Some Types of Carbon-based Nanomaterials as Contrast Agents for Photoacoustic Tomography
Kateryna Dubyk, Lesia Chepela, Sergei Alekseev, Andrey Kuzmich, Boris Zousman, Olga Levinson, Aleksey Rozhin, Alain Geloen, Mykola Isaiev, Vladimir Lysenko

To cite this version:

HAL Id: hal-02921216
https://hal.science/hal-02921216
Submitted on 26 Aug 2020
Some Types of Carbon-based Nanomaterials as Contrast Agents for Photoacoustic Tomography

Kateryna Dubyk1,2, Lesia Chepela1, Sergei Alekseev1,2, Andrey Kuzmich1,2, Boris Zouman3, Olga Levinson4, Aleksey Rozhin4, Alain Geloen5, Mykola Isaiev6, Vladimir Lysenko7,8

1 Taras Shevchenko National University of Kyiv, 64/13, Volodymyrska St., 01601 Kyiv, Ukraine
2 Science Park Kyiv Taras Shevchenko University, 60, Volodymyrska St., 01033 Kyiv, Ukraine
3 Ray Techniques Ltd., Hebrew University of Jerusalem, P.O.B. 39162, Israel
4 Aston Institute of Photonic Technologies, Aston University, Aston Triangle, Birmingham B4 7ET, United Kingdom
5 University of Lyon, CarMeN Laboratory, UMR INSERM 1060, INSa de Lyon, University of Lyon, 69361 Lyon, France
6 Université de Lorraine, CNRS, LEMTA (UMR 7563), Nancy, F-54000, France
7 Nanotechnology Institute of Lyon (INL), UMR CNRS 5270, University of Lyon, 69361 Lyon, France
8 Light Matter Institute, UMR-5306, Claude Bernard University of Lyon/CNRS, Université de Lyon, 69622 Villeurbanne cedex, France

(Received 29 April 2020; revised manuscript received 20 August 2020; published online 25 August 2020)

This paper is devoted to the study of various carbon-based nanomaterials as photoacoustic contrast agents. The research work was performed on agarose-based tissue phantom containing inclusions with and without carbon-based nanomaterials. The inclusion was created with the higher density compared to phantom in order to simulate a tumor. A specially designed photoacoustic probe was introduced for measuring a level of photoacoustic signal and its enhancement caused by the nanoinclusions presence. The probe consists of a buffer for time separation of the signal coming from the excitation source, piezoelectric transducer, and amplifier. A point-by-point measurement of the signal was performed to obtain a two-dimensional map from magnitude of photoacoustic signal and phase delay of the signal registration. From phase delay the 3D photoacoustic images were reconstructed by evaluation of the depth coordinate based on the tissue sound velocity. As an excitation source the light radiation from Nd:YAG laser with a 16 ns pulse duration and a 1064 nm wavelength was used. Firstly, we considered tissue phantom with a tumor covered by graphene oxide as a reference one. It has been shown that the use of graphene oxide leads to significant improvement of the image contrast. Further, the tumours labelled with nanodiamonds (NDs) and carbon fluoroxide (CFO) nanoparticles (NPs) were studied systematically. Amplitude of the photoacoustic signals registered from such tumor phantoms are one order of magnitude lower than the signal ensured by graphene oxide. All three types of the studied carbon-based nanomaterials (GO, NDs, CFO) give stable photoacoustic signal, this allows to consider them as good candidates for further in-vitro experiments in photoacoustic imaging for biological applications. The dependences of the signal level as a function of the NPs concentration were measured for types of NPs. Considering much more efficient penetration of NDs and CFO NPs inside the cells as well as their extremely low cytotoxicity, these both types of carbon nanomaterials could be used for further in-vivo experiments.

Keywords: Carbon nanomaterials, Tissue phantoms, 3D imaging, Photoacoustic tomography.

