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Polymer grafting on nitrene functionalized green silica *via* “grafting from” and “grafting to” approaches through enhanced spin capturing polymerization and a 1,3-dipolar cycloaddition reaction†

Lukkumanul Hakkim N.  and Leena Nebhani *

The nitrene functionality is proven to have the potential to enable polymer conjugation *via* enhanced spin-capturing polymerization (ESCP) and nitrene-mediated radical coupling (NMRC); however, reactions using nitrene have been less explored for surface functionalization of solid substrates. In addition, nitrene has a unique ability to undergo a 1,3-dipolar cycloaddition reaction which has not been explored for polymer conjugation and polymer “grafting from” solid substrates. In this work, nitrene-functionalized silica derived from rice husk ash has been used for polymer grafting *via* “grafting from” and “grafting to” through spin capturing and 1,3-dipolar cycloaddition reactions. The ability of the nitrene functionality to undergo 1,3-dipolar cycloaddition was monitored using styrene and nitrene, nitrene-functionalized silica and isobornyl acrylate, as well as nitrene-functionalized polystyrene and a polystyrene macromonomer. The nitrene functionality was introduced onto the mercaptopropyl functionalized silica surfaces synthesized *via* a co-condensation reaction. The extent of polymer grafting *via* ESCP has been analyzed at three different conversions. The effect of the surface area of silica and the method of functionalization on polymer grafting density has been studied. The efficiency of polymer grafting over the nitrene functionality on the surface indicates the potential of surface nitrene functionalities to undergo polymer grafting *via* “grafting from” and “grafting to” approaches.

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1. Introduction

Polymer grafting on silica can be performed by chemical or physical methods. The chemical covalent grafting can be *via* either a “grafting to” or a “grafting from” approach. “Grafting to” involves the pre-synthesis of polymers followed by attaching to the silica surface, whereas in the “grafting from” approach, the polymerization starts from the surface of the silica.¹ The “grafting from” approach of polymer grafting has been reported over silica surfaces *via* conventional free radical polymerization,² atom transfer radical polymerization (ATRP),^{3–6} reversible addition–fragmentation chain transfer polymerization (RAFT),^{7–10} and nitroxide-mediated polymerization (NMP),^{11–14} and the “grafting to” approach has been reported using click reactions, for example, the nitroxide mediated radical coupling (NRC)¹⁵ reaction and the nitrene mediated radical coupling

reaction (NMRC).¹⁶ The grafting of suitable silanes over silica surfaces capable of polymerization can be carried out *via* co-condensation or post-modification. The co-condensation method^{17–22} is a one-step method of synthesis of functionalized silica from its precursor, whereas the post-modification^{23–29} method is a two-step process that includes the synthesis of silica followed by functionalization.

Nitrene is a reactive functionality that contains *N*-oxide of an imine, and can be synthesized *via* the oxidation of secondary amine, imine, and hydroxylamine, and the condensation of aldehyde and hydroxylamine.³⁰ The nitrene functionality has been proven to be able to graft polymer chains *via* enhanced spin capturing polymerization (ESCP)³¹ and nitrene mediated radical coupling (NMRC),^{16,32} and undergo 1,3 dipolar cycloaddition with the alkene functionality. The ESCP method of polymer grafting is a “grafting from” approach and the NMRC and 1,3-dipolar cycloaddition methods are “grafting to” approaches of synthesis. The ESCP and NMRC methods can yield the grafting of two growing polymer chains per nitrene functionality, whereas the 1,3-dipolar cycloaddition method can yield the grafting of single polymer chains per

Department of Materials Science and Engineering, Indian Institute of Technology Delhi, Hauz Khas, New Delhi 110016, India. E-mail: Leena.Nebhani@mse.iitd.ac.in
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nitron functionality. The alkoxyamine group formed *via* polymer grafting can undergo nitroxide-mediated polymerization at a higher temperature. The introduction of the nitron functionality onto a silica surface can make the silica capable of having polymers grafted *via* ESCP, NMRC, and 1,3-dipolar cycloaddition.³³

In this study, we have introduced the nitron functionality onto mercaptopropyl functionalized silica samples synthesized *via* a thiol-ene reaction. The mercaptopropyl functionalized silica samples were synthesized *via* co-condensation from sodium silicate derived from RHA. The nitron-functionalized silica samples were subjected to polystyrene grafting studies through “grafting from” and “grafting to” approaches *via* ESCP and 1,3-dipolar cycloaddition reactions. The ESCP was carried out using a styrene monomer, and 1,3-dipolar cycloaddition was performed using a polystyrene macromonomer synthesized *via* HBr elimination of a polystyrene macroinitiator. The extent of polymer grafting using ESCP has been analyzed at three different conversions, and the effect of the surface area and method of functionalization on polymer grafting has been studied. The synthesized nitron-functionalized samples were observed to be efficient in the grafting of the polymer chains *via* “grafting from” and “grafting to” approaches.

2. Experimental methods

2.1. Synthesis of nitron functionalized silica particles

5 g of mercaptopropyl functionalized silica (for the method of synthesis refer to ESI section S1†) was dispersed in 50 ml of 1 : 1 methanol and THF solvent mixture. 0.01 equivalent of triphenylphosphine and 4-(methacryloyloxy)benzaldehyde phenylnitron (same equivalent as equivalent of 3-mercaptopropyl triethoxy silane used in the preparation of mercaptopropyl functionalized silica, for the method of synthesis refer to ESI sections S2 and S3†) were added. The reaction mixture was stirred for 72 h at 40 °C. The nitron functionalized samples were centrifuged and washed with THF to remove triphenylphosphine and unreacted 4-(methacryloyloxy)benzaldehyde phenylnitron. The nitron functionalized silica particles were dried under vacuum at 40 °C and characterized using FTIR

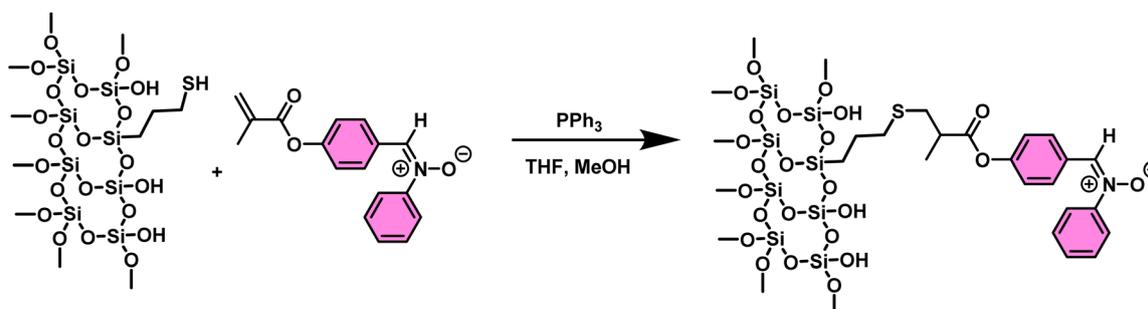
spectroscopy, TGA, ¹³C solid-state NMR spectroscopy, solid-state UV-Visible spectroscopy, and XPS analysis.

