

Description

The present invention relates to crosslinkable thermoplastic polymeric compositions which can be thermoformed, for example by extrusion, and subsequently crosslinked to give products having improved properties.

More particularly the present invention relates to thermoplastic polymeric compositions which are inherently adapted for crosslinking by thermal or chemical treatment subsequent to being thermoformed into finished or partially finished products. Examples of known compositions of this type are those comprising silane modified polymers which are adapted to be crosslinked by treatment with water, generally in the presence of a silanol condensation catalyst, subsequent to thermoforming of the silane modified polymer.

FR-A-2 226 410 relates to a peroxide-curable composition comprising an ethylene-propylene rubber and a fluorine containing ethylene polymer. The composition when cured is said to have excellent heat and oil resistance.

US-A-3 125 547 relates to a melt extrudable thermoplastic polymer of an olefin, having the formula $\text{RCH}=\text{CH}_2$, where R is a radical selected from the group consisting of hydrogen and alkyl radicals having from 1 to 8 carbon atoms, containing from 0.005 to 2 weight percent of a fluorocarbon polymer.

Thermoplastic silane-modified polymers are organic polymers containing hydrolysable groups (attached to the silicon) that can be crosslinked by hydrolysis, e.g. by the action of water, preferably in the presence of a silanol condensation catalyst. Crosslinkable silane-modified copolymers can be manufactured, for example, by copolymerising unsaturated organic monomers with unsaturated silane compounds containing hydrolysable groups. Examples of copolymers of this type are described in GB-A-2028831 and GB-A-2039513 which disclose the preparation of crosslinkable copolymers of ethylene and an ethylenically unsaturated silane compound by copolymerising the monomers at relatively high temperatures and pressures in the presence of a radical polymerisation initiator. Further examples of copolymers of this type are described in GB-A-1415194 which discloses the preparation of crosslinkable copolymer by contacting ethylene, optionally with other olefinically unsaturated comonomer, and a terminally unsaturated silane compound with certain defined Ziegler catalysts under polymerisation conditions which preferably employ relatively low temperatures and pressures.

Another known method for making such crosslinkable silane-modified organic polymers comprises graft-polymerising an ethylenically unsaturated silane compound on to an organic polymer preferably in the presence of a free radical initiator. Examples of this method are disclosed in GB-A-1357549, GB-A-1234034 and GB-A-1286460. The graft-polymerisation method forms the basis of the well-known "SIOPLAS" (RTM) commercial technique for making crosslinkable thermoplastic polymers. In the well known single-step "MONOSIL" (RTM) process for making crosslinkable compositions of this type an ethylenically unsaturated silane compound is graft-polymerised on to an organic polymer in the presence of a free radical initiator and a silanol condensation catalyst. In this process the grafting reaction is performed simultaneously with the fabrication of the polymeric article, for example, by feeding the organic polymer, the unsaturated silane compound, the initiator and the silanol condensation catalyst (optionally with conventional additives) to an extruder wherein the grafting reaction occurs and a crosslinkable product is extruded directly.

Crosslinkable organic polymers can be fabricated to form a large variety of useful articles by conventional thermoforming techniques, for example, extrusion, injection moulding, blow-moulding and film-blowing processes.

A problem encountered with crosslinkable thermoplastic compositions of the aforescribed types is that, during thermoforming of such compositions into useful articles, premature crosslinking can take place, for example, due to thermal or mechanical effects or to the presence of moisture or other impurities. Premature crosslinking can also occur in "dead" spaces in extruders or dies, e.g. spaces where eddies form in the melt flow pattern or where the melt can stagnate. Premature crosslinking in the thermoplastic polymeric melt causes imperfections in the final thermoformed article, for example, visible surface defects such as "fish-eyes" and streaking in extruded products.

It is an object of the present invention to provide an improved crosslinkable thermoplastic polymeric composition. It is a further object of the invention to provide a crosslinkable thermoplastic polymeric composition which can be extruded to provide products having improved surface finish.

Accordingly the present invention provides a thermoformable and crosslinkable polymeric composition comprising (a) a crosslinkable thermoplastic component which is a silane-modified organic polymer having hydrolysable groups attached to the silicon atoms thereof, (b) 0.002 to 2.0 weight %, based on the weight of the crosslinkable component of a fluorocarbon polymer and (c) a silanol condensation catalyst.

