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(54) **Fuel composition containing an aliphatic amine and a poly (oxyalkylene) monool**

Ein aliphatisches Amin und eine Polyoxyalkylenmonool enthaltende Brennstoffzusammensetzung

Composition de combustible contenant une amine aliphatique et un polyoxyalkylenemonool

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US-A- 5 522 906

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Description

BACKGROUND OF THE INVENTION

[0001] This invention relates to a novel fuel additive composition. More particularly, this invention relates to a fuel composition containing an aliphatic amine and a poly(oxyalkylene) monool.

[0002] It is well known that automobile engines tend to form deposits on the surface of engine components, such as carburetor ports, throttle bodies, fuel injectors, intake ports and intake valves, due to the oxidation and polymerization of hydrocarbon fuel. These deposits, even when present in relatively minor amounts, often cause noticeable driveability problems, such as stalling and poor acceleration. Moreover, engine deposits can significantly increase an automobile's fuel consumption and production of exhaust pollutants. Therefore, the development of effective fuel detergents or "deposit control" additives to prevent or control such deposits is of considerable importance and numerous such materials are known in the art.

[0003] For example, U.S. Patent No. 3,438,757 to Honnen et al. discloses branched chain aliphatic hydrocarbon N-substituted amines and alkylene polyamines having a molecular weight in the range of about 425 to 10,000, preferably about 450 to 5,000, which are useful as detergents and dispersants in hydrocarbon liquid fuels for internal combustion engines.

[0004] U.S. Patent No. 3,756,793 to Robinson discloses a fuel composition containing minor amounts of (a) a polyamine which is the reaction product of a halohydrocarbon having an average molecular weight between 600 to 2500 and an alkylene polyamine, and (b) an organic substance having a viscosity between 20 and 2500 cs. at 200C. This patent further discloses that a wide variety of compounds are suitable as the organic substance, including polyamines, amides, and esters or mixtures of esters, such as aliphatic diesters of dibasic aliphatic carboxylic acids. Preferred materials for use as the organic substance are described in this patent as polymers or copolymers having an average molecular weight of 300 to 5,000 which are selected from hydrocarbons, substituted hydrocarbons containing oxygen and substituted hydrocarbons containing oxygen and nitrogen. Most preferred polymeric compounds are described in this patent as polyalkylene oxides and polyether glycols.

[0005] U.S. Patent No. 5,004,478 to Vogel et al. discloses a motor fuel for internal combustion engines which contains an additive comprising (a) an amino- or amino-containing detergent and (b) a base oil which is a mixture of (1) a polyether based on propylene oxide or butylene oxide and having a molecular weight not less than 500, and (2) an ester of a monocarboxylic or polycarboxylic acid and an alkanol or polyol.

[0006] U.S. Patent No. 5,089,028 to Abramo et al. discloses a fuel composition containing an additive which comprises the combination of (1) a polyalkenyl succinimide, (2) a polyalkylene polymer, such as polyisobutylene or polypropylene, (3) an ester of an aliphatic or aromatic carboxylic acid, and (4) a polyether, such as polybutylene oxide, polypropylene or a polybutylene/polypropylene copolymer. The additive may also contain an optional amount of a mineral oil or a synthetic oil.

[0007] U.S. Patent No. 5,242,469 to Sakakibara et al. discloses a gasoline additive composition comprising (a) a monoester, diester or polyolester, and (b) a dispersant selected from (1) a monosuccinimide, (2) a bis-succinimide, (3) an alkylamine having a polyolefin polymer as an alkyl group and an average molecular weight of 500-5,000, and (4) a benzylamine derivative having an average molecular weight of 500-5,000. The additive composition may additionally contain a polyoxyalkylene glycol or its derivative and/or a lubricant oil fraction.

[0008] U.S. Patent No. 4,877,416 to Campbell discloses a fuel composition which contains (a) from about 0.001 to 1.0 percent by weight of a hydrocarbyl-substituted amine or polyamine having an average molecular weight of about 750 to 10,000 and at least one basic nitrogen atom, and (b) a hydrocarbyl-terminated poly(oxyalkylene) monool having an average molecular weight of about 500 to 5,000, wherein the weight percent of the hydrocarbyl-terminated poly (oxyalkylene) monool in the fuel composition ranges from about 0.01 to 100 times the amount of hydrocarbyl-substituted amine or polyamine.

[0009] U.S. Patent No. 5,006,130 to Aiello et al. discloses an unleaded gasoline composition containing a mixture of (a) about 2.5 parts per million by weight or higher of basic nitrogen in the form of an oil-soluble aliphatic alkylene polyamine containing at least one olefinic polymer chain, said polyamine having a molecular weight of about 600 to 10,000, and (b) from about 75 to about 125 parts per million by weight based on the fuel composition of certain oil-soluble olefinic polymers, a poly(oxyalkylene) alcohol, glycol or polyol or a mono or di-ether thereof, non-aromatic naphthenic or paraffinic oils or polyalphaolefins. This patent further teaches that, as a matter of practicality, the basic nitrogen content of the aliphatic polyamine component is usually about 4.0 or below and that this generally corresponds to a concentration of about 100 to 160 ppm when the aliphatic polyamine is a 1050 molecular weight aliphatic diamine, such as -polyisobutenyl N'-N'-dimethy-1,3-diaminopropane.

