

US Patent & Trademark Office

Patent Public Search | Text View

United States Patent	12384882
Kind Code	B2
Date of Patent	August 12, 2025
Inventor(s)	Nakajima; So et al.

Polyalkylene glycol compound

Abstract

A polyalkylene glycol-based compound of formula (1): ##STR00001## may be one in which R.sup.1 is a monovalent hydrocarbon group having 1 to 32 carbon atoms, a monovalent aromatic hydrocarbon group having 6 to 42 ring carbon atoms, a monovalent acyl group having 2 to 32 carbon atoms, or a monovalent oxygen-containing hydrocarbon group having 2 to 32 carbon atoms; R.sup.2 is a monovalent hydrocarbon group having 1 to 32 carbon atoms, a monovalent aromatic hydrocarbon group having 6 to 42 ring carbon atoms, a monovalent acyl group having 2 to 32 carbon atoms, a monovalent oxygen-containing hydrocarbon group having 2 to 32 carbon atoms, or a hydrogen atom; R.sup.3 is a divalent hydrocarbon group having 4 carbon atoms; R.sup.4 is a divalent hydrocarbon group having 2 or 3 carbon atoms; m and n are respectively numbers between 1 and 40 and 0 and 20; and $m/(m+n) \geq 0.5$.

Inventors:	Nakajima; So (Edogawa-ku, JP), Yoshida; Yukio (Ichihara, JP)
Applicant:	IDEMITSU KOSAN CO., LTD. (Chiyoda-ku, JP)
Family ID:	1000008750437
Assignee:	IDEMITSU KOSAN CO., LTD. (Chiyoda-ku, JP)
Appl. No.:	17/418083
Filed (or PCT Filed):	December 26, 2019
PCT No.:	PCT/JP2019/051166
PCT Pub. No.:	WO2020/138308
PCT Pub. Date:	July 02, 2020

Prior Publication Data

Document Identifier	Publication Date
US 20220081513 A1	Mar. 17, 2022

Foreign Application Priority Data

JP

2018-247423

Dec. 28, 2018

Publication Classification

Int. Cl.: C08G65/331 (20060101); C07C41/02 (20060101); C07C41/16 (20060101); C07C43/11 (20060101); C07C43/13 (20060101); C07C43/15 (20060101); C07C43/178 (20060101); C07C43/215 (20060101); C08G65/04 (20060101); C08G65/16 (20060101); C08K11/00 (20060101); C10M105/18 (20060101); C10M107/34 (20060101); C10M169/04 (20060101); C10N30/02 (20060101); C10N30/06 (20060101); C10N30/10 (20060101); C10N30/12 (20060101); C10N30/14 (20060101); C10N30/18 (20060101); C10N20/02 (20060101); C10N20/04 (20060101); C10N40/16 (20060101); C10N40/30 (20060101)

U.S. Cl.:

CPC C08G65/331 (20130101); C07C41/02 (20130101); C07C41/16 (20130101); C07C43/11 (20130101); C07C43/13 (20130101); C07C43/15 (20130101); C07C43/1785 (20130101); C07C43/215 (20130101); C08G65/04 (20130101); C08G65/16 (20130101); C08K11/00 (20130101); C10M105/18 (20130101); C10M107/34 (20130101); C10M169/04 (20130101); C07C41/02 (20130101); C07C43/11 (20130101); C07C41/16 (20130101); C07C43/11 (20130101); C10M2209/1033 (20130101); C10M2209/1065 (20130101); C10M2209/1085 (20130101); C10N2020/02 (20130101); C10N2020/04 (20130101); C10N2030/02 (20130101); C10N2030/06 (20130101); C10N2030/10 (20130101); C10N2030/12 (20130101); C10N2030/14 (20130101); C10N2030/18 (20130101); C10N2040/16 (20130101); C10N2040/30 (20130101); C10M2209/1065 (20130101); C10M2209/1055 (20130101); C10M2209/1085 (20130101); C10M2209/1065 (20130101); C10M2209/1045 (20130101); C10M2209/1085 (20130101)

Field of Classification Search

CPC: C08G (65/331); C08G (65/16); C08G (65/04); C07C (43/15); C07C (43/1785); C07C (43/215); C07C (41/02); C07C (41/16); C07C (43/11); C07C (43/13); C08K (11/00); C10M (107/34); C10M (169/04); C10M (2209/1033); C10M (2209/1065); C10M (2209/1085); C10M (105/18); C10N (2030/02); C10N (2030/06); C10N (2030/10); C10N (2030/12); C10N (2030/14); C10N (2030/18); C10N (2020/02); C10N (2020/04); C10N (2040/16); C10N (2040/30)

References Cited

U.S. PATENT DOCUMENTS

Patent No.	Issued Date	Patentee Name	U.S. Cl.	CPC
3834935	12/1973	Symm et al.	252/8.84	D06M 15/53
4970295	12/1989	Schuchardt	N/A	N/A
5416240	12/1994	Weyer et al.	N/A	N/A
5648557	12/1996	Wei	N/A	N/A
5659068	12/1996	Weyer et al.	N/A	N/A
5741946	12/1997	Wei	N/A	N/A

5907054	12/1998	Setoyama et al.	N/A	N/A
2011/0218322	12/2010	Nakamoto et al.	N/A	N/A

FOREIGN PATENT DOCUMENTS

Patent No.	Application Date	Country	CPC
2-248426	12/1989	JP	N/A
4-130188	12/1990	JP	N/A
5-17567	12/1992	JP	N/A
5-70586	12/1992	JP	N/A
10-53647	12/1997	JP	N/A
10-508334	12/1997	JP	N/A
2000-169874	12/1999	JP	N/A
2005-281216	12/2004	JP	N/A
WO 2010/055866	12/2009	JP	N/A
2011-195725	12/2010	JP	N/A
2014-534316	12/2013	JP	N/A

OTHER PUBLICATIONS

Combined Chinese Office Action and Search Report issued Nov. 8, 2023, in corresponding Chinese Patent Application No. 201980086731.3 (with English Translation), 20 pages. cited by applicant
International Search Report mailed on Mar. 17, 2020 in PCT/JP2019/051166 filed on Dec. 26, 2019 (citing references AA-AG, AM-AU, therein, 3 pages). cited by applicant

Japanese Office Action issued on Jun. 28, 2022 in Japanese Patent Application No. 2018-247423 (with unedited computer generated English Translation), 11 pages. cited by applicant

Extended European Search Report issued May 27, 2022 in European Patent Application No. 19901476.2, 8 pages. cited by applicant

Office Action issued Mar. 24, 2023, in corresponding European Patent Application No. 19 901 476.2, 5 pages. cited by applicant

Office Action issued Mar. 11, 2024, in corresponding European Patent Application No. 19 901 476.2, 4 pages. cited by applicant

Korean Office Action issued Sep. 20, 2024 in Korean Patent Application No. 10-2021-7018822 (with unedited computer-generated English translation). 16 pages. cited by applicant

Chinese Office Action issued Nov. 14, 2024 in Chinese Patent Application No. 201980086731.3 (with English translation), 14 pages. cited by applicant

Combined Chinese Office Action and Search Report issued Sep. 7, 2024, in corresponding Chinese Patent Application No. 201980086731.3 (with English translation), citing document 15 therein, 19 pages. cited by applicant

Primary Examiner: Sergeant; Rabon A

Attorney, Agent or Firm: Oblon, McClelland, Maier & Neustadt, L.L.P.

Background/Summary

CROSS-REFERENCE TO RELATED APPLICATIONS

(1) The present application is the national stage of international application PCT/JP2019/051166, filed on Dec. 26, 2019, and claims the benefit of the filing date of Japanese Appl. No. 2018-247423, filed on Dec. 28, 2018.

TECHNICAL FIELD

(2) The present invention relates to a polyalkylene glycol-based compound. In more detail, the present invention relates to a polyalkylene glycol-based compound and a lubricating oil composition containing the polyalkylene glycol-based compound.

BACKGROUND ART

(3) As synthetic oils to be used for a lubricating oil composition, a polyalkylene glycol-based compound is known (see PTL 1). For example, for a car air conditioner of belt drive system which is mounted in a gasoline engine car or a diesel engine car, a lubricating oil composition using a polyalkylene glycol-based compound as the synthetic oil is broadly utilized.

