

Synthesis of Novel Hexathiolated Squalene and Its Thiol-Ene Photopolymerization with Unsaturated Monomers

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ABSTRACT

In this work is described the synthesis of a multifunctional thiolated squalene. Thiol-ene coupling reactions were employed to functionalize the six double bonds of squalene, using thioacetic acid. Hydrolysis of the resulting thioacetates, rendered the corresponding hexathiolated squalene SQ6SH. This compound was further photopolymerized separately with triallyl cyanurate, pentaerythritol triacrylate and diethyleneglycol divinyl ether. Real Time FTIR kinetics revealed that homopolymerization of the ene monomers took place in addition to the thiol-ene photopolymerization. Flexible films were obtained when SQ6SH was photopolymerized in bulk with the above mentioned unsaturated monomers.

Keywords: Squalene; Thiol-Ene; Photopolymerization; Biomaterials; Kinetics

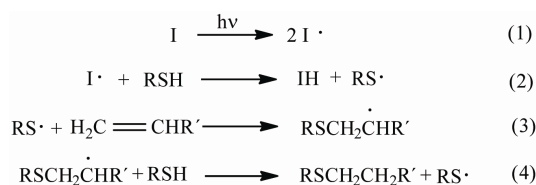
1. Introduction

The search for new biobased raw materials to prepare polymers, has become an important issue due to increasing depletion of oil sources. A wide array of compounds like, terpenes [1,2], vegetable oils [3,4], cellulosic materials [5,6], carbohydrates [7,8], etc., have been used either as they are found naturally or chemically modified, to prepare polymeric materials.

Squalene is a triterpenoid found in shark liver, in wheat germ, and in olive oil. It is also present in the human body at 5% - 8% w/w in the body fat [9]. It is produced industrially from the olive oil and it has found uses in the cosmetics industry and as skin humectant. Polyunsaturation of this compound make it a very attractive candidate to use it in the preparation of biobased polymers.

The “Click” thiol-ene Chemistry has acquired renewed interest due to the inherent characteristic of this kind of reactions. For instance, these reactions are very rapid, proceed almost quantitatively, and are not inhibited either by oxygen or humidity [10-12]. If monofunctional thiols and alkenes are irradiated with UV light, occurs thiol-ene coupling, but if difunctional or multifunctional thiols and alkenes are used, photopolymerization takes place.

In **Scheme 1** is depicted the mechanism of thiol-ene photopolymerizations [13]. Initiation involves the photo-



Scheme 1. Mechanism of thiol-ene photopolymerization.

lysis of the photoinitiator when the photocurable system is irradiated with UV light. Then the produced primary radicals abstract the hydrogen atom of thiol groups forming thiyl radicals. These radicals can react with the double bonds of the unsaturated compound forming a secondary radical which in turn can abstract the hydrogen atom of a second molecule of the thiol generating a new thiyl radical to repeat the cycle.

In continuing with our effort to utilize renewable starting materials to prepare monomers for thiol-ene photopolymerization, the main objective of this research was to develop a methodology to prepare thiols using biobased multiunsaturated compounds, such as vitamin A, polyisoprene and squalene. This last compound was preferred to start with, because of its low molecular weight and ease of handling. The hexathiolated squalene was synthesized and subsequently photopolymerized with unsaturated monomers. Kinetics of photopolymerization was determined by means of Real-Time FTIR spectroscopy.

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2. Experimental Section

2.1. Materials and Equipment

Squalene (SQ), thiol acetic acid (TAA), sodium hydroxide, 2,2-Dimethoxy-2-phenylacetophenone (DMPA), diethyleneglycol divinyl ether (DEGVE), triallyl cyanurate (TAC), pentaerythritol triacrylate (PETA), were all reagent grade and purchased from Aldrich Co (Toluca, Mexico). In **Scheme 2** are shown the chemical structures of SQ, of the prepared compound 2, 6, 10, 15, 19, 23, hexamethyl tetracosane-3, 7, 11, 14, 18, 22-hexathiol (SQ6SH) and those of the unsaturated monomers mentioned above. Routine infrared spectra and photopolymerization kinetics were performed on a Magna Nicolet 550 Infrared spectrometer (Middleton, WI). NMR spectra were obtained using a 300 MHz Jeol NMR spectrometer. Perkin Elmer Elemental Analyzer Series II CHNS/O 2400 (Waltham, MA).

