

Description**FIELD OF THE INVENTION**

This invention relates to 4-methyl-1-pentene polymer compositions and more particularly to 4-methyl-1-pentene polymer compositions excellent in rigidity at high temperatures, toughness at low temperatures, thermoformability such as vacuum formability, surface hardness impact strength and the like properties.

BACKGROUND OF THE INVENTION

It is well known that when polyolefin such as polyethylene or polypropylene is incorporated with inorganic fillers such as calcium carbonate, barium sulfate and talc, the thus incorporated polyolefin is improved in rigidity. On one hand, 4-methyl-1-pentene polymer which is a kind of polyolefin is excellent in heat resistance, chemical resistance, transparency, mold release characteristics, etc., as compared with other polyolefins, but from the viewpoint of molecular structure there was such a problem that because of its low melt tension, the polymer is difficult to thermoform. Further, there was another problem that the incorporation into 4-methyl-1-pentene polymer of such inorganic fillers as mentioned above results in improved rigidity at high temperatures, but the toughness at low temperatures of the polymer thus incorporated comes to decrease thereby.

DE-3307948-A1 discloses a composition for use as a diaphragm material for loudspeakers comprising a 4-methylpentene homopolymer a homopolymer of another α -olefin and a reinforcement of the key material, which composition is said to have improved film forming and thermal moulding characteristics.

The present invention seeks to solve the problems associated with the prior art and to provide 4-methyl-1-pentene polymer compositions having excellent thermoformability, for example, by means of, for example, vacuum forming or injection blow molding, and having excellent rigidity at high temperature, toughness at low temperature and impact strength.

The 4-methyl-1-pentene polymer composition of the present invention comprises 10 to 70 parts by weight of 4-methyl-1-pentene polymer (A), 10 to 70 parts by weight of propylene polymer (B), 10 to 70 parts by weight of inorganic filler (C) and 20-3 parts by weight of α -olefin copolymer (D) which is a random copolymer of at least 2 kinds of α -olefins, said random copolymer having a melt index of 0.1-50 g/10 min, a density of 0.850-0.900 g/cm³ and a crystallinity index as measured by X-ray of less than 40%.

Because of their components as defined above, the 4-methyl-1-pentene polymer compositions are excellent in thermoformability, rigidity at high temperatures, toughness at low temperatures, impact strength and the like properties.

DETAILED DESCRIPTION OF THE INVENTION

The components of the 4-methyl-1-pentene polymer compositions of the present invention are illustrated below in detail.

The 4-methyl-1-pentene polymer (A) used in the present invention is usually crystalline resin and includes a homopolymer of 4-methyl-1-pentene or copolymers of methyl-1-pentene and other α -olefins of 2-20 carbon atoms, for example, ethylene, propylene, 1-butene, 1-hexene, 1-octene, 1-decene, 1-tetradecene, 1-octadecene, etc., said copolymers usually being polymers consisting essentially of 4-methyl-1-pentene in an amount of at least 85 mol%. Usually, this 4-methyl-1-pentene polymer (A) has a melt flow rate (MFR) measured by ASTM D 1238 (temperature: 260° C, load: 5kg) of 0.05-500 g/10 min, though this rate may be varied according to the purpose for which the composition containing this polymer is used.

The propylene polymer (B) used in the present invention is usually a crystalline resin and includes a homopolymer of propylene or copolymers of propylene and less than 30% by weight of at least one other α -olefin having 2-20 carbon atoms, for example, ethylene, 1-butene, 1-hexene, 4-methyl-1-pentene, etc. or terpolymers of propylene, ethylene and less than 5% by weight of a polyene component, for example, 5-ethylidene-2-norbornene, 5-methylene-2-norbornene or 1,4-hexadiene. Usually, this propylene polymer (B) has a melting point of at least 130° C and a MFR of 0.1-80 g/10 min (ASTM D 1238 L).

