

PROCESS FOR THE PREPARATION OF A FLEXIBLE THERMOPLASTIC
COMPOSITE FILAMENT CONTAINING CONTINUOUS FIBRES.

The present invention relates to a process for the
preparation of a flexible thermoplastic composite
5 filament containing continuous fibres.

More specifically, the present invention relates
to a process for the preparation of a flexible thermo-
plastic composite filament consisting of bundles of
continuous fibres of any kind, either mineral, organic
10 or metallic, impregnated with a thermoplastic polyole-
finic resin in powder form and/or in fibres and covered
with an external flexible thermoplastic sheath which is
also polyolefinic.

In many applications, formerly dominated by the
15 use of metallic materials, composite materials have
been proposed basically consisting of a polymeric
matrix and an inorganic reinforcement, mainly glass or
carbon fibres. In conditions which require particularly
high performances, from the point of view of mechanical

resistance, composite materials are proposed with a long or, above all, continuous fibre.

The first type of composite material with a long or continuous fibre, developed as an alternative to metallic materials, was characterized in that it had a matrix consisting of thermosetting polymers, of which epoxy resins, unsaturated polyester resins and bismaleimides are typical examples.

End products made of these materials are obtained by impregnating and/or pre-impregnating the fibres with the monomer and/or a prepolymer and subsequently forming the material with technologies such as manual stratification, winding, pultrusion and moulding in a press or autoclave.

In each case the fibres are passed through solutions of the thermosetting resin with consequent problems of environment and process costs due to the necessity of arranging the reuse and/or storage of the solvents used.

To overcome this type of inconvenience, composite materials have quite recently been introduced which use thermoplastic polymers as matrix. In this case the relative transformation technology involves, if the composite is a long or continuous fibre, the preliminary formation of a semifinished product.

An example of the preparation of a semifinished product of this kind is described in U.S. patent 3.742.106, which discloses the formation of a composite material by the impregnation of a continuous reinforcing filament in a thermoplastic melt.

Another example of semifinished product for composites with a thermoplastic matrix is described in Journal of Applied Polymer Science - Applied Polymer Symposium, vol. 47, page 501, 1991, which discloses the preparation of a continuous filament by the spinning of a mix consisting of reinforcing and thermoplastic fibres.

U.S. patent 4.614.678 describes the formation of a composite by the impregnation of a bundle of reinforcement fibres with thermoplastic powders and the covering of the bundle with a sheath which is also thermoplastic.

More specifically, the process described in this patent comprises:

- 20 - unrolling a continuous filament, consisting of a bundle of fibres, for example glass or carbon fibres, from a bobbin;
- feeding the filament to a closed chamber, inside which a thermoplastic polymer in powder form is maintained in a fluidized state by means of gas;

- opening, inside the chamber, the bundle of filaments basically into single fibres;
- drawing the opened filament through the fluidized polymer for a time which is sufficient for particles of polymer to be deposited around the single fibres;
- reclosing the filament and enclosing it with a protective flexible sheath of thermoplastic polymer preferably of the same kind as the powder polymer.

After obtaining the semifinished products, units are produced with consolidation techniques which can involve heating and pressure application, as described for example in various items of the Encyclopedia of Composites, prepared by S. M. Lee, published by VCH Publishers, New York, 1990-1991, or the pultrusion of the semifinished products themselves.

The use of thermoplastic composites undoubtedly produces improvements from an environmental point of view as no solvents are involved in the production cycles. Problems do arise however in the transformation into end products, above all when the composite material consists of a continuous glass multifibre filament impregnated and covered with a polyolefin such as polyethylene. In this case, there are problems of

compatibilization between the polyolefinic matrix and reinforcement.

It is known, in fact, that to improve adhesion between a reinforcement fibre, particularly glass
5 fibre, and a polymeric matrix, it is necessary to modify the surface of the fibre with substances, for example of the silanic or aminic type, which, by interacting between the reactive sites present on the fibre and reactive groups present in the polymer,
10 produce strong bonds which considerably improve the compatibility between matrix and reinforcement. This surface treatment of the fibre, known as chemical finishing, is amply described in literature, for example in "Reinforced Plastics and Elastomers. Recent
15 Developments", M.W.Ranney, Noyes Data Corporation, 1977, 4.

