

ARTICLE

Effect of nanomaterials on the absorbed dose during an X-ray exposure

F. Benlakhdar, A.S.A. Dib* and A.H. Belbachir

Université des Sciences et de la Technologie d'Oran Mohamed Boudiaf, USTO-MB, BP 1505 EL M'naouer, 31000 Oran, Algeria.

Received 6 April 2016 – Accepted 20 September 2016

Abstract – This work is based on a numerical study using the Monte Carlo code Geant4 of an X-ray exposure of a human head containing a tumor at the center. Our principal goal was to investigate the effect of inserting nanomaterials into the tumor on the absorbed dose. Gold is the most popular nanomaterial used in nanomedicine. In our simulation, we also focused on the other nanomaterials platinum, gadolinium and silver. Our results show that gold and platinum increase the absorbed dose in the tumor by up to 55%; the most suitable X-ray energy is between 20 keV and 140 keV. Both of these nanomaterials present the same advantages in X-ray therapy. Moreover, gadolinium increases the absorbed dose by up to 50%, while silver increases the absorbed dose by up to 20%. However, for both of these nanomaterials, the best X-ray energy to use is between 20 keV and 80 keV.

Keywords: Radiation dose / human organ / tumors / X-ray

1 Introduction

Nowadays, nanotechnology in cancer treatments has become a reality, providing new tools, from earlier diagnostics and improved imaging to better, more efficient, and more targeted therapies. One of the most promising ways to achieve this goal is the use of bionanomaterials. In recent years, the *in vitro* and *in vivo* biocompatibility of gold nanomaterials has been examined in depth (Connor *et al.*, 2005; Brun *et al.*, 2008; Boisselier and Astruc, 2009); because of its high atomic number (Z) and strong photoelectric coefficient, gold has been investigated as a radiosensitizing nanomaterial inside cancer cells to improve the dose in radiotherapy (Herold *et al.*, 2000; Cho, 2005). An ideal radiation therapy treatment delivers a sufficiently high dose to cancerous cells without adverse effects to normal tissues. In this field, many studies have suggested using Monte Carlo simulation (Casta *et al.*, 2014; Berrezoug *et al.*, 2015). These studies explain the increase in destruction of cancerous cells by the electrons emitted from nanoparticles (Butterworth *et al.*, 2008; Brun *et al.*, 2009). Theoretical studies were characterized to evaluate numerous aspects of the application of gold nanoparticles for external beam radiotherapy or brachytherapy (Bahreyni Toossi *et al.*, 2012; Delaram *et al.*, 2013). Their capacity to absorb high quantities of X-ray radiation can be used to enhance cancer radiation therapy or increase imaging contrast in diagnostic computed tomography (Sanche, 2009). The application of Monte Carlo code in radiotherapy has been validated by several researchers. Calculation of the depth dose was evaluated by Noblet *et al.* (2016), who

confirmed good agreement between simulations and measurements, with uncertainties estimated at 1%.

In a MC study, the interaction of X-rays with single gold nanoparticles (GNPs) with different diameters was considered. It was shown that larger sizes of nanoparticle offer a higher dose (Leung *et al.*, 2011). In other research, the energy deposition due to secondary electrons produced from GNPs and the microscopic dose enhancement around GNPs were quantified (Jones *et al.*, 2010). The MC results obtained by McMahan (McMahon *et al.*, 2011; Lechtman *et al.*, 2011, 2013; Garnica-Garza *et al.*, 2013) are in good agreement with experimentally observed cell killing by the combination of X-rays and GNPs. In an experimental study (Rahman *et al.*, 2009) where kilovolt X-ray radiation and megavolt electron radiation were used in the presence of GNPs in bovine aortic endothelial cells, it was found that the cell damage increases with the increase in the GNP concentration. Nanoparticles (NPs) can be delivered orally or intravenously. Because tumor cells are bigger and more numerous than normal cells, they have the capacity to consume more substances. Therefore, each tumor cell has the ability to contain more than one NP. Several studies have investigated the effects of inserting GNPs into tumor cells (Chow *et al.*, 2012; Ricketts *et al.*, 2012).