CITATION LIST

Patent Literature

(4) PTL 1: JP-T-2014-534316

SUMMARY OF INVENTION

Technical Problem

(5) Now, hybrid cars and electric cars are becoming widespread recently. Following this, the car air conditioner which is mounted in a hybrid car or an electric car is also shifting from a belt drive system toward a motor drive system. In the car air conditioner of a motor drive system (car air conditioner of a motor drive system in which an electric compressor and a motor are integrated), the motor is dipped in a lubricating oil composition, and therefore, a winding for motor comes into direct contact with the lubricating oil composition. Then, the foregoing lubricating oil composition is required to be excellent in electric insulation.

(6) However, the conventional polyalkylene glycol-based compounds are not satisfactory in the electric insulation.

(7) An object of the present invention is to provide a polyalkylene glycol-based compound which is excellent in electric insulation.

Solution to Problem

(8) The present inventors have found that a polyalkylene glycol-based compound having a specified structure is excellent in electric insulation, thereby leading to accomplishment of the present invention.

(9) Specifically, the present invention relates to the following [1] to [5].

(10) [1] A polyalkylene glycol-based compound represented by the following general formula (1);

(11) ##STR00002##

wherein R^{sup.1} represents a monovalent hydrocarbon group having 1 to 32 carbon atoms, a monovalent aromatic hydrocarbon group having 6 to 42 ring carbon atoms, a monovalent acyl group having 2 to 32 carbon atoms, or a monovalent oxygen-containing hydrocarbon group having 2 to 32 carbon atoms; R^{sup.2} represents a monovalent hydrocarbon group having 1 to 32 carbon atoms, a monovalent aromatic hydrocarbon group having 6 to 42 ring carbon atoms, a monovalent acyl group having 2 to 32 carbon atoms, a monovalent oxygen-containing hydrocarbon group having 2 to 32 carbon atoms, or a hydrogen atom; R^{sup.3} represents a divalent hydrocarbon group having 4 carbon atoms; R^{sup.4} represents a divalent hydrocarbon group having 2 or 3 carbon atoms; m represents a number of 1 to 40; n represents a number of 0 to 20; and m/(m+) \geq 0.5.

(12) [2] The polyalkylene glycol-based compound as set forth in the above [1], wherein when the carbon number of each of the monovalent hydrocarbon group, the monovalent aromatic hydrocarbon group, the monovalent acyl group, and the monovalent oxygen-containing hydrocarbon group, which are selected as R^{sup.1} and R^{sup.2}, is 16 or less, then a hydroxyl value is 50 mgKOH/g or less.

(13) [3] The polyalkylene glycol-based compound as set forth in the above [1] or [2], wherein a volume resistivity at 25° C. is 0.0030 T Ω .Math.m or more.

(14) [4] A lubricating oil composition including the polyalkylene glycol-based compound as set forth in any one of the above [1] to [3].

(15) [5] The lubricating oil composition as set forth in the above [4], further including at least one

additive selected from an antioxidant, an oiliness improver, an oxygen scavenger, an extreme pressure agent, a copper deactivator, a rust inhibitor, an anti-foaming agent, and a viscosity index improver.

Advantageous Effects of Invention

(16) In accordance with the present invention, it is possible to provide a polyalkylene glycol-based compound which is excellent in electric insulation.

Description

DESCRIPTION OF EMBODIMENTS

(1) [Polyalkylene Glycol-Based Compound]

(2) The polyalkylene glycol-based compound of the present invention is represented by the following general formula (1).

(3) ##STR00003##

(4) In the general formula (1), R.sup.1 represents a monovalent hydrocarbon group having 1 to 32 carbon atoms, a monovalent aromatic hydrocarbon group having 6 to 42 ring carbon atoms, a monovalent acyl group having 2 to 32 carbon atoms, or a monovalent oxygen-containing hydrocarbon group having 2 to 32 carbon atoms; R.sup.2 represents a monovalent hydrocarbon group having 1 to 32 carbon atoms, a monovalent aromatic hydrocarbon group having 6 to 42 ring carbon atoms, a monovalent acyl group having 2 to 32 carbon atoms, a monovalent oxygen-containing hydrocarbon group having 2 to 32 carbon atoms, or a hydrogen atom; R.sup.3 represents a divalent hydrocarbon group having 4 carbon atoms; R.sup.4 represents a divalent hydrocarbon group having 2 or 3 carbon atoms; m represents a number of 1 to 40; n represents a number of 0 to 20; and $m/(m+n) \geq 0.5$.

(5) The polyalkylene glycol-based compound represented by the general formula (1) is high in volume resistivity and excellent in electric insulation.

(6) In this specification, among the compounds represented by the general formula (1), one in which R.sup.2 is a hydrogen atom is occasionally referred to as "one-end blocked polyalkylene glycol-based compound". In addition, among the compounds represented by the general formula (1), one in which R.sup.2 is not a hydrogen atom is occasionally referred to as "both-end blocked polyalkylene glycol-based compound".

(7) The polyalkylene glycol-based compound may be a "one-end blocked polyalkylene glycol-based compound" alone, may be a "both-end blocked polyalkylene glycol-based compound" alone, or may be a mixture of a "one-end blocked polyalkylene glycol-based compound" and a "both-end blocked polyalkylene glycol-based compound". In the present invention, these are named generically as "polyalkylene glycol-based compound". In addition, in this specification, the "polyalkylene glycol-based compound" is also occasionally abbreviated as "PAG".

(8) From the viewpoint of more improving the volume resistivity of the PAG, the polyalkylene glycol-based compound according to an embodiment of the present invention is preferably a "both-end blocked polyalkylene glycol-based compound" alone, or a mixture of a "one-end blocked polyalkylene glycol-based compound" and a "both-end blocked polyalkylene glycol-based compound"; and more preferably a "both-end blocked polyalkylene glycol-based compound" alone.

(9) R.sup.1, R.sup.2, R.sup.3, R.sup.4, m, n, and $m/(m+n)$ in the general formula (1) are hereunder described in detail.

(10) <R.sup.1 and R.sup.2>

(11) In the general formula (1), R.sup.1 represents a monovalent hydrocarbon group having 1 to 32 carbon atoms, a monovalent aromatic hydrocarbon group having 6 to 42 ring carbon atoms, a monovalent acyl group having 2 to 32 carbon atoms, or a monovalent oxygen-containing

hydrocarbon group having 2 to 32 carbon atoms.

(12) In the general formula (1), R.sup.2 represents a monovalent hydrocarbon group having 1 to 32 carbon atoms, a monovalent aromatic hydrocarbon group having 6 to 42 ring carbon atoms, a monovalent acyl group having 2 to 32 carbon atoms, a monovalent oxygen-containing hydrocarbon group having 2 to 32 carbon atoms, or a hydrogen atom.

(13) R.sup.1 and R.sup.2 may be the same as or different from each other.

(14) In this specification, the “hydrocarbon group” means a group constituted of only a carbon atom and a hydrogen atom.

(15) ((Monovalent Hydrocarbon Group Having 1 to 32 Carbon Atoms))

(16) The monovalent hydrocarbon group having 1 to 32 carbon atoms, which is selected as R.sup.1 and R.sup.2, is selected from a monovalent aliphatic hydrocarbon group having 1 to 32 carbon atoms and a monovalent hydrocarbon group having 2 to 6 bonding sites and having 2 to 32 carbon atoms.

(17) (Monovalent Aliphatic Hydrocarbon Group)

(18) The monovalent aliphatic hydrocarbon group having 1 to 32 carbon atoms, which is selected as R.sup.1 and R.sup.2, may be linear or branched, and it may also be cyclic in the case where the carbon number thereof is 3 or more.

(19) Here, from the viewpoint of more improving the volume resistivity of the PAG, the foregoing monovalent aliphatic hydrocarbon group is preferably an alkyl group.

(20) Specific examples of the alkyl group include various alkyl groups, such as a methyl group, an ethyl group, various propyl groups, various butyl groups, various pentyl groups, various hexyl groups, various heptyl groups, various octyl groups, various nonyl groups, various decyl groups, various undecyl groups, various dodecyl groups, various tridecyl groups, various tetradecyl groups, various pentadecyl groups, various hexadecyl groups, various heptadecyl group, and various octadecyl groups.