In the case of the polyolefins, however, owing to the substantial absence of polar or reactive groups in the polymeric chain, the sizing treatment of the fibre
20 alone is not sufficient to guarantee the necessary adhesion between matrix and reinforcement. To overcome this inconvenience a proposal has been made to modify the structure of the olefinic polymer by grafting onto the polymeric chain monomers having polar groups such
25 as maleic acid or anhydride, acrylic or methacrylic

acids, etc. The modification operation however has proved to be valid when the polyolefins are mixed in an extruder with short fibres. In this case, in fact, by carrying out the functionalization of the polymer
5 directly in an extruder or by adding a prefunctionalized polymer to the mixture to be treated, it is possible to have a dynamic contact, guaranteed by the extruder itself, between fibre and polymeric matrix which favours the chemical bond between the size of the
10 fibre and the polar groups grafted to the polyolefin.

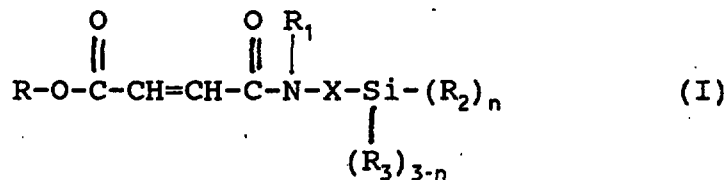
On the contrary, in the case of composites with a continuous fibre the functionalization of the polyolefin has proved to be ineffective as subsequent transformation technologies of the semifinished products do
15 not permit the above dynamic contact necessary for the compatibilization between reinforcement and polymeric matrix. On the basis of this, the use of polyolefins, and in particular polyethylene, for the preparation of thermoplastic composites reinforced with continuous
20 glass fibres has had little success in the past.

The Applicant has now found a process for preparing a flexible thermoplastic composite filament with a polyolefinic base and containing continuous fibres which, being capable to favour and improve the adhesion
25 between fibrous reinforcement and polymeric matrix

during the moulding phase, allow the production of end-products having high mechanical performances.

The present invention therefore relates to a process for the preparation of a flexible, thermoplastic composite filament, containing continuous inorganic fibres basically consisting of a flexible sheath in polyolefinic resin which covers a bundle of fibres impregnated with a polyolefinic resin, which comprises:

- a) unrolling a filament, consisting of a bundle of continuous inorganic fibres, from a bobbin;
- b) opening the bundle into the single fibres;
- c) wetting the open fibres with a solution basically consisting of:
 - i) an organic solvent;
 - ii) 1-30% by weight, preferably 3-15%, of at least one additive selected from those having the general formula:



or from ethylenically unsaturated organic compounds containing at least one anhydride, carboxylic or epoxidic function and triallylcyanurate

or trimethylallylisocyanurate;

and wherein R and R₁ represent a hydrogen atom or alkylic radical containing from 1 to 8 carbon atoms, R₂ represents an alkoxylic radical containing from 1 to 6 carbon atoms or a halogen, R₃ represents an alkyl radical containing from 1 to 8 carbon atoms, X represents a C₆-C₂₀ aromatic bivalent radical or cycloalkyl radical or C₄-C₁₀ heterocyclic radical or a radical having the formula
$$\begin{array}{c} -(-\text{CH}-)_m- \\ | \\ \text{R}_4 \end{array}$$
 wherein R₄ is a hydrogen atom or

a C₁-C₁₈ alkyl radical, n is an integer between 1 and 3, extremes included, m is an integer between 1 and 4, extremes included;

iii) 0-10% by weight, preferably 0.5-5%, of at least one organic peroxide;

d) evaporating the solvent from the wet fibre;

e) impregnating the open filament with a polyolefinic resin in powder form or in the form of a continuous fibre;

f) reclosing the impregnated filament and applying a protective flexible thermoplastic sheath around it in polyolefinic resin preferably of the same kind as the powder polyolefinic resin.

