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Synthesis of Tris(4-Methoxyphenyl)Phenylsilane Using Phenylsilane and 4-Iodo Anisole Catalyzed by Palladium Complex

Aldes Lesbani^{1*}, Risfidian Mohadi¹, Eliza¹, and Mardiyanto²1.

Department of Chemistry, Faculty of Mathematics and Natural Sciences, Universitas Sriwijaya, Ogan Ilir 30662, Sumatera Selatan, Indonesia 2. Department of Pharmacy, Faculty of Mathematics and Natural Sciences, Universitas Sriwijaya, Ogan Ilir 30662, Sumatera Selatan, Indonesia *E-mail: aldeslesbani@pps.unsri.ac.id **Abstract**The synthesis of tris(4-methoxyphenyl)phenylsilane through the coupling reaction of phenylsilane and 4-iodoanisole using palladium tertiary tributylphosphine as a catalyst was carried out through the determination of the base, solvent, and reaction time. The results showed that tris(4-methoxyphenyl)phenylsilane can be synthesized to form maximum yield using 1,4-diazabicyclo[2.2.2]octane (DABCO) as a base, tetrahydrofuran as a solvent, and a 5-day reaction time. The highest yield of tris(4-methoxyphenyl)phenylsilane was 35%.

Abstrak

Sintesis Tris(4-Metoksifenil)Fenilsilan Menggunakan Fenilsilan dan 4-Iodida Anisol dengan Katalis Senyawa Kompleks Paladium. Sintesis tris(4-metoksifenil)fenilsilan melalui reaksi kopling antara fenilsilan dan 4-iodida anisol menggunakan senyawa paladium tersier tributilfosfin sebagai katalis telah dilakukan berdasarkan penentuan senyawa basa, pelarut, dan waktu reaksi kopling. Hasil penelitian menunjukkan bahwa senyawa tris(4-metoksifenil)fenilsilan dapat disintesis menghasilkan persentase produk maksimum pada penggunaan senyawa 1,4-diazabisiko[2,2,2]oktan (DABCO) sebagai basa, tetrahidrofuran sebagai pelarut, dan waktu reaksi kopling selama 5 hari. Persentase tertinggi produk senyawa tris(4-metoksifenil)fenilsilan adalah 35%. **Keywords:** 4-iodoanisole, palladium, phenylsilane, tris(4-methoxyphenyl)phenylsilane

1. Introduction The coupling reaction is one of the most useful techniques for synthesizing valuable chemical materials [1]. This reaction has been developed in couplings such as Suzuki-Miyaura, Negishi, Heck, and Sonogashira. Prof Suzuki and Prof. Negishi won the Nobel Prize in 2010 for research of palladium as catalyst for coupling reaction in organic synthesis [2]. The coupling reaction occurs when a carbon-carbon chain is combined using a powerful catalyst such as a complex of transition metals [3]. The example this reaction is asymmetric synthesis of drugs with chirality in the compound [4], which is still developed until now. The development of the coupling reaction has included various substrates depending on molecule targets. Previous research using trialkoxysilane and aryl halide as reactants was carried out by Manoso et al. [5], Denmark et al. [6], and Murata et al. [7]. This reaction included coupling aryl halide with organosilane. Although the coupling reaction has been done using several materials, it is limited due to by-products formed by aryl halide. Yamanoi et al. [8] also found by-products when using aryl halide as a reactant. By changing a halide of chloride or bromide to iodide, the by-product percentage may be significantly reduced. Thus, aryl iodide is an appropriate substrate for coupling with organosilica [9]. Lesbani et al. applied the coupling reaction of aryl iodide for the synthesis of pesticides [10]. The synthesis of materials with unique photoluminescence was also reported by Lesbani et

al., who coupled aryl halide with bulky organosilica [11]. Its unique photoluminescence properties include a violet-blue color under irradiation of UV at 365 nm and a quantum yield of 0.55. Furthermore, Lesbani et al.

Lesbani, et al. *Makara J. Sci.* March 2014 | Vol. 18 | No. 114 enlarged the scope of coupling reactions by using not only aryl iodides [12–13] but also organosilicas [14–15] as starting materials. These researchers found that, although the various aryl iodides and organosilica can be reacted using palladium catalyst through the coupling reaction, there was a slight increase in the yield. Therefore, the optimization of the coupling reaction in this case is vital to increase the coupling product yield. This present research examined the optimization of reactions of phenylsilane and 4-iodoanisole to tris(4-methoxyphenyl) phenylsilane by evaluating bases, solvents, and reaction times. The tris(4-methoxyphenyl) phenylsilane was synthesized using 1,4-diazabicyclo[2, 2, 2]octane (DABCO) as a base and tetrahydrofuran as a solvent, and a 5-day reaction time.

