

Process for the Synthesis of Benzo[b]thiophenes

The present invention is directed to new processes for the synthesis of benzo[b]thiophenes, in particular 2-aryl-benzo[b]thiophenes.

Benzo[b]thiophenes have been prepared by a number of different synthetic routes. One of the most widely used methods is the oxidative cyclization of o-mercaptocinnamic acids. This route is limited to the preparation of benzo[b]-thiophene-2-carboxylates. 2-Phenylbenzo[b]thiophenes are prepared by acid-catalyzed cyclization of 2-phenylthioacetaldehyde dialkyl acetals. Unsubstituted benzo[b]thiophenes are prepared by catalytic condensation of styrene and sulfur. 3-Substituted benzo[b]thiophenes are prepared by acid-catalyzed cyclization of arylthiomethyl ketones; however, this route is limited to the preparation of 3-alkylbenzo[b]thiophenes. See Campaigne, "Thiophenes and their Benzo Derivatives: (iii) Synthesis and Applications," in **Comprehensive Heterocyclic Chemistry** (Katritzky and Rees, eds.), Volume IV, Part III, 863-934 (1984). 3-Chloro-2-phenylbenzo[b]thiophene is prepared by the reaction of diphenylacetylene with sulfur dichloride. Barton and Zika, *J. Org. Chem.*, **35**, 1729-1733 (1970). Benzo[b]thiophenes have also been prepared by pyrolysis of styryl sulfoxides. However, low yields and extremely high temperatures make this route unsuitable for production-scale syntheses. See Ando, *J. Chem. Soc., Chem. Comm.*, 704-705 (1975).

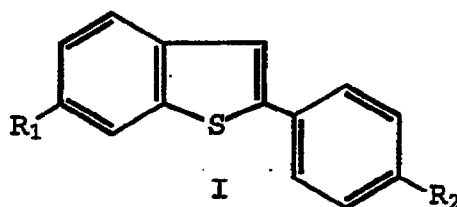
One process of the present invention for preparing benzo[b]thiophenes requires an intermediate sulfenic acid derivative. Sulfenic acids have been postulated as key intermediates in a variety of chemical reactions; however, very few examples exist of the isolation of these compounds. See Shelton and Davis, *J. Am. Chem. Soc.*, **89**(3), 718-719 (1968) and Davis et al., *J. Am. Chem. Soc.*, **100**, 2844 (1978). Sulfenic acids have been generated *in situ*, and intramolecularly or intermolecularly cyclized with olefins and acetylenes. See Mazzanti et al., *J. Chem. Soc., Perkin*

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Trans. I, 3299-3004 (1944) and Davis et al., *J. Org. Chem.*, **45**, 1650-1653 (1980). A series of trimethylsilyl arenesulfenates have been prepared from the corresponding *N*-benzylidenearenesulfinamides; however, the yield of the
5 trimethylsilyl ester was generally very low. Davis et al., *J. Org. Chem.*, **45**, 1650-1653 (1980).

The preparation of 6-hydroxy-2-(4-hydroxyphenyl)benzo-
[b]thiophenes was described in U.S. Patent Nos. 4,133,814 and
4,380,635. One process described in these patents is the
10 acid-catalyzed intramolecular cyclization/rearrangement of
 α -(3-methoxyphenylthio)-4-methoxyacetophenone. The reaction
of this starting compound in neat polyphosphoric acid at
about 85°C to about 90°C gives an approximate 3:1 mixture of
two regioisomeric products: 6-methoxy-2-(4-methoxyphenyl)-
15 benzo[b]thiophene and 4-methoxy-2-(4-methoxyphenyl)benzo[b]-
thiophene. These isomeric benzo[b]thiophenes co-precipitate
from the reaction mixture, producing a mixture containing
both compounds. To obtain a single regioisomer, the
regioisomers must be separated, such as by chromatography or
20 fractional crystallization. Therefore, there currently
exists a need for an efficient and regiospecific synthesis of
2-arylbenzo[b]thiophenes from readily available starting
materials.

25 The present invention is directed to processes for the
synthesis of benzo[b]thiophenes. Specifically, the present
invention is directed to a process for preparing a compound
of the formula



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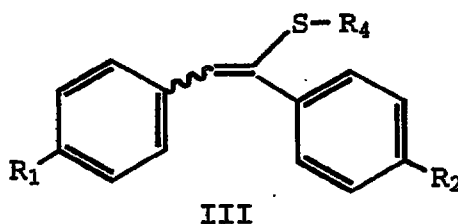
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wherein:

R₁ is hydrogen, C₁-C₄ alkoxy, arylalkoxy, halo, or amino;
and

R₂ is hydrogen, C₁-C₄ alkoxy, arylalkoxy, halo, or amino;

5 which comprises cyclizing in the presence of an acid catalyst
a compound of the formula



10 wherein:

R₁ and R₂ are as defined above;

R₄ is OSi(R)₃, NR₅R₆, or SR₈;

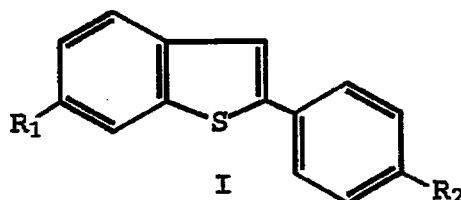
each R is independently C₁-C₆ alkyl, aryl, or arylalkyl;

R₅ and R₆ are independently hydrogen, C₁-C₆ alkyl,

15 arylalkyl, or aryl, or R₅ and R₆ together with the nitrogen
atom form a ring selected from piperidine, pyrrolidine,
morpholine, or hexamethylimine; and

R₈ is C₁-C₆ alkyl, aryl, or arylalkyl.

Another aspect of the present invention is a second
20 process for the synthesis of benzo[b]thiophenes. Specifically,
the present invention is directed to a process for preparing a
compound of the formula



25 wherein:

R₁ is hydrogen, C₁-C₄ alkoxy, arylalkoxy, halo, amino;

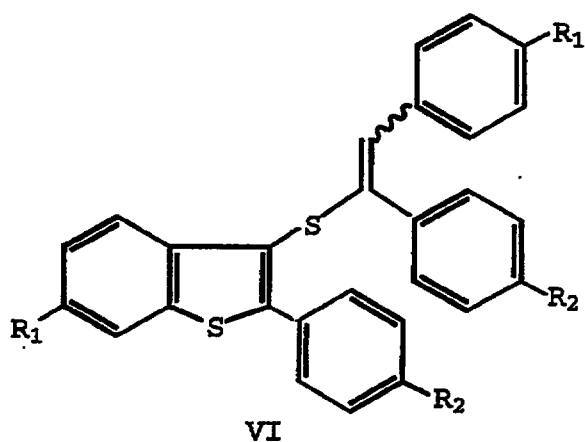
and

R₂ is hydrogen, C₁-C₄ alkoxy, arylalkoxy, halo, amino;

which comprising treating a compound of the formula

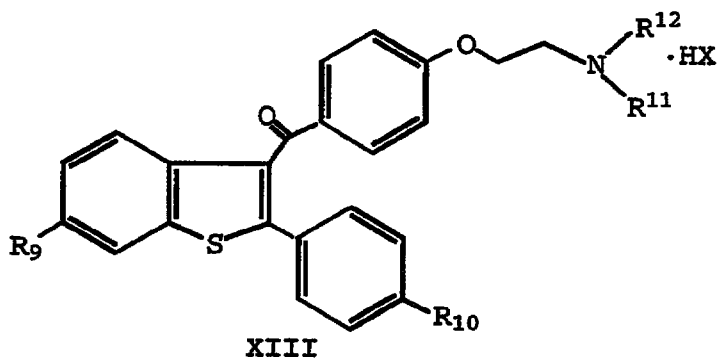
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wherein R₁ and R₂ are as defined above, with an acid catalyst.
 The present invention is also directed to the formula VI
 5 compounds, as well as, processes for their preparation.

Another aspect of the present invention is a process for
 the synthesis of a compound of the formula



10

wherein:

R₉ is hydrogen, halo, amino, or hydroxyl;

R₁₀ is hydrogen, halo, amino, or hydroxyl;

R₁₁ and R₁₂ are independently C₁-C₄ alkyl, or R₁₁ and R₁₂
 15 together with the adjacent nitrogen atom form a heterocyclic
 ring selected from the group consisting of pyrrolidino,
 piperidino, hexamethyleneimino, and morpholino; and

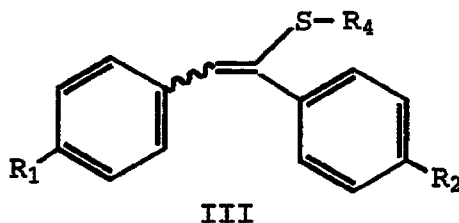
HX is HCl or HBr;

comprising the steps of:

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(a) cyclizing in the presence of an acid catalyst a compound of the formula



5 wherein:

R_1 is hydrogen, C_1 - C_4 alkoxy, arylalkoxy, halo, or amino;

R_2 is hydrogen, C_1 - C_4 alkoxy, arylalkoxy, halo, or amino;

and

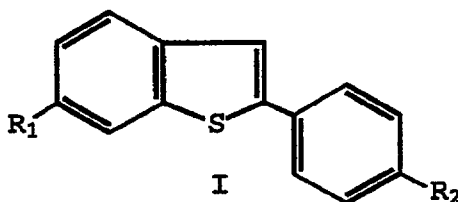
R_4 is $OSi(R)_3$, NR_5R_6 , or SR_8 ;

10 each R is independently C_1 - C_6 alkyl, aryl, or arylalkyl;

R_5 and R_6 are independently hydrogen, C_1 - C_6 alkyl, or aryl, or R_5 and R_6 together with the nitrogen atom form a ring selected from piperidine, pyrrolidine, morpholine, and hexamethylimine; and

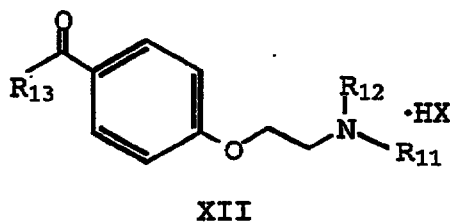
15 R_8 is C_1 - C_6 alkyl, aryl, or arylalkyl;

to prepare a benzothiophene compound of the formula



20 wherein R_1 and R_2 are as defined above;

(b) acylating said benzothiophene compound with an acylating agent of the formula



25

wherein:

R₁₁, R₁₂, and HX are as defined previously; and

R₁₃ is chloro, bromo, or hydroxyl; in the presence of BX'₃, wherein X' is chloro or bromo;

5

(c) when R₁ and/or R₂ is C₁-C₄ alkoxy or arylalkoxy, dealkylating one or more phenolic groups of the acylation product of step (b) by reacting with additional BX'₃, wherein X' is as defined above; and

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(d) isolating the formula XIII compound;

The term "acid catalyst" represents a Lewis acid or a Brønsted acid. Representative Lewis acids are zinc chloride, zinc iodide, aluminum chloride, and aluminum bromide. Representative Brønsted acids include: inorganic acids, such as sulfuric and phosphoric acids; carboxylic acids, such as acetic and trifluoroacetic acids; sulfonic acids, such as methanesulfonic, benzenesulfonic, 1-naphthalenesulfonic, 1-butanesulfonic, ethanesulfonic, 4-ethylbenzenesulfonic, 1-hexanesulfonic, 1,5-naphthalenedisulfonic, 1-octanesulfonic, camphorsulfonic, trifluoromethanesulfonic, and *p*-toluenesulfonic acids; and polymeric arylsulfonic acids, such as Nafion®, Amberlyst®, or Amberlite®. The preferred acids for use in catalyzing the processes of the present invention are sulfonic or polymeric sulfonic acids. More preferably, the acid catalysts are sulfonic acids, such as methanesulfonic acid, benzenesulfonic acid, camphorsulfonic acid, and *p*-toluenesulfonic acid. The most preferred acid catalyst is *p*-toluenesulfonic acid.

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In the above formula, the term "C₁-C₄ alkoxy" represents groups such as methoxy, ethoxy, *n*-propoxy, isopropoxy, *n*-butoxy, *t*-butoxy, and like groups. The term "halo" refers to fluoro, chloro, bromo, or iodo groups.