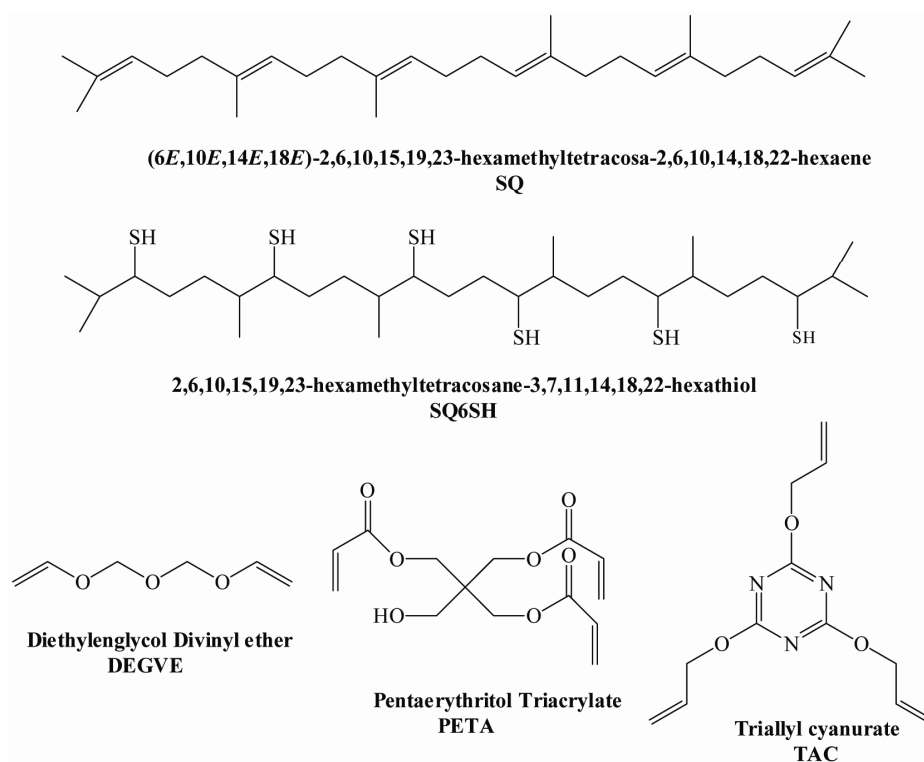
2.2. Synthesis of the Hexathioacetate of Squalene (SQ6TA)

In quartz tube were weighed 1 g (2.43×10^{-3} mol) of SQ, 2.22 g (2.92×10^{-2} mol) of TAA and 1.25×10^{-2} g (4.86×10^{-5} mol) of DMPA. Once that all components were added to the tube, it was placed in an UV light oven provided with a 300 W Fusion lamp. The tube was irradiated during 15 minutes and after this time the tube was allowed to cool to room temperature. The reaction mixture

was dissolved in chloroform and extracted with 10% NaOH w/w (3×30 mL) in order to eliminate the excess of TAA. Then, the organic phase was washed with distilled water and subsequently dried with anhydrous sodium sulfate. The solvent was rotoevaporated and the crude SQ6TA was obtained at 85% yield.

2.3. Synthesis of SQ6SH

In a three neck round-bottomed flask provided with condenser, thermometer, nitrogen inlet and magnetic stirring, were added 5.5 g (6.34×10^{-3} mol) of the crude SQ6TA, 1.1 g (2.75×10^{-2} mol) of powdered NaOH and 40 mL of methanol. Vigorous stirring was needed to dissolve the sodium hydroxide. Then, the reaction mixture was refluxed by 5.5 hours under nitrogen atmosphere. Afterwards, the solvent of reaction mixture was evaporated and the residue was dissolved in water and acidified with a 0.1 N HCL solution. The aqueous phase was extracted with chloroform, (3×30 mL) and the resulting organic phase was washed with distilled water, dried with anhydrous sodium sulfate and rotoevaporated to eliminate the solvent. The crude product was further purified by column chromatography using silica gel as support and a mixture hexane: ethyl acetate 98:2 as eluent. A yellowish liquid with astringent odor was obtained at 77% yield. Elemental Analysis: Theory C 58.57, H 10.16, S 31.27, Experimental C 58.22, H 9.96, S 31.06.



Scheme 2. Chemical structures of squalene (SQ), of synthesized compound SQ6SH and of ene monomers used in this study.

