

ACCEPTED MANUSCRIPT

Efficient second harmonic generation by para-nitroaniline embedded in electro-spun polymeric nanofibres

To cite this article before publication: Hugo Gonçalves *et al* 2018 *J. Phys. D: Appl. Phys.* in press <https://doi.org/10.1088/1361-6463/aaa9da>

Manuscript version: Accepted Manuscript

Accepted Manuscript is “the version of the article accepted for publication including all changes made as a result of the peer review process, and which may also include the addition to the article by IOP Publishing of a header, an article ID, a cover sheet and/or an ‘Accepted Manuscript’ watermark, but excluding any other editing, typesetting or other changes made by IOP Publishing and/or its licensors”

This Accepted Manuscript is © 2018 IOP Publishing Ltd.

During the embargo period (the 12 month period from the publication of the Version of Record of this article), the Accepted Manuscript is fully protected by copyright and cannot be reused or reposted elsewhere.

As the Version of Record of this article is going to be / has been published on a subscription basis, this Accepted Manuscript is available for reuse under a CC BY-NC-ND 3.0 licence after the 12 month embargo period.

After the embargo period, everyone is permitted to use copy and redistribute this article for non-commercial purposes only, provided that they adhere to all the terms of the licence <https://creativecommons.org/licenses/by-nc-nd/3.0>

Although reasonable endeavours have been taken to obtain all necessary permissions from third parties to include their copyrighted content within this article, their full citation and copyright line may not be present in this Accepted Manuscript version. Before using any content from this article, please refer to the Version of Record on IOPscience once published for full citation and copyright details, as permissions will likely be required. All third party content is fully copyright protected, unless specifically stated otherwise in the figure caption in the Version of Record.

View the [article online](#) for updates and enhancements.

1
2
3
4
5
6
7
8
9
10 Efficient second harmonic generation by *para*-nitroaniline
11
12
13 embedded in electro-spun polymeric nanofibres
14
15
16
17

18 **Hugo Gonçalves¹, Inês Saavedra¹, Rute AS Ferreira², PE Lopes³, Etelvina de**
19 **Matos Gomes¹, Michael Belsley¹**

20
21
22 ¹ Centre of Physics, University of Minho, Campus de Gualtar, 4710-057 Braga,
23 Portugal

24
25 ² Department of Physics, CICECO – Aveiro Institute of Materials, University of
26 Aveiro, 3810-193 Aveiro, Portugal

27
28 ³ Institute for Polymers and Composites/i3N, University of Minho, Guimarães,
29 Portugal

30 E-mail: belsley@fisica.uminho.pt
31
32
33
34
35

36 **Abstract:** Intense well polarized second harmonic light was generated by
37 Poly(methyl methacrylate) nanofibres with embedded *para*-Nitroaniline
38 nanocrystals. Subwavelength diameter fibres were electro-spun using a 1:2 weight
39 ratio of chromophore to polymer. Analysis of the generated second harmonic
40 light indicates that the *para*-Nitroaniline molecules, which nominally crystallize in
41 the centrosymmetric space group, were organized into noncentrosymmetric
42 structures leading to a second order susceptibility dominated by a single tensor
43 element. Under the best deposition conditions, the nanofibers display an effective
44 nonlinear optical susceptibility approximately two orders of magnitude greater
45 than that of potassium dihydrogen phosphate. Generalizing this approach to a
46 broad range of organic molecules with strong individual molecular second order
47 nonlinear responses, but which nominally form centrosymmetric organic crystals,
48 could open a new pathway for the fabrication of efficient sub-micron sized second
49 harmonic light generators.
50
51
52
53

54 **PACS:** **42.65.-k** Nonlinear optics; **42.65.An** Optical susceptibility; hyperpolarizability,
55 **42.65.Ky** Frequency conversion; harmonic generation; **42.70.Jk** Polymers and organics
56
57
58
59
60

1. Introduction

New materials with tailored linear and nonlinear optical (NLO) properties are actively being sought to extend the frontiers that limit many photonic devices. Currently the majority of commercial nonlinear photonic components are based on inorganic materials [1]. Despite the successful development of many new inorganic crystalline materials over the past several decades, the applications of these materials is often limited by the magnitude of the available optical nonlinearities as well as the associated fabrication costs. In particular, efficient generation of second harmonic light by sub-micron sized systems is not routinely available from inorganic materials. Recently, several different processing approaches to extend the range of available nonlinear optical materials have been explored, including self-assembly [2,3], poling [4], lithography [5,6], molecular beam epitaxy [7] and electrospinning deposition [8-12]. Concurrently, researchers have developed a variety of alternative building blocks such as metallic nanoparticles [13], semi-conductors quantum dots [14], 2D-Materials [15] and tailored organic molecules [16].

In particular, organic systems have been widely investigated for nonlinear optical applications. They possess several attractive features including their low cost, fast, large nonlinear responses over a broad frequency range and tailorability [4,16]. A paradigmatic example is the organic molecule of *para*-Nitroaniline (*p*NA) [17-20]. It has a delocalized π -electron system and unsaturated bridge linking a donor amino group (NH_2) and an acceptor nitro group (NO_2). The delocalization of the π -electrons leads to an asymmetric charge distribution resulting in a strong molecular hyperpolarizability (β) [21]. Unfortunately, because it crystallizes in a centrosymmetric space group ($P2_1/n$) [17], the bulk second order nonlinear response of *p*NA crystals vanishes. This commonly occurs in donor- π bridge-acceptor organic crystals: the dominant dipole-dipole interactions result in a pairwise, side-by-side antiparallel molecular alignment within the unit cell [22]. Experimentally, several different approaches have sought to achieve useful SHG when employing *p*NA as the active molecular building block [10,18,20,23]. These range from modifying the surface of *p*NA crystals [18] to the incorporation of the organic dye within carbon nanotube structures [24].

Recently we reported observing strong SHG from *p*NA embedded in electro-spun poly(L-lactic acid) (PLLA) polymer nanofibres [10]. Here we report on similar results obtained using Poly(methyl methacrylate) (PMMA) as the polymeric host. PLLA, being a chiral polymer, induces a higher degree of orientation alignment to the *p*NA nanocrystals within the electrospun fibres. The intensity of the dominant x-ray diffraction peak corresponding to the [202] plane relative to that of the [111] or [311] planes in PLLA fibres is more than four times greater than what we observe for the PMMA fibres. Nonetheless, as

we will show, the effective second order nonlinear constant, d_{eff} is roughly the same for *p*NA embedded electrospun PLLA and PMMA fibres. However, contrary to the situation with PLLA, the *p*NA-PMMA fibres produce an intense well-polarized second harmonic response dominated by a single tensor element of the effective second order susceptibility.

Electrospinning deposition is a versatile technique with deposition parameters that can be tuned to induce long range molecular order organic molecular dopants, leading to enhanced optical and NLO coefficients [13,25]. The sub wavelength size of the resultant hybrid nanostructures promotes localized coherent interactions in the transverse direction and allows as well as wave-guiding effects along the longitudinal direction [26]. Furthermore, the technique is relatively low cost and can be easily scaled-up to an industrial level [9].

In this study, *p*NA molecules were embedded in a one dimensional (1D) nanostructured PMMA matrix, using different deposition rates during the electrospinning process. With this technique, the growth of the *p*NA crystals is encouraged along the longitudinal axis. Strong coherent SHG radiation was produced when 100fs pulses from a mode-locked Ti:Sapphire laser operating at a central wavelength of 800nm were focused onto these nanofibres. The polarization dependence of the SHG generated by these fibres allows us to comment on how the electrospinning deposition rate influences the degree of *p*NA alignment with the fibres. The complementary characterization techniques of Scanning Electron Microscope (SEM), X-ray diffraction, Raman spectroscopy, Fluorescence Lifetime Imaging Microscopy (FLIM) and Hyperspectral Microscopy were employed to obtain insights regarding the local nanostructured environment of the embedded *p*NA molecules.

2. Methods

2.1 Materials

Para-nitroaniline (*p*NA) was purchased from Sigma Chemical Co and used as received. Poly(methyl methacrylate) (PMMA, Mw 120000) was purchased from Alfa Aesar. The raw materials were dissolved in dimethyl formamide (DMF) and toluene at a 1:2 volume ratio. DMF ($\text{C}_3\text{H}_7\text{NO}$, 99.8%) and toluene ($\text{C}_6\text{H}_5\text{CH}_3$, 99.8%) were purchased from Sigma-Aldrich as used as received. The relative weight ratio of *p*NA to PMMA in the final precursor solution was 1:2. The prepared solutions were stirred for several hours under ambient conditions prior to deposition by electrospinning.

2.2 Nanofibre Production

The polymeric nanofibres were electro-spun. The precursor solution was loaded into a syringe with its needle connected to the anode of a high voltage power supply (Spellmann CZE2000). To produce the in-

plane aligned fibre arrays, 17 kV were applied to the needle while the grounded rotating drum collector was rotated at 100 rpm. In this technique, the nanofibres are the result of a rapid solidification of a jet of polymeric solution created by the intense electric field between the tip and the support. The nanofibres are deposited on a glass slide attached to a rotating drum collector, while the distance between anode and collector was 12 cm. Precursor solution flow rates of 0.05 mL/h, 0.40 mL/h and 0.75 mL/h were used. These rates were controlled by a dedicated syringe pump.