The present invention further provides a process for making a crosslinkable thermoformable composi-

tion comprising blending together components comprising (a) a crosslinkable thermoplastic component which is a silane-modified organic polymer having hydrolysable groups attached to the silicon atoms thereof, (b) 0.002 to 2.0 weight %, based on the weight of the crosslinkable component of a fluorocarbon polymer and (c) a silanol condensation catalyst.

5 The present invention further provides a process for the production of a crosslinkable thermoplastic article comprising thermoforming the article from a thermoformable and crosslinkable polymeric composition as described above.

The fluorocarbon polymer employed in the composition of the present invention is suitably capable of forming a melt extrudable blend with the crosslinkable component of the composition. Preferably the 10 fluorocarbon polymer melts or softens at a temperature in the range 100-300 °C, more preferably in the range 150 to 250 °C. Fluorocarbon homo- and copolymers especially suitable for use in the composition of the present invention are disclosed in United States patent No. 3125547. Such homo- and copolymers have an atomic ratio of fluorine to carbon of at least 1:2. They comprise, for example, homo and copolymers of one or more of vinylidene fluoride, vinyl fluoride, chlorotrifluoro ethylene, tetrafluoroethylene and hexa- 15 afluoropropylene. Copolymers of these fluorinated olefins with unfluorinated olefins can be employed in the composition of the present invention provided the copolymer contains an atomic ratio of fluorine to carbon of at least 1:2. Specific examples of fluorocarbon polymers which can be suitably employed in the present invention are telomers of tetrafluoroethylene, telomers of chlorotrifluoroethylene and fluorocarbon elastomers, for example copolymers of hexafluoropropylene and vinylidene fluoride.

20 The quantity of fluorocarbon polymer employed in the composition of the present invention is in the range 0.002 to 2.0 weight %, preferably 0.03 to 0.4 weight % based on the weight of the crosslinkable component. It is particularly preferred to use a quantity of fluorocarbon polymer in the range 100 to 450 ppm based on the weight of the crosslinkable component.

The fluorocarbon polymer can be incorporated into the crosslinkable composition of the present 25 invention using any of the blending or mixing techniques conventionally employed in the art. For example, the fluorocarbon polymer can be incorporated directly into crosslinkable silane modified polymer. Alternatively, the fluorocarbon polymer can be blended with thermoplastic polymer, for example low density polyethylene, to form a masterbatch concentrate which can then be blended or compounded into the crosslinkable component or into a precursor of the crosslinkable component.

30 The silane modified polymer is crosslinkable by the action of water or other chemical agency.

The silane modified polymer can be, for example, a copolymer prepared by copolymerising an olefinically unsaturated compound with an unsaturated silane compound having hydrolysable groups, a graft copolymer prepared by grafting an unsaturated silane compound having hydrolysable silane groups on to an organic polymer or a copolymer modified to contain hydrolysable silane groups by a transesterification 35 process (see for example EP4752). Preferred silane modified polymers are copolymers prepared by copolymerising ethylene, optionally together with one or more alpha-olefins, vinyl esters, alkyl(meth)acrylates, unsaturated nitriles or unsaturated ethers (e.g. vinyl ethers), with an unsaturated silane compound in the presence of a free radical initiator. Also preferred are graft copolymers prepared by grafting an unsaturated silane compound on to polyethylene or on to a copolymer of ethylene with one or more alpha- 40 olefins, vinyl esters, alkyl(meth)acrylates, unsaturated nitriles or unsaturated ethers (e.g. vinyl ethers), by heating the polyethylene, or copolymer of ethylene, with the unsaturated silane compound in the presence of a free radical initiator, for example, an organic peroxide. The polyethylene or copolymer of ethylene can comprise, for example, low density polyethylene, low density ethylene hydrocarbon copolymers (e.g. LLDPE), high density polyethylene, ethylene/ethyl acrylate copolymer, ethylene/vinyl acetate copolymer or 45 ethylene propylene rubber (EPR).

The silane modified polymer can be formed "in situ" in the presence of the other component(s) of the composition of the present invention, for example, by grafting an unsaturated silane compound on to polyethylene or on to a copolymer of ethylene of the type described above in the presence of a free radical polymerisation initiator.

50 The silane compound copolymerised with the ethylene, or graft copolymerised with the polyethylene or copolymer of ethylene, is preferably a compound having the general formula $R^1SiR^2_nY_{3-n}$ wherein R^1 represents an ethylenically unsaturated hydrocarbyl or hydrocarbyloxy group; R^2 represents an aliphatic saturated hydrocarbyl group; Y represents a hydrolysable organic group; and n represents zero, 1 or 2. Preferred unsaturated silane compounds for use in making the copolymer or graft copolymer are vinyl 55 trimethoxy silane, vinyl triethoxy silane and vinyl triacetoxysilane.