(21) Here, in this description, the term “various XXX groups” (“XXX” is a substituent name) is to include all isomers that are considered as the XXX group. For example, the term “various alkyl groups” expresses “linear, branched, or cyclic alkyl groups”. In consequence, so far as the term “various propyl groups” is concerned, it expresses “a n-propyl group, an isopropyl group, and a cyclopropyl group”, and so on. In addition, so far as the term “various butyl groups” is concerned, it expresses “a n-butyl group, a sec-butyl group, an isobutyl group, a tert-butyl group, and a cyclobutyl group”, and so on.

(22) From the viewpoint of more improving the volume resistivity of the PAG, the carbon number of the monovalent aliphatic hydrocarbon group is preferably 1 to 24, more preferably 1 to 18, still more preferably 1 to 12, yet still more preferably 1 to 6, even yet still more preferably 1 to 3, even still more preferably 1 to 2, and even still more further preferably 1.

(23) Of these, a linear alkyl group is preferred from the viewpoint of more improving the volume resistivity of the PAG.

(24) (Monovalent Hydrocarbon Group Having 2 to 6 Bonding Sites)

(25) R.sup.1 and R.sup.2 can be each provided with plural bonding site via an adjacent oxygen atom in the general formula (1).

(26) The monovalent hydrocarbon group having 2 to 6 bonding sites and having 2 to 32 carbon atoms, which is selected as R.sup.1 and R.sup.2, may be chained or may be cyclic.

(27) The monovalent hydrocarbon group having 2 bonding sites is preferably an aliphatic hydrocarbon group, and specific examples thereof include an ethylene group, a propylene group, a butylene group, a pentylene group, a hexylene group, a heptylene group, an octylene group, a nonylene group, a decylene group, a cyclopentylene group, and a cyclohexylene group. Specific examples of other monovalent hydrocarbon group having 2 bonding sites include a residue resulting from eliminating a hydroxy group from a bisphenol, such as bisphenol, bisphenol F, and bisphenol A.

- (28) The monovalent hydrocarbon group having 3 to 6 bonding sites is preferably an aliphatic hydrocarbon group, and examples thereof include a residue resulting from eliminating a hydroxy group from a polyhydric alcohol, such as trimethylolpropane, glycerin, pentaerythritol, sorbitol, 1,2,3-trihydroxycyclohexane, and 1,3,5-trihydroxycyclohexane.
- (29) ((Monovalent Aromatic Hydrocarbon Group Having 6 to 42 Ring Carbon Atoms))
- (30) The monovalent aromatic hydrocarbon group having 6 to 42 ring carbon atoms, which is selected as R.sup.1 and R.sup.2, is, for example, selected from those represented by the following general formulae (2) to (7).
- (31) ##STR00004##
- (32) In the general formulae (2) to (7), R's are each independently a methyl group, an ethyl group, or a vinyl group. A's each independently represent a group represented by the following general formula (8).
- (33) ##STR00005##
- (34) In the general formula (2), t is an integer of 0 to 5; and a is an integer of 0 to 2, provided that $(t+a) \leq 5$.
- (35) In the general formula (3), (u_1+u_2) is an integer of 0 to 7; and $(1+b_2)$ is an integer of 0 to 2, provided that $(u_1+u_2+b_1+b_2) \leq 7$.
- (36) In the general formula (4), $(v_1+v_2+v_3)$ is an integer of 0 to 9; and $(c_1+c_2+c_3)$ is an integer of 0 to 2, provided that $(v_1+v_2+v_3+c_1+c_2+c_3) \leq 9$.
- (37) In the general formula (5), $(w_1+w_2+w_3)$ is an integer of 0 to 9; and $(d_1+d_2+d_3)$ is an integer of 0 to 2, provided that $(w_1+w_2+w_3+d_1+d_2+d_3) \leq 9$.
- (38) In the general formula (6), $(x_1+x_2+x_3)$ is an integer of 0 to 9; and $(e_1+e_2+e_3)$ is an integer of 0 to 2, provided that $(x_1+x_2+x_3+e_1+e_2+e_3) \leq 9$.
- (39) In the general formula (7), $(y_1+y_2+y_3)$ is an integer of 0 to 9; and $(f_1+f_2+f_3)$ is an integer of 0 to 2, provided that $(y_1+y_2+y_3+f_1+f_2+f_3) \leq 9$.
- (40) In the general formulae (2) to (7), * represents a bonding position to the oxygen atom in the general formula (1).
- (41) In the general formula (8), z is an integer of 0 to 2; L.sup.1 is a vinylene group; and Ar.sup.1 is one kind selected from the general formulae (2) to (7) (however, when one kind selected from the general formulae (2) to (7) is Ar.sup.1, then * represents a bonding position to L.sup.1 in the general formula (8)).
- (42) In this specification, the number of ring carbon atoms of the aromatic hydrocarbon group which is selected as R.sup.1 and R.sup.2 expresses the number of carbon atoms constituting the foregoing ring itself of the compound of a structure in which the carbon atoms are bonded in a ring form. In the case where the foregoing ring is substituted with a substituent, when the substituent is an aromatic hydrocarbon group, the carbon number of the foregoing aromatic hydrocarbon group is included in the number of ring carbon atoms, too.
- (43) In the general formulae (2) to (7), the solid lines from the substituents R and A toward the aromatic ring express bonds of the substituents R and A to the carbon atoms constituting the aromatic ring and mean that the substituents R and A may be bonded to the carbon atoms constituting the aromatic ring at arbitrary positions, respectively.
- (44) In improving the volume resistivity of the PAG, with respect to the case where a, b₁, b₂, c₁, c₂, c₃, d₁, d₂, d₃, e₁, e₂, e₃, f₁, f₂, and f₃ are 0, first of all, preferred embodiments regarding R.sup.1 and R.sup.2 are hereunder described on the basis of the general formulae (2) to (7).
- (45) (Case where a, b₁, b₂, c₁, c₂, c₃, d₁, d₂, d₃, e₁, e₂, e₃, f₁, f₂, and f₃ are 0)
- (46) In the group represented by the general formula (2), t is preferably 0 to 2, more preferably 0 to 1, and still more preferably 0 (namely, unsubstituted). R is preferably a methyl group or a vinyl group. In this specification, the case of t=1 means that one R is bonded to one carbon atom forming the aromatic ring. The case of t=2 means that two R's are bonded to two carbon atoms forming the aromatic ring, respectively.

(47) Specific examples of the group represented by the general formula (2) include a phenyl group, a tolyl group, a dimethylphenyl group, and styryl group. Of these, a phenyl group, a tolyl group, and a styryl group are preferred, and a phenyl group is more preferred.

(48) In the compound of the general formula (3), (u_1+u_2) is preferably 0 to 4, more preferably 0 to 2, still more preferably 0 to 1, and yet still more preferably 0 (namely, unsubstituted). In addition, R is preferably a methyl group or a vinyl group.

(49) Specific examples of the group represented by the general formula (3) include a naphthyl group, a methylnaphthyl group, a dimethylnaphthyl group, and a vinylnaphthyl group. Of these, a naphthyl group, a methylnaphthyl group, and a vinylnaphthyl group are preferred, and a naphthyl group is more preferred.

(50) In the compounds of the general formulae (4) and (5), $(v_1+v_2+v_3)$ and $(w_1+w_2+w_3)$ are preferably 0 to 5, more preferably 0 to 3, still more preferably 0 to 2, yet still more preferably 0 to 1, and even yet still more preferably 0 (namely, unsubstituted). In addition, R is preferably a methyl group or a vinyl group.

(51) Specific examples of the groups represented by the general formulae (4) and (5) include an anthracenyl group, a methylantracenyl group, a dimethylantracenyl group, and a vinylantracenyl group. Of these, an anthracenyl group, a methylantracenyl group, and a vinylantracenyl group are preferred, and an anthracenyl group is more preferred.

(52) In the compounds of the general formulae (6) and (7), $(x_1+x_2+x_3)$ and $(y_1+y_2+y_3)$ are preferably 0 to 5, more preferably 0 to 3, still more preferably 0 to 2, yet still more preferably 0 to 1, and even yet still more preferably 0 (namely, unsubstituted). In addition, R is preferably a methyl group or a vinyl group.