2.2. Polymer grafting studies on nitron functionalized silica particles *via* enhanced spin capturing polymerization

Enhanced spin capturing polymerization studies were carried out on nitron functionalized silica samples using the styrene monomer along with the AIBN initiator at 60 °C.³¹ The concentration of AIBN was set to 4×10^{-2} mol L⁻¹ and the surface nitron to monomer ratio was maintained at 1 : 3000 yielding the AIBN to surface nitron to monomer ratio of 1 : 0.07 : 219. The nitron-to-monomer ratio was set to obtain an optimum surface-to-volume ratio in order to reduce silica aggregation. 250 mg of nitron functionalized silica was added to styrene along with the AIBN initiator. The reaction mixture was bath sonicated for 5 minutes, purged under nitrogen, and stirred at 60 °C for the required time. The conversion was monitored using ¹H NMR spectroscopy. The reaction mixture was immediately centrifuged and washed using chloroform after quenching. The free polymers from the supernatant were precipitated using cold methanol for SEC analysis. The polymer grafted samples were repeatedly washed with chloroform until no cloudiness of the supernatant in cold methanol was observed. The polymer grafted samples were dried at 50 °C under vacuum and characterized using FTIR spectroscopy and TGA analysis. The molecular weight of the grafted samples was determined using SEC *via* cleaving the polymers from the surface by transesterification using *p*-toluene sulfonic acid and methanol in toluene.

2.3. Polymer grafting studies on nitron functionalized silica particles *via* 1,3-dipolar cycloaddition of polystyrene macromonomers

A series of model 1,3-dipolar cycloaddition reactions were carried out to confirm the ability of nitron functionalities to form cycloadducts both in solution and on the surface (refer to ESI section S9† for the methods of model 1,3-dipolar cycloadditions). 100 mg of nitron functionalized silica was dispersed in 3 ml of toluene *via* bath sonication. 0.25 mmol of polystyrene macromonomers (refer to ESI section S8† for the method of synthesis) was introduced into the reaction mixture.



Scheme 1 Synthesis of nitron functionalized (NRHA) silica by a thiol-ene reaction.

The reaction mixture was purged with nitrogen for 15 minutes and stirred at 80 °C for 36 h. The reaction mixture was cooled down, centrifuged, and washed repeatedly using toluene, chloroform, and dried under vacuum. The polymer grafted samples were characterized using FTIR, TGA and XPS analysis.

3. Results and discussion

3.1. Synthesis of nitrone functionalized silica particles

As shown in Scheme 1, nitrone functionalized silica was synthesized *via* a thiol-ene reaction between *N*-(4-methacryloyloxy-

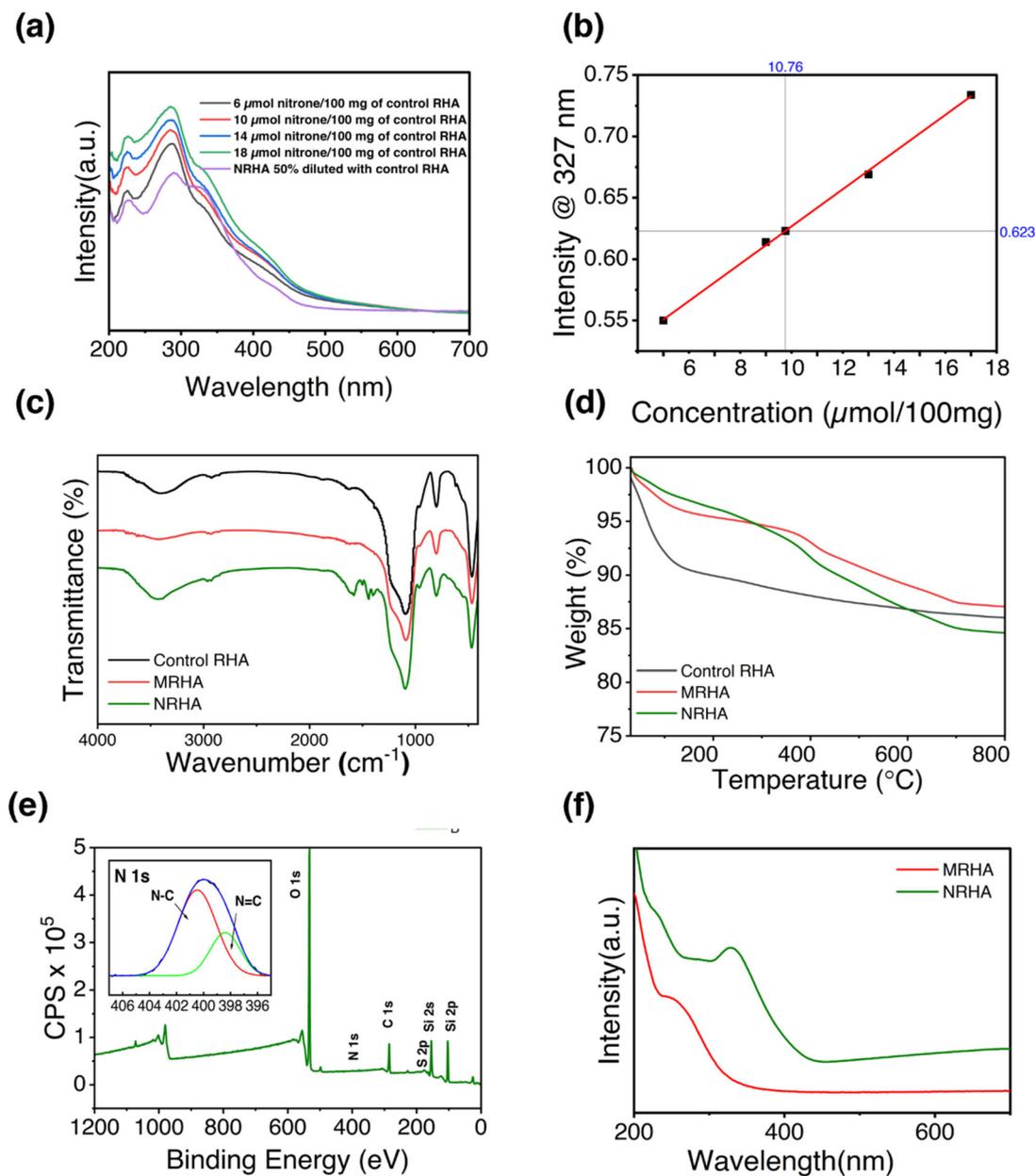
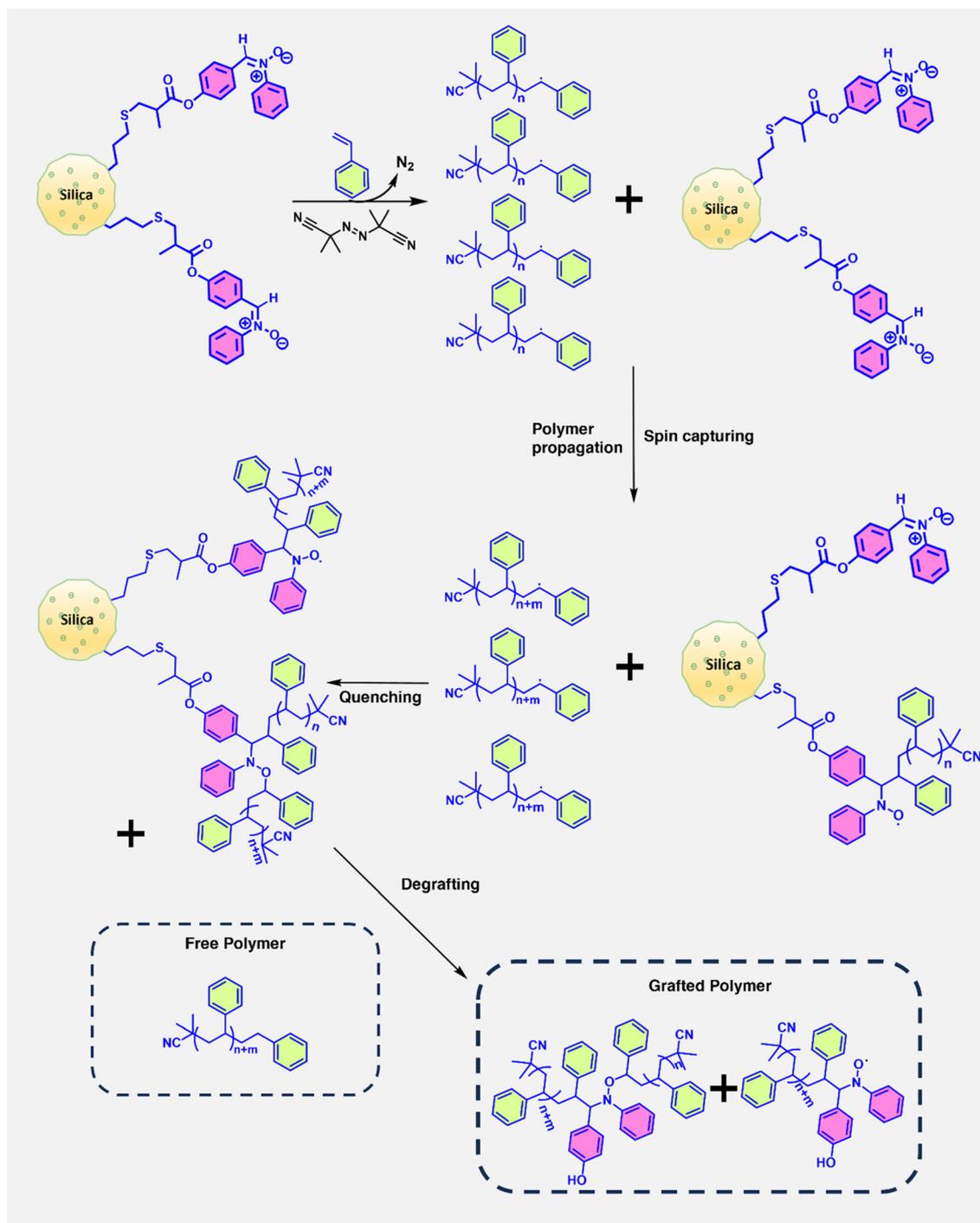