DOI: 10.21272/jnep.12(4).04033 PACS numbers: 81.05.Uw, 42.30.Wb

1. INTRODUCTION

Optical imaging is one of the main tools for visualization in medicine, which is based on interaction of electromagnetic radiation with a studied object. The advantage of optical techniques is the relatively high resolution (1-10 μm) which increases with decrease in wavelength. However, strong optical absorption of biological tissues in the range of short wavelengths is a crucial limitation. An optimal spectral range corresponds to near-infrared region, but in this region strong light scattering leads to significant reduction of optical image quality especially for the objects localized at depths larger than the scattering length (~3 cm). On the other hand, ultrasound imaging based on propagation of acoustic waves in bio-tissues can overcome this limitation. The ultrasound imaging quality is poorer and less resolved in comparison with optical one. The photoacoustic tomography aims to use the advantages of both optical and ultrasound imaging approaches [1, 2].

Photoacoustic effect is a generation of pressure perturbations in a medium as a result of interaction of the medium with modulated intensity electromagnetic radiation. The main mechanism of the photoacoustic transformation is based on the heating of the medium because of non-radiative relaxation of photoexcited excitations which is followed by the thermelastic expansion of the medium. It should be emphasized that the significant advantage of photoacoustic technique is the sensitivity only to absorbed portion of electromagnetic radiation. Let us note that the study of photoacoustic phenomena in multiphase systems is a complicated issue [3-5]. As it was mentioned above, the absorption coefficient of biological tissues for the near-infrared wavelengths and their corresponding photoacoustic responses is rather low. The use of nanoparticles (NPs) as contrast agents significantly improves photoacoustic imaging [6] and allows to perform a deeper imaging within the tissue with enhanced contrast. The good choice corresponds
to the NPs with peak absorption in the NIR range, where optical attenuation by tissues is relatively low. Additionally, nanomaterials due to their excellent biocompatibility provide a versatile platform for theranostics to perform simultaneous therapy and diagnostics [7].

Carbon nanomaterials have significant therapeutic potential because of their biocompatibility, low toxicity, easy penetration inside living cells, photoluminescence features [8]. Furthermore, most of carbon nanomaterials possess strong absorption in the infrared or near infrared regions, which is useful for photothermal therapy of cancer [9, 10]. In the biomedical field, carbon nanomaterials were highly anticipated to provide unique and new opportunities for the development of novel biosensors, nanocarriers for drug delivery, cell imaging and photo-therapy treatment of cancer [11, 12].

The study of efficiency of various NPs as contrast agents in phantom tissues is an important preliminary step for the following in-vivo diagnostics [13, 14]. Additionally, such phantoms allow testing and optimization of new methods in a controlled way, comparing performances of photoacoustic imaging, ultrasound systems as well as to assist in the development of new acoustic, ultrasound transducers, systems or diagnostic techniques. A large number of phantoms were already developed to mimic optical and acoustic properties of biological tissues [15]. In particular, agarose gel is widely used as a phantom of human tissue because its elastic characteristics are similar to those of human body [16].

Our paper is devoted to the study of different carbon-based nanomaterials as photoacoustic sensitizers. More specifically, we consider nanodiamonds (NDs) and carbon fluorooxide (CFO) NPs. Additionally, graphene oxide (GO) contrast agent was used for reference measurements. We have fabricated an agarose-based tissue phantom with inclusions of a higher density to mimic a tumor site. The inclusion was covered by nano-carbon materials before its incorporation inside the tissue phantoms. 3D photoacoustic tomography of the inclusion with and without carbon nanomaterials was performed to estimate the level of image contrast improvement.

2. MATERIALS AND METHODS

2.1 Sample Preparation

In our work, gels with 2 w/v % agarose concentrations were prepared to mimic a soft tissue [17]. Three types of agarose-based phantom samples were prepared for our measurements: (i) homogenous tissue phantom; (ii) tissue phantom with tumor; (iii) tissue phantom with tumor covered by carbon nanomaterials. Tissue phantom and tumor were fabricated from agarose gel. Density difference of the inclusions and the phantom were set by variation of concentration of agarose powder, and it was around 30%. Size of bulk inclusions 5 × 3 mm, 10 × 6 mm, 20 × 10 mm should correspond to cancer sites at different development stage of the cancer. Schematically, a series of the fabricated phantom samples is illustrated in Fig. 1.