(53) Specific examples of the groups represented by the general formulae (6) and (7) include a phenanthrenyl group, a methylphenanthrenyl group, a dimethylphenanthrenyl group, and a vinylphenanthrenyl group. Of these, a phenanthrenyl group, a methylphenanthrenyl group, and a vinylphenanthrenyl group are preferred, and a phenanthrenyl group is more preferred.

(54) In the general formulae (2) to (7), when plural R's are existent, the respective R's may be the same as or different from each other.

(55) Next, with respect to the case where a, b₁, b₂, c₁, c₂, c₃, d₁, d₂, d₃, e₁, e₂, e₃, f₁, f₂, and f₃ are not 0, preferred embodiments regarding R^{sup.1} and R^{sup.2} are described on the basis of the general formulae (2) to (8).

(56) (Case where a, b₁, b₂, c₁, c₂, c₃, d₁, d₂, d₃, e₁, e₂, e₃, f₁, f₂, and f₃ are not 0)

(57) In the general formula (8), though z is preferably 0 (namely, it is meant that the carbon atoms constituting the aromatic ring of the general formulae (2) to (7) are bonded directly to Ar^{sup.1}) or 1, it is more preferably 1.

(58) In the general formulae (2) to (7), the total number of the substituents A (namely, a, (b₁+b₂), (c₁+c₂+c₃), (d₁+d₂+d₃), (e₁+e₂+e₃), and (f₁+f₂+f₃)) is 1 or 2.

(59) In the case where the general formulae (2) to (7) have the substituent R, the group represented by the general formula (2) or (3) is preferred, and the group represented by the general formula (2) is more preferred. Ar^{sup.1} is preferably the group represented by the general formula (2) or (3), and more preferably the group represented by the general formula (2).

(60) In the general formulae (2) to (7), when plural A's are existent, the respective R's may be the same as or different from each other.

(61) In the general formula (8), the general formulae (2) to (7) as Ar^{sup.1} may have a substituent A. The foregoing substituent A is also represented by the general formula (8), and preferred examples of L^{sup.1} and Ar^{sup.1} are also the same as those mentioned above.

(62) When the general formulae (2) to (7) have the substituent A, preferred specific examples thereof include a biphenyl group, a binaphthyl group, and a group derived from distyrylbenzene. Of these, a group derived from distyrylbenzene is preferred. The "group derived from distyrylbenzene" means a group in which in distyrylbenzene, the hydrogen atom is eliminated at a

position serving as a bonding point to the oxygen atom. The bonding point may be the carbon atom in the central benzene ring of distyrylbenzene or may be the carbon atom of the end benzene ring. (63) Even in the case where a, b1, b2, c1, c2, c3, d1, d2, d3, e1, e2, e3, f1, f2, and f3 are not 0, the substituent R may be existent in the general formulae (2) to (7). In this case, the total number of the substituents R (t, (u1+u2), (v1+v2+v3), (w1+w2+w3), (x1+x2+x3)), and (y1+y2+y3) and the kinds thereof are the same as those mentioned above.

(64) ((Monovalent Acyl Group Having 2 to 32 Carbon Atoms))

(65) The hydrocarbon group moiety of the monovalent acyl group having 2 to 32 carbon atoms, which is selected as R.sup.1 and R.sup.2, may be linear, branched, or cyclic. The hydrocarbon group moiety of the monovalent acyl group is preferably an alkyl group, and specific examples of the alkyl group include the aforementioned specific examples.

(66) ((Monovalent Oxygen-Containing Hydrocarbon Group Having 2 to 32 Carbon Atoms))

(67) Examples of the oxygen-containing hydrocarbon group having 2 to 32 carbon atoms, which is selected as R.sup.1 and R.sup.2, include a chained aliphatic group and a cyclic aliphatic group, each having an ether bond. Specific examples thereof include a tetrahydrofurfuryl group.

(68) ((Preferred Substituent Selected as R.sup.1 and R.sup.2))

(69) The substituents which are selected as R.sup.1 and R.sup.2 are each independently preferably a monovalent hydrocarbon group having 1 to 32 carbon atoms or a monovalent aromatic hydrocarbon group having the number of ring carbon atoms of 6 to 42 (preferably 6 to 30, more preferably 6 to 24, and still more preferably 6 to 20). Of these, a monovalent aliphatic hydrocarbon group having 1 to 32 carbon atoms or a monovalent aromatic hydrocarbon group having 6 to 42 ring carbon atoms is preferred, and a monovalent aliphatic hydrocarbon group having 1 to 32 carbon atoms is more preferred.

(70) Preferably, at least one of R.sup.1 and R.sup.2 is an alkyl group; more preferably, at least one of R.sup.1 and R.sup.2 is an alkyl group which has 1 to 3 carbon atoms and/or is linear; still more preferably, both R.sup.1 and R.sup.2 are an alkyl group which has 1 to 3 carbon atoms and/or is linear; and yet still more preferably, both R.sup.1 and R.sup.2 are a methyl group.

(71) In this way, in view of the fact that both R.sup.1 and R.sup.2 are an alkyl group having a small carbon atom number, in the case where the PAG is used upon being mixed with a refrigerant, favorable compatibility with the refrigerant may be exhibited.

(72) <R.sup.3>

(73) In the general formula (1), R.sup.3 represents a divalent hydrocarbon group having 4 carbon atoms.

(74) Examples of the divalent hydrocarbon group having 4 carbon atoms, which may be selected as R.sup.3, include various butylene groups. The butylene group may be linear or may be branched.

(75) As for the conventional PAG's, ethylene oxide having 2 carbon atoms or propylene oxide having 3 carbon atoms was used as the monomer. But, in the case where a monomer unit of the PAG is formed of only ethylene oxide or propylene oxide, the volume resistivity of the PAG is liable to become low. For that reason, in a car air conditioner of a motor drive system (car air conditioner of a motor drive system in which an electric compressor and a motor are integrated), there is a concern that when a lubricating oil composition constituted of the PAG comes into direct contact with a winding for motor, a risk for safety to be caused due to an electric leakage, or the like is generated.

(76) In contrast, as in the present invention, when R.sup.3 in the general formula (1) is a divalent hydrocarbon group having 4 carbon atoms, the volume resistivity of the PAG is enhanced, whereby the electric insulation can be made excellent, and a risk for safety to be caused due to an electric leakage, or the like can be inhibited. In consequence, it becomes possible to use the PAG as the lubricating oil composition in a car air conditioner of a motor drive system.

(77) <R.sup.4>

(78) In the general formula (1), R.sup.4 represents a divalent hydrocarbon group having 2 or 3

carbon atoms.

(79) Examples of the divalent hydrocarbon group having 2 or 3 carbon atoms, which may be selected as R.sup.4, include an ethylene group and various propylene groups.

(80) <m>

(81) In the general formula (1), m is a number of 1 to 40.

(82) The number of m is an average value of the number of the (R.sup.3O) unit and is also an average addition molar number of the (R.sup.3O) unit.

(83) In the case where plural (R.sup.3O) units are existent, namely, in the case of $m \geq 2$, the respective (R.sup.3O) units may be the same as or different from each other.

(84) In the case where the respective (R.sup.3O) units are different from each other, the respective (R.sup.3O) units may be subjected to random addition or may be subjected to block addition.

However, the random addition is preferred from the viewpoint of handling properties.

(85) From the viewpoint of more improving the volume resistivity of the PAG, the number of m is preferably 5 to 35, more preferably 10 to 30, and still more preferably 10 to 20.

(86) <n>

(87) In the general formula (1), n is a number of 0 to 20.

(88) The number of n is an average value of the number of the (R.sup.4O) unit and is also an average addition molar number of the (R.sup.4O) unit.

(89) In the case where plural (R.sup.4O) units are existent, namely, in the case of $n \geq 2$, the respective (R.sup.4O) units may be the same as or different from each other. In the case where the respective (R.sup.4O) units are different from each other, the respective (R.sup.4O) units may be subjected to random addition or may be subjected to block addition. However, the random addition is preferred from the viewpoint of handling properties.

(90) From the viewpoint of more improving the volume resistivity of the PAG, the number of n is desirably smaller, and it is preferably 0 to 10, more preferably 0 to 5, still more preferably 0 to 3, yet still more preferably 0 to 2, even yet still more preferably 0 to 1, and even still more preferably 0.