Fig. 1 Solid-state UV-visible spectra of nitrone functionalized RHA silica samples with different nitrone concentrations. The different nitrone concentrations were prepared by mixing 6 μmol, 10 μmol, 14 μmol and 18 μmol of 4-(methacryloyloxy)benzaldehyde phenylnitron in 100 mg of control RHA silica. The nitrone functionalized RHA (NRHA) silica was also diluted to 50% using control RHA silica to fit inside the prepared nitrone concentration range. (a) The solid-state UV-visible spectra for all the concentrations prepared and (b) linear fitting of concentrations with intensity at $\lambda = 327$ nm for the grafting density determination. The values obtained were μmol per 100 mg of samples, which later were converted to $\mu\text{mol g}^{-1}$. (c) FTIR spectra of control RHA silica, mercaptopropyl functionalized RHA (MRHA) silica and nitrone functionalized RHA (NRHA) silica and (d) TGA of control RHA silica, mercaptopropyl functionalized RHA (MRHA) silica and nitrone functionalized RHA (NRHA) silica. (e) XPS wide scan and N 1s narrow scan for nitrone functionalized RHA (NRHA) silica. (f) UV-visible spectra of mercaptopropyl functionalized RHA (MRHA) silica and nitrone functionalized RHA (NRHA) silica, and the peak observed at 327 nm confirms the presence of nitrone functionalities in NRHA silica.

benzylidene)aniline-*N*-oxide (refer to ESI section S3† for synthesis) and mercaptopropyl functionalized silica particles. The synthesized nitron functionalized silica samples were characterized using FTIR analysis, and the band at 1442 cm^{-1} corres-

ponds to alkyl C-H bending vibration (Fig. 1) and the band at 2925 cm^{-1} corresponds to C-H stretching confirming the organic functionalization. The thermogravimetric analysis of nitron functionalized silica shows an increase in weight loss



Scheme 2 Schematic representation of the surface nitron functionality in enhanced spin capturing polymerization. The polymer chains can be grafted at different propagation points, and the remaining polymers in the solution continue to propagate until quenching. The maximum spin capturing of polymer radicals over the surface was observed at the time of quenching, especially at higher conversion, and they were cleaved using the transesterification reaction for SEC analysis. The free polymers in the solution were quenched *via* regular termination mechanisms of free radical polymerization and were collected immediately *via* centrifugation and precipitation of the supernatant in cold methanol for SEC analysis.

compared to its mercaptopropyl functionalized silica precursor; however, the formation of char from 4-(methacryloyloxy)benzaldehyde phenylnitron limits the accuracy in calculating nitron grafting density and hence, the nitron grafting density was calculated from the calibration of solid-state UV-visible spectroscopy analysis (Fig. 1). Different concentrations of nitron functionalities were prepared by mixing $6 \mu\text{mol g}^{-1}$, $10 \mu\text{mol g}^{-1}$, $14 \mu\text{mol g}^{-1}$ and $18 \mu\text{mol g}^{-1}$ *N*-(4-methacryloyloxybenzylidene)aniline-*N*-oxide in 100 mg of non-functionalized silica precipitated from rice husk ash (control RHA silica). The nitron functionalized silica samples were further diluted to 50% using control RHA silica to fit inside the calibrated region. The nitron grafting density obtained for nitron functionalized RHA silica was $215 \mu\text{mol g}^{-1}$. The X-ray photoelectron spectroscopy (XPS) of nitron functionalized silica samples provides the peaks corresponding to Si 2p, Si 2s, S 2p, C 1s, N 1s and O 1s (Fig. 1), and the binding energy observed for N 1s was deconvoluted and the peak corresponding to C=N was observed in the range of 398 eV along with the peaks corresponding to C-N, confirming the presence of the nitron functionality over the silica surface.

3.2. “Grafting from” approach for the synthesis of polystyrene grafted silica via enhanced spin capturing polymerization

Enhanced spin capturing polymerization using the styrene was performed to determine the efficiency of surface nitron radicals in capturing the growing polymer chain (Scheme 2). To determine the ultimate grafting density, maximum polymer radicals need to be initiated in the system, and to achieve the maximum active polymer radicals, a very high concentration of the initiator ($4 \times 10^{-2} \text{ mol L}^{-1}$) to styrene was used, which is 6.5 mg of AIBN per ml of styrene. This rapidly initiates the polymerization and results in a solution having a very high concentration of growing polymer radicals with a similar molecular weight range, resulting in a lower dispersity. A low molecular weight tail is also observed in SEC traces corresponding to the later activated initiators (Fig. 2). The intensity of the low molecular weight tail in SEC traces was observed to decrease with an increase in conversion due to the propagation of later activated low molecular weight polymer chains into the range of initially activated chains, further decreasing the dispersity.

The polystyrene grafted rice husk ash silica via enhanced spin capturing polymerization (PSRHA) was synthesized at three different conversions. The conversions obtained were 25%, 68% and 80% as determined using ^1H NMR spectroscopy (Fig. 3). The qualitative confirmation of grafted polystyrene on nitron functionalized RHA was made using FTIR analysis, and bands corresponding to the grafted polystyrene were observed to increase with an increase in conversion (refer to ESI section S4† for a discussion). The quantitative determination of polystyrene grafting was made using TGA and the organic weight loss for the grafted polystyrene was observed to increase with an increase in conversion (Fig. 3). 9.85%, 22.94% and 34.68% organic fraction weight losses corresponding to grafted polystyrene were observed for the conversions at 25%, 68% and 80% polymerization, respectively.