2. Methods **Materials and equipment.** All experiments were carried out using Schlenk techniques under an argon or nitrogen atmosphere in a vacuum system. Chemicals 4-iodoanisole, palladium tertiary tributylphosphine, diethyl ether, tetrahydrofuran, dimethoxyethane, 1,4-dioxane, tetrahydrofuran, triethylamine, diisopropylethylamine, potassium phosphate, TLC plate, ethyl acetate, and dichloromethane were obtained from Merck. 1,2,2,6,6-Pentamethylpiperidine, N-methylpiperidine, N-methylmorpholine, 1,8-diazabicyclo[5,4,0]undec-7-ene, quinoline, 2,6-lutidine, cyclopentyl methyl ether, and pyridine were obtained from Aldrich, and 1,4-diazabicyclo-[2,2,2]octane (DABCO) was obtained from TCI. All chemical were used immediately following purchase without further treatments. NMR spectra were recorded on JEOL 500 MHz for ^1H , and 125 MHz for ^{13}C in CDCl_3 . GC-MS was performed on Shimadzu GCMS QP 2100. The sample was diluted in acetone before analysis. Synthesis and characterization of tris(4-Methoxy phenyl)phenylsilane. Synthesis of tris(4-methoxyphenyl) phenylsilane was carried out at room temperature according to the literature [12] as follows: The solid reactants (i.e., palladium tertiary tributylphosphine 0.05 mmol) and base (i.e., potassium phosphate 5 mmol and 4-iodoanisole 3.5 mmol) were poured into a 100-ml Schlenk flask equipped with a magnetic stirrer. The flask was stirred for 30 min and vacuum processed for 15 min to remove co-existent gas. Then, 3 ml of tetrahydrofuran was added as a solvent. The reaction was monitored using a TLC plate for 1-2 h intervals. The reaction was quenched using 40 ml water when a new spot in the TLC plate appeared. The product was isolated using column chromatography on silica and ethyl acetate as eluent. Pure product was characterized using GC-MS, ^1H , and ^{13}C NMR spectroscopies.

3. Results and Discussion The coupling reaction between 4-iodoanisole with phenylsilane, presented in Fig. 1, was carried out at room temperature. The effect of temperature was not studied due to the volatility of the starting materials. According to Fig. 1, the formation of tris(4-methoxyphenyl)phenylsilane depends on the base, solvent, and reaction time. In the first trial of this research, the effect of several bases in the reaction of 4-iodoanisole with phenylsilane were studied. The results of the influence of base are shown in Table 1. Base is important for the reaction in Fig. 1 due to substitution of aryl iodide to the

organosilica moiety. Theoretical calculations revealed that at least a 3-mol or mmol base is needed to remove three hydrogens in the reaction. Thus, aryl iodide reacts with organosilica to form the product. Table 1 shows that there is no product Table 1. Optimization of Base Time (day) Base % Yield^a

Base	Time (day)	% Yield ^a
Triethylamine	7	4
Diisopropylethylamine	10	5
1,4-diazabicyclo[2,2,2]octane (DABCO)	35	35
1,2,2,2,6,6-pentamethylpiperidine	1	4
N-methyl piperidine	9	4
N-methyl morpholine	5	5
1,8-diazabicyclo[5,4,0]undec-7-en	0	3
Quinuclidine	14	5
2,6-lutidine	0	5
Pyridine	0	5
Potassium phosphate	Trace	5
Without base	0	0