The composite filament of the present invention

has a high flexibility which allows it to be twisted, woven to form highly flexible mats or to be used as such to form articles of any kind or dimension, according to the filament winding technique. The filament can
5 have a fibre content of up to 70% in volume, preferably between 40 and 60%.

Any kind of inorganic fibre can be used to prepare the composite filament of the present invention. For example glass fibres with a titer of between 150 and
10 4000 tex.

The opening of the bundle of inorganic continuous fibres can be carried out with a mechanical roll system with a small diameter, a vibrating, pneumatic system etc. When the bundle has been opened, this is ready for
15 the wetting step of the fibres with the solution. This operation can be carried out either by immersing the opened bundle in the solution or drawing the group through two layers of fabric soaked in the solution itself.

20 The organic solvent suitable for preparing the solution preferably has a low boiling point and must be such as to dissolve both the products of group (ii) and the peroxides of group (iii). Preferred solvents are aliphatic hydrocarbons, such as pentane and hexane,
25 ketones such as acetone, alcohols, such as methanol,

ethanol and isopropyl alcohol, esters such as methylacetate or ethylacetate, etc.

Products having general formula (I) which can be used in the process of the present invention are those
5 wherein R and R₁ are hydrogen atoms; R₂ is -OCH₃, -OC₂H₅, -OC₃H₇, -OC₄H₉, X is a propyl; R₄ is a hydrogen atom; n is 2 or 3; m is equal to 3. These products are known in literature and available on the market. They can also be synthesized with well-known methods described, for
10 example, in U.S. patent 4.429.064.

Products with general formula (I) which can be used in the present process are: ethyl triethoxysilane, chloropropyl trimethoxysilane, vinyl triethoxysilane, vinyl trichlorosilane, vinyl triacetoxysilane, vinyl
15 tri(methoxyethoxy)silane, etc.

Finally, examples of ethylenically unsaturated organic compounds containing at least one anhydride, carboxylic or epoxidic function are unsaturated anhydrides such as maleic anhydride, itaconic anhydride,
20 citraconic anhydride, succinic alkenyl anhydrides with a C₄-C₁₈ alkenylic residue; unsaturated carboxylic acids such as maleic, crotonic, cinnamic, itaconic, acrylic, methacrylic acids; unsaturated epoxides such as glycidyl acrylate, glycidyl methacrylate; etc. Maleic
25 anhydride and acrylic acid are preferred.

Organic peroxides are products used as radicalic initiators in grafting reactions of polyfunctional compounds with polymeric matrixes and have an average half time of between 10 and 200 seconds at temperatures of between 100 and 240°C at which the grafting reactions generally take place. Examples of peroxides, according to the present invention, are: dicumyl peroxide, 2,5-bis(ter-butylperoxy)-2,5-dimethylhexane, ter-butylperoxybenzoate, ter-butylcumyl peroxide, 1,3-bis(ter-butylperoxy)isopropylbenzene, dibutyl peroxide, 2,5-dimethyl-2,5-bis(ter-butylperoxy)hexane, etc. These peroxides are commercially available and are prepared by AKZO CHEMIE under the trade-names Perkadox BC, Trigonox 101, Trigonox C, Trigonox T, Perkadox 14-90, Trigonox B, Trigonox 45.

When the solvent has been evaporated from the bundle, this is then ready for the impregnation of the fibres with the polyolefinic thermoplastic resin in powder form or in the form of continuous fibre.

Any system capable of impregnating continuous fibres with a powder can be used in the process of the present invention.

A valid impregnation system can be that with a fluid bed described in U.S. patent 4.614.678 which involves passing the open, dry bundle through the

powder maintained in a fluid state.

The fluid bed consists of a sealed chamber inside which the polyolefinic resin in powder form is maintained in suspension by air or nitrogen circulation.

5 The particle size of the powder is not critical but it is preferable to have an average dimension comparable to the diameter of each single fibre. The average diameter of the single particles is generally between 10 and 100 micrometers.

10 To favour impregnation, the open filament is guided inside the fluid bed with metallic pulleys preferably made of aluminium and asymmetrically arranged inside the sealed chamber, and at least one of these can be electrically connected to earth. The
15 number of pulleys is generally between 2 and 6.