In practice, X-rays are generally used as a radiosensitizer with the use of the biomarker agent for diagnosis of tumors. In this paper, we are interested in the effect of some nanomaterials on the absorbed dose. Therefore, we focused our simulation on platinum, gadolinium, silver and gold, the most popular nanomaterials in nanomedicine research. Consequently, our study is composed of two parts; the first part is devoted to a simulation of a human head exposed to X-rays. The principal goal

* dibanis70@gmail.com

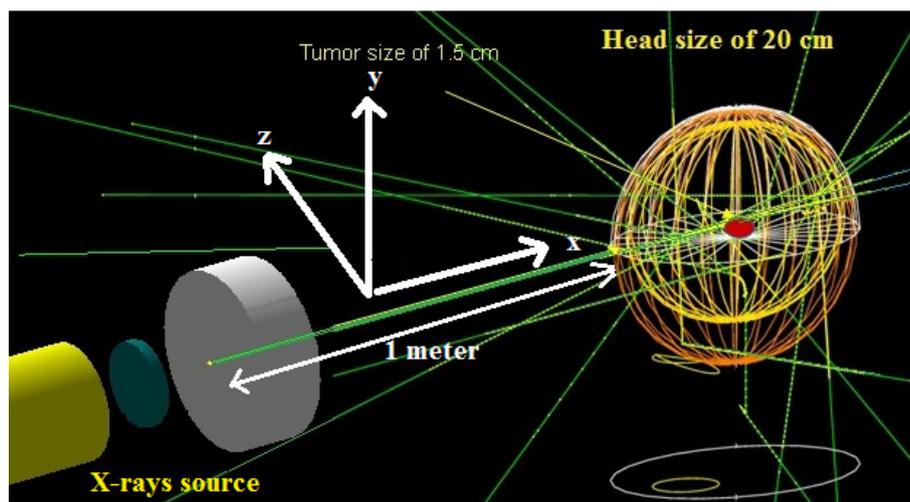


Figure 1. Monte Carlo simulation of X-ray radiation beam interaction with a human head. The radiation source is placed 1 meter from the head. The energy of the radiation beam ranged between 20 keV and 200 keV.

of this simulation is to investigate the effect of inserting nanomaterials on the absorbed dose in a tumor inside a head. The second part of our study investigates in depth the X-ray beam interaction with a nanoparticle inside a water box.

2 Methods and geometry

Our main study was to investigate the effect of bionanomaterials (bioNMs) injected into a tumor during X-ray radiation. We simulated a spherical tumor localized in the center of a human head. Then, the human head was exposed to isotropic X-ray energy ranging between 20 keV and 200 keV. Like most medical X-ray scanners used in radiation therapy, the radiation source is placed 1 meter from the patient (see Figure 1).

2.1 Monte Carlo simulations

Geant4 (Agostinelli *et al.*, 2003; Allison *et al.*, 2006) is a platform for the simulation of the passage of particles through matter using Monte Carlo methods. It is used in various application domains, including high energy physics, astrophysics and space science, and medical physics. In this work, the low electromagnetic energy package was used together with the Geant4-DNA extensions (Chauvie *et al.*, 2007; Incerti *et al.*, 2016).

2.2 Materials and geometry

2.2.1 Simulation of a tumor inside a human head

In this part of our paper, we carried out a Monte Carlo simulation of an X-ray exposure of a tumor inside a human head; we were interested in calculating the absorbed dose in the tumor with and without nanoparticles. The tumor was assumed to have a spherical shape with a diameter of 1.5 cm, and to be localized at the center of the head (see Figure 1). 1.5 cm for