^a % isolated yield b. % yield calculated from GC-MS data

Synthesis of Tris(4-Methoxyphenyl)Phenylsilane Using Phenylsilane Makara J. Sci. March 2014 | Vol. 18 | No. 115 in the absence of base (entry 12). These results indicated that the base is crucial in the coupling of aryl iodide with organosilica. Inorganic bases, such as potassium phosphate (entry 11), gave trace amounts of tris(4-methoxyphenyl)phenylsilane. Various organic bases that are available commercially were tested in this reaction (entries 1–10) and resulted in several product percentages. Aliphatic amines, such as triethylamine and diisopropylamine, generated product in the range of 7–10% (entries 1, 2). This percentage is still low for the catalytic reaction process, although this reaction type is difficult. Thus, by using cyclic, aromatic, and bicyclo in this research (entries 3–10), the scope of organic bases was enlarged. All cyclic, aromatic, and bicyclo used in this research contained nitrogen in their structures to react with hydrogen from silane in organosilica. Cyclic and aromatic bases, such as piperidine, quinuclidine, lutidine, and pyridine, generated coupling product of less than 14%. By using a bicyclo base compound, such as 1,4-diazabicyclo[2,2,2]octane (DABCO), the yield was slightly increased to 35%. DABCO was used as a base for the next optimization for the reaction. The reaction in entry 8 (quinuclidine) was stopped after 3 days, although the yield showed a slight increase. The formation of the product on the first and second days was similar and kinetically slow [16]. The characterization of the product was carried out using ¹H, ¹³C NMR, and GC-MS [17]. After the isolation process, the pure compound of tris(4-methoxyphenyl)phenylsilane was dissolved in chloroform (d-1). The ¹H NMR spectrum of tris(4-methoxyphenyl)phenylsilane is shown in Fig. 2. The ¹H NMR spectrum indicates 6 equivalent protons in the product at chemical shifts 3.82 ppm (9H); 6.92 ppm (d, 6H, J = 8.6 Hz); 7.35 ppm (q, 2H, J = 7.3 Hz); 7.40 ppm (m, 1H); 7.47 ppm (d, 6H, J = 8.6 Hz); and 7.53 ppm (q, 2H, J = 6.4 Hz). Six equivalent protons in the product were distributed as methoxy, aromatic methoxy substituent, and aromatic of organosilane. The results are consistent with the structure of the product shown in Fig. 1. The ¹³C NMR spectrum of the product is shown in Fig. 3. All the carbon in tris(4-methoxyphenyl)phenylsilane had 9 equivalents at chemical shifts 55.0 ppm (CH₃); 113.6 ppm (CH); 125.6 ppm (Cq); 127.7 ppm (CH); 129.3 ppm (CH); 135.4 ppm (Cq); 136.2 ppm (CH); 137.8 ppm (CH); and 160.7 ppm (Cq). Theoretical structural prediction reveals that the structure of tris(4-methoxyphenyl)phenylsilane, as shown in Figure 1, has 9 carbon equivalents. Thus, the results of ¹³C NMR are in accordance with the theoretical prediction. The characterization using the GC-MS-produced spectrum is shown in Fig. 4. The molecular ion peak of tris(4-methoxyphenyl)phenylsilane was at m/z 426. The calculation of molecular mass was 426, atomic mass number 12 for carbon, 1 for hydrogen, 16 for oxygen, and 28 for silica. The characterization data showed that compound tris(4-methoxyphenyl)phenylsilane is purely synthesized by coupling 4-iodoanisole with phenyl silane using palladium tertiary tributylphosphine as the catalyst. Various solvents (especially nonaqueous) are

widely available for organic synthesis [18]. These solvents are generally water-free, because the universal properties of water can destroy the phenylsilane functional group before it reacts with 4-iodoanisole [19]. However, oxygen atoms bound in the solvent (such as in the ether group) have special solvent properties. This study investigated several solvents in the ether group. Cyclic and aliphatic ether were used as separate solvents in the reaction between 4-iodoanisole and phenylsilane. Table 2 shows the percentage of coupling product obtained using DABCO as base in an ether-based solvent.

SiH₃COISiOCH₃OCH₃OCH₃+DABCO/THF, 5d[Pd(P(tBu)₃)₂] Figure 1. Coupling Reaction between 4-iodoanisole with Phenylsilane

Lesbani, et al. Makara J. Sci. March 2014 | Vol. 18 | No. 116 Figure 2. ¹H NMR Spectrum of Tris(4-methoxyphenyl)phenylsilane Figure 3. ¹³C NMR Spectrum of Tris(4-methoxyphenyl)phenylsilane Aliphatic ether (entries 2, 5), such as dimethoxyethane and diethyl ether, did not strongly affect the product yield. Cyclic ether, such as tetrahydrofuran, 1,4-dioxane, tetrahydropyran, and cyclopentyl methyl ether (entries 1, 3, 4, 6), affected the formation a small amount. Similar results were generated (shown in Table 1) when DABCO was used as the base. The effect of reaction time on the formation of tris(4-methoxyphenyl)phenylsilane from 4-iodoanisole and phenyl silane is shown in Table 3 using DABCO as a base and tetrahydrofuran as a solvent. The reaction time was prolonged to 11 days using dimethoxyethane, 1,4-dioxane, and tetrahydropyran as solvents, since the product formation remained very low after 5–6 days. The reaction occurred slowly, and after a 3-day reaction time, only 10% of the product was formed. The reaction time was extended to 5 days, and the amount of the product formed was similar to Table 1. After extending the reaction time to 6 days, there was a slight decrease in yield from 35% to 34%. In the catalytic and synthetic objects, this percentage was equal, so a 5-day reaction time for optimum formation of tris(4-methoxyphenyl)-phenylsilane was determined to be optimal.