[0010] U.S. Patent No. 5,405,419 to Ansari et al. discloses a fuel additive composition comprising (a) a fuel-soluble aliphatic hydrocarbyl-substituted amine having at least one basic nitrogen atom wherein the hydrocarbyl group has a number average molecular weight of about 700 to 3,000; (b) a polyolefin polymer of a C_2 to C_6 monolefin, wherein the

polymer has a number average molecular weight of about 350 to 3,000; and (c) a hydrocarby-terminated poly(oxyalkylene) monool having an average molecular weight of about 500 to 5,000. This patent further teaches that fuel compositions containing these additives will generally contain about 50 to 500 ppm by weight of the aliphatic amine, about 50 to 1,000 ppm by weight of the polyolefin and about 50 to 1,000 ppm by weight of the poly(oxyalkylene) monool. This patent also discloses that fuel compositions containing 125 ppm each of aliphatic amine, polyolefin and poly (oxyalkylene) monool provide better deposit control performance than compositions containing 125 ppm of aliphatic amine plus 125 ppm of poly(oxyalkylene) monool. U.S. Patent No. 5,522,906 discloses gasoline compositions comprising (a) gasoline, (b) a deposit inhibitor or a detergent containing a basic nitrogen atom, (c) a carrier oil, and (d) one or more compounds selected from (d1) an ester of a fatty acid and an alylene oxide addition compound to bisphenol, (d2) a compound obtained by the ester exchange reaction of an alcohol and a triglyceride-type fat or oil, (d3) an aliphatic or aromatic carboxylic acid, (d4) a metal salt of an aliphatic or aromatic carboxylic acid, (d5) an ester of an aliphatic or aromatic carboxylic acid and an alcohol, and (d6) an ester of boric acid. The compositions are able to remove deposits in fuel intake systems and to keep systems clean.

SUMMARY OF THE INVENTION

[0011] It has now been discovered that the unique combination of an aliphatic hydrocarbyl-substituted amine and a poly(oxyalkylene) monool, when employed at very low concentrations as a fuel additive composition for hydrocarbon fuels, provides excellent control of engine deposits, especially engine intake valve deposits.

[0012] According to the present invention, there is provide a fuel composition comprising a major amount of hydrocarbons boiling in the gasoline or diesel range and from 85 to below 145 parts per million by weight of a deposit control additive composition comprising: (a) 50 to 70 parts per million by weight of a fuel-soluble aliphatic hydrocarbyl-substituted amine of the formula:

$$R_3NH\text{-}(R_4\text{-}NH)_n\text{-}H$$

wherein R_3 is a hydrocarbyl group derived from polyisobutylene having a number average molecular weight of 900 to 1,500; R_4 is an alkylene group of from 2 to 6 carbon atoms; and n is an integer of from 0 to 10; and (b) 35 to below 75 parts per million by weight of a hydrocarbyl-terminated poly(oxyalkylene) monool having an average molecular weight of 500 to 5,000, wherein the oxyalkylene group is a C_2 to C_5 oxyalkylene group and the hydrocarbyl group is a C_1 to C_{30} hydrocarbyl group; wherein the ratio of the poly(oxyalkylene) monool to the aliphatic amine in the range of 0:5:1 to 1.5:1. ***[deletion(s)]***

[0013] Preferably, the fuel composition of the present invention will contain about 40 to 70 parts per million by weight of the hydrocarbyl-terminated poly(oxyalkylene) monool.

[0014] The present invention further provides a method for reducing engine deposits in an internal combustion engine which comprises operating the engine with the novel fuel composition of the present invention.

[0015] Among other factors, the present invention is based on the surprising discovery that the unique combination of an aliphatic amine and a poly(oxyalkylene) monool unexpectedly provides excellent deposit control performance at low concentrations not heretofore recognized in the art as effective for optimum deposit control.

DETAILED DESCRIPTION OF THE INVENTION

[0016] As noted above, the fuel composition of the present invention contains an aliphatic hydrocarbyl-substituted amine and a hydrocarbyl-terminated poly(oxyalkylene) monool. These compounds are described in further detail below.

A. The Aliphatic Hydrocarbyl-Substituted Amine

[0017] The fuel-soluble aliphatic hydrocarbyl-substituted amine component of the present fuel composition is as defined in claim 1. Typically, such aliphatic amines will be of sufficient molecular weight so as to be nonvolatile at normal engine intake valve operating temperatures, which are generally in the range of about 175°C to 300°C.

[0018] In most instances, the branched-chain hydrocarbyl amines are not a pure single product, but rather a mixture of compounds having an average molecular weight. Usually, the range of molecular weights will be relatively narrow and peaked near the indicated molecular weight.

[0019] The amine component of the branched-chain hydrocarbyl amines may be derived from ammonia, a monoamine or a polyamine. The monoamine or polyamine component embodies a broad class of amines having from 1 to about 12 amine nitrogen atoms and from 1 to about 40 carbon atoms with a carbon to nitrogen ratio between about 1:1 and 10:1. Generally, the monoamine will contain from 1 to about 40 carbon atoms and the polyamine will contain

from 2 to about 12 amine nitrogen atoms and from 2 to about 40 carbon atoms. In most instances, the amine component is not a pure single product, but rather a mixture of compounds having a major quantity of the designated amine. For the more complicated polyamines, the compositions will be a mixture of amines having as the major product the compound indicated and having minor amounts of analogous compounds. Suitable monoamines and polyamines are described more fully below.

[0020] When the amine component is a polyamine, it will preferably be a polyalkylene polyamine, including alkylenediamine. Preferably, the alkylene group will contain from 2 to 6 carbon atoms, more preferably from 2 to 3 carbon atoms. Examples of such polyamines include ethylene diamine, diethylene triamine, triethylene tetramine and tetraethylene pentamine. Preferred polyamines are ethylene diamine and diethylene triamine.

[0021] Particularly preferred branched-chain hydrocarbyl amines include polyisobutenyl ethylene diamine and polyisobutyl amine, wherein the polyisobutyl group is substantially saturated and the amine moiety is derived from ammonia.

[0022] The aliphatic hydrocarbyl amines employed in the fuel composition of the invention are prepared by conventional procedures known in the art. Such aliphatic hydrocarbyl amines and their preparations are described in detail in U.S. Patent Nos. 3,438,757; 3,565,804; 3,574,576; 3,848,056; 3,960,515; and 4,832,702.

[0023] Typically, the hydrocarbyl-substituted amines employed in this invention are prepared by reacting a hydrocarbyl halide, such as a hydrocarbyl chloride, with ammonia or a primary or secondary amine to produce the hydrocarby-substituted amine.

[0024] As noted above, the amine component of the presently employed hydrocarbyl-substituted amine is derived from a nitrogen-containing compound selected from ammonia, a monoamine having from 1 to about 40 carbon atoms, and a polyamine having from 2 to about 12 amine nitrogen atoms and from 2 to about 40 carbon atoms. The nitrogen-containing compound is reacted with a hydrocarbyl halide to produce the hydrocarbyl-substituted amine fuel additive finding use within the scope of the present invention. The amine component provides a hydrocarbyl amine reaction product with, on average, at least about one basic nitrogen atom per product molecule, i.e., a nitrogen atom titratable by a strong acid.

