



Imaging dielectric relaxation in nanostructured polymers by frequency modulation electrostatic force microscopy

Clément Riedel, R. Sweeney, N. E. Israeloff, Richard Arinero, Gustavo Ariel Schwartz, Angel Alegría, Philippe Tordjeman, Juan Colmenero

► To cite this version:

Clément Riedel, R. Sweeney, N. E. Israeloff, Richard Arinero, Gustavo Ariel Schwartz, et al.. Imaging dielectric relaxation in nanostructured polymers by frequency modulation electrostatic force microscopy. *Applied Physics Letters*, 2010, vol. 96, 10.1063/1.3431288 . hal-00909650

HAL Id: hal-00909650

<https://hal.science/hal-00909650>

Submitted on 26 Nov 2013

HAL is a multi-disciplinary open access archive for the deposit and dissemination of scientific research documents, whether they are published or not. The documents may come from teaching and research institutions in France or abroad, or from public or private research centers.

L'archive ouverte pluridisciplinaire **HAL**, est destinée au dépôt et à la diffusion de documents scientifiques de niveau recherche, publiés ou non, émanant des établissements d'enseignement et de recherche français ou étrangers, des laboratoires publics ou privés.



Open Archive TOULOUSE Archive Ouverte (OATAO)

OATAO is an open access repository that collects the work of Toulouse researchers and makes it freely available over the web where possible.

This is an author-deposited version published in : <http://oatao.univ-toulouse.fr/>
Eprints ID : 10270

To link to this article : doi:10.1063/1.3431288
URL : <http://dx.doi.org/10.1063/1.3431288>

To cite this version : Riedel, Clément and Sweeney, R. and Israeloff, N. E. and Arinero, Richard and Schwartz, Gustavo Ariel and Alegría, Angel and Tordjeman, Philippe and Colmenero, Juan Imaging dielectric relaxation in nanostructured polymers by frequency modulation electrostatic force microscopy. (2010) Applied Physics Letters, vol. 96 (n° 21). ISSN 0003-6951
--

Any correspondence concerning this service should be sent to the repository administrator: staff-oatao@listes-diff.inp-toulouse.fr

Imaging dielectric relaxation in nanostructured polymers by frequency modulation electrostatic force microscopy

C. Riedel,^{1,2,a)} R. Sweeney,³ N. E. Israeloff,³ R. Arinero,¹ G. A. Schwartz,⁴ A. Alegria,² Ph. Tordjeman,⁵ and J. Colmenero^{2,4}

¹*Institut d'Electronique du Sud (IES), UMR CNRS 5214, Université Montpellier II, CC 082, Place E. Bataillon, 34095 Montpellier Cedex, France*

²*Departamento de Física de Materiales, Facultad de Química, UPV/EHU, Apartado 1072, 20080 San Sebastián, Spain and Donostia International Physics Center, Paseo Manuel de Lardizabal 4, 20018 San Sebastián, Spain*

³*Department of Physics, Northeastern University, Boston, Massachusetts 02115, USA*

⁴*Centro de Física de Materiales CSIC-UPV/EHU, Edificio Korta, 20018 San Sebastián, Spain*

⁵*Institut de Mécanique des Fluides (IMFT), Université de Toulouse–CNRS, 1 Allée du Professeur Camille Soula, 31400 Toulouse, France*

We have developed a method for imaging the temperature-frequency dependence of the dynamics of nanostructured polymer films with spatial resolution. This method provides images with dielectric compositional contrast well decoupled from topography. Using frequency-modulation electrostatic-force-microscopy, we probe the local frequency-dependent (0.1–100 Hz) dielectric response through measurement of the amplitude and phase of the force gradient in response to an oscillating applied electric field. When the phase is imaged at fixed frequency, it reveals the spatial variation in dielectric losses, i.e., the spatial variation in molecular/dipolar dynamics, with 40 nm lateral resolution. This is demonstrated by using as a model system; a phase separated polystyrene/polyvinyl-acetate (PVAc) blend. We show that nanoscale dynamic domains of PVAc are clearly identifiable in phase images as those which light-up in a band of temperature, reflecting the variations in the molecular/dipolar dynamics approaching the glass transition temperature of PVAc.