Silane modified polymers particularly preferred for use as the crosslinkable component in the composition of the present invention are copolymers prepared by copolymerising ethylene, optionally together with up to 40% by weight (based on ethylene) of additional monomer selected from one or more vinyl esters or

(meth)acrylates, and an unsaturated silane compound selected from vinyl trialkoxy silanes and vinyl triacetoxysilane, under a pressure of from 500 to 4000 bars and at a temperature in the range 150 to 400 °C in the presence of a free radical polymerisation initiator.

The silane modified polymer is preferably a copolymer or a graft copolymer containing 0.1-10 wt %, most preferably 0.5 to 5 wt % of copolymerised units of the silane compound.

For further details of silane modified polymers suitable for use in the present invention reference may be made to GB-A-2028831, GB-A-2039513, GB-A-1357549, GB-A-1415194, GB-A-1286480, GB-A-1234034 US-A-3225018 and EP4752.

In the process of the present invention the thermoforming can be, for example, extrusion, extrusion coating, blow moulding, injection moulding, film-blowing or slot-casting. The process is particularly suitable for the production of extruded or extrusion-coated articles, for example pipe, tubing and insulated wire and cable.

In a particularly preferred embodiment of the present invention a masterbatch is made up comprising a masterbatch base polymer, for example LDPE, LLDPE or ethylene alkyl acrylate copolymer, together with the fluorocarbon polymer and optional additives, for example, antioxidants, fillers, metal deactivators (e.g. salicylaldehyde oximes), lubricants, water-tree inhibitors, foaming agent, flame retardants and pigments.

The present invention also includes a masterbatch for adding to the crosslinkable silane-modified organic polymer, said masterbatch comprising the fluorocarbon polymer and the silanol condensation catalyst optionally with a thermoplastic base polymer. The hydrocarbyl groups in the dihydrocarbyl tin (IV) carboxylate are preferably alkyl groups containing 1-12 carbon atoms, for example propyl, butyl, hexyl, octyl or decyl. The carboxylate groups in the tin compound can be provided by, for example, aliphatic or aromatic mono- or dicarboxylic acid. Preferred dihydrocarbyl tin compounds include dibutyl tin dilaurate, dibutyl tin dipalmitate, dibutyl tin distearate, dioctyl tin dilaurate and dibutyl tin maleate. Dihydrocarbyl tin (IV) carboxylates wherein the carboxylate function is provided by a dicarboxylic acid are particularly preferred (e.g. dibutyl tin maleate). Generally speaking, the quantity of dihydrocarbyl tin carboxylate employed should be sufficient to give a concentration in the crosslinkable composition in the range 0.001 to 3.0 moles, preferably 0.003 to 0.05 moles per mole of silyl units in the crosslinkable component. The quantity of the tin compound in the masterbatch will generally lie in the range 0.1 to 70 weight %, preferably 0.3 to 20 weight % based on the weight of the masterbatch. The quantity of fluorocarbon polymer in the masterbatch can be, for example, 0.01 to 95 weight %, preferably 0.1 to 20 weight % based on the weight of the masterbatch. The optional thermoplastic base polymer can be any suitable organic polymer which is compatible with the crosslinkable component and can be melt blended therewith. Examples of such base polymers are conventional low density polyethylene, ethylene/vinyl acetate copolymer, ethylene/ethyl acrylate copolymer, linear low density polyethylene and high density PE.

The composition and process of the present invention can be used to make crosslinkable thermoformed articles having for example improved surface characteristics and reduced incidence of gel defects.

The invention is illustrated with reference to the following Examples.

Examples 1-3 and Comparative Test A

In Examples 1, 2 and 3 Viton A fluoroelastomer (DuPont) was incorporated into a masterbatch in varying proportions. In the comparative test no Viton A was incorporated into the masterbatch. The exact formulations of the masterbatch are shown in Table 1. The ingredients shown in Table 1 were blended together to make a masterbatch using a Werner and Pfleiderer ZSK 30 twin screw extruder at a screw speed of 200 revolutions per minute. The total quantity of composition prepared in each Example and in the Test was 10kg per run. The extruder barrel temperature was about 140 °C close to the hopper zone, increasing to about 190 °C at the extruder head. Extrusion was carried out to give a strand which was chopped to give the pelleted masterbatch.