[0025] Preferably, the amine component is derived from a polyamine having from 2 to about 12 amine nitrogen atoms and from 2 to about 40 carbon atoms. The polyamine preferably has a carbon-to-nitrogen ratio of from about 1:1 to 10:1.

[0026] The polyamine may be substituted with substituents selected from (a) hydrogen, (b) hydrocarbyl groups of from 1 to about 10 carbon atoms, (c) acyl groups of from 2 to about 10 carbon atoms, and (d) monoketo, monohydroxy, mononitro, monocyano, lower alkyl and lower alkoxy derivatives of (b) and (c). "Lower", as used in terms like lower alkyl or lower alkoxy, means a group containing from 1 to about 6 carbon atoms. At least one of the substituents on one of the basic nitrogen atoms of the polyamine is hydrogen, e.g., at least one of the basic nitrogen atoms of the polyamine is a primary or secondary amino nitrogen.

[0027] Hydrocarbyl, as used in describing the polyamine moiety on the aliphatic amine employed in this invention, denotes an organic radical composed of carbon and hydrogen which may be aliphatic, alicyclic, aromatic or combinations thereof, e.g., aralkyl. Preferably, the hydrocarbyl group will be relatively free of aliphatic unsaturation, i.e., ethylenic and acetylenic, particularly acetylenic unsaturation. The substituted polyamines of the present invention are generally, but not necessarily, N-substituted polyamines. Exemplary hydrocarbyl groups and substituted hydrocarbyl groups include alkyls such as methyl, ethyl, propyl, butyl, isobutyl, pentyl, hexyl, octyl, etc., alkenyls such as propenyl, isobutenyl, hexenyl, octenyl, etc., hydroxyalkyls, such as 2-hydroxyethyl, 3-hydroxypropyl, hydroxy-isopropyl, 4-hydroxybutyl, etc., ketoalkyls, such as 2-ketopropyl, 6-ketooctyl, etc., alkoxy and lower alkenoxy alkyls, such as ethoxyethyl, ethoxypropyl, propoxyethyl, propoxypropyl, diethyleneoxymethyl, triethyleneoxyethyl, tetraethyleneoxyethyl, diethyleneoxyhexyl, etc. The aforementioned acyl groups (c) are such as propionyl, acetyl, etc. The more preferred substituents are hydrogen, C_1-C_6 alkyls and C_1-C_6 hydroxyalkyls.

[0028] In a substituted polyamine, the substituents are found at any atom capable of receiving them. The substituted atoms, e.g., substituted nitrogen atoms, are generally geometrically unequivalent, and consequently the substituted amines finding use in the present invention can be mixtures of mono- and poly-substituted polyamines with substituent groups situated at equivalent and/or unequivalent atoms.

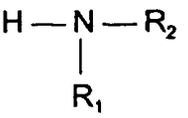
[0029] The more preferred polyamine finding use within the scope of the present invention is a polyalkylene polyamine, including alkylene diamine, and including substituted polyamines, e.g., alkyl and hydroxyalkyl-substituted polyalkylene polyamine. Preferably, the alkylene group contains from 2 to 6 carbon atoms, there being preferably from 2 to 3 carbon atoms between the nitrogen atoms. Such groups are exemplified by ethylene, 1,2-propylene, 2,2-dimethylpropylene, trimethylene, 1,3,2-hydroxypropylene, etc. Examples of such polyamines include ethylene diamine, diethylene triamine, di(trimethylene) triamine, dipropylene triamine, triethylene tetraamine, tripropylene tetraamine, tetraethylene pentamine, and pentaethylene hexamine. Such amines encompass isomers such as branched-chain polyamines and previously-mentioned substituted polyamines, including hydroxy- and hydrocarbyl-substituted polyamines. Among the polyalkylene polyamines, those containing 2-12 amino nitrogen atoms and 2-24 carbon atoms are especially preferred, and the C_2-C_3 alkylene polyamines are most preferred, that is, ethylene diamine, polyethylene

polyamine, propylene diamine and polypropylene polyamine, and in particular, the lower polyalkylene polyamines, e. g., ethylene diamine, dipropylene triamine, etc. Particularly preferred polyalkylene polyamines are ethylene diamine and diethylene triamine.

[0030] The amine component of the presently employed aliphatic amine fuel additive also may be derived from heterocyclic polyamines, heterocyclic substituted amines and substituted heterocyclic compounds, wherein the heterocycle comprises one or more 5-6 membered rings containing oxygen and/or nitrogen. Such heterocyclic rings may be saturated or unsaturated and substituted with groups selected from the aforementioned (a), (b), (c) and (d). The heterocyclic compounds are exemplified by piperazines, such as 2-methylpiperazine, N-(2-hydroxyethyl)-piperazine, 1,2-bis-(N-piperazinyl)ethane and N,N'-bis(N-piperazinyl)piperazine, 2-methylimidazoline, 3-aminopiperidine, 3-aminopyridine, N-(3-aminopropyl)-morpholine, etc. Among the heterocyclic compounds, the piperazines are preferred.

[0031] Typical polyamines that can be used to form the aliphatic amine additives employed in this invention by reaction with a hydrocarbyl halide include the following: ethylene diamine, 1,2-propylene diamine, 1,3-propylene diamine, diethylene triamine, triethylene tetramine, hexamethylene diamine, tetraethylene pentamine, dimethylaminopropylene diamine, N-(beta-aminoethyl)piperazine, N-(beta-aminoethyl)piperidine, 3-amino-N-ethylpiperidine, N-(beta-aminoethyl) morpholine, N,N'-di(beta-aminoethyl)piperazine, N,N'-di(beta-aminoethyl)imidazolidone-2, N-(beta-cyanoethyl) ethane-1,2-diamine, 1-amino-3,6,9-triazaoctadecane, 1-amino-3,6-diaza-9-oxadecane, N-(beta-aminoethyl) diethanolamine, N'-acetylmethyl-N-(beta-aminoethyl) ethane-1,2-diamine, N-acetonyl-1,2-propanediamine, N-(beta-nitroethyl)-1,3-propane diamine, 1,3-dimethyl-5(beta-aminoethyl)hexahydrotriazine, N-(beta-aminoethyl)-hexahydrotriazine, 5-(beta-aminoethyl)-1,3,5-dioxazine, 2-(2-aminoethylamino)ethanol, and 2-[2-(2-aminoethylamino) ethylamino] ethanol.