Broadband dielectric spectroscopy (BDS) is a well established technique that enables measurement of molecular dynamics of bulk materials with permanent molecular dipoles. Recent advances have made it possible to probe the dynamics of thin polymer films with a thicknesses as low as 5 nm, enabling the study of interfaces.^{1,2} However, BDS has poor spatial resolution and the interpretation of the results from nanostructured polymers is always model dependent.³ Concerning local dielectric characterization, on the one hand, various atomic force microscopy based methods have been developed to image the dc dielectric constant ϵ at nanoscale. Fumagalli *et al.*⁴ have developed the so-called “nanoscale capacitance microscopy,” where the microscope is equipped with a subattofarad low-frequency capacitance detector. The same group also proposed a method based on the detection of the dc electrostatic force to image the dielectric constant of a purple membrane patch.⁵ By measuring the dc electrostatic force gradient, we have been able to obtain a quantitative value of ϵ in one point⁶ and a map of the dielectric constant of a nanostructured polymer film.⁷ On the other hand, in previous work we have shown that the frequency response of the real and imaginary parts of the dielectric constant [$\epsilon^*(\omega) = \epsilon' - i\epsilon''$] can be investigated using frequency modulation electrostatic force microscopy (FM-EFM).^{8,9} Measurements made in one spatial point gives direct information about the local dynamics of the sample area probed by the tip sample interaction. In the following, we will detail how we have been able to combine imaging

and measurement of the frequency dependence of the dielectric response to map the dipolar/molecular dynamics of a nanostructured thin film of a phase separated polymer blend.

In order to maximize contrast, we have chosen to work with polystyrene (PS-[C₈H₈]_n—M_w=70 950 g/mol) and poly(vinyl-acetate) (PVAc-[C₄H₆O₂]_n—M_w=33 200 g/mol) for which the bulk dielectric response has been previously well characterized.^{10–12} A solution containing 25% PVAc and 75% PS (by mass) and having a total concentration of 1% polymer in toluene has been spin-coated over a conductive film (Au) on a mica substrate. The samples are dried at 70 °C for 1 h under ambient conditions and annealed at 35 °C overnight in vacuum to remove any trace of solvent. The resulting polymer films consist of a 25 nm thick matrix with island-like features having an average diameter of about 300 nm and a maximum height of about 25 nm above the reference level of the matrix (Fig. 1). In the investigated range of frequency (0.1–100 Hz), we expect to detect the so-called dielectric alpha-relaxation of PVAc (correlated with segmental motion¹²) slightly above its glass transition temperature; T_g~35 °C. Therefore we have explored the temperature range between 25 and 50 °C in order to follow the evolution of this dynamic process. The T_g of PS being ~105 °C, no measurable relaxation is expected in our range of temperature and frequency. These two polymers are immiscible, even at the nanoscale and form phase separated domains.

Experiments were carried out by FM-EFM under ultrahigh vacuum (UHV) with a variable temperature stage (RHK UHV 350). The temperature was measured with a small thermocouple clamped to the sample surface.

^{a)} Author to whom correspondence should be addressed. Electronic mail: riedel@ies.univ-montp2.fr.

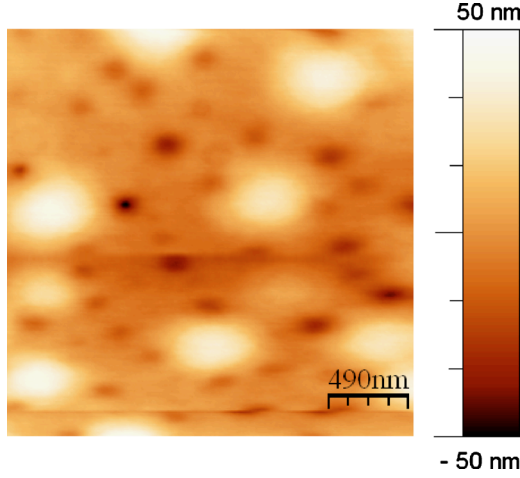


FIG. 1. (Color online) Topography of the thin PS/PVAc polymer film. The thickness of the matrix is approximately 25 nm, whereas islands rise to 50 nm from the substrate.

