Reduction of filler networking in silica based elastomeric nanocomposites with exfoliated organo-montmorillonite

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Montmorillonite with dimethyl di(hydrogenatedtalloyl) ammonium as the compensating cation was added to a silica based elastomeric composite and the hybrid filler system led to reduction of filler networking phenomenon, better stability of elastic modulus with temperature, enhancement of stresses at all elongations, improvement of ultimate properties. This composite was based on a blend of natural rubber, poly(1,4-cis-isoprene) and poly(sty-rene-co-butadiene) from anionic polymerization and contained 70 parts per hundred rubber (phr) of silica. The organically modified clay (OC) was below the threshold required to establish an hybrid OC-silica filler network. Such threshold (about 7 phr) was estimated by preparing silica based nanocomposites containing various amounts of OC and determining shear storage and loss moduli as a function of strain amplitude. This work dem-onstrates that exfoliated OC favour lower dissipation of energy of silica based elastomeric composites under dy-namic mechanical stresses and paves the way for further large scale applications.

Keywords: Silica Montmorillonite Organoclay Nanocomposite Filler networking

1. Introduction

Silica is characterized by the polar surface rich of oxygen atoms, in-volved in siloxane and silanol functional groups (Legrand, 1998). Pre-cipitated silica (Chevalier and Morawski, 1984; Legrand, 1998), having high specific surface area and activity, has thus poor dispersibility in apolar matrices and easily forms filler networks in polymer melts and elastomers (Donnet and Custodero, 2005). Compatibilization of precip-itated silica with hydrocarbon polymers is achieved by using molecules, such as organosilanes, that react with silanols, shielding the silica sur-face and that, in some cases, form covalent bonds with the polymer chains (Leblanc, 2002; Donnet and Custodero, 2005; Fröhlich et al., 2005). Composites based on diene elastomers, precipitated silica and organosilanes reactive with the polymer are the preferred ones for large scale highly demanding dynamic-mechanical applications such as the one in tyre compounds, when low dissipation of energy is re-quired (Leblanc, 2002; Donnet and Custodero, 2005; Fröhlich et al., 2005). In fact, the efficient distribution and dispersion of the silica/silane system and its interaction with the polymer matrix substantially reduce the filler networking phenomenon, that is known as the main source of energy dissipation. The most important application of precipitated silica is indeed in the field of elastomeric composites. Research is

Article history: Received 15 June 2016 Received in revised form 12 September 2016 Accepted 12 September 2016 Available online 24 September 2016 continuously performed both in academic and industrial laboratories in order to achieve further reduction of the filler networking in such sil-ica based composites.

Cationic clay minerals (Bergaya and Lagaly, 2013) have been widely used over the last decades for the preparation of both thermoplastic and elastomeric composite materials (LeBaron et al., 1999; Alexandre and Dubois, 2000; Ray and Okamoto, 2003; Annabi-Bergaya, 2008; Chen et al., 2008; Galimberti, 2011a). Their compatibilization with apolar poly-mer matrices is successfully achieved through the exchange reaction of the alkali and alkali-earth cations with organophilic ammonium salts (Lagaly et al., 2013). Clay polymer nanocomposites (CPN) contain-ing the organically modified clays (OC) are characterized by improved mechanical reinforcement, impermeability and thermal stability (Paul and Robeson, 2008; Galimberti et al., 2013a). It is also acknowledged that OC give rise to remarkable filler networking in polymer melts and elastomers (Ramorino et al., 2009; Galimberti et al., 2011b; Galimberti et al., 2012), that means to pronounced Payne effect (Payne, 1962). The addition of OC to silica based composites could thus improve impor-tant properties, but could also lead to an undesired enhancement of fill-er networking phenomenon.

It is thus worthwhile to study the filler networking phenomenon and the static and dynamic-mechanical properties of polymer melts and elastomers based on silica and OC. In the scientific literature, many papers are available on elastomeric nanocomposites based on OC and carbon black (CB) (Maiti et al., 2005; Cataldo, 2007; Praveen et al., 2009; Galimberti et al., 2009, 2012, 2013b; Sridhar et al., 2009; Das

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et al., 2010; Chattopadhyay et al., 2011; Gopi et al., 2011; Malas and Das, 2012). It has been shown that OC favours the filler networking phenom-enon, leading to the formation of hybrid OC-CB filler networks (Galimberti et al., 2012, 2013b).