(91) The (R.sup.3O) unit and the (R.sup.4O) unit may be subjected to random addition or may be subjected to block addition. However, the random addition is preferred from the viewpoint of handling properties.

(92) <m/(m+n)>

(93) Here, in the general formula (1), $m/(m+n)$ is 0.5 or more.

(94) The value of " $m/(m+n)$ " is a value corresponding to the proportion at which the (R.sup.3O) unit occupies relative to the total number of the (R.sup.3O) unit and the (R.sup.4O) unit. When this value is less than 0.5, the volume resistivity of the PAG becomes insufficient, and the electric insulation is inferior.

(95) From the viewpoint of more improving the volume resistivity of the PAG, the value of " $m/(m+n)$ " is preferably 0.6 or more, more preferably 0.7 or more, still more preferably 0.8 or more, yet still more preferably 0.9 or more, even yet still more preferably 0.95 or more, and even still more preferably 1.0.

(96) [Physical Properties of Polyalkylene Glycol-Based Compound]

(97) The volume resistivity, the hydroxyl value, the mass average molecular weight, the number average molecular weight, the 40° C. kinematic viscosity, the 100° C. kinematic viscosity, and the viscosity index of the polyalkylene glycol-based compound according to an embodiment of the present invention are hereunder described.

(98) <Volume Resistivity>

(99) The volume resistivity at 25° C. of the PAG according to an embodiment of the present invention is preferably 0.0030 TΩ.Math.m or more, more preferably 0.0050 Tam or more, still more preferably 0.0100 TΩ.Math.m or more, yet still more preferably 0.0200 TΩ.Math.m or more, even yet still more preferably 0.0400 TΩ.Math.m or more, even still more preferably 0.0600

TΩ.Math.m or more, and even still more further preferably 0.0800 TΩ.Math.m or more, and it is typically 1 TΩ.Math.m or less.

(100) In this specification, the volume resistivity at 25° C. is a value obtained through measurement at room temperature (25° C.) in conformity with JIS C2101-24 (volume resistivity test).

(101) <Hydroxyl Value>

(102) In the case where the carbon number of each of the aforementioned monovalent hydrocarbon group, the aforementioned monovalent aromatic hydrocarbon group, the aforementioned monovalent acyl group, and the aforementioned monovalent oxygen-containing hydrocarbon group, which are selected as R.sup.1 and R.sup.2, is 16 or less, the hydroxyl value of the PAG according to an embodiment of the present invention is preferably 50 mgKOH/g or less, more preferably 45 mgKOH/g or less, still more preferably 30 mgKOH/g or less, yet still more preferably 10 mgKOH/g or less, and even yet still more preferably 5 mgKOH/g or less. In addition, it is typically 0.1 mgKOH/g or more. The carbon number of the monovalent aromatic hydrocarbon group as referred to herein means the sum total of the ring carbon number and the carbon number of the substituent.

(103) The hydroxyl value of the PAG is an index expressing the blocked state of the hydroxy groups at the both ends of the PAG by the substituent. In the case where the carbon number of each of the aforementioned monovalent hydrocarbon group, the aforementioned monovalent aromatic hydrocarbon group, the aforementioned monovalent acyl group, and the aforementioned monovalent oxygen-containing hydrocarbon group, which are selected as R.sup.1 and R.sup.2, is 16 or less, so far as the hydroxyl value is 50 mgKOH/g or less, the PAG in which the both ends are not blocked is not substantially existent in the mixture. Then, as the hydroxyl value becomes smaller than this value, the proportion at which the “both-end blocked polyalkylene glycol-based compound” occupies in the mixture of the “one-end blocked polyalkylene glycol-based compound” and the “both-end blocked polyalkylene glycol-based compound” increases, and the volume resistivity of the PAG is improved.

(104) In the PAG according to an embodiment of the present invention, in the case where the carbon number of each of the aforementioned monovalent hydrocarbon group, the aforementioned monovalent aromatic hydrocarbon group, the aforementioned monovalent acyl group, and the aforementioned monovalent oxygen-containing hydrocarbon group, which are selected as R.sup.1 and R.sup.2, is more than 16, so far as the hydroxyl value is 65 mgKOH/g or less, the PAG in which the both ends are not blocked is not substantially existent in the mixture. Then, as the hydroxyl value becomes smaller than this value, the proportion at which the “both-end blocked polyalkylene glycol-based compound” occupies in the mixture of the “one-end blocked polyalkylene glycol-based compound” and the “both-end blocked polyalkylene glycol-based compound” increases, and the volume resistivity of the PAG is improved.

(105) In this specification, the hydroxyl value is a value through measurement by the neutralization titration method in conformity with JIS K0070.

(106) <Mass Average Molecular Weight (Mw) and Number Average Molecular Weight (Mn)>

(107) The mass average molecular weight (Mw) of the polyalkylene glycol-based compound according to an embodiment of the present invention is preferably 500 to 5,000, more preferably 700 to 3,000, and still more preferably 800 to 2,000.

(108) The number average molecular weight (Mn) of the polyalkylene glycol-based compound according to an embodiment of the present invention is preferably 500 to 5,000, more preferably 600 to 2,500, and still more preferably 700 to 1,800.

(109) The mass average molecular weight (Mw) and the number average molecular weight (Mn) can be measured by the method described in the section of Examples as mentioned later.

(110) <40° C. Kinematic Viscosity, 100° C. Kinematic Viscosity, and Viscosity Index>

(111) The 40° C. kinematic viscosity of the polyalkylene glycol-based compound according to an embodiment of the present invention is preferably 10 to 400 mm.sup.2/s, more preferably 20 to 300

mm.sup.2/s, still more preferably 20 to 200 mm.sup.2/s, and yet still more preferably 30 to 100 mm.sup.2/s.

(112) The 100° C. kinematic viscosity of the polyalkylene glycol-based compound according to an embodiment of the present invention is preferably 2.0 to 30 mm.sup.2/s, more preferably 3.0 to 25 mm.sup.2/s, still more preferably 4.0 to 20 mm.sup.2/s, and yet still more preferably 5.0 to 18 mm.sup.2/s.

(113) The viscosity index of the polyalkylene glycol-based compound according to an embodiment of the present invention is preferably 70 to 250, more preferably 80 to 230, and still more preferably 90 to 220.

(114) The 40° C. kinematic viscosity, the 100° C. kinematic viscosity, and the viscosity index are each a value measured and calculated by using a glass-made capillary viscometer in conformity with JIS K2283-2000.

(115) [Production Method of Polyalkylene Glycol-Based Compound]

(116) The production method of the polyalkylene glycol-based compound of the present invention is not particularly limited.

(117) As for the production method of the polyalkylene glycol-based compound according to an embodiment of the present invention, the one-end blocked polyalkylene glycol-based compound can be produced by addition-polymerizing an oxirane monomer in the presence of an alkali metal alkoxide and then removing the alkali metal ion. In the case of introducing the group having the substituent A, such as a group derived from distyrylbenzene, a compound in which a hydroxy group is introduced into the compound serving as a constituent source of the substituent A may be used in combination with the alkali metal alkoxide, such as sodium methoxide. For example, in the case of introducing a group derived from distyrylbenzene as the substituent A, a compound having a hydroxy group added to a benzene ring of trans, trans-1-styryl-4-styrylbenzene may be used in combination.

(118) The alkali metal alkoxide is obtained through alkoxylation of an alcohol with a hydride of an alkaline metal. The alcohol to be used is changed according to R.sup.1 of the general formula (1). As the alkali metal, at least one selected from sodium, potassium, and the like is used, with sodium being preferred.

(119) As the oxirane monomer, butylene oxide is used, and taking into consideration the “value of n” and the “value of m/(m+n)” of the general formula (1), ethylene oxide and/or propylene oxide are/is further used. In the case of using ethylene oxide and/or propylene oxide in addition to butylene oxide, blending proportions of these oxides are determined taking into consideration the “value of n” and the “value of m” of the general formula (1).

(120) On the occasion of addition-polymerizing the oxirane monomer in the presence of an alkali metal alkoxide, for example, the reaction temperature is 85 to 125° C., the reaction time is 8 to 24 hours, and the reaction pressure is 0.1 to 3 MPa.