Nanosensors ATEC EFM cantilevers with 4:1 aspect ratio and 25 nm tip radius, spring-constant $k=2$ N/m, are used so that V_{dc} and V_{ac} bias voltages can be applied to the tip. In FM-EFM, the cantilever is oscillated at its resonance frequency, $f_0=70$ kHz, and the resonance frequency shift, δf , due to tip-sample interaction forces, is detected with very high resolution using a Nanosurf Easy-PLL Phase-Locked-Loop detector. This parameter is used as a feedback for controlling the tip-sample distance z and allows us to obtain the topography as shown in Fig. 1. Additional signals are those measured as a result of the electrodynamic interaction between the conducting tip and the polymer sample. As the force gradient dF/dz is proportional to the square of the voltage, a sinusoidal voltage $V_{ac}=V_0 \sin(\omega t)$ produces a 2ω component. This component can be detected as a modulation in the δf signal using a lock-in amplifier. The obtained $V_{2\omega}$ signal is related to dF/dz by the following: $V_{2\omega}=(A/4k) \times (df/dz)f_0$, where A is the instrument gain.⁸ As the force gradient is a function of the dielectric permittivity,⁶⁻⁹ any measurable dielectric losses in the material will be detected as a phase shift in the $V_{2\omega}$ signal.

Two different experiments were performed to study successively the thin polymer blend. The first is similar to that describe in Refs. 8 and 9 and consisted in recording the phase in one point at different frequencies and constant temperatures. During the acquisition of the dielectric spectrum, the feedback of the PLL detector (typically $\delta f_0=15$ Hz) was turned off in order to avoid any coupling with the corresponding electrical frequency. We apply a $V_{ac}=V_0 \sin(\omega t)$ voltage at different frequencies ($\omega/2\pi$) varying from 0.1 to 120 Hz. The frequency range is limited at low frequencies only by the time of acquisition and at high frequencies by the bandwidth of the FM electronics. A preliminary step consists of measuring the reference phase of $V_{2\omega}$, Φ_{ref} , at 19 °C where both polymers are in the glassy state and have no measurable dielectric relaxation. The difference between this reference phase and the phase measured in the experiments, $\Delta\Phi=\Phi_{ref}-\Phi$, is used to characterize the dielectric dissipation. During the relaxation, due to the loss of energy to align the molecular dipoles in the field, a peak is observed in $\Delta\Phi(\omega)$. Using BDS experiments on bulk PVAc, the macroscopic phase of the dielectric permittivity, $\text{invtan}(\epsilon''/\epsilon')$,

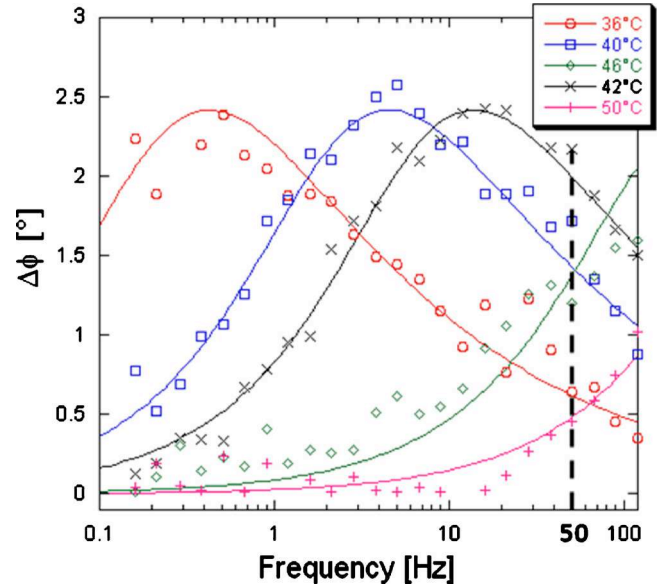


FIG. 2. (Color online) Phase shift measured as a function of the frequency over an island of PVAc. Statistical error is 0.25 for $f > 1$ Hz and 0.35 for $f \leq 1$ Hz. Lines are fits derived from a KWW model with amplitude and shape parameters fixed for all curves. The dashed line indicates the imaging frequency of Fig. 3.

peaks at about 16° for frequencies of 0.1–100 Hz, similar to what was found for $\Delta\Phi(\omega)$ of thick films ($1 \mu\text{m}$) of PVAc using the present FM-EFM technique.⁸ For thinner films of PVAc, we found that the peak in $\Delta\Phi(\omega)$ is considerably reduced, to $\sim 3^\circ$ for 50 nm films. This can be understood by recognizing that the force gradient is a complex function of the dielectric permittivity, film thickness, tip radius, and tip height that requires numerical modeling.⁷ Whereas the macroscopic phase is proportional to fractional increase in permittivity, $\Delta\epsilon/\epsilon$, upon warming through the glass transition, the $\Delta\Phi(\omega)$ we measure here is proportional to the fractional increase in force gradient, $\Delta(dF/dz)/(dF/dz)$. This fraction is found in simulations to be reduced with reduced film thickness due to a relatively larger constant background dF/dz .