2.4. Determination of Kinetics of Thiol-Ene Photopolymerizations by RT-FTIR

RT-FTIR was used to monitor the kinetics of photopolymerization of the synthesized monomer SQ6SH with a Nicolet Magna 550 FT-IR spectrometer equipped with a DTGS detector fitted with a UVEXS model SCU 110 mercury lamp. The intensity of the UV irradiation was measured with a UV Process Supply Inc. Control cure radiometer. Thiol-ene formulations were prepared by mixing 0.5 g of SQ6SH with each of the ene monomers in stoichiometric ratios, considering the number of functional groups. DMPA was added to each formulation at 1 mol %. All kinetics experiments were conducted at 25 °C. The course of the photopolymerization was followed by simultaneously monitoring the decrease of the peaks of the corresponding functional groups. For instance, the thiol was monitored following the infrared absorption band at 2560 cm⁻¹ due to its S-H group. Allyl group conversion was monitored using the carbon-carbon double bond absorption peak at 1646 cm⁻¹. Each kinetic run was carried out a minimum of five times. Data were collected at a rate of one spectrum per second and processed with the OMNIC Series software. Conversions were calculated using the ratio of peak areas to the peak area prior to photopolymerization. The kinetic parameter R_p/M_0 , was determined from the initial slopes of the irradiation time-conversion curves according to Equation (5):

$$R_p/M_0 = ([\text{conversion}]t_2 - [\text{conversion}]t_1)/(t_2 - t_1) \quad (5)$$

where R_p and M_0 are respectively the rate of photopolymerization and the initial monomer concentration as well as the conversions are as determined from the curves at irradiation times t_1 and t_2 . The obtained profile of diminution of the absorbance of a determined peak, using the *Series* software included in OMNIC, is processed in excel, by using the Equation (6), where C_t is the conversion at any time, B_0 is the absorbance of the peak at zero time and B_t is the absorbance at any given time

$$C_t = [B_0 - B_t/B_0] \times 100 \quad (6)$$

In **Table 1** are shown the amounts and ratios of the comonomers used in the determination of the photopoly-

Table 1. Amounts and ratios of SQ6SH and of the three ene monomers. SQ6SH was formulated separately with each of the three monomers and 3 mol % of DMPA.

	SQ6SH	DEGVE	TAC	PETA	DMPA
Equation	1	3	2	3	0.03
mol	3.25E-04	9.75E-04	6.50E-04	6.50E-04	9.75E-06
MW (g/mol)	615.2	158.2	249.27	298.30	256.30
g	0.2	0.15	0.16	0.19	0.002

merization kinetics by RT-FTIR.

2.5. Study by DSC

The Tg's of the prepared polymers were measured on DSC2920 thermal analyzer (TA instruments, Inc.) with a heating rate of 10 °C/min under nitrogen atmosphere. The average weight of test samples was 3 - 5 mg.