The intimate interaction between CB and a nanofiller has been justi-fied on the basis of the zeta potential (ZP) values (Bhattacharya and Bhowmick, 2010): in the pH range of elastomeric composite, ZP of CB is positive, whereas those of nanofillers are negative (Gates, 2004; Lin et al., 2006; Sabah et al., 2007; Xu et al., 2007; Kvande et al., 2008). It was commented that ZP can be also applied to polymer melts (Brochard-Wyart and de Gennes, 2000; Flores et al., 2007) and the affin-ity of clay minerals for CB was reported (Feller et al., 2004; Konishi and Cakmak, 2005; Galimberti et al., 2012).

The affinity of clay minerals for silica is intuitive. Moreover, positive values of ZP have been reported for silica (Anderson and Zukoski, 2007). The addition of OC to silica based composite could thus lead to the en-hancement of the filler networking phenomenon. Only a few papers are available on elastomeric CPN containing silica and OC: they are based on poly(1,4-cis-isoprene) from *Hevea brasiliensis* (natural rubber, NR) (Rattanasom and Prasertsri, 2012) and poly(styrene-*co*-butadiene)(SBR) (Kim et al., 2009, 2011, 2013). The presence of OC was shown to enhance the tensile moduli and to worsen the ultimate properties (Kim et al., 2011) and it was reported that OC does not form filler–filler net-works with silica (Kim et al., 2013). The amount of OC was kept constant and the content of silica was not higher than 53 parts per hundred rub-ber (phr) (Kim et al., 2013).

Aim of this work was to study the filler networking phenomenon in elastomeric nanocomposites containing precipitated silica and OC. Montmorillonite (Mt) with dimethyl di-hydrogenated talloyl (2HT) as the compensating cation was used as the OC. Poly(1,4-cisisoprene) from Ziegler-Natta catalysis (PI) and a blend of PI, SBR and poly(1,4-cis-butadiene) were used as the polymer matrix and nanocomposites were prepared via melt blending. Structure of nanocomposites contain-ing OC and silica was investigated by means of transmission electron microscopy (TEM) and X-ray diffraction (XRD) analysis. Dynamic-me-chanical moduli were determined through measurements in the torsion and in the axial mode. Stresses at various elongations and ultimate properties were assessed with quasi-static measurements.

2. Experimental

2.1. Materials

2.1.1. Organoclay and ammonium salt

OC was Mt with dimethyl di(hydrogenated talloyl) ammonium (2HT) as the compensating cation (Dellite® 67G from Laviosa Chimica Mineraria S.p.A.). Mt and the ammonium moiety were 55 and 45 as mass %, respectively. The ammonium salt was dimethyl dihydrogenat-ed talloyl ammonium chloride (2HTCl) (Arquad® HC Pastilles from Akzo Nobel). Typical value of chemical purity was 98.5 mass %. The alkyl chain distribution, as mass%, was: $<C_{12} = 1$, $C_{14} = 4$, $C_{16} = 31$, $C_{18} = 64$.

2.1.2. Silica

Precipitated silica was Ultrasil 7000 from Evonik, with 270 g/L den-sity and 175 m²/g as specific surface area, determined through nitrogen absorption.

2.1.3. Polymers

Synthetic poly(1,4-*cis*-isoprene) (PI) (SKI3 from Nizhnekamskneftechim Export) had 70 Mooney Units (MU) as Mooney viscosity ($M_{L(1 + 4)}$ at 100 °C). Poly(1,4-*cis*-butadiene) (BR) (Buna CB 25 from Lanxsess) had 44 MU as Mooney viscosity ($M_{L(1 + 4)}$ at 100 °C). Poly(styrene-*co*-butadiene) from anionic solution polymerization (sSBR) (SOL R 72612 from Versalis) had 25 mass% of styrene, 50

mass% of vinyl groups, 36.8 phr of aliphatic extension oil and 48 MU as Mooney viscosity ($M_{L(1 + 4)}$ at 100 °C).