Figure 2 shows $\Delta\Phi(\omega)$ over an island at different temperatures. The fitting lines were obtained using a Kohlrausch–Williams–Watts response function, $\exp(-t/\tau)^\beta$,^{13,14} as a convenient way to describe the experimental result. Maximum value of $\Delta\Phi$ and shape parameter ($\beta=0.42$) were fixed by fitting the data measured at 40 °C. Keeping these parameters constant and using only a free timescale parameter we obtained a rather satisfactory description of the data, i.e., the shape of the peak does not depend on temperature within the uncertainties. The peak frequencies and shape of these peaks are nearly identical to those found in pure PVAc thick films. Measurements at the same temperatures have been performed over the matrix and the response (not shown) is always flat ($\Delta\Phi < 0.1^\circ$). Therefore, the frequency spectrum permits us to characterize locally the dynamical properties of the sample, which shows that the matrix is composed of PS whereas islands are dynamic domains of PVAc, in agreement with Ref. 7 which imaged static dielectric response.

The second experiment involved scanning the sample surface while a 50 Hz V_{ac} is applied to the tip. In order to keep a constant tip-sample distance, the feedback of the PLL

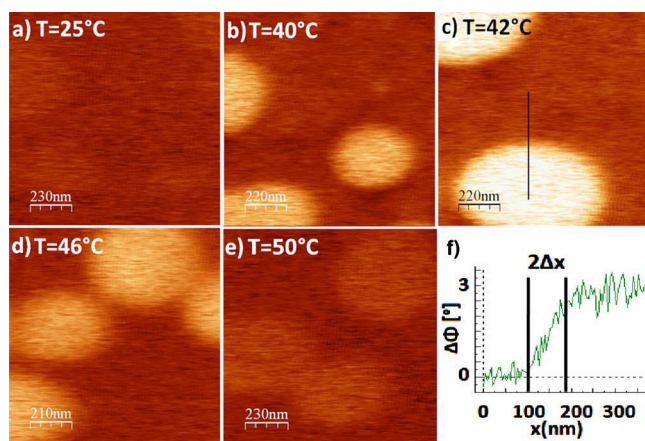


FIG. 3. (Color online) Image of the phase shift recorded at 50 Hz on a PS/PVAc film for various temperatures. Brighter areas correspond to higher losses, indicative of segmental relaxation on ~ 3 ms time scales in PVAc. The profile recorded on the black line at 42 °C is shown in (f). The phase jump at the PS/PVAc boundary gives an estimated lateral resolution, $\Delta x \sim 40$ nm

is used during the scan. Figure 3 presents images of $\Delta\Phi$ recorded at different temperatures. All images have been off-set to the same phase-shift color scale using WSXM.¹⁵ The phase recorded on the flat matrix areas is always near zero, as expected for PS. On the other hand all values measured on the PVAc islands are, within experimental errors, the same as those measured using the frequency sweep in one single point at 50 Hz (Fig. 2). At 25 °C, the image is homogeneous; there is no dielectric relaxation occurring in the islands of PVAc. This demonstrates that this method is only sensitive to dipolar relaxation and not topographical effects. When the temperature is increased to 36 °C we begin to see strong contrast in the images as the high frequency wing of the dielectric loss peak starts to contribute at 50 Hz in the PVAc regions. The maximum contrast is found at 42 °C. Further increasing the temperature reduces the contrast as the loss peak shifts to higher frequencies and only the lower frequency contributions are detected. As the shape of the relaxation does not depend significantly on temperature, the contrast observed is only attributed to the passage of dynamics associated with the alpha-relaxation through our measurement window. Figure 3(f) shows the profile of the phase recorded at 42 °C across the interface between a PVAc island and the PS matrix [as shown in Fig. 3(c)]. We can estimate an upper limit of the spatial resolution (corresponding to half of the distance necessary to achieve the transition between the dielectric level of PVAc and PS) Δx around ~ 40 nm. This corresponds to the state-of-the-art resolution in EFM imaging when using standard probes, and it is also close to the theoretical resolution under our working conditions.^{16,17}