The inorganic filler (C) used in the present invention includes inorganic substances such as calcium carbonate, magnesium carbonate, hydrous basic magnesium carbonate, kaolin clay, pyrophyllite, talc, calcined clay, diatomaceous earth, barium sulfate, etc. Of these inorganic substances, particularly preferred is talc.

The α -olefin copolymer (D) used in the present invention is a random copolymer of at least two kinds of α -olefins. This α -olefin copolymer has a crystallinity index as measured by X-ray of less than 40%.

preferably 1-30%, and is amorphous or low crystalline. This α -olefin copolymer (D) has a melt index or melt flow rate of 0.1-50 g/10 min, preferably 2-8 g/10 min (ASTM D 1238 E), a density of 0.850-0.900 g/cm³, preferably 0.875-0.895 g/cm³ and preferably has a melting point of 40-130 °C, more preferably 60-80 °C.

Concretely speaking, this α -olefin copolymer (D) is composed of at least two kinds of α -olefins selected from α -olefins of 2-20 carbon atoms such as ethylene, propylene, 1-butene, 1-dodecene, 1-tetradecene, 1-hexadecene, 1-octadecene, and 1-eicosene. Random copolymers of such α -olefins as mentioned above include concretely ethylene/propylene copolymers, ethylene/1-butene copolymers, ethylene/1-hexene copolymers, ethylene/1-octene copolymers, ethylene/1-decene copolymers, ethylene/1-hexadecene copolymers, propylene/1-butene copolymers, propylene/1-hexene copolymers, propylene/1-octene copolymers, propylene/1-decene copolymers, 1-butene/1-octene copolymers, 1-hexene/1-octene copolymers, ethylene/propylene/1-hexene copolymers, ethylene/1-butene/1-hexene copolymers, etc.

In the 4-methyl-1-pentene polymer compositions of the present invention, there are present the 4-methyl-1-pentene polymer (A) in an amount of 70-10, preferably 70-30 parts by weight, the propylene polymer (B) in an amount of 70-10, preferably 70-30 parts by weight, the inorganic filler (C) in an amount of 70-10, preferably 50-30 parts by weight, and the α -olefin copolymer (D) in an amount of 20-3, preferably 15-5 parts by weight, provided that the sum of (A), (B), (C) and (D) totals to 100 parts by weight.

In preparing the 4-methyl-1-pentene polymer composition of the present invention, the use of the 4-methyl-1-pentene polymer (A) in an amount of 70-10 parts by weight is selected, because the resulting composition becomes good in heat-resistant rigidity and impact strength. Similarly, the use of the propylene polymer (B) in an amount of 70-10 parts by weight is selected, because the resulting composition becomes tough and, is excellent in heat-resistant rigidity. The use of the inorganic filler (C) in an amount of 70-10 parts by weight is similarly selected, because the resulting composition becomes excellent in heat-resistance rigidity and, toughness.

In preparing the 4-methyl-1-pentene polymer composition of the present invention, the use of the α -olefin copolymer (D) in an amount of 20-3 parts by weight improves impact strength and maintains a good heat-resistant rigidity.

The 4-methyl-1-pentene polymer compositions of the present invention as illustrated above may be prepared by mixing, melt kneading, granulating or pulverizing a mixture comprising the 4-methyl-1-pentene polymer (A), propylene polymer (B), inorganic filler (C) and α -olefin copolymer (D) in the specific proportion as defined above by various known methods, for example, those using a V-blender, ribbon blender, Henschel mixer, tumbler blender, extruder, single-screw extruder, double-screw extruder, kneader or Banbury mixer.

The 4-methyl-1-pentene polymer compositions of the present invention may be incorporated with various additives commonly added to polyolefin such as weathering agents, heat stabilizers, rust preventives, slip agents, nucleating agents, pigments, dyes or surfactants, in such an amount as will not hinder the object of the present invention.

