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Confinement Effects at Nanoscale in Natural Rubber Composites: Influence on Macroscopic Properties

Bindu Patanair^{a,b}, Sabu Thomas^a, Abitha V. K^a, Allisson Saiter-Fourcin^{b,*}
a: School of Chemical Sciences, School of Energy Materials and International and Inter-University Centre for Nanoscience and Nanotechnology, Mahatma Gandhi University, Priyadarshini Hills P.O, Kottayam, Kerala, India 686 560
b: Normandie Univ, UNIROUEN, INSA Rouen, CNRS, GPM, 76000 Rouen, France
*Corresponding author. tel: 00332 32 95 50 86 fax: 00332 32 95 50 82
E-mail address: <u>allison.saiter@univ-rouen.fr</u>

ABSTRACT:

The study of molecular level interactions in elastomer composites has got very scant attention even though a large number of studies are going on in this topic. This work embodies the understanding of confinement effects in natural rubber (NR)/ ZnO composites from micro to nano length scales, and their influence on macroscopic properties of the composites. The interactions between fillers and matrix are characterized from the Donth's approach, which allows estimating the cooperativity size N_{α} at the glass transition, N_{α} being directly related to the filler-matrix interactions. The improved properties of NR-nano ZnO composites can be attributed to microstructural and morphological changes due to nano ZnO in the NR matrix. A correlation between N_{α} , the constrained volume C_v and the thickness of immobilized polymer chains χ_m at the glass transition has also been established. Thus, this work proves that the Donth's approach is a powerful probe to estimate the enhance of mechanical properties in nanocomposites from calorimetric investigations.

Keywords: Natural rubber, Composites, Mechanical Properties, Glass transition, Cooperativity length, Confinement, Donth's approach

INTRODUCTION

Elastomers are special kind of amorphous polymers containing soft and flexible chains. They have low modulus and strength compared to other types of polymers, and hence the inclusion of fillers is essential for obtaining substantial improvements in the physical and mechanical properties. The effects of various types of fillers in the micrometer size range such as carbon black, calcium carbonate, glass fibers, talc, expanded graphite (EG) etc. on elastomer matrix properties are widely studied [1]. However, optimal performance of the final micro composite could be achieved at higher filler concentrations. This causes the viscosity enhancement, and thus creates difficulties in the processing. Nanofillers offer an ultimate solution to these problems due to their high specific surface area and hence fewer quantities are required. It has been observed that the addition of just a very small quantity of nano-sized fillers like carbon nanotubes (CNTs), graphene, clay, polyhedral oligomericsilsequioxane (POSS) etc. is sufficient for improving the properties of virgin polymers to a great extent when compared to micro sized fillers [2-4]. Novel engineered materials from natural rubber (NR) with new fibrous nanofillers with improved barrier properties were prepared for use in tire inner-liner applications [5] and the composites of polyethylene (PE) with NR has higher level of biodegradability under environmental conditions [6]. Moreover, the high surface to volume ratio and other superior properties of nanoparticles help the stress transfer from the matrix to the reinforcement, and assume good adhesion at the interface. But still both micro as well as nano composites are not devoid of deterioration in properties and structural defects. This depends on the mode of filler dispersion in the matrices [7-10] and therefore modification of filler surfaces is often practiced prior to compounding.

Confinement effects of elastomer molecules in composite systems can regulate composite properties and thus their applications [10, 11]. The orientation of elastomer chains in the presence of micro as well as nanofillers is one of the key factors affecting mechanical properties. In our previous study [12], we proposed a core-shell morphology model and constrained polymer model to explain the effect of constrained polymer chains in the NR-nano ZnO nanocomposite system. In the current study, the confinement effect due to the size differences of fillers (micro and nano ZnO) will be quantified by focusing on the amorphous phase, and by using the Donth's approach [13] from which it is possible to estimate a cooperative rearranging region (CRR) in terms of structural unit number noted N_{α} . N_{α} represents the number of structural units which move together at the glass transition. Thus, a CRR allows to illustrate the well-known notion of cooperative movement at the glass

transition. N_{α} can be calculated from the equation 1. All the parameters in this equation can be estimated from MT-DSC experiments.