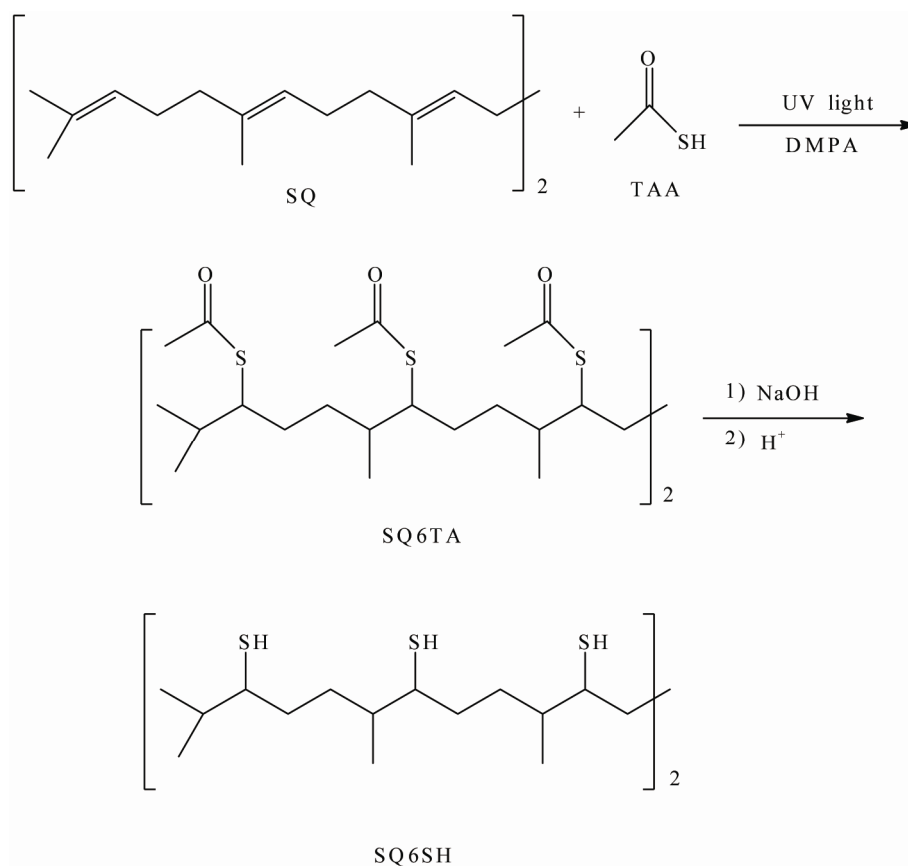
3. Results and Discussion

3.1. Synthesis of SQ6SH

The development of monomers based on renewable source derived materials has been one of the main interests of our research group. Previously, we have reported the use of vegetable oil [14], sucrose [15-17] and isosorbide [18], as starting materials to prepare epoxy monomers for cationic photopolymerization, and allylated carbohydrates to use them as comonomers in thiol-ene photopolymerization. To go further to obtain purely biobased polymers using the thiol-ene photopolymerization technique, we needed to prepare multifunctional thiols derived from natural sources. The "Click" thiol-ene coupling between an unsaturated compound such as SQ with TAA under UV irradiation, and subsequent hydrolyzation of the obtained thioacetate, proved to be an effective method to synthesize the hexafunctional thiol (see **Scheme 3**). Here, is worth mentioning that hydrothiolation of internal double bonds is not as fast as that of terminal double bonds. Hoyle, *et al.*, [19,20] found that hydrothiolation of 1-hexene is eight times faster than that of trans-2-hexene and 18 times faster than of trans-3-hexene. This difference in reactivity was attributed to steric effects and also to the reversibility of the addition reaction of the thyl radical to the double bond. As in our case, all double bonds of SQ are internal, adjacent to a methyl group and of type trans, they were not very reactive, therefore, it was necessary to add a 100% excess of TAA to force the completion of the reaction. **Figure 1** shows the ¹H NMR spectrum of SQ6TA. It can be observed the presence of the peak at 2.3 ppm corresponding to the methyl of the thioacetate groups and the absence of the signals at 5 ppm corresponding to the double bonds of SQ. The proton NMR spectrum of SQ6SH depicted in **Figure 2** reveals the disappearing of the peak at 2.3 ppm. Also, in **Figure 3** are shown the IR spectra of the starting material SQ, of the intermediate compound SQ6TA and that of the final product where the SH peak is marked at 2560 cm⁻¹.

3.2. Photopolymerization Studies

The prepared SQ6SH was photopolymerized with three types of unsaturated monomers, namely, vinyl ether (DEGVE), allyl isocyanurate (TAC) and pentaerithritol



Scheme 3. Methodology to prepare SQ6SH.

triacrylate (PETA). All three compounds were selected based on their different reactivities. The reactivity depends on the substitution in the double bond group. The higher the electron density in the double bonds the higher the reactivity. **Figure 4** shows the kinetics curves obtained by RT-FTIR of the photopolymerization of the photocurable system SQ6SH/TAC. It was observed that this photocurable system was rather reactive, with the double bonds groups being consumed more rapidly and with higher conversion than the thiol groups. As in the step-growth polymerization, both thiol and ene compounds must be consumed at the same time, it is clear that two polymerization mechanisms are occurring concurrently: the thiol-ene photopolymerization and the radical homopolymerization of TAC. On one hand, although it has been reported that allylic bonds can homopolymerize to some extent [21], in our case, monomer TAC polymerize readily achieving 42% conversion in the first 20 seconds and 75% after 120 seconds. On the other hand, low conversion of thiol group (25%) was produced in the same run as a result of the multifunctionality of SQ6SH. As usual, in thiol-ene photopolymerizations, if multifunctional enes and thiols are used, crosslinked polymers are obtained. The higher the multifunctionality, the higher the crosslink density in the produced polythioether. Lower