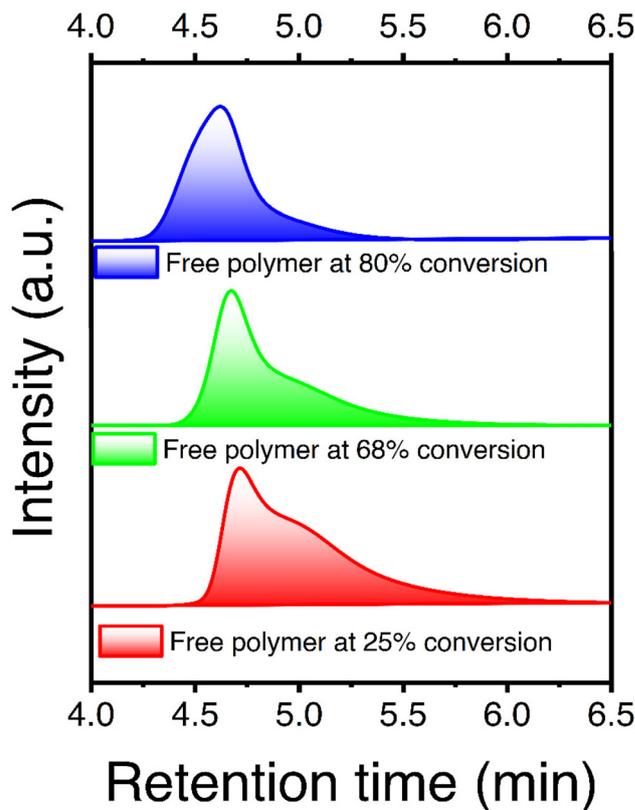


Fig. 2 SEC traces of free polymers at 25%, 68%, and 80% conversion. The very high AIBN concentration provides rapid initiation resulting in a very high concentration of a high molecular weight polymer chain followed by a low molecular weight tail corresponding to the later activated initiators. The molecular weight was observed to increase with an increase in conversion along with the lowering of the intensity corresponding to low molecular weight polymer concentration, which is due to the propagation of later activated low molecular weight polymer chains into the range of initially activated chains.

The SEC traces of free polymers from the solution and the cleaved grafted polymers from polystyrene grafted nitron functionalized RHA (PSRHA) (Fig. 4) for three different conversions are given (Table 1). Since the polymer initiation and propagation were independently carried out in the solution without the role of the surface nitron functionality, the maximum grafting was observed at the time of quenching, resulting in a significant population of grafted polymers having the molecular weight of free polymers.

The SEC trace of grafted polymers observed at a lower conversion of 25% indicates two different populations of molecular weights 3000 g mol^{-1} and 85000 g mol^{-1} due to the grafting of growing polymer chains at different propagation points at the time of quenching, eq (7) and (8) in Scheme 3. At higher conversions of 68% and 80%, the maximum molecular weight observed for grafted polymer chains corresponds to the free polymer molecular weight at quenching due to the restricted mobility and steric hindrance of growing polymer radicals, eq (4) in Scheme 3.

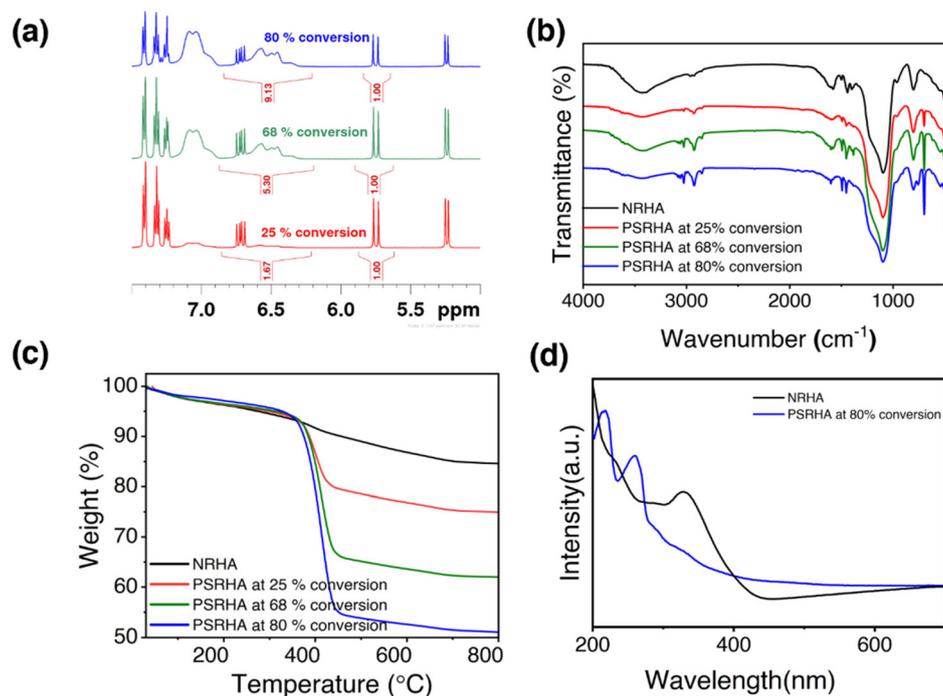


Fig. 3 (a) Conversion of three different reactions was determined using ^1H NMR spectroscopy with respect to the area under the vinylic proton of styrene observed at $\delta = 5.7$ ppm to the broad peak corresponding to the aromatic proton of polystyrene observed at $\delta = 6.5$ ppm. The conversions obtained are 25%, 68% and 80%. (b) FTIR spectra of polystyrene grafted on nitrene functionalized RHA (NRHA) silica using ESCP (PSRHA) at 25%, 68% and 80% conversions. The intensity of the bands corresponding to the aromatic C–H stretching at 3022 cm^{-1} and out of plane C–H bending at 694 cm^{-1} was observed to increase with an increase of polystyrene grafting. (c) TGA plots for polystyrene grafted nitrene functionalized RHA (NRHA) silica synthesized using ESCP (PSRHA) at 25%, 68%, and 80% conversion and the weight loss corresponding to the grafted polystyrene was observed to increase with an increase in conversion, which corresponds to the increased polystyrene grafting. (d) Solid-state UV-visible spectra of nitrene functionalized RHA (NRHA) silica and polystyrene grafted silica synthesized using ESCP (PSRHA) at 80% conversion. The band corresponding to the nitrene functionality was observed to decrease upon polymer grafting.

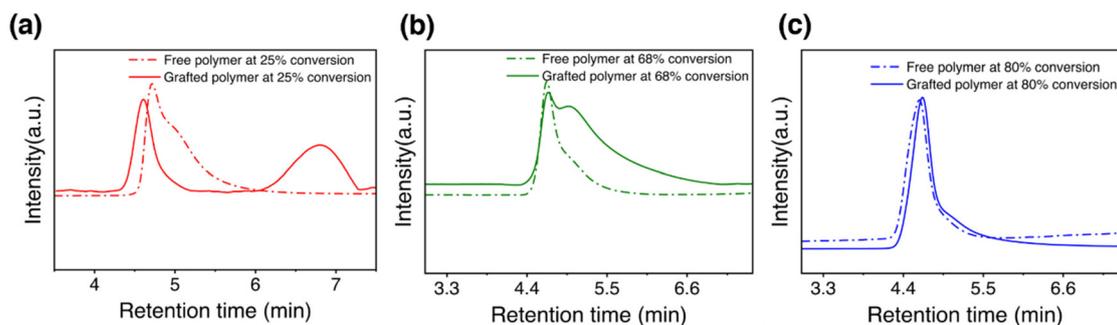


Fig. 4 SEC traces for free polymers from solution (FP) and grafted polymers (GP) from polystyrene grafted silica synthesized *via* ECPS (PSRHA) at (a) 25% conversion, (b) 68% conversion, and (c) 80% conversion. The free polymers were obtained by precipitating the decant in cold methanol after centrifugation and the grafted polymers were cleaved by transesterification of polystyrene grafted silica synthesized *via* ESCP using *p*-toluenesulfonic acid and methanol in toluene. The broader dispersity and multiple populations of grafted polystyrene than that of the single population of free polymers was observed due to the ability of surface nitrene functionalities to capture and terminate the growing polymer radicals at any stage of propagation. The molecular weight of grafted polystyrene observed at 25% conversion was $85\,000\text{ g mol}^{-1}$, with respect to the $38\,000\text{ g mol}^{-1}$ of free polymers, which confirms the potential of nitrene functionalities to capture two polymer radicals.