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The term "C₁-C₆ alkyl" represents a straight or branched alkyl chain having from one to six carbon atoms. Typical C₁-C₆ alkyl groups include methyl, ethyl, *n*-propyl, isopropyl,

n-butyl, isobutyl, *sec*-butyl, *t*-butyl, *n*-pentyl, isopentyl, *n*-hexyl, 2-methylpentyl, and the like. The term "C₁-C₄ alkyl" represents a straight or branched alkyl chain having from one to four carbon atoms, and includes methyl, ethyl, *n*-propyl, isopropyl, *n*-butyl, *sec*-butyl, *i*-butyl, and *t*-butyl.

The term "aryl" represents groups such as phenyl and substituted phenyl. The term "substituted phenyl" represents a phenyl group substituted with one or more moieties chosen from the group consisting of halo, hydroxy, nitro, C₁-C₄ alkyl, C₁-C₄ alkoxy, trichloromethyl, and trifluoromethyl. Examples of a substituted phenyl group include 4-chlorophenyl, 2,6-dichlorophenyl, 2,5-dichlorophenyl, 3,4-dichlorophenyl, 3-chlorophenyl, 3-bromophenyl, 4-bromophenyl, 3,4-dibromophenyl, 3-chloro-4-fluorophenyl, 2-fluorophenyl, 4-hydroxyphenyl, 3-hydroxyphenyl, 2,4-dihydroxyphenyl, 3-nitrophenyl, 4-nitrophenyl, 2,4-dinitrophenyl, 4-methylphenyl, 4-ethylphenyl, 4-methoxyphenyl, 4-propylphenyl, 4-*n*-butylphenyl, 4-*t*-butylphenyl, 3-fluoro-2-methylphenyl, 2,3-difluorophenyl, 2,6-difluorophenyl, 2,6-dimethylphenyl, 2-fluoro-5-methylphenyl, 2,4,6-trifluorophenyl, 2-trifluoromethylphenyl, 2-chloro-5-trifluoromethylphenyl, 3,5-bis-(trifluoromethyl)phenyl, 2-methoxyphenyl, 3-methoxyphenyl, 3,5-dimethoxyphenyl, 4-hydroxy-3-methylphenyl, 3,5-dimethyl-4-hydroxyphenyl, 2-methyl-4-nitrophenyl, 4-methoxy-2-nitrophenyl, and the like.

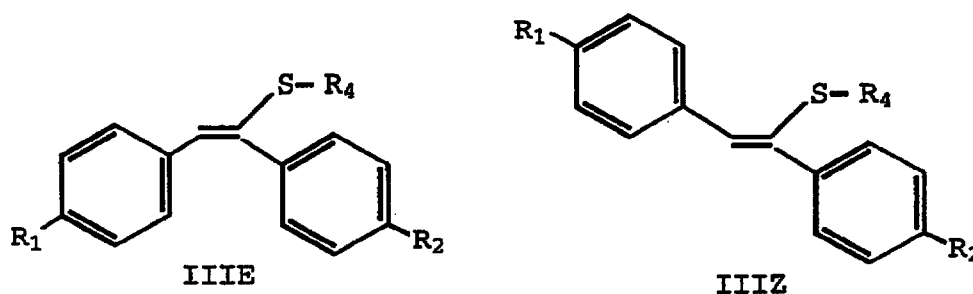
The term "arylalkyl" represents a C₁-C₄ alkyl group bearing one or more aryl groups. Representatives of this group include benzyl, *o*-nitrobenzyl, *p*-nitrobenzyl, *p*-halobenzyl (such as *p*-chlorobenzyl, *p*-bromobenzyl, *p*-iodobenzyl), 1-phenylethyl, 2-phenylethyl, 3-phenylpropyl, 4-phenylbutyl, 2-methyl-2-phenylpropyl, (2,6-dichlorophenyl)methyl, bis(2,6-dichlorophenyl)methyl, (4-hydroxyphenyl)methyl, (2,4-dinitrophenyl)methyl, diphenylmethyl, triphenylmethyl, (*p*-methoxyphenyl)-diphenylmethyl, bis(*p*-methoxyphenyl)methyl, bis(2-nitrophenyl)methyl, and the like.

The term "arylalkoxy" represents a C₁-C₄ alkoxy group bearing one or more aryl groups. Representatives of this group include benzyloxy, o-nitrobenzyloxy, p-nitrobenzyloxy, p-halobenzyloxy (such as p-chlorobenzyloxy, p-bromobenzyloxy, p-iodobenzyloxy), 1-phenylethoxy, 2-phenylethoxy, 3-phenylpropoxy, 4-phenylbutoxy, 2-methyl-2-phenylpropoxy, (2,6-dichlorophenyl)methoxy, bis(2,6-dichlorophenyl)methoxy, (4-hydroxyphenyl)methoxy, (2,4-dinitrophenyl)methoxy, diphenylmethoxy, triphenylmethoxy, (p-methoxyphenyl)-diphenylmethoxy, bis(p-methoxyphenyl)methoxy, bis(2-nitrophenyl)methoxy, and the like.

The term "thermally-labile or acid-labile C₂-C₁₀ alkyl, C₄-C₁₀ alkenyl, or aryl(C₁-C₁₀ alkyl) group" represents a group that is readily removed from the sulfoxide (SO) group under heating or by treatment with the acid catalyst. The thermally-labile or acid-labile C₂-C₁₀ alkyl groups are straight or branched alkyl chains having from two to ten carbon atoms and having at least one beta-hydrogen atom. Representative thermally-labile or acid-labile C₂-C₁₀ alkyl groups include ethyl, n-propyl, i-propyl, 1,1-dimethylpropyl, n-butyl, sec-butyl, t-butyl, 1,1-dimethylbutyl, 2-methylbutyl, 3-methylbutyl, 1-methylbutyl, 1,2-dimethylbutyl, 1,3-dimethylbutyl, 2,4-dimethylbutyl, 3,3-dimethylbutyl, n-pentyl, 1-methylpentyl, 2-methylpentyl, 3-methylpentyl, 4-methylpentyl, n-hexyl, and the like. The thermally-labile or acid-labile C₄-C₁₀ alkenyl groups are straight or branched alkenyl chains having from four to ten carbon atoms, at least one site of unsaturation, and either a beta-hydrogen or delta-hydrogen atom. Representative thermally-labile or acid-labile C₄-C₁₀ alkenyl groups include 2-butenyl, 3-butenyl, 2-methyl-2-butenyl, 3-methyl-2-butenyl, 2-methyl-3-butenyl, 2-pentenyl, 3-pentenyl, 4-pentenyl, 2-methyl-2-pentenyl, 3-methyl-2-pentenyl, 4-methyl-2-pentenyl, 2-methyl-3-pentenyl, 3-methyl-3-pentenyl, 4-methyl-3-pentenyl, 2-methyl-4-pentenyl, 3-methyl-4-pentenyl, 4-methyl-4-pentenyl, 2-hexenyl, 3-hexenyl, 4-hexenyl, 5-hexenyl, and the like. The term thermally-labile or acid-labile aryl(C₁-

C₁₀ alkyl) represents thermally-labile or acid-labile C₂-C₁₀ alkyl groups additionally containing one or more aryl groups and aryl-substituted methyl groups. Representative aryl(C₁-C₁₀ alkyl) groups include benzyl, diphenylmethyl, triphenylmethyl, *p*-methoxybenzyl, 2-phenylethyl, 2-phenylpropyl, 3-phenylpropyl, and the like.

The formula III compounds exist in two regioisomeric forms: the **E** isomer and the **Z** isomer. The process of the present invention uses individually the **E** and **Z** isomers, or mixtures thereof, of the formula III compounds. These **E** and **Z** regioisomers are represented by the following structures:



15

One group of compounds that are useful in the processes of the present invention are sulfenate silyl esters. In particular, the formula III compounds, where R₄ is OSi(R)₃ and each R is independently C₁-C₆ alkyl, aryl, or arylalkyl, and the formula IV compounds are silyl esters of sulfenic acids. The preferred sulfenate silyl esters are abbreviated using nomenclature well recognized in the chemical arts, as shown in the following table.

20

Table 1

	<u>abbreviation</u>	<u>silyl group</u>
	TMS	trimethylsilyl
	TES	triethylsilyl
5	TIPS	triisopropylsilyl
	DMIPS	dimethylisopropylsilyl
	DEIPS	diethylisopropylsilyl
	TDS	dimethylhexylsilyl
	TBDMS	t-butyldimethylsilyl
10	TBDPS	t-butyldiphenylsilyl
	TBS	tribenzylsilyl
	TPS	triphenylsilyl
	DPMS	diphenylmethylsilyl
	<u>TBMPs</u>	<u>t-butyldi(methoxyphenyl)silyl</u>

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The term "silylating reagent" represents a compound, or a combination of compounds, used to convert the intermediate sulfenic acid to a sulfenate silyl ester. Representative silylating reagents include bis(trialkylsilyl)ureas, such as

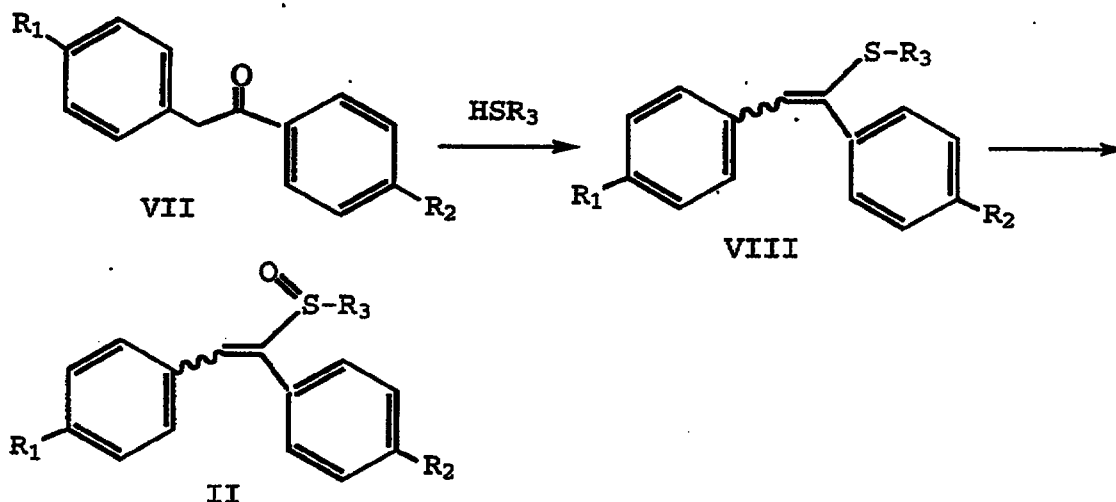
20 1,3-bis(trimethylsilyl)urea, 1,3-bis(triethylsilyl)urea, 1,3-bis(dimethylisopropylsilyl)urea, 1,3-bis(triisopropylsilyl)urea, 1,3-bis(diethylisopropylsilyl)urea, 1,3-bis(dimethylhexylsilyl)urea, and 1,3-bis(t-butyldimethylsilyl)urea; bis(triaryl)silyl)ureas, such as 1,3-bis-

25 (triphenylsilyl)urea; bis(diarylalkylsilyl)ureas, such as 1,3-bis(diphenylmethylsilyl)urea and 1,3-bis(t-butyldiphenylsilyl)urea; and hexaalkyldisilylzanones, such as hexamethyldisilylthane; or combination of a hexaalkyldisilylthane and a catalytic amount of a chlorotrialkylsilane,

30 such as chlorotrimethylsilane.

The starting compounds for the processes of the present invention can be prepared by a number of routes. One method for preparing the formula II compounds is shown in Scheme 1.

Scheme 1



5 Generally, a formula VII compound is converted to a styryl sulfide by reaction with a mercaptan of the formula HSR_3 in the presence of a Lewis acid. The formula VIII compound is then oxidized to a styryl sulfoxide, a compound of formula II compound.