2.1.4. Other ingredients for composites preparation

Aliphatic oil catenex SNR was from Shell; bis(triethoxysilylpropyl) tetrasulfide (TESPT) (Si69) was from Evonik. ZnO was from Zincol Ossidi; stearic acid (chemical composition: C16 + C18 = 95 mass%, C12 + C14 = 5 mass%) was from Sogis. *N*-(1,3-dimethylbutyl)-*N*'-phenyl-*p*-phenylendiamine (6PPD) was from Flexsys, 2,5-dimethyl-2,5-di(*t*-butylperoxy)hexane (DCUP) supported on Silica/CaCO₃ (45 mass%) was from Arkema Inc., and sulphur was from Solfotecnica. Benzothiazyl-2-cyclohexylsulfenamide (CBS), diphenyl guanidine (DPG), and *N*-(cyclohexylthio)phthalimide (CTP), were from Flexsys.

2.2. Preparation of composites and nanocomposites

2.2.1. PI based nanocomposites containing silica and OC

Formulations are in Table 1. A PI/silica masterbatch was firstly prepared. In the first step of the procedure, 738 g of PI were introduced into the Banbury® type internal mixer (1050 mL mixing chamber) at 60 °C and masticated for 30 s with rotors rotating at 75 rpm. In the sec-ond step, 517 g of silica and 51.7 g of TESPT were added, temperature was rapidly increased to 150 °C, mixing was carried out for further 240 s and the resulting PI/silica masterbatch was discharged at 150 °C. This masterbatch was then used for the preparation of all the nanocom-posites. For the composite with only silica as the filler (OC-0), the PI/sil-ica masterbatch was introduced into the Brabender® type internal mixer (50 mL mixing chamber) and peroxide was added, performing the mixing at 50 °C for 120 s. For the nanocomposites containing silica and OC, the PI/silica masterbatch was introduced into the Brabender® type internal mixer at 80 °C and masticated at 30 rpm for 60 s. OC was then added and mixing was carried out for 240 s, discharging the nano-composite at about 90 °C. The nanocomposite with OC was finally fed at room temperature to the Brabender® mixer and the peroxide was added, performing the mixing at 50 °C for 120 s. Curing was carried out at 170 °C for 20 min under a pressure of 150 bar. Nanocomposites were labelled as OC-X, where X indicates the amount of OC expressed in phr (part per hundred rubber).

2.2.2. PI based composites containing silica and 2HTCl

The same formulations reported in Table 1 were adopted, except that in place of OC a molar amount of 2HTCl equal to the molar amount of 2HT in OC was added. The amounts of 2HTCl in phr were: 0.22, 0.49, 0.76, 1.03, 1.53, 2.03, 3.60, 6.75, 8.10. The PI/silica masterbatch was in-troduced into Brabender® type internal mixer at 80 °C and masticated at 30 rpm for 60 s. The ammonium salt was then added and mixing was carried out for 240 s, discharging the composite at about 90 °C. The resultant composite was finally fed at room temperature to the mixer and peroxide was added, performing the mixing at 50 °C for 120 s. Curing was carried out at 170 °C for 20 min under a pressure of 150 bar. Composites were labelled as 2HTCl-Y, where Y indicates the amount of 2HTCl expressed in phr.

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Formulations of PI based nanocomposites containing silica and OC. ^{a-c}	
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	OC-	OC-	0C-	OC-	OC-	0C-	0C-	OC-	OC-	OC-	OC-
	0	0.5	1.1	1.7	2.3	3.4	4.5	8	12	15	18
PI	100	100	100	100	100	100	100	100	100	100	100
Silica	70	70	70	70	70	70	70	70	70	70	70
OC	0	0.5	1.1	1.7	2.3	3.4	4.5	8.0	12.0	15.0	18.0

^a OC = Mt/2HT.

Table

^b Amount of ingredients are indicated in phr (part per hundred rubber).

^c Other ingredients: TESPT 7, DCUP 1.4.

2.2.3. PI-BR-sSBR based nanocomposites containing silica and OC

Formulations are in Table 2. In the first step of the procedure, polymers were introduced into in the Banbury® type internal mixer at 60 °C and were masticated for 30 s with rotors rotating at 75 rpm. In the second step, silica or silica and OC were then added, together with silane TESPT, stearic acid and the oil, to prepare the silica based composite or the nanocomposite, respectively. Temperature was rapidly increased to 110 °C and mixing time of the second step was 240 s. The resulting masterbatches were discharged at about 110 °C, and stored at room temperature for about 16 h. Afterwards, the addition of ZnO and 6PPD was performed by mixing at 50 °C for 120 s. The remaining ingredients (S, CBS, DPG, CTP) were then added, performing a further mixing at 50 °C for 180 s. The compounds were finally homogenised by passing them 5 times on a two roll mill, with the front roll rotating at 30 rpm and the back roll at 38 rpm. Curing of the composites was carried out at 170 °C for 20 min under a pressure of 150 bar. Composites were labelled as PI-BR-sSBR-OC-X, where X indicates the amount of OC expressed in phr.

