

PROCESS FOR PREPARING A PARA-BRANCHED ALKYL-SUBSTITUTED
HYDROXYAROMATIC COMPOUND

PRIORITY

[0001] This application claims the benefit under 35 U.S.C. §119 to Provisional Application Serial No. 61/898,413, filed on October 31, 2013, the contents of which are incorporated by reference herein

BACKGROUND OF THE INVENTION

1. Technical Field

[0002] The present invention generally relates to a process for preparing a para-branched mono-alkyl-substituted hydroxyaromatic compound.

2. Description of the Related Art

[0003] Long chain para-alkylphenols lead to high-performance surfactants that are widely used in the manufacturing of lubricating oil additives. It is well known to prepare alkylphenols by alkylating phenol with branched olefins over ionic exchange resin catalysts.

[0004] Branched para-alkylphenates are a key component for making detergents for use in marine lubricants. One manufacturing process for preparing branched para-alkylphenates involves alkylating regioselectively an excess of phenol with a branched olefin in presence of an ionic exchange resin catalyst. Under the conditions used, the acidic resin heavily favors the formation of the desired para-alkylphenol isomers (95%) at the expense of the ortho-isomer (5%), which has a tendency to drastically decrease the surfactant performance. Subsequent steps of basification of branched para-alkylphenates followed by vulcanization and overbasing treatment of the resulting calcium alkylphenate (CAP) complete the sequence.

[0005] The resulting CAP detergent can contain a substantial amount of the residual monalkylphenol as an ortho alkylphenol isomer (e.g., tetrapropenylphenol ("TPP")). The presence of the bulky aliphatic chain in close proximity to the hydroxyl group may prevent the ortho-alkylphenol to react further, and consequently remain as is in the final product. One potential approach to decrease the amount of this residue is to increase the already high para selectivity (95%) of the phenol alkylation to prepare TPP. By improving the regioselectivity of the alkylation of phenol in favor of the para isomer, less residual TPP may remain in the finished product.

[0006] U.S. Patent No. 3,953,538 discloses an alkylation process in which a stream of an olefinic material is mixed with an acid stream and polymerized to cause formation of a

polymeric diluent for the high strength acid which is initially charged to the alkylation process.

[0007] U.S. Patent No. 4,225,737 (“the ‘737 patent”) discloses a process for the alkylation of an aromatic hydrocarbon with an olefin-acting alkylating agent. The process disclosed in the ‘737 patent involves (a) commingling an aromatic hydrocarbon and a first portion of the olefin-acting alkylating agent at alkylation reaction conditions in a first alkylation reaction zone in contact with a hydrofluoric acid catalyst; (b) separating the effluent from the first alkylation reaction zone into an acid phase and a hydrocarbon phase comprising alkylate and unreacted aromatic hydrocarbon; (c) commingling the hydrocarbon phase with a second portion of the olefin-acting alkylating agent at alkylation reaction conditions in a second alkylation reaction zone in contact with the acid phase transferred from the first to said second reaction zone by maintaining a pressure differential between the first and second zones of from about 5 to about 20 psig.; (d) separating the effluent from the second alkylation reaction zone into an acid phase and a hydrocarbon phase and recycling the former to the first alkylation reaction zone; and, (e) recovering an aromatic alkylation product from the last-mentioned hydrocarbon phase.

[0008] U.S. Patent No. 4,536,301 (“the ‘301 patent”) discloses a surfactant slug used to recover residual oil in subterranean reservoirs. The slug disclosed in the ‘301 patent comprises a mixture of (1) from about 1 to about 10% of a sulfonate of a mixture of mono- and dialkyl-substituted aromatic hydrocarbon which has been obtained by the alkylation of an aromatic hydrocarbon with an olefinic hydrocarbon in the presence of a hydrogen fluoride catalyst; (2) a lower alkyl alcohol which possesses from about 3 to about 6 carbon atoms; and (3) a nonionic cosurfactant comprising an ethoxylated n-alcohol which possesses from about 12 to about 15 carbon atoms.

[0009] U.S. Patent No. 4,816,185 discloses reaction products of C₉-C₃₀ alkylbenzenes with styrene and sulfonated derivatives thereof and processes for preparing such products and derivatives. The sulfonate salts of reaction products are especially useful as detergents.

[0010] U.S. Patent No. 5,750,818 discloses a process for the liquid phase alkylation in an alkylation reactor of a hydrocarbon substrate with an olefinic alkylating agent in the presence of an acid alkylation catalyst at least one hydrocarbon having a lower boiling point than the hydrocarbon substrate and with a substantial stoichiometric excess of the hydrocarbon substrate over the alkylating agent to form a liquid product mixture.

[0011] U.S. Patent No. 6,054,419 discloses a mixture of alkyl aryl sulfonates of superalkalinized alkaline earth metals comprising (a) 50 to 85% by weight of a mono alkyl

phenyl sulfonate with a C14 to C40 linear chain wherein the molar proportion of phenyl sulfonate substituent in position 1 or position 2 is between 0 and 13% and (b) 15 to 50% by weight of a heavy alkyl aryl sulfonate, wherein the aryl radical is phenyl or not, and the alkyl chains are either two linear alkyl chains with a total number of carbon atoms of 16 to 40, or one or a plurality of branched alkyl chains with on average a total number of carbon atoms of 15 to 48.

[0012] U.S. Patent No. 6,551,967 discloses a low overbased alkaline earth metal alkylaryl sulfonate having a Total Base Number of from about 2 to about 30, a dialkylate content of 0% to about 25% and a monoalkylate content of about 75% to about 90% or more, wherein the alkylaryl moiety is alkyltoluene or alkylbenzene in which the alkyl group is a C₁₅-C₂₁ branched chain alkyl group derived from a propylene oligomer are useful as lubricating oil additives.

[0013] U.S. Patent No. 6,989,355 discloses an under-neutralized alkylxylene sulfonic acid composition for enhanced oil recovery processes. The patent also discloses a method for enhancing the recovery of oil from a subterranean reservoir which method employs the under-neutralized alkylxylene sulfonic acid compositions. The under-neutralized alkylxylene sulfonic acid compositions are employed in an aqueous media. The method optionally employs suitable co-surfactants, such as alcohols, alcohol ethers, polyalkylene glycols, poly(oxyalkylene)glycols and/or poly(oxyalkylene)glycol ethers.

[0014] It is desirable to provide an improved process for preparing a para-branched mono-alkyl-substituted hydroxyaromatic compound.

SUMMARY OF THE INVENTION

[0015] In accordance with a first embodiment of the present invention, there is provided a process for preparing an isomeric mixture comprising a major amount of a para-branched mono-alkyl-substituted hydroxyaromatic compound, the process comprising the steps of:

[0016] (a) providing an isomeric mixture comprising a major amount of a para-di(alkyl-substituted)aromatic compound; wherein a first alkyl substituent is a C₃ to C₈ alkyl moiety and a second alkyl substituent is a C_{4+n} to C_{8+n} branched alkyl moiety, wherein n is 0 to 42 and further wherein the second alkyl substituent is at least one carbon atom greater than the first alkyl substituent;

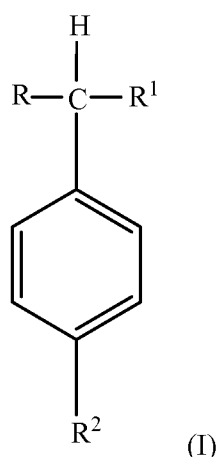
[0017] (b) subjecting the isomeric mixture comprising a major amount of a para-di(alkyl-substituted)aromatic compound to oxidation conditions in the presence of an oxygen-

containing source, thereby converting the first alkyl substituent which is a C₃ to C₈ alkyl moiety to a hydroperoxide-containing substituted moiety to produce an isomeric mixture comprising a major amount of a para-branched alkyl-substituted, hydroperoxide-containing substituted aromatic compound; and

[0018] (c) converting the hydroperoxide-containing substituted moiety to a hydroxyl moiety thereby providing an isomeric mixture comprising a major amount of a para-branched mono-alkyl-substituted hydroxyaromatic compound.