The pelleted masterbatch (5 parts by weight) was dry blended with 95 parts by weight of a silyl modified polymer prepared by copolymerising ethylene with vinyl trimethoxy silane under high temperature and high pressure conditions using a free radical initiator. The silyl modified polymer (EVTMS) contained 1.8 weight % of copolymerised vinyl trimethoxy silane, had a melt index (190 °C, 2.16 Kg load) of 0.7 and a density of 923 Kg/m³.

The concentrations of fluoroelastomer in Examples 1, 2 and 3 were thus 500, 250 and 100 ppm respectively.

The dry blend was fed into the hopper of a Francis-Shaw Extruder equipped with 2½ inch screw having L:D of 23:1 and a die diameter of 2.2 mm through which a copper wire of 0.6 mm diameter passed. The die temperature was 200 °C and the screw speed was 50 RPM. The coated wire was taken off at 200

metres/minute producing a cable of external diameter 2.05 mm. It was observed that less die drool formed in Examples 1, 2 and 3 than in Comparative Test A. The extruder cable was cured by immersion in a water bath thermostatted at 80° C.

Visual examination of the cables produced using the composition of the present invention (Examples 1, 2 and 3) showed the surface of the cables to be smooth and glossy (1 and 2 better than 3) even after 1 hour's extrusion time. On the other hand, cable produced from the composition of Comparative Test A exhibited a matt finish and a more grainy surface.

All samples showed essentially identical tensile and thermo mechanical properties indicating that the fluoroelastomer does not impair the crosslinking reaction.

The incorporation of the fluoroelastomer provides a significant reduction in extruder power.

TABLE 1

**Masterbatch
Formulations**

	MB1	MB2	MB3	MBA
Viton A	1.00	0.50	0.20	-
Dibutyltin maleate	0.75	0.75	0.75	0.75
Slip Agent	1.5	1.5	1.5	1.5
Antioxidant	6.0	6.0	6.0	6.0
LDPE	90.75	91.25	91.55	91.75

		Extruder Amps	Head Pressure	Surface Finish
Example 1	EVTMS + MB1(5%)	23	140	Excellent glossy
Example 2	" + MB2(5%)	24	150	Excellent glossy
Example 3	" + MB3(5%)	26	155	Good
Test A	" + MBA(5%)	26	158	Good Matt

Claims

1. A thermoformable and crosslinkable polymeric composition comprising (a) a crosslinkable thermoplastic component which is a silane-modified organic polymer having hydrolysable groups attached to the silicon atoms thereof, (b) 0.002 to 2.0 weight %, based on the weight of the crosslinkable component of a fluorocarbon polymer and (c) a silanol condensation catalyst.
2. A composition as claimed in Claim 1 wherein the fluorocarbon polymer melts or softens at a temperature in the range 100 to 300° C.
3. A composition as claimed in Claim 1 or 2 wherein the fluorocarbon polymer has an atomic ratio of fluorine to carbon of at least 1:2.
4. A composition as claimed in any of Claims 1 to 3 wherein the fluorocarbon polymer is a homo- or copolymer of one or more of vinylidene fluoride, vinyl fluoride, chlorotrifluoro ethylene, tetrafluoroethylene and hexafluoropropylene.
5. A composition as claimed in any of Claims 1 to 4 wherein the fluorocarbon polymer is a copolymer of hexafluoropropylene and vinylidene fluoride.



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3. A composition as claimed in Claim 1 or 2 wherein the fluorocarbon polymer has an atomic ratio of fluorine to carbon of at least 1:2.
4. A composition as claimed in any of Claims 1 to 3 wherein the fluorocarbon polymer is a homo- or copolymer of one or more of vinylidene fluoride, vinyl fluoride, chlorotrifluoro ethylene, tetrafluoroethylene and hexafluoropropylene.
5. A composition as claimed in any of Claims 1 to 4 wherein the fluorocarbon polymer is a copolymer of hexafluoropropylene and vinylidene fluoride.