[0032] Alternatively, the amine component of the presently employed aliphatic hydrocarbyl-substituted amine may be derived from an amine having the formula:



wherein R_1 and R_2 are independently selected from the group consisting of hydrogen and hydrocarbyl of 1 to about 20 carbon atoms and, when taken together, R_1 and R_2 may form one or more 5- or 6-membered rings containing up to about 20 carbon atoms. Preferably, R_1 is hydrogen and R_2 is a hydrocarbyl group having 1 to about 10 carbon atoms. More preferably, R_1 and R_2 are hydrogen. The hydrocarbyl groups may be straight-chain or branched and may be aliphatic, alicyclic, aromatic or combinations thereof. The hydrocarbyl groups may also contain one or more oxygen atoms.

[0033] An amine of the above formula is defined as a "secondary amine" when both R_1 and R_2 are hydrocarbyl. When R_1 is hydrogen and R_2 is hydrocarbyl, the amine is defined as a "primary amine"; and when both R_1 and R_2 are hydrogen, the amine is ammonia.

[0034] Primary amines useful in preparing the aliphatic hydrocarbyl-substituted amine fuel additives of the present invention contain 1 nitrogen atom and 1 to about 20 carbon atoms, preferably 1 to 10 carbon atoms. The primary amine may also contain one or more oxygen atoms.

[0035] Preferably, the hydrocarbyl group of the primary amine is methyl, ethyl, propyl, butyl, pentyl, hexyl, octyl, 2-hydroxyethyl or 2-methoxyethyl. More preferably, the hydrocarbyl group is methyl, ethyl or propyl.

[0036] Typical primary amines are exemplified by N-methylamine, N-ethylamine, N-n-propylamine, N-isopropylamine, N-n-butylamine, N-isobutylamine, N-sec-butylamine, N-tert-butylamine, N-n-pentylamine, N-cyclopentylamine, N-n-hexylamine, N-cyclohexylamine, N-octylamine, N-decylamine, N-dodecylamine, N-octadecylamine, N-benzylamine, N-(2-phenylethyl)amine, 2-aminoethanol, 3-amino-1-proponal, 2-(2-aminoethoxy)ethanol, N-(2-methoxyethyl)amine, N-(2-ethoxyethyl)amine, and the like. Preferred primary amines are N-methylamine, N-ethylamine and N-n-propylamine.

[0037] The amine component of the presently employed aliphatic hydrocarbyl-substituted amine fuel additive may also be derived from a secondary amine. The hydrocarbyl groups of the secondary amine may be the same or different and will generally contain 1 to about 20 carbon atoms, preferably 1 to about 10 carbon atoms. One or both of the hydrocarbyl groups may also contain one or more oxygen atoms.

[0038] Preferably, the hydrocarbyl groups of the secondary amine are independently selected from the group consisting of methyl, ethyl, propyl, butyl, pentyl, hexyl, 2-hydroxyethyl and 2-methoxyethyl. More preferably, the hydrocarbyl groups are methyl, ethyl or propyl.

[0039] Typical secondary amines which may be used in this invention include N,N-dimethylamine, N,N-diethylamine, N,N-di-n-propylamine, N,N-diisopropylamine, N,N-di-n-butylamine, N,N-di-sec-butylamine, N,N-di-n-pentylamine, N,N-di-n-hexylamine, N,N-dicyclohexylamine, N,N-dioctylamine, N-ethyl-N-methylamine, N-methyl-N-n-propylamine, N-n-butyl-N-methylamine, N-methyl-N-octylamine, N-ethyl-N-isopropylamine, N-ethyl-N-octylamine, N,N-di(2-hydroxyethyl)amine, N,N-di(3-hydroxypropyl)amine, N,N-di(ethoxyethyl)amine, N,N-di(propoxyethyl)amine, and the like. Preferred secondary amines are N,N-dimethylamine, N,N-diethylamine and N,N-di-n-propylamine.

[0040] Cyclic secondary amines may also be employed to form the aliphatic amine additives of this invention. In such cyclic compounds, R_1 and R_2 of the formula hereinabove, when taken together, form one or more 5- or 6-membered rings containing up to about 20 carbon atoms. The ring containing the amine nitrogen atom is generally saturated, but may be fused to one or more saturated or unsaturated rings. The rings may be substituted with hydrocarbyl groups of from 1 to about 10 carbon atoms and may contain one or more oxygen atoms.

[0041] Suitable cyclic secondary amines include piperidine, 4-methylpiperidine, pyrrolidine, morpholine, 2,6-dimethylmorpholine, and the like.

[0042] In many instances, the amine component is not a single compound but a mixture in which one or several compounds predominate with the average composition indicated. For example, tetraethylene pentamine prepared by the polymerization of aziridine or the reaction of dichloroethylene and ammonia will have both lower and higher amine members, e.g., triethylene tetraamine, substituted piperazines and pentaethylene hexamine, but the composition will be mainly tetraethylene pentamine and the empirical formula of the total amine composition will closely approximate that of tetraethylene pentamine. Finally, in preparing the compounds of this invention using a polyamine, where the various nitrogen atoms of the polyamine are not geometrically equivalent, several substitutional isomers are possible and are encompassed within the final product. Methods of preparation of amines and their reactions are detailed in Sidgewick's "The Organic Chemistry of Nitrogen", Clarendon Press, Oxford, 1966; Noller's "Chemistry of Organic Compounds", Saunders, Philadelphia, 2nd Ed., 1957; and Kirk-Othmer's "Encyclopedia of Chemical Technology", 2nd Ed., especially Volume 2, pp. 99-116.