Another system of impregnation involves immersing and drawing the open bundle through a static bed consisting of the polyolefinic resin in powder form contained in a basin subjected to continuous vibration.

20 The term static bed as used in the present invention and in the claims refers to a bed or a layer of powder wherein the single particles can move freely horizontally whereas the vertical movements are exclusively limited to the thickness of the layer.

25 The vibrations of the basin containing the polymer

in powder form can be obtained mechanically, by oscillating arms, or by means of an electromagnetic head. The intensity of the vibrations is not very high but must be such as to prevent the formation of preferential channels inside the powder during the passage of the bundle.

The impregnation of the inorganic fibres of the bundle with the polyolefinic resin in the form of continuous fibre can be carried out with the method known as "commingling" and described for example in European patent application publication 138.294.

Polyolefinic resins which are particularly suitable for the process of the present invention are polyethylene and polypropylene. More particularly preferred are the polyethylene homopolymer or copolymer of ethylene with 2-25% by weight of at least another C_3-C_{10} olefin such as propylene, butene-1, hexene-1, octene-1, etc. The polyolefinic resin has a density of between 0.85 and 0.97 g/cm³, preferably between 0.92 and 0.96 g/cm³. The Melt Flow Index of the polyolefinic resin is generally between 5 and 40 g/10 min.

When the impregnation step has been completed, a protective sheath is applied to the filament. The flexible sheath which covers the bundle of fibres is present in such a way as to have a weight ratio between

impregnating material and sheath of between 1:2 and 2:1.

The material of which the sheath is formed is selected from the thermoplastic polyolefins previously listed and can be of the same type or different from that selected for impregnation. Sheaths made of polyolefins which are not only different but also with melting points higher or lower than those of the polyolefins impregnating the bundle, can also be used.

10 The sheath is applied to the bundle of impregnated fibres with the known extrusion techniques described in European patent application publications 188.939 and 190.522.

According to these techniques the bundle impregnated with powder or polyolefinic fibres is coaxially passed through a tubular die, with a basically circular section, connected to an extruder. The plastic material, extruded with the tubular die, is drawn by a viscous effect during the passage of the bundle and, after a few seconds, cools to form the sheath.

It is also possible to have a composite filament with a transversal section basically elliptic in shape. In this case it is sufficient to have a tubular die with a geometrically rhombus-shaped transversal section.

An alternative method for having the transversal section of the composite filament with a basically elliptical shape involves crushing the impregnated bundle of fibres with a pair of rolls and extruding the sheath by means of a tubular die with a circular section having a diameter which is greater than that of the maximum thickness of the crushed bundle.

The composite material obtained with the process of the present invention has a high flexibility which allows it to be woven to form highly flexible mats or to be used as such to form articles of any shape or dimension, in accordance with the "filament winding" technique.

More specifically the composite material of the invention is suitable to be used as an alternative to metals and relative alloys in the building, transport, airspace industries, etc.

The process for the preparation of the flexible composite materials of the present invention can be better illustrated by referring to the drawing of the enclosed figure, which represents an operative but not limiting example.

A bundle of continuous fibres (1) is unrolled from a bobbin (2) and is fed to sealed chamber (3) through the seal opening (4).

Before entering the chamber, the bundle is opened, basically into its single fibres, using two or more cylinders (5), wetted by means of a passage between two rolls (5') covered with a fabric soaked with the solution of the present invention and then dried with a jet of air not shown in the figure.

Inside the chamber (3), a thermoplastic polyolefin in powder form (6) is maintained in a fluid state by a stream of air or nitrogen fed through the pipe (7).

10 The open bundle is drawn through the fluid bed by means of pulleys (8), (8') and (8'') and, during this passage, the fibres are impregnated with the powder. To favour the impregnation, the pulleys are electrically connected to earth (9, 9' and 9'').

15 The bundle impregnated with the powder leaves the chamber (3) and is reclosed through the seal opening (10) having at the top a tubular die (11), fed by an extruder (12), suitable for covering the impregnated bundle with a flexible sheath made of thermoplastic
20 polyolefin.