2.3. Characterization of OC and nanocomposites

2.3.1. X-Ray diffraction

Wide-angle X-ray diffraction (WAXD) patterns with nickel filtered Cu-K α radiation were obtained in reflection, with an automatic Bruker D8 Advance diffractometer. Patterns were recorded in 2°–80° as the 2 θ range, being 2 θ the peak diffraction angle.

2.3.2. Transmission electron microscopy (TEM)

TEM analysis of nanocomposites was performed with a Zeiss EM 900 microscope applying an accelerating voltage of 80 kV. Ultrathin sections (about 50 nm thick) were prepared at -130 °C by using a Leica EM FCS cryoultramicrotome.

2.3.3. Mooney viscosity measurements

Mooney viscosity was determined with a Mooney MV 2000 E Viscometer from Alpha Technologies, following the experimental procedure described by ASTM D1646. In particular, tests were run at 100 °C, using a rotor with a 38.1 mm diameter rotating at 0.209 rad/s. Rubber samples were preheated for 1 min at 100 °C and the rotor was allowed to rotate for 4 min before measuring the viscosity value, indicated in Mooney Units (MU).

2.3.4. MDR rheometric analysis

It was conducted using a Monsanto MDR rheometer, the test being carried out at 170 °C, for 30 min, at an oscillation frequency of 1.66 Hz (100 oscillations per min) and an oscillation amplitude of \pm 0.5°. The following parameters were obtained: M_L is the minimum torque measured; M_H is the maximum torque measured; ts_x is the time needed to have a torque equal to $(M_L + x)$; t_x is the time at x% of vulcanization required to have a torque value equal to $M_L + (x/100)(M_H - M_L)$.

2.3.5. Tensile and Hardness measurements

Both of them were performed on vulcanized samples. Tensile tests were carried out according to Standard ISO 37:2005, with a

Table 2				
Formulations of nanocom	posites based on P	I, BR and sSBR v	with silica and	OC as fillers. ^{a,b}

	OC-0	OC-3
sSBR	45	45
PI	25	20
BR	30	30
Silica	70	70
OC	0	3

^a Amount of ingredients are indicated in phr.

^b Other ingredients: TESPT 5.2, stearic acid 2, oil 25, ZnO 3, 6PPD 2, Sulphur 1.2, CBS 1.5, DPG 2, CTP 0.5.

dynamometer (Zwick Roell Z010) with optical extensometer. The clamps rate was 1 mm/min and the chamber load was 10 kN. Stresses at 50, 100 and 300% elongation (σ_{50} , σ_{100} and σ_{300} , respectively), stress at break (σ_b), elongation at break (ϵ_B) and the energy required to break were measured. The hardness in IRHD degrees (at 23 °C and at 70 °C) was measured according to Standard ISO 48:1994.

2.3.6. Dynamic mechanical measurements

2.3.6.1. In the shear mode. Monsanto R.P.A. 2000 rheometer was used. Samples were first kept in the instrument at 120 °C for 10 min. A strain sweep (0.28-25% strain) was then performed, on raw samples, at 120 °C. Afterwards, the curing was carried out at 170 °C for 30 min, with a frequency of 1.667 Hz and an angle of 6.98% (0.5 arc). Cured samples were left in the instrument for 10 min at 50 °C. Data reported in this manuscript were finally obtained with a strain sweep (0.28-25% strain) performed at 50 °C with a frequency of 1 Hz.

2.3.6.2. In the axial mode. Dynamic-mechanical properties were measured on the crosslinked elastomeric compositions using an Instron dynamic device in the traction-compression mode according to the following procedure. Specimen with a cylindrical form (length = 25 mm; diameter = 12 mm), kept at the fixed temperature (10, 23, 70 and 100 °C) for the whole duration of the test, was compression-preloaded with 25% longitudinal deformation with respect to the initial length and was then submitted to a dynamic sinusoidal strain having an amplitude of \pm 3.5% with respect to the length under pre-load, with a 100 Hz frequency.