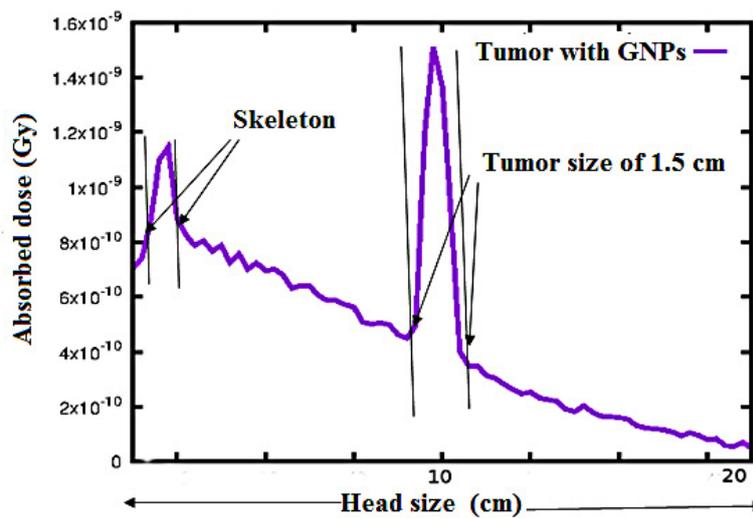
the diameter of the tumor was chosen so as to see the effect of inserting nanomaterials into a deep tumor inside a human head. The best way to inject NMs into a tumor localized in a sensitive area such as the human head is by the bloodstream. Several teams (Carmeliet and Jain, 2000; Avnesh *et al.*, 2013) studied the development of blood vessels in tumor cells; they noted that the blood vessels are more concentrated in the center of a tumor. Consequently, nanomaterials were distributed inside the tumor in a non-homogeneous way and concentrated more toward the center of the tumor. In this simulation, the concentration of nanomaterials in the tumor was assumed to range between 0 and 4%, and they are not present outside the tumor.

The geometry of the human head is composed essentially of a skeleton of 0.8 cm thickness and brain matter. Then, a soft 0.2-cm-thick tissue covers the skeleton. The tumor is placed in the center of the head. The chemical compositions and densities of the skeleton, brain, soft tissue and tumor are taken from the Geant4 database (see Table 1).

Figure 2 shows the plot of the absorbed dose in the head when exposed to X-rays of 80 keV. In this plot, the human head is exposed to an X-ray source placed 1 m from the patient. In this part of our simulation, we used the low electromagnetic energy package with cut-off energy of 250 eV and the step range of 0.01 mm. To obtain results with good accuracy, 10^8 gamma rays were emitted from the radiation source; the calculation time took more than three days on a HP Z800 Workstation. This figure shows that the absorbed dose increases considerably with the insertion of GNPs into the tumor. For more details, Figure 3 presents the calculated absorbed dose in the tumor resulting from the addition of different NMs at the same concentration and during an X-ray exposure with the energy ranging between 20 keV and 200 keV. As can be seen, in this X-ray energy range the plots of the absorbed dose in the tumor when adding gold or platinum NPs are similar and present two peaks. The first peak occurs for X-ray energy around 50 keV. At this energy value, the absorbed dose is slightly higher when adding gold NPs rather than platinum NPs. The second peak

Table 1. Chemical compositions and densities of the materials used in our geometry.

Materials and densities	Chemical compositions
Soft tissue (0.9869 g cm^{-3})	H(10.47%); C(23.02%); N(2.34%); O(63.21%); Na(0.1%); Mg(0.015%); P(0.24%); S(0.22%); Cl(0.14%); K(0.2%); Ca(9.91%); Fe(0.0063%); Zn(0.0032%); Rb(0.00057%); Sr(0.000034%); Pb(0.000016%); Zr(0.00008%)
Skeleton (1.4862 g cm^{-3})	H(7.04%); C(22.79%); N(3.87%); O(48.56%); Na(0.32%); Mg(0.11%); P(6.94%); S(0.17%); Cl(0.14%); K(0.15%); Ca(9.91%); Fe(0.008%); Zn(0.0048%); Sr(0.0032%); Pb(0.0011%)
Brain (1.040 g cm^{-3})	H(10.70%); C(14.50%); N(2.20%); O(71.20%); Na(0.20%); P(0.40%); S(0.20%); Cl(0.30%); K(0.30%)
Tumor (1.256 g cm^{-3})	H(10.70%); C(14.50%); N(2.20%); O(71.20%); Na(0.20%); P(0.40%); S(0.20%); Cl(0.30%); K(0.30%)