conversions of comonomers are achieved as a result of the higher amount of trapped functional groups in the tridimensional crosslinked network. One of the main characteristics of the thiol-ene photopolymerization is the delayed onset of gel formation in comparison with the radical photopolymerization of acrylates. Jacobine and collaborators [22] proposed an equation (Equation (7)) to calculate the gel point of thiol-ene systems considering the functionality of comonomers:

$$\alpha = \sqrt{\frac{1}{r(f_{thiol} - 1)(f_{ene} - 1)}} \quad (7)$$

where α is the gel point, r is the thiol-ene molar ratio based on functional groups, f_{thiol} is the thiol functionality and f_{ene} is the ene functionality. In the case of our photocurable system TAC/SQ6SH, as the functionality of thiol compound is 6 and that of the ene compound is 3, the gel point obtained by using Equation (1), is 22%, which corresponds fairly with the observed conversion of SQ6SH. The difference between the conversion of the thiol and that of the ene compound correspond to the amount of homopolymerization of TAC.

Figure 5 depicts the kinetics curves of double bonds of DEGVE and of thiol groups of SQ6SH. It was observed

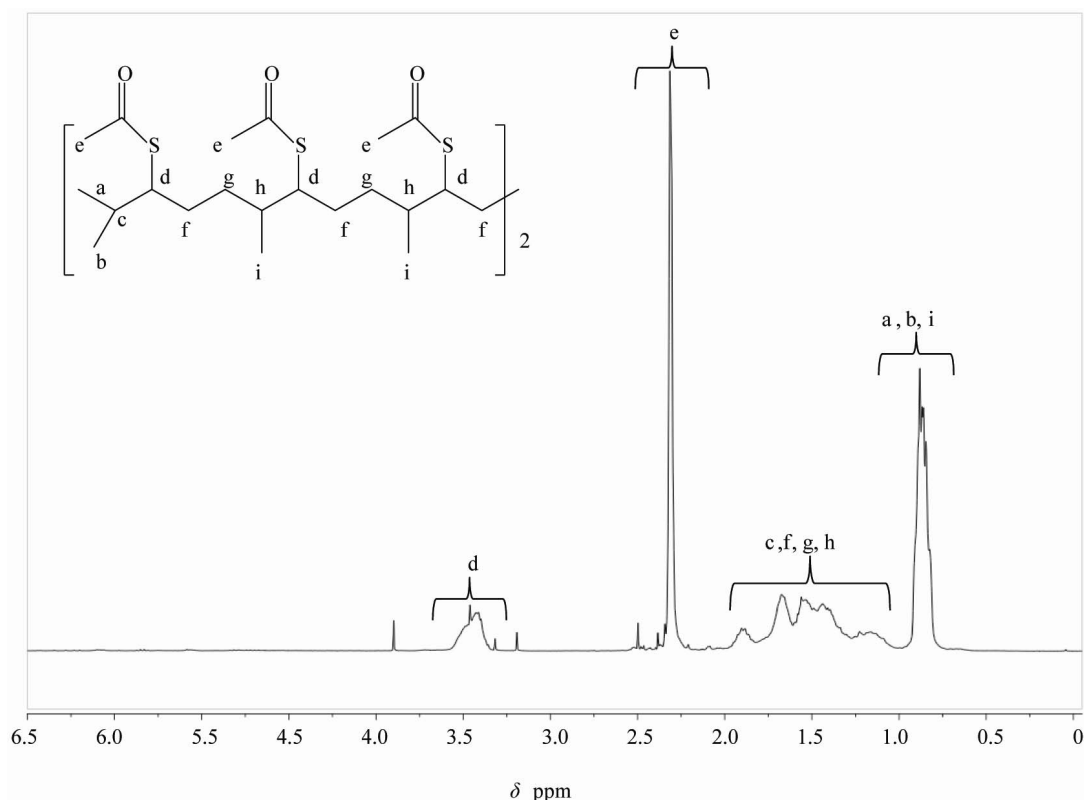


Figure 1. ^1H NMR spectrum of SQ6TA run in CDCl_3 .

a similar situation than in the case of TAC, as in this case it also took place homopolymerization of vinyl ether in addition to the step-growth thiol-ene photopolymerization, although it was slightly slower than the formulation with TAC. The hexafunctionality of SQ6SH and the bifunctionality of DEGVE resulted in 25% conversion of the thiol groups (theory 25%).