The composite filament thus prepared is passed through the pulley (13) to favour the air cooling, then, guided by the rolls (14) is collected onto the bobbin (15).

25 The following examples which are illustrative but

not limiting provide a better understanding of the present invention and its embodiment.

EXAMPLES 1-10

A bundle of glass fibres of 1200 tex of OCF type
5 R25B is unrolled from a bobbin at a rate of 20 m/min
and basically opened into its single fibres by passage
through a mechanical opener consisting of 4 rolls of 20
mm in diameter. The open bundle is uniformly wetted by
passing it between two layers of fabric soaked with an
10 acetone solution containing maleic anhydride (10% by
weight) and dicumyl peroxide (2% by weight) and dried
with an air jet.

The open bundle is subsequently impregnated by
passing it through a fluid bed obtained by bubbling in
15 1600 l/hour of nitrogen through 900 g of polyethylene
(ERACLENE MP 80 of Enichem S.p.A.) in powder form
contained in a sealed chamber.

Inside the sealed chamber, the open bundle is
guided by three pulleys asymmetrically arranged in such
20 a way that there is always a stretch of filament of
about 0.5 metres immersed in the fluid bed.

At the end of the impregnation, a sheath consist-
ing of polyethylene of the type ERACLENE MS 80 of
Enichem, is applied to the bundle.

25 The sheath is applied to the filament by a Maille-

fer 30 type extruder equipped with a die of the type used in the "sleeving" technique. The diameter of the die is 3.5 mm. The diameter of the chuck is 2.9mm. The diameter of the filament guide is 2.0 mm. The diameter
5 of the die inlet is 2.4 mm.

With the composite filament thus prepared, test samples were prepared for determining the mechanical characteristics of the moulded material.

Unidirectional plates of 28x28 cm were made with
10 the filament, which were stratified with the sequence $[0^\circ, 0^\circ, 90^\circ, 0^\circ]_{\text{symmetric}}$ and compression moulded at 190 and 210°C.

Table I summarizes the final results. For the determination of the shear resistance, reference was
15 made to the method described in "Developments in the Science and Technology of Composite Materials" ECCM5, B.Lauke et al, 1992, page 313, modified for flat test samples having dimensions of 10x10x3.3 mm.

Example	Damping	Fibre wt (%)	Sheath wt (%)	Powder wt (%)	Flexural Strength (*) (MPa)		Shear Resistance (MPa)	
					190°C	210°C	190°C	210°C
1	-	64.4	25.5	10.1	147	158	9.8	10.0
2	AM 10%	48.0	30.9	21.0	220	228	9.2	9.6
3	SM 5%	53.6	24.7	21.7	268	320	11.7	10.3
4	VTES 5%	59.3	27.8	13.0	227	285	9.2	9.6
5	TAIC 5%	57.1	21.9	21.1	226	254	7.5	9.1
6	AM 10%-DCP 2%	51.3	29.9	18.8	278	300	13.3	12.5
7	SM 5%-DCP 1%	57.7	24.2	18.1	244	270	10.4	10.2
8	VTES5%-DCP1%	63.0	21.1	15.9	271	268	9.2	8.3
9	TAIC 5%-DCP1%	58.9	25.3	15.8	244	238	11.3	8.2
10	Prosil 5%	54.2	32.4	13.5	280	291	11.1	11.3

*ASTM D790M, distance ratio between supports and 32/1 thickness

AM: maleic anhydride

SM: maleamic silane

TAIC: Triallylisocyanurate

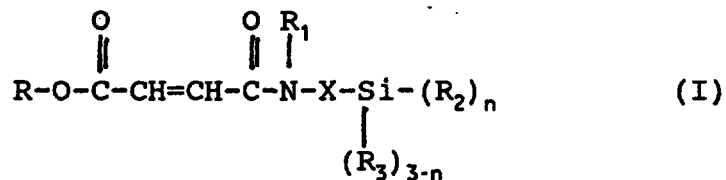
VTES: Vinyltriethoxysilane

DCP: Dicumyl

CLAIMS

1) Process for the preparation of a flexible, thermo-plastic composite filament, containing continuous inorganic fibres basically consisting of a flexible sheath in polyolefine resin which covers a bundle of fibres impregnated with a polyolefine resin, which comprises:

- a) unrolling a filament, consisting of a bundle of continuous inorganic fibres, from a bobbin;
- b) opening the bundle into the single fibres;
- c) wetting the open fibres with a solution basically consisting of:
 - i) an organic solvent;
 - ii) 1-30% by weight of at least one additive selected from those having the general formula:



or from ethylenically unsaturated organic compounds containing at least one anhydride, carboxylic or epoxidic function and



trialllylcyanurate or trimethylallylisocyanurate;

and wherein R and R₁ represent a hydrogen atom or alkylic radical containing from 1 to 8 carbon atoms, R₂ represents an alkoxylic radical containing from 1 to 6 carbon atoms or a halogen, R₃ represents an alkyl radical containing from 1 to 8 carbon atoms, X represents a C₆-C₂₀ aromatic bivalent radical or cycloalkyl radical or C₄-C₁₀ heterocyclic radical or a radical having the formula

$$\begin{array}{c} \text{-(CH-)}_m\text{-} \\ \quad \quad \quad \diagdown \\ \quad \quad \quad \text{R}_4 \end{array}$$

wherein R₄ is a hydrogen atom or

a C₁-C₁₈ alkyl radical, n is an integer between 1 and 3, extremes included, m is an integer between 1 and 4, extremes included;

iii) 0-10% by weight, preferably 0.5-5%, of at least one organic peroxide;

- d) evaporating the solvent from the wet fibre;
- e) impregnating the open filament with a polyolefine resin in powder form or in the form of a continuous fibre;
- f) reclosing the impregnated filament and applying a protective flexible thermoplastic sheath around it in polyolefine resin pref-



* C L N S *

erably of the same kind as the powder polyol-
efine resin.

- 2) Process according to claim 1, wherein the compos-
ite filament has a fibre content of up to 70% in
5 volume.
- 3) Process according to claims 1 or 2, wherein the
products having general formula (I) are those
wherein R and R₁ are hydrogen atoms; R₂ is -OCH₃,
-OC₂H₅, -OC₃H₇, -OC₄H₉; X is a propyl; R₄ is a hydro-
10 gen atom; n is 2 or 3; m is equal to 3.
- 4) Process according to any of the previous claims,
wherein the products having general formula (II)
are selected from: ethyl triethoxysilane, chloro-
propyl trimethoxysilane, vinyl triethoxysilane,
15 vinyl trichlorosilane, vinyl triacetoxysilane,
vinyl tri(methoxyethoxy)silane.
- 5) Process according to any of the previous claims,
wherein the ethylenically unsaturated organic
compounds containing at least one anhydride,
20 carboxylic or epoxidic function are selected from:
unsaturated anhydrides such as maleic anhydride,
itaconic anhydride, citraconic anhydride, succinic
alkenyl anhydrides with a C₄-C₁₈ alkenylic residue;
unsaturated carboxylic acids such as maleic,
25 crotonic, cinnamic, itaconic, acrylic, methacrylic



acids; unsaturated epoxides such as glycidyl acrylate, glycidyl methacrylate.

- 6) Process according to any of the previous claims, wherein the organic peroxides are selected from:
- 5 dicumyl peroxide, 2,5-bis(ter-butylperoxy)-2,5-dimethylhexane, ter-butylperoxybenzoate, ter-butylcumyl peroxide, 1,3-bis(ter-butylperoxy)isopropylbenzene, dibutyl peroxide, 2,5-dimethyl-2,5-bis(ter-butylperoxy)hexine.
- 10 7) Process according to any of the previous claims, wherein the polyolefine resins are polyethylene and polypropylene.
- 8) Process according to claim 7, wherein the polyolefine resin is the polyethylene homopolymer or
- 15 copolymer of ethylene with 2-25% by weight of at least another C_3 - C_{10} olefin.
- 9) Process according to any of the previous claims, wherein the material of which the sheath is made is selected from thermoplastic polyolefins.