Because of their excellent rigidity at high temperature, toughness at low temperatures, thermoformability such as vacuum formability or surface hardness, the 4-methyl-1-pentene polymer compositions of the present invention are suitable for use in parts of household electrical appliances, tablewares for electron range cooking or industrial parts.

The present invention is illustrated below in detail with reference to examples, but it should be construed that the invention is in no way limited to those examples.

Comparative Example 1

Using a Henschel mixer, a mixture was prepared by mixing together 35 parts by weight of 4-methyl-1-pentene/1-decene crystalline copolymer (1-decene content: 3% by weight, melting point: 236 °C, MFR: 26g/10min), 35 parts by weight of propylene/ethylene crystalline block copolymer (ethylene content: 20% by weight, density: 0.91g/cm³ MFR: 0.5g/10min), 30 parts by weight of talc (a product of Matsumura Sangyo, sold under a trade name Crown P-2), 0.25 part by weight of tetrakis [methylene-3-(3,5-di-tert-butyl-4-hydroxyphenyl)propionate] methane (a product of Musashino-Geigy, sold under a trade name Iruganox 1010: hereinafter abbreviated to TPM), and 0.25 part by weight of stearic acid-monoglyceride (a product of Kao Soap, sold under a trade name Atomos #150; hereinafter abbreviated to SMG). The resulting mixture was melt kneaded with a 40 mm ϕ extruder to obtain composition-I.

With an injection molding machine (M-100 Type manufactured and sold by Meiki Seisakusho) operated at the molding temperature of 300 °C and the mold temperature of 60 °C, the composition-I thus obtained was then molded into a test specimen, 120 mm x 20 mm x 3 mm, for measuring heat deformation temperature (HDT), an ASTM TYPE IV specimen for tensile test, a test specimen for Izod impact test, and

into a square sheet, 100 mm x 100 mm x 2 mm, for surface hardness test, respectively. These test specimens thus prepared were evaluated by the following procedures.

Tensile test: Measurement was conducted in accordance with ASTM D638, provided that the ASTM TYPE IV specimen was used.

Heat deformation temperature: Measurement was conducted in accordance with ASTM D648, provided that the load applied was 66 psi.

Izod impact test: Measurement was conducted in accordance with ASTM D256.

Surface hardness: Measurement was conducted in accordance with JIS K-5401.

Separately, the composition-I was molded at a press temperature of 260 °C and a press pressure of 80 kg/cm² into a film of 300 μm in thickness.

The film thus obtained was co-biaxially oriented at 260 °C, whereupon a biaxially oriented film of a draw ratio of 3 x 3 was obtained.

The results obtained are shown in Table 1.

15 Comparative Examples 2-4

Comparative Example 1 was repeated except that the amounts of the components used therein were changed to those as shown in Table 1.

The results obtained are shown in Table 1.

20 Comparative Examples 5-8

Comparative Example 1 was prepared except that the amounts of the components used therein were changed to those as shown in Table 1.

The results obtained are shown in Table 1.

Table 1

	Comparative Example 1	Comparative Example 2	Comparative Example 3	Comparative Example 4	Comparative Example 5	Comparative Example 6	Comparative Example 7	Comparative Example 8
4-methyl-1-pentene/1-decene crystalline copolymer	35	55	25	20	50	10	90	5
Propylene/ethylene crystalline block copolymer	35	25	25	50	50	10	5	90
Talc	30	20	50	30	-	80	5	5
Stress at yield (kg/cm ²)	285	280	290	280	230	320	233	237
Elongation at yield (%)	8	10	7	10	15	Less than 1	10	20
Heat deformation temperature (°C)	136	135	145	130	85	180	95	92
Izod impact strength without notch (kg.cm/cm ²)	15	20	10	25	20	Less than 1	8	16
Surface hardness (Pencil hardness)	H	H	H	H	3B	2H	B	B
Result of biaxial orientation								
Minimum orientation temperature (°C)	240	240	240	235	230	-	235	195
Draw ratio	3 x 3	3 x 3	3 x 3	3 x 3	4 x 4	-	3 x 3	4 x 4
Evaluation	⊙	⊙	⊙	⊙	⊙	x' (not stretchable)	⊙ (irregular stretching)	⊙