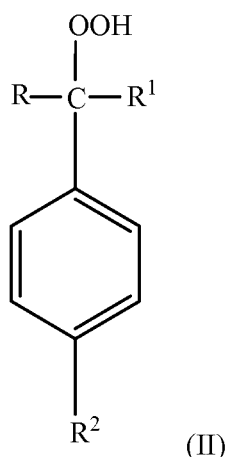
[0019] In accordance with a second embodiment of the present invention, there is provided a process comprising the steps of:

[0020] (a) providing an isomeric mixture comprising a major amount of a para-di(alkyl-substituted)aromatic compound of formula I:



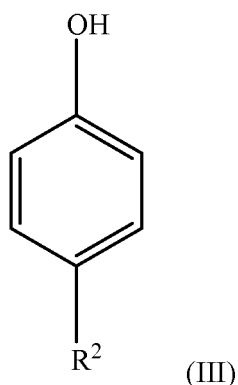
wherein R and R¹ are the same or different alkyl group such that the combined number of carbon atoms is no more than 8 carbon atoms, and R² is a branched alkyl group of from 4 to 50 carbon atoms, wherein R² is at least one carbon atom greater than the combined number of carbon atoms of R and R¹;

[0021] (b) subjecting the isomeric mixture comprising a major amount of the para-di(alkyl-substituted) aromatic compound of formula I to oxidation conditions in the presence of oxygen-containing source to produce an isomeric mixture comprising a major amount of a para-branched alkyl-substituted, hydroperoxide-containing substituted aromatic compound of formula II:



wherein R, R¹ and R² have the aforesaid meanings; and

[0022] (c) converting the hydroperoxide-containing substituted moiety of the para-branched alkyl-substituted, hydroperoxide-containing substituted aromatic compound of formula II to a hydroxyl group thereby providing an isomeric mixture comprising a major amount of a para-branched mono-alkyl-substituted hydroxyaromatic compound of formula III:



wherein R² has the aforesaid meaning.

[0023] The process of the present invention advantageously provides an isomeric mixture containing a major amount of a para-branched mono-alkyl-substituted hydroxyaromatic compound that can be prepared in a simple, cost efficient manner with a relatively low amount of ortho isomer formation. This is an unexpected improvement in that past efforts have not been able to produce such an isomeric mixture with a relatively low amount of the ortho isomer formation.

DETAILED DESCRIPTION OF THE PREFERRED EMBODIMENTS

[0024] Prior to discussing the invention in further detail, the following terms will be defined:

[0025] DEFINITIONS

[0026] As used herein, the following terms have the following meanings, unless expressly stated to the contrary:

[0027] The term “Total Base Number” or “TBN” as used herein refers to the amount of base equivalent to milligrams of KOH in 1 gram of sample. Thus, higher TBN numbers reflect more alkaline products, and therefore a greater alkalinity reserve. The TBN of a sample can be determined by ASTM Test No. D2896-11 issued May 15, 2011 or any other equivalent procedure.

[0028] The term “phenate” means a metal salt of a phenol.

[0029] The term “alkylphenate” means a metal salt of an alkylphenol.

[0030] The term “alkylphenol” means a phenol having an alkyl substituent which has a sufficient number of carbon atoms to impart oil solubility to the phenol.

[0031] The term “lime” refers to calcium hydroxide, also known as slaked lime or hydrated lime.

[0032] The term “metal” means alkali metals, alkaline earth metals, or mixtures thereof.

[0033] The term “alkaline earth metal” refers to calcium, barium, magnesium, and strontium.

[0034] The term “alkali metal” refers to lithium, sodium, potassium, rubidium, and cesium.

[0035] The term “metal base” refers to a metal hydroxide, metal oxide, metal alkoxides and the like and mixtures thereof, wherein the metal is an alkaline earth metal or alkali metal.

[0036] The term “overbased” refers to a class of metal salts or complexes. These materials have also been referred to as “basic”, “superbased”, “hyperbased”, “complexes”, “metal complexes”, “high-metal containing salts”, and the like. Overbased products are metal salts or complexes characterized by a metal content in excess of that which would be present according to the stoichiometry of the metal and the particular acidic organic compound reacted with the metal, e.g., a carboxylic acid. Suitable overbasing metals include alkaline earth metals such as magnesium, calcium, barium, and strontium. Suitable overbasing metals can be provided from the corresponding metal hydroxides, for example, calcium hydroxide and magnesium hydroxide provide the source for the alkaline earth metals calcium and magnesium, respectively. Additional overbasing can be achieved by the addition of acidic overbasing compounds for example, carbon dioxide and boric acid.

[0037] The terms “alkenyl succinic acid or anhydride” and “alkyl succinic acid or anhydride” may be used interchangeably.

[0038] The term “isomerized alpha olefin (IAO)” as used herein refers to an alpha olefin that has been subjected to isomerization conditions which results in an alteration of the distribution of the olefin species present and/or the introduction of alkyl branching along the alkyl chain.

[0039] Partially Branched Linear Olefins – The term “partially branched linear olefins” refers to a class of linear olefins comprising less than one alkyl branch per straight chain containing the double bond, wherein the alkyl branch may be a methyl group or higher. Partially branched linear olefins may also contain double-bond isomerized olefin.

[0040] Branched Olefins – The term “branched olefins” refers to a class of olefins comprising one or more alkyl branches per linear straight chain containing the double bond, wherein the alkyl branch may be a methyl group or higher.

[0041] Double-Bond Isomerized linear Olefins - The term “double-bond isomerized linear olefins” refers to a class of linear olefins comprising more than 5% of olefins in which the carbon-carbon double bond is not terminal (i.e., the double bond is not located between the first and second carbon atoms of the chain).

[0042] The present invention is directed to a process for preparing an isomeric mixture comprising a major amount of a para-branched alkyl-substituted hydroxyaromatic compound. In general, the process involves the steps of (a) providing an isomeric mixture comprising a major amount of a para-di(alkyl-substituted)aromatic compound; wherein a first alkyl substituent is a C₃ to C₈ alkyl moiety and a second alkyl substituent is a C_{4+n} to C_{8+n} branched alkyl moiety, wherein n is 0 to 42 and further wherein the second alkyl substituent is at least one carbon atom greater than the first alkyl substituent; (b) subjecting the isomeric mixture comprising a major amount of a para-di(alkyl-substituted)aromatic compound to oxidation conditions in the presence of an oxygen-containing source, thereby converting the first alkyl substituent which is a C₃ to C₈ alkyl moiety to a hydroperoxide-containing substituted moiety to produce an isomeric mixture comprising a major amount of a para-branched alkyl-substituted, hydroperoxide-substituted aromatic compound; and (c) converting the hydroperoxide-containing substituted moiety to a hydroxyl group thereby providing an isomeric mixture comprising a major amount of a para-branched alkyl-substituted hydroxyaromatic compound.