10 More specifically, a formula VII compound, wherein R_1 and R_2 are as defined above, is treated with a Lewis acid, such as titanium(IV) chloride. This reaction is carried out in an anhydrous organic solvent, such as dry tetrahydrofuran, at a temperature of about 0°C to about 35°C . After about 15

15 minutes to about one hour, the reaction mixture is treated with an amine base and a mercaptan of the formula HSR_3 , where R_3 is a thermally-labile or acid-labile C_1 - C_{10} alkyl, C_4 - C_{10} alkenyl, or aryl(C_1 - C_{10} alkyl) group. Preferably, the mercaptan and amine base are added as a solution in the

20 reaction solvent. A representative amine base is triethylamine. After the addition of the mercaptan and amine base, the reaction is generally heated to a temperature of about 35°C to about 65°C , preferably at about 50°C . The products of this reaction can be purified using techniques

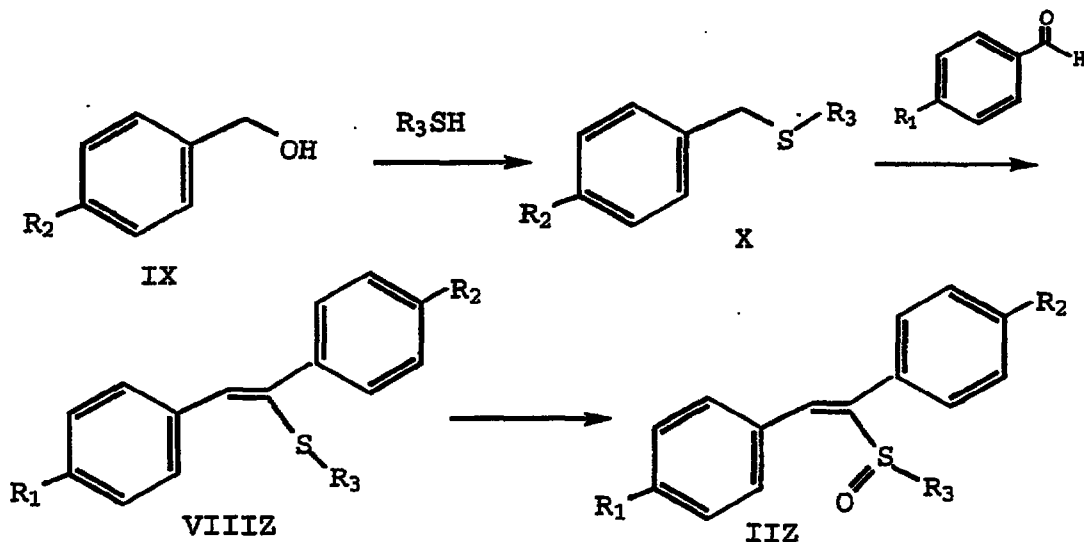
25 well known in the chemical arts, such as by crystallization or chromatography.

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The formula VIII compound, where R_1 , R_2 , and R_3 are as defined above, is then oxidized to produce the formula II compounds. Suitable oxidizing agents for this reaction are peracids, such as peracetic acid and *m*-chloroperoxybenzoic acid, and hydrogen peroxide. This oxidation reaction is typically run in an organic solvent, such as toluene, methylene chloride, chloroform, or carbontetrachloride. When a peracid is used as the oxidant, the reaction is generally carried out at a temperature of about -30°C to about 15°C , preferably at about -20°C . The products of the reaction are easily purified by recrystallization. When R_3 is *t*-butyl, the crystalline product of this reaction sequence is the **E** regioisomer of formula II.

When R_3 has a tertiary carbon adjacent to the sulfur atom, the **Z** regioisomer of the formula II compounds can be prepared selectively by a second route as shown in Scheme II.

Scheme 2



Generally, a benzyl alcohol, a formula IX compound, is
 5 reacted with a mercaptan of the formula R_3SH to produce a
 benzyl sulfide, a formula X compound. This benzyl sulfide is
 reacted with a strong base, forming a benzylic anion, which
 is condensed with a benzaldehyde. This condensation product
 is reacted with an acid chloride and the resulting
 10 intermediate treated with a second strong base to produce a
 styryl sulfide, a formula VIII Z compound. This styryl
 sulfide is then oxidized with an oxidizing agent to produce
 the formula II Z compound.

The first step in the synthesis of the Z styryl
 15 sulfoxide compounds is the conversion of a benzyl alcohol to
 a benzyl sulfide, formula X compound. The reaction of the
 formula IX compound, where R_2 is as defined above, with a
 mercaptan of the formula R_3SH , wherein R_3 is a thermally-
 labile or acid-labile C_2 - C_{10} alkyl, C_4 - C_{10} alkenyl, or
 20 aryl(C_1 - C_{10} alkyl) group having a tertiary carbon atom
 adjacent to the sulfur atom, in the presence of a Lewis acid
 produces the benzyl sulfide, a formula X compound. Suitable
 Lewis acids for this transformation are zinc bromide, zinc
 chloride, zinc iodide, ferric chloride, titanium(IV)
 25 chloride, aluminum trichloride, and aluminum tribromide,
 preferably zinc iodide. The reaction is generally carried

out in an organic solvent, such as 1,2-dichloroethane or methylene chloride. When the reaction is carried out at room temperature, the reaction is complete after about 18 hours.

The benzyl sulfide is reacted with a strong base to form a benzylic anion. Suitable strong bases for this reaction include metal alkoxides, such as sodium methoxide, sodium ethoxide, lithium ethoxide, lithium *t*-butoxide, and potassium *t*-butoxide; sodium hydride; and alkylolithiums, such as *n*-butyllithium, *t*-butyllithium, *sec*-butyllithium, and methyllithium. The preferred strong base for this reaction is *n*-butyllithium. The preferred solvent for this reaction is dry tetrahydrofuran. When *n*-butyllithium is used as the strong base, the reaction is carried out at a temperature of about -35°C to about -15°C.

The benzylic anion is condensed with a benzaldehyde to prepare an intermediate condensation product. The benzaldehyde has the general formula $p\text{-R}_1(\text{C}_6\text{H}_4)\text{CHO}$, wherein R_1 is hydrogen, $\text{C}_1\text{-C}_4$ alkoxy, arylalkoxy, halo, or amino. Preferably, the benzylic anion is prepared and the condensation product is formed *in situ* by adding the benzaldehyde to the cold solution of the benzylic anion.

The condensation product is treated with an acid chloride to produce an intermediate compound. Representative acid chlorides include acyl chlorides, such as acetyl chloride and benzoyl chloride; sulfonyl chlorides, such as methanesulfonyl chloride, benzenesulfonyl chloride, 1-butanesulfonyl chloride, ethanesulfonyl chloride, isopropylsulfonyl chloride, and *p*-toluenesulfonyl chloride; alkoxycarbonyl chlorides, such as methoxycarbonyl chloride and benzyloxycarbonyl chloride; and dialkylaminocarbonyl chlorides, such as *N,N*-dimethylaminocarbonyl chloride; preferably a sulfonyl chloride. Preferably, methanesulfonyl chloride is added to the reaction mixture shortly after formation of the condensation product.

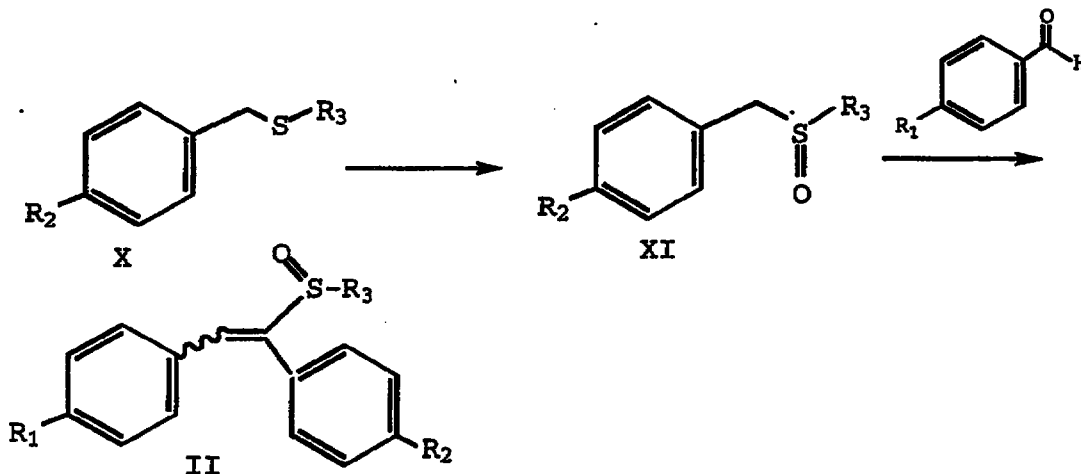
This intermediate compound is reacted with a second strong base to produce a styryl sulfide, a formula VIIIZ compound where R_1 , R_2 , and R_3 are as defined above. Suitable

strong bases for this reaction include metal alkoxides, such as sodium methoxide, sodium ethoxide, lithium ethoxide, lithium *t*-butoxide, and potassium *t*-butoxide; sodium hydride; alkyllithiums, such as *n*-butyllithium, *t*-butyllithium, *sec*-butyllithium, and methyllithium; and metal amides, such as sodium amide, magnesium diisopropylamide, and lithium diisopropylamide. The preferred strong base for this reaction is potassium *t*-butoxide. Generally, this reaction is carried out at about 15°C to about room temperature, preferably at room temperature.

The styryl sulfide is oxidized to prepare the corresponding styryl sulfoxide. Suitable oxidizing agents for this reaction are peracids, such as peracetic acid and *m*-chloroperoxybenzoic acid; organic peroxides, such as *t*-butyl peroxide; and hydrogen peroxide. Preferably the oxidizing agent is peracetic acid. This oxidation is typically carried out in an organic solvent, such as toluene, benzene, xylene, methanol, ethanol, methylacetate, ethylacetate, methylene chloride, 1,2-dichloroethane, or chloroform; preferably methylene chloride. This oxidation can be carried out at a temperature of about -40°C to about 0°C.

Alternatively, when R₃ has a tertiary carbon adjacent to the sulfur atom, the benzyl sulfide intermediate (formula X compound) can be used to produce a mixture of *E* and *Z* isomers of the styryl sulfoxides, the formula II compounds. This synthesis is outlined in Scheme 3.

Scheme 3



5 The benzyl sulfide, prepared as described above, is oxidized to produce the corresponding benzyl sulfoxide. This benzyl sulfoxide is reacted with a strong base, and the resulting anion condensed with a benzaldehyde. The condensation product is reacted with an acid chloride and the
 10 resulting intermediate compound reacted with a second strong base to produce the styryl sulfoxide.

 The benzyl sulfide, the formula X compound, wherein R_2 is as defined above and R_3 is a thermally-labile or acid-labile C_2 - C_{10} alkyl, C_4 - C_{10} alkenyl, or aryl(C_1 - C_{10} alkyl) group having a tertiary carbon atom adjacent to the sulfur atom, is oxidized to produce the corresponding benzyl sulfoxide, formula XI compound. Suitable oxidizing agents for this reaction are peracids, such as peracetic acid and *m*-chloroperoxybenzoic acid; organic peroxides, such as *t*-butyl peroxide; and hydrogen peroxide. Preferably the oxidizing
 20 agent is peracetic acid. This oxidation is typically carried out in an organic solvent, such as toluene, benzene, xylene, methanol, ethanol, methylacetate, ethylacetate, methylene chloride, 1,2-dichloroethane, or chloroform; preferably at a
 25 temperature of about -30°C to about 5°C .

 The benzyl sulfoxide, formula XI compound wherein R_2 and R_3 are as defined above, is reacted with a strong base to

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produce a benzylic anion. Suitable strong bases for this reaction include metal alkoxides, such as sodium methoxide, sodium ethoxide, lithium ethoxide, lithium *t*-butoxide, and potassium *t*-butoxide; sodium hydride; alkylolithiums, such as *n*-butyllithium, *t*-butyllithium, *sec*-butyllithium, and methylolithium; and metal amides, such as sodium amide, magnesium diisopropylamide, and lithium diisopropylamide. The preferred base for this transformation is *n*-butyllithium. This deprotonation reaction is carried out in a dry organic solvent, such as tetrahydrofuran or 1,2-dimethoxyethane, at a temperature of about -25°C.

The benzylic anion is condensed, without isolation, with a benzaldehyde compound of the formula $p\text{-R}_1(\text{C}_6\text{H}_4)\text{CHO}$, wherein R_1 is as defined above. Preferably, about one equivalent of the benzaldehyde is added to the cold solution prepared as described in the preceding paragraph. The resulting diastereomeric mixture of condensation products may be isolated, or preferably used in the next step without isolation.