The kinetics curves of the system PETA/SQ6SH are shown in **Figure 6**. Although PETA is also trifunctional, the achieved conversion for the thiol groups was 17.5%. This apparently slightly lower conversion could be explained by the higher reactivity of PETA, as the homopolymerization of this monomer was somewhat faster than the other two ene monomers. In this way, the chances to react with the thiol are reduced to some extent.

3.3. DSC of Produced Films

The formulations of SQ6SH with the three unsaturated monomers were photopolymerized in bulk, as thin films on glass slides. Flexible films were obtained with sub-zero T_g 's. The values of T_g of the obtained polymers are shown in **Table 2**. In the resulting mixture of polymers, the inherent high mobility of the aliphatic hexathiolated squalene backbone added to the flexibility imparted by the thioether groups resulted in the formation of polymers

Table 2. T_g 's of the polymers derived from SQ6SH with the three ene monomers and from their homopolymers. The T_g was determined by DSC considering the second heating of the sample.

Sample	T_g °C
SQ6SH/TAC	-22.61
SQ6SH/PETA	-25.12
SQ6SH/DEGVE	-20.29
TAC	28.75
PETA	25.24
DEGVE	41.02

with T_g 's around -20°C . For comparison pristine Ene monomers were also polymerized in bulk and the T_g of these polymers was also determined. It can be seen that the T_g of the photopolymers are around 50°C - 60°C higher than that of the polymers obtained using the SQ6SH, due to the already mentioned higher flexibility of the formed polythioethers.

4. Conclusion

A novel biobased multifunctional thiolated comonomer was prepared using SQ as starting material. The comonomer was produced in two stages. First, SQ was irradiated in the presence of TAA to produce the hexathio-