The absence of free nitrene functionalities in the solution results in free radical polymerization kinetics in the solution and radical capturing at different degrees of polymerization on the surface. This ultimately results in

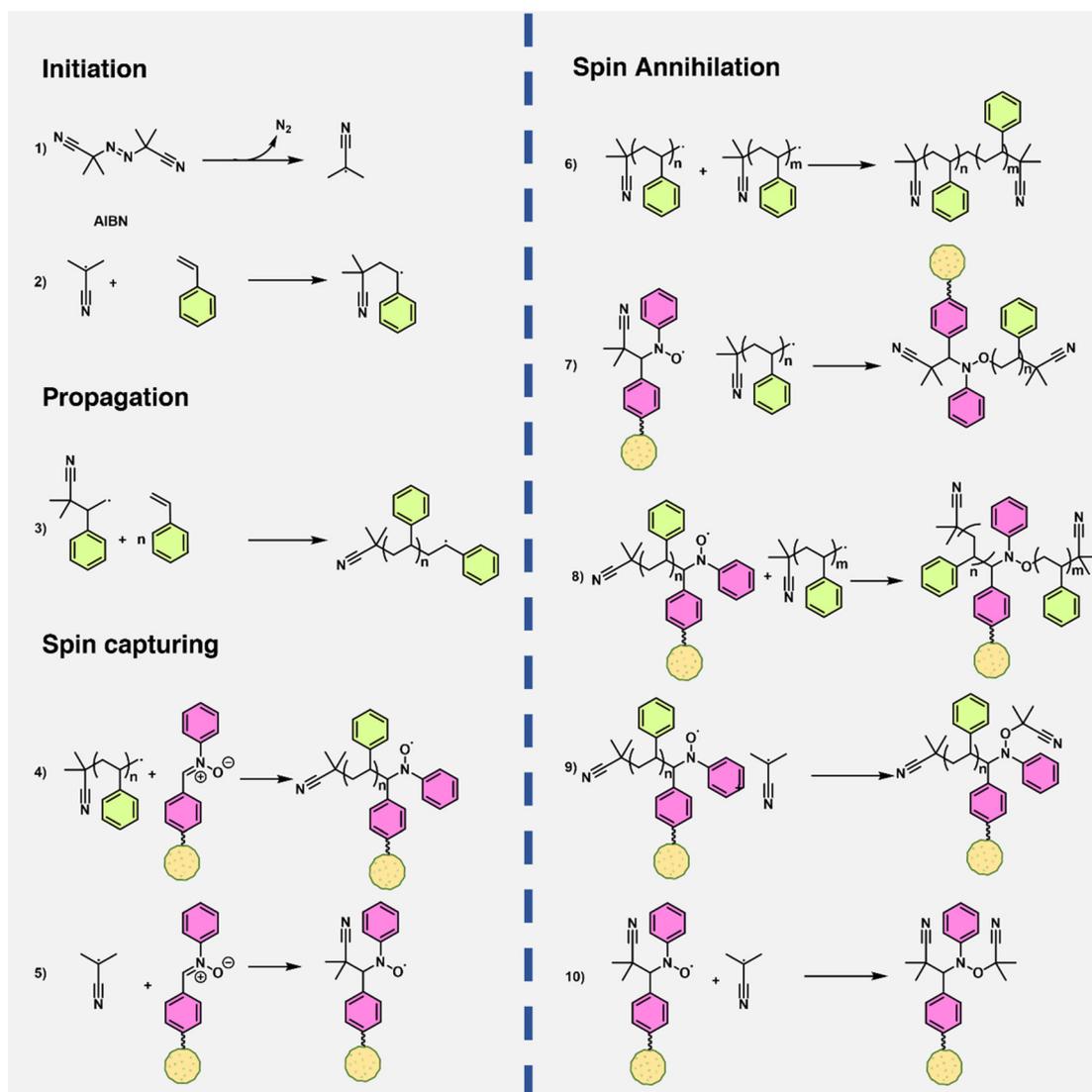
different SEC populations for both free polymers and grafted polymers.

Due to the multiple populations observed at 25% conversion, corresponding to two polymer chains grafting

Table 1 Molecular weights and grafting density of polystyrene grafted nitron functionalized silica

Sample	Conversion determined using $^1\text{H NMR}$ (%)	TGA weight loss due to grafted polymers (%)	M_n FP (g mol^{-1})/ Dispersity	M_n GP (g mol^{-1})/ Dispersity	Grafting density ($\mu\text{mol g}^{-1}$)
PSRHA	25	13.05	38 000/1.34	85 000/1.07 3500/1.09	3.4
	68	26.14	52 000/1.24	29 000/1.56	9.1
	80	37.88	81 000/1.15	56 000/1.35	6.8

The grafting density was determined in $\mu\text{mol g}^{-1}$ units due to the inconsistent surface area of RHA silica as a result of high agglomeration, and the grafting density was determined using eq (1) (refer to the ESI†).



Scheme 3 Schematic representation of the initiation, propagation, spin capturing, and spin annihilation of enhanced spin capturing polymerization over surface nitron functionalities.³⁴ Eq (7), (8), and (9) result in different populations of grafted polymers on the surface. Maximum grafting was observed at quenching, especially at higher conversions. At higher conversion, where chain movements are restricted due to viscosity and steric hindrance, significant grafting can happen according to eq (4) with the maximum molecular weight of grafted chains being in the range of that of free polymers which can be observed in SEC traces.

on one nitron functionality, the grafting density was determined using the free polymer molecular weight, and a polymer grafting density of $3.4 \mu\text{mol g}^{-1}$ was obtained. The

grafting density was significantly increased for 68% conversion to $9.1 \mu\text{mol g}^{-1}$, followed by a decrease to $6.8 \mu\text{mol g}^{-1}$ for 80% conversion. However, the organic weight loss

corresponding to the grafted polymers was observed to increase with an increase in conversion due to the grafting of longer polymer chains with conversion. The maximum grafting density observed was $9.1 \mu\text{mol g}^{-1}$ at 68% conversion despite having a surface nitron grafting density of $215 \mu\text{mol g}^{-1}$ due to the steric hindrance of growing polymer radicals with surface nitron functionalities.

In addition to nitron functionalized RHA silica, similar experiments have been performed on commercially available precipitated silica and commercially available RHA silica. These results have been discussed in the ESI (Fig. S13–S16†).

3.3. “Grafting to” approach for the synthesis of polystyrene grafted silica via 1,3-dipolar cycloaddition of polystyrene macromonomers

The “grafting to” approach for the synthesis of polystyrene grafted silica was carried out via 1,3-dipolar cycloaddition of polystyrene macromonomers over surface nitron functionalities (Scheme 6). The polystyrene macromonomer having a molecular weight of 2600 g mol^{-1} and dispersity 1.07 was synthesized via HBr elimination of a polystyrene macroinitiator synthesized using ATRP (refer to ESI sections S7 and S8† for the method and characterization). Prior to the polymer grafting studies, a series of model reactions were carried out to determine the ability of the nitron functionality in solution

and on the surface to undergo 1,3-dipolar cycloaddition and the potential of the polystyrene macromonomer to undergo 1,3-dipolar cycloaddition reactions. The model reactions include the synthesis of *N*-1-diphenylmethanimine oxide and its cycloaddition reaction with styrene, synthesis of nitron functionalized polystyrene and its cycloaddition reaction with the polystyrene macromonomer and the cycloaddition reaction of nitron functionalized RHA silica with the isobornyl acrylate monomer.