[0043] Aliphatic hydrocarbyl-substituted amines suitable for use in the present invention are hydrocarbyl-substituted polyalkylene polyamines having the formula:

$$R_3NH\text{—}(R_4\text{—}NH)_n\text{—}H$$

wherein R_3 is a hydrocarbyl group derived from polyisobutylene having a number average molecular weight of 900 to 1,500; R_4 is alkylene of from 2 to 6 carbon atoms; and n is an integer of from 0 to about 10.

[0044] Preferably,
R_4 is alkylene of from 2 to 3 carbon atoms and n is preferably an integer of from 1 to 6.

B. The Hydrocarbyl-Terminated Poly(oxyalkylene) Monool

[0045] The hydrocarbyl-terminated poly(oxyalkylene) polymers employed in the present invention are monohydroxy compounds, i.e., alcohols, often termed monohydroxy polyethers, or polyalkylene glycol monohydrocarbylethers, or "capped" poly(oxyalkylene) glycols and are to be distinguished from the poly(oxyalkylene) glycols (diols), or polyols, which are not hydrocarbyl-terminated, i.e., not capped. The hydrocarbyl-terminated poly(oxyalkylene) alcohols are produced by the addition of lower alkylene oxides, such as ethylene oxide, propylene oxide, the butylene oxides, or the pentylene oxides to the hydroxy compound R_5OH under polymerization conditions, wherein R_5 is the hydrocarbyl group which caps the poly(oxyalkylene) chain. Methods of production and properties of these polymers are disclosed in U.S. Patent Nos. 2,841,479 and 2,782,240 and Kirk-Othmer's "Encyclopedia of Chemical Technology", 2nd Ed., Volume 19, p. 507. In the polymerization reaction, a single type of alkylene oxide may be employed, e.g., propylene oxide, in which case the product is a homopolymer, e.g., a poly(oxyalkylene) propanol. However, copolymers are equally satisfactory and random copolymers are readily prepared by contacting the hydroxyl-containing compound with a mixture of alkylene oxides, such as a mixture of propylene and butylene oxides. Block copolymers of oxyalkylene units also provide satisfactory poly(oxyalkylene) polymers for the practice of the present invention. Random polymers are more easily prepared when the reactivities of the oxides are relatively equal. In certain cases, when ethylene oxide is copolymerized with other oxides, the higher reaction rate of ethylene oxide makes the preparation of random copolymers difficult. In either case, block copolymers can be prepared. Block copolymers are prepared by contacting the hydroxyl-containing compound with first one alkylene oxide, then the others in any order, or repetitively, under polymerization conditions. A particular block copolymer is represented by a polymer prepared by polymerizing propylene oxide on a suitable monohydroxy compound to form a poly(oxypropylene) alcohol and then polymerizing butylene oxide on the poly(oxyalkylene) alcohol.

[0046] In general, the poly(oxyalkylene) polymers are mixtures of compounds that differ in polymer chain length.

However, their properties closely approximate those of the polymer represented by the average composition and molecular weight.

[0047] The polyethers employed in this invention can be represented by the formula:

$$R^5O—(R_6O)_p—H$$

wherein R_5 is a hydrocarbyl group of from 1 to 30 carbon atoms; R_6 is a C_2 to C_5 alkylene group; and p is an integer such that the molecular weight of the polyether is from about 500 to about 5,000.

[0048] Preferably, R_6 is a C_3 or C_4 alkylene group.

[0049] Preferably, R_5 is a C_7-C_{30} alkylphenyl group. Most preferably, R_5 is dodecylphenyl.

[0050] Preferably, the polyether has a molecular weight of from about 750 to about 3,000; and more preferably from about 900 to about 1,500.

Fuel Compositions

[0051] The fuel additive composition employed in the present invention will generally be employed in a hydrocarbon distillate fuel boiling in the gasoline or diesel range. The proper concentration of this additive composition necessary in order to achieve the desired detergency and dispersancy varies depending upon the type of fuel employed, the presence of other detergents, dispersants and other additives, etc. Generally, however, from about 85 to below about 145 ppm by weight, preferably from about 90 to 140 ppm, of the present additive composition per part of base fuel is needed to achieve the best results.

[0052] In terms of individual components, fuel compositions containing the additive composition employed in the invention will generally contain about 50 to 70 ppm by weight of the aliphatic hydrocarbyl-substituted amine and about 35 to below about 75 ppm, preferably about 40 to 70 ppm, by weight of the poly(oxyalkylene) monool. The ratio of poly (oxyalkylene) monool to aliphatic amine (monool:amine) will generally be in the range of about 0.5:1 to about 1.5:1.

[0053] The deposit control fuel additive composition employed in the invention may be formulated as a concentrate, using an inert stable oleophilic (i.e., dissolves in gasoline or diesel fuel) organic solvent boiling in the range of about 150°F to 400°F (about 65°C to 205°C). Preferably, an aliphatic or an aromatic hydrocarbon solvent is used, such as benzene, toluene, xylene or higher-boiling aromatics or aromatic thinners. Aliphatic alcohols of about 3 to 8 carbon atoms, such as isopropanol, isobutylcarbinol, n-butanol and the like, in combination with hydrocarbon solvents are also suitable for use with the detergent-dispersant additive. In the concentrate, the amount of the presently employed additive composition will be ordinarily at least 10% by weight and generally not exceed 90% by weight, preferably 40 to 85 weight percent and most preferably from 50 to 80 weight percent.

[0054] In gasoline fuels, other fuel additives may be employed with the additives used in the present invention, including, for example, oxygenates, such as t-butyl methyl ether, antiknock agents, such as methylcyclopentadienyl manganese tricarbonyl, and other dispersants/detergents, such as various hydrocarbyl amines, hydrocarbyl poly(oxyalkylene) amines, or succinimides. Also included may be lead scavengers, such as aryl halides, e.g., dichlorobenzene, or alkyl halides, e.g., ethylene dibromide. Additionally, antioxidants, metal deactivators, pour point depressants, corrosion inhibitors, demulsifiers and anti-valve seat recession agents may be present. The gasoline fuels may also contain amounts of other fuels such as, for example, methanol.

