Image-guided treatment using an X-ray therapy unit and gold nanoparticles: Test of concept
Cindy Le Loirec, Dominique Chambellan, David Tisseur

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Gold nanoparticles (GNPs) have the potential to enhance the radiation dose locally in conjunction with kV X-rays used for radiotherapy. As for other radiotherapy modalities, the absorbed dose needs to be controlled. To do that, it is an advantage to know the distribution of GNPs. However, no effective imaging tool exists to determine the GNP distribution in vivo. Various approaches have been proposed to determine the concentration of GNPs and its distribution in a tumour and in other organs and tissues. X-ray fluorescence computed tomography (XFCT) is a promising imaging technique to do that. A new experimental device based on the XFCT technique allowing the in vivo control of GNP