu_{\text{Si-O-Si}}$). $M_w = 1.2 \times 10^3$, PDI = 1.2.

$\text{H}_2(\text{cyclo-hexyl})\text{SiO}[(\text{cyclo-hexyl})(\text{H})\text{SiO}]_n\text{Si}(\text{cyclo-hexyl})\text{H}_2$, 4. ^1H NMR (CDCl_3): δ 0.54 (br, 1H, Si-CH), 1.37 (br, 4H, CH_2), 1.9 (br, 6H), 4.42 (1H, Si-H); $^{29}\text{Si}\{^1\text{H}\}$ NMR (CDCl_3): δ -17.6 (H_2SiO), -33.17, -35.92, -37.09 (HSiO); IR (KBr cm^{-1}): 2957, 2924 ($\nu_{\text{C-H}}$, aliphatic), 2160 ($\nu_{\text{Si-H}}$), 1461 ($\delta_{\text{C-H}}$), 1085 ($\nu_{\text{Si-O-Si}}$). $M_w = 1.1 \times 10^3$, PDI = 1.4.

$\text{H}_2\text{RSiO}[\text{RHSiO}]_n\text{SiRH}_2$ ($\text{R} = \text{Et}_3\text{SiCH}_2\text{CH}_2$), 5. ^1H NMR (CDCl_3): δ 0.54 (br, 8H, SiCH_2), 1.12 (br, 11H, SiCH_2CH_2 , CH_3),



4.66 (Si-H); $^{29}\text{Si}\{^1\text{H}\}$ NMR: δ 8.63, 8.40 $\{(\text{C}_2\text{H}_5)_3\text{Si}\}$, -19.2 (H_2SiO), -32.2, -34.5 (HSiO); IR (KBr, cm^{-1}): 2953, 2908 ($\nu_{\text{C-H}}$, aromatic), 2140 ($\nu_{\text{Si-H}}$), 1414 ($\delta_{\text{C-H}}$), 1075 ($\nu_{\text{Si-O-Si}}$). $M_w = 1.1 \times 10^3$, PDI = 1.1.

$\text{H}_2\text{RSiO}[\text{RHSiO}]_n\text{SiRH}_2$ ($\text{R} = \text{PhMe}_2\text{SiCH}_2\text{CH}_2$), **6**. ^1H NMR (CDCl_3): δ 0.50–0.64 (br, 8H, SiMe_2 , SiCH_2), 1.02 (br, 2H, SiCH_2CH_2), 4.58 (br, Si-H), 7.24 (br, 5H, SiPh); $^{29}\text{Si}\{^1\text{H}\}$ NMR (CDCl_3): δ -5.36 (Ph_2MeSi), -34.96 (HSiO); IR (KBr, cm^{-1}): 3070, 3052 ($\nu_{\text{C-H}}$, aromatic), 2150 ($\nu_{\text{Si-H}}$), 1421 ($\delta_{\text{C-H}}$), 1081 ($\nu_{\text{Si-O-Si}}$). $M_w = 1.8 \times 10^3$, PDI = 1.3.

$[\text{Ph}_2(\text{H})\text{Si}]_2\text{O}$ [product from Ph_2SiH_2], **7**. ^1H NMR (CDCl_3 , δ): 7.63, 7.31, (m, 20H, Si-Ph), 5.59 (s, 2H, Si-H); $^{29}\text{Si}\{^1\text{H}\}$ NMR (CDCl_3 , δ): -19.8. IR (KBr, cm^{-1}): 3062, 3011 ($\nu_{\text{C-H}}$, aromatic), 2135 ($\nu_{\text{Si-H}}$), 1428 ($\delta_{\text{C-H}}$), 1067 ($\nu_{\text{Si-O-Si}}$); GC-MS: retention time (t_R) = 31.5 min, m/z calcd, 382; found, 382.

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