**Figure 2.** The absorbed dose in the head during head irradiation using an 80-keV X-ray beam.

occurs for X-ray energy of about 80 keV in the case of platinum NPs and about 85 keV in the case of gold NPs. The plots of the absorbed dose in the tumor when adding silver or gadolinium NPs present a maximum for X-ray energy around 50 keV. In Figure 3, it can be seen that both gold and platinum nanomaterials present the same advantages. Their insertion into the tumor results in an increase in the absorbed dose in the tumor of more than 55%. On the other hand, gadolinium enhances the absorbed dose in the tumor by up to 50%, 2.5 times more than silver. Comparing the effect of these nanomaterials on the absorbed dose in the tumor, gold and platinum nanomaterials increase the absorbed dose significantly, especially in the X-ray energy ranging from about 40 keV to 150 keV. However, gadolinium, gold and platinum nanomaterials are more efficient than silver in the X-ray energy ranging between 40 keV and 80 keV. To explain the results found in Figure 3, we suggest investigating the effect of nanomaterials used previously with an isotropic source X-ray on a microscopic scale.

2.2.2 Simulation of one type of nanomaterial inside a water box

In this part of our work, we studied the interaction of an X-ray beam with a water box containing one type of nanopar-

ticle. The water box size was 1 mm^3 and the nanomaterial particle diameter was taken to be 10 nm (see Figure 4). The simulation was based on the Geant4_DNA package (Chauvie *et al.*, 2007; Incerti *et al.*, 2016). The step range was taken to be 1 nanometer, and the radiation beam 10^8 gamma. In this case, the calculation time took 24 h on a HP Z800 Workstation. The interaction of X-rays with metallic nanoparticles generates several physical processes; some of these processes are photoelectric effects, ionization and electron multiscattering based on Goudsmit-Saunderson angular distribution and the Lewis spatial displacement theories (Kadri *et al.*, 2009).

The radiation beam is emitted in the x -axis, and the presence of one type of nanomaterial inside the water box generates secondary electrons and photons. As Figure 4 shows, the green lines define secondary X-rays, and the red lines define secondary electrons. The yellow dots represent the step range of secondary electrons and photons. For an external X-ray energy of 100 keV, the step range of secondary electrons is around a few micrometers outside the GNP.

In particular, for heavy atoms, the Auger effect is the most interesting physical process in this interaction. An Auger electron is ejected from the inner electron shell of the metal, which creates a vacancy. When an upper electron fills the vacancy, this gives rise to the emission of a photon, called X-ray fluorescence. The Auger electron spectrum is represented in

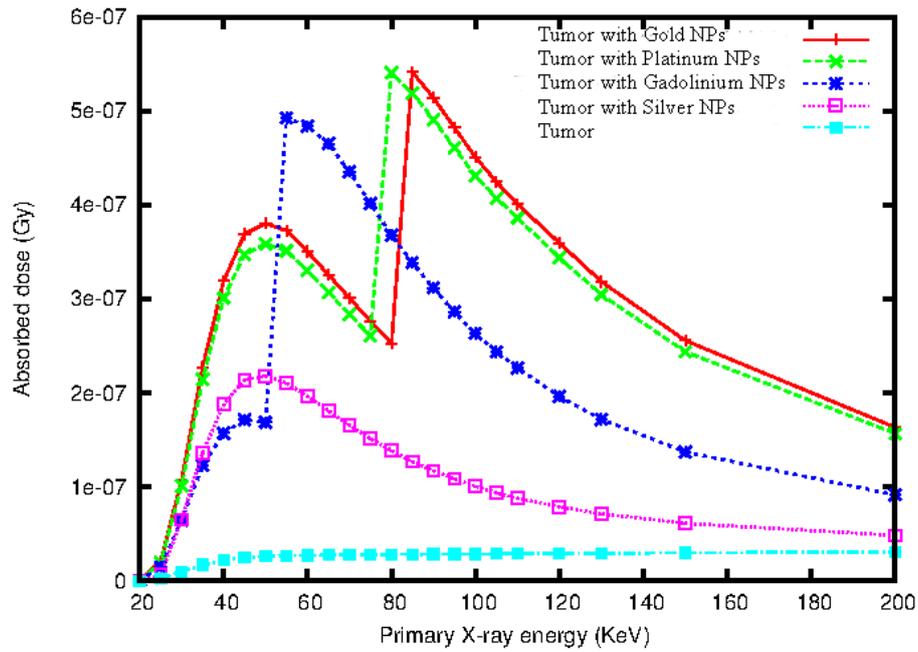


Figure 3. Plots of the absorbed dose within the tumor versus the X-ray energy (ranging from 20 keV to 200 keV) for different types of NPs with a non-homogeneous concentration ranging between 0 and 4%. The tumor is localized within the brain.