2.3 Material Characterization

The morphology, size and shape of *p*NA-PMMA nanofibers was verified by using a Nova Nano SEM 200 Scanning Electron Microscope (SEM) operated at an accelerating voltage of 10 kV. A homebuilt Fluorescent Lifetime Image Microscope (FLIM) [27] was used as a complementary analysis of the morphology and alignment of the *p*NA-PMMA nanofibres. FLIM analysis gives information about the individual SHG response of each nanofibre.

Crystallinity and crystallographic orientation of the *p*NA nanocrystals inside the nanostructured fibres was studied by X-ray diffraction. The X-ray diffraction pattern of nanofibres was measured from $\theta - 2\theta$ scans recorded between 10 and 60° on a Philips PW-1710 X-ray diffractometer using Cu-K α radiation of wavelength 1.5406 Å. The lattice planes parallel to the substrate surface were determined from the q vector perpendicular to the nanofibres mat plane and thus perpendicular to the longitudinal nanofibre axes.

A high-resolution Raman spectrometer, Horiba LabRAM HR Evolution confocal microscope using laser excitation at 532 nm (2.33 eV), was employed to characterize the local molecular order and orientation of the *p*NA inside of the PMMA host matrix. A 100x objective lens was used to focus the laser onto the sample. Polarized Raman spectra were obtained at room temperature between 600-1800cm $^{-1}$.

The bright field microscopic images were recorded using an Olympus BX51 microscope (10x, 50x, 100x objectives), in the transmission mode, equipped with a digital CCD camera (Retiga 4000R, QImaging) used to capture the microphotographs (exposure time: 0.1 s) of the samples under white light illumination via a DC regulated illuminator (DC-950, Fiber-Lite). The dark field images were acquired by replacing the standard microscope condenser, used for bright field transmission mode, by the CytoViVa enhanced darkfield illumination system. The hyperspectral microscope images were obtained with a digital camera (IPX-2M30, Imperx) coupled to a spectrograph (V10E 2/3", Specim, 30 μ m slit, nominal spectral range of 400–1000 nm, and nominal spectral resolution of 2.73 nm). A UV-IR Cut Filter (IF 486, Schneider

Kreuznach) with transmission range from 390 to 690 nm was placed in front of the cameras. Each pixel field-of-view on the hyperspectral images corresponds to an area of $128 \times 128 \text{ nm}^2$ on the sample plane.

2.4 Polarimetry measurements

The SHG response of the *p*NA-PMMA nanofibres was analysed by polarimetry measurements using the experimental set-up shown schematically in figure 1. The incident fundamental light at frequency ω was provided by a femtosecond Ti:Sapphire mode-locked laser (Coherent Mira 900F) pumped by a frequency doubled CW Neodymium-laser (Coherent Verdi 5W), with a 100 femtosecond (fs) temporal pulse width and a central wavelength of 800nm. The measurement consists in illuminating *p*NA-PMMA nanofibres with linearly polarized fundamental light and analysing the polarization of the transmitted SHG field. An achromatic half-wave plate ($\lambda/2$) is placed before a 50mm focal length plano-convex lens, in order to continuously vary the polarization direction of the incident light ($0 - 360^\circ$). The generated SHG field is collected by a microscope objective $100\times$ Mitutoyo Plan Infinity-Corrected Long working distance objective (NA=0.7). A set of filters was used to eliminate the transmitted fundamental beam intensity and select the SHG field with a central frequency of 2ω . A polarizer was used to analyse the polarization of the generated SHG light. The polarizer was fixed to be either parallel (q-p configuration) or perpendicular (q-s configuration) to the nanofibre's longitudinal axis. The analysed SHG field was recorded by focusing the beam onto a multimode fibre optic cable coupled to the entrance slit of a 0.3m imaging spectrograph Andor Shamrock SR-303i, with 0.20nm resolution.

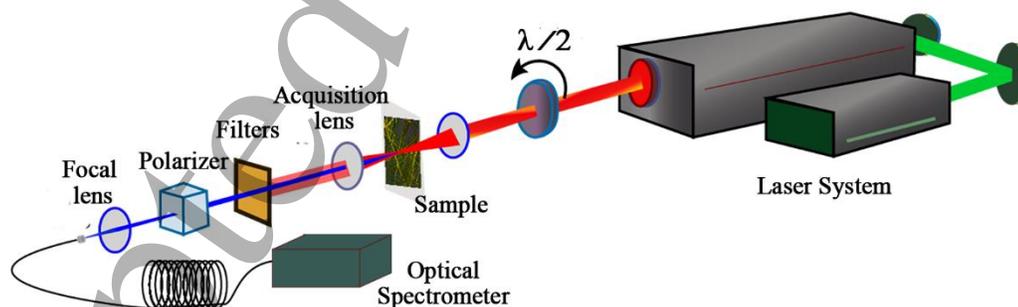


Figure 1. Polarimetry Setup. The sample is excited by a linearly polarized Ti:Sapphire laser beam. The polarization of the incident beam is continuously varied using an achromatic half-wave wave plate (WP – $\lambda/2$). After passing through a fixed polarization analyser the second harmonic signal is detected using a cooled CCD array coupled to a 0.3 m spectrograph.

3. Results and Discussion

3.1 Structural, Morphology and filled content analysis

The morphology, size and shape of the *p*NA-PMMA nanofibers were investigated using SEM as shown in figure 2 and summarized in table 1. One clearly sees that the electrospun fibres present a homogeneous cylindrical morphology with sub-micron diameters. A crucial aspect is the absence of *p*NA crystals grown outside of the PMMA fibres confirming that the SHG signal comes from *p*NA molecules located inside the polymeric matrix. No external beads are observed [28].

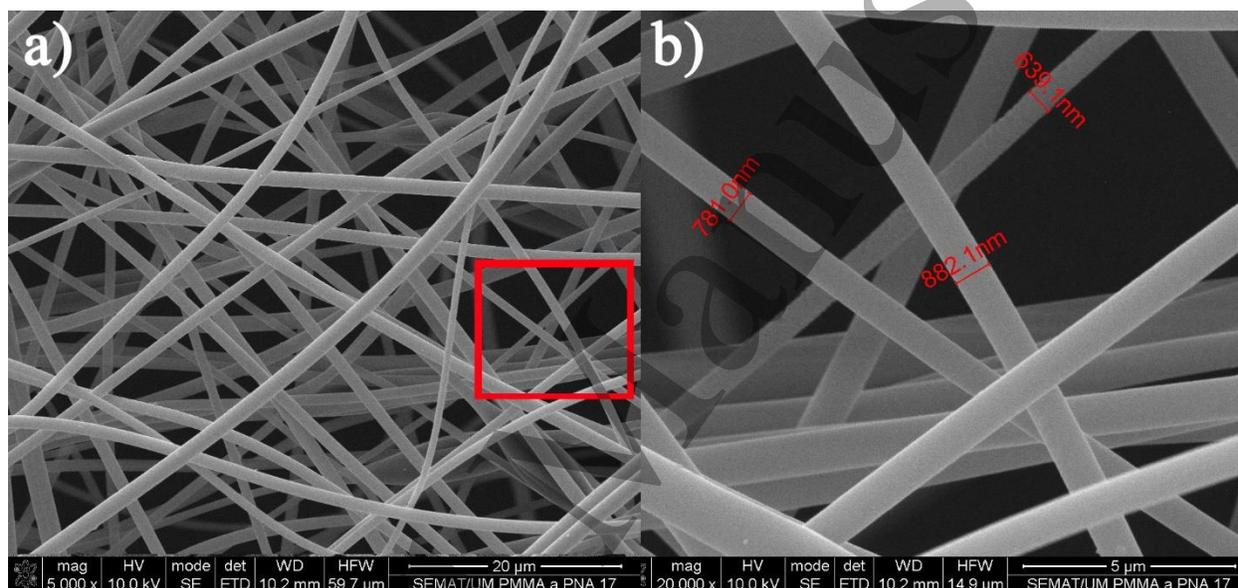


Figure 2. SEM image of *p*NA-PMMA nanofibres (a) deposited by electrospinning using a deposition rate of 0.05 mL/h. (b) Detail of the selected area in the left image with an indication of a few selected nanofibre diameters.

The high spatial and time resolution of the FLIM microscopy setup allows the measurement of the SHG signal of each individual nanofibre (figure 3). Illuminating the nanofibres by the femtosecond laser with a central wavelength of 800 nm the resulting second harmonic response (SHG) near 400 nm is collected and recorded. Each pixel of the image in figure 3 is coloured and corresponds to a single decay profile at the respective pixel's spatial coordinate. The overall temporal response is represented in the histogram. The prompt SHG emission is characteristic of a scattering process in a well-defined temporal window with a width that is close to the instrumental response time of the detection system. This eliminates the possibility of other effects, such as fluorescence induced by the absorption of two photons. FLIM microscopy demonstrates that each individual *p*NA-PMMA nanofibre presents a second order nonlinear response.