[0043] In step (a), an isomeric mixture comprising a major amount of a para-di(alkyl-substituted)aromatic compound; a first alkyl substituent is a C₃ to C₈ alkyl moiety and a

second alkyl substituent is a C_{4+n} to C_{8+n} branched alkyl moiety, wherein n is 0 to 42 and further wherein the second alkyl substituent is at least one carbon atom greater than the first alkyl substituent is provided. In general, the isomeric mixture is obtained by alkylating a monoalkylaromatic compound with one or more branched alkylating agents in the presence of an acidic catalyst under reaction conditions sufficient to form an isomeric mixture of dialkylated benzenes which is, as noted hereinbefore, enriched in the para dialkylated benzene isomer and further enriched in the para-2-phenylalkane and para-3-phenylalkane isomers.

[0044] In general, the starting monoalkylaromatic compound may be obtained commercially. Suitable types of monoalkylaromatic compounds include, but are not limited to, cumene and the like and mixtures thereof. Alternatively, the starting monoalkylaromatic compound can be prepared by methods well known in the art, e.g., contacting benzene with a suitable alkylating agent under reaction conditions such that a monoalkylaromatic is formed. For example, the alkylating agent may be propylene, butene, hexene or octene, thereby producing, respectively, cumene, butylbenzene, hexylbenzene, or octylbenzene.

[0045] The one or more branched alkylating agents for use in alkylating the monoalkylaromatic compound may include one or more branched alpha olefins. Suitable one or more branched alpha olefins include polyolefins which may be derived from C_3 or higher monoolefins (i.e., propylene oligomers, butylenes oligomers, or co-oligomers etc.). In one embodiment, a mixture of branched alpha olefins includes propylene oligomers or butylenes oligomers or mixtures thereof.

[0046] In one embodiment of the present invention, the one or more branched olefins include normal alpha olefins which have been isomerized using, for example, a solid or liquid acid catalyst. One type of acidic catalyst is a solid catalyst having at least one metal oxide and having an average pore size of less than 5.5 angstroms. Another type is a molecular sieve with a one-dimensional pore system, such as SM-3, MAPO-11, SAPO-11, SSZ-32, ZSM-23, MAPO-39, SAPO-39, ZSM-22 and SSZ-20. Other possible solid acidic catalysts useful for isomerization include ZSM-35, SUZ4, NU-23, NU-87 and natural or synthetic ferrierites. These molecular sieves are well-known in the art and are discussed in Rosemarie Szostak's Handbook of Molecular Sieves (New York, Van Nostrand Reinhold, 1992) and in U.S. Pat. No. 5,282,858, which is hereby incorporated by reference for all purposes. Another type of isomerization catalyst that can be used is iron pentacarbonyl ($Fe(CO)_5$).

[0047] The isomerization process may be carried out in batch or continuous mode. The process temperatures can range from 50°C to 250°C. In the batch mode, a typical

method is to use a stirred autoclave or glass flask, which may be heated to the desired reaction temperature. A continuous process is most efficiently carried out in a fixed bed process. Space rates in a fixed bed process can range from 0.1 to 10 or more WHSV. In a fixed bed process, the catalyst is charged to the reactor and activated or dried at a temperature of at least 150°C under vacuum or flowing inert, dry gas. After activation, the catalyst is cooled to the desired reaction temperature and a flow of the olefin is introduced. The reactor effluent containing the partially branched, isomerized olefin is collected. The resulting partially-branched isomerized olefin contains a different olefin distribution (alpha-olefin, beta-olefin, internal-olefin, trisubstituted olefin and vinylidene-olefin) and branching content than the un-isomerized olefin and conditions are chosen in order to obtain the appropriate structure regarding the level of double bonds between carbon 1 and carbon 2 of the alkyl chain of the olefin (alpha-olefin content).

[0048] Suitable normal alpha olefins include, but are not limited to, 1-hexene, 1-nonene, 1-decene, 1-dodecene and the like and mixtures thereof. In one embodiment, the branched alkylating agent is a mixture of normal alpha olefins selected from olefins having from about 10 to about 30 carbon atoms per molecule. In one embodiment, the branched alkylating agent is a mixture of normal alpha olefins selected from olefins having from about 12 to about 30 carbon atoms per molecule. In one embodiment, branched alkylating agent is a mixture of normal alpha olefins selected from olefins having from about 18 to about 30 carbon atoms per molecule. In one embodiment, the branched alkylating agent is a mixture of normal alpha olefins selected from olefins having from about 20 to about 24 carbon atoms per molecule.

[0049] Alkylation of the monoalkylaromatic compound with the one or more branched alkylating agents is advantageously carried out in the presence of an acidic alkylation catalyst. Useful acidic alkylation catalysts include, by way of example, zeolite catalysts, Lewis and Brønsted acid catalysts, solid acid catalysts and the like and mixtures thereof. Suitable zeolite catalysts include natural zeolites, synthetic zeolites, and the like and mixtures thereof. In particular, zeolite, which is an inorganic crystalline porous compound containing silicon and aluminum, is a solid acid substance suitable for the present invention in terms of heat resistance and the target selectivity for cumene. Zeolite preferably has pores having approximately the same size as the molecular size of cumene and has a 6, 8, 10, 11, 12, or 14-membered ring structure, and more preferably a 12-membered ring structure. Examples of zeolite having a 12-membered ring structure include a Y type, a USY type, a

mordenite type, a dealuminated mordenite type, a beta type, an MCM-22 type, and an MCM-56 type.

[0050] In one embodiment, suitable zeolites include Y zeolite, beta, SSZ-25, SSZ-26, and SSZ-33. Other possible catalysts include L zeolite, mordenite, boggsite, cloverite, VPI-5, MCM-41, MCM-36, SAPO-8, SAPO-5, MAPO-36, SAPO40, SAPO-41, MAPSO-46, CoAPO-50, hexagonal faujasite (EMC-2), gmelinite, mazzite (omega zeolite), offretite, ZSM-18, ZSM-12. Some of these catalysts are discussed in Rosemarie Szostak's Handbook of Molecular Sieves (New York, Van Nostrand Reinhold, 1992).

[0051] In one embodiment, the acidic catalyst is an acid mordenite zeolite. Any acid mordenite zeolite may be used as the alkylation catalyst in the alkylation step, provided that the isomeric mixture is formed comprising a major amount of the para-di(alkyl-substituted)aromatic compound. A suitable acid mordenite zeolite catalyst and its preparation are described in U.S. Pat. No. 5,004,841, the contents of which are incorporated herein by reference. Typically these zeolite catalysts may be purchased from, for example, BASF, W.R. Grace & Co. and the like. Useful Lewis acid catalysts include, but are not limited to, aluminum trichloride, aluminum tribromide, aluminum triiodide, boron trifluoride, boron tribromide, boron triiodide, Amberlyst® 36 and Amberlyst® 70, and the like.

[0052] Useful acidic clays may be derived from naturally occurring or synthetic materials. One skilled in the art would realize that there are a number of such clays that are known to be alkylation catalysts. Examples of such acidic clays include montmorillonite, laponite, and saponite. Pillared clays may also be used as catalysts.