$$N_{\alpha} = \frac{N_{A} \left[\left(\frac{1}{c_{p,glass}(T_{\alpha})} \right) - \left(\frac{1}{c_{p,liquid}(T_{\alpha})} \right) \right]}{M_{o}(\delta T)^{2}} k_{B} T_{\alpha}^{2}$$
(1)

where T_{α} is the glass transition temperature (in K), k_B the Boltzmann constant, C_p the heat capacity at constant pressure (in J/(g.K)), M₀ the molar mass of one structural unit (68.12 g/mol), and δT (in K) the mean temperature fluctuation related to the glass transition temperature of an average CRR. All these quantities are directly extracted from MT-DSC curves. All the details about this extraction are clearly explained in reference [13]. The notion of CRR has been first introduced by Adam and Gibbs [14], which defined this as a subsystem, which can rearrange its configuration to another one independently of its environment. Each CRR presents subsystem having its own glass transition temperature and its own free volume, linked to its own relaxation time. The Donth's approach which allows estimating the CRR size, is based on calorimetric measurements, and has been often used to study the effect of fillers on molecular mobility in nanocomposites [15-20]. It has been proved that this approach allows to characterize the interactions between fillers and matrices, and then to predict if the fillers improve the macroscopic properties for industrial applications. This approach is a powerful way to study and quantify the filler/matrix interactions in composites. As far as we know, this is the first time that this model is applied to quantify the filler/rubber interactions in natural rubber composites, and to understand and better explain why the mechanical properties are enhanced or not.

MATERIALS AND METHODS

The preparation of ZnO nanoparticles, its characterization and its composites were described in detail in ref. [12, 21]. Synthesis of ZnO nanoparticles from zinc chloride and chitosan is a new synthetic approach [22] and in the present procedure we optimized the reaction conditions to facilitate the formation of Zn–chitosan complex and also to increase the yield of the ZnO nanoparticles. Since chitosan is a linear polymine with reactive amino groups and hydroxyl groups, and hence can easily form chelates with transition metal ions. In the present reaction, chitosan acts as a chelating agent for Zn^{2+} ions to form Zn–chitosan complex. This is an ionexchange method to precipitate Zn(OH)₂. By mixing both 5 % solution of ZnCl₂ in 1 % acetic acid (*ie.* 5 g of ZnCl₂ in 100 ml 1 % acetic acid) with 1 % solution of chitosan in 1 % acetic acid and the whole mixture was stirred for 21 h. After this, stoichiometric amount of NaOH (5 %) was added drop wise to the above reaction mixture with constant stirring. The whole mixture was then allowed to digest for 24 h at room temperature. During this time, OH⁻ and Cl⁻ ions were diffused through the medium and white gel-like precipitate of Zn(OH)₂ was formed. This was filtered and washed thoroughly with distilled water to remove unreacted chitosan and other by-product like NaCl. This was then dried at 100 °C and annealed at 550 °C for 4 h. in a muffle furnace to get ZnO nanocrystals. The yield of ZnO nanocrystals obtained by this method is about 90 %. In this modified method, the reaction can be carried out under moderate conditions, yield of the product is good and particles of nanometer size ZnO can be attained.

Preparation of NR/nano ZnO Composites

Natural rubber (NR) (100 g) was masticated in a two-roll mill; after that, 1 g (or 1 phr) of nano ZnO and 2.5 g of co-activator stearic acid were added and again masticated. Milling was continued by the successive addition of the accelerators N-cyclohexylbenzothiazole- 2-sulphenamide (CBS) (0.6 g) and tetramethyl thiuram disulphide (TMTD) (0.1 g), and the vulcanizing agent sulfur (2.5 g). The NR composites with varying weight percentages of nano ZnO, *viz.* 2, and 3 g (or 2 and 3 phr), were also prepared in the same manner. In general practice for comparison, 5 g (or 5 phr) of micro ZnO is used as the standard cure activator for 100g NR instead of nano ZnO. The weights of all other reactants remain intact. All weights are taken in parts per 100 g of rubber (phr).

Curing characteristics of the compounded vulcanizate (prepared composites) were tested by using Prescott moving die Rheometer according to the ASTM D 5283 standard at high pressure and temperature 150° C for 15 min. All samples were cured as mentioned above and these cured samples were used for further analyses.

The nanocomposites, viz. NR-1 phr nano, NR-2 phr nano and NR-3 phr nano were already characterized by SEM, TEM and AFM [12]. Cryo fractured surface of the samples coated with a thin film of carbon was used for SEM analysis. SEM images were taken with the help of a JEOL 840 analyzer with a W-filament, secondary E-T, and a solid-state backscattered electron detector. Cryo-microtome sliced samples were used for TEM analysis. Cryo-microtome sliced samples of thickness about 100–150 nm at -110 °C were prepared with the help of a Leica Ultracut UCT (Leica Microsystems GmbH, Vienna, Austria) instrument with sharp glass knives. The cryo sections were collected in sucrose solution and directly placed on a copper grid of 300 mesh. TEM was performed with a JEOL 2100, Japan instrument, operated at an accelerating voltage of 200 kV.