Example 1

Using a Wenschel mixer, a mixture was prepared by mixing together 35 parts by weight of 4-methyl-1-pentene /1-decene crystalline copolymer (1-decene content: 3% by weight, melting point: 236 °C, MFR: 26g/10min), 30 parts by weight of propylene/ethylene crystalline block copolymer (ethylene content: 20% by weight, density: 0.91g/cc, MFR: 0.5g/10min), 30 parts by weight of talc (Crown P-2), 5 parts by weight of ethylene/1-butene random copolymer (ethylene content 80% by weight MFR (ASTM D 1238 E, 4g/10min), density 0.896g/cm³), 0.25 part by weight of tetrakis [methylene-3-(3,5-di-tert-butyl-4-hydroxyphenyl)-propionate] methane (TPM) and 0.25 part by weight of stearic acid monoglyceride (Atomos #150). The resulting mixture was melt kneaded with a 40 mmØ extruder to obtain composition-II.

With an injection molding machine (M-100 Type of Meiki Seisakusho) operated at the molding temperature of 300 °C and the mold temperature of 60 °C, the composition-II thus obtained was then molded into a test specimen, 120 mm x 20 mm x 3 mm, measuring heat deformation temperature (HDT), an ASTM TYPE IV specimen for tensile test, a test specimen for Izod impact test, and into a square sheet, 100 mm x 100 mm x 2 mm, for surface hardness test, respectively. These test specimens thus prepared were evaluated by the same procedures as in Example 1.

Separately, the composition-II was molded at a press temperature of 260 °C and a press pressure of 80 kg/cm² into a film of 400 µm in thickness.

The film thus obtained was co-biaxially oriented at 235 °C, whereupon a biaxially oriented film of a draw ratio of 3 x 3 was obtained.

The results obtained are shown in Table 2.

Examples 2-4

Example 1 was repeated except that the amount of the components used therein were changed to those as shown in Table 2.

The results obtained are shown in Table 2.

Comparative Examples 9-11

Example 1 was repeated except that the amounts of the components used therein were changed to those as shown in Table 2.

The results obtained are shown in Table 2.

For reference's sake, the results obtained in Comparative Example 1 are shown also in Table 2.

Table 2

	Example 1	Example 2	Example 3	Example 4	Comparative Example 5	Comparative Example 9	Comparative Example 10	Comparative Example 11
4-ethyl-1-pentene/1-decene crystalline copolymer	38	45	25	20	50	10	85	5
Propylene/ethylene crystalline block copolymer	30	25	25	20	50	5	5	85
Talc	30	20	45	50	-	80	5	5
Ethylene/1-butene random copolymer	5	10	5	10	-	5	5	5
Stress at yield (kg/cm ²)	280	275	285	290	230	315	230	235
Elongation at yield (%)	10	15	8	10	15	1	12	25
Heat deformation temperature (°C)	135	130	140	145	85	175	95	90
Izod impact strength without notch (kg.cm/cm ²)	20	35	15	30	20	1	10	18
Surface hardness (Pencil hardness)	H	H	H	H	3B	2H	B	B
Result of biaxial orientation								
Minimum orientation temperature (°C)	235	235	235	240	230	-	235	190
Draw ratio	3 x 3	3 x 3	3 x 3	3 x 3	4 x 4	-	-	4 x 4
Evaluation	⊙	⊙	⊙	⊙	⊙	x (not stretchable)	Δ - x (irregular stretching)	⊙