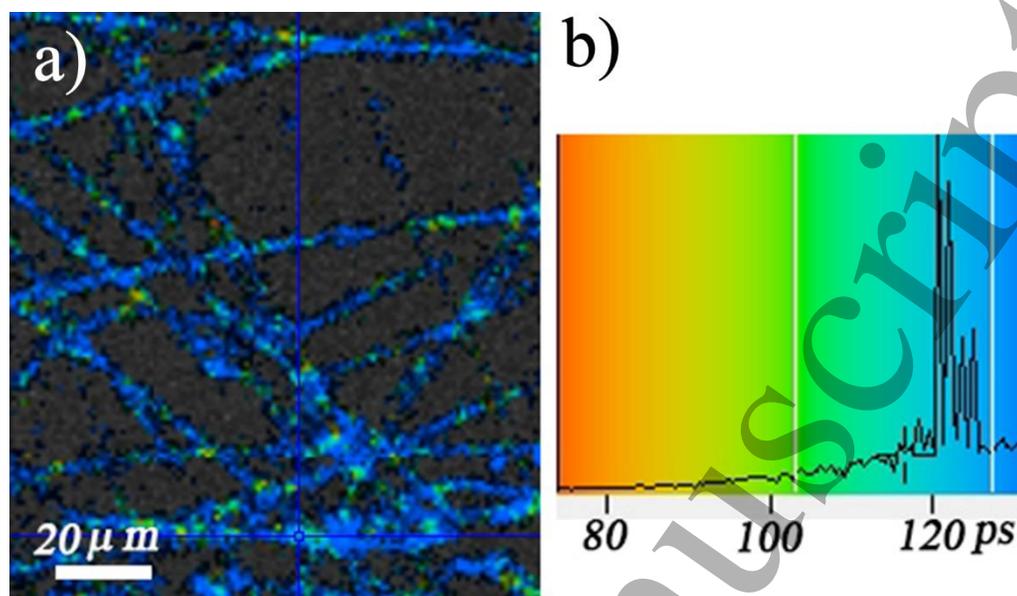


Figure 3. (a) Coloured FLIM microscope image of *p*NA-PMMA nanofibres using a deposition rate of 0.05 mL/h. The image consists of 128x128 pixels (b) Colour encoded temporal histogram of the SHG response. The sharp peaks in the histogram, with widths close to the instrument response time of the system, are a characteristic of a prompt nonlinear effect.

The mean diameter of the *p*NA-PMMA nanofibres tends to diminish as the deposition rate increases (table 1). Furthermore, the distribution of diameters broadens at higher deposition rates.

Table 1. Mean diameter of the *p*NA-PMMA nanofibres as a function of the deposition rate.

Deposition rate (mL/h)	Mean Fibre Diameter (nm)
0.05	786 ± 45
0.40	594 ± 64
0.75	578 ± 140

X-ray diffraction spectra acquired on *p*NA-PMMA fibre mats for each of the deposition rates listed in table 1 are shown in figure 4. The acquired diffraction patterns show evidence of well aligned nanocrystals dominated by a strong peak corresponding to the plane with the Miller indices (202) which is parallel to *p*NA natural cleavage plane (101) [19]. After performing a baseline subtraction, a Lorentzian was fit to the (202) peak for the different deposition rates as shown in the inset of figure 4.

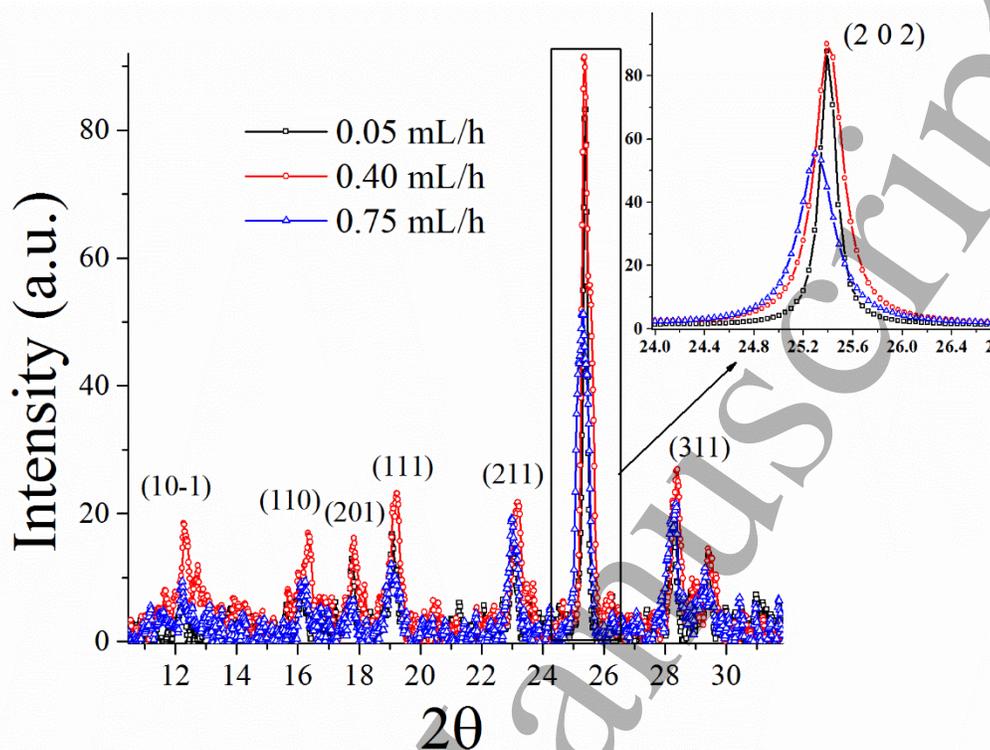


Figure 4. The X-Ray diffraction peaks of the PMMA nanofibres doped with *p*NA, deposited at different rates. (Inset) Lorentzian fit to the peak corresponding to dominant peak with the Miller indices (202).

The Lorentzian fit allows us to quantify the broadening of the (202) diffraction peak as the deposition rate is increased. This broadening could be due either to a reduction in the mean size of the *p*NA nanocrystals or an increase in strains to which they are subjected. When size effects dominate the broadening, the Scherrer equation [29] can be used to relate the width of the *hkl* diffraction peak in radians, to the mean crystal size through the equation:

$$\beta_{hkl} = \left(\frac{k\lambda}{D \cos(\theta)} \right). \quad (1)$$

Here *D* is the crystal size in the direction perpendicular to the lattice planes, λ is the wavelength of the radiation (1.5406Å), θ is the diffraction angle in radians and *k* numerical factor frequently referred to as the crystallite-shape factor. In the absence of detailed shape information, *k*=0.9 is generally accepted as a good approximation. This allows us to estimate a minimum size for the *p*NA crystals.

Table 2. The experimental values of β_{202} obtained by the Lorentzian peak fit of cleavage plane with the Miller indices (202) for the different deposition rate and the corresponding estimate for lower limit of the mean crystal size using equation 1.

Deposition rate (mL/h)	β_{202} (radians)	Minimum <i>p</i> NA crystal size (nm)
0.05	0.15	54.26
0.40	0.28	29.07
0.75	0.37	22.01

The experimental values of β_{202} and the corresponding minimum crystal sizes listed in table 2 together with the SEM results presented in table 1, suggest that nanofibres deposited at higher rates lead to thinner nanofibres containing smaller embedded *p*NA nanocrystals. Generally, thinner nanofibres are the result of a fast and efficient solvent evaporation [9] as the nanofibres solidify on their way to the grounded collection substrate. Another possible consequence of the rapid solidification can be the increase of the surface tension generating micro-strains, which in turn broadens the diffraction peaks. For a deposition rate of 0.75 mL/h the peak broadening is accompanied by a decrease of the peak area and maximum peak height. This suggests a lower degree of crystallinity accompanied by some combination of a decrease in the average size of the nanocrystals and an increase in the presence of micro-strains. The 0.05 mL/h deposition rate leads to a significantly larger lower limit on the mean crystal size. At this deposition rate the average nanofibre diameter is larger, consistent with a slower evaporation rate which should lead to more uniform crystallization within the nanofibres. The preferential orientation corresponds to the plane with Miller indices (202) which is parallel to the natural cleavage plane (101). Theoretical calculations of surface quadratic susceptibilities in *p*NA crystals [19] indicate that (101) and (202) surface terminations correspond to large quadratic susceptibilities and consequently strong surface nonlinear responses.

Hyperspectral microscopy was used to ascertain whether the concentration of *p*NA varied significantly along the fibres. Figure 5a shows an optical image of *p*NA fibres on a microscope slide. The fibre region was magnified (area marked with a rectangle) and the resulting hyperspectral image of a few *p*NA fibres under white light excitation is depicted in figure 5c. The hyperspectral curves collected at a few selected points along the fibres (within pixels of size 128x128 nm²) are spectrally uniform consistent with an overall homogenous concentration of embedded *p*NA molecules. We note that the intensity variations

present in figure 5a, primarily occur in regions of overlapping fibres which increases the amount of light scattered.