The surface morphology of rubber nanocomposites was obtained by AFM, images were taken by using a Veeco di Innova AFM microscope. The samples were kept in phase contrast tapping mode with a frequency of 258 kHz and a scan rate of 1 Hz. The AFM measurements were carried out in air at ambient conditions (25 $^{\circ}$ C).

The mechanical properties were studied by using a Zwick 1474 Universal testing machine at room temperature as per ASTM D 412 standard. Dumbbell shaped samples were punched out from compression molded sheets (length \times width \times thickness =120 \times 30 \times 2 mm) along the mill direction using a dumbbell die and were used at a testing cross head speed of 500mm/min with the cell load of 1kN. The software, testXpert is used for data processing.

The dynamic mechanic analysis (**DMA**) was also carried out [12] using NETZSCH DMA 242 machine in the temperature range -110 °C to 110 °C, at a heating rate of 2 °C/min and three different frequencies 10.00, 1.00 and 0.10 Hz. Rectangular test specimens having sample dimensions $60 \times 10 \times 2$ mm (length × width × thickness) were kept in a dual cantilever mode and measurements were performed in air atmosphere. Liquid nitrogen was used as the cooling medium.

Classical (DSC) and modulated temperature differential scanning calorimetry (MT-DSC) experiments were conducted on a thermal analysis instrument heat flow calorimeter (DSC Q100 TA instrument®), coupled with a Refrigerated Cooling System (RCS). The experiments were conducted under inert nitrogen atmosphere with a flow rate of 50 ml/min. The apparatus was calibrated for heat flow, temperature and baseline using standard Tzero technology. For temperature and heat flow calibrations, standards as Indium were used. The calibration of the specific heat capacity was carried out using Sapphire as a reference.

The DSC analyses of all the samples were carried out by cooling from 10 °C down to -90 °C at a cooling rate of 5 °C/min, permitting first to erase the thermal history of the sample (as we started the experiments above the glass transition domain), second to optimize the thermal contact between sample and pan (as the sample is in its rubbery state), and also to follow the temperature ramp in a broad domain. All the MT-DSC analyses were carried out with the following protocol: oscillation amplitude of 1 °C, oscillation period of 120 s and cooling rate of 0.5 °C/min from 10 °C down to -90 °C. This protocol corresponds to the optimal one to characterize the glass transition domain. We could precise that all the calorimetric curves were normalized to the NR weight.

RESULTS AND DISCUSSION

The DSC curves are presented in Figure 1. As usually observed for rubber systems, the glass transition signature from DSC investigations is not clear, even if a change in baseline (baseline step) can be observed around -65°C proving the existence of glass transition region as drawn in Figure 1. However, it is not possible to estimate with good accuracy the parameters characterizing the glass transition from these experiments, i.e. the glass transition temperature T_g and the heat capacity step at T_g noted $\Delta Cp_{(Tg)}$. That is why we performed MT-DSC experiments. The curves obtained by this way are presented in Figures 2 and 3. We may note that the Rev Cp plots correspond to the heat capacity values of the samples. Thus, it is now possible to estimate with a very good accuracy the values characterizing the glass transition domain. All these values are given in Table 1. From the equation 1, it is possible to calculate the CRR size and all the necessary values are extracted from experimental data presented in Figures 2 and 3. We may note that T_a corresponds to the temperature at the peak maxima in Figure 3, and δT is proportional to the width at the middle height of these peaks. Details about δT extraction are given in [13]. The N_a values are given in Table 1.

Concerning the variations in T_{α} , we clearly observed an increase for the micro-composite and for all the nanocomposites. In this case, the molecular mobility at the glass transition is alleviated by the presence of filler in comparison to the neat NR sample. $\Delta C_p(T_{\alpha})$ values being quasi constant, we can conclude the amorphous content remains constant whatever the content and the filler nature.



Figure 1.Heat Flow curves obtained by classical DSC for neat NR and the different composites



Figure 2. Reversing Cp curves obtained by MT-DSC experiments for neat NR and the different composites



Figure 3. Derivative of the Reversing Cp curves obtained by MT-DSC experiments for neat NR and the different composites