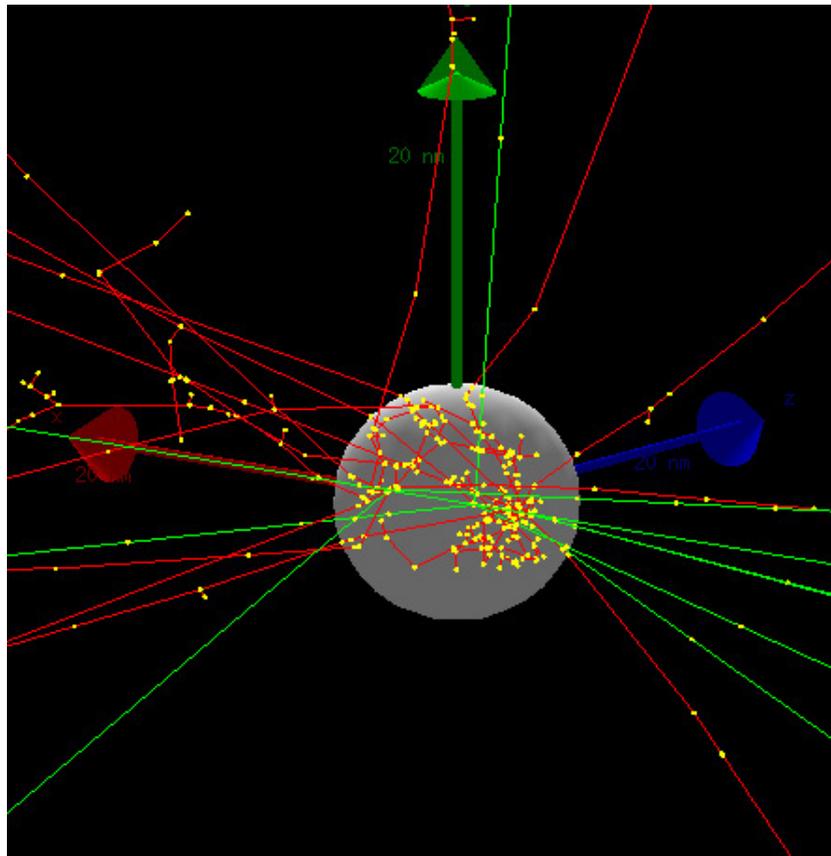


Figure 4. Simulation of X-ray beam interaction with a spherical GNP of a diameter of 10 nanometers in a water box with sides of 1 mm^3 . The green lines define secondary X-rays, and the red lines define secondary electrons. The yellow dots represent the step range of secondary electrons. In this figure the X-ray energy is 100 keV.

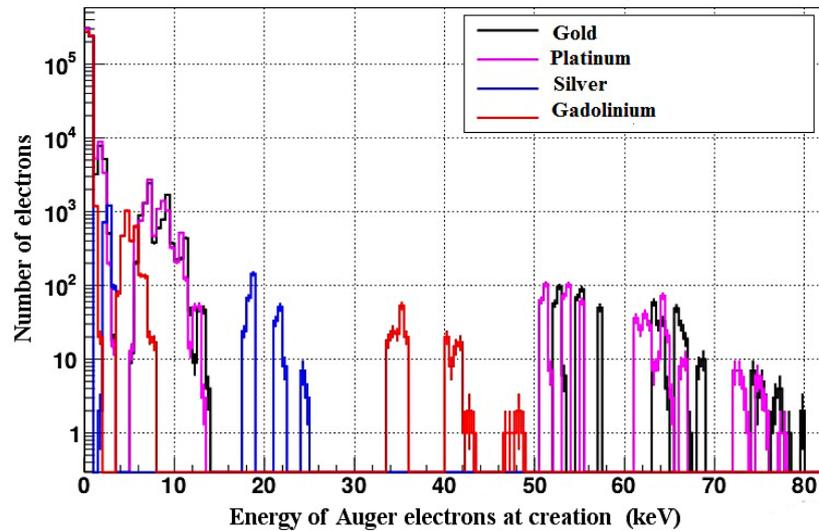


Figure 5. Spectrum of Auger electrons at an X-ray energy of 100 keV.