3. Results and discussion

Structure of PI/OC/silica nanocomposites, whose formulations are in Table 1, was investigated by means of XRD and TEM analysis. Fig. 1 shows the XRD patterns of pristine OC clay (a) and of OC-15 nanocom-posite (b).

In the pattern of pristine OC clay, (001), (002) and (003) reflections, due to periodicities perpendicular to the structural layers, are visible at 2.5, 4.7 and 7.2 as 2 θ angle values. From the 2 θ angle value of (001) re-flection, the 001 basal spacing can be evaluated equal to 3.5 nm. In con-sideration of the thickness of the clay layer (\approx 1 nm) and of the length of 2HT (\approx 2.5 nm) (Osman et al., 2002), the tilting angle of the hydro-carbon chains in the paraffin-type arrangement can be evaluated to be not far from 60°. Reflections due to OC clay are not present in the pattern of the nanocomposite. As reported in the literature (Galimberti et al., 2007, 2011b) such amount of OC is high enough to be detectable in



Fig. 1. XRD patterns of Mt/2HT as OC (a) and of OC-15 nanocomposite (b).

XRD diffractograms. The absence of reflections seem thus to suggest the occurring of OC exfoliation.

TEM micrographs were taken on OC-8 and OC-15 nanocomposites and are shown in Fig. 2a,b and Fig. 2c,d, respectively.

OC distribution is similar in the two nanocomposites. In all of the mi-crographs, OC aggregates and isolated layers appear evenly distributed: they are visible both in the elastomer matrix and in between silica ag-gregates. Many isolated layers are visible at high magnification, suggest-ing a high level of OC exfoliation, in agreement with the results obtained from XRD analysis. The micrograph taken at high magnification on OC-15 (Fig. 2d) reveals that both layers and stacks seem to act as bridges among silica aggregates, forming an almost continuous hybrid network.

The study of the filler networking phenomenon moved from the Guth equation (Eq. (1)) (Guth, 1945a, 1945b):

$$G_c/G_m = 1 + 0.67f\Phi + 1.62f^2\Phi^2 \tag{1}$$

where G_c is the elastic modulus of the composite, G_m is the elastic modulus of the neat elastomer, Φ is the filler volume fraction and the quadratic term accounts for the mutual disturbance of filler particles. f, known as shape factor, is the ratio of length to thickness of particles or aggregates The ratio G_c/G_m is the enhancement of the neat elastomer modulus, that occurs when the elastomer is filled with rigid filler particles having largely higher elastic modulus, so that the elastic energy is stored in the elastomer. This model was applied in previous studies on nanocomposites based on Mt/2HT as the OC in NR (Ramorino et al., 2009) and in PI (Galimberti et al., 2012) as the neat polymer matrices. Initial modulus values were obtained either as the slope at the origin of stress-strain curves (Ramorino et al., 2009; Galimberti et al., 2012)

Table 3

Data of dynamic storage modulus, G', and dynamic loss modulus, G'', obtained for crosslinked OC/Silica/Pl nanocomposites of Table 1.

	OC- 0.5	0C- 1.1	OC- 1.7	0C- 2.3	0C- 3.4	0C- 4.5	OC- 8	0C- 12	0C- 15	OC- 18
OC (phr)	0.5	1.1	1.7	2.3	3.4	4.5	8	12	15	18
$G'c^{a}$ (MPa)	2.9	2.9	2.9	2.8	2.8	2.9	3.2	3.4	3.6	3.6
$G'c^{\mathrm{a}} - G'm^{\mathrm{b}}$ (MPa)	-	2.6	2.5	-	2.1	2.0	1.6	-	-	-

^a $G'c = G'_{\gamma min}$ (0.3%) of OC-X nanocomposites.

^b $G'm = G'_{\gamma min}$ (0.3%) of PI/Silica/2HTCl composites.

or as dynamic shear storage *G'* modulus at minimum deformation (Galimberti et al., 2012). Modulus enhancement as a function of the OC content was found to be strongly not linear, suggesting two different regimes for the mechanical reinforcement. By reporting the excess of modulus as a function of the OC content in a double logarithmic plot, ap-plying the Huber-Vilgis method originally developed for CB based com-posites, the OC percolation threshold was calculated at about 8 phr in NR and at about 6 phr in IR.