Schemes of phantoms fabrication procedure are shown in Fig 2. The first step includes tumor fabrication (Fig 2a). For this purpose, agarose powder (A9539, Sigma Aldrich) was dissolved in water heated to 95 °C and stirred for 20 min. Concentration of agarose powder in gel was about 2 wt. %. The solution was poured into a mold to perform inclusion with a desired shape. Then, it was left for cooling and solidification for 2 h at room temperature. After that, inclusions were removed from the mold.

![Fig. 1 – Series of heterogeneous tissue phantoms](image1)

![Fig. 2 – Schemes of tissue phantom fabrication: (a) tumor fabrication; (b) covering of tumor by nanomaterials; (c) incorporation of tumor in tissue phantom](image2)
Finally, the third step was the incorporation of the tumor in a tissue phantom (Fig 2c). The inclusions were incorporated in agarose gel (concentration is equal to 1.5 wt.%) and left for cooling and solidification for 12 h at room temperature. The thickness of final tissue phantoms is about 20 mm. Photos of fabricated tissue phantom with tumor without contrast agents (a); tumor covered by NPs (b); tumor covered by CFO (c); tumor covered by GO (d) are shown in Fig. 3.

![Photos of fabricated tissue phantoms with tumor](image)

**Fig. 3** – Photos of fabricated tissue phantoms with tumor: (a) without contrast agents; (b) tumor covered by NPs; (c) tumor covered by CFO; (d) tumor covered by GO. Concentration of carbon nanomaterials for all tissue phantoms is equal to 0.1 mg/ml

3. EXPERIMENTAL DETAILS

3.1 Excitation

The sketch of the used experimental setup is shown schematically in Fig. 4. The light radiation from a Q-switched Nd:YAG laser with a 16 ns pulse duration and a 1064 nm wavelength was used as an excitation source. Pulse energy was tuned by filters to avoid any sample damage. The intensity of the laser beam was controlled with a photodiode and laser fluency was equal to 300 mJ/cm². The diameter of laser beam focusing was approximately equal to 1-2 mm.

3.2 Registration

The beam was directed toward a photoacoustic buffer shown in Fig. 4a. The photoacoustic buffer consists of a transparent buffer (30 mm thick glass piece) and a piezoelectric sensor being in rigid contact with the buffer. The photoacoustic buffer was contacted with tissue phantom through a transmission gel. The piezoelectric transducer PIC151 (Piezo Technology) with a ring shape has outer/dinner diameters and thickness equal to 10/20 mm and 1 mm, respectively, and frequency 2 MHz. Generated photoacoustic signals were amplified by Tecsonics SAP Amplifier and detected by a digital oscilloscope Tektronix TDS 2024B. The final oscillogram was averaged over 64 pulses. Typical shape of photoacoustic response from the studied phantom with tumor covered by nanomaterials is shown in Fig. 4b. To obtain 3D imaging of a sample, peak-to-peak amplitude of the photoacoustic signal (A) was measured point by point with 2D x-y stage. The scanning area was about 12 × 12 mm with a step of 250 μm. The third (z) coordinate was reconstructed by measuring the times of arrival of the ultrasound waves at the studied sample with the use of previously determined speed of sound in phantom tissue.