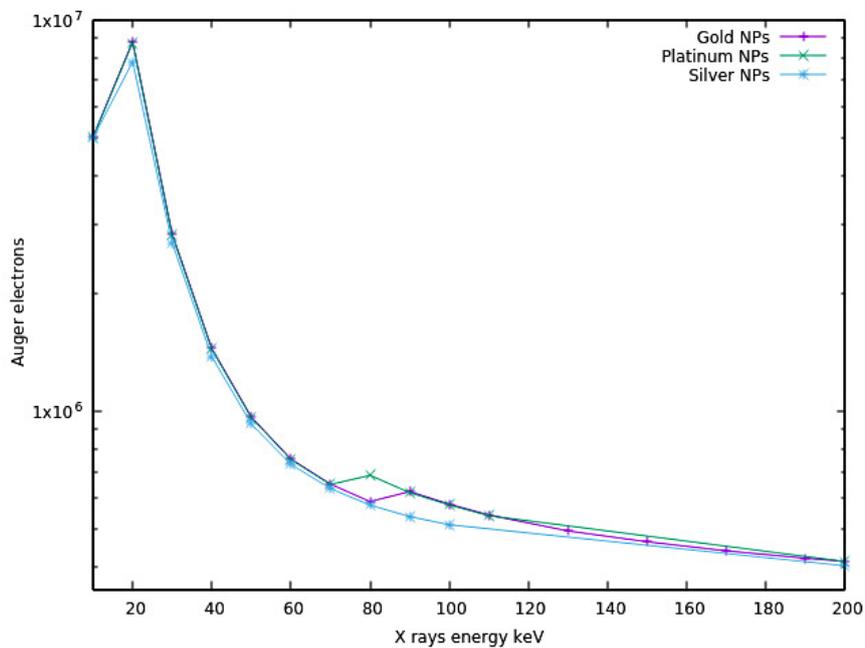


Figure 6. Number of Auger electrons produced by X-ray energy ranging from 10 keV to 200 keV.

Figure 5 in the case of different metallic nanoparticle types. In this figure, it can be seen that platinum and gold nanomaterials present the same Auger electron spectrum below 20 keV, and slightly shifted in the energy range between 50 keV and 80 keV. These spectra clearly show the existence of two energy peaks in Figure 3 in the case of gold and platinum nanomaterials. Furthermore, gadolinium presents two regions of discrete energy; the first is below 9 keV and the second is between 35 keV and 50 keV. Silver presents an energy spectrum of the electrons emitted ranging between 15 keV and 25 keV. Moreover, we plotted the number of emitted Auger electrons as a function of the X-ray energy (see Figure 6). As can be seen, this plot presents a maximum number of emitted Auger electrons at 20 keV. This result is not in contradiction with the

previous results in Figure 3; with the presence of nanomaterials, the absorbed dose increases in function of the energy of secondary electrons and photons. As Figure 5 shows, gold and platinum nanomaterials are the most favorable to enhance the dose absorption inside a tissue, with a slight difference between them.

3 Results and discussion

Over the last few decades, a large number of nanomaterial delivery systems have been developed for radiation therapy, including organic and inorganic materials. Many nanoparticles are currently in the preclinical stages of development. The use

of gold nanoparticles in radiation therapy is increasing exponentially, especially in imaging and tumor diagnosis applications, because of their biocompatibility and ability to convert energy radiation into heat (Abadeer *et al.*, 2016). In this paper, we carried out a Monte Carlo simulation based on the Geant4 code of an external X-ray beam irradiating a human head tumor, and we investigated the effect of inserting metal nanoparticles into this tumor in order to enhance this therapy.

Our results show that the presence of gold or platinum nanomaterials in the tumor considerably increases the absorbed dose in the tumor (Figures 2 and 3). Indeed, a density of 1% of these nanomaterials in the tumor is enough to enhance the absorbed dose by up to 55%. In order to increase the absorbed dose in the case of a deep tumor, as considered in our simulation, with the presence of nanoparticles in the tumor, the energy of the X-ray should be between 40 keV and 120 keV. The Auger electron spectrum (Figure 5) clearly shows that both gold and platinum nanomaterials interact in the same way during an X-ray exposure. Gadolinium and silver nanomaterials were also considered in this study. Gadolinium enhances the absorbed dose by up to 50%, while silver enhances the absorbed dose by up to 20%. As Figure 3 shows, with the use of these two nanomaterials, the absorbed dose is more enhanced in the X-ray energy range between 40 keV and 80 keV.