Claims

1. A composition comprising 10 to 70 parts by weight of 4-methyl-1-pentene polymer (A), 10 to 70 parts by weight of a propylene polymer (B), 10 to 70 parts by weight of an inorganic filler (C) and 3 to 20 parts by weight of an α -olefin copolymer (D) which is a random copolymer of at least two kinds of α -olefins, said random copolymer having a melt flow rate of 0.1 to 50 g/10 min, a density of 0.850 to 0.900 g/cm³ and a crystallinity index as measured by X-ray of less than 40%, the sum of (A), (B), (C) and (D) totalling 100 parts by weight.
2. A composition according to claim 1 wherein (A) is a homopolymer of 4-methyl-1-pentene or a copolymer of 4-methyl-1-pentene and at least one other α -olefin of 2 to 20 carbon atoms.
3. A composition according to claim 1 or 2 wherein (B) is a homopolymer of propylene or a copolymer of propylene and at least one other α -olefin of 2 to 20 carbon atoms.
4. A composition according to claim 1, 2, or 3 wherein (C) is calcium carbonate, magnesium carbonate, hydrous basic magnesium carbonate, kaolin clay, pyrophyllite, talc, calcined clay, diatomaceous earth or barium sulfate.
5. A composition according to claim 1, 2, 3, or 4 wherein (D) is an ethylene/propylene copolymer, ethylene/1-butene copolymer, ethylene/1-hexene copolymer, ethylene/1-octene copolymer, ethylene/1-hexadecene copolymer, propylene/1-butene copolymer, propylene/1-hexene copolymer, propylene/1-octene copolymer, propylene/1-decene copolymer, 1-butene/1-octene copolymer, ethylene/propylene/1-hexene copolymer or ethylene/1-butene/1-hexene copolymer.
6. A composition according to any preceding claim which contains 30 to 70 parts by weight of (A).
7. A composition according to any preceding claim which contains 30 to 70 parts by weight of (B).
8. A composition according to any preceding claim which contains 30 to 50 parts by weight of (C).
9. A composition according to any preceding claim which contain 5 to 15 parts by weight of (D).
10. Shaped articles obtained by thermoforming a composition as claimed in any preceding claim.

Patentansprüche

1. Masse, umfassend 10 bis 70 Gew.-Teile 4-Methyl-1-penten-polymer (A), 10 bis 70 Gew.-Teile eines Propylenpolymers (B), 10 bis 70 Gew.-Teile eines anorganischen Füllstoffs (C) und 3 bis 20 Gew.-Teile eines α -Olefin-Copolymers (D), das ein statistisches Copolymer aus mindestens zwei Arten von α -Olefinen ist, wobei das statistische Copolymer eine Schmelzflußrate von 0,1 bis 50 g/10 min, eine Dichte von 0,850 bis 0,900 g/cm³ und einen Kristallinitätsindex, gemessen durch Röntgenstrahlen, von weniger als 40% besitzt, und die Summe von (A), (B), (C) und (D) insgesamt 100 Gew.-Teile ausmacht.
2. Masse nach Anspruch 1, wobei (A) ein Homopolymer aus 4-Methyl-1-penten oder ein Copolymer aus 4-Methyl-1-penten und mindestens einem anderen α -Olefin mit 2 bis 20 Kohlenstoffatomen ist.
3. Masse nach Anspruch 1 oder 2, wobei (B) ein Homopolymer aus Propylen oder ein Copolymer aus Propylen und mindestens einem anderen α -Olefin mit 2 bis 20 Kohlenstoffatomen ist.
4. Masse nach Anspruch 1, 2 oder 3, wobei (C) Calciumcarbonat, Magnesiumcarbonat, wasserhaltiges basisches Magnesiumcarbonat, Kaolinton, Pyrophyllit, Talkum, calcinierter Ton, Diatomeenerde oder Bariumsulfat ist.
5. Masse nach Anspruch 1, 2, 3 oder 4, wobei (D) ein Ethylen/Propylen-Copolymer, Ethylen/1-Buten-Copolymer, Ethylen/1-Hexen-Copolymer, Ethylen/1-Octen-Copolymer, Ethylen/1-Hexadecan-Copolymer, Propylen/1-Buten-Copolymer, Propylen/1-Hexen-Copolymer, Propylen/1-Octen-Copolymer, Propylen/1-Decen-Copolymer, 1-Buten/1-Octen-Copolymer, Ethylen/Propylen/1-Hexen-Copolymer oder Ethylen/1-



Buten/1-Hexen-Copolymer ist.