Synthesis of Tris(4-Methoxyphenyl)Phenylsilane Using Phenylsilane Makara J. Sci. March 2014 | Vol. 18 | No. 117 Figure 4. Mass Spectrum of Tris(4-methoxyphenyl)phenylsilane Table 2.

Optimization of Solvent Time (day) Solvent % Yield

5 Tetrahydrofuran 35 11

Dimethoxyethane 8 11 1,4-dioxane 10 11 Tetrahydropyran 18 6 Diethyl ether 13 5

Cyclopentyl methyl ether 20 a. % isolated yield Table 3. Optimization of Reaction Time

Time (day) % Yield 3 10 5 35 6 34 a. % isolated yield

4. Conclusions The coupling reaction between phenylsilane and 4-iodoanisole using palladium tertiary tributylphosphine yielded tris(4-methoxyphenyl)phenylsilane at 35% by optimization of the base, solvent, and reaction time.

The maximum yield was obtained using DABCO as the base, THF as the solvent, and a 5-day reaction time.

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Synthesis of Tris(4-Methoxyphenyl)Phenylsilane Using Phenylsilane and 4-Iodo Anisole Catalyzed by Palladium Complex

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Abstract

The synthesis of tris(4-methoxyphenyl)phenylsilane through the coupling reaction of phenylsilane and 4-iodoanisole using palladium tertiary tributylphosphine as a catalyst was carried out through the determination of the base, solvent, and reaction time. The results showed that tris(4-methoxyphenyl)phenylsilane can be synthesized to form maximum yield using 1,4-diazabicyclo[2,2,2]octane (DABCO) as a base, tetrahydrofuran as a solvent, and a 5-day reaction time. The highest yield of tris(4-methoxyphenyl)phenylsilane was 35%.

Abstrak

Sintesis Tris(4-Metoksifenil)Fenilsilan Menggunakan Fenilsilan dan 4-Iodida Anisol dengan Katalis Senyawa Kompleks Paladium. Sintesis tris(4-metoksifenil)fenilsilan melalui reaksi kopling antara fenilsilan dan 4-iodida anisol menggunakan senyawa paladium tersier tributilfosfin sebagai katalis telah dilakukan berdasarkan penentuan senyawa basa, pelarut, dan waktu reaksi kopling. Hasil penelitian menunjukkan bahwa senyawa tris(4-metoksifenil)fenilsilan dapat disintesis menghasilkan persentase produk maksimum pada penggunaan senyawa 1,4-diazabisiko[2,2,2]oktan (DABCO) sebagai basa, tetrahidrofuran sebagai pelarut, dan waktu reaksi kopling selama 5 hari. Persentase tertinggi produk senyawa tris(4-metoksifenil)fenilsilan adalah 35%.

Keywords: 4-iodoanisole, palladium, phenylsilane, tris(4-methoxyphenyl)phenylsilane

1. Introduction

The coupling reaction is one of the most useful techniques for synthesizing valuable chemical materials [1]. This reaction has been developed in couplings such as Suzuki-Miyaura, Negishi, Heck, and Sonogashira. Prof Suzuki and Prof. Negishi won the Nobel Prize in 2010 for research of palladium as catalyst for coupling reaction in organic synthesis [2]. The coupling reaction occurs when a carbon-carbon chain is combined using a powerful catalyst such as a complex of transition metals [3]. The example this reaction is asymmetric synthesis of drugs with chirality in the compound [4], which is still developed until now.

The development of the coupling reaction has included various substrates depending on molecule targets. Previous research using trialkylsilane and aryl halide

as reactants was carried out by Manoso *et al.* [5], Denmark *et al.* [6], and Murata *et al.* [7]. This reaction included coupling aryl halide with organosilane. Although the coupling reaction has been done using several materials, it is limited due to by-products formed by aryl halide. Yamanoi *et al.* [8] also found by-products when using aryl halide as a reactant. By changing a halide of chloride or bromide to iodide, the by-product percentage may be significantly reduced. Thus, aryl iodide is an appropriate substrate for coupling with organosilica [9]. Lesbani *et al.* applied the coupling reaction of aryl iodide for the synthesis of pesticides [10]. The synthesis of materials with unique photoluminescence was also reported by Lesbani *et al.*, who coupled aryl halide with bulky organosilica [11]. Its unique photoluminescence properties include a violet-blue color under irradiation of UV at 365 nm and a quantum yield of 0.55. Furthermore, Lesbani *et al.*

enlarged the scope of coupling reactions by using not only aryl iodides [12–13] but also organosilicas [14–15] as starting materials. These researchers found that, although the various aryl iodides and organosilica can be reacted using palladium catalyst through the coupling reaction, there was a slight increase in the yield. Therefore, the optimization of the coupling reaction in this case is vital to increase the coupling product yield.