The ability of the nitron functionality to undergo 1,3-dipolar cycloaddition in solution was confirmed using the reaction of benzaldehyde phenylnitron (*N*-1-diphenylmethanimine oxide) with the styrene monomer (refer to ESI S9† for the method). The $^1\text{H NMR}$ analysis confirms the formation of the cycloadduct with the presence of peaks at $\delta = 2.5\text{--}3.5 \text{ ppm}$ and $4.5\text{--}5.5 \text{ ppm}$ (Fig. 5).

The reaction between the polystyrene macromonomer and the polystyrene with nitron end groups confirms the ability of the polystyrene macromonomer to undergo a 1,3-dipolar cycloaddition reaction. The polystyrene with nitron end groups was synthesized via a thiol-ene reaction between -SH functionalized polystyrene and 4-(methacryloyloxy)benzaldehyde phenylnitron which was subjected to the 1,3-dipolar cycloaddition reaction with the polystyrene macromonomer (Scheme 4). The HBr elimination of bromine terminated polystyrene synthesized via the ATRP method yields the ene terminated polystyrene macromonomer.

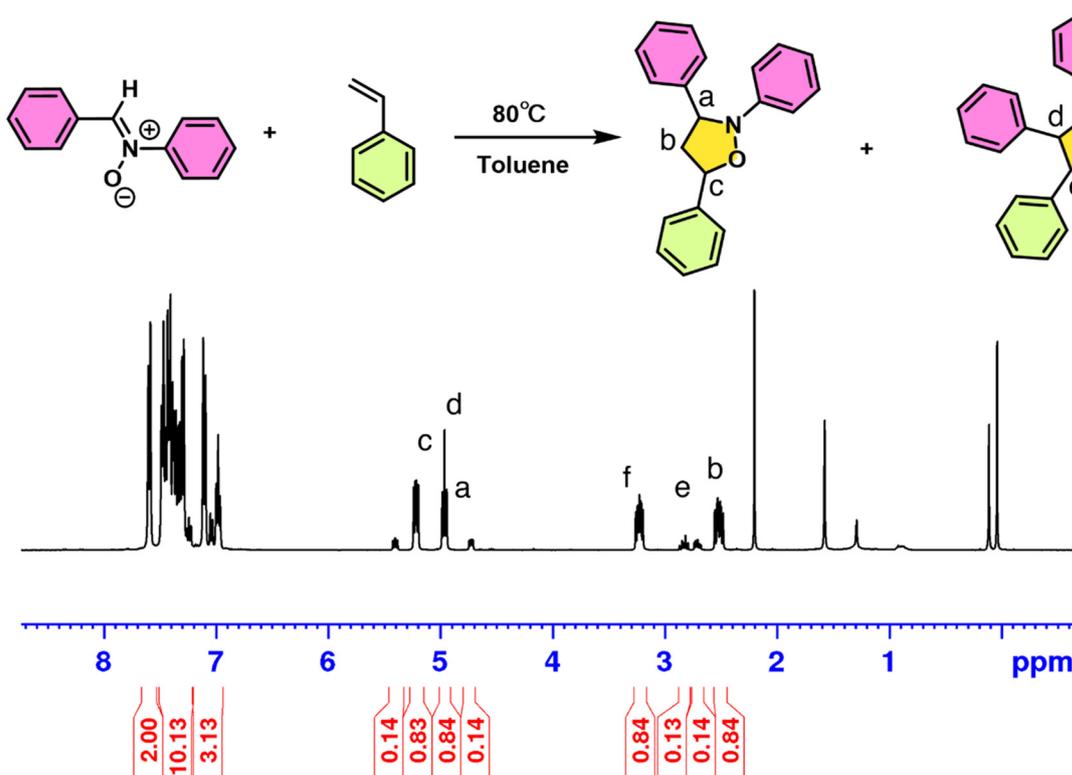
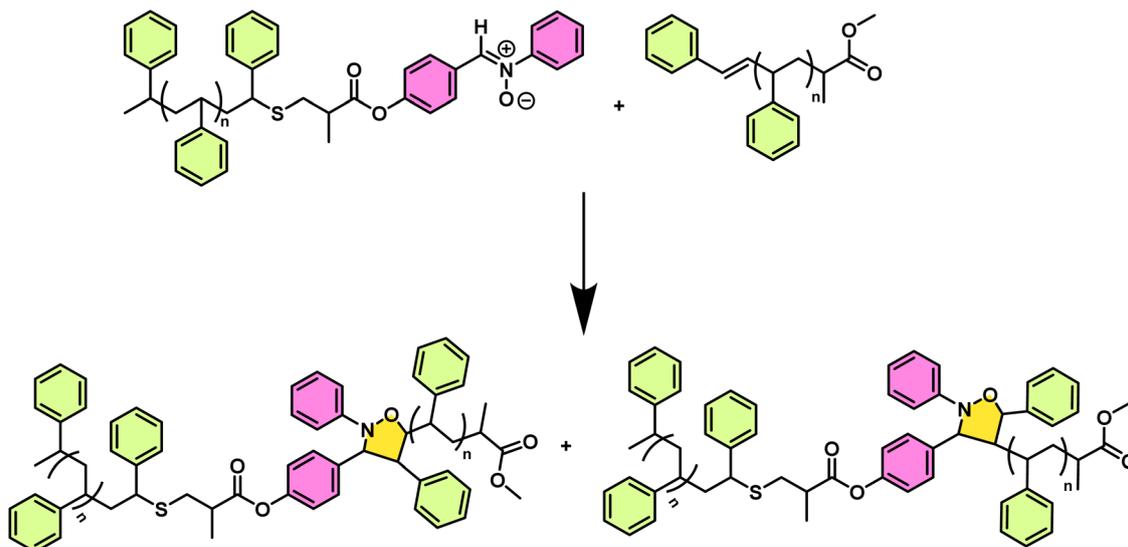


Fig. 5 $^1\text{H NMR}$ spectrum of the cycloadduct formed from styrene and *N*-1-diphenylmethanimine oxide and the peaks present at $\delta = 2.5 \text{ ppm--}3.5 \text{ ppm}$ and $4.5 \text{ ppm--}5.5 \text{ ppm}$ confirms the formation of the cycloadduct.



Scheme 4 1,3-Dipolar cycloaddition of benzaldehyde phenylnitronne functionalized polystyrene with the polystyrene macromonomer.

The presence of peaks corresponding to the benzaldehyde phenylnitronne functionality in ^1H NMR in the range of $\delta = 7.5\text{--}8.0$ ppm confirms the presence of the nitronne functionality (Fig. 6) which further disappears along with the proton corre-

sponding to the alkene functionality upon the cycloaddition reaction with the polystyrene macromonomer.