In the present work, the filler networking phenomenon was investi-gated by analyzing the dependence on OC content of dynamic shear storage G' and loss G'' moduli of crosslinked nanocomposites. Crosslinking was performed with peroxide, used in place of sulphur based crosslinking system, that is known to interact with OC. The ammonium cations increase the mobility of sulphur accelerating anionic species and amines, that arise from thermal decomposition of the ammonium cations, react with sulphur (Giannini et al., 2011). Acceleration of sulphur based crosslinking reaction and increase of crosslinking net-work density have been documented (Avalos et al., 2008; Maiti et al.,



Fig. 2. TEM micrographs of OC-8 (a,b) and OC-15 (c,d) nanocomposites.





Fig. 5. Dynamic loss modulus in shear mode at 1 Hz plotted as a function of dynamic shear modulus for crosslinked nanocomposites of Table 1.

Fig. 3. Dynamic storage modulus in shear mode at 1 Hz plotted as a function of the strain amplitude for crosslinked nanocomposites of Table 1.

2008; Verdejo et al., 2011). *G*'and *G*" moduli were measured as a function of the strain amplitude, with strain sweep tests. *G*' modulus at min-imum deformation ($G'_{\gamma min}$) was obtained and was taken as initial modulus value. Values of $G'_{\gamma min}$ are reported in Table 3. Fig. 3 shows the dependence of *G*' modulus on the strain amplitude for the crosslinked nanocomposites.

It appears that curves of nanocomposites with low OC content (\leq 4.5 phr) cross over the curves of nanocomposites with higher OC content (\geq 8 phr), at strain amplitudes close to 5%. Moreover, crosslinking curves shown in Fig. 4 reveal that the maximum modulus M_H decreases as the OC content increases.

These findings could be interpreted considering that the lipophilic ammonium cation acts as plasticizer, reducing the modulus of the



Fig. 4. Rheometer curves for nanocomposites of Table 1.

matrix. However, other hypotheses could be taken into account. OC could affect the strength of filler network and of polymer-filler interac-tion at high strain amplitudes. The ionic liquid could also have an effect on the efficiency of silanization. However, recently a phosphonium ionic liquid has been reported to enhance the efficiency of the silanization re-action between silica and silane TESPT, behaving as a catalyst (Tang et al., 2015)."

Fig. 5 shows, in the Cole-Cole plot, the dependence of *G*" on *G*' mod-ulus, for crosslinked OC-X samples.

At a given G' value, G'' values are quite similar for nanocomposites with OC content from 0.5 to 4.5 phr, whereas they appreciably and con-sistently increase when the OC content is at least 8 phr. This suggests that, at a given OC content, the filler networking phenomenon has a clear enhancement. The amount of OC required to have such enhance-ment could be estimated adopting the approach reported in the litera-ture and commented above, correlating the initial modulus enhancement, given by the difference $\Delta G'$ between the modulus G'_c of



Fig. 6. Double logarithmic plot of the excess of G' modulus $\Delta G' = G'_c - G'_m$ (see text) as a function of OC volume fraction (Huber–Vilgis plot).



Fig. 7. Nominal stress/nominal strain curves for silica based nanocomposites of Table 1.

the composite containing OC and the modulus G'_m of the matrix ($\Delta G' = G'_c - G'_m$), with the nanofiller content. However, it is evident from what shown in Fig. 3 and in Fig. 4 that the G'_m modulus of the PI/silica com-posite (without OC) can not be taken as the reference modulus for all the nanocomposites. To overcome this problem, analogous composites were prepared feeding only the ammonium salt 2HTCl, in the absence of Mt. G'_m values were thus determined from strain sweep experiments in the shear mode. These experiments revealed that 2HTCl led to the reduction of G' modulus, both at minimum and at large strain, and to the reduction of $\Delta G'$. G' values at minimum strain, $G'_{\gamma min}$ (0.3%), were taken as the initial modulus of the composites and were used to determine the excess of the initial modulus $\Delta G'$. Values of the excess of modulus $\Delta G' = G'_c - G'_m$, where G'_m is the $G'_{\gamma min}$ of the neat matrix containing the ammonium salt 2HTCl, were reported in Table 3.