This present research examined the optimization of reactions of phenylsilane and 4-iodoanisole to tris(4-methoxyphenyl) phenylsilane by evaluating bases, solvents, and reaction times. The tris(4-methoxyphenyl) phenylsilane was synthesized using 1,4-diazabicyclo[2, 2,2]octane (DABCO) as a base and tetrahydrofuran as a solvent, and a 5-day reaction time.

2. Methods

Materials and equipment. All experiments were carried out using Schlenk techniques under an argon or nitrogen atmosphere in a vacuum system. Chemicals 4-iodoanisole, palladium tertiary tributylphosphine, diethyl ether, tetrahydrofuran, dimethoxyethane, 1,4-dioxane, tetrahydropyran, triethylamine, diisopropylethylamine, potassium phosphate, TLC plate, ethyl acetate, and dichloromethane were obtained from Merck. 1,2,2,6,6-Pentamethylpiperidine, N-methylpiperidine, N-methylmorpholine, 1,8-diazabicyclo[5,4,0]undec-7en, quinuclidine, 2,6-lutidine, cyclopentyl methyl ether, and pyridine were obtained from Aldrich, and 1,4-diazabicyclo-[2,2,2]octane (DABCO) was obtained from TCI. All chemical were used immediately following purchase without further treatments. NMR spectra were recorded on JEOL 500 MHz for ^1H , and 125 MHz for ^{13}C in CDCl_3 . GC-MS was performed on Shimadzu GCMS QP 2100. The sample was diluted in acetone before analysis.

Synthesis and characterization of tris(4-Methoxyphenyl)phenylsilane. Synthesis of tris(4-methoxyphenyl) phenylsilane was carried out at room temperature according to the literature [12] as follows: The solid reactants (i.e., palladium tertiary tributylphosphine 0.05 mmol) and base (i.e., potassium phosphate 5 mmol and 4-iodoanisole 3.5 mmol) were poured into a 100-ml Schlenk flask equipped with a magnetic stirrer. The flask was stirred for 30 min and vacuum processed for 15 min to remove co-existent gas. Then, 3 ml of tetrahydrofuran was added as a solvent. The reaction was monitored using a TLC plate for 1-2 h intervals. The reaction was quenched using 10 ml water when a new spot in the TLC plate appeared. The product was isolated using column chromatography on silica and ethyl acetate as eluent. Pure product was characterized using GC-MS, ^1H , and ^{13}C NMR spectroscopies.

3. Results and Discussion

The coupling reaction between 4-iodoanisole with phenylsilane, presented in Fig. 1, was carried out at room temperature. The effect of temperature was not studied due to the volatility of the starting materials. According to Fig. 1, the formation of tris(4-methoxyphenyl)phenylsilane depends on the base, solvent, and reaction time. In the first trial of this research, the effect of several bases in the reaction of 4-iodoanisole with phenylsilane were studied. The results of the influence of base are shown in Table 1.

Base is important for the reaction in Fig. 1 due to substitution of aryl iodide to the organosilica moiety. Theoretical calculations revealed that at least a 3-mol or mmol base is needed to remove three hydrogens in the reaction. Thus, aryl iodide reacts with organosilica to form the product. Table 1 shows that there is no product

Table 1. Optimization of Base

Time (day)	Base	% Yield ^a
4	Triethylamine	7
4	Diisopropylethylamine	10
5	1,4-diazabicyclo[2,2,2]octane (DABCO)	35
5	1,2,2,2,6,6-pentamethylpiperidine	1
4	N-methyl piperidine	9
4	N-methyl morpholine	5
5	1,8-diazabicyclo[5,4,0]undec-7en	0 ^b
3	Quinuclidine	14
5	2,6-lutidine	0 ^b
5	Pyridine	0 ^b
5	Potassium phosphate	Trace ^b
5	Without base	0 ^b