[0055] Additional fuel additives which may be present include fuel injector inhibitors, low molecular weight fuel injector detergents, and carburetor detergents, such as a low molecular weight hydrocarbyl amine, including polyamines, having a molecular weight below 700, such as oleyl amine or a low molecular weight polyisobutenyl ethylene diamine, for example, where the polyisobutenyl group has a number average molecular weight of about 420.

[0056] In diesel fuels, other well-known additives can be employed, such as pour point depressants, flow improverse, cetane improvers, and the like. The diesel fuels can also include other fuels such as, for example, methanol.

[0057] A fuel-soluble, nonvolatile carrier fluid or oil may also be used with the fuel additive composition employed in this invention. The carrier fluid is a chemically inert hydrocarbon-soluble liquid vehicle which substantially increases the nonvolatile residue (NVR), or solvent-free liquid fraction of the fuel additive composition while not overwhelmingly contributing to octane requirement increase. The carrier fluid may be a natural or synthetic oil, such as mineral oil or refined petroleum oils.

[0058] These carrier fluids are believed to act as a carrier for the fuel additives of the present invention and to assist in removing and retarding deposits. The carrier fluid may also exhibit synergistic deposit control properties when used in combination with a fuel additive composition employed in this invention.

[0059] The carrier fluids are typically employed in amounts ranging from about 25 to about 5,000 ppm by weight of the hydrocarbon fuel, preferably from about 35 to 800 ppm of the fuel. Preferably, the ratio of carrier fluid to deposit control additive will range from about 0.2:1 to about 10:1, more preferably from about 0.5:1 to about 3:1.

[0060] When employed in a fuel concentrate, carrier fluids will generally be present in amounts ranging from about 10 to about 60 weight percent, preferably from 20 to 40 weight percent.

[0061] The following examples are presented to illustrate specific embodiments of this invention and are not to be construed in any way as limiting the scope of the invention.

EXAMPLES

Example 1

BMW Mileage Accumulator Test

[0062] The fuel composition of the present invention was tested in a BMW vehicle to evaluate its intake valve deposit control performance. The BMW was equipped with a 1.8 liter, port fuel injected, 4-cylinder overhead cam engine manufactured by BMW. Vehicle preparations were conducted in accordance with ASTM D 5500, unless specified otherwise.

[0063] Mileage accumulation was conducted on a chassis dynamometer using the driving cycle specified in Table I.

Table I

Test Method Driving Cycle			
Mode	Total Distance %	Average Speed mph	Distance miles
AMA City Portion	10	36	80
Suburban	20	40	166
Highway	70	65	560
Total	100	45	800

[0064] At the end of each test run, the intake valves were removed, washed with hexane, and weighed. The previously determined weights of the clean valves were subtracted from the weights of the valves at the end of the run. The difference between the two weights is the weight of the intake valve deposit (IVD).

[0065] The base fuel employed was a regular octane unleaded gasoline containing no deposit control additive. The test compounds were admixed with the base fuel to give the concentrations indicated in the tables.

[0066] The test was run for 5,000 miles on the test fuel. The amount of carbonaceous deposit in milligrams on the intake valves is reported for each of the test samples in Table II.

Table II

BMW 5.000 Mile Results				
Sample	Amine[a], ppma	Monool[b], ppm	Ratio (Monool/Amine)	Intake Valve Deposit Weight, mg
Base Fuel				281
No. 1[c]	80	115	1.4	44
No. 2[c]	30	115	3.8	254
No. 3[c]	80	30	0.38	117
No. 4[c]	30	30	1	345
No. 5	68.5	57	0.83	57
No. 6	60	40	0.67	54

[a] Polyisobutenyl (1300 MW) ethylene diamine, in parts per million actives (ppma).

[b] Dodecylphenyl poly (oxypropylene) monool having an average molecular weight of about 1000, in parts per million (ppm).

[c] Comparative samples.

[0067] The data in Table II demonstrates that the combination of aliphatic amine and poly(oxyalkylene) monool additives at very low concentrations in fuels (Sample Nos. 5 and 6) provides excellent deposit control performance which is significantly better than would be predicted from a linear combination of these additives.

[0068] Additional testing was obtained in a 10,000 mile BMW test as specified in ASTM D 5500. An average deposit weight of 100 milligrams per valve or less at the conclusion of the 10,000 mile test meets BMW requirements for unlimited mileage acceptance. The results are set forth in Table III.

Table III

BMW 10.000 Mile Results				
Sample	Amine[a], ppma	Monool[b], ppm	Ratio (Monool/Amine)	Intake Valve Deposit Weight, mg
Base Fuel A				158
Base Fuel A				281
Base Fuel A				238
Base Fuel B				360
Base Fuel B				300
No. 1[c]	53	42.5	0.8	75
No. 2[d]	53	42.5	0.8	86

[a] Polyisobutenyl (1300 MW) ethylene diamine, in parts per million actives (ppma).

[b] Dodecylphenyl poly (oxypropylene) monool having an average molecular weight of about 1000, in parts per million (ppm).

[c] Formulated in Base Fuel A.

[d] Formulated in Base Fuel B.

[0069] The data in Table III illustrates the significant reduction in intake valve deposits provided by the fuel composition of the present invention (Sample Nos. 1 and 2) compared to the base fuel and shows that the instant fuel composition passes the BMW unlimited mileage test at a very low concentration of additives.

Claims

1. A fuel composition comprising a major amount of hydrocarbons boiling in the gasoline or diesel range and from 85 to below 145 parts per million by weight of a deposit control additive composition comprising:

(a) 50 to 70 parts per million by weight of a fuel-soluble aliphatic hydrocarbyl-substituted amine of the formula:

$$R_3NH\text{-}(R_4\text{-}NH)_n\text{-}H$$

wherein R_3 is a hydrocarbyl group derived from polyisobutylene having a number average molecular weight of 900 to 1,500; R_4 is an alkylene group of from 2 to 6 carbon atoms; and n is an integer of from 0 to 10; and

(b) 35 to below 75 parts per million by weight of a hydrocarbyl-terminated poly(oxyalkylene) monool having an average molecular weight of 500 to 5,000, wherein the oxyalkylene group is a C_2 to C_5 oxyalkylene group and the hydrocarbyl group is a C_1 to C_{30} hydrocarbyl group;

wherein the ratio of the poly (oxyalkylene) monool to the aliphatic amine in the range of 0.5:1 to 1.5:1.