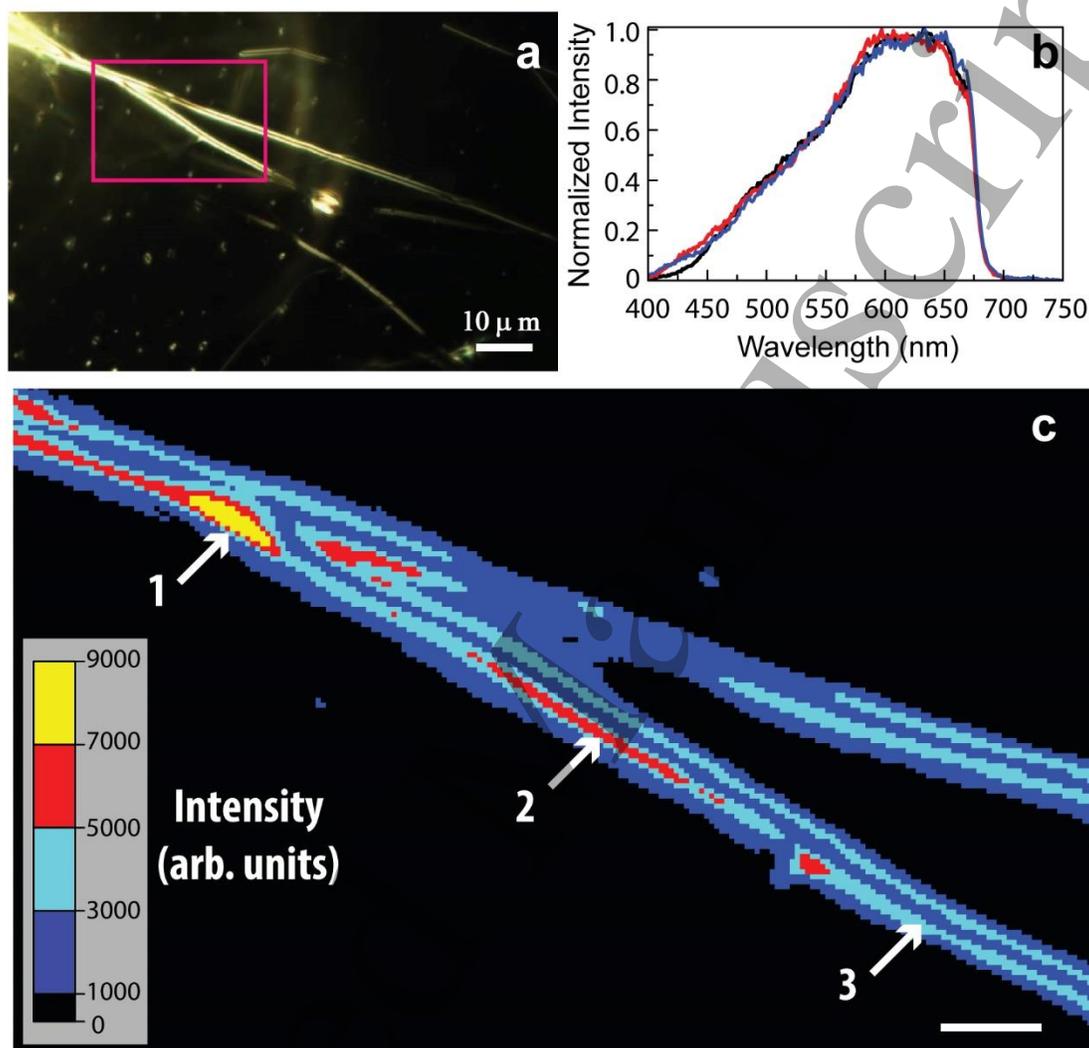


Figure 5. (a) Optical and (c) hyperspectral microscopy images of *p*NA fibres in transmission dark-field mode (b) Hyperspectral spectrum measured within the single pixel areas (1-3 in c).

Raman spectroscopy was used to better understand the nature of the *p*NA embedded within the polymeric fibres. Several peaks in the Raman spectrum have nearly the same frequencies in a thin sample of bulk crystalline *p*NA and the nanofibres, both acquired at room temperature. The most prominent of these peaks are listed in table 3.

In figure 6, the Raman bands in bulk crystalline *p*NA (solid line), and *p*NA-PMMA nanofibres (dashed line) are shown for the spectral region between 600-1700 cm^{-1} . Intense lines at 1282 cm^{-1} , 1314 cm^{-1} and

1338 cm^{-1} characterize the Raman spectrum of bulk *p*NA crystals [32,33]. The most striking feature in figure 6 is the inexistence in the *p*NA-PMMA nanofibre Raman spectra of the band centred at 1282 cm^{-1} , which is the most intense peak in bulk crystalline *p*NA. Harrand [32] has identified this peak as resulting from a superposition of the benzene ring symmetric stretching mode, ω_3 , with a combination line arising from another ring stretching mode and a torsion of the amino group, $\omega_5 + \tau_{\text{NH}_2}$. The hydrogen bonding between the nitro and amine groups of neighboring molecules induces π -electron charge concentration in the C-N bonds as well as between the bonds connecting unsubstituted carbons in the benzene ring. This produces a quinodal-like structure that leads to an enhanced Raman response. In contrast, the 1282 cm^{-1} peak is absent for *p*NA in solution where the lack of hydrogen bonding results in a more aromatic π -electron charge distribution. The absence of this prominent peak in the *p*NA-nanofibres suggests that the normal hydrogen bonds between the nitro and amine groups are somehow weakened by the electro-spinning production technique. Alternatively, it might be evidence that the majority of *p*NA molecules within the fibre are not incorporated into nanocrystals, but are rather embedded in a less organized structures. Nonetheless, the x-ray diffraction spectra does indicate that a significant fraction of *p*NA molecules do crystallize with their 202 planes oriented within the fibre mat.

Table 3. Raman frequency of the coincident peaks in bulk *p*NA and the electro-spun nanofibres together with the identification of the corresponding vibrational mode¹.

Raman Shift (cm^{-1})	Relative Intensity (Nanofibres)	Relative Intensity (Thin Crystal)	Vibrational modes ^a
860	0.34	0.24	$\delta(\text{NO}_2)$
1109	0.34	0.12	$\nu(\text{C-NO}_2)$
1137	0.12	0.08	$\phi(13)$
1178	0.12	0.07	$\phi(9a)$
1397	0.07	0.08	$\phi(19b)$
1452	0.09	0.06	$\phi(19a)$
1507	0.11	0.06	$\nu_{\text{as}}(\text{NO}_2)$
1598	0.20	0.08	$\phi(8a)$

^a ν =stretching, δ =in-plane deformation, γ =out-of-plane deformation, ϕ =vibrations of benzene; nomenclature according to G. Varsanyi [30].

The doublet 1313/1332 cm^{-1} band, that is characteristic of *p*NA in solution and assigned predominately to symmetric NO_2 stretch, is unexpectedly the strongest band in *p*NA nanofibre Raman spectra. For *p*NA in solution, the relative intensities of the doublet varies with the solvent polarity. Interestingly, the observed ratio in the *p*NA nanofibres is similar to that of *p*NA in acetonitrile (a highly polar solvent), suggesting that *p*NA molecules inside the PMMA matrix

find themselves in a polar environment. This might indicate the presence of a residual electric field from the electro-spinning process that was frozen into the nanofibres as they solidified during the flight from the needle to the collecting drum. There is also a noticeable enhancement in intensity of the of *p*NA nanofibre Raman peaks at 860 cm^{-1} which corresponds to NO_2 in plane deformation (“scissors”); at 1109 cm^{-1} corresponding to a phenyl- NO_2 stretch; and at 1598 cm^{-1} belonging to a phenyl ring stretching mode.