a. % isolated yield

b. % yield calculated from GC-MS data

in the absence of base (entry 12). These results indicated that the base is crucial in the coupling of aryl iodide with organosilica. Inorganic bases, such as potassium phosphate (entry 11), gave trace amounts of tris(4-methoxyphenyl)phenylsilane. Various organic bases that are available commercially were tested in this reaction (entries 1–10) and resulted in several product percentages. Aliphatic amines, such as triethylamine and diisopropylamine, generated product in the range of 7–10% (entries 1, 2). This percentage is still low for the catalytic reaction process, although this reaction type is difficult. Thus, by using cyclic, aromatic, and bicyclo in this research (entries 3–10), the scope of organic bases was enlarged. All cyclic, aromatic, and bicyclo used in this research contained nitrogen in their structures to react with hydrogen from silane in organosilica. Cyclic and aromatic bases, such as piperidine, quinuclidine, lutidine, and pyridine, generated coupling product of less than 14%. By using a bicyclo base compound, such as 1,4-diazabicyclo[2,2,2]octane (DABCO), the yield was slightly increased to 35%. DABCO was used as a base for the next optimization for the reaction. The reaction in entry 8 (quinuclidine) was stopped after 3 days, although the yield showed a slight increase. The formation of the product on the first and second days was similar and kinetically slow [16].

The characterization of the product was carried out using ^1H , ^{13}C NMR, and GC-MS [17]. After the isolation process, the pure compound of tris(4-methoxyphenyl)phenylsilane was dissolved in chloroform (*d*-1). The ^1H NMR spectrum of tris(4-methoxyphenyl)phenylsilane is shown in Fig. 2. The ^1H NMR spectrum indicated 6 equivalent protons in the product at chemical shifts 3.82 ppm (s, 9H); 6.92 ppm (d, 6H, $J = 8.6$ Hz); 7.35 ppm (q, 2H, $J = 7.3$ Hz); 7.40 ppm (m, 1H); 7.47 ppm (d, 6H, $J = 8.6$ Hz); and 7.53 ppm (q, 2H, $J = 6.4$ Hz). Six equivalent protons in the product were distributed as

methoxy, aromatic methoxy substituent, and aromatic of organosilane. The results are consistent with the structure of the product shown in Fig. 1. The ^{13}C NMR spectrum of the product is shown in Fig. 3. All the carbon in tris(4-methoxyphenyl)phenylsilane had 9 equivalents at chemical shifts 55.0 ppm (CH_3); 113.6 ppm (CH); 125.6 ppm (Cq); 127.7 ppm (CH); 129.3 ppm (CH); 135.4 ppm (Cq); 136.2 ppm (CH); 137.8 ppm (CH); and 160.7 ppm (Cq). Theoretical structural prediction reveals that the structure of tris(4-methoxyphenyl)phenylsilane, as shown in Figure 1, has 9 carbon equivalents. Thus, the results of ^{13}C NMR are in accordance with the theoretical prediction.

The characterization using the GC-MS-produced spectrum is shown in Fig. 4. The molecular ion peak of tris(4-methoxyphenyl)phenylsilane was at m/z 426. The calculation of molecular mass was 426, atomic mass number 12 for carbon, 1 for hydrogen, 16 for oxygen, and 28 for silica. The characterization data showed that compound tris(4-methoxyphenyl)phenylsilane is purely synthesized by coupling 4-iodoanisole with phenyl silane using palladium tertiary tributylphosphine as the catalyst.

Various solvents (especially nonaqueous) are widely available for organic synthesis [18]. These solvents are generally water-free, because the universal properties of water can destroy the phenylsilane functional group before it reacts with 4-iodoanisole [19]. However, oxygen atoms bound in the solvent (such as in the ether group) have special solvent properties. This study investigated several solvents in the ether group. Cyclic and aliphatic ether were used as separate solvents in the reaction between 4-iodoanisole and phenylsilane. Table 2 shows the percentage of coupling product obtained using DABCO as base in an ether-based solvent.