2. The fuel composition according to claim 1, wherein the aliphatic amine of component (a) is a polyisobutyl amine, wherein the polyisobutyl group is substantially saturated and the amine moiety is derived from ammonia.

3. The fuel composition according to Claim 1, wherein the amine moiety of the aliphatic amine is derived from a polyamine having from 2 to 12 amine nitrogen atoms and from 2 to 40 carbon atoms.

4. The fuel composition according to Claim 3, wherein the polyamine is a polyalkylene polyamine having 2 to 12 amine nitrogen atoms and 2 to 24 carbon atoms.

5. The fuel composition according to Claim 4, wherein the polyalkylene polyamine is selected from the group consisting of ethylene diamine, diethylene triamine, triethylene tetramine and tetraethylene pentamine.

6. The fuel composition according to Claim 5, wherein the polyalkylene polyamine is ethylene diamine or diethylene triamine

7. The fuel composition according to Claim 6, wherein the aliphatic amine of component (a) is a polyisobutenyl ethylene diamine.

8. The fuel composition according to Claim 1, wherein the hydrocarbyl-terminated poly(oxyalkylene) monool of component (b) has an average molecular weight of 900 to 1500.

9. The fuel composition according to Claim 1, wherein the oxyalkylene group of the hydrocarbyl-terminated poly (oxyalkylene) monool of component (b) is a C_3 to C_4 oxyalkylene group.

10. The fuel composition according to Claim 9, wherein the oxyalkylene group of the hydrocarbyl-terminated poly (oxyalkylene) monool of component (b) is a C_3 oxypropylene group.

11. The fuel composition according to Claim 9, wherein the oxyalkylene group of the hydrocarbyl-terminated poly (oxyalkylene) monool of component (b) is a C_4 oxybutylene group.

12. The fuel composition according to Claim 1, wherein the hydrocarbyl group of the hydrocarbyl-terminated poly (oxyalkylene) monool of component (b) is a C_7 to C_{30} alkylphenyl group.

13. The fuel composition according to Claim 1, wherein the fuel composition contains 40 to 70 parts per million by weight of the hydrocarbyl-terminated poly(oxyalkylene) monool of component (b).

14. The fuel composition according to Claim 1, wherein component (a) is a polyisobutenyl amine, wherein the amine moiety is derived from ethylene diamine or diethylene triamine, and component (b) is a C_7 to C_{30} alkylphenyl-terminated poly(oxypropylene) or poly(oxybutylene) monool.

15. A method for reducing engine deposits in an internal combustion engine which comprises operating the engine with a fuel composition as claimed in any preceding claim.

Revendications

1. Composition de carburant comprenant une quantité dominante d'hydrocarbures bouillant dans la plage de l'essence ou du carburant diesel et 85 à moins de 145 parties par million en poids d'une composition d'additifs limitant les dépôts comprenant :

   (a) 50 à 70 parties par million en poids d'une amine à substituant hydrocarbyle aliphatique, soluble dans les carburants, de formule :

   $$R_3NH\text{-}(R_4\text{-}NH)_n\text{-}H$$

   dans laquelle R_3 représente un groupe hydrocarbyle dérivé du polyisobutylène, ayant une moyenne en nombre du poids moléculaire de 900 à 1500 ; R_4 représente un groupe alkylène de 2 à 6 atomes de carbone ; et n représente un nombre entier de 0 à 10 ; et
   (b) 35 à moins de 75 parties par million en poids d'un polyoxyalkylène-monool à terminaison hydrocarbyle, ayant un poids moléculaire moyen de 500 à 5000, dans lequel le groupe oxyalkylène est un groupe oxyalkylène en C_2 à C_5 et le groupe hydrocarbyle est un groupe hydrocarbyle en C_1 à C_{30} ;

   dans laquelle le rapport du polyoxyalkylène-monool à l'amine aliphatique est compris dans l'intervalle de 0,5:1 à 1,5:1.

2. Composition de carburant suivant la revendication 1, dans laquelle l'amine aliphatique du constituant (a) est une polyisobutylamine, dans laquelle le groupe polyisobutyle est substantiellement saturé et le groupement amine est dérivé de l'ammoniac.

3. Composition de carburant suivant la revendication 1, dans laquelle le groupement amine de l'amine aliphatique est dérivé d'une polyamine ayant 2 à 12 atomes d'azote d'amine et 2 à 40 atomes de carbone.

4. Composition de carburant suivant la revendication 3, dans laquelle la polyamine est une polyalkylènepolyamine ayant 2 à 12 atomes d'azote d'amine et 2 à 24 atomes de carbone.

5. Composition de carburant suivant la revendication 4, dans laquelle la polyalkylènepolyamine est choisie dans le groupe consistant en éthylènediamine, diéthylènetriamine, triéthylènetétramine et tétraéthylènepentamine.

6. Composition de carburant suivant la revendication 5, dans laquelle la polyalkylènepolyamine est l'éthylènediamine ou la diéthylènetriamine.

7. Composition de carburant suivant la revendication 6, dans laquelle l'amine aliphatique du constituant (a) est la polyisobuténylétheylènediamine.

8. Composition de carburant suivant la revendication 1, dans laquelle le polyoxyalkylène-monool à terminaison hydrocarbyle du constituant (b) a un poids moléculaire moyen de 900 à 1500.

9. Composition de carburant suivant la revendication 1, dans laquelle le groupe oxyalkylène du polyoxyalkylène-monool à terminaison hydrocarbyle du constituant (b) est un groupe oxyalkylène en C_3 ou C_4.

10. Composition de carburant suivant la revendication 9, dans laquelle le groupe oxyalkylène du polyoxyalkylène-monool à terminaison hydrocarbyle du constituant (b) est un groupe oxypropylène en C_3.