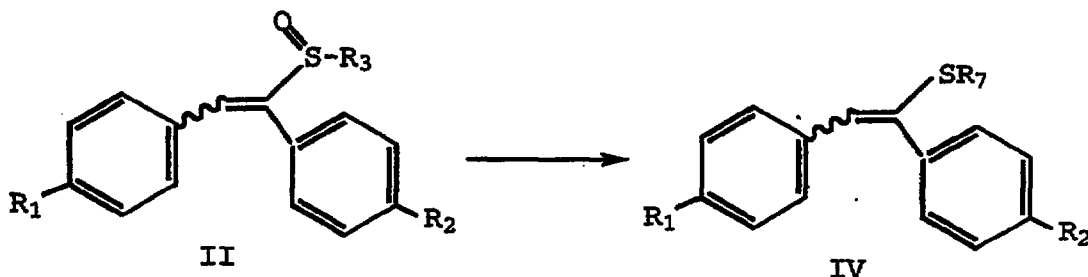
The condensation product is reacted with an acid chloride to produce an intermediate compound. The condensation product is optionally treated with a base, such as *n*-butyllithium, and reacted with an acid chloride. Representative acid chlorides include acyl chlorides, such as acetyl chloride and benzoyl chloride; sulfonyl chlorides, such as methanesulfonyl chloride, benzenesulfonyl chloride, 1-butan sulfonyl chloride, ethanesulfonyl chloride, isopropylsulfonyl chloride, and *p*-toluenesulfonyl chloride; alkoxycarbonyl chlorides, such as methoxycarbonyl chloride and benzyloxycarbonyl chloride; and dialkylaminocarbonyl chlorides, such as *N,N*-dimethylaminocarbonyl chloride; preferably a sulfonyl chloride. The acid chloride is added to the cold reaction mixture, then the resulting mixture is allowed to warm to room temperature. Preferably, methanesulfonyl chloride is added to the reaction mixture shortly after formation of the condensation product, which eliminates the need to add additional base.

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The resulting intermediate compound is reacted with a second strong base to produce the E and Z styryl sulfoxides, formula II compounds where R₁, R₂, and R₃ are as defined above. Representative second strong bases for this elimination reaction include metal alkoxides, such as sodium methoxide, sodium ethoxide, lithium ethoxide, lithium t-butoxide, and potassium t-butoxide; sodium hydride; alkyllithiums, such as n-butyllithium, t-butyllithium, sec-butyllithium, and methyllithium; and metal amides, such as sodium amide, magnesium diisopropylamide, and lithium diisopropylamide. The preferred base for this transformation is potassium t-butoxide. Preferably, a 20% excess, such as 1.2 equivalents, of the second base are added. Generally, this reaction is carried out at a temperature of about 15°C to about room temperature, preferably at room temperature.

The compounds of the present invention can be prepared from the formula II compounds. The novel sulfenate silyl esters are prepared from the styryl sulfoxides as shown in Scheme 4.

Scheme 4



5 Generally, the sulfenamide silyl esters, where R_1 , R_2 , and R_7 are as defined above and R_3 is a thermally-labile or acid-labile $\text{C}_1\text{-C}_{10}$ alkyl, $\text{C}_4\text{-C}_{10}$ alkenyl, or aryl($\text{C}_1\text{-C}_{10}$ alkyl) group, are prepared by reacting a formula II compound with a silylating reagent. Suitable solvents for this reaction

10 include benzene, toluene, xylene, and high-boiling, halogenated hydrocarbon solvents, having a boiling point greater than or equal to 80°C , such as 1,1,2-trichloroethane. Suitable silylating reagents include bis(trialkylsilyl)ureas, such as 1,3-bis(trimethylsilyl)urea, 1,3-

15 bis(triethylsilyl)urea, 1,3-bis(dimethylisopropylsilyl)-urea, 1,3-bis(*t*-butyl-dimethylsilyl)urea; bis(triarylsilyl)-ureas, such as 1,3-bis(triphenylsilyl)urea; bis(dialkylaryl-silyl)ureas, such as 1,3-bis(diphenylmethylsilyl)urea; and hexaalkyldisilylzanones, such as hexamethyldisilylthane; or

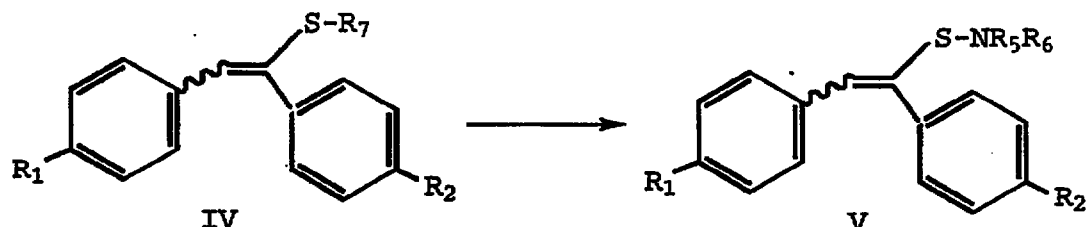
20 combination of a hexaalkyldisilylthane and a catalytic amount of a chlorotrialkylsilane, such as chlorotrimethylsilane. For best results, the final concentration, after complete addition, of the formula II compound is about 0.001 M to about 0.5 M. Preferably, a slight excess, such as ten

25 percent, of the silylating reagent is used. This reaction can be carried out at about 80°C to about 140°C for about ten minutes to about two hours. Because the *Z* isomer reacts much faster than the corresponding *E* isomer, the use of only the *Z* isomer as the starting compound requires less time for

30 complete transformation.

The novel sulfenamides are prepared from the sulfenamide silyl esters as shown in Scheme 5.

Scheme 5



5

Generally, the sulfenate silyl ester, where R₁, R₂, and R₇ are as defined above, is prepared from the styryl sulfoxide and, preferably without isolation or purification, reacted with an amine of the formula HNR₅R₆, wherein R₅ and R₆ as defined above. Typically, the sulfenate silyl ester is prepared, the reaction solution cooled to about 0°C to about 50°C, and treated with the amine. Preferably, one to two equivalents of the amine are used. The conversion from the silyl ester to the sulfenamide is typically complete after about two hours to about eight hours. The resulting sulfenamides can be purified using standard organic techniques, such as silica-gel chromatography.

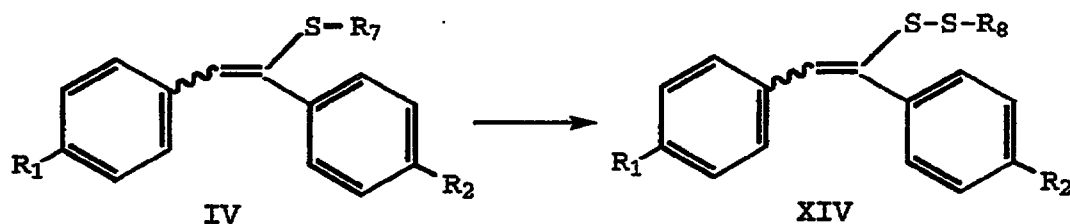
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The novel disulfides are prepared from the sulfenate silyl esters as shown in Scheme 6.

20

Scheme 6



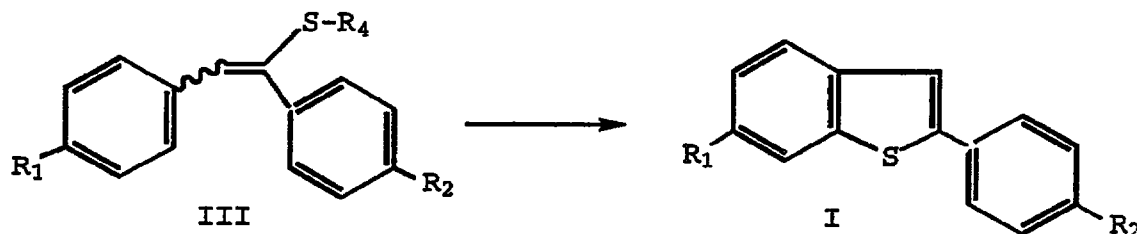
25

Generally, the sulfenate silyl ester, where R₁, R₂, and R₇ are as defined above, is prepared from the styryl sulfoxide and, preferably without isolation or purification, reacted with a mercaptan of the formula HSR₈, where R₈ is as defined above, in the presence of an amine base. Preferably,

the sulfenate silyl ester is prepared, the reaction solution allowed to cool to room temperature, and the reaction mixture treated with a solution containing the mercaptan and amine base. The solvent for this mercaptan/amine solution is the same as the solvent for the sulfenate silyl ester-containing mixture. Representative amine bases include triethylamine, diisopropylethylamine, pyridine, morpholine, *N*-methyl-morpholine, and collidine. The conversion of the sulfenate silyl ester is typically complete after about one to about eight hours. The resulting disulfides can be purified using standard organic techniques, such as silica-gel chromatography.

The intermediate sulfenate silyl esters, sulfenamides, and disulfides are useful for the synthesis of 2-arylbenzo-[b]thiophenes as shown in Scheme 7.

Scheme 7



20

Generally, the sulfenate silyl esters, sulfenamides, or disulfides are treated with acid catalysts to produce the formula I compounds. Suitable acid catalysts for this reaction include Lewis acids or Brønsted acids. Representative Lewis acids include zinc chloride, zinc iodide, aluminum chloride, and aluminum bromide. Representative Brønsted acids include inorganic acids, such as sulfuric and phosphoric acids; carboxylic acids, such as acetic and trifluoroacetic acids; sulfonic acids, such as methanesulfonic, benzenesulfonic, 1-naphthalenesulfonic, 1-butanesulfonic, ethanesulfonic, 4-ethylbenzenesulfonic, 1-hexanesulfonic, 1,5-naphthalenedisulfonic, 1-octanesulfonic, camphorsulfonic, trifluoromethanesulfonic, and *p*-toluene-

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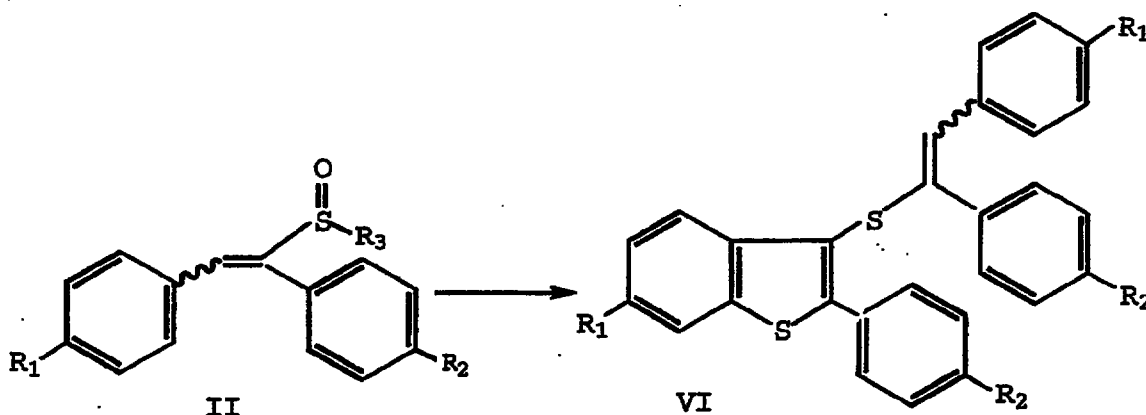
-22-

sulfonic acids; and polymeric arylsulfonic acids, such as Nafion®, Amberlyst®, or Amberlite®. The more preferred acid catalysts are sulfonic acids, such as methanesulfonic acid, benzenesulfonic acid, camphorsulfonic, and *p*-

5 *p*-toluenesulfonic acid. The most preferred acid catalyst is *p*-toluenesulfonic acid. Typically, a solution of the acid catalyst in an organic solvent, such as toluene, benzene, xylene, or a high-boiling halogenated hydrocarbon solvent, such as 1,1,2-trichloroethane, is heated to about 80°C to
10 about 140°C, and treated with a solution of the sulfenate silyl ester, sulfenamide, or disulfide in the same solvent. An excess amount of the acid catalyst is used, preferably three equivalents of the acid. For best results, the final concentration of the starting compound is about 0.01 M to
15 about 0.2 M, preferably 0.05 M. Furthermore, best yields are obtained when the sulfenate silyl ester is slowly added to the heated acid solution over a period of about 15 minutes to about three hours. For best results, residual water is removed from the reaction solution by the use of a Dean-Stark
20 trap or Soxhlet extractor.

The styryl sulfoxides are also useful for the preparation of a benzothiophene styryl sulfide as shown in Scheme 8.