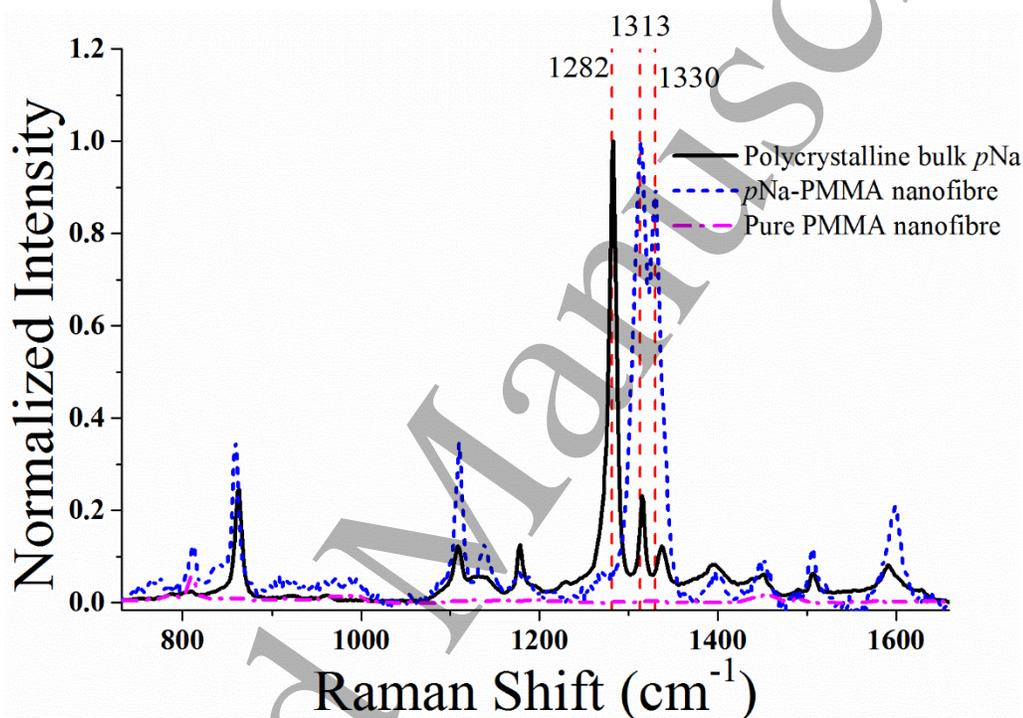


Figure 6. Raman spectra of a bulk sample of polycrystalline *p*NA, a *p*NA-PMMA nanofibre and a pure PMMA nanofibre (without *p*NA) between 600 cm^{-1} and 1700 cm^{-1} . The two spectra containing *p*NA were normalized to have the same maximum peak height, while the PMMA nanofibre spectrum is reported using the same scale as the *p*NA-PMMA nanofibre spectrum.

The Raman spectra of un-doped PMMA nanofibres is strongly polarized as shown in figure 7. The polymer chains are found to preferentially align along the nanofibre direction. The peak at 810 cm^{-1} corresponds to the stretching mode of the $\nu(\text{C-O-C})$ bond; its intensity decreases for crossed polarization, indicating an alignment of the PMMA scaffold and consequently an anisotropic environment for the embedded *p*NA molecules. This may explain why the doublet intensity ratio observed for *p*NA-PMMA nanofibres is similar to that of *p*NA in a polar solvent.

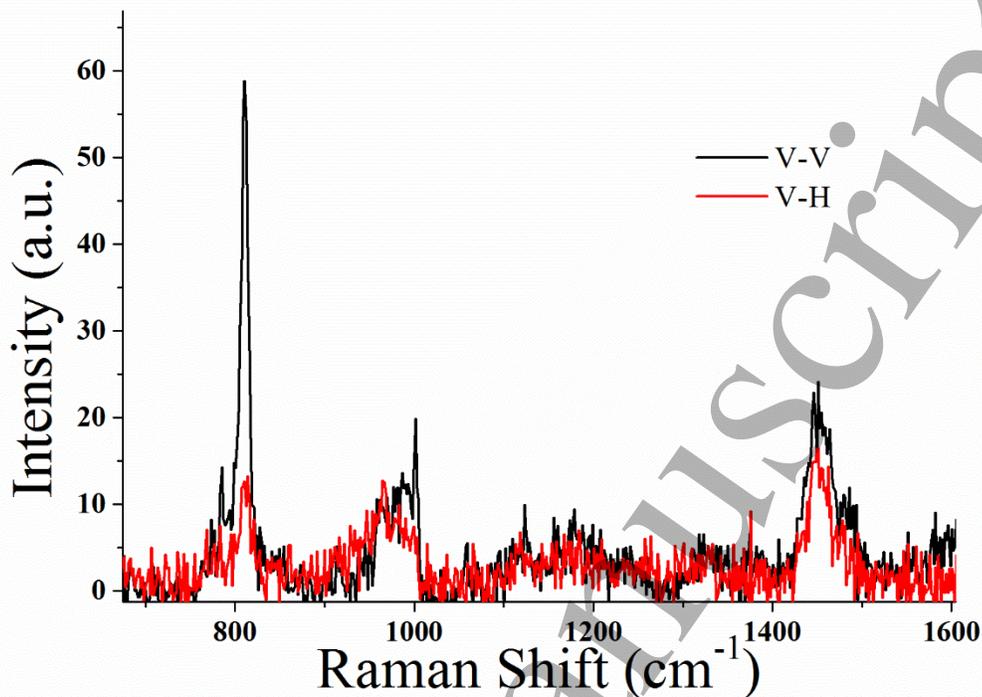


Figure 7. Raman spectra of PMMA nanofibres obtained with polarization configuration V-V and V-H.

3.2 Second harmonic polarimetry

The polarization properties of the second harmonic light was studied to assess the degree of orientational order of the *p*NA molecules embedded within the nanofibres, using the experimental set-up of figure 1. While the polarization of the incident field varies continuously, the SHG field is collected in the so-called q-p and q-s configurations, i.e. with the analyser parallel and perpendicular to the longitudinal axis of the fibre respectively. A strong signal SHG resulted when the polarization of the incident fundamental light and the analyser were both aligned parallel to the selected fibre.

The SHG response of the *p*NA-PMMA nanofibres shows a quadratic dependence as a function of the incident power (figure 8) as expected for a second order nonlinear effect. The r-squared coefficient for the quadratic fit is 0.998.

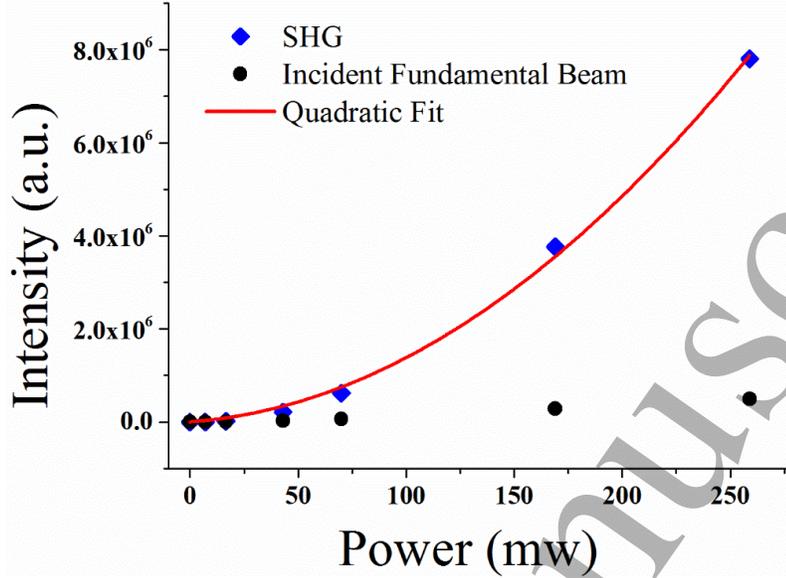


Figure 8. The SHG response of the *p*NA-PMMA nanofibres using a deposition rate of 0.05 mL/h as a function of the average incident power of the fundamental beam (repetition rate of 76 MHz).

The polarimetry curves display a polarization dependence that is characteristic of a SHG response dominated by a single diagonal second-order susceptibility tensor element, figure 9. This is in contrast to the curves obtained for *p*NA-PLLA fibres in which the q-s curves showed a lower but measurable cosine-squared like q-s response [10].

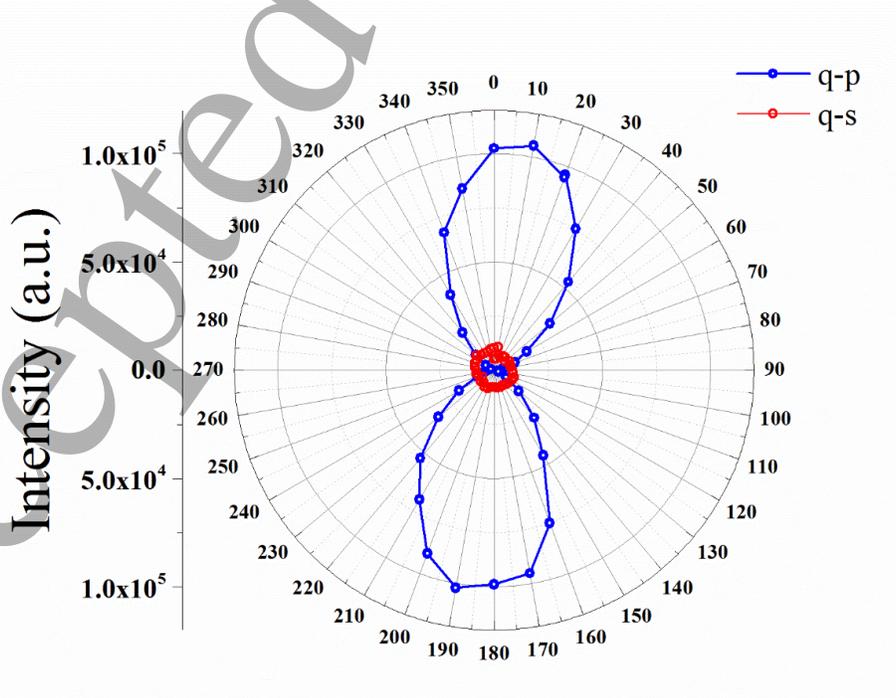


Figure 9. Polar plot of the SHG polarimetry data of *p*NA-PMMA nanofibres produced using a deposition rate of 0.05 mL/h.