[0053] The catalysts used in the process of the present invention may be shaped or formed into tablets, extrudates or any other shape, such as beads, using procedures well known in the prior art. The preparation of extrudates requires the presence of a binder, such as alumina. The tableted catalysts do not require the presence of a binder, but a binder may be present in a tableted zeolite catalyst. The crystalline zeolite powder may be compressed to form a tablet.

[0054] In general, the acidic catalyst is present in an amount ranging from about 1 wt. % to about 70 wt. %, based on the total weight of the alkylation reaction mixture. In one embodiment, the acidic catalyst is present in an amount ranging from about 5 to about 50 wt. %, based on the total weight of the alkylation reaction mixture.

[0055] The reaction conditions for the alkylation depend upon the type of catalyst used, and any suitable set of reaction conditions that result in an isomeric mixture comprising a major amount of a para-di(alkyl-substituted)aromatic compound. In general, the reaction

temperature will be the boiling temperature of the light solvent. In one embodiment, the reaction temperature for the alkylation reaction will be in the range of about 0°C to about 150°C. In another embodiment, the reaction temperature for the alkylation reaction will be in the range of about 90°C to about 140°C. The reaction pressure will generally be atmospheric, although higher or lower pressures may be employed. The alkylation process can be practiced in a batchwise, continuous or semi-continuous manner.

[0056] The molar ratio of the monoalkylaromatic compound to the one or more branched alkylating agents may vary from about 0.5:2 to about 2:0.5. In one embodiment, the molar ratio of the monoalkylaromatic compound to the one or more branched alkylating agents is in the range of about 2:1 to about 1:2. In another embodiment, the molar ratio of the monoalkylaromatic compound to the one or more branched alkylating agents is in the range of about 1.1:1 to about 1:1.1.

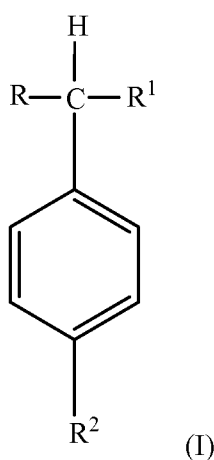
[0057] The residence time in the reactor is a time that is sufficient to convert a substantial portion of the one or more branched alkylating agents to a para-di(alkyl)substituted aromatic compound. In one embodiment, the time required is generally from about 2 hours to about 30 hours. A more precise residence time may be determined by those skilled in the art using batch stirred tank reactors to measure the kinetics of the alkylation process.

[0058] The alkylation reaction may be carried out neat or in the presence of a solvent which is inert to the reaction of the monoalkylaromatic compound and the one or more branched alkylating agents. When employed, a typical solvent is hexane.

[0059] Upon completion of the reaction, the desired isomeric mixture is obtained comprising a major amount of a para-di(alkyl-substituted)aromatic compound. In one embodiment, the isomeric mixture will contain about 60 mole % or greater of the para-di(alkyl-substituted)aromatic compound with the remaining isomeric mixture comprising ortho isomers of the di(alkyl-substituted)aromatic compound. In another embodiment, the isomeric mixture will contain 80 mole % or greater of the para-di(alkyl-substituted)aromatic compound with the remaining isomeric mixture comprising ortho isomers of the di(alkyl-substituted)aromatic compound. In yet another embodiment, the isomeric mixture will contain 90 mole % or greater of the para-di(alkyl-substituted)aromatic compounds with the remaining isomeric mixture comprising ortho isomers of the di(alkyl-substituted)aromatic compound. In still yet another embodiment, the isomeric mixture will contain 95 mole % or greater of the para-di(alkyl-substituted)aromatic compounds with the remaining isomeric mixture comprising ortho isomers of the di(alkyl-substituted)aromatic compound. In still yet

another embodiment, the resulting isomeric mixture will contain 99 mole % or greater of the para-di(alkyl-substituted)aromatic compounds with the remaining isomeric mixture comprising ortho isomers of the di(alkyl-substituted)aromatic compound. In one embodiment, the isomeric mixture will contain no more than about four structural isomers of the para-di(alkyl-substituted) aromatic compound. In one embodiment, the isomeric mixture will contain no more than about two structural isomers of the para-di(alkyl-substituted)aromatic compound.

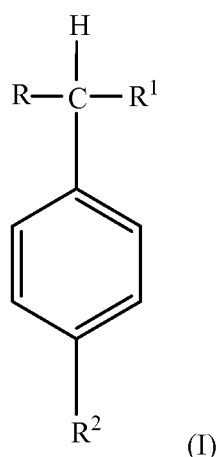
[0060] In one embodiment, an isomeric mixture is obtained comprising a major amount of a para-di(alkyl-substituted)aromatic compound of formula I:



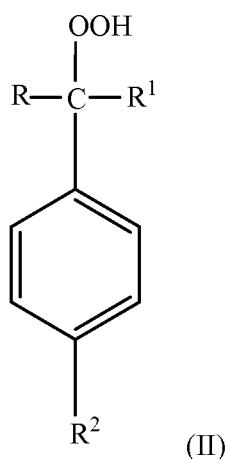
wherein R and R¹ are the same or different alkyl group such that the combined number of carbon atoms is no more than 8 carbon atoms, and R² is a branched alkyl group of from 4 to 50 carbon atoms, wherein R² is at least one carbon atom greater than the combined number of carbon atoms of R and R¹. In one embodiment, R² is a branched group derived from one or more branched alpha olefins as discussed hereinabove. In one embodiment, the mixture will contain no more than about four structural isomers of formula I. In one embodiment, the mixture will contain no more than about two structural isomers of formula I.

[0061] Step (b) of the process of the present invention subjects the isomeric mixture comprising a major amount of a para-di(alkyl-substituted)aromatic compound of step (a) to oxidation conditions in the presence of an oxygen-containing source, thereby converting the first alkyl substituent which is a C₃ to C₈ alkyl moiety to a hydroperoxide-containing substituted moiety to produce an isomeric mixture comprising a major amount of a para-branched alkyl-substituted, hydroperoxide-containing substituted aromatic compound.

[0062] In one embodiment, the isomeric mixture comprising a major amount of the para-di(alkyl-substituted)aromatic compound of formula I:



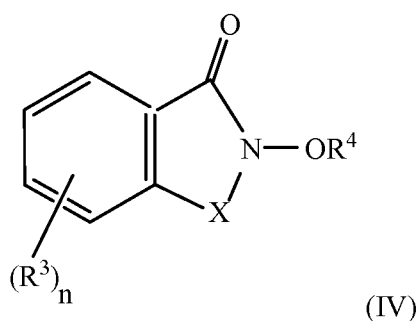
wherein R, R¹ and R² have the aforesaid meanings, is subjected to oxidation conditions in the presence of an oxygen-containing source to produce an isomeric mixture comprising a major amount of a para-branched alkyl-substituted, hydroperoxide-containing substituted aromatic compound of formula II:



wherein R, R¹ and R² have the aforesaid meanings. This oxidation process can take place in one or more oxidation reactor(s). The oxidation reactor(s) may be batch reactor(s) or continuous reactor(s).