The Huber-Vilgis method was then applied to the hybrid filler system, by reporting the modulus excess $\Delta G'$, as a function of OC content in a double logarithmic plot. The Huber Vilgis plot shown in Fig. 6 reveals two different regimes for the mechanical reinforcement, the switch from one to the other occurring at 7.4 phr as OC content. Such a shift could be interpreted with the formation of continuous silica-OC hybrid filler network.

This interpretation appears to be supported by the TEM micrographs shown in Fig. 2. In fact, a continuous network formed by silica and OC layers and aggregates is visible in the pictures of OC-15 (Fig. 2d).

Silica based nanocomposites presented in this manuscript have fea-tures similar to those based on CB, previously reported (Galimberti et al., 2012): they are based on the same polymer and OC, silica and CB have the same volume fraction. The existence of two regimes for the mechanical reinforcement was verified also for these OC/CB

Table 4

Viscosity data and crosslinking reaction parameters for PI-BR-sSBR-OC-X samples of Table 2.

	PI-BR-sSBR-OC-0	PI-BR-sSBR-OC-3
MU ^a	85.7	75.2
M _L [dNm]	4.2	3.3
M _H [dNm]	17.0	15.9
t _{s1} [min]	1.410	1.450
t ₉₀ [min]	4.790	4.200

^a From Mooney measurements.

Table 5

 $G'_{(0.5\%)}$ modulus and $\Delta G'_{(0.5-10\%)}$ for PI-BR-sSBR-OC-X samples of Table 2.

	PI-BR-sSBR-OC-0	PI-BR-sSBR-OC-3
<i>G</i> ′ _(0.5%) [MPa]	4.4	3.7
$\Delta G'_{(0.5-10\%)}$ [MPa]	2.9	2.2

nanocomposites (Galimberti et al., 2012), and the shift from one regime to the other was found to occur for OC content of about 3 phr. Larger amount of OC is thus required to form a continuous hybrid filler net-work with silica rather than with CB. This finding can be justified with the dispersion of OC in silica domains, thanks to their affinity, and indi-cates the possibility of using larger amounts of OC without enhancing the filler networking phenomenon in silica based nanocomposites.

Tensile properties of nanocomposites of Table 1, containing 3.4, 12 and 18 phr of OC, were investigated by means of quasi-static measure-ments. Fig. 7 shows the obtained stress strain curves.

Nanocomposites with OC show higher break energy than the composite with only silica as the filler. The largest increase, from 34.7 to 38.3 J/cm³, was obtained when the OC amount (3.4 phr) was below the percolation threshold. Values of initial modulus were calculated from the stress strain curves: also quasi-static measurements confirm that appreciable filler networking occurs only for OC content above the percolation threshold.

Results discussed so far show that the addition of OC to a silica based elastomeric composites leads to the improvement of the composite end properties, without affecting the dynamic-mechanical behaviour, that means without enhancing the filler networking phenomenon, provided that the OC amount is not high enough to give rise to a continuous hy-brid filler network with silica.

To verify the effect of the polymer matrix on the filler networking phenomenon of OC/silica hybrid filler system, composite was prepared with a blend of PI, BR and SBR from anionic solution polymerization (sSBR). Silica was used in the same amount as in the composite discussed above and the OC content (3 phr) was lower than the one re-quired to have the formation of the hybrid filler network. The effect of the crosslinking system was investigated by using a traditional sulphur based system, in place of peroxide. Formulations are shown in Table 3. Table 4 reports data on viscosity and parameters of the crosslinking re-action obtained for PI-BR-sSBR-OC-X samples of Table 2.

OC caused the reduction of the composite viscosity, as shown by Mooney viscosity and M_L values, whereas let the crosslinking reaction substantially unaffected. The latter finding could be due to the low amount of OC in the nanocomposite and also to the interaction of the amine arising from the ammonium decomposition with silica rather with sulphur.

Dynamic-mechanical moduli were determined in the shear mode, from strain sweep tests. G' at minimum strain $(G'_{(0.5\%)})$ and $\Delta G'_{(0.5-10\%)}$ data are in Table 5.

It is evident that OC leads to the reduction of the filler networking phenomenon.