Table 1. Values of the glass transition temperature T_{α} , the heat capacity step at T_{α} $\Delta Cp_{(T\alpha)}$, the number of structural units per CRR N_{α} , the constrained volume C_{ν} , and the thickness of immobilized polymer layer at the interface χ_m for neat NR and the different

composites

samples	$T_{\alpha} (^{\circ}C)$	$\Delta Cp_{(T\alpha)}(J/(g.^{\circ}C))$	Να	C _v (%) [12]	χ _m (nm)
Neat NR	-63.5 ± 0.5	0.39 ± 0.02	188	0.0	0.0
NR-5phr micro	-61.5 ± 0.5	0.35 ± 0.02	119		0.058
NR-1 phr nano	-61.5 ± 0.5	$0.37{\pm}0.02$	112	9.5	0.042
NR-2 phr nano	-62.0 ± 0.5	0.32 ± 0.02	105	17.6	0.163
NR-3 phr nano	-61.0 ± 0.5	0.34 ± 0.02		4.8	0.102

aamnlaa	Tensile strength	Tensile Modulus	Elongation at	
samples	(MPa)	(MPa) 300%	the break (%)	
Neat NR	2.04	0.47	801	
NR-1 phr nano	18.17	0.87	1403	
NR-2phr nano	25.56	2.28	637	
NR-3phr nano	6.68	1.37	873	

 Table 2. Tensile Strength, Tensile Modulus, Elongation at the Break of rubber nanocomposites [12]

Table 2 gives the evolution of the mechanical properties (Tensile strength, Tensile Modulus, Elongation at the break) for nanocomposites in comparison to neat NR sample. The tensile strength and tensile modulus increase with the addition of the nano ZnO from 1 to 2 phr, and decrease on further addition (i.e. 3 phr) of nano ZnO; whereas the elongation at the break of rubber nanocomposites decreases from 1 to 2 phr of nano ZnO, and increases on further addition (i.e. 3 phr) of nano ZnO. A comparison of the reported mechanical properties of the NR-micro ZnO filled composite [23] with that of NR-2 phr nanocomposite shows that about 27% enhancement in the tensile strength, 78% enhancement in the modulus (300%), and 48% decrease in the elongation at the break were observed for the NR-2 phr nanocomposite. The decrease in the value of tensile strength and tensile modulus for NR-3 phr nanocomposite is due to the aggregation of nanoparticles at higher nanofiller concentration as evidenced from TEM images (Figures 4 a,b) [12]. The enhancement of mechanical properties of nanocomposites can be attributed to the homogeneous dispersion of nano ZnO, and strong nanofiller-rubber matrix interactions exist in them. For NR-3 phr nanocomposite, filler-filler interaction is more pronounced than nanofiller-rubber matrix interactions.



Figure. 4(a) TEM image of NR-2 phr nano (b)TEM image of NR-3 phr nano [12]

Considering the CRR size evolution, the value obtained for the micro composite (i.e. NR-5 phr micro) in comparison with neat NR proves the decrease of the cooperativity due to the presence of filler (N_{α} significantly decreases from 188 down to 119), despite the less increase in terms of glass transition temperature (from -63,5°C up to -61,5°C). Indeed with 5 phr of microfiller, there are greater interactions between filler and matrix inducing the CRR decrease, but the presence of microfiller hampers the chain mobility, involving a low T_{α} increase. Concerning the nanocomposites, as for NR-3 phr nanocomposite aggregations of nanoparticles have been evidenced from TEM images (Figure 4b), we only compare the evolution between neat NR and nanocomposites up to 2% of filler. We may note the higher the wt. % of nanofiller, the lower

the CRR size, proving that the filler-matrix interactions are more and more pronounced. From the CRR size values, it seems the best nanocomposite in terms of filler/matrix interactions is NR-2 phr nano. As reported in our previous work, ZnO acts as a cure activator in rubber compounds: instead of 5 phr of micro ZnO, 2 phr of nano ZnO is sufficient to bring the properties of 5 phr of micro ZnO in NR composites [12]. Furthermore, the best mechanical properties are observed for 2 phr of nano ZnO sample due to the stronger interactions between filler and matrix. Then, the predictions related to the CRR size seems to be in good agreement with the previous work [12].

DMA was performed to investigate the nature of the constrained polymer region in NR matrix. The segmental motion of rubber chains is progressively restricted when nano ZnO particles are homogeneously dispersed in the rubber matrix and hence the volume of the constrained rubber chains (C_v) also increases as in the case of NR-1 phr nanocomposite and NR-2 phr nanocomposite. The constrained polymer region is the significant factor which establishes the reinforcement mechanism in NR nanocomposites. Moreover, as shown in Table 1, the constrained volume values noted C_v and calculated from DMA investigations [12] are inversely proportional to the CRR size for NR-1 phr nanocomposite and NR-2 phr nanocomposite, proving the correlation of the CRR size evolution with macroscopic properties like mechanical ones.