Our simulation results show that to increase the dose absorption in a deep tumor inside a sensitive area such as the human head, gold and platinum are the most appropriate for use during X-ray therapy. Furthermore, inserting a density of 1% of these nanomaterials inside a tumor with a size of 1.5 cm is sufficient to increase the absorbed dose by more than 55%. However, to avoid toxicity and adverse effects when using these nanometals, we propose to study in a future work the relationship between the tumor size and the percentage of nanomaterials to insert into the tumor.

References

- Abadeer N.S., Murphy C.J. (2016) Recent Progress in Cancer Thermal Therapy Using Gold Nanoparticles, *J. Phys. Chem. C* **120** (9), 4691-4716.
- Agostinelli S. *et al.* (2003) Geant4- a simulation toolkit, *Nucl. Instrum. Methods Phys. Res. A* **506**, 250-303.
- Allison J. *et al.* (2006) Geant4 developments and applications, *IEEE Transactions on Nuclear Science* **53**, 270-278.
- Avnesh S., Thakor M.D., Sanjiv S., Gambhir, M.D. (2013) Nano oncology: The future of cancer diagnosis and therapy, *CA: A Cancer Journal for Clinicians*, DOI:10.3322/caac.21199.
- Bahreyni Toossi M.T. *et al.* (2012) A Monte Carlo study on tissue dose enhancement in brachytherapy: a comparison between gadolinium and gold nanoparticles, *Australas. Phys. Eng. Sci. Med.* **35**, 177-185.
- Berrezoug A., Dib A.S.A., Belbachir A.H. (2015) Enhanced X-ray absorption by using gold nanoparticles in a biological tissue, *Radioprotection* **50** (4), 281-285.
- Boisselier E., Astruc D. (2009) Gold nanoparticles in nanomedicine: preparations, imaging, diagnostics, therapies and toxicity, *Chem. Soc. Rev.* **38**, 1759-1782.
- Brun E., Simon-Deckers A., Carriere M., Sanche L., Sicard-Roselli C. (2008) Experimental Evidence of Gold Nanoparticle Radiosensitization in vitro, *Radioprotection* **43** (5), 185.
- Brun E., Cloutier P., Sicard-Roselli C., Fromm M., Sanche L. (2009) Damage induced to dna by low-energy (0–30 eV) electrons under vacuum and atmospheric conditions, *J. Phys. Chem. B* **113** (29), 10008-10013.
- Butterworth K.T. *et al.* (2008) Gold nanoparticles: from nanomedicine to nano sensing, *Nanotechnol. Sci. Appl.* **1**, 4566.
- Carmeliet P., Jain R.K. (2000) Angiogenesis in cancer and other diseases, *Nature* **407** (6801), 249-257.
- Casta R., Champeaux J.P., Sence M., Moretto-Capelle P., Cafarelli P., Amsellem A., Sicard-Roselli C. (2014) Electronic emission of radio-sensitizing gold nanoparticles under X-ray irradiation: experiment and simulations, *J. Nanopart. Res.* **16** (4), 2348.
- Chauvie S., Francis Z., Guatelli S., Incerti S., Mascialino B., Moretto P., Nieminen P., Pia M.G. (2007) Geant4 physics processes for microdosimetry simulation: design foundation and implementation of the first set of models, *IEEE Transactions on Nuclear Science* **54** (6), 2619-2628.
- Cho S.H. (2005) Estimation of tumour dose enhancement due to gold nanoparticles during typical radiation treatments: a preliminary Monte Carlo study, *Phys. Med. Biol.* **50**, N163-N173.
- Chow J.C.L., Leung M.K.K., Jaffray D.A. (2012) Monte Carlo simulation on a gold nanoparticle irradiated by electron beams, *Phys. Med. Biol.* **57**, 3323-3331.
- Connor E.E., Mwamuka J., Gole A., Murphy C.J., Wyatt M.D. (2005) Gold nanoparticles are taken up by human cells but do not cause acute cytotoxicity, *Small* **1**, 325-327.
- Delaram P., Mahdi G., Mehdi M. (2013) Tumor dose enhancement by gold nanoparticles in a 6 MV photon beam: a Monte Carlo study on the size effect of nanoparticles, *NUKLEONIKA* **58** (2), 275-280.
- Garnica-Garza H. (2013) Microdosimetry of X-ray irradiated gold nanoparticles, *Radiat. Prot. Dosim.* **155**, 59-63.
- Herold D.M. *et al.* (2000) Gold micro-spheres: a selective technique for producing biologically effective dose enhancement, *Int. J. Radiat. Biol.* **76**, 1357-1364.
- Incerti S., Suerfu B., Xu J., Ivantchenko V., Mantero A., Brown J.M.C., Bernal M.A., Francis Z., Karamitros M., Tran H.N. (2016) Simulation of Auger electron emission from nanometer-size gold targets using the Geant4 Monte Carlo simulation toolkit, *Nucl. Instrum. Methods Phys. Res. B* **372**, 91-101.
- Jones B.L., Krishnan S., Cho S.H. (2010) Estimation of microscopic dose enhancement factor around gold nanoparticles by Monte Carlo calculations, *Med. Phys.* **37**, 3809-3816.
- Kadri O., Ivanchenko V., Gharbi F., Trabelsi A. (2009) Incorporation of the Goudsmit-Saunderson electron transport theory in the Geant4 Monte Carlo code, *Nucl. Instrum. Methods Phys. Res. B* **267** (23-24), 3624-3632.
- Lechtman E., Chattopadhyay N., Cai Z., Mashouf S., Reilly R., Pignol J.P. (2011) Implications on clinical scenario of gold nanoparticle radiosensitization in regards to photon energy, nanoparticle size, concentration and location, *Phys. Med. Biol.* **56** (15), 4631-4647.
- Lechtman E., Mashouf S., Chattopadhyay N., Keller B., Lai P., Cai Z., Reilly R., J-P P (2013) A Monte Carlo-based model of gold nanoparticle radiosensitization accounting for increased radiobiological effectiveness, *Phys. Med. Biol.* **58**, 3075-3087.