6. A composition as claimed in any of the preceding claims wherein the quantity of fluorocarbon employed is in the range 0.03 to 0.4 weight % based on the weight of the crosslinkable component.
7. A composition as claimed in any one of Claims 1 to 5 wherein the quantity of the fluorocarbon polymer is in the range 100 to 450 ppm based on the weight of the crosslinkable component.
8. A composition as claimed in any one of claims 1 to 7 wherein the silane modified organic polymer is a copolymer prepared by copolymerising an olefinically unsaturated compound with an unsaturated silane compound having hydrolysable groups.
9. A composition as claimed in any one of claims 1 to 7 wherein the silane modified organic polymer is prepared by grafting an unsaturated silane compound having hydrolysable silane groups on to an organic polymer or copolymer selected from polyethylene or a copolymer of ethylene with one or more alpha olefins, vinyl esters, alkyl (meth)acrylates, unsaturated nitriles or unsaturated ethers.
10. A composition as claimed in claim 8 or 9 wherein the unsaturated silane compound is selected from vinyl trimethoxy silane, vinyl triethoxy silane and vinyl triacetoxysilane.
11. A composition as claimed in claim 8 or 9 wherein the copolymer or graft copolymer contains 0.1 to 10 weight % of copolymerised units of silane compound.
12. An extruded or extrusion-coated article comprising the product of extruding and subsequently thermally or chemically crosslinking a composition as claimed in any one of Claims 1 to 10.
13. A process for making a crosslinkable thermoformable composition comprising blending together components comprising (a) crosslinkable thermoplastic component which is a silane-modified organic polymer having hydrolysable groups attached to the silicon atoms thereof, (b) 0.002 to 2.0 weight %, based on the weight of the crosslinkable component of a fluorocarbon polymer and (c) a silanol condensation catalyst.

Revendications

1. Composition polymère thermoformable et réticulable comprenant (a) un composant thermoplastique réticulable, qui est un polymère organique modifié par un silane ayant des groupes hydrolysables fixés à l'atome de silicium de celui-ci, (b) 0,002 à 2,0 % en poids, sur la base du poids du composant réticulable d'un polymère fluorocarboné et (c) un catalyseur de condensation à base de silanol.
2. Composition suivant la revendication 1, dans laquelle le polymère fluorocarboné fond ou se ramollit à une température comprise entre 100 et 300 °C.
3. Composition suivant la revendication 1 ou 2, dans laquelle le polymère fluorocarboné a un rapport atomique fluor-carbone d'au moins 1:2.
4. Composition suivant l'une des revendications 1 à 3, dans laquelle le polymère fluorocarboné est un homopolymère ou copolymère d'un ou plusieurs composé(s) le fluorure de vinylidène, le fluorure de vinyle, le chlorotrifluoroéthylène, le tétrafluoroéthylène et l'hexafluoropropylène.
5. Composition suivant l'une des revendications 1 à 4, dans laquelle le polymère fluorocarboné est un copolymère d'hexafluoropropylène et de fluorure de vinylidène.
6. Composition suivant l'une ou l'autre des revendications précédentes, dans laquelle la quantité de polymère fluorocarboné utilisée est de l'ordre de 0,03 à 0,4 % en poids, sur la base du poids du composant réticulable.
7. Composition suivant l'une des revendications 1 à 5, dans laquelle la quantité de polymère fluorocarboné est comprise entre 100 et 450 ppm, sur la base du poids du composant réticulable.
8. Composition suivant l'une ou l'autre des revendications 1 à 7, dans laquelle le polymère organique



modifié par un silane est un copolymère préparé par copolymérisation d'un composé oléfiniquement non-saturé avec un composé de silane non-saturé ayant des groupes hydrolysables.

- 5 9. Composition suivant l'une des revendications 1 à 7, dans laquelle le polymère organique modifié par un silane est préparé par greffage d'un composé de silane non-saturé ayant des groupes silane hydrolysables sur un polymère ou copolymère organique sélectionné parmi le polyéthylène ou un copolymère d'éthylène avec un ou plusieurs des composés suivants : alpha-oléfines, esters vinyliques, (méth) acrylates d'alcoyle, nitriles non-saturés ou éthers non-saturés.
- 10 10. Composition suivant la revendication 8 ou 9, dans laquelle le composé de silane non-saturé est sélectionné parmi le triméthoxy-silane de vinyle, le triéthoxy-silane de vinyle et le triacétoxy-silane de vinyle.
- 15 11. Composition suivant la revendication 8 ou 9, dans laquelle le copolymère ou copolymère greffé contient de 0,1 à 10% en poids d'unités de silane copolymérisées.
- 20 12. Article extrudé ou revêtu par extrusion comprenant le produit de l'extrusion et de la réticulation thermique ou chimique ultérieure d'une composition suivant l'une des revendications 1 à 10.
- 25 13. Procédé de préparation d'une composition thermoformable réticulable comprenant le mélange de composés comprenant (a) un composé thermoplastique réticulable, qui est un polymère organique modifié par un silane ayant des groupes hydrolysables fixés à l'atome de silicium de celui-ci, (b) 0,002 à 2,0 % en poids, sur base du poids du composant réticulable, d'un polymère fluorocarboné et (c) un catalyseur de condensation à base de silanol.