[0063] The isomeric mixture comprising a major amount of the para-di(alkyl-substituted)aromatic compound is oxidized in the presence of an oxygen-containing source such as a gas, for example, air. While a solvent may be added to the isomeric mixture, oxidation is typically carried out in the absence of any solvent, other than the mixture itself.

[0064] Oxidation can also take place in the presence of at least one substituted cyclic imide of formula IV:



in which X represents a carbonyl (CO) group or a sulfonyl (SO₂) group, n is 0, 1, 2, 3 or 4, R^{sup.3} is one or several groups selected from a hydrogen atom, a halogen atom, an alkyl group, an alkoxy group, an amino group and R⁴ is a hydrogen atom, an alkaline metal cation or an alkaline earth metal cation. Representative examples of the cyclic imide of formula (IV) include N-hydroxyphthalimide, 4-amino-N-hydroxyphthalimide, 3-amino-N-hydroxyphthalimide, tetrabromo-N-hydroxyphthalimide, tetrachloro-N-hydroxyphthalimide, N-hydroxysaccharin and mixtures thereof. In one embodiment, oxidation can take place in the presence of N,N',N''-trihydroxyisocyanuric acid (THICA). The cyclic imide of formula (IV) or THICA may be used in an amount of from about 0.0001 mol % to about 15 mol %, or from about 0.001 mol % to about 10 mol %, relative to the amount of the para-di(alkyl-substituted) aromatic compound.

[0065] In one embodiment, the cyclic imide of formula (IV) or THICA is used in a form in which it has been deposited or fixed chemically on a support, for example, silica, alumina, a zeolite, a polymer (e.g. polystyrene resin) or a mixture thereof.

[0066] The oxidation mixture may also include a free radical initiator, such as a peroxy compound or azo compound. Representative examples of such compounds include, but are not limited to, cumene hydroperoxide, sec-butylbenzene hydroperoxide and the like and mixtures thereof. If used, the free radical initiator can be present in an amount ranging from about 0.1 wt. % to about 5 wt. %.

[0067] Suitable oxidation conditions for step (b) include a temperature ranging from about 70°C to about 300°C, or from about 90°C to about 130°C, or from about 100°C to about 125°C, or from about 105°C to about 120°C and at a pressure of from about 101 to about 2026 kPa (about 1 to about 20 atmospheres), or from about 101 kPa to about 500 kPa (1 to 5 atmospheres), or from about 101 kPa to about 150 kPa (1 to 1.5 atmospheres).

[0068] A basic agent, such as an alkali metal carbonate, e.g., sodium carbonate, an alkali metal bicarbonate, e.g., sodium bicarbonate, or ammonia may also be added to react with any acidic by-products that may form during oxidation. In addition, an aqueous phase

may be introduced, which can help dissolve basic compounds, such as sodium carbonate. The per-pass conversion in the oxidation step is preferably kept below 50%, to minimize the formation of byproducts. The oxidation reaction can be conveniently conducted in a catalytic distillation unit and the hydroperoxide of formula (II) produced may be concentrated by distilling off the unreacted alkylbenzene of formula (I) prior to the cleavage step.

[0069] Optionally, oxidation of the mixture may also take place in the presence of a transition metal co-catalyst, such as a cobalt, manganese or copper-containing catalyst, e.g., cobaltous oxide (formula Co_3O_4).

[0070] Oxidation is an exothermic reaction, and the heat of reaction is removed from the oxidation reaction mixture during the reaction. For example, heat is removed by vaporization of hydrocarbon, product and water, if water is present in the oxidation mixture, into the air passing through the reactor(s). If necessary, external heat exchangers can be used to cool the vaporized products and recirculate them to the oxidation reactor(s).

[0071] In one embodiment, oxidation is carried out in an entrained flow reactor. For example, a bubble column reactor is employed. This type of reactor is well known in the field of chemical engineering. In particular, this type of reactor has multiple injection zones, which improves mass transfer.

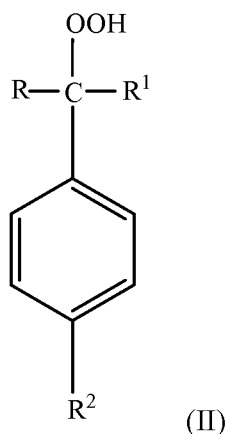
[0072] Step (c) of the process of the present invention converts the hydroperoxide-containing substituted moiety of the para-branched alkyl-substituted, hydroperoxide-containing substituted aromatic compound to a hydroxyl moiety thereby providing an isomeric mixture comprising a major amount of a para-branched mono-alkyl-substituted hydroxyaromatic compound, i.e., the hydroperoxide-containing substituted moiety is cleaved into a hydroxyl moiety. In general, cleavage is carried out in a cleavage reactor or reactor zone, operating, for example, as a plug flow reactor, a plug flow reactor with recycle or a continuous stirred tank reactor.

[0073] This cleavage reaction is effected by contacting the isomeric mixture comprising a major amount of the para-branched alkyl-substituted, hydroperoxide-containing substituted aromatic compound in a liquid phase with a catalyst at a temperature ranging from about 20°C to about 200°C, or from about 40°C to about 120°C, a pressure of about 101 kPa to about 5 MPa (about 1 to about 50 atmospheres), or from about 101 kPa to 3 MPa (about 1 to about 30 atmospheres) or from about 101 kPa to about 1 MPa (about 1 to 10 atmospheres) or from about 101 kPa to about 500 kPa (about 1 to about 5 atmospheres), and a liquid hourly space velocity (LHSV) based on the hydroperoxide of about 0.1 hr^{-1} to about 10 hr^{-1} , or about 1 hr^{-1} to about 5 hr^{-1} . The isomeric mixture comprising a major amount of the para-branched

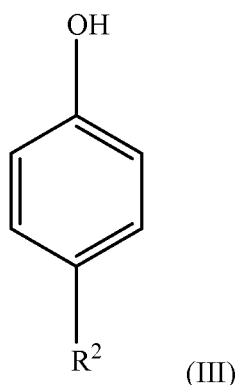
alkyl-substituted, hydroperoxide-containing substituted aromatic compound can be diluted in an organic solvent inert to the cleavage reaction, such as a ketone, e.g., acetone, methyl ethyl ketone and the like, to assist in heat removal. The cleavage reaction is conveniently conducted in a catalytic distillation unit.

[0074] The catalyst employed in the cleavage step can be a homogeneous catalyst or a heterogeneous catalyst. Suitable homogeneous cleavage catalysts include, but are not limited to, sulfuric acid, perchloric acid, phosphoric acid, hydrochloric acid, p-toluenesulfonic acid, ferric chloride, boron trifluoride, sulfur dioxide, sulfur trioxide and the like and mixtures thereof. In one preferred embodiment, sulfuric acid is the homogeneous cleavage catalyst. Suitable heterogeneous catalysts include a smectite clay, such as an acidic montmorillonite silica-alumina clay, as described in U.S. Pat. No. 4,870,217, the contents of which are incorporated herein by reference.

[0075] In one embodiment, the hydroperoxide-containing substituted moiety of the para-branched alkyl-substituted, hydroperoxide-containing substituted aromatic compound of formula II:



wherein R, R¹ and R² have the aforesaid meanings is converted to a hydroxyl moiety thereby providing an isomeric mixture comprising a major amount of a para-branched mono-alkyl-substituted hydroxyaromatic compound of formula III:



wherein R^2 has the aforesaid meanings.