25 **Scheme 8**

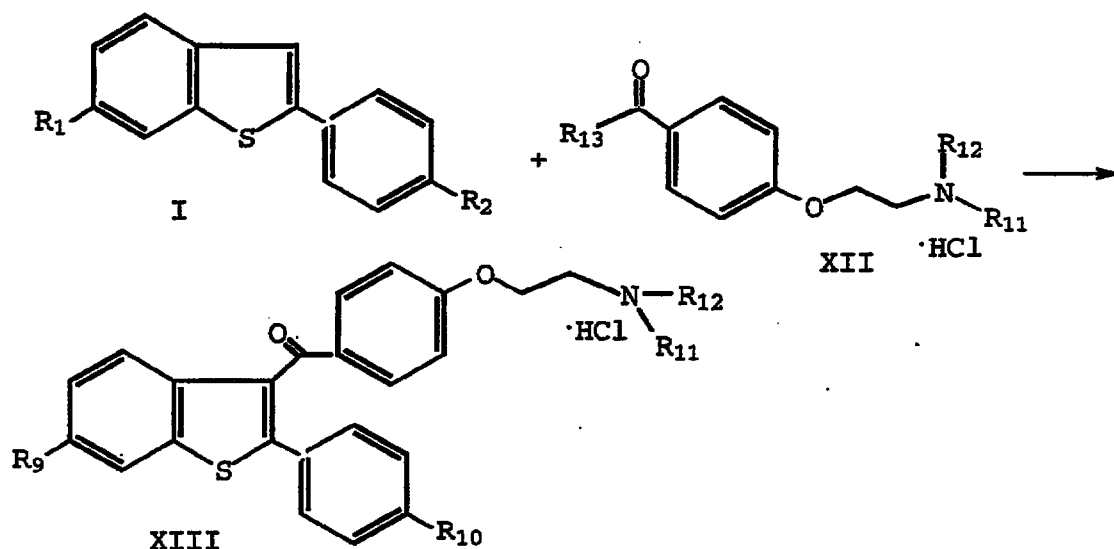


These benzothiophene styryl sulfides, where R_1 and R_2 are as defined above, are prepared from the styryl sulfoxides. Generally, a solution of the styryl sulfoxide, where R_1 and R_2 are as defined above and R_3 is a thermally-labile or acid-labile C_2 - C_{10} alkyl, C_4 - C_{10} alkenyl, or aryl(C_1 - C_{10} alkyl) group, is added to a solution of an acid catalyst at a temperature of about 100°C to about 140°C , where the acid catalyst is defined above. The concentration of acid catalyst is dependent on the final concentration of the formula II compound and the rate of addition of the formula II compound. When the styryl sulfoxide is at a final concentration of about 0.2 M and is added over six hours, the acid concentration is about 0.002 M. When the styryl sulfoxide is at a final concentration of about 0.05 M and is added over 30 minutes, the acid concentration is about 0.025 M. Significant quantities of the formula VI compounds are present in the reaction after about one to two hours. Longer reaction times lead to the production of the formula I compounds.

These formula VI compounds may be subsequently converted to the formula I compounds by treatment with additional acid, such as about 0.5 to about three equivalents, and heating to about 100°C to about 140°C . The concentration of the formula VI compound is in the range of about 0.01 M to about 0.5 M. Suitable solvents for both the formation of the formula VI compounds and their conversion to formula I compounds include toluene, xylene, and 1,2-dichloroethane.

The formula I compounds are useful as intermediates in the synthesis of a series of 3-aryl-2-arylbenzo[b]-thiophenes. U.S. Patent Nos. 4,133,814 and 4,418,068, which are incorporated herein by reference, described these 3-aryl-2-arylbenzo[b]thiophenes, as well as methods for their preparation from the formula I compounds. An improved synthesis of a group of these 3-aryl-2-arylbenzo[b]-thiophenes from the formula I compounds, wherein R_1 and R_2 are hydrogen, C_1 - C_4 alkoxy, or arylalkoxy is outlined in Scheme 9.

Scheme 9



5 The Formula I compound, wherein R₁ and R₂ are hydrogen, C₁-C₄ alkoxy, or arylalkoxy, is acylated with the formula XII compound, wherein R₁₃ is chloro or hydroxy, in the presence of boron trichloride or boron tribromide; boron trichloride is preferred. The reaction can be carried out in a variety of
 10 organic solvents, such as chloroform, methylene chloride, 1,2-dichloroethane, 1,2,3-dichloropropane, 1,1,2,2-tetrachloroethane, 1,2-dichlorobenzene, chlorobenzene, and fluorobenzene. The preferred solvent for this synthesis is 1,2-dichloroethane. The reaction is carried out at a
 15 temperature of about -10°C to about 25°C, preferably at 0°C. The reaction is best carried out at a concentration of the benzothiophene formula I compound of about 0.2 M to about 1.0 M. The acylation reaction is generally complete after about two hours to about eight hours.

20 When R₁ and/or R₂ is a C₁-C₄ alkoxy or arylalkoxy group, the acylated benzothiophene preferably is converted to a formula XIII compound, wherein R₅ and/or R₆ are hydroxy, without isolation of the product from the acylation reaction. This conversion is performed by adding additional boron
 25 trichloride or boron tribromide and heating the reaction

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mixture. Preferably, two to five molar equivalents of boron trichloride are added to the reaction mixture, most preferably three molar equivalents. This reaction is carried out at a temperature of about 25°C to about 40°C, preferably at 35°C. The reaction is generally complete after about 4 hours to about 48 hours.

The acylation reaction or acylation/dealkylation reaction is quenched with an alcohol or a mixture of alcohols. Suitable alcohols for use in quenching the reaction include methanol, ethanol, and isopropanol. Preferably, the acylation/dealkylation reaction mixture is added to a 95:5 mixture of ethanol and methanol (3A ethanol). The 3A ethanol can be at room temperature or heated to reflux, preferably at reflux. When the quench is performed in this manner, the Formula XIII compound conveniently crystallizes from the resulting alcoholic mixture. Generally, 1.25 mL to 3.75 mL of alcohol per millimole of the benzothiophene starting material are used.

The following examples further illustrate the present invention. The examples are not intended to be limiting to the scope of the invention in any respect, and should not be so construed. All experiments were run under positive pressure of dry nitrogen. All solvents and reagents were used as obtained. The percentages are generally calculated on a weight (w/w) basis; except for high performance liquid chromatography (HPLC) solvents which are calculated on a volume (v/v) basis. Proton nuclear magnetic resonance (^1H NMR) spectra and ^{13}C nuclear magnetic resonance (^{13}C NMR) spectra were obtained on a Bruker AC-300 FTNMR spectrometer at 300.135 MHz or at 75.469 MHz for proton and carbon, respectively, or a GE QE-300 spectrometer at 300.15 MHz. Silica-gel flash chromatography was performed as described by Still et al. using Silica Gel 60 (230-400 mesh, E. Merck). Still et al., *J. Org. Chem.*, **43**, 2923 (1978). Elemental analyses for carbon, hydrogen, and nitrogen were determined on a Control Equipment Corporation 440 Elemental Analyzer. Elemental analyses for sulfur were determined on a Brinkman

Colorimetric Elemental Analyzer. Melting points were determined in open glass capillaries on a Mel-Temp II melting point apparatus or a Mettler FP62 Automatic instrument, and are uncorrected. Field desorption mass spectra (FDMS) were obtained using a Varian Instruments VG 70-SE or VG ZAB-3F mass spectrometer. High resolution free atom bombardment mass spectra (FABMS) were obtained using a Varian Instruments VG ZAB-2SE mass spectrometer.

The *in situ* yields of 6-methoxy-2-(4-methoxyphenyl)-benzo[b]thiophene were determined by high performance liquid chromatography (HPLC) in comparison to an authentic sample of this compound prepared by published synthetic routes. See U.S. Patent No. 4,133,814. Generally, samples of the reaction mixture was diluted with acetonitrile and the diluted sample assayed by HPLC using a Zorbax® RX-C8 column (4.6 mm x 25 cm) with UV detection (280 nm). The following linear-gradient solvent system was used for this analysis:

Gradient Solvent System

Time (min)	A (%)	B (%)
0	50	50
2	50	50
20	20	80
35	20	80
37	50	50
45	50	50

A: 0.01 M aqueous sodium phosphate (pH 2.0)

B. acetonitrile

The amount (percentages) of 6-hydroxy-2-(4-hydroxy-phenyl)-3-[4-(2-piperidinoethoxy)benzoyl]benzo[b]thiophene hydrochloride in the crystalline material (potency) was determined by the following method. A sample of the crystalline solid (5 mg) was weighed into a 100-mL volumetric flask, and dissolved in a 70/30 (v/v) mixture of 75 mM

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potassium phosphate buffer (pH 2.0) and acetonitrile. An aliquot of this solution (10 μ L) was assayed by high performance liquid chromatography, using a Zorbax® Rx-C8 column (25 cm x 4.6 mm ID, 5 μ particle) and UV detection (280 nm). The following gradient solvent system was used:

Gradient Solvent System (Potency)

	<u>Time (min)</u>	<u>A (%)</u>	<u>B (%)</u>
10	0	70	30
	12	70	30
	14	25	75
	16	70	30
	25	70	30

15

A: 75 mM KH₂PO₄ buffer (pH 2.0)

B: acetonitrile

The percentage of 6-hydroxy-2-(4-hydroxyphenyl)-3-[4-(2-piperidinoethoxy)benzoyl]benzo[b]thiophene hydrochloride in the sample was calculated using the peak area, slope (m), and intercept (b) of the calibration curve with the following equation:

$$\% \text{ potency} = \frac{\text{peak area} - b}{m} \times \frac{\text{sample volume (mL)}}{\text{sample weight (mg)}}$$

25

The amount (percentage) of solvent, such as 1,2-dichloroethane, present in the crystalline material was determined by gas chromatography. A sample of the crystalline solid (50 mg) was weighed into a 10-mL volumetric flask, and dissolved in a solution of 2-butanol (0.025 mg/mL) in dimethylsulfoxide. A sample of this solution was analyzed on a gas chromatograph using a DB Wax column (30 m x 0.53 mm ID, 1 μ particle), with a column flow of 10 mL/min and flame ionization detection. The column temperature was heated from

35

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35°C to 230°C over a 12 minute period. The amount of solvent was determined by comparison to the internal standard (2-butanol).

5

Example 1***E*-t-Butyl 4,4'-Dimethoxystilbenyl Sulfoxide****A. Preparation of *E*-t-Butyl 4,4'-Dimethoxystilbenyl Sulfide**

10 A solution of desoxyanisoin (12.82 g) in tetrahydrofuran (100 mL) was treated with titanium (IV) chloride (10.43 g). During the dropwise addition of titanium (IV) chloride, the reaction mixture was cooled to maintain the temperature below 35°C. Upon complete addition, the resulting mixture was
15 stirred at 30°C. After an additional 30 minutes, this mixture was treated with a solution of 2-methyl-2-propane-thiol (6.76 mL) and triethylamine (16.70 mL) in tetrahydrofuran (15 mL). The resulting mixture was stirred at 50°C. After two hours, the mixture was added to ten percent sodium
20 carbonate (500 mL). The resulting mixture was extracted with methylene chloride. The combined methylene chloride extracts were dried over magnesium sulfate, filtered, and concentrated in vacuo to give 17.2 g of an oil, which crystallized upon cooling to room temperature. This crystalline material was
25 recrystallized from hot ethanol to give 12.3 g of the title compound. Melting point 71-73°C.

Analysis calculated for $C_{20}H_{24}O_2S$: C, 73.13; H, 7.36; S, 9.76. Found: C, 73.37; H, 7.51; S, 9.87.

30

B. Preparation of *E*-t-Butyl 4,4'-Dimethoxystilbenyl Sulfoxide

The crystalline compound prepared as described in Example 1A was dissolved in toluene (150 mL), and the
35 resulting solution cooled to about -20°C. The cold solution was treated with peracetic acid (32% w/w in dilute acetic acid, 1.24 g) over ten minutes. The resulting mixture was

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extracted with saturated sodium sulfite and brine. The organic phase was concentrated *in vacuo*. The residue was recrystallized from ethyl acetate/heptane to give 14.11 g of the title compound. Melting point 104°C (dec).