(121) Examples of a method for removing the alkali metal ion from the reaction product include a method of using an ion exchange resin. Specifically, there is exemplified a method in which a solution in which the reaction product is dissolved in a mixed solvent of water and methanol is passed through a column having a cation exchange resin filled therein and then passed through a column having an anion exchange resin filled therein.

(122) Thereafter, by removing the mixed solvent by means of distillation or the like, the one-end blocked polyalkylene glycol-based compound can be produced.

(123) The both-end blocked polyalkylene glycol-based compound is obtained by reacting the one-end blocked polyalkylene glycol compound with an alkali metal compound, such as sodium methoxide, to substitute the hydrogen atom of the hydroxy group with the alkali metal and subsequently, reacting this with a compound represented by the following general formula (9) to substitute the alkali metal moiety with R.sup.2.

X—R.sup.2 (9)

(124) In the general formula (9), X is at least one selected from a fluorine atom, a chlorine atom, a bromine atom, and an iodine atom; and R^{sup.2} is the same as that in the general formula (1) (provided that a hydrogen atom is excluded).

(125) On the occasion of reacting the one-end blocked polyalkylene glycol compound with the alkali metal compound, such as sodium methoxide, for example, the reaction temperature is 20 to 60° C., the reaction time is 1 to 2 hours, and the reaction pressure is 0.1 to 1 MPa.

(126) On the occasion of reacting the one-end blocked polyalkylene glycol compound with the alkali metal compound, such as sodium methoxide and then reacting with the compound represented by the general formula (9) to substitute the alkali metal moiety with R^{sup.2}, for example, the reaction temperature is 60 to 90° C., the reaction time is 8 to 24 hours, and the reaction pressure is 0.1 to 1 MPa.

(127) Examples of a method for removing an alkali metal halide from the reaction product include a method in which a solution in which the reaction product is dissolved in a mixed solvent of water and methanol is passed through a column having a cation exchange resin filled therein and then passed through a column having an anion exchange resin filled therein.

(128) Thereafter, by removing the mixed solvent by means of distillation or the like, the both-end blocked polyalkylene glycol-based compound can be produced.

(129) In the reaction product, the proportion at which the both-end blocked polyalkylene glycol-based compound occupies relative to the total amount of the one-end blocked polyalkylene glycol-based compound and the both-end blocked polyalkylene glycol-based compound can be regulated by the blending amount of the alkali metal compound, such as sodium methoxide, relative to the one-end blocked polyalkylene glycol compound on the occasion of reacting the one-end blocked polyalkylene glycol compound with the alkali metal compound, the blending amount of the compound represented by the general formula (1) relative to the one-end blocked polyalkylene glycol compound, the reaction time, the reaction temperature, the reaction pressure, and so on.

(130) [Lubricating Oil Composition]

(131) The lubricating oil composition of the present invention contains the aforementioned PAG. In view of the fact that the lubricating oil composition contains the PAG, the electric insulation is improved.

(132) In the lubricating oil composition, the content of the PAG is typically 50 to 100% by mass, preferably 60 to 100% by mass, more preferably 70 to 100% by mass, yet still more preferably 80 to 100% by mass, and even yet still more preferably 90 to 100% by mass on the basis of the whole amount (100% by mass) of the lubricating oil composition.

(133) The lubricating oil composition may further contain at least one additive selected from an antioxidant, an oiliness improver, an oxygen scavenger, an extreme pressure agent, a copper deactivator, a rust inhibitor, an anti-foaming agent, and a viscosity index improver within the range that does not hinder the effect of the PAG.

(134) The lubricating oil composition may be one composed of only the PAG according to an application.

(135) <Lubricating Oil Composition for Refrigerator>

(136) The lubricating oil composition is preferably used as a lubricating oil composition for refrigerator (hereinafter also referred to as “refrigerator oil”). That is, the foregoing refrigerator oil is used upon being filled together with a refrigerant in the interior of a refrigerator, and for example, it is used for lubricating a sliding part of a compressor provided in the refrigerator, or the like. In the following description, the mixture of the refrigerator oil and the refrigerant is also referred to as a “refrigerator oil composition”.

(137) In view of the fact that the refrigerator oil containing the PAG is excellent in electric insulation, a risk for safety to be caused due to an electric leakage, or the like can be inhibited. In consequence, the refrigerator oil is preferably used as a refrigerator oil in a car air conditioner of a motor drive system. As a matter of course, the refrigerator oil can also be used in a car air

conditioner of a belt drive system, an air-conditioning machine, an icebox, an automatic vending machine, a showcase, a refrigeration system, a hot water supplying system, or a heating system, other than the car air conditioner of a motor drive system.

(138) As the refrigerant which is used upon being mixed with the refrigerator oil, there is exemplified at least one selected from an unsaturated fluorinated hydrocarbon compound (hereinafter also referred to as “HFO refrigerant”), a saturated fluorinated hydrocarbon compound (hereinafter also referred to as “HFC refrigerant”), a hydrocarbon-based refrigerant (hereinafter also referred to as “HC refrigerant”), carbon dioxide, and ammonia. Of these, a refrigerant including an HFO refrigerant is preferably used.

(139) (HFO Refrigerant)

(140) Examples of the HFO refrigerant include compounds having a carbon-carbon double bond, such as fluorides of a linear or branched chained olefin having 2 or more and 6 or less carbon atoms; and a cyclic olefin having 4 or more and 6 or less carbon atoms.

(141) More specifically, examples thereof include an ethylene having 1 or more and 3 or less (preferably 3) fluorine atoms introduced thereinto; a propene having 1 or more and 5 or less fluorine atoms introduced thereinto; a butene having 1 or more and 7 or less fluorine atoms introduced thereinto; a pentene having 1 or more and 9 or less fluorine atoms introduced thereinto; a hexene having 1 or more and 11 or less fluorine atoms introduced thereinto; a cyclobutene having 1 or more and 5 or less fluorine atoms introduced thereinto; a cyclopentene having 1 or more and 7 or less fluorine atoms introduced thereinto; and a cyclohexene having 1 or more and 9 or less fluorine atoms introduced thereinto.

(142) Of these HFO refrigerants, a fluoride of a propene is preferred, a propene having 3 or more and 5 or less fluorine atoms introduced thereinto is more preferred, and a propene having 4 fluorine atoms introduced thereinto is still more preferred.

(143) Suitable examples of the HFO refrigerant include 1,2,3,3,3-pentafluoropropene (R1225ye), 2,3,3,3-tetrafluoropropene (R1234yf), 1,3,3,3-tetrafluoropropene (R1234ze), 1,2,3,3-tetrafluoropropene (R1234yz), 1,1,2-trifluoroethylene (R1123), and (Z)-1,1,1,4,4,4-hexafluoro-2-butene (R1336mzz(Z)). Of these HFO refrigerants, at least one selected from the group consisting of R1234yf, R1234ze, R1123, and R1336mzz(Z) is preferred; at least one selected from the group consisting of R1234yf, R1234ze, and R1336mzz(Z) is more preferred; and at least one selected from the group consisting of R1234yf and R1234ze is still more preferred.

(144) (HFC Refrigerant)

(145) The HFC refrigerant is preferably a fluoride of an alkane having 1 or more and 4 or less carbon atoms, more preferably a fluoride of an alkane having 1 or more and 3 or less carbon atoms, and still more preferably a fluoride of an alkane having 1 or 2 carbon atoms (methane or ethane). Examples of the fluoride of methane or ethane include trifluoromethane (R23), difluoromethane (R32), 1,1-difluoroethane (R152a), 1,1,1-trifluoroethane (R143a), 1,1,2-trifluoroethane (R143), 1,1,1,2-tetrafluoroethane (R134a), 1,1,2,2-tetrafluoroethane (R134), and 1,1,1,2,2-pentafluoroethane (R125). Of these HFC refrigerants, at least one selected from the group consisting of R32, R134a, and R125 is preferred, and R32 is more preferred.