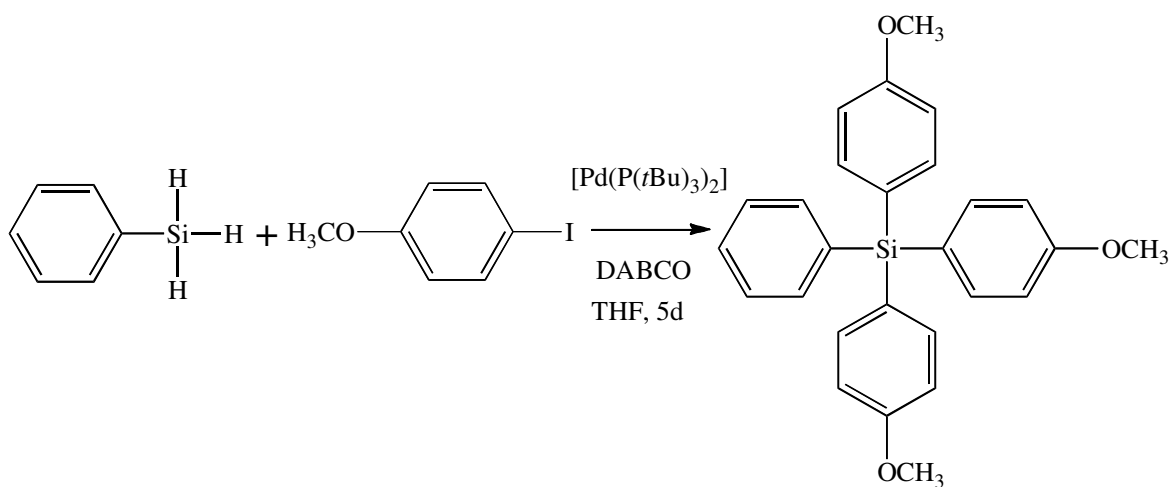


Figure 1. Coupling Reaction between 4-iodoanisole with Phenylsilane

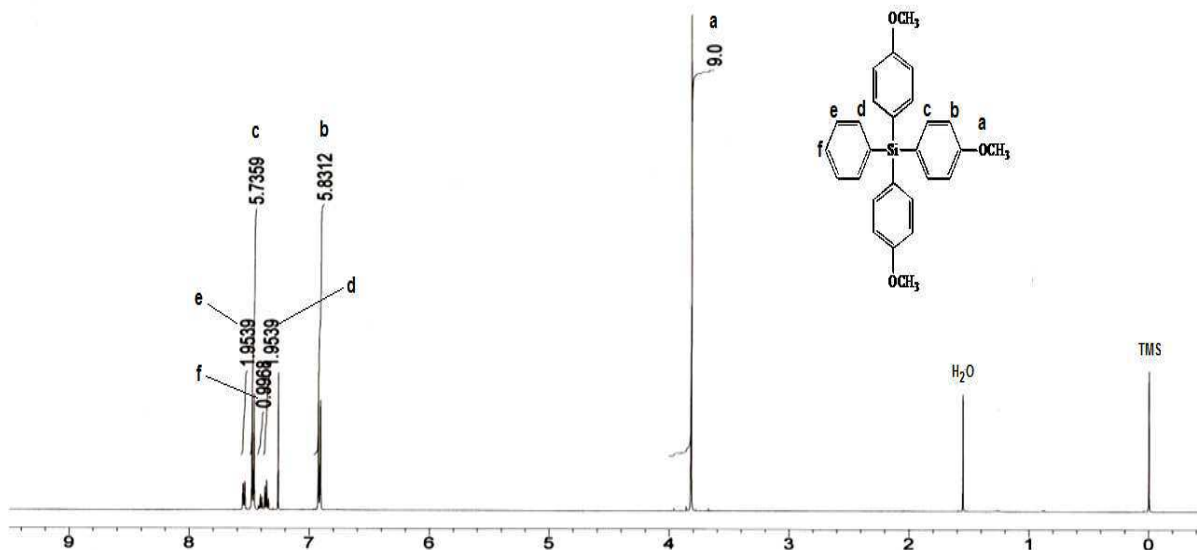


Figure 2. ¹H NMR Spectrum of Tris(4-methoxyphenyl)phenylsilane

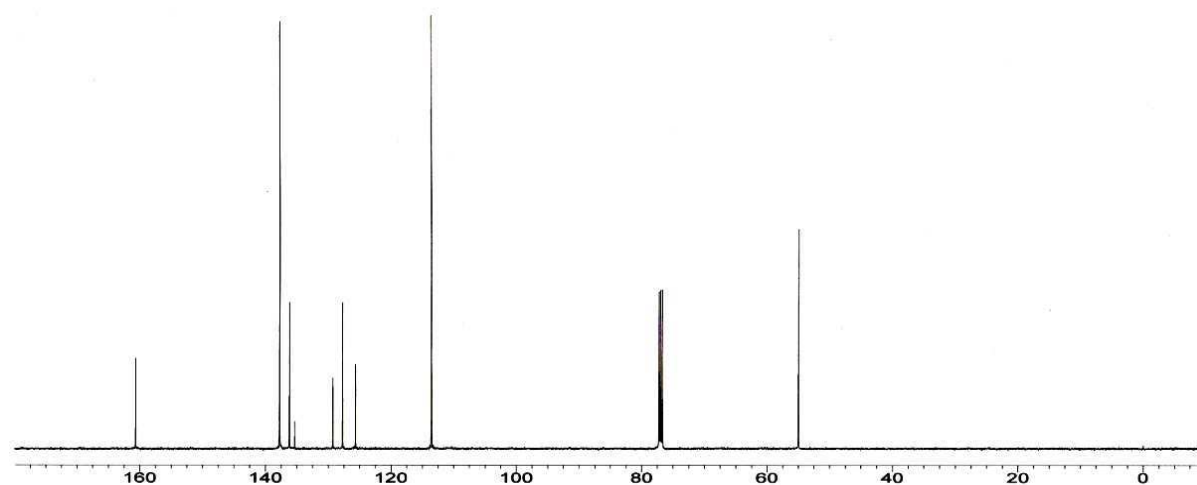


Figure 3. ¹³C NMR Spectrum of Tris(4-methoxyphenyl)phenylsilane

Aliphatic ether (entries 2, 5), such as dimethoxyethane and diethyl ether, did not strongly affect the product yield. Cyclic ether, such as tetrahydrofuran, 1,4-dioxane, tetrahydropyran, and cyclopentyl methyl ether (entries 1,3,4, 6), affected the formation a small amount. Similar results were generated (shown in Table 1) when DABCO was used as the base. The effect of reaction time on the formation of tris(4-methoxyphenyl)phenylsilane from 4-iodoanisole and phenyl silane is shown in Table 3 using DABCO as a base and tetrahydrofuran as a solvent. The reaction time was prolonged to 11 days using dimethoxyethane, 1,4-dioxane, and tetrahydropyran

as solvents, since the product formation remained very low after 5–6 days.