5 Analysis calculated for $C_{20}H_{24}O_3S$: C, 69.74; H, 7.02; S, 9.31. Found: C, 69.47; H, 7.04; S, 9.54.

Example 2

Z-t-Butyl 4,4'-Dimethoxystilbenyl Sulfoxide

10 A. Preparation of t-Butyl 4-Methoxybenzyl Sulfide

A mixture of 4-methoxybenzyl alcohol (10.13 g) and zinc iodide (11.7 g) in 1,2-dichloroethane (120 mL) was treated with 2-methyl-2-propanethiol (9.92 mL) in one portion. The resulting mixture was stirred at room temperature. After about 18 hours, the reaction was diluted with water (100 mL) and methylene chloride (100 mL). The organic phase was removed, dried over magnesium sulfate, filtered, and concentrated *in vacuo* to give 14.4 g of an oil.

20 1H NMR ($CDCl_3$): δ 7.28 (d, 2H), 6.85 (d, 2H), 3.77 (s, 3H), 3.73 (s, 2H), 1.36 (s, 9H).

^{13}C NMR ($CDCl_3$): δ 130, 114, 56, 35, 32.

Analysis calculated for $C_{12}H_{18}OS$: C, 68.52; H, 8.63. Found: C, 68.80; H, 8.67.

25

B. Preparation of Z-t-Butyl 4,4'-Dimethoxystilbenyl Sulfide

A solution of the compound prepared as described in Example 2A (2.51 g) in tetrahydrofuran (50 mL) was cooled to about -20°C. This cold solution was treated with a solution of n-butyllithium in hexane (1.6 M, 7.47 mL) over ten minutes. The resulting solution was allowed to warm to about 0°C over 35 minutes. This cold solution was treated with p-anisaldehyde (1.46 mL). After an additional 15 minutes, the reaction solution was treated with methanesulfonyl chloride (0.95 mL). The resulting reaction was allowed to warm to room temperature. After an additional 45 minutes, the

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reaction mixture was treated with a solution of potassium *t*-butoxide in tetrahydrofuran (1.0 M, 12.0 mL). After an additional 45 minutes, the reaction was quenched by the addition of 1N hydrochloric acid (12.0 mL). The organic
5 phase was separated, dried over magnesium sulfate, filtered, and concentrated to an oil (4.4 g).

^1H NMR (CDCl_3): δ 7.95 (d, H), 7.05 (s, H), 6.9 (d, H), 6.8 (dd, 2H), 3.75 (s, 3H), 0.95 (s, 9H).

^{13}C NMR (CDCl_3): δ 153, 139, 137, 114, 56, 32.

10

C. Preparation of *Z*-*t*-Butyl 4,4'-Dimethoxystilbenyl Sulfoxide

The compound from Example 2B was converted to the title
15 compound using the procedure substantially as described in Example 1B.

^1H NMR (CDCl_3): δ 7.61 (d, H), 7.56 (d, H), 7.1 (s, H), 6.9 (dd, 2H), 3.83 (s, 3H), 1.05 (s, 9H).

^{13}C NMR (CDCl_3): δ 142, 132.5, 131, 118, 117, 56, 24.

20 Analysis calculated for $\text{C}_{20}\text{H}_{24}\text{O}_3\text{S}$: C, 69.74; H, 7.02.
Found: C, 69.98; H, 6.94.

Example 3

E and *Z*-*t*-Butyl 4,4'-Dimethoxystilbenyl Sulfoxide

25 A. Preparation of *t*-Butyl 4-Methoxybenzyl Sulfide

A mixture of 4-methoxybenzyl alcohol (10.13 g) and zinc iodide (11.7 g) in 1,2-dichloroethane (120 mL) was treated with 2-methyl-2-propanethiol (9.92 mL) in one portion. The
30 resulting mixture was stirred at room temperature. After about 18 hours, the reaction was diluted with water (100 mL) and methylene chloride (100 mL). The organic phase was removed, dried over magnesium sulfate, filtered, and concentrated *in vacuo* to give 14.4 g of an oil.

35 ^1H NMR (CDCl_3): δ 7.28 (d, 2H), 6.85 (d, 2H), 3.77 (s, 3H), 3.73 (s, 2H), 1.36 (s, 9H).

^{13}C NMR (CDCl_3): δ 130, 114, 56, 35, 32.

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Analysis calculated for $C_{12}H_{18}OS$: C, 68.52; H, 8.63.
Found: C, 68.80; H, 8.67.

B. Preparation of *t*-Butyl 4-Methoxybenzyl Sulfoxide

5

A solution of the compound prepared as described in Example 3A (14.4 g) in 1,2-dichloroethane (50 mL) was cooled to about 5°C and the cold solution treated with peracetic acid (32% w/w in dilute acetic acid, 14.2 mL) over 30 minutes. Upon complete addition of the peracetic acid, the reaction was treated with brine and sodium bicarbonate. The organic phase was removed, dried over magnesium sulfate, filtered, and concentrated to a yellow precipitate. This residue was treated with hexane (100 mL) and the resulting mixture stirred at room temperature. After about 18 hours, the mixture was filtered and the solids washed with hexane (100 mL). The solid material was dried in vacuo to give 14.07 g of the title compound. Melting point 124-126°C.

1H NMR ($CDCl_3$): δ 7.26 (d, 2H), 6.89 (d, 2H), 3.79 (d, H), 3.78 (s, 3H), 3.58 (d, H), 1.3 (s, 9H).

^{13}C NMR ($CDCl_3$): δ 132, 114, 56, 53, 23.

Analysis calculated for $C_{12}H_{18}O_2S$: C, 63.68; H, 8.02.
Found: C, 63.72; H, 7.93.

25 C. Preparation of *E* and *Z*-*t*-Butyl 4,4'-Dimethoxystilbenyl Sulfoxide

A solution of the compound prepared as described in Example 3B (10.0 g) in tetrahydrofuran (140 mL) was cooled to about -30° to -25°C (dry ice/acetone bath). This cold solution was treated with *n*-butyllithium in cyclohexane (1.6 M, 27.65 mL) over 25 minutes. After stirring for 35 minutes, the reaction mixture was treated with *p*-anisaldehyde (5.4 mL). The dry ice/acetone bath was removed and the reaction allowed to warm to about 20°C. This mixture was treated with methanesulfonyl chloride (3.5 mL). The temperature of the reaction rose from about 20° to about 35°C

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upon addition of the methanesulfonyl chloride. The mixture was cooled to about 25°C, then treated with potassium t-butoxide in tetrahydrofuran (1 M, 50.9 mL). After stirring for an additional 35 minutes, the reaction was treated with
5 1N hydrochloric acid (51.0 mL). The phases were separated, and the organic layer dried over magnesium sulfate, filtered, and concentrated to an oil (16.67 g). This material was used in the next step without further purification. The carbon and proton NMR spectra were similar to that obtained for the
10 compound prepared as described in Examples 1 and 2.

Example 4

E and Z-Trimethylsilyl 4,4'-Dimethoxystilbenyl Sulfenate

15 A mixture of the compound prepared as described in Example 1 (350 mg) and 1,3-bis(trimethylsilyl)urea (116 mg) in toluene (11 mL) was heated to reflux. After 1.5 hours, the reaction mixture was allowed to cool to room temperature, filtered, and the filtrate concentrated in vacuo to give a
20 7:1 mixture of E/Z regioisomers of the title compounds.

FDMS: m/z = 361 (M+1).

E Isomer:

25 ¹H NMR (d₆-benzene): δ 7.39 (d, 2H), 7.10 (d, 2H), 6.68 (d, 2H), 6.68 (s, 1H), 6.57 (d, 2H), 3.18 (s, 3H), 3.17 (s, 3H), 0.23 (s, 9H).

Z Isomer:

30 ¹H NMR (d₆-benzene): δ 7.71 (d, 2H), 7.31 (d, 2H), 6.85 (d, 2H), 6.79 (d, 2H), 6.60 (s, 1H), 3.28 (s, 3H), 3.26 (s, 3H), -0.05 (s, 9H).

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Example 5**E and Z-Trimethylsilyl 4,4'-Dimethoxystilbenyl Sulfenate**

A mixture of the compound prepared as described in Example 2 and 1,3-bis(trimethylsilyl)urea in toluene was heated to reflux. After ten minutes, the mixture was allowed to cool, filtered, and concentrated in vacuo to give a 7:1 mixture of E/Z regioisomers of the title compounds.

E Isomer:

¹³C NMR (d₆-benzene, 8°C): δ 160.49, 158.53, 141.54, 131.97, 129.91, 129.65, 125.59, 116.41, 114.68, 113.98, 54.56, -0.09.

Example 6**E and Z-N,N-Dimethyl-4,4'-Dimethoxystilbenyl Sulfenamide**

A mixture of the compound prepared as described in Example 1 (1.74 g) and 1,3-bis(trimethylsilyl)urea (578 mg) in toluene (54 mL) was heated to reflux. After 1.5 hours, the reaction was allowed to cool to room temperature, and treated with dimethylamine (2.80 mL, 2.0 M in tetrahydrofuran). After an additional two hours, the reaction solution was evaporated to dryness to give a 7:1 mixture of E/Z regioisomers of the title compounds. This residual mixture was purified using silica-gel flash chromatography, eluting with a mixture of ethyl acetate/hexane (9:1), to give 1.06 g of the title compounds as an 8:1 mixture of E/Z regioisomers.

FDMS: m/z = 315 (M⁺).

Analysis calculated for C₁₈H₂₁NO₂S: C, 68.54; H, 6.71; N, 4.44. Found: C, 68.40; H, 6.69; N, 4.22.

E Isomer:

¹H NMR (d₆-benzene): δ 7.44 (d, 2H), 7.11 (d, 2H), 6.99 (s, 1H), 6.71 (d, 2H), 6.56 (d, 2H), 3.22 (s, 3H), 3.18 (s, 3H), 2.66 (s, 6H).

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¹³C NMR (d₆-benzene): δ 160.00, 158.83, 139.70, 131.48, 130.78, 130.51, 129.94, 123.77, 114.55, 113.97, 54.63, 54.61, 48.17.

5

Z Isomer:

¹H NMR (d₆-benzene): δ 7.61 (d, 4H), 6.82 (d, 2H), 6.80 (d, 2H), 6.80 (s, 1H), 3.32 (s, 3H), 3.27 (s, 3H), 2.41 (s, 6H).

10

¹³C NMR (d₆-benzene): δ 159.89, 159.30, 139.76, 136.46, 131.94, 131.82, 130.22, 130.20, 113.83, 113.76, 54.81, 54.73, 48.61.

15

Example 7

E and Z-N-Benzyl-4,4'-Dimethoxystilbenyl Sulfenamide

A mixture of the compound prepared as described in Example 1 (1.74 g) and 1,3-bis(trimethylsilyl)urea (578 mg) in toluene (54 mL) was heated to reflux. After 1.5 hours, the reaction was allowed to cool to room temperature, and treated with benzylamine (0.575 mL). After an additional two hours, the reaction solution was evaporated to dryness to give a 7:1 mixture **E/Z** of regioisomers of the title compounds. This residual mixture was purified using silica-gel flash chromatography, eluting with a mixture of ethyl acetate/hexane (7:1), to give 1.06 g of the title compounds as a 6:1 mixture of **E/Z** regioisomers.

Analysis calculated for C₂₃H₂₃NO₂S: C, 73.18; H, 6.14; N, 3.71. Found: C, 73.16; H, 6.18; N, 3.50

E Isomer:

¹H NMR (d₆-benzene): δ 7.41 (d, 2H), 7.13 (d, 2H), 7.12-7.03 (m, 5H), 6.87 (s, 1H), 6.71 (d, 2H), 6.59 (d, 2H), 3.89 (d, 2H), 3.23 (s, 3H), 3.20 (s, 3H), 2.71 (t, 1H).

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¹³C NMR (d₆-benzene): δ 159.98, 158.91, 140.53, 139.77, 131.45, 130.50, 129.87, 128.77, 128.66, 128.59, 127.53, 123.10, 114.74, 114.02, 56.14, 54.69, 54.64.

5 **Z Isomer:**

¹H NMR (d₆-benzene): δ 7.59 (d, 2H), 7.53 (d, 2H), 7.01-6.91 (m, 5H), 6.83 (s, 1H), 6.79 (d, 2H), 6.77 (d, 2H), 3.62 (d, 2H), 3.31 (s, 3H), 3.27 (s, 3H), 2.82 (t, 1H).