Patentansprüche

- 30 1. Warmformbare und vernetzbare polymere Zusammensetzung, umfassend
 - (a) eine vernetzbare thermoplastische Komponente, welche ein Silan-modifiziertes organisches Polymer ist, das an seine Siliziumatome gebundene hydrolysierbare Gruppen besitzt,
 - (b) 0,002 bis 2,0 Gew.-%, bezogen auf das Gewicht der vernetzbaren Komponente, eines Fluorkohlenstoffpolymers und
 - (c) einen Silanolkondensationskatalysator.
- 35 2. Zusammensetzung nach Anspruch 1, worin das Fluorkohlenstoffpolymer bei einer Temperatur im Bereich von 100 bis 300 °C schmilzt oder erweicht.
- 40 3. Zusammensetzung nach Anspruch 1 oder 2, worin das Fluorkohlenstoffpolymer ein Atomverhältnis von Fluor zu Kohlenstoff von mindestens 1:2 besitzt.
- 45 4. Zusammensetzung nach einem der Ansprüche 1 bis 3, worin das Fluorkohlenstoffpolymer ein Homo- oder Copolymer aus einer oder mehreren der Verbindungen Vinylidenfluorid, Vinylfluorid, Chlortrifluorethylen, Tetrafluorethylen und Hexafluorpropylen ist.
- 50 5. Zusammensetzung nach einem der Ansprüche 1 bis 4, worin das Fluorkohlenstoffpolymer ein Copolymer aus Hexafluorpropylen und Vinylidenfluorid ist.
6. Zusammensetzung nach einem der vorstehenden Ansprüche, worin die angewandte Menge an Fluorkohlenstoffpolymer im Bereich von 0,03 bis 0,4 Gew.-%, bezogen auf das Gewicht der vernetzbaren Komponente, liegt.
7. Zusammensetzung nach einem der Ansprüche 1 bis 5, worin die Menge des Fluorkohlenstoffpolymers im Bereich von 100 bis 450 ppm, bezogen auf das Gewicht der vernetzbaren Komponente, beträgt.
- 55 8. Zusammensetzung nach einem der Ansprüche 1 bis 7, worin das Silan-modifizierte organische Polymer ein Copolymer ist, welches durch Copolymerisation einer olefinisch ungesättigten Verbindung mit einer ungesättigten Silanverbindung, welche hydrolysierbare Gruppen besitzt, hergestellt wird.



9. Zusammensetzung nach einem der Ansprüche 1 bis 7, worin das Silan-modifizierte organische Polymer durch Pfropfen einer ungesättigten Silanverbindung, welche hydrolysierbare Silangruppen enthält, auf ein organisches Polymer oder Copolymer hergestellt wird, welches unter Polyethylen oder einem Copolymer aus Ethylen und einem oder mehreren alpha-Olefinen, Vinylestern, Alkyl(meth)acrylaten, ungesättigten Nitrilen oder ungesättigten Estern ausgewählt ist.
10. Zusammensetzung nach Anspruch 8 oder 9, worin die ungesättigte Silanverbindung von Vinyltrimethoxysilan, Vinyltriethoxysilan und Vinyltriacetoxysilan ausgewählt ist.
11. Zusammensetzung nach Anspruch 8 oder 9, worin das Copolymer oder das Pfropfcopolymer 0,1 bis 10 Gew.-% an copolymerisierten Einheiten der Silanverbindung enthält.
12. Extrudierter oder extrusionsbeschichteter Gegenstand, bestehend aus dem Produkt des Extrudierens und des darauffolgenden thermischen oder chemischen Vernetzens einer Zusammensetzung nach einem der Ansprüche 1 bis 10.
13. Verfahren zur Herstellung einer vernetzbaren, warmformbaren Zusammensetzung, umfassend das Zusammenmischen von Komponenten, welche sich zusammensetzen aus
- (a) einer vernetzbaren thermoplastischen Komponente, welche ein Silan-modifiziertes organisches Polymer ist, das an seine Siliziumatome gebundene hydrolysierbare Gruppen besitzt,
 - (b) 0,002 bis 2,0 Gew.-%, bezogen auf das Gewicht der vernetzbaren Komponente, eines Fluorkohlenstoffpolymers und
 - (c) einem Silanolkondensationskatalysator.