A model 1,3-dipolar cycloaddition over nitronne functionalized silica using isobornyl acrylate was performed to confirm

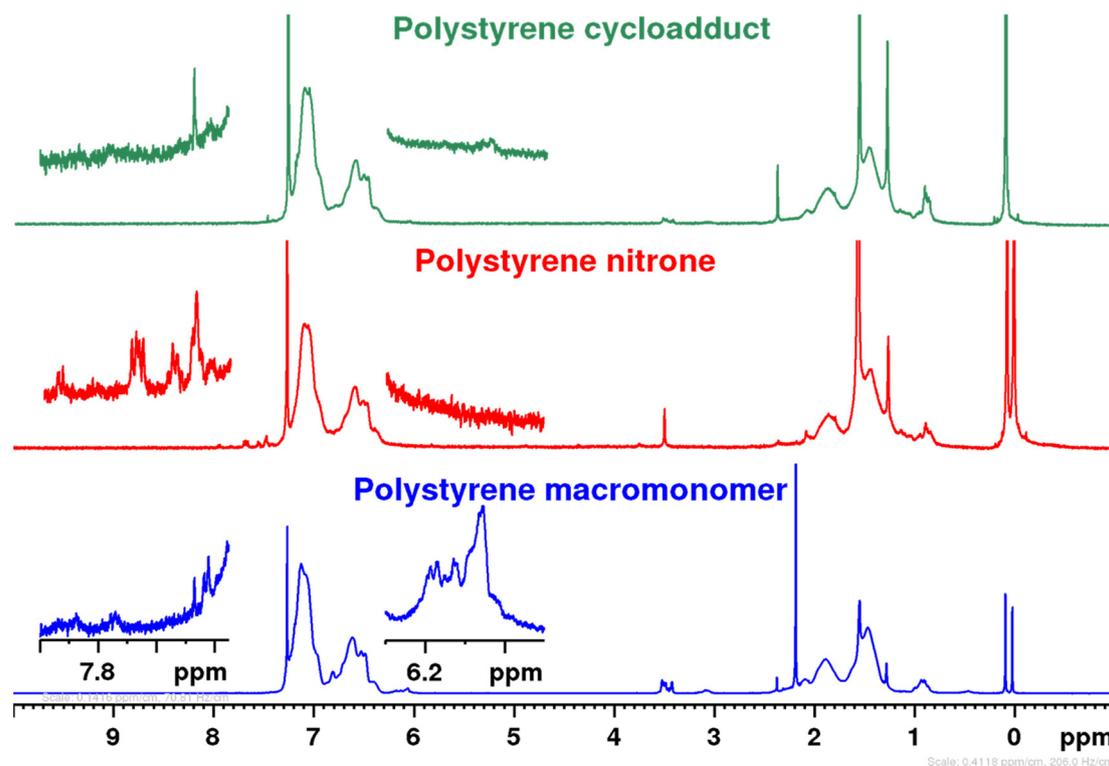


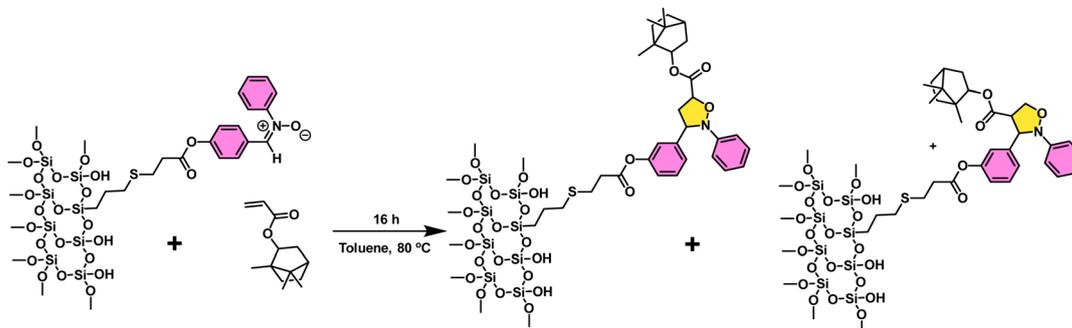
Fig. 6 ^1H NMR spectra of the cycloadduct formed from the polystyrene macromonomer and nitronne functionalized polystyrene. The disappearance of peaks in the region of 7.8 ppm to 8.4 ppm in polystyrene nitronne and 5.9 ppm to 6.3 ppm in the polystyrene macromonomer for the polystyrene cycloadduct confirms the utilization of the nitronne functionality in the 1,3-dipolar cycloaddition reaction.

the potential of surface nitrene functionalities to undergo cycloaddition (Scheme 5).

The monomer isobornyl acrylate was chosen over styrene due to the higher molecular weight, which can impart a notable weight loss in TGA (Fig. 7). The additional weight loss observed below 200 °C for isobornyl acrylate grafted samples is possibly due to the residual toluene present on the silica surface resulting due to low temperature drying to protect the remaining surface nitrene functionalities from hydrolysis. An additional weight loss was observed for isobornyl acrylate

grafted samples with significant reduction in intensity at $\lambda = 327$ nm in solid-state UV-visible analysis (Fig. 7) which confirms the potential of the surface nitrene functionality to undergo the 1,3-dipolar cycloaddition reaction.

After confirming the ability to undergo 1,3-dipolar cycloaddition of the surface nitrene functionality and the polystyrene macromonomer using different model reactions, polystyrene macromonomers were grafted on a nitrene functionalized silica surface *via* the “grafting to” method through 1,3-dipolar cycloaddition (Scheme 6).



Scheme 5 1,3-Dipolar cycloaddition of nitrene functionalized RHA silica with isobornyl acrylate.

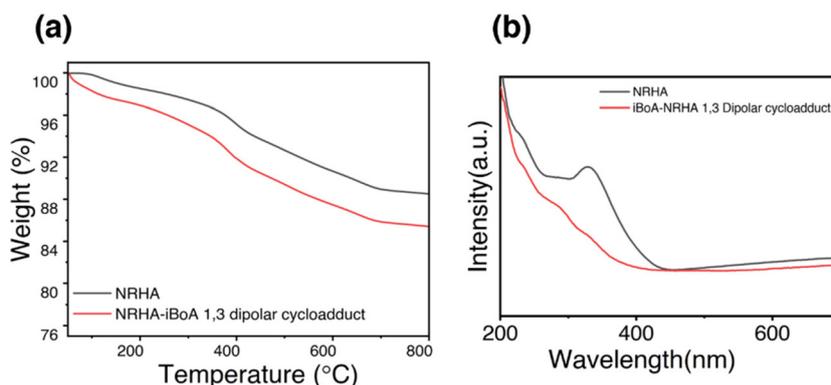
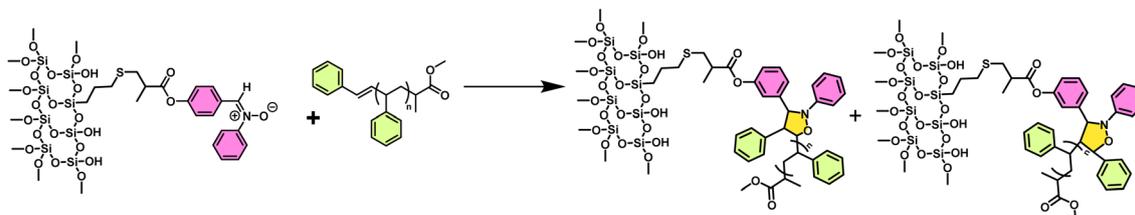


Fig. 7 (a) Thermogravimetric analysis of nitrene functionalized RHA (NRHA) silica and the 1,3-dipolar cycloadduct of isobornyl acrylate over nitrene functionalized RHA silica (NRHA-iBoA). The residual toluene due to the controlled drying yields NRHA-iBoA with higher initial weight loss; however, an additional weight loss was observed corresponding to the grafted isobornyl acrylate. (b) Solid-state UV-visible spectra of nitrene functionalized RHA (NRHA) silica and the 1,3-dipolar cycloadduct of isobornyl acrylate over nitrene functionalized RHA silica (iBoA-NRHA). The disappearance of the band at $\lambda = 327$ nm corresponding to the nitrene functionality confirms the utilization of the surface nitrene functionality in the formation of the cycloadduct.