Table 6

E', *tan* δ at different temperatures and $\Delta E'_{23-70}$ _C values for PI-BR-sSBR-OC-X samples of Table 2.

	PI-BR-sSBR-OC-0	PI-BR-sSBR-OC-3
E' @ 0 °C	11.7	10.9
tanδ @ 0 °C	0.40	0.38
E' @ 10 °C	9.6	9.2
tanδ @ 10 °C	0.35	0.32
E' @ 23 °C	8.3	8.0
tanδ @ 23 °C	0.27	0.24
E' @ 70 °C	5.9	6.0
tanδ 70 °C	0.17	0.16
E' @ 100 °C	5.5	5.6
tanδ @ 100 °C	0.16	0.13
$\Delta E'_{23-70°C}$	6.1	5.2



Fig. 8. Nominal stress/nominal strain curves for PI-BR-sSBR-OC-X samples of Table 2.

Dynamic-mechanical measurements were performed also in the axial mode, in wide range of temperatures, from 0 to 100 °C. Data of dy-namic storage modulus (E') and loss factor (tan δ) are in Table 6.

The $tan\delta$ value is calculated as the ratio between loss (E'') and storage modulus (E'). E' values of the OC based nanocomposite are lower at low temperatures (from 0 °C to 23 °C) and slightly higher at high temperatures (70 °C and 100 °C): the reduction of the storage modulus with the increase of temperature is more pronounced in the absence of OC. Moreover, $tan \delta$ values are similar at low temperatures in the absence and in the presence of OC and become appreciably lower for the OC based nanocomposite as the temperature increases. All these findings are clear indication of the reduction of the filler network, caused by OC.

Fig. 8 shows the stress strain curves for PI-BR-sSBR-OC-X samples of Table 2.

OC brought to higher stresses at all the elongations: σ_{100} was enhanced from 1.3 to 1.4 MPa and σ_{300} from 8.3 to 9.2 MPa.

Ultimate properties were improved by OC: the stress at break (σ_B) was enhanced from 13.3 to 14.2 MPa, the break energy from 29.4 to 31.7 MPa, while the elongation at break remained substantially unchanged (485 vs 477). Hardness was measured at 23 °C and 70 °C, detecting 74.3 and 71.9 as the hardness values at 23 °C for PI-BR-sSBR-OC-0 and PI-BR-sSBR-OC-3, respectively, with a reduction at 70 °C of 5.7 and 5.2 points.

All data collected for the PI-BR-sSBR-OC-X samples appear highly consistent and show that OC, used in amount lower than the one required to form the hybrid filler network with silica, in a nanocomposite crosslinked with a sulphur based system, leads to appreciable reduction of the filler networking phenomenon. To justify these results one should also take into account the role played by the ingredients, such as ZnO, stearic acid, sulphenamides, that are added to perform the sulphur based crosslinking.

4. Conclusions

This work demonstrates that the addition, to a silica based elastomeric composite, of an organically modified clay, a montmorillonite with a compensating ammonium cation bearing long chain alkenyl substituents, brings about reduction of filler networking phenomenon, lower reduction of elastic modulus as the temperature increases, improvement of stresses at every elongation and of ultimate properties such as the break energy. The ammonium cation 2HTCl, used in silica based compounds in the absence of montmorillonite, appears to be able to bring about the reduction of the filler networking, but also of the mechanical reinforcement. To achieve the abovementioned positive effects, OC should be used in a low amount, below the threshold content at which continuous hybrid OC/silica filler network is formed. Crosslinking system based on sulphur appears to be helpful, as it promotes the OC dispersion in the polymer matrix.

Results presented in this work seem to arise from the synergy among different aspects. The nanoclay promotes the mechanical reinforcement and its exfoliation allows to establish interaction between the cation bearing long chain alkenyl substituents and silica, leading to lower filler networking. The interaction between the cation and the sulphur based crosslinking system, most likely via amine formation, is likely to promote higher interface crosslinking density. The interest of results here reported is due to the great importance of silica based elastomeric composites, that are extensively used, in tyre compounds, because they allow a substantial reduction of energy dissipation and thus of fuel consumption and carbon footprint, because of the low filler networking phenomenon. The OC/silica nanocomposites studied in this work pave the way for further reduction of such phenomenon and thus for larger diffusion of silica based elastomeric composites.

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