[0076] The resulting isomeric mixture obtained from the process of the present invention comprises a major amount of a para-branched mono-alkyl-substituted hydroxyaromatic compound. In one embodiment, the resulting isomeric mixture will contain about 60 mole % or greater of the para-branched mono-alkyl-substituted hydroxyaromatic compound with the remaining isomeric mixture comprising ortho isomers of the mono-alkyl-substituted hydroxyaromatic compound. In another embodiment, the resulting isomeric mixture will contain 80 mole % or greater of the para-branched mono-alkyl-substituted hydroxyaromatic compound with the remaining isomeric mixture comprising ortho isomers of the mono-alkyl-substituted hydroxyaromatic compound. In yet another embodiment, the resulting isomeric mixture will contain 90 mole % or greater of the para-branched mono-alkyl-substituted hydroxyaromatic compound with the remaining isomeric mixture comprising ortho isomers of the mono-alkyl-substituted hydroxyaromatic compound. In still yet another embodiment, the resulting isomeric mixture will contain 95 mole % or greater of the para-branched mono-alkyl-substituted hydroxyaromatic compound with the remaining isomeric mixture comprising ortho isomers of the mono-alkyl-substituted hydroxyaromatic compound. In still yet another embodiment, the resulting isomeric mixture will contain 99 mole % or greater of the para-branched mono-alkyl-substituted hydroxyaromatic compound with the remaining isomeric mixture comprising ortho isomers of the mono-alkyl-substituted hydroxyaromatic compound. In one embodiment, the resulting isomeric mixture will contain no more than about four structural isomers of para-branched mono-alkyl-substituted hydroxyaromatic compound. In one embodiment, the resulting isomeric mixture will contain no more than about two structural isomers of para-branched mono-alkyl-substituted hydroxyaromatic compound.

[0077] If desired, the para-branched mono-alkyl-substituted hydroxyaromatic compound may be subsequently sulfurized and neutralized in any order to provide a salt of a

sulfurized para-branched mono-alkyl-substituted hydroxyaromatic composition. The sulfurization and neutralization steps may be performed in any order so as to provide the salt of the sulfurized para-branched mono-alkyl-substituted hydroxyaromatic composition. Alternatively, the neutralization and sulfurization steps may be carried out simultaneously.

[0078] In general, sulfurization is carried out by contacting the para-branched mono-alkyl-substituted hydroxyaromatic compound with a sulfur source which introduces S_x bridging groups between the para-branched mono-alkyl-substituted hydroxyaromatic compounds, wherein x is 1 to 7, in the presence of a base. Any suitable sulfur source can be used such as, for example, elemental sulfur or a halide thereof such as sulfur monochloride or sulfur dichloride, hydrogen sulfide, sulfur dioxide and sodium sulfide hydrates. The sulfur can be employed either as molten sulfur or as a solid (e.g., powder or particulate) or as a solid suspension in a compatible hydrocarbon liquid.

[0079] The base catalyzes the reaction to incorporate sulfur onto the para-branched alkyl-substituted hydroxyaromatic compound. A suitable base includes, but is not limited to, NaOH, KOH, Ca(OH)₂ and the like and mixtures thereof.

[0080] The base is generally employed at from about 0.5 to about 5 moles per mole of the para-branched alkyl-substituted hydroxyaromatic compound in the reaction system. In one embodiment, the base is employed at from about 1 to about 1.5 moles per mole of the para-branched mono-alkyl-substituted hydroxyaromatic compound in the reaction system. The base can be added to the reaction mixture as a solid or a liquid.

[0081] Sulfur is generally employed at from about 0.5 to about 4 moles per mole of the para-branched alkyl-substituted hydroxyaromatic compound in the reaction system. In one embodiment, sulfur is employed at from about 0.8 to 2 moles per mole of the para-branched mono-alkyl-substituted hydroxyaromatic compound. In one embodiment, sulfur is employed at from about 1 to 1.5 moles per mole of para-branched mono-alkyl-substituted hydroxyaromatic compound.

[0082] The temperature range in which the sulfurization reaction is carried out is generally from about 120°C to about 200°C. In one embodiment, the temperature range is from about 160°C to about 180°C. The reaction can be conducted under atmospheric pressure (or slightly lower) or at elevated pressures. In one embodiment the reaction is carried out under vacuum to facilitate H₂S elimination. The exact pressure developed during the reaction is dependent upon such factors as the design and operation of the system, the reaction temperature, and the vapor pressure of the reactants and products and it may vary

during the course of the reaction. In one embodiment, the process pressures are at atmospheric to about 20 mm Hg.

[0083] Neutralization of the sulfurized or unsulfurized para-branched mono-alkyl-substituted hydroxyaromatic compound may be carried out in a continuous or batch process by any method known to a person skilled in the art. Numerous methods are known in the art to neutralize the sulfurized or unsulfurized para-branched mono-alkyl-substituted hydroxyaromatic compound and to produce basic phenates by incorporation of a source of base. In general, neutralization can be carried out by contacting the sulfurized or unsulfurized para-branched alkyl-substituted hydroxyaromatic compound with a metal base under reactive conditions, preferably in an inert-compatible liquid hydrocarbon diluent. If desired, the reaction can be conducted under an inert gas, typically nitrogen. The metal base may be added either in a single addition or in a plurality of additions at intermediate points during the reaction.

[0084] Suitable metal basic compounds include hydroxides, oxides or alkoxides of the metal such as (1) an alkali metal salt derived from a metal base selected from an alkali hydroxide, alkali oxide or an alkali alkoxide, or (2) an alkaline earth metal salt derived from a metal base selected from an alkaline earth hydroxide, alkaline earth oxide or alkaline earth alkoxide. Representative examples of metal basic compounds with hydroxide functionality include lithium hydroxide, potassium hydroxide, sodium hydroxide, magnesium hydroxide, calcium hydroxide, barium hydroxide, aluminum hydroxide and the like. Representative examples of metal basic compounds with oxide functionality include lithium oxide, magnesium oxide, calcium oxide, barium oxide and the like. In one embodiment, the alkaline earth metal base is slaked lime (calcium hydroxide), because of its handling convenience and cost versus, for example, calcium oxide.

[0085] Neutralization is typically conducted in a suitable solvent or diluents oil, such as toluene, xylene and commonly with a promoter such as an alcohol, e.g., a C₁ to C₁₆ alcohol, such as methanol, decyl alcohol, or 2-ethyl hexanol; a diol, e.g., C₂ to C₄ alkylene glycols, such as ethylene glycol; and/or carboxylic acids. Suitable diluent oils include naphthenic oils and mixed oils, e.g., paraffinic oils such as 100 neutral oil. The quantity of solvent or diluent oil used is such that the amount of solvent or oil in the final product constitutes from about 25% to about 65% by weight of the final product, preferably from about 30% to about 50%. For example, the source of alkaline earth metal is added in excess as a slurry (i.e., as a pre-mixture of source of an alkaline earth metal lime, solvent or diluent

oil) and then reacted with the sulfurized or unsulfurized para-branched alkyl-substituted hydroxyaromatic compound.