In summary, we have developed a method that permits imaging the dynamics of nanostructured polymers with nanometer spatial resolution. This method of imaging dielectric losses is well decoupled from topography and can provide images with dielectric compositional contrast. For PS/PVAc polymer blends, strong phase separation was found. The glassy dynamics of the raised island-like domains was found

to be nearly identical to that of pure PVAc films of similar thickness, whereas between these domains no dielectric response was found, consistent with pure PS behavior. Similar results were found on thicker films of PS/PVAc, though weak PVAc-like response was found in the PS regions indicative of possible buried PVAc domains. With further numerical modeling it may be possible to use this technique to analyze the three-dimensional structure of such films. The method described here makes it possible to study dielectric relaxation and molecular dynamics with nanometer scale spatial resolution in nanostructured polymers, nanocomposites, and other multicomponent soft materials, addressing questions of utmost importance such as the role of interfacial dynamics on the properties of such systems. Extending this method to ambient conditions would enable the study of soft hydrated materials such as proteins, and cell membranes, and investigate processes such as membrane potential formation, action potential propagation, or ion membrane transport.¹⁸ Work in this direction is in progress.

This work was supported by the Donostia Internacional Physics Center (DIPC), the Basque Country Government (Ref. No. IT-436-07, Depto. Educación, Universidades e Investigación), the Spanish Ministry of Science and Innovation (Grant No. MAT 2007-63681), European Community (SOFTCOMP program), and the National Science Foundation (Grant No. NSF DMR-0606090). C.R. and R.S. have equally contributed to the work.

¹A. Serghei, M. Tress, and F. Kremer, *Macromolecules* **39**, 9385 (2006).

²S. Napolitano, V. Lupascu, and M. Wübbenhorst, *Macromolecules* **41**, 1061 (2008).

³R. Lund, L. Willner, A. Alegría, J. Colmenero, and D. Richter, *Macromolecules* **41**, 511 (2008).

⁴L. Fumagalli, G. Ferrari, M. Sampietro, and G. Gomila, *Nano Lett.* **9**, 1604 (2009).

⁵G. Gramse, I. Casuso, J. Toset, L. Fumagalli, and G. Gomila, *Nanotechnology* **20**, 395702 (2009).

⁶C. Riedel, R. Arinero, P. Tordjeman, M. Ramonda, G. Lévêque, G. A. Schwartz, D. G. de Oteya, A. Alegría, and J. Colmenero, *J. Appl. Phys.* **106**, 024315 (2009).

⁷C. Riedel, R. Arinero, P. Tordjeman, G. Lévêque, G. A. Schwartz, A. Alegría, and J. Colmenero, *Phys. Rev. E* **81**, 010801(R) (2010).

⁸P. S. Crider, M. R. Majewski, J. Zhang, H. Oukris, and N. E. Israeloff, *Appl. Phys. Lett.* **91**, 013102 (2007).

⁹P. S. Crider, M. R. Majewski, J. Zhang, H. Oukris, and N. E. Israeloff, *J. Chem. Phys.* **128**, 044908 (2008).

¹⁰G. A. Schwartz, E. Tellechea, J. Colmenero, and A. Alegría, *J. Non-Cryst. Solids* **351**, 2616 (2005).

¹¹G. A. Schwartz, J. Colmenero, and A. Alegría, *J. Non-Cryst. Solids* **353**, 4298 (2007).

¹²M. Tyagi, A. Alegría, and J. Colmenero, *J. Chem. Phys.* **122**, 244909 (2005).

¹³F. Kohlrausch, *Pogg. Ann. Phys.* **119**, 352 (1863).

¹⁴G. Williams and D. C. Watts, *Trans. Faraday Soc.* **66**, 80 (1970).

¹⁵I. Horcas, R. Fernandez, J. M. Gomez-Rodriguez, J. Colchero, J. Gomez-Herrero, and A. M. Baro, *Rev. Sci. Instrum.* **78**, 013705 (2007).

¹⁶S. Gómez-Moñivas, L. S. Froufe, R. Carminati, J. J. Greffet, and J. J. Saenz, *Nanotechnology* **12**, 496 (2001).

¹⁷B. Bhushan and H. Fuchs, *Applied Scanning Probe Methods II* (Springer, New York, 2003), p. 312.

¹⁸J. Malmivuo and J. R. Plonsey, *Bioelectromagnetism: Principles and Applications of Bioelectric and Biomagnetic Field* (Oxford University Press, New York, 1995).