(146) (HC Refrigerant)

(147) The hydrocarbon-based refrigerant is preferably a hydrocarbon having 1 or more and 8 or less carbon atoms, more preferably a hydrocarbon having 1 or more and 5 or less carbon atoms, and still more preferably a hydrocarbon having 3 or more and 5 or less carbon atoms. When the carbon number is 8 or less, a boiling point of the refrigerant does not become excessively high, and hence, such is preferred as the refrigerant. As the hydrocarbon-based refrigerant, there is exemplified at least one selected from the group consisting of methane, ethane, ethylene, propane (R290), cyclopropane, propylene, n-butane, isobutane (R600a), 2-methylbutane, n-pentane, isopentane, cyclopentane, isobutane, and n-butane.

(148) (Content Ratio of Refrigerator Oil and Refrigerant)

(149) In the refrigerator oil composition, a content ratio of the refrigerator oil and the refrigerant ((refrigerator oil)/refrigerant) is preferably 1/99 to 99/1, more preferably 1/99 to 90/10, still more preferably 5/95 to 70/30, and yet still more preferably 5/95 to 60/40 in terms of a mass ratio. By controlling the content ratio of the refrigerator oil and the refrigerant to the aforementioned range, lubricity and suitable refrigeration ability in the refrigerator are exhibited.

(150) [Moisture Content in Refrigerator Oil Composition]

(151) The moisture content of the refrigerator oil composition according to an embodiment of the present invention is preferably 800 ppm by mass or less, more preferably 700 ppm by mass or less, still more preferably 500 ppm by mass or less, and yet still more preferably 300 ppm by mass or less. In the refrigerator oil composition according to an embodiment of the present invention, though a lower limit value of the moisture content is not particularly limited, it is preferably 50 ppm by mass or more.

(152) (Other Application of Lubricating Oil Composition)

(153) The lubricating oil composition may also be used for a transmission, a shock absorber, various gear apparatuses, various bearing mechanisms, other various industrial apparatuses, and so on, other than the refrigerator application.

(154) (Apparatus)

(155) As the apparatus according to an embodiment of the present invention, there is exemplified at least one selected from a car air conditioner of a motor drive system (car air conditioner of a motor drive system in which an electric compressor and a motor are integrated), a car air conditioner of a belt drive system, an air-conditioning machine, an icebox, an automatic vending machine, a showcase, a refrigeration system, a hot water supplying system, and a heating system, each having the lubricating oil composition filled therein.

(156) In addition, examples of the apparatus according to an embodiment of the present invention include an industrial apparatus having the lubricating oil composition filled therein. As the industrial apparatus, there is exemplified at least one selected from a transmission, a shock absorber, various gear structures, and various bearing mechanisms.

EXAMPLES

(157) The present invention is hereunder specifically described by reference to Examples, but it should be construed that the present invention is not limited to the following Examples.

Production Examples

(158) PAG's were prepared by the following Production Examples.

Production Example A1

(159) (Step A1-1)

(160) A 200 mL-volume stainless steel-made autoclave installed with a stirrer and a liquid-introducing tube (hereinafter also referred to simply as "autoclave") was charged with 3.0 g (0.056 mol) of powdered sodium methoxide and then hermetically sealed. Then, an autoclave temperature was raised to 105° C., and 95 g (1.32 mol) of butylene oxide was injected into the autoclave from the liquid-introducing tube under stirring over 9 hours, to obtain a reaction product. The reaction product was subjected to temperature lowering to room temperature (25° C.), and this was then dissolved in 100 mL of water and 200 mL of methanol, to prepare a solution of the reaction product. The foregoing solution was passed through a column having 200 mL of a cation exchange resin filled therein and then passed through a column having 200 mL of an anion exchange resin filled therein, to remove the sodium ion from the reaction product. Subsequently, the methanol and water were distilled off from the solution after passing through the columns, and the residue was dried under reduced pressure with a vacuum pump (0.4 mmHg) at 100° C. for 1 hour, to obtain 91 g of a one-end blocked polyalkylene glycol-based compound A1 (R_{sup.1}=methyl group (Me), R_{sup.2}=H, R_{sup.3}=butylene group (monomer: butylene oxide (BO), hereinafter also referred to as "C4"), m=15, n=0).

(161) (Step A1-2)

(162) In a 300 mL-volume glass-made three-necked flask installed with a stirrer and a distillation head (hereinafter also referred to simply as “flask”), 50 g of the one-end blocked polyalkylene glycol-based compound A1 obtained in Step A1-1 and 80 mL of toluene were charged and then stirred while heating to distil off about 20 mL of the toluene, thereby removing the moisture remained in the charged one-end blocked polyalkylene glycol-based compound A1. Subsequently, after temperature lowering to room temperature (25° C.), 25 g of a methanol solution of sodium methoxide (sodium methoxide concentration: 28% by mass) (sodium methoxide: 0.13 mol) was charged in the flask and heated to distill off the methanol and about 20 mL of the toluene.

(163) After temperature lowering to room temperature (25° C.), the contents in the flask were transferred into the autoclave, and the autoclave was charged with 36.8 g (0.26 mol) of methyl iodide and then hermetically sealed. Subsequently, an autoclave temperature was raised under stirring from 50° C. to 70° C. over 4.5 hours, the temperature was then raised to 85° C. over 3 hours, and the reaction was performed at 85° C. for 12 hours, to obtain a reaction product. The reaction product was subjected to temperature lowering to room temperature (25° C.), and this was then dissolved in 100 mL of water and 200 mL of methanol, to prepare a solution of the reaction product. The foregoing solution was passed through a column having 200 mL of a cation exchange resin filled therein and then passed through a column having 200 mL of an anion exchange resin filled therein. Subsequently, the methanol and water were distilled off from the solution after passing through the columns, and the residue was dried under reduced pressure with a vacuum pump (0.1 mmHg) at 100° C. for 1 hour, to obtain 42.5 g of PAG-A1.

(164) As for the PAG-A1, the fact that an infrared absorption spectrum (3,450 cm.^{sup.}-1) assigned to the hydroxy group vanished reveals R.^{sup.}2=Me.

Production Example A2

(165) (Step A2-1)

(166) The same procedures as in Step A1-1 of Production Example A1 were followed, except that the amount of the powdered sodium methoxide was changed to 2.8 g, to obtain 90 g of a one-end blocked polyalkylene glycol-based compound A2 (R.^{sup.}1=Me, R.^{sup.}2=H, R.^{sup.}3=C4, m=17, n=0).

(167) (Step A2-2)

(168) The same procedures as in Step A1-2 of Production Example A1 were followed, except that the reaction time with methyl iodide at 85° C. was changed to 4 hours, to obtain 42 g of PAG-A2.

(169) From the measurement results regarding the hydroxyl value as shown in Table 1, as for the PAG-A2, it is noted that a compound of R.^{sup.}2=Me and a compound of R.^{sup.}2=H coexist.

Production Example A3

(170) (Step A3-1)

(171) The same procedures as in Step A1-1 of Production Example A1 were followed, except that the amount of the powdered sodium methoxide was changed to 2.7 g, to obtain 90 g of a one-end blocked polyalkylene glycol-based compound A3 (R.^{sup.}1=Me, R.^{sup.}2=H, R.^{sup.}3=C4, m=18, n=0).

(172) (Step A3-2)

(173) The same procedures as in Step A1-2 of Production Example A1 were followed, except that the reaction time with methyl iodide at 85° C. was changed to 2 hours, to obtain 45 g of PAG-A3.

(174) From the measurement results regarding the hydroxyl value as shown in Table 1, as for the PAG-A3, it is noted that though a compound of R.^{sup.}2=Me and a compound of R.^{sup.}2=H coexist, the proportion at which the compound of R.^{sup.}2=Me occupies is lower than that of the PAG-A2.

Production Example A4

(175) (Step A4-1)

(176) The same procedures as in Step A1-1 of Production Example A1 were followed, except that the amount of the powdered sodium methoxide was changed to 4.0 g, and that in order to introduce

a group (DSB) derived from distyrylbenzene into R.sup.1 together with the foregoing powdered sodium methoxide, 40 g of a compound having a hydroxy group added to a benzene ring of trans, trans-1-styryl-4-styrylbenzene was charged, to obtain 110 g of a one-end blocked polyalkylene glycol-based compound A4 (R.sup.1=DSB, R.sup.2=H, R.sup.3=C4, m=6, n=0).

(177) (Step A4-2)

(178) The same procedures as in Step A1-2 of Production Example A1 were followed, except that the methyl iodide was changed to 110 g of distyrylbenzene chloride (compound in which a chlorine atom is added to the benzene ring of trans, trans-1-styryl-4-styrylbenzene), to obtain 180 g of PAG-A4.