[0086] The neutralization reaction between the metal base and the sulfurized or unsulfurized para-branched mono-alkyl-substituted hydroxyaromatic compound is typically conducted at temperatures above room temperature (20°C). In one embodiment, neutralization can be carried out at a temperature between about 20°C and about 150°C. It is however preferred to carry the neutralization at low temperature. In one embodiment, neutralization can be carried out at a temperature of between about 25°C and about 30°C. The neutralization reaction itself should take place for a period of time of from about 5 to about 60 minutes. If desired, the neutralization reaction is carried out in the presence of a promoter such as ethylene glycol, formic acid, acetic acid, and the like and mixtures thereof.

[0087] Upon completion of the sulfurizing and neutralizing of the para-branched mono-alkyl-substituted hydroxyaromatic compound, a neutral salt of a sulfurized para-branched mono-alkyl-substituted hydroxyaromatic composition is obtained. If desired, the neutral salt of a sulfurized para-branched mono-alkyl-substituted hydroxyaromatic composition can be overbased to provide an overbased salt of a sulfurized para-branched alkyl-substituted hydroxyaromatic composition. Overbasing can be carried out either during or after one of the sulfurization and neutralization steps and by any method known by a person skilled in the art. Alternatively, sulfurization, neutralization and overbasing can be carried out simultaneously. In general, the overbasing is carried out by reaction with an acidic overbasing compound such as, for example, carbon dioxide or boric acid. In one embodiment, an overbasing process is by way of carbonation, i.e., a reaction with carbon dioxide. Such carbonation can be conveniently effected by addition of solvents such as aromatic solvents, alcohols or a polyols, typically an alkylene diol, e.g., ethylene glycol. Conveniently, the reaction is conducted by the simple expedient of bubbling gaseous carbon dioxide through the reaction mixture. Excess solvents and any water formed during the overbasing reaction can be conveniently removed by distillation either during or after the reaction.

[0088] In one embodiment, the overbasing reaction is carried out in a reactor by reacting the salt of the sulfurized para-branched mono-alkyl-substituted hydroxyaromatic composition with a source of an alkaline earth metal such as lime (i.e., an alkaline earth metal hydroxide) in the presence of carbon dioxide, and in the presence of an aromatic solvent (e.g., xylene), and a hydrocarbyl alcohol such as methanol. Conveniently, the reaction is conducted by the simple expedient of bubbling gaseous carbon dioxide through the reaction

mixture. The carbon dioxide may be introduced over a period of about 1 hour to about 3 hours, at a temperature ranging from about 30°C to about 60°C. The degree of overbasing may be controlled by the quantity of the source of an alkaline earth metal, carbon dioxide and the reactants added to the reaction mixture and the reaction conditions used during the carbonation process.

[0089] In another embodiment, the overbasing reaction can be carried out between 140°C and 180°C in the presence of a polyol, typically an alkylene diol, e.g., ethylene glycol, and/or alkanols, e.g., C₆ to C₁₆ alkanols, such as decyl alcohols, 2-ethyl hexanol. Excess solvent and any water formed during the overbasing reaction can be conveniently removed by distillation either during or after the reaction.

[0090] The overbased salt of a sulfurized para-branched mono-alkyl-substituted hydroxyaromatic composition may have a TBN of from about 50 to about 500.

[0091] The resulting neutral or overbased salt of the sulfurized para-branched mono-alkyl-substituted hydroxyaromatic composition is advantageously employed in a lubricating oil composition comprising at least a major amount of an oil of lubricating viscosity. The lubricating oil compositions may also contain other conventional additives that can impart or improve any desirable property of the lubricating oil composition in which these additives are dispersed or dissolved. Any additive known to a person of ordinary skill in the art may be used in the lubricating oil compositions disclosed herein. Some suitable additives have been described in Mortier et al., "Chemistry and Technology of Lubricants," 2nd Edition, London, Springer, (1996); and Leslie R. Rudnick, "Lubricant Additives: Chemistry and Applications," New York, Marcel Dekker (2003), both of which are incorporated herein by reference. For example, the lubricating oil compositions can be blended with antioxidants, anti-wear agents, detergents such as metal detergents, rust inhibitors, dehazing agents, demulsifying agents, metal deactivating agents, friction modifiers, pour point depressants, antifoaming agents, co-solvents, package compatibilisers, corrosion-inhibitors, ashless dispersants, dyes, extreme pressure agents and the like and mixtures thereof. A variety of the additives are known and commercially available. These additives, or their analogous compounds, can be employed for the preparation of the lubricating oil compositions of the invention by the usual blending procedures.

[0092] The following non-limiting examples are illustrative of the present invention.

[0093] In the examples, cumene (99% pure) was purchased from Acros and was used as received.. The C₉-C₁₄ branched olefin blend was from ChevronPhillips Company LLC, Cedar Bayou, Texas. The mordenite extrudates were from Engelhardt (now BASF) and were

pre-dried (approximately 120°C) overnight prior to use. The NMR experiments were performed on Bruker Advance III. Deuterated solvents for NMR were purchased from ACROS and stored under nitrogen upon receiving. GC/MS analytical experiments were run on a Hewlett Packard 5973 GC-Mass selective detector equipped with a Zebtron capillary GC column ZB-1HT Inferno 15 m (length) x 0.25 mm (I.D) x 0.10 µM (thickness).

EXAMPLE 1

[0094] Preparation of para-alkyl cumene isomers.

[0095] Reactions were performed in a round bottom neck flask equipped with a reflux condenser. According to the experimental procedure described in US patent 5,468,407, a mixture of mordenite (14g), cumene (10mL, 8.62g, 1.1 eq.) and propylene tetramer (10.84g, 13.2mL, 1eq.) was heated at 140°C until the reaction was finished (olefin residual concentration monitored by GC). The reaction mixture was then cooled down to room temperature. Mordenite extrudates were then separated by filtration and washed with pentane (50mL). The organic phase was then concentrated under vacuum to afford 14.8 g (79% yield) of a large amount of *p*-monoalkyl cumene isomers as confirmed by GC/MS.

EXAMPLE 2

[0096] Preparation of alkylcumene hydroperoxide.

[0097] Into a three-neck round bottom flask was added 15.4 g of the *p*-monoalkyl cumene isomers of Example 1 followed by 0.33 g of cumene hydroperoxide. After 5 min of mixing, 0.55 g of cobaltosic oxide was added to the mixture with vigorous stirring. The flask was sealed and oxygen was introduced through a sparger at 200 sccm rate. The flask was then placed into an oil bath. The mixture was heated to 98°C in 20 min and kept at this temperature for 6 hrs. During the reaction, small kinetic samples were withdrawn from the flask periodically to monitor the conversion by FTIR. Upon completion, the flask was cooled down to room temperature under oxygen. The liquid products were separated from the catalyst by filtration and analyzed by FTIR. The product was used for the next step without further purification

EXAMPLE 3

[0098] Acid cleavage of *p*-monoalkyl cumene hydroperoxide isomers to *p*-alkyl phenol products.

[0099] The reaction was carried out in 100 ml round bottom flask under agitation with a magnetic stirrer. The flask had a distillation link with Liebig condenser and a receiver flask attached to it. A catalyst mixture was first prepared by adding 0.5 ml of water to 50 ml of acetone followed by 1 ml of concentrated sulfuric acid. Next, 20 ml of the mixture was transferred into the round bottom flask. In the following step, 6 ml of acetone was added to 6 ml of the product obtained by the procedure of the Example 2. The solution was added slowly to the flask containing the acid catalyst at room temperature. The flask was heated up in a mineral oil bath to 60°C, the temperature which is close to a boiling point of the mixture. The acetone was distilled out from the flask. The vacuum was applied and the temperature was slowly increased to 70°C to assure the completeness of the reaction. 4.9 g of product was recovered. It had a dark red-brown color. The reaction product was analyzed by GC/MS. The mass fragmentation pattern and retention time comparison with known samples determined para-C₁₂ branched alkyl phenol yield of about 18% based on the total product mixture.