When the polarization direction of the incident field is continuously varied, the intensity of the SHG field decreases and reaches a minimum at 90°. This minimum can be a function of the orientation, micro-strain and/or size homogeneity of the nanocrystals inside the nanofibres [34]. The very good alignment of the nanocrystals leads to a nearly vanishing SHG field at cross polarization. The maximum intensity of the SHG radiation for the different deposition rates represented in Figure 10 was normalized to the same value in order to explore the variation with the deposition rate of the minimum intensity (at crossed polarization) of the SHG (inset). The lowest cross-polarization intensity corresponds to the 0.05 mL/h deposition rate, consistent with the X-Ray diffraction data. This is the deposition rate that leads to a narrower diffraction peak indicating larger crystals or less micro-strain and consequently a larger degree of *p*NA crystal alignment.

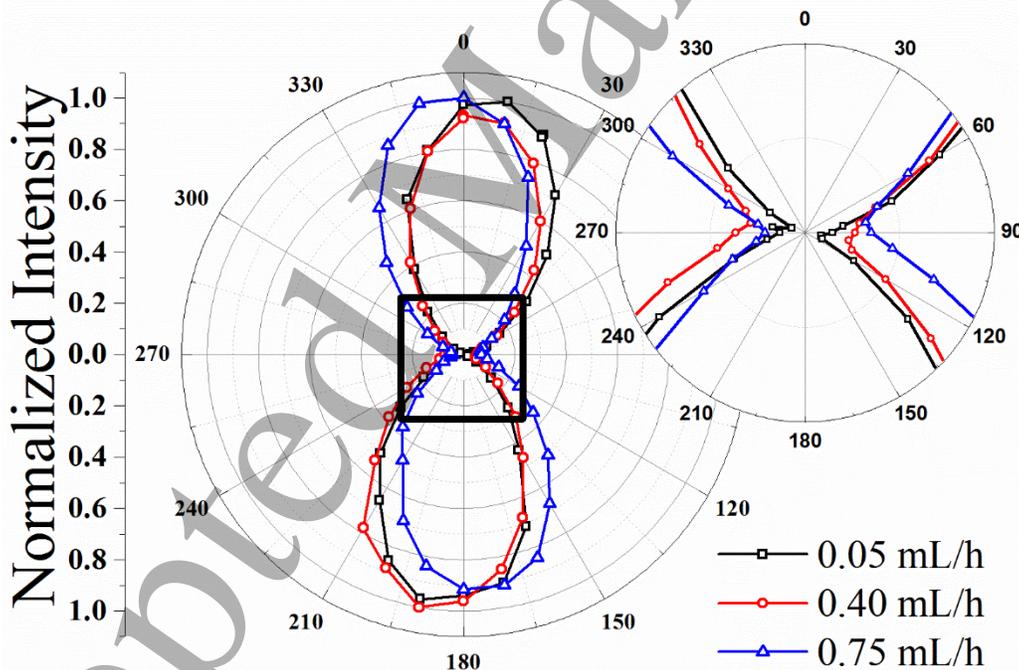


Figure 10. Polar plots of the normalized q-p SHG polarimetry data of PMMA nanofibres doped with *p*NA for the 3 different deposition rates. The close resemblance to a cosine squared polarization dependence is indicative of well-oriented SHG generators. (Inset) Variation of the minimum of SHG field near the position of crossed polarizer and analyser.

In figure 11 the second harmonic response of a *p*NA embedded PMMA nanofiber mat taken at two different times separated by interval of one year is shown. Roughly 80% of the original

response has been maintained, indicating good temporal stability for the second harmonic response of the electro-spun fibres.

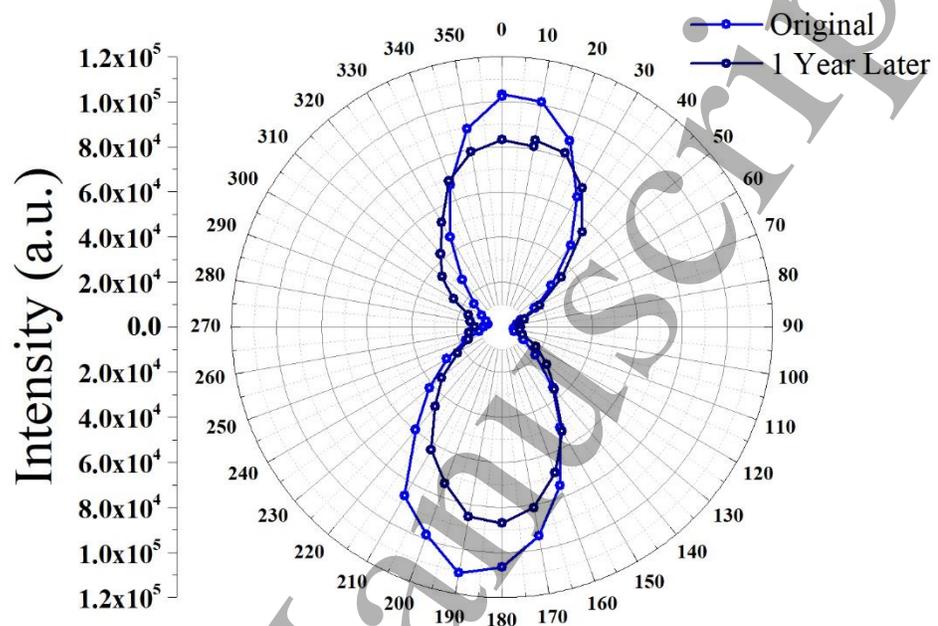


Figure 11. Polar plots of the integrated SHG polarimetry data of PMMA nanofibres doped with *p*NA produced at a deposition rate of 0.4mL/h. The two polarimetry curves were taken at an interval of one year.

3.3 Effective nonlinear optical coefficient

In this section, we explore two potential explanations for observed second harmonic signal from the *p*NA nanofibres. One possibility is that the signal is due to the surface polarization of the *p*NA nanocrystals embedded within the nanofibres. We focus our attention on the nanofibres deposited at speed of 0.05 mL/h, which have the strongest observed SHG response. The SHG signal generated by a 4 mm thick potassium dihydrogen phosphate (KDP) crystal under the same excitation and detection conditions is used to calibrate the overall detection efficiency of our system.

The incident fundamental light consisted of pulses with an energy of approximately 0.2 nJ, a FWHM duration of 100 fs and a $1/e^2$ diameter of 0.8 mm. A 50 mm focal length plano-convex lens focused the fundamental beam onto the fibre or KDP crystal. We estimate that the diameter of the focused fundamental beam is roughly 65 μm within the nanofibre or crystal with a corresponding confocal length,

1
2
3
4
5
6
7
8
9
10
11
12
13
14
15
16
17
18
19
20
21
22
23
24
25
26
27
28
29
30
31
32
33
34
35
36
37
38
39
40
41
42
43
44
45
46
47
48
49
50
51
52
53
54
55
56
57
58
59
60

$b = k_1 D^2 / 4 \approx 1.2 \text{ cm}$. Here k_1 is the magnitude of the wave vector for the fundamental beam in the medium, while D is the 1/e2 diameter at the focus. The expression for the SHG light intensity from a bulk crystal can be obtained from the solution of the nonlinear wave equation. When the confocal length of the incident fundamental beam is much greater than the crystal thickness the second harmonic power, $\mathcal{P}_{2\omega}^{KDP}$, generated by the crystal is well described by solution obtained using the plane wave approximation [35,36]:

$$\mathcal{P}_{2\omega}^{KDP} = \frac{2\omega^2}{\varepsilon_0 n_1^2 n_2 c^3} [d_{eff}]^2 \frac{L^2}{\mathcal{A}} \text{sinc}^2\left(\frac{\Delta k L}{2}\right) \mathcal{P}_\omega^2 \quad (2)$$

Here $\Delta k = 2k_1 - k_2$ is the wave vector mismatch between the fundamental beam (k_1) and the second harmonic beam (k_2) within the medium, \mathcal{A} is area of the focused beam waist and d_{eff} is the effective nonlinear coefficient. For type I phase matching in KDP at the fundamental wavelength of 800nm, the appropriate second order coefficient is $d_{eff} = \sin(\theta_{pm}) d_{36}$, where the phase matching angle is $\theta_{pm} = 44.9^\circ$ and $d_{36} = 0.39 \text{ pm} \cdot \text{V}^{-1}$ [36].

The birefringence of the KDP crystal induces a spatial walk-off between the wave-vector propagation and the direction of energy flow of the second harmonic beam that propagates as an extraordinary ray. In KDP at the phase matching orientation for 800nm incident light, this walk-off angle is 29.5 mrad [37]. Furthermore, the dispersion of KDP imposes a group velocity mismatch of approximately 80 fs/mm [37] between the fundamental and second harmonic beams leading to a further temporal walk-off effect. Wang and Weiner [38] have developed a theoretical expression for the reduction in SHG efficiency in the presence of simultaneous spatial and temporal walk-off while taking into account the diffraction effects associated with Gaussian beam propagation. Using their expression we estimate that under our conditions these combined effects will reduce the SHG efficiency by roughly a factor of 3 from that predicted by the continuous plane-wave approximation implicit in equation (2).