10 ¹³C NMR (d₆-benzene): δ 160.05, 159.14, 140.48, 139.27, 132.50, 131.32, 130.04, 129.86, 128.87, 128.58, 128.46, 127.49, 114.48, 114.00, 56.23, 54.90, 54.78.

Example 8

15 6-Methoxy-2-(4-methoxyphenyl)benzo[b]thiophene

A solution of p-toluenesulfonic acid monohydrate (552 mg) was added to toluene (15 mL) and heated to reflux, and water was removed by allowing it to collect in a Dean-
20 Stark trap. This refluxing solution was treated with a solution of the regioisomeric compounds prepared as described in Example 4 (523 mg) in toluene (15 mL) over 15 minutes. Upon complete addition, an aliquot was removed for HPLC analysis. This analysis showed a 46.6% *in situ* yield of the
25 title compound.

Example 9

6-Methoxy-2-(4-methoxyphenyl)benzo[b]thiophene

30 A solution of p-toluenesulfonic acid monohydrate (1.26 g) in toluene (20 mL) was heated to reflux, and water was removed by allowing it to collect in a Dean-Stark trap. A solution of the regioisomeric compounds prepared as described in Example 6 (650 mg) in toluene (9 mL) was added
35 to the refluxing acid solution over 1.8 hours. The reaction solution was treated with ethanol (10 mL), and the resulting

-36-

mixture allowed to cool to room temperature. The resulting slurry was stirred at room temperature. After about 18 hours, the mixture was cooled to about 5°C, and filtered to give 290 mg of the title compound. Melting point 199-200°C.

5

¹H NMR (d₆-DMSO): δ 7.67 (d, 1H), 7.64 (d, 2H), 7.61 (s, 1H), 7.52 (d, 1H), 7.01 (d, 2H), 6.98 (dd, 1H), 3.81 (s, 3H), 3.79 (s, 3H).

10 Analysis calculated for C₁₆H₁₄O₂S: C, 71.09; H, 5.22.
Found: C, 71.09; H, 5.27.

Example 10

15 E and Z-3-(4, 4'-Dimethoxystilbenyl sulfide)-6-methoxy-2-(4-methoxyphenyl)benzo[b]thiophene

A solution of p-toluenesulfonic acid monohydrate (552 mg) in toluene (111 mL) was heated to reflux, and water was removed by allowing it to collect in a Dean-Stark trap.

20 A solution of the compound prepared as described in Example 1 (10 g) in toluene (34 mL) was added to the refluxing acid solution over six hours. After an additional two hours, the mixture was cooled to 0°C. After an additional 18 hours, the cold mixture was filtered to remove the precipitated 6-methoxy-2-(4-methoxyphenyl)benzo[b]thiophene.

25 The filtrate was extracted with an equal volume of saturated sodium bicarbonate solution. The organic phase was separated, dried over sodium sulfate, filtered, and concentrated in vacuo to give 4.8 g of an orange oil. This oil was divided into two

30 parts and each purified using silica-gel flash chromatography, eluting with hexane/ethyl acetate (3.5:1). The fractions contained in the desired regioisomers were concentrated to an oil. This oil was treated with diethyl ether to selectively crystallize the early-eluting regio-

35 isomer (155 mg). The mother liquor from these crystallizations were enriched in the late-eluting regioisomer.

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Early-eluting Isomer

¹H NMR (CDCl₃): δ 7.71 (d, 2H), 7.64 (d, 1H), 7.46 (d, 2H),
7.06 (d, 1H), 6.94 (d, 2H), 6.92 (d, 2H), 6.90 (m, 1H), 6.85
(d, 2H), 6.59 (s, 1H), 6.45 (d, 2H), 3.86 (s, 3H), 3.85
5 (s, 3H), 3.80 (s, 3H), 3.66 (s, 3H).

High resolution FABMS calculated for C₃₂H₂₉O₄S₂ (MH⁺)
541.1507. Found: 541.1491.

10 Late-eluting Isomer

¹H NMR (CDCl₃): δ 7.90 (d, 1H), 7.62 (d, 2H), 7.24 (1H), 7.08
(d, 2H), 7.02 (dd, 1H), 6.96 (d, 2H), 6.74-6.71 (d, 2H), 6.70
(d, 2H), 6.55 (d, 2H), 6.21 (s, 1H), 3.86 (s, 3H), 3.85
(s, 3H), 3.76 (s, 3H), 3.67 (s, 3H).

15

FDMS: m/z = 540 (m⁺)

Example 11

6-Methoxy-2-(4-methoxyphenyl)benzo[b]thiophene

20

The compound (early-eluting isomer) prepared as
described in Example 10 (125 mg) was added to a refluxing
solution of *p*-toluenesulfonic acid monohydrate (4.2 mg) in
toluene (1.5 mL). After six hours, methanesulfonic acid
25 (7.5 μL) was added to the reaction mixture. After an
additional hour, the reaction mixture was allowed to cool to
room temperature. The resulting mixture was diluted with
acetonitrile and assayed by HPLC, showing a 71.1% *in situ*
yield of the title compound.

30

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Example 12**6-Hydroxy-2-(4-hydroxyphenyl)-3-[4-(2-piperidinoethoxy)-
benzoyl]benzo[b]thiophene Hydrochloride
1,2-Dichloroethane Solvate**5 **A. Preparation of Ethyl 4-(2-Piperidinoethoxy)benzoate**

A mixture of ethyl 4-hydroxybenzoate (8.31 g), 1-(2-chloroethyl)piperidine monohydrochloride (10.13 g), potassium carbonate (16.59 g), and methyl ethyl ketone (60 mL) was
10 heated to 80°C. After one hour, the mixture was cooled to about 55°C and treated with additional 1-(2-chloroethyl)-piperidine monohydrochloride (0.92 g). The resulting mixture was heated to 80°C. The reaction was monitored by thin layer chromatography (TLC), using silica-gel plates and ethyl
15 acetate/acetonitrile/triethylamine (10:6:1, v/v). Additional portions of 1-(2-chloroethyl)piperidine hydrochloride are added until the starting 4-hydroxybenzoate ester is consumed. Upon complete reaction, the reaction mixture was treated with water (60 mL) and allowed to cool to room temperature. The
20 aqueous layer was discarded and the organic layer concentrated in vacuo at 40°C and 40 mm Hg. The resulting oil was used in the next step without further purification.

25 **B. Preparation of 4-(2-Piperidinoethoxy)benzoic
Acid Hydrochloride**

A solution of the compound prepared as described in Example 12A (about 13.87 g) in methanol (30 mL) was treated with 5 N sodium hydroxide (15 mL), and heated to 40°C. After
30 4 1/2 hours, water (40 mL) was added. The resulting mixture was cooled to 5-10°C, and concentrated hydrochloric acid (18 mL) was added slowly. The title compound crystallized during acidification. This crystalline product was collected by filtration, and dried in vacuo at 40-50°C to give 83%
35 yield of the title compound. Melting point 270-271°C.

C. Preparation of 4-(2-Piperidinoethoxy)benzoyl
Chloride Hydrochloride

A solution of the compound prepared as described in
5 Example 12B (30.01 g) and dimethylformamide (2 mL) in
methylene chloride (500 mL) was treated with oxalyl chloride
(10.5 mL) over a 30-35 minute period. After stirring for
about 18 hours, the reaction was assayed for completion by
HPLC analysis. Additional oxalyl chloride may be added to
10 the reaction if the starting carboxylic acid is present.
Upon completion, the reaction solution was evaporated to
dryness in vacuo. The residue was dissolved in methylene
chloride (200 mL), and the resulting solution evaporated to
dryness. This dissolution/evaporation procedure was repeated
15 to give the title compound as a solid.

D. Preparation of 6-Hydroxy-2-(4-hydroxyphenyl)-3-[4-(2-
piperidinoethoxy)benzoyl]benzo[b]thiophene Hydrochloride
1,2-Dichloroethane Solvate

20 A mixture of the compound prepared as described in
Example 8 or 9 (2.92 g), the compound prepared as described
in Example 12C (3.45 g), and 1,2-dichloroethane (52 mL) was
cooled to about 0°C. Boron trichloride gas was condensed
25 into a cold graduated cylinder (2.8 mL), and added to the
cold mixture described above. After eight hours at 0°C, the
reaction mixture was treated with additional boron
trichloride (2.8 mL). The resulting solution was heated to
35°C. After 16 hours, the reaction was complete.

30 Methanol (30 mL) was treated with the reaction mixture
from above over a 20-minute period, causing the methanol to
reflux. The resulting slurry was stirred at 25°C. After one
hour, the crystalline product was filtered, washed with cold
methanol (8 mL), and dried at 40°C in vacuo to give 5.14 g of
35 the title compound. Melting point 225°C.

Potency (HPLC): 86.8%

1,2-Dichloroethane (gas chromatography): 6.5%

Example 13**6-Methoxy-2-(4-methoxyphenyl)benzo[b]thiophene**

5 A solution of *p*-toluenesulfonic acid monohydrate (1.05 g) in toluene (20 mL) was heated to reflux, and water was removed by allowing it to collect in a Dean-Stark trap. A solution of the regioisomeric compounds prepared as described in Example 7 (780 mg) in toluene (9 mL) was added
10 to the refluxing acid solution over ten minutes. After one hour, the reaction solution was treated with ethanol (10 mL), and the resulting mixture allowed to cool to room temperature. The resulting slurry was stirred at room temperature. After about 18 hours, the mixture was filtered
15 to give 149 mg of the title compound. Melting point 199-200°C.

Analysis calculated for C₁₆H₁₄O₂S: C, 71.09; H, 5.22.
Found: C, 71.05; H, 5.22.

20

Example 14***E* and *Z*-4,4'-Dimethoxystilbenyl Ethyl Disulfide**

A solution of the regioisomeric compounds prepared as
25 described in Example 4 (1.83 g) in toluene (54 mL) was treated with ethanethiol (0.433 mL) and triethylamine (0.715 mL). After about 2.5 hours at room temperature, the reaction solution was evaporated to dryness *in vacuo* to give a mixture of regioisomers. The residue was purified using
30 silica-gel chromatography, eluting with ethyl acetate/hexane (9:1), to give 1.14 g of a 5.7:1 mixture of *E/Z* regioisomers of the title compounds.

Analysis calculated for C₁₈H₂₀O₂S₂: C, 65.03; H, 6.06.
35 Found: C, 65.32; H, 6.28.

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E Isomer:

¹H NMR (d₆-benzene): δ 7.35 (d, 2H), 7.19 (s, 1H), 7.05 (d, 2H), 6.72 (d, 2H), 6.54 (d, 2H), 3.21 (s, 3H), 3.14 (s, 3H), 2.39 (q, 2H), 1.09 (t, 3H).

¹³C NMR (d₆-benzene): δ 160.09, 159.16, 135.95, 131.71, 130.61, 130.16, 129.48, 126.88, 114.54, 113.99, 54.64, 54.61, 32.29, 14.33.

10

Z Isomer:

¹H NMR (d₆-benzene): δ 7.67 (d, 2H), 7.58 (d, 2H), 6.90 (s, 1H), 6.83 (d, 2H), 6.80 (d, 2H), 3.30 (s, 3H), 3.28 (s, 3H), 2.26 (q, 2H), 0.94 (t, 3H).

15

¹³C NMR (d₆-benzene): δ 159.98, 159.53, 137.58, 134.03, 132.79, 131.69, 130.45, 113.91, 113.87, 54.79, 54.73, 32.61, 14.25.

20

Example 15**6-Methoxy-2-(4-methoxyphenyl)benzo[b]thiophene**

A solution of *p*-toluenesulfonic acid monohydrate (1.21 g) in toluene (20 mL) was heated to reflux, and water was removed by allowing it to collect in a Dean-Stark trap. A solution of the regioisomeric compounds prepared as described in Example 14 (685 mg, 5.7:1 regioisomeric mixture) in toluene (9 mL) was added to the refluxing acid solution over 1.8 hours. An aliquot of the mixture was analyzed by HPLC, showing a 23.2% *in situ* yield of the title compound.