![Experimental setup of pulse-induced photoacoustic response](image)

**Fig. 4** – Experimental setup of pulse-induced photoacoustic response in tissue phantom (a). Typical shape of photoacoustic response from tissue phantom with tumor covered by GO (concentration is equal to 0.1 mg/ml) (b)

4. RESULTS AND DISCUSSION

Experiments were performed for all types of agarose-based phantoms shown in Fig. 1. Photoacoustic measurements were carried out for (i) homogeneous tissue phantoms without any inclusions and (ii) tissues with tumor without carbon nanomaterials (Fig. 5). All measurements have three repetitions. It is obvious that level of acoustic signal for homogeneous samples without inclusion (a) and for phantoms with inclusions of different sizes (b) is small. It is worth noting that despite the low level of the signal from considered samples, the signal registered from tissue phantom with inclusion of higher agarose density is still distinguishable bigger. These experimental facts additionally confirm the necessity of the introduction of contrast agents inside tissue phantoms to increase the amplitude of the detected photoacoustic signal.

We have measured photoacoustic signals for phantoms with the same inclusion size, but for different concentrations of GO to obtain a calibration dependence of signal amplitude on the concentration of NPs that cover the tumor. Thus, Fig. 6 compares averaged photoacoustic responses for heterogeneous tissue phantoms with various concentrations of NPs.

Calibration dependence of amplitude of photoacoustic signal on the concentration of NPs, which cover the tumor, is represented in Fig. 7. The range of concentration of GO is equal to 0.001-0.1 mg/ml. As one can see, an experimental dependence is linear. Thus, by measuring the signal level, we can evaluate the concentration of NPs which are present in the sample.

After that, photoacoustic measurements of cancer tissues with carbon nanomaterials (GO, Graphenea) were performed. The concentration of GO is equal to...
0.025 mg/ml. Fig. 8 demonstrates typical photoacoustic signal for tissue phantoms with different size of tumors covered by NPs. Amplitude of signals for all sizes of tumor inclusions has approximately the same level. It also should be noted, that since the inclusions are in a healthy phantom at different depths due to their various sizes, then the photoacoustic response should arrive with a delay. This feature is clearly observed experimentally (Fig. 8). Particularly, in the case of the smallest switch-on size, the photoacoustic response is recorded approximately in 12 µs, while the signal for the largest bulk inclusions arrives approximately in 10 µs after the laser pulse irradiation.

These features allow us to construct 3D images of heterogeneous tissue phantoms with inclusions covered with various carbon nanomaterials.

Fig. 5 – Fragment of oscillogram of photoacoustic measurements for a homogenous tissue phantom (a) and a tissue phantom with different size of tumor, but without contrast agents (b)

Fig. 6 – Typical photoacoustic signal for tissue phantoms with tumor covered by GO with different concentrations

Fig. 7 – Experimental dependence of the signal amplitude on the concentration of GO which covers the tumor. Solid line is the linear approximation of experimental data

Fig. 8 – Typical photoacoustic signal for tissue phantoms with tumor covered by GO with the same concentration

Fig. 9 – Amplitude of photoacoustic responses for agarose-based phantom tissues with tumor covered by carbon nanomaterials with different concentration. The selected area shows the concentration of nanomaterials that were selected for reconstruction of 3D images

Since, a level of acoustical signal is small for a homogenous tissue phantom and a tissue phantom with tumor and without contrast agents, 3D tomographies were reconstructed for tissue phantom with tumor covered by carbon nanomaterials.

Firstly, peak-to-peak averaged amplitude of the photoacoustic signal was measured for several concentrations of carbon NPs varying in the range 0.01 mg/ml to 1 mg/ml. Fig. 9 shows an experimental increase in amplitude with an increase in concentration of various photoacoustic contrast agents. GO leads to more important enhancement of the photoacoustic response in comparison with NDs and CFO. Therefore, tissue phantom with
tumor covered by GO is the most preferable candidate for in-vitro and ex-vivo studies. Nevertheless, degree of toxicity and typical size of the GO NPs (~ 600 nm) are not really suitable for in-vivo experiments. Fig. 10 demonstrates reconstructed 3D photoacoustic tomography of the tissue phantom with tumor covered by GO (concentration is equal to 0.05 mg/ml). As seen in Fig. 10, the experimental 3D imaging fully agrees with real photo (Fig. 3d). Particularly, GO is mainly concentrated in the center of the phantom tumor, which corresponds to the maximum photoacoustic signal amplitude. It should be mentioned that the signals detected from the different phantom tumors covered with GO with the same concentration have the same amplitude.