We couple the second harmonic light generated by the crystal or the nanofibre to the 0.3 m imaging spectrograph (Andor Shamrock SR-303i) via a multimode optical fibre bundle by employing a 100× infinity corrected long working length microscope objective from Mitutoyo (Plan Apo). This objective has a numerical aperture of 0.7 and an effective focal length of 2 mm both large enough to fully collect the second harmonic light generated by the crystal.

On other hand, following the calculations of Brevet [39] the SHG power arising from a single surface of a pNA crystal is given by,

$$\mathcal{P}_{2\omega}^{pNa} = \frac{\omega^2}{2\varepsilon_0 n_1^2 n_2 c^3} [\chi^{(2s)}]^2 \frac{1}{\mathcal{A}} \mathcal{P}_\omega^2 \quad (3)$$

Here $\chi^{(2s)}$ is the quadratic response of pNA nanocrystal's surface polarization and we have ignored small correction factors related to the Fresnel reflection and transmission coefficients. The ratio of second harmonic signals experimentally measured using our set-up for KDP and the pNA nanofibres is roughly $P_{2\omega}^{KDP} \approx 150 P_{2\omega}^{pNa}$. Comparing equations (2) and (3) and ignoring the possibly different refractive index contributions, while including the factor of 3 reduction due to the spatial and temporal walk-off in the KDP crystal, we can estimate the effective surface quadratic response under the assumption that the observed pNA nanofibre SHG signal is due to a single pNA nanocrystal via the expression,

$$\chi^{(2s)} = d_{eff}^{KDP} L \sqrt{\frac{\mathcal{P}_{2\omega}^{pNa}}{3 \mathcal{P}_{2\omega}^{KDP}}} \approx 5.2 \times 10^{-17} \text{ m}^2 / \text{V} . \quad (4)$$

Here we have taken into account that a single crystal has two surfaces that will contribute coherently to the second harmonic electric field, leading to a multiplicative factor of four in right hand side of equation (3). Malagoli and Munn [19] predict that the dominant quadratic surface susceptibility tensor element for a pNA surface oriented along the (202) direction is $\chi_{yyy}^{(2s)} = 1.93 \times 10^{-22} \text{ m}^2 / \text{V}$, more than five orders of magnitude lower than the above estimate. This would seem to rule out simple surface SHG as the sole explanation for the observed SHG signal.

A possible alternative mechanism is that the strong electric fields applied during the electrospinning process manage to align a significant number of individual pNA molecules during the ejection process and this orientation is frozen into the nanofibres during the rapid solidification that occurs as the fibres travel from the needle to the collection drum. This possibility is consistent with the Raman spectra that indicate the majority of pNA molecules find themselves effectively in a liquid state dispersed within a polar environment. In many ways this process might be viewed as an accelerated form of electric field poling. This would then allow one to make use of the high molecular polarizability of individual pNA molecules.

Assuming, for the sake of argument, that all the embedded pNA molecules were appropriately aligned by the applied electric field, the order of magnitude of the resultant second order susceptibility can be easily estimated through the relation,

$$\varepsilon_0 \chi_{pNa,eff}^{(2)}(-2\omega; \omega, \omega) = f(2\omega) f^2(\omega) N \langle \beta(-2\omega; \omega, \omega) \rangle \quad (5)$$

Here N is the number density of pNA molecules, while $\langle \beta \rangle$ is the average value of first molecular hyperpolarizability. The quantities $f(2\omega)$ and $f(\omega)$ represent the local field factors due to the presence of the surrounding PMMA matrix. Within the Lorentz approximation these factors can be estimated according to the relation

$$f(\omega) = (\epsilon_\omega + 2) / 3 \quad (6)$$

Together they result in an overall increase by a factor of 1.24 as estimated using the tabulated refractive indices of PMMA at the fundamental (1.484) and second harmonic (1.502) wavelengths. In this case the second harmonic generation mimics that of a crystalline material with the effective nonlinear coefficient d_{eff} in equation (2) replaced by $f(2\omega)f^2(\omega)N\langle\beta\rangle/\epsilon_0$. Assuming that the density of pNA molecules in the nanofibres is approximately that of pNA in the crystalline phase ($6.3 \times 10^{27} \text{ m}^{-3}$) and disregarding the possible differences in the refractive indices, we can estimate the value of the mean molecular hyperpolarizability that would be required to generate the magnitude of SHG signals we observe,

$$\langle \beta_{eff}^{pNa} \rangle = \frac{\epsilon_0}{f(2\omega)f^2(\omega)N_{pNa}} \left(\frac{L_{KDP}}{t_{pNa}} \right) \sqrt{\frac{\mathcal{P}_{2\omega}^{pNa}}{3\mathcal{P}_{2\omega}^{KDP}}} d_{eff}^{KDP}} \quad (4)$$

Where t is the thickness of a typical pNA nanofibre, roughly 800 nm and we have taken into account the reduction in SHG efficiency of the KDP crystal by a factor of 3 due to spatial and temporal beam walk-off. Substituting the appropriate values into the above equation, we find that $\langle \beta_{eff}^{pNa} \rangle \approx 7 \times 10^{-50} \text{ Cm}^3 / \text{V}^2$.

Huyskens et al. [40] have measured the average first hyperpolarizability of pNA in a variety of solvents. From their extensive list, the solvent that is chemically most similar to PMMA is methyl acetate, for which they measure a value of $9.5 \times 10^{-50} \text{ Cm}^3 / \text{V}^2$ at an incident wavelength of 1064nm when corrected for the most recent value based on second harmonic generation in quartz [41]. This agrees with the above order of magnitude estimate, suggesting that a large fraction of the second order optical response of the individual pNA molecules was translated into a macroscopic response of the doped electro-spun polymeric fibres.

The equivalent effective second order susceptibility can be estimated to be $\chi_{eff}^{(2)} \approx 65 \text{ pm} / \text{V}$, close to that estimated for pNA PLLA electrospun fibres [10] and the same magnitude as was reported by Miyazaki et al. [42] for guest-host mixtures of pNA and poly(ϵ - caprolactone). However, our electro-spinning method has the added advantage of producing fibres with a well-defined alignment and good optical quality leading to a coherent and strongly polarized response.

4. Conclusions

In this study, we have shown it is possible to obtain a strong macroscopic second harmonic response from *p*NA molecules embedded in PMMA electro-spun fibres with sub-micron diameters. Low deposition rates favour stronger SHG responses. The magnitude of the effect indicates that it is not simply a result of symmetry breaking at the *p*NA-polymer surfaces, but rather suggests that a non-centrosymmetric arrangement has been induced, possibly as the result of strong hydrogen bonds between the *p*NA molecules and the polymeric host. We speculate that the high electric fields experienced during extrusion in the electro-spinning process, coupled to the unusually large ground state *pNA* electric dipole moment of nearly 7 Debye may lead to highly aligned molecular structures that are subsequently frozen into the 1D nanostructure by the rapid solidification that occurs. This would in essence, amount to a highly accelerated version of electric field poling commonly done to align organic molecules with strong nonlinear responses in polymer substrates. However contrary to normal electric field poling, the electro-spun fibres retain their ability to generate significant second harmonic light over timescales extending to one year.

The electro-spinning technique should be extendable to other organic molecules with large dipole moments and strong second order nonlinear optical responses. Given the large number of NLO active organic compounds that crystallize in centrosymmetric structures; this technique has the potential to open up a new pathway to fabricate sub-wavelength photonic materials with large second order nonlinear optical responses.

5. Acknowledgements

Hugo Gonçalves thanks the Portuguese Foundation for Science and Technology (FCT) for the support under grant PD/BD/111873/2015. The equipment used to characterize the second harmonic response of the electro-spun fibres was acquired within the framework of the Portuguese National Program for Scientific Re-equipment, contract REEQ-25/FIS/2005 with funds from POCI 2010 (FEDER) and FCT. The Raman microscope was acquired through the project nSTeP Nanostructured Systems for Tailored Performance, NORTE-07-0124-FEDER-000039, ON.2. This work was in part developed in the scope of the projects CICECO – Aveiro Institute of Materials (UID/CTM/50011/2013), financed by national funds through the Fundação para a Ciência e a Tecnologia/Ministério da Educação e Ciência (FCT/MEC) and co-financed by FEDER under the PT2020 Partnership Agreement. The authors are grateful to A. M. P. Botas (University of Aveiro) for help in acquiring the hyperspectral microscopy data.