(179) From the measurement results regarding the hydroxyl value as shown in Table 1, as for the PAG-A4, it is noted that a compound of R.sup.2=DSB and a compound of R.sup.2=H coexist.

Production Example A5

(180) (Step A5-1)

(181) The same procedures as in Step A1-1 of Production Example A1 were followed, except that the amount of the powdered sodium methoxide was changed to 4.0 g, and that 30 g of oleyl alcohol was charged together with the foregoing powdered sodium methoxide, to obtain 100 g of a one-end blocked polyalkylene glycol-based compound A5 (R.sup.1=oleyl group (Oleyl), R.sup.2=H, R.sup.3=C4, m=7, n=0).

(182) (Step A5-2)

(183) The same procedures as in Step A1-2 of Production Example A1 were followed, except that the methyl iodide was changed to 100 g of oleyl chloride, to obtain 165 g of PAG-A5.

(184) From the measurement results regarding the hydroxyl value as shown in Table 1, as for the PAG-A5, it is noted that a compound of R.sup.2=Oleyl and a compound of R.sup.2=H coexist.

Production Example A6

(185) (Step A6-1)

(186) The same procedures as in Step A1-1 of Production Example A1 were followed, except that the amount of the butylene oxide was changed to 47.5 g, and that 38.3 g of propylene oxide was injected together with the foregoing butylene oxide, to obtain 81 g of a one-end blocked polyalkylene glycol-based compound A6 (R.sup.1=Me, R.sup.2=H, R.sup.3=C4, R.sup.4=propylene group (monomer: propylene oxide, hereinafter also referred to as "C3"), m=9, n=9).

(187) (Step A6-2)

(188) The same procedures as in Step A1-2 of Production Example A1 were followed to obtain 35 g of PAG-A6.

(189) From the measurement results regarding the hydroxyl value as shown in Table 1, as for the PAG-A6, it is noted that a compound of R.sup.2=Me and a compound of R.sup.2=H coexist.

Production Example B1

(190) (Step B1-1)

(191) The same procedures as in Step A1-1 of Production Example A1 were followed, except that the butylene oxide was changed to 77 g of propylene oxide, to obtain 75 g of a one-end blocked polyalkylene glycol-based compound B1 (R.sup.4=Me, R.sup.2=H, R.sup.4=C3, m=0, n=19).

(192) (Step B1-2)

(193) The same procedures as in Step A1-2 of Production Example A1 were followed to obtain 38 g of PAG-B1.

(194) As for the PAG-B1, the fact that an infrared absorption spectrum (3,450 cm.sup.-1) assigned to the hydroxy group vanished reveals R.sup.2=Me.

Production Example 2

(195) (Step B2-1)

(196) The same procedures as in Step B1-1 of Production Example B1 were followed to obtain 75 g of a one-end blocked polyalkylene glycol-based compound B2 (R.sup.1=Me, R.sup.2=H,

R.sup.4=C3, m=0, n=19).

(197) (Step B2-2)

(198) The same procedures as in Step B1-2 of Production Example B1 were followed, except that the reaction time with methyl iodide at 85° C. was changed to 6 hours, to obtain 45 g of PAG-B2.

(199) From the measurement results regarding the hydroxyl value as shown in Table 1, as for the PAG-B2, it is noted that a compound of R.sup.2=Me and a compound of R.sup.2=H coexist.

Production Example B3

(200) (Step B3-1)

(201) The same procedures as in Step B1-1 of Production Example B1 were followed, except that the amount of the powdered sodium methoxide was changed to 2.7 g, to obtain 74 g of a one-end blocked polyalkylene glycol-based compound B3 (R.sup.1=Me, R.sup.2=H, R.sup.4=C3, m=0, n=21).

(202) (Step B3-2)

(203) The same procedures as in Step B1-2 of Production Example B1 were followed, except that the reaction time with methyl iodide at 85° C. was changed to 1 hour, to obtain 51 g of PAG-B3.

(204) From the measurement results regarding the hydroxyl value as shown in Table 1, as for the PAG-B3, it is noted that though a compound of R.sup.2=Me and a compound of R.sup.2=H coexist, the proportion at which the compound of R.sup.2=Me occupies is lower than that of the PAG-B2.

Production Example B4

(205) (Step B4-1)

(206) The same procedures as in Step B1-1 of Production Example B1 were followed, except that the amount of the propylene oxide was changed to 61.6 g, and that 11.5 g of ethylene oxide was injected together with the foregoing propylene oxide, to obtain 73 g of a one-end blocked polyalkylene glycol-based compound B4 (R.sup.1=Me, R.sup.2=H, R.sup.4=ethylene group (monomer: ethylene oxide (EO), hereinafter also referred to as "C2") and C3, m=0, n=20).

(207) (Step B4-2)

(208) The same procedures as in Step B1-2 of Production Example B1 were followed, except that the reaction time with methyl iodide at 85° C. was changed to 6 hours, to obtain 57 g of PAG-B4.

(209) From the measurement results regarding the hydroxyl value as shown in Table 1, as for the PAG-B4, it is noted that a compound of R.sup.2=Me and a compound of R.sup.2=H coexist.

Examples 1 to 6 and Comparative Examples 1 to 4

(210) With respect to PAG-A1 to PAG-A6 and PAG-B1 to PAG-B4 synthesized in Production Examples A1 to A6 and Comparative Examples B1 to B4, respectively, the mass average molecular weight (Mw), the number average molecular weight (Mn), the 40° C. kinematic viscosity, the 100° C. kinematic viscosity, the viscosity index, the hydroxyl value, and the volume resistivity were measured or calculated.

(211) <Mass Average Molecular Weight (Mw) and Number Average Molecular Weight (Mn)>

(212) The mass average molecular weight (Mw) and the number average molecular weight (Mn) were measured by means of gel permeation chromatography (GPC). As for the GPC, the measurement was performed by using connected two columns: TSKgel Super Multipore HZ-M, manufactured by Tosoh Corporation, using tetrahydrofuran as an eluent, and using a reflective index detector as a detector, thereby determining the mass average molecular weight (Mw) and the number average molecular weight (Mn) while using polystyrene as a standard sample.

(213) <40° C. Kinematic Viscosity, 100° C. Kinematic Viscosity, and Viscosity Index>

(214) The 40° C. kinematic viscosity and the 100° C. kinematic viscosity of the polyalkylene glycol-based compound were measured using a glass-made capillary viscometer in conformity with JIS K2283-2000, thereby calculating the viscosity index of the polyalkylene glycol-based compound.

(215) <Hydroxyl Value>

1. A lubricating oil composition comprising: a polyalkylene glycol-based compound represented by formula (1): ##STR00006## and at least one additive selected from an antioxidant, an oiliness improver, an oxygen scavenger, an extreme pressure agent, a copper deactivator, a rust inhibitor, an anti-foaming agent, and a viscosity index improver; wherein R.sup.1 represents a monovalent aliphatic hydrocarbon group having 1 to 32 carbon atoms, a monovalent aromatic hydrocarbon group having 6 to 42 ring carbon atoms, a monovalent acyl group having 2 to 32 carbon atoms, or a monovalent oxygen-containing hydrocarbon group having 2 to 32 carbon atoms; R.sup.2 represents a monovalent aliphatic hydrocarbon group having 1 to 32 carbon atoms, a monovalent aromatic hydrocarbon group having 6 to 42 ring carbon atoms, a monovalent acyl group having 2 to 32 carbon atoms, a monovalent oxygen-containing hydrocarbon group having 2 to 32 carbon atoms, or a hydrogen atom; R.sup.3 represents a divalent hydrocarbon group having 4 carbon atoms; R.sup.4 represents a divalent hydrocarbon group having 2 or 3 carbon atoms; m represents a number of 1 to 40; n represents a number of 0 to 20; and $m/(m+n) \geq 0.6$, and wherein a volume resistivity of the polyalkylene glycol-based compound represented by formula (1) at 25° C. is 0.0050 TΩ.Math.m or

more, and a 40° C. kinematic viscosity of the polyalkylene glycol-based compound represented by formula (1) is 30 to 400 mm.²/s.