It is well established that the presence of nanoparticles in a polymer network would create entangled or constrained state of the polymer chains. More precisely, when nanoparticles are incorporated in amorphous polymers viz. elastomers, the mobile amorphous chains in the original polymer are changed into two components *viz*. mobile and rigid amorphous components (Schemes 1a, b). Several research reports revealed that the presence of a very thin layer (a few nanometers of thickness) of interfacial polymer with restricted mobility is detected around the filler particles, while the rest of the polymer follows bulk dynamics [24].



Scheme 1a. Formation of rigid amorphous fractions in amorphous polymers: (a) before filler and (b) after the addition of nanofiller



Scheme 1b. Immobilized polymer chains on the surface of ZnO nanoparticles forming a rigid amorphous part

This aspect has been well explained in various models *viz*. three-layer models [25], two-layer models [24, 26] or a continuous distribution of glass transition temperatures as a function of distance from the particle surface [27]. The nature and behavior of the polymer-filler interface can create a major influence on the macroscopic properties. Moreover, studies revealed that different segmental mobility have been observed close to the filler surface, where segmental relaxation times are significantly longer than those of the unfilled polymer [28].

But in nanocomposites due to strong polymer-filler interactions, the amorphous content is supposed to change to mobile and rigid amorphous portions [29]. In the present case, NR has no affinity towards ZnO, unless the presence of a coupling agent; it acts as a cure activator in the sulphur vulcanization and curing process [12]. The interaction of the polymer chains with the filler particles surface is a kind of secondary interactions *viz*. weak Van der Waals forces [12]. Based on the above reports, it was understood that the polymer chains are immobilized in the rigid amorphous content by the filler surface. So, Δ Cp measurements can be used as a tool to predict the rigid amorphous content of the nanocomposites [20, 30].

Figure 5 presents the variations of the Δ Cp or normalized heat capacity Δ Cp/ Δ Cp⁰ with wt. % of ZnO, where Δ Cp⁰ is the heat capacity of the NR, and Δ Cp the heat capacity value of each nanocomposite given in Table 1. The Δ Cp or normalised Δ Cp values show a drastic decrease from the reference line (additivity line) indicating strong polymer-filler interaction. The Δ Cp or normalized heat capacity Δ Cp/ Δ Cp⁰ value decreases with wt. % of ZnO and this can be explained in terms of the formation of an immobilized layer of polymer around the filler particles. The fraction of immobilized polymer can then be calculated by the following equation:

$$\chi_{\rm m} = 1 - \frac{\Delta C_{\rm p}}{\Delta C_{\rm p}^{\rm 0} \left(1 - W_{\rm ZnO}\right)} \tag{2}$$

Where W_{ZnO} is the weight fraction of ZnO. The accuracy obtained on Δ Cp measurements thanks to MT-DSC investigations allows us to estimate the fraction of immobilized polymer χ_m with a very good accuracy. The values obtained are given in Table 1. The thickness of immobilized polymer layer increases with wt. % of ZnO, it is revealed that the strongest rubbernano ZnO interactions are present in NR-2 phr nano ZnO composite. The reason for the enhanced mechanical properties is the strong rubber-nano ZnO interactions, proved from the calculated values of C_v, χ_m and CRR size. As NR-3 phr nano ZnO nanocomposite exhibits agglomerated filler particles in the matrix (see Figure 4b) a lower value of constrained volume

 C_v is obtained (Table 1), which establishes that more filler-filler interactions are pronounced in the composite than filler-polymer interactions.



Figure 5. ΔC_p as a function of wt.% of ZnO; insert: Normalised ΔC_p as a function of wt.% of ZnO

CONCLUSIONS

In this work, the glass transition characteristics of NR composites, i.e. the glass transition temperature and the heat capacity step at this temperature, have been obtained with a very good accuracy through MT-DSC investigations in comparison to the classical DSC experiments. This work highlights the interest of using the Donth's approach as a probe for understanding the chain mobility in rubber systems. Indeed, we proved a good correlation between the CRR size evolution, and the evolution of the mechanical properties as a function of filler content: the lower the CRR size, the higher the fraction of immobilized polymer chains and stronger interactions between fillers and matrix, and hence the enhancement in the mechanical properties. Furthermore, the evolution of the CRR size at the glass transition in terms of number (N_{α}) is consistent with the evolution of the constrained volume (C_v) and the thickness of immobilized polymer chains at the interface (χ_m) measured at the same temperature for nanocomposites with homogeneous dispersion of ZnO nanoparticles: the lower the N_{α} value,

the higher the C_v and χ_m values. Thus, the Donth's approach is a powerful probe to estimate the enhancement of mechanical properties in nanocomposites from calorimetric investigations.

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