6. Masse nach einem der vorangehenden Ansprüche, die 30 bis 70 Gew.-Teile (A) enthält.
- 5 7. Masse nach einem der vorangehenden Ansprüche, die 30 bis 70 Gew.-Teile (B) enthält.
8. Masse nach einem der vorangehenden Ansprüche, die 30 bis 50 Gew.-Teile (C) enthält.
9. Masse nach einem der vorangehenden Ansprüche, die 5 bis 15 Gew.-Teile (D) enthält.
- 10 10. Formkörper, erhalten durch thermisches Formen einer Masse nach einem der vorangehenden Ansprüche.

Revendications

- 15 1. Composition comprenant de 10 à 70 parties en poids d'un polymère (A) de 4-méthyl-1-pentène, de 10 à 70 parties en poids d'un polymère (B) de propylène, de 10 à 70 parties en poids d'une charge minérale (C) et de 3 à 20 parties en poids d'un copolymère (D) d' α -oléfines qui est un copolymère statistique d'au moins deux types d' α -oléfines, ce copolymère statistique présentant un indice de fluidité à chaud de 0,1 à 50 g/10 min, une masse volumique de 0,850 à 0,900 g/cm³ et un taux de cristallinité, mesuré par diffraction des rayons X, inférieur à 40 %, la somme des constituants (A), (B), (C) et (D) représentant 100 parties en poids.
- 20 2. Composition conforme à la revendication 1, dans laquelle le polymère (A) est un homopolymère du 4-méthyl-1-pentène ou un copolymère du 4-méthyl-1-pentène et d'au moins une autre α -oléfine comportant de 2 à 20 atomes de carbone.
- 25 3. Composition conforme à la revendication 1 ou 2, dans laquelle le polymère (B) est un homopolymère du propylène ou un copolymère du propylène et d'au moins une autre α -oléfine comportant de 2 à 20 atomes de carbone.
- 30 4. Composition conforme à la revendication 1, 2 ou 3, dans laquelle la charge minérale (C) est du carbonate de calcium, du carbonate de magnésium, du carbonate de magnésium basique hydraté, du kaolin, de la pyrophyllite, du talc, une argile calcinée, de la terre à diatomées ou du sulfate de baryum.
- 35 5. Composition conforme à la revendication 1, 2, 3 ou 4, dans laquelle le copolymère (D) est un copolymère d'éthylène et de propylène, un copolymère d'éthylène et de 1-butène, un copolymère d'éthylène et de 1-hexène, un copolymère d'éthylène et de 1-octène, un copolymère d'éthylène et de 1-hexadécène, un copolymère de propylène et de 1-butène, un copolymère de propylène et de 1-hexène, un copolymère de propylène et de 1-octène, un copolymère de propylène et de 1-décène, un copolymère de 1-butène et de 1-octène, un copolymère d'éthylène, de propylène et de 1-hexène, ou un copolymère d'éthylène, de 1-butène et de 1-hexène.
- 40 6. Composition conforme à l'une des revendications précédentes, qui contient de 30 à 70 parties en poids de polymère (A).
- 45 7. Composition conforme à l'une des revendications précédentes, qui contient de 30 à 70 parties en poids de polymère (B).
- 50 8. Composition conforme à l'une des revendications précédentes, qui contient de 30 à 50 parties en poids de charge minérale (C).
- 55 9. Composition conforme à l'une des revendications précédentes, qui contient de 5 à 15 parties en poids de copolymère (D).
10. Objets façonnés, obtenus par thermoformage d'une composition conforme à l'une des revendications précédentes.