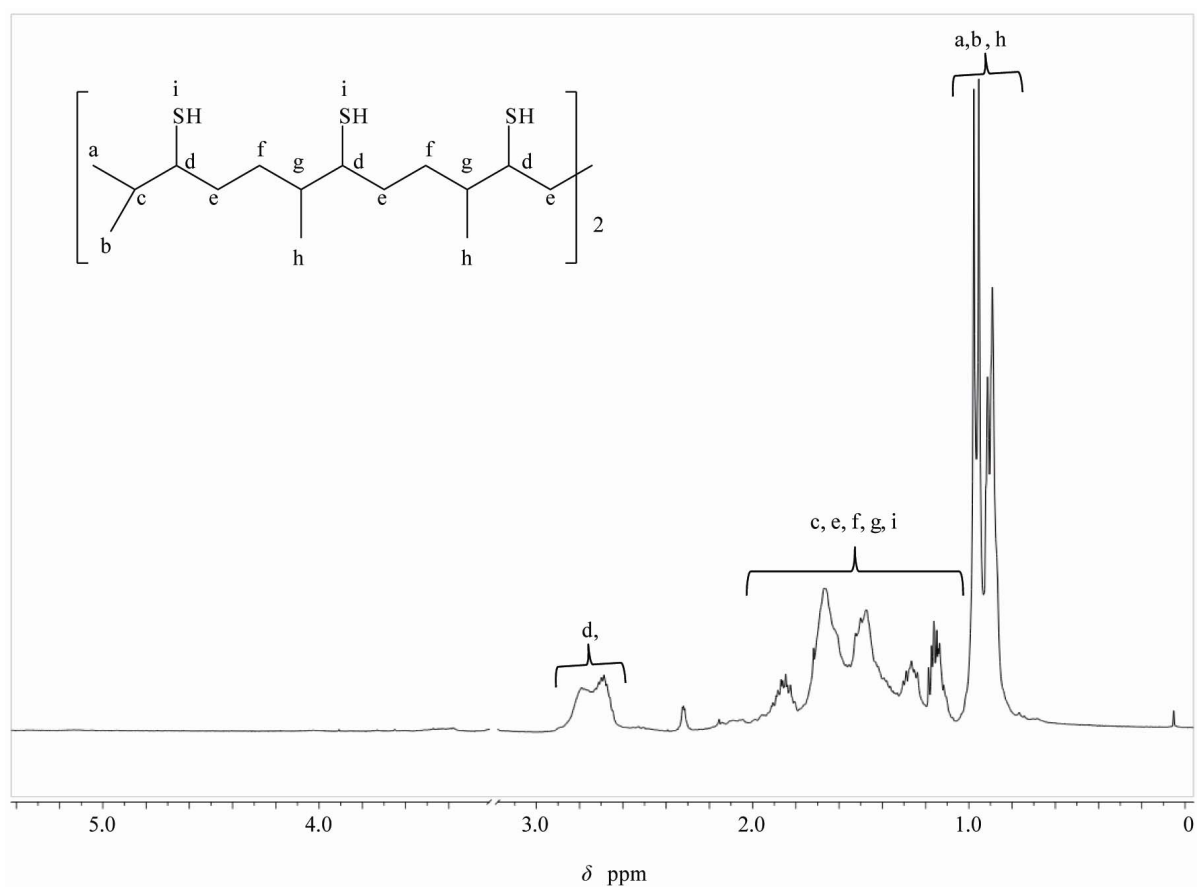


Figure 2. ^1H NMR spectrum of SQ6SH run in CDCl_3 .

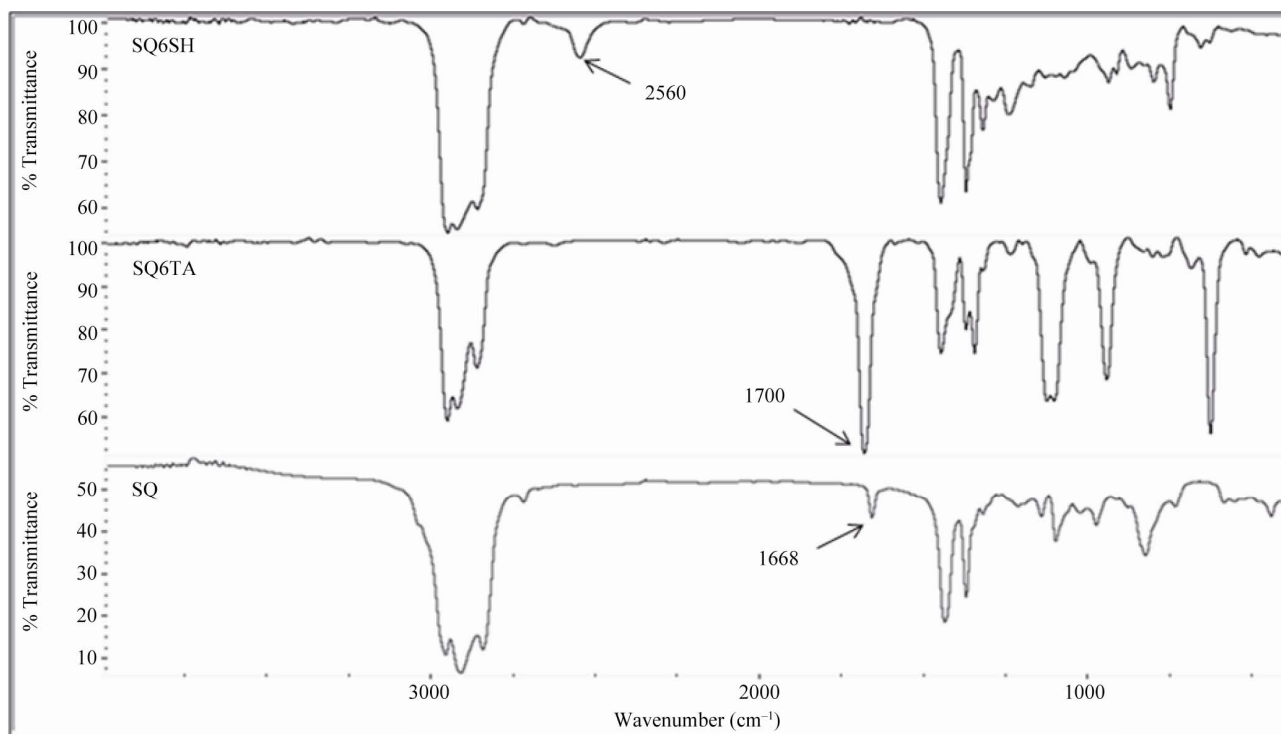


Figure 3. Comparison of FTIR spectra of SQ, SQ6TA and SQ6SH.