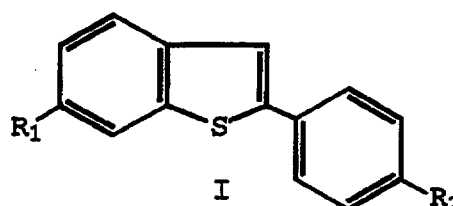
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We claim:

1. A process for preparing a compound of the formula

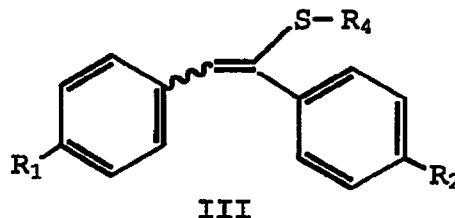
5



wherein:

10 R₁ is hydrogen, C₁-C₄ alkoxy, arylalkoxy, halo, or amino; and

R₂ is hydrogen, C₁-C₄ alkoxy, arylalkoxy, halo, or amino; which comprises cyclizing in the presence of an acid catalyst a compound of the formula



15

wherein:

R₁ and R₂ are as defined above;

R₄ is OSi(R)₃, NR₅R₆, or SR₈;

20 each R is independently C₁-C₆ alkyl, aryl, or arylalkyl;

R₅ and R₆ are independently hydrogen, C₁-C₆ alkyl, arylalkyl, or aryl; or R₅ and R₆ together with the nitrogen atom form a ring selected from piperidine, pyrrolidine, morpholine, and hexamethylimine; and

25 R₈ is C₁-C₆ alkyl, aryl, or arylalkyl.

2. The process of Claim 1 wherein:

R₁ is hydrogen, C₁-C₄ alkoxy, or arylalkoxy; and

R₂ is hydrogen, C₁-C₄ alkoxy, or arylalkoxy.



3. The process of Claim 2 wherein the acid catalyst is selected from the group consisting of methanesulfonic acid, benzenesulfonic acid, 1-naphthalenesulfonic acid, 1-
5 butanesulfonic acid, ethanesulfonic acid, 4-ethylbenzenesulfonic acid, 1-hexanesulfonic acid, 1,5-naphthalenedisulfonic acid, 1-octanesulfonic acid, camphorsulfonic acid, trifluoromethanesulfonic acid, p-toluenesulfonic acid, Nafion®, Amberlyst®, and Amberlite®.
- 10 4. The process of Claim 3 wherein the acid catalyst is selected from the group consisting of methanesulfonic acid, benzenesulfonic acid, camphorsulfonic acid, p-toluenesulfonic acid, Nafion®, Amberlyst®, and Amberlite®.
- 15 5. The process of Claim 4 wherein the acid catalyst is selected from the group consisting of methanesulfonic acid, p-toluenesulfonic acid, Nafion®, Amberlyst®, and Amberlite®.
- 20 6. The process of Claim 5 wherein:
R₄ is OSi(R)₃; and
each R is independently C₁-C₆ alkyl, aryl, or arylalkyl.
- 25 7. The process of Claim 6 wherein R₄ is OTMS, OTES, OTIPS, ODMIPS, ODEIPS, OTDS, OTBDMS, OTBDPS, OTBS, OTPS, ODPMS, or OTEMPS.
- 30 8. The process of Claim 7 wherein R₄ is OTMS, OTES, ODMIPS, ODEIPS, OTBDMS, OTBS, or OTPS.
- 35 9. The process of Claim 8 wherein R₄ is OTMS.
10. The process of Claim 9 wherein the acid catalyst is p-toluenesulfonic acid.
11. The process of Claim 5 wherein:
R₄ is NR₅R₆; and



R₅ and R₆ are independently hydrogen, C₁-C₆ alkyl, arylalkyl, or aryl; or R₅ and R₆ together with the nitrogen atom form a ring selected from piperidine, pyrrolidine, morpholine, and hexamethyimine.

5

12. The process of Claim 11 wherein R₅ and R₆ are independently hydrogen, C₁-C₆ alkyl, or arylalkyl; or R₅ and R₆ together with the nitrogen atom form a ring selected from piperidine and pyrrolidine.

10

13. The process of Claim 12 wherein R₅ and R₆ are methyl, or R₅ is hydrogen and R₆ is benzyl.

14. The process of Claim 13 wherein the acid catalyst is p-toluenesulfonic acid.

15

15. The process of Claim 5 wherein:
R₄ is SR₈; and
R₈ is C₁-C₆ alkyl, aryl, or arylalkyl.

20

16. The process of Claim 15 wherein R₈ is C₁-C₆ alkyl or arylalkyl.

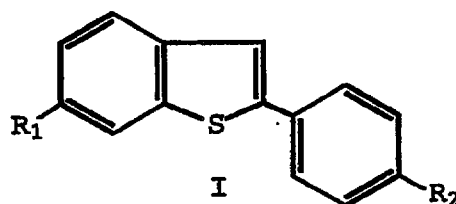
17. The process of Claim 16 wherein R₈ is C₁-C₆ alkyl.

25

18. The process of Claim 17 wherein the acid catalyst is p-toluenesulfonic acid.

19. A process for preparing a compound of the formula

30



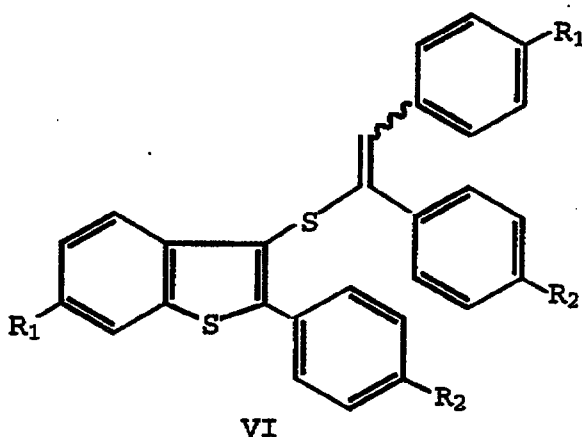
wherein:



R₁ is hydrogen, C₁-C₄ alkoxy, arylalkoxy, halo, amino;
and

R₂ is hydrogen, C₁-C₄ alkoxy, arylalkoxy, halo, amino;
which comprises treating a compound of the formula

5



wherein R₁ and R₂ are as defined above, with an acid.

10 20. The process of Claim 19 wherein:

R₁ is hydrogen, C₁-C₄ alkoxy, or arylalkoxy; and

R₂ is hydrogen, C₁-C₄ alkoxy, or arylalkoxy.

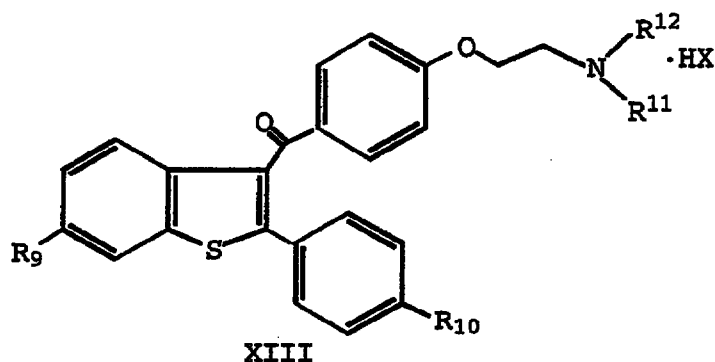
15 21. The process of Claim 20 wherein the acid catalyst is
selected from the group consisting of methanesulfonic acid,
benzenesulfonic acid, camphorsulfonic acid, p-toluenesulfonic
acid, Nafion®, Amberlyst®, and Amberlite®.

20 22. The process of Claim 21 wherein R₁ and R₂ are C₁-C₄
alkoxy.

23. The process of Claim 22 wherein the acid catalyst is p-
toluenesulfonic acid.

25 24. The process of Claim 23 wherein R₁ and R₂ are methoxy.

25. A process for preparing a compound of the formula



wherein:

R₉ is hydrogen, halo, amino, or hydroxyl;

5 R₁₀ is hydrogen, halo, amino, or hydroxyl;

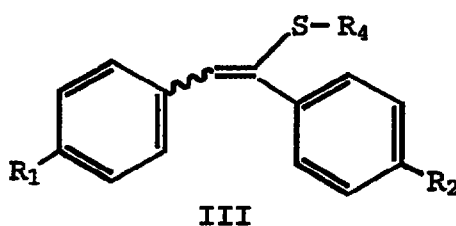
R₁₁ and R₁₂ are independently C₁-C₄ alkyl, or R₁₁ and R₁₂ together with the adjacent nitrogen atom form a heterocyclic ring selected from the group consisting of pyrrolidino, piperidino, hexamethyleneimino, and morpholino;

10 and

HX is HCl or HBr;

comprising the steps of:

(a) cyclizing in the presence of an acid catalyst a compound
15 of the formula



wherein:

R₁ is hydrogen, C₁-C₄ alkoxy, arylalkoxy, halo, or
20 amino;

R₂ is hydrogen, C₁-C₄ alkoxy, arylalkoxy, halo, or amino; and

R₄ is OSi(R)₃, NR₅R₆, or SR₈;

each R is independently C₁-C₆ alkyl, aryl, or arylalkyl;

25 R₅ and R₆ are independently hydrogen, C₁-C₆ alkyl, or aryl, or R₅ and R₆ together with the nitrogen atom form a

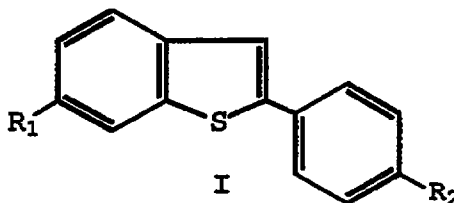


ring selected from piperidine, pyrrolidine, morpholine, and hexamethylimine; and

R₈ is C₁-C₆ alkyl, aryl, or arylalkyl;

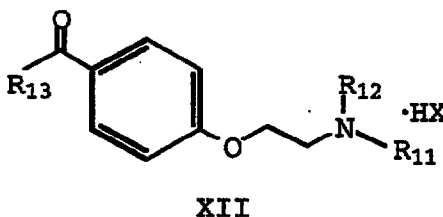
to prepare a benzothiophene compound of the formula

5



wherein R₁ and R₂ are as defined above;

- 10 (b) acylating said benzothiophene compound with an acylating agent of the formula



wherein:

- 15 R₁₁, R₁₂, and HX are as defined previously; and
R₁₃ is chloro, bromo, or hydroxyl; in the presence of BX'₃, wherein X' is chloro or bromo; and

- (c) when R₁ and/or R₂ is C₁-C₄ alkoxy or arylalkoxy,
20 dealkylating one or more phenolic groups of the acylation product of step (b) by reacting with additional BX'₃, wherein X' is as defined above.

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INTERNATIONAL APPLICATION PUBLISHED UNDER THE PATENT COOPERATION TREATY (PCT)

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(21) International Application Number: PCT/US96/09477 (22) International Filing Date: 4 June 1996 (04.06.96) (30) Priority Data: 08/481,015 7 June 1995 (07.06.95) US (60) Parent Application or Grant (63) Related by Continuation US 08/481,015 (CON) Filed on 7 June 1995 (07.06.95) (71) Applicant (for all designated States except US): ELI LILLY & COMPANY [US/US]; Lilly Corporate Center, Indianapolis, IN 46285 (US). (72) Inventors; and (75) Inventors/Applicants (for US only): HOARD, David, W. [US/US]; 440 David Drive, Greenwood, IN 46142 (US). LUKE, Wayne, D. [US/US]; 208 Jennings Street, West Lafayette, IN 47906 (US). (74) Agents: STRODE, Janelle, D. et al.; Eli Lilly & Company, Lilly Corporate Center, Indianapolis, IN 46285 (US).		(51) Designated States: AL, AM, AT, AU, AZ, BB, BG, BR, BY, CA, CH, CN, CZ, DE, DK, EE, ES, FI, GB, GE, HU, IL, IS, JP, KE, KG, KP, KR, KZ, LK, LR, LS, LT, LU, LV, MD, MG, MK, MN, MW, MX, NO, NZ, PL, PT, RO, RU, SD, SE, SG, SI, SK, TJ, TM, TR, TT, UA, UG, US, UZ, VN, ARIPO patent (KE, LS, MW, SD, SZ, UG), Eurasian patent (AM, AZ, BY, KG, KZ, MD, RU, TJ, TM), European patent (AT, BE, CH, DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE), OAPI patent (BF, BJ, CF, CG, CI, CM, GA, GN, ML, MR, NE, SN, TD, TG). Published With international search report.	
(54) Title: PROCESS FOR THE SYNTHESIS OF BENZO[b]THIOPHENES			
(57) Abstract The present invention is directed to new processes for the synthesis of 2-aryl benzo[b]thiophenes.			