References

- [1] Garmire E 2013 Nonlinear optics in daily life *Opt. Express* **21** 30532–44
- [2] Rosenne S, Grinvald E, Shirman E, Neeman L, Dutta S, Bar-Elli O, Ben-Zvi R, Oksenberg E, Milko P, Kalchenko V, Weissman H, Oron D and Rybtchinski B, 2015 Self-assembled organic nanocrystals with strong nonlinear optical response *Nano Lett.* **15** 7232–37
- [3] Xu J, Semin S, Niedzialek D, Kouwer PHJ, Fron E, Coutino E, Savoini M, Li Y, Hofkens J, Uji-I H, Beljonne D, Rasing T and Rowan AE 2013 Self-assembled organic microfibers for nonlinear optics *Adv. Mater.* **25** 2084–9
- [4] Ray PC and Leszczynski 2006 J Nonlinear optical properties of highly conjugated push-pull porphyrin Aggregates: role of intermolecular interaction *Chem. Phys. Lett.* **419** 578–83
- [5] Jeon S, Malyarchuk V, Rogers JA and Wiederrecht, GP Fabricating three-dimensional nanostructures using two photon lithography in a single exposure step *Opt. Express* **14** 2300–08
- [6] Koh AL, Fernández-Domínguez AI, McComb DW, Maier SA and Yang JKW 2011 High-resolution mapping of electron-beam-excited plasmon modes in lithographically defined gold nanostructures *Nano Lett.* **11** 1323–30
- [7] Shambat G, Ellis B, Petykiewicz J, Mayer M; Majumdar A, Sarmiento T, Harris J, Haller EE, and Vuckovic J 2012 Electrically driven photonic crystal nanocavity devices *IEEE J. Sel. Top. Quantum Electron.* **18** 1700–10
- [8] Isakov DV., de Matos Gomes E, Vieira LG, Dekola T, Belsley MS and Almeida BG 2011 Oriented single-crystal-like molecular arrangement of optically nonlinear 2-methyl-4-nitroaniline in electrospun nanofibers *ACS Nano* **5** 73–8
- [9] Bhardwaj N and Kundu SC 2010 Electrospinning: a fascinating fiber fabrication technique *Biotechnol. Adv.* **28** 325–47
- [10] Isakov DV, Belsley MS, de Matos Gomes E, Gonçalves H, Schellenberg P and Almeida B G 2014 Intense optical second harmonic generation from centrosymmetric nanocrystalline para-nitroaniline *Appl. Phys. Lett.* **104** 181903
- [11] Isakov D, de Matos Gomes E, Belsley MS, Almeida B and Cerca N 2012 Strong enhancement of second harmonic generation in 2-methyl-4-nitroaniline nanofibers *Nanoscale* **4** 4978
- [12] Chronakis IS 2005 Novel nanocomposites and nanoceramics based on polymer nanofibers using electrospinning process - a review *J. Mater. Process. Technol.* **167** 283–93
- [13] Li J, Li H, Hu H, Zhao Y and Wang Q 2015 Preparation and application of polymer nano-fiber doped with nano-particles *Opt. Mater.* **40** 49–56
- [14] Ren ML, Liu W, Aspetti CO, Sun L and Agarwal R 2014 Enhanced second-harmonic Generation from metal-integrated semiconductor nanowires via highly confined whispering gallery modes *Nat. Commun.* **5** 5432
- [15] Kravets VG, Marshall OP, Nair, RR, Thackray B, Zhukov A, Leng J and Grigorenko AN 2015 Engineering optical properties of a graphene oxide metamaterial assembled in microfluidic channels *Opt. Express* **23** 1265

- [16] Verbiest T, Houbrechts S, Kauranen M, Clays K and Persoons A 1997 Second-order nonlinear optical materials: recent advances in chromophore design *J. Mater. Chem.* **7** 2175–89
- [17] Shkir M, Riscob B, Hasmuddin M, Singh P, Ganesh V, Wahab MA, Dieguez E and Bhagavannarayana G 2014 Optical spectroscopy, crystalline perfection, etching and mechanical studies on p-nitroaniline (pNA) single crystals *Opt. Mater. (Amst.)*. **36** 675–81
- [18] Kobayashi H and Kotani M 1994 Study of single crystal surface with second-harmonic generation : p -nitroaniline *Mol. Cryst. Liq. Cryst. Sci. Technol. Sect. A. Mol. Cryst. Liq. Cryst.* **252** 277–81
- [19] Malagoli M and Munn, RW 2000 Microscopic calculation of surface-induced second-harmonic generation in crystals of para-nitroaniline *J. Chem. Phys.* **112** 6757–62
- [20] Marchewka MK, Ratajczak H and Debrus S 2003 Infrared spectrum and nonlinear optical properties of p-nitroaniline-l-tartaric acid (2:1) molecular complex *J. Nonlinear Opt. Phys. Mater.* **12** 113–21
- (21) Kosenkov D and Slipchenko LV. 2011 Solvent effects on the electronic transitions of p-nitroaniline: a QM/EFP study *J. Phys. Chem. A* **115** 392–401
- [22] Kuzyk M G, Pérez-Moreno J and Shafei S 2013 Sum rules and scaling in nonlinear optics *Physics Reports*. **529** 297–398
- [23] Kato M, Kiguchi M, Sugita N and Taniguchi Y 1997 Second-order nonlinearity of mixtures Including p-nitroaniline derivatives *J. Phys. Chem. B*, **101** 8856–59
- [24] Cambré S, Campo J, Beirnaert C, Verlact C, Cool P and Wenseleers W 2015 Asymmetric dyes align inside carbon nanotubes to yield a large nonlinear optical response *Nat. Nanotechnol.* **10** 248–52
- [25] Zhang CL and Yu SH 2014 Nanoparticles meet electrospinning: recent advances and future prospects *Chem. Soc. Rev.* **43** 4423–48
- [26] Garreau A and Duvail JL 2014 Recent advances in optically active polymer-based nanowires and nanotubes *Adv. Opt. Mater.* **2** 1122–40
- [27] Bernardo C, Belsley M, de Matos Gomes E, Gonçalves H, Isakov D, Liebold F, Pereira E, Pires V, Samantilleke A, Vasilevskiy M and Schellenberg P 2014 A versatile fluorescence lifetime imaging system for scanning large areas with high time and spatial resolution *Second International Conference on Applications of Photonics and Optics, Proc. SPIE* vol 9286 (Aveiro, Portugal, 26-30 May 2014) eds. Martins Costa MFPC and Nogueira, RN (Bellingham SPIE) pp 928637–8
- [28] Piperno S, Lozzi L, Rastelli R, Passacantando M and Santucci S 2006 PMMA nanofibers production by electrospinning *Appl. Surf. Sci.* **252** 5583–6
- [29] Zak AK, Majid, WHA, Abrishami ME and Yousefi R 2011 X-Ray analysis of ZnO nanoparticles by Williamson-Hall and size-strain plot methods *Solid State Sci.* **13** 251–6
- [30] Varsányi G 1969 *Vibrational Spectra of Benzene Derivatives*, 1st edition (New York, Academic Press) 1969
- [31] Marlow F, Hill W, Caro J and Finger G 1993 Raman-study on p-nitroaniline in channels of the molecular-sieve alpo(4)-5 *J. Raman Spectrosc.* **24** 603–8

- 1
2
3 [32] Harrand M 1975 Raman study on para nitroaniline single crystal. II: internal vibrations *J. Raman Spectrosc.* **4** 53–73
4
5
6 [33] Harrand M 1979 Raman study on p-nitroaniline: molecular structure in the molten phase *J. Raman Spectrosc.* **8** 161–4
7
8
9 [34] Brasselet S, le Floc'h V, Treussart F, Roch, JF, Zyss J, Botzung-Appert E and Ibanez A 2004 In situ diagnostics of the crystalline nature of single organic nanocrystals by nonlinear microscopy. *Phys. Rev. Lett.* **92** 207401–1
10
11
12
13 [35] Boyd, RW 2003 *Nonlinear Optics* 3rd. ed. (Rochester Academic Press)
14
15 [36] Nikogosyan DN 1991 *Handbook of Nonlinear Optical Crystals*, 3rd Revised ed. (Berlin, Springer Verlag)
16
17
18 [37] SNLO nonlinear optics code available from A. V. Smith, AS-Photonics, Albuquerque, NM
19
20 [38] Wang H and Weiner AM 2003 Efficiency of short-pulse type I second harmonic generation with simultaneous spatial walk-off, temporal walk-off and pump depletion *IEEE J. Quantum Electronics* **39** 1600
21
22
23
24 [39] Brevet PF 1996 Phenomenological three-layer model for surface second-harmonic generation at the interface between two centrosymmetric media *J. Chem. Soc. Faraday Trans.* **92** 4547
25
26
27 [40] Huyskens FL, Huyskens PL and Persoons AP 1998 Solvent dependence of the first hyperpolarizability of p-nitroanilines: differences between nonspecific dipole–dipole interactions and solute–solvent H-bonds *J. Chem. Phys.* **108** 8161
28
29
30
31 [41] Reis H 2006 Problems in the comparison of theoretical and experimental hyperpolarizabilities revisited *J. Chem. Phys.* **125** 014506
32
33
34 [42] Miyazaki T, Wantabe T, and Miyata S 1988 Highly efficient second harmonic generation in p-nitroaniline / poly(lactone) systems *Japanese J. of Appl. Phys.* **27** L1724
35
36
37
38
39
40
41
42
43
44
45
46
47
48
49
50
51
52
53
54
55
56
57
58
59
60