[00100] The reaction product was also analyzed by HPLC (Column: Beckman Ultrasphere, Cyano phase, 80A pore size 3 micron, 4.6x75 mm -Part# 237573) using the following conditions: Pump flow = 2.0 ml/min, UV spectrophotometer wavelength = 280 nm, Column Thermostat = 40°C, Injection size = 5.0 µl of 0.10 (+/- 0.02) gm of sample diluted to 10-ml with cyclohexane. Isocratic elution with 90/10 cyclohexane / 2.5 % ethanol in cyclohexane. The ortho isomer elutes between 1 and 1.5 minutes and the para-isomer between 1.75-2.9 minutes. The sample was found to contain 96 % para-alkylphenol and 4.0 % ortho-alkylphenol. The HPLC analysis indicated 96/4 para- to ortho-alkyl phenol ratio.

[00101] It will be understood that various modifications may be made to the embodiments disclosed herein. Therefore the above description should not be construed as limiting, but merely as exemplifications of preferred embodiments. For example, the functions described above and implemented as the best mode for operating the present invention are for illustration purposes only. Other arrangements and methods may be implemented by those skilled in the art without departing from the scope and spirit of this invention. Moreover, those skilled in the art will envision other modifications within the scope and spirit of the claims appended hereto.

WHAT IS CLAIMED IS:

1. A process for preparing an isomeric mixture comprising a major amount of a para-branched mono-alkyl-substituted hydroxyaromatic compound, the process comprising the steps of:

(a) providing an isomeric mixture comprising a major amount of a para-di(alkyl-substituted)aromatic compound; wherein a first alkyl substituent is a C₃ to C₈ alkyl moiety and a second alkyl substituent is a C_{4+n} to C_{8+n} branched alkyl moiety, wherein n is 0 to 42 and further wherein the second alkyl substituent is at least one carbon atom greater than the first alkyl substituent;

(b) subjecting the isomeric mixture comprising a major amount of a para-di(alkyl-substituted)aromatic compound to oxidation conditions in the presence of an oxygen-containing source, thereby converting the first alkyl substituent which is a C₃ to C₈ alkyl moiety to a hydroperoxide-containing substituted moiety to produce a mixture comprising a major amount of a para-branched alkyl-substituted, hydroperoxide-containing substituted aromatic compound; and

(c) converting the hydroperoxide-containing substituted moiety to a hydroxyl moiety thereby providing a mixture comprising a major amount of a para-branched mono-alkyl-substituted hydroxyaromatic compound.

2. The process according to Claim 1, wherein the second alkyl substituent is derived from a C₄ to C₅₀ branched alpha olefin.

3. The process according to Claims 1 or 2, wherein the mixture of step (a) is obtained from the reaction of a monoalkylaromatic compound and one or more branched alkylating agents in the presence of an acidic catalyst.

4. The process according to Claim 3, wherein the acidic catalyst is an acidic zeolite catalyst and the molar ratio of the monoalkylaromatic compound to the one or more branched alkylating agents is in the range of about 0.5:2 to about 2:0.5.

5. The process according to Claims 1-4, wherein the oxygen-containing source is air.

6. The process according to Claims 1-5, wherein step (b) is carried out in the presence of a free radical initiator.

7. The process according to Claim 6, wherein the free radical initiator is a peroxy compound or azo compound.

8. The process according to Claims 1-7, wherein the oxidation conditions for step (b) comprise a temperature ranging from about 70°C to about 200°C.

9. The process of according to Claims 1-8, wherein step (c) is carried out in the presence of a catalyst.

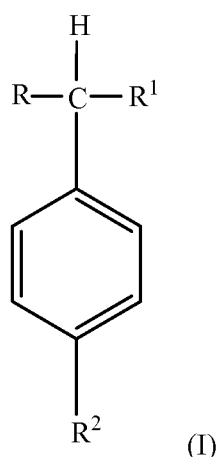
10. The process according to Claim 9, wherein the catalyst is selected from the group consisting of sulfuric acid, perchloric acid, phosphoric acid, hydrochloric acid, p-toluenesulfonic acid, ferric chloride, boron trifluoride, sulfur dioxide and sulfur trioxide.

11. The process according to Claims 1-10, wherein the isomeric mixture contains about 95 mole % or greater of the para-branched mono-alkyl-substituted hydroxyaromatic compounds with the remaining isomeric mixture comprising ortho isomers of the mono-alkyl-substituted hydroxyaromatic compound.

12. The process according to Claims 1-11, wherein the isomeric mixture contains about 99 mole % or greater of the para-branched mono-alkyl-substituted hydroxyaromatic compounds with the remaining isomeric mixture comprising ortho isomers of the mono-alkyl-substituted hydroxyaromatic compound.

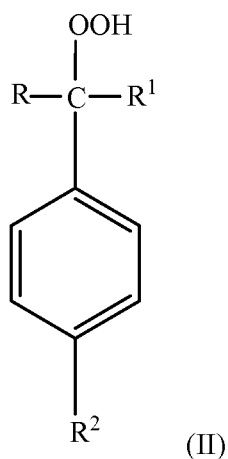
13. A process comprising the steps of:

(a) providing an isomeric mixture comprising a major amount of a para-di(alkyl-substituted)aromatic compound of formula I:



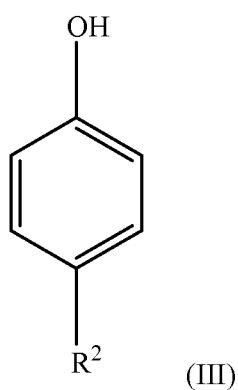
wherein R and R¹ are the same or different alkyl group such that the combined number of carbon atoms is no more than 8 carbon atoms, and R² is a branched alkyl group of from 4 to 50 carbon atoms, wherein R² is at least one carbon atom greater than the combined number of carbon atoms of R and R¹;

(b) subjecting the isomeric mixture comprising a major amount of the para-di(alkyl-substituted)aromatic compound of formula I to oxidation conditions in the presence of an oxygen-containing source to produce an isomeric mixture comprising a major amount of a para-branched alkyl-substituted, hydroperoxide-containing substituted aromatic compound of formula II:



wherein R, R¹ and R² have the aforesated meanings; and

(c) converting the hydroperoxide-containing substituted moiety of the para-branched alkyl-substituted, hydroperoxide-containing substituted aromatic compound of formula II to a hydroxyl moiety thereby providing an isomeric mixture comprising a major amount of a para-branched mono-alkyl-substituted hydroxyaromatic compound of formula III:



wherein R^2 has the aforestated meaning.

14. An isomeric mixture comprising a major amount of a para-branched mono-alkyl-substituted hydroxyaromatic compound prepared by the process according to Claims 1-12.

15. A lubricating oil composition comprising (a) a major amount of an oil of lubricating viscosity and (b) the isomeric mixture of Claim 14.