Scheme 6 1,3-Dipolar cycloaddition of the polystyrene macromonomer over nitrene functionalized (NRHA) silica.

The qualitative confirmation of polystyrene grafting over nitrene functionalized RHA silica *via* 1,3-dipolar cycloaddition was made using FTIR analysis which showed aromatic C–H and out of plane bending vibration at 694 cm^{-1} . The quantification of polystyrene grafting carried out *via* TGA (Fig. 8) gives an additional weight loss corresponding to the degradation of polystyrene grafted *via* 1,3-dipolar cycloaddition reaction.

The XPS analysis of polystyrene grafted RHA silica *via* 1,3-dipolar cycloaddition (Fig. 9) shows characteristics peaks for N 1s corresponding to C=N (398 eV) in nitrene functionalized silica sample along with the formation of a small N 1s peak at a higher binding energy (404 eV) corresponding to the cycloadduct formed, which confirms the formation of a stable C–N bond resulting from polystyrene grafting and

proves the ability of surface nitrene radicals to capture macromonomers.

The synthesized nitrene functionalized silica samples have stability up to $100\text{ }^{\circ}\text{C}$ due to the acidic nature of silica which can hydrolyse the nitrene functionality, so all the polymer grafting studies were carried out up to $80\text{ }^{\circ}\text{C}$. The polymer grafting density was limited regardless of nitrene grafting density which is possibly due to the steric hindrance. The polymer grafted samples were subjected to solid-state UV-visible spectroscopy, and it was observed that the intensity of the peak at $\lambda = 327\text{ nm}$ reduced but it did not completely disappear for polystyrene grafted silica samples (refer to ESI Fig. S24[†]). The study establishes a method for synthesizing nitrene functionalized silica and its efficiency in polymer grafting *via* both “grafting from” and “grafting to”

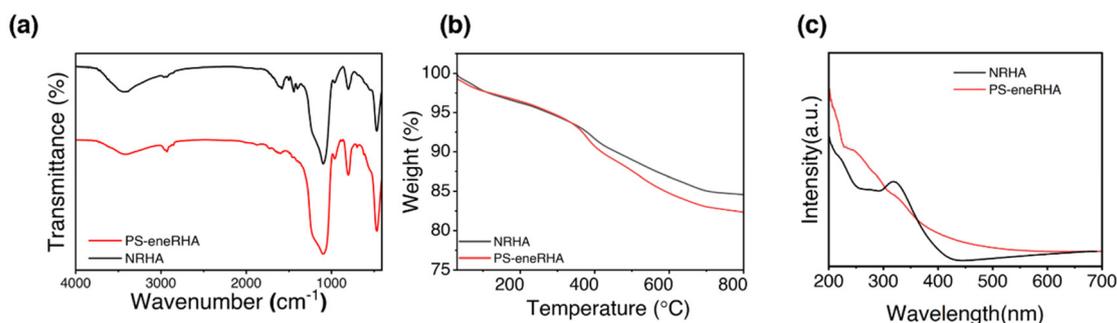


Fig. 8 (a) FTIR analysis of nitrene functionalized RHA (NRHA) silica and the 1,3-dipolar cycloadduct of the polystyrene macromonomer over nitrene functionalized RHA silica (PS-eneRHA). (b) Thermogravimetric analysis of NRHA and PS-eneRHA, and the weight loss corresponding to the grafted polystyrene was lower than that of the polystyrene grafted samples synthesized *via* ESCP, which is due to the lower molecular weight of the polystyrene macromonomer. (c) Solid-state UV-visible spectra of nitrene functionalized RHA (NRHA) silica and polystyrene grafted silica synthesized *via* 1,3-dipolar cycloaddition (PS-eneRHA). The band corresponding to the nitrene functionality was observed to decrease in intensity after polymer grafting.

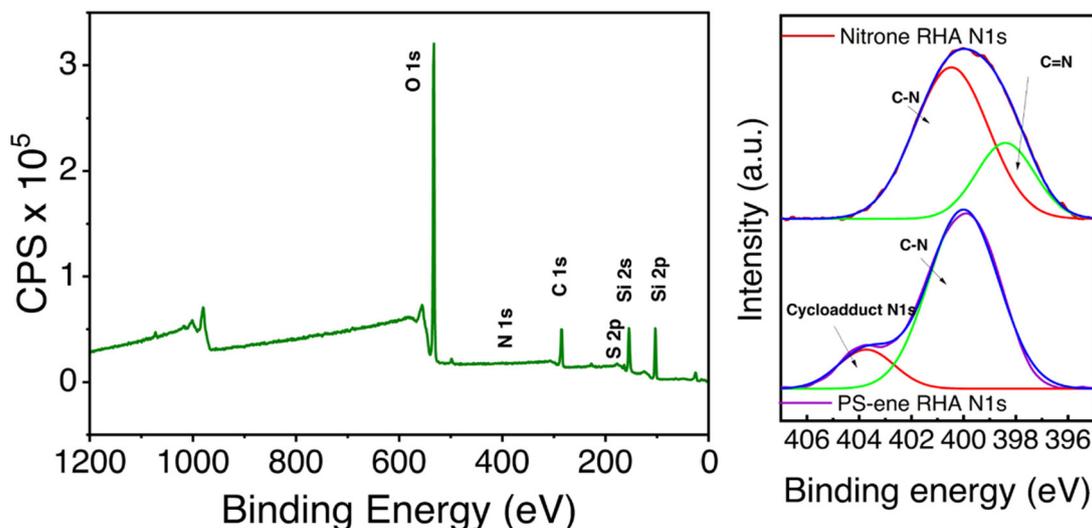


Fig. 9 XPS plot of wide scan and deconvoluted N 1s scan for PS-eneRHA, and polymer grafting results in the formation of a small peak at 404 eV corresponding to nitrogen present in cycloadduct along with the peaks corresponding to the C–N functionality at 400 eV.

approaches using ESCP and a 1,3-dipolar cycloaddition reaction.

The efficiency of surface nitrene radicals on different types of silica has also been studied using commercial precipitated silica and commercial RHA silica samples synthesized *via* post-modification and the grafting densities have been compared with those of co-condensed nitrene functionalized RHA silica samples. The co-condensed nitrene RHA samples were observed to have a higher polymer grafting density maximum of $9.1 \mu\text{mol g}^{-1}$. The effect of the surface area and method of functionalization on polymer grafting has been discussed for all three different types of nitrene functionalized silica samples (refer to the ESI†).

4. Conclusion

The nitrene functionality has been introduced *via* a thiol-ene reaction on silica synthesized from sodium silicate derived from rice husk ash. The presence of active nitrene functionalities on the surface was determined using solid-state UV-visible spectroscopy. The nitrene grafting density on the silica surface was observed to be $215 \mu\text{mol g}^{-1}$ for nitrene functionalized RHA silica synthesized *via* co-condensation. The efficiency of the surface nitrene functionality towards polymer grafting was studied *via* enhanced spin capturing polymerization using styrene, and 1,3-dipolar cycloaddition using the polystyrene macromonomer. The nitrene functionalized silica was observed to have the potential for grafting polymer chains. The extent of grafting studied *via* ESCP indicates that the co-condensed samples (NRHA) have a better surface area and higher nitrene loading and they were observed to have a maximum polymer grafting density of $9.1 \mu\text{mol g}^{-1}$. The “grafting to” method of functionalization *via* 1,3-dipolar cycloaddition yields a grafting density of $6.8 \mu\text{mol g}^{-1}$. The efficiency of the nitrene functionality in polymer grafting *via* ESCP and 1,3-dipolar cycloaddition proves the potential of the surface nitrene functionality to undergo polymer grafting *via* “grafting from” and “grafting to” methods.

Conflicts of interest

There are no conflicts to declare.

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