The images of tissue phantoms with tumor covered by NDs and CFO were also reconstructed (Fig. 11). As one can state from the figure, the signal level is one order of magnitude less compared to the case of GO. On the other hand, the experimental amplitude is quite sufficient to record a stable laser ultrasound signal with an excellent repeatability.

Since all three types of the studied nanomaterials (GO, NDs, CFO) give stable photoacoustic responses, this allows to consider them as excellent candidates for further experiments in photoacoustic imaging of biological objects. Despite the fact that the use of GO ensures the highest amplitude of the photoacoustic signal and the better visual contrast of the phantom tissue images, it is necessary to take into account various additional factors (toxicity, cell penetration, clearance rate) for future use of this kind of nanomaterials for in-vivo experiments.

10 % and 5 % of the response obtained with application of GO. Nevertheless, even such photoacoustic signals generated by CFO and NDs are quite sufficient to visualize a tumor phantom at a selected excitation wavelength. In general, all the considered carbon-based nanomaterials are excellent candidates for ex-vivo study. In our opinion, only NDs and CFO are much more suitable for further in-vivo experiments due to their efficient penetration inside the cells and low cytotoxicity.

5. CONCLUSIONS

In conclusions, 3D photoacoustic tomography can be carried out with the use of nanocarbon-based sensitizers having different structures: (i) CFO NPs with molecular nature; (ii) GO as a 2D nanomaterial and (iii) 3D NDs. GO leads to a more important photoacoustic response in comparison with NDs and CFO. In particular, for the case of NDs and CFO covering the tumor, the magnitude of the photoacoustic signals is about, respectively,
Порівняльний аналіз застосування деяких карбових матеріалів як контрастних речовин для фотоакустичної томографії

Kateryna Dubyk1,2, Lesia Chepela1, Sergei Alekseev1,2, Andrey Kuzmic1,2, Boris Zousman3, Olga Levinson4, Aleksey Rozhin4, Alain Geloen5, Mykola Isaiev6, Vladimir Lysenko7,8

1 Taras Shevchenko National University of Kyiv, 64/13, Volodymyrska St., 01601 Kyiv, Ukraine
2 Science Park Kyiv Taras Shevchenko University, 60, Volodymyrska St., 01033 Kyiv, Ukraine
3 Ray Techniques Ltd, Hebrew University of Jerusalem, P.O.B. 39162, Israel
4 Aston Institute of Photonic Technologies, Aston University, Aston Triangle, Birmingham B4 7ET, United Kingdom
5 University of Lyon, CarMeN Laboratory, UMR INSERM 1060, INSA de Lyon, University of Lyon, 69361 Lyon, France
6 Université de Lorraine, CNRS, LEMTA (UMR 7563), Nancy, F-54000, France
7 Nanotechnology Institute of Lyon (INL), UMR CNRS 5270, University of Lyon, 69621 Lyon, France
8 Light Matter Institute, UMR-5396, Claude Bernard University of Lyon, CNRS, Université de Lyon 69622 Villeurbanne cedex, France

Ця стаття присвячена вивченню різних наноматеріалів на основі вуглецю як фотоакустичних контрастних речовин. Дослідницька робота проводилася на фантомі тканини на основі агарози, що містить графен (GO) як контрастний речовину, що дозволяє вважати їх хорошими кандидатами для подальших експериментів in vivo у фотоакустичному візуалізації для біологічних застосувань. Залежності частоти сигналу від концентрації наночастинок визначають для окремих типів наночастинок. Враховуючи набагато ефективніше проникнення NDs та CFO наночастинок всередину клітин, а також їх надзвичайно низьку цитотоксичність, ці обидва типи вуглецевих наноматеріалів можуть бути використані для подальших експериментів in vivo.

Ключові слова: Вуглецеві наноматеріали, Фантоми тканини, 3D-зображення, Фотоакустична томографія.