11. Composition de carburant suivant la revendication 9, dans laquelle le groupe oxyalkylène du polyoxyalkylène-monool à terminaison hydrocarbyle du constituant (b) est un groupe oxybutylène en C_4.

12. Composition de carburant suivant la revendication 1, dans laquelle le groupe hydrocarbyle du polyoxyalkylène-monool à terminaison hydrocarbyle du constituant (b) est un groupe alkylphényle en C_7 à C_{30}.

13. Composition de carburant suivant la revendication 1, qui contient 40 à 70 parties par million en poids du polyoxyalkylène-monool à terminaison hydrocarbyle du constituant (b).

14. Composition de carburant suivant la revendication 1, dans laquelle le constituant (a) est une polyisobutylamine, dans laquelle le groupement amine est dérivé de l'éthylènediamine ou de la diéthylènetriamine, et le constituant (b) est un polyoxypropylène- ou polyoxybutylène-monool à terminaison alkylphényle en C_7 à C_{30}.

15. Procédé pour réduire les dépôts dans un moteur à combustion interne, qui comprend le fonctionnement du moteur avec une composition de carburant suivant l'une quelconque des revendications précédentes.

Patentansprüche

1. Kraftstoffzusammensetzung, umfassend eine größere Menge Kohlenwasserstoffe, die im Benzin- oder Dieselbereich sieden, und 85 bis weniger als 145 Gewichtsteile pro Million einer Ablagerungen bekämpfenden Additivzusammensetzung, umfassend:

(a) 50 bis 70 Gewichtsteile pro Million eines kraftstofflöslichen Amins, das mit einem aliphatischen Kohlenwasserstoff substituiert ist und das die Formel besitzt:

$$R_3NH\text{-}(R_4\text{-}NH)_n\text{-}H,$$

wobei R_3 ein von einem Polyisobutylen abgeleiteter Kohlenwasserstoffrest mit einem Molekulargewichtszahlenmittel von 900 bis 1500 ist; R_4 ein Alkylenrest mit 2 bis 6 Kohlenstoffatomen ist; und n eine ganze Zahl von 0 bis 10 ist; und

(b) 35 bis weniger als 75 Gewichtsteile pro Million eines kohlenwasserstoffendständigen Poly(oxyalkylen) monools mit einem durchschnittlichen Molekulargewicht von 500 bis 5000, wobei der Oxyalkylen-Rest ein C_2- bis C_5-Oxyalkylenrest ist, und der Kohlenwasserstoffrest ein C_1- bis C_{30}-Kohlenwasserstoffrest ist;

wobei das Verhältnis von Poly(oxyalkylen)monool zu aliphatischem Amin im Bereich von 0,5:1 bis 1,5:1 ist.

2. Kraftstoffzusammensetzung nach Anspruch 1, wobei das aliphatische Amin der Komponente (a) ein Polyisobutylamin ist, wobei der Polyisobutylrest im Wesentlichen gesättigt ist und die Amineinheit von Ammoniak hergeleitet ist.

3. Kraftstoffzusammensetzung nach Anspruch 1, wobei die Amineinheit des aliphatischen Amins von einem Polyamin mit 2 bis 12 Amin-Stickstoffatomen und 2 bis 40 Kohlenstoffatomen hergeleitet ist.

4. Kraftstoffzusammensetzung nach Anspruch 3, wobei das Polyamin ein Polyalkylen-Polyamin mit 2 bis 12 Amin-Stickstoffatomen und 2 bis 24 Kohlenstoffatomen ist.

5. Kraftstoffzusammensetzung nach Anspruch 4, wobei das Polyalkylen-Polyamin ausgewählt ist aus der Gruppe Ethylendiamin, Diethylentriamin, Triethylentetramin und Tetraethylenpentamin.

6. Kraftstoffzusammensetzung nach Anspruch 5, wobei das Polyalkylen-Polyamin Ethylendiamin oder Diethylentriamin ist.

7. Kraftstoffzusammensetzung nach Anspruch 6, wobei das aliphatische Amin der Komponente (a) ein Polyisobutenylethylendiamin ist.

8. Kraftstoffzusammensetzung nach Anspruch 1, wobei das kohlenwasserstoffendständige Poly(oxyalkylen)monool der Komponente (b) ein durchschnittliches Molekulargewicht von 900 bis 1500 hat.

9. Kraftstoffzusammensetzung nach Anspruch 1, wobei der Oxyalkylen-Rest des kohlenwasserstoffendständigen Poly(oxyalkylen)monools der Komponente (b) ein C_3bis C_4-Oxyalkylen-Rest ist.

10. Kraftstoffzusammensetzung nach Anspruch 9, wobei der Oxyalkylenrest des kohlenwasserstoffendständigen Poly (oxyalkylen)monools der Komponente (b) ein C_3-Oxypropylen-Rest ist.

11. Kraftstoffzusammensetzung nach Anspruch 9, wobei der Oxyalkylenrest des kohlenwasserstoffendständigen Poly (oxyalkylen)monools der Komponente (b) ein C_4-Oxybutylen-Rest ist.

12. Kraftstoffzusammensetzung nach Anspruch 1, wobei der Kohlenwasserstoff-Rest des kohlenwasserstoffendständigen Poly(oxyalkylen)monools der Komponente (b) ein C_7- bis C_{30}-Alkylphenylrest ist.

13. Kraftstoffzusammensetzung nach Anspruch 1, wobei die Kraftstoffzusammensetzung 40 bis 70 Gewichtsteile pro Million eines kohlenwasserstoffendständigen Poly(oxyalkylen)monools der Komponente (b) enthält.

14. Kraftstoffzusammensetzung nach Anspruch 1, wobei die Komponente (a) ein Polyisobutenylamin ist, wobei die Amineinheit von Ethylendiamin oder Diethylentriamin hergeleitet ist und Komponente (b) ein Poly(oxypropylen)- oder Poly(oxybutylen)monool mit C_7- bis C_{30}-Alkylphenyl-Ende ist.

15. Verfahren zur Verringerung von Motorablagerungen in einem Verbrennungsmotor, umfassend den Betrieb des Motors mit einer Kraftstoffzusammensetzung nach irgend einem vorhergehenden Anspruch.