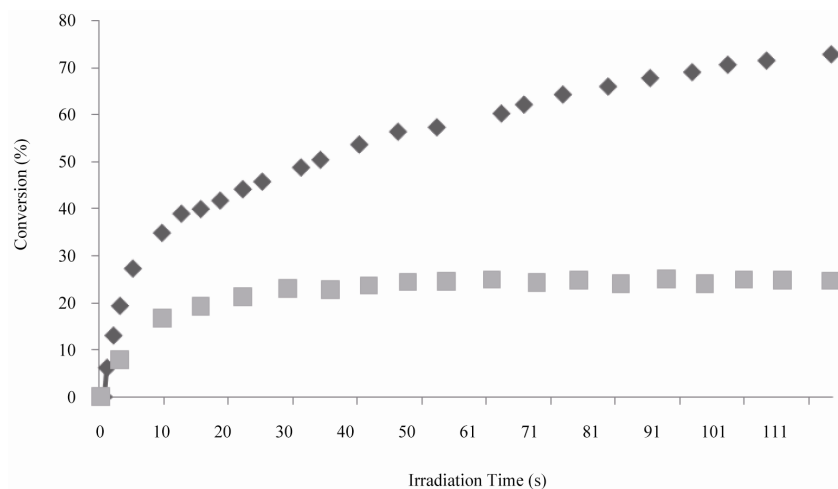


Figure 4. Comparison of conversion vs time curves for thiol groups of SQ6SH (■) and double bonds of TAC, (◆), UV light intensity was 15 mW/cm².

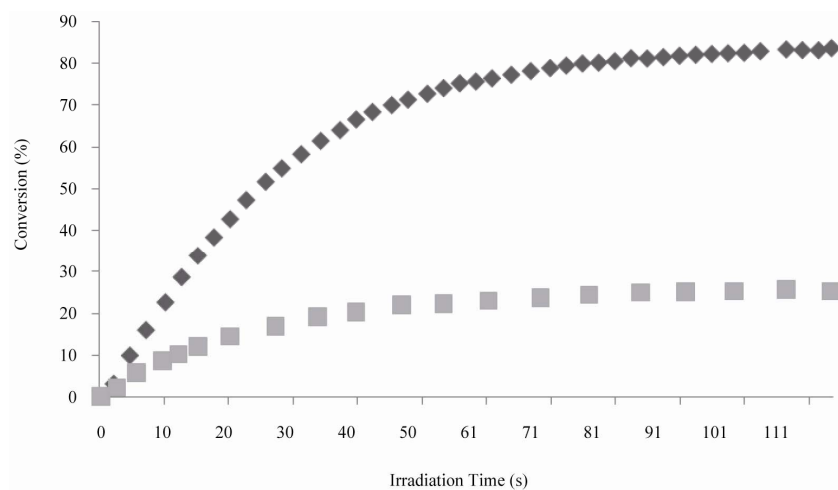


Figure 5. Comparison of conversion vs time curves for thiol groups of SQ6SH (■) and double bonds of DEGVE (◆), UV light intensity was 15 mW/cm².

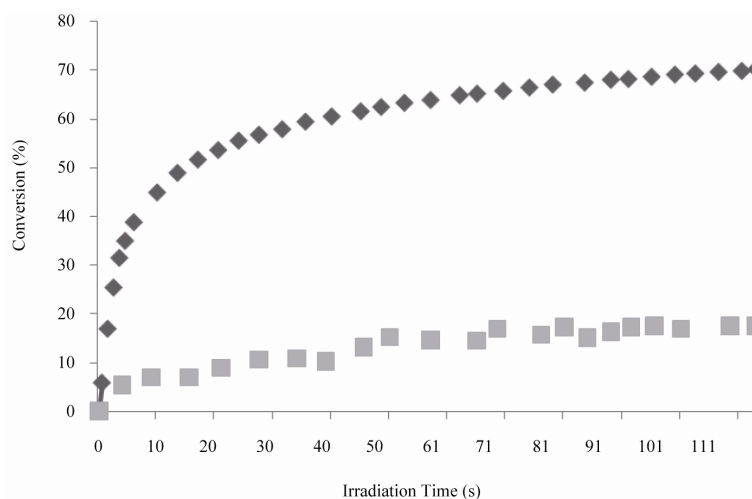


Figure 6. Comparison of Conversion vs Time curves for thiol groups of SQ6SH (■) and double bonds of PETA, (◆), UV light intensity was 15 mW/cm².

ester. Then in the second stage the thioester was hydrolyzed in basic methanolic solution to obtain the hexathiolated monomer SQ6SH. Photopolymerization of this compound separately with three unsaturated monomer resulted in the formation of very flexible polythioether films with T_g 's in the range -20°C to -25°C . The photocurable systems involving the prepared multifunctional thiol were rather reactive observing in all cases, that both homopolymerization and step-growth thiol-ene photopolymerization occurred concurrently.

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