- Leung M.K. *et al.* (2011) Irradiation of gold nanoparticles by X-rays: Monte Carlo simulation of dose enhancements and the spatial properties of the secondary electrons production, *Med. Phys.* **38**, 624-631.
- McMahon S.J., Hyland W.B., Muir M.F., Coulter J.A., Jain S., Butterworth K.T., Schettino G., Dickson G.R., Hounsell A.R., OSullivan J.M., Prise K.M., Hirst D.G., Currell F.J. (2011) Biological consequences of nanoscale energy deposition near irradiated heavy atom nanoparticles, *Sci. Rep.* **1**, 18.
- Noblet C., Chiavassa S., Smekens F., Sarrut D., Passal V., Suhard J., Lisbona A., Paris F., Delpon G. (2016) Validation of fast Monte Carlo dose calculation in small animal radiotherapy with EBT3 radiochromic films, *Phys. Med. Biol.* **61**, 3521-3535.
- Rahman W.N., Bishara N., Ackerly T., Fa He C., Jackson P., Wong C. (2009) Enhancement of radiation effects by gold nanoparticles for superficial radiation therapy, *Nanomedicine: Nanotechnology, Biology, and Medicine* **5**, 136-142.
- Ricketts K., Castoldi A., Guazzoni C., Ozkan C., Christodoulou C., Gibson A.P., Royle G.J. (2012) A quantitative X-ray detection system for gold nanoparticle tumour biomarkers, *Phys. Med. Biol.* **57**, 5543-5555.
- Sanche L. (2009) Role of secondary low energy electrons in radiobiology and chemoradiation therapy of cancer, *Chem. Phys. Lett.* **474**, 16.

Cite this article as: F. Benlakhdar, A.S.A. Dib, A.H. Belbachir. Effect of nanomaterials on the absorbed dose during an X-ray exposure. *Radioprotection* 51(4), 279-285 (2016).