The reaction occurred slowly, and after a 3-day reaction time, only 10% of the product was formed. The reaction time was extended to 5 days, and the amount of the product formed was similar to Table 1. After extending the reaction time to 6 days, there was a slight decrease in yield from 35% to 34%. In the catalytic and synthetic objects, this percentage was equal, so a 5-day reaction time for optimum formation of tris(4-methoxyphenyl)phenylsilane was determined to be optimal.

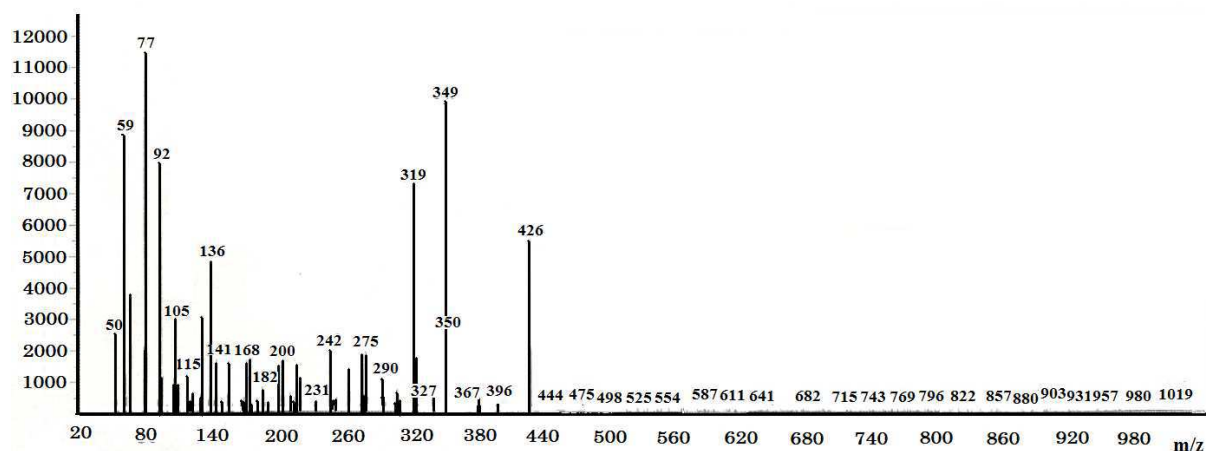


Figure 4. Mass Spectrum of Tris(4-methoxyphenyl)phenylsilane

Table 2. Optimization of Solvent

Time (day)	Solvent	% Yield ^a
5	Tetrahydrofuran	35
11	Dimethoxyethane	8
11	1,4-dioxane	10
11	Tetrahydropyran	18
6	Diethyl ether	13
5	Cyclopentyl methyl ether	20

a. % isolated yield

Table 3. Optimization of Reaction Time

Time (day)	% Yield
3	10
5	35
6	34

a. % isolated yield

4. Conclusions

The coupling reaction between phenylsilane and 4-iodoanisole using palladium tertiary tributylphosphine yielded tris(4-methoxyphenyl)phenylsilane at 35% by optimization of the base, solvent, and reaction time. The maximum yield was obtained using DABCO as the base, THF as the solvent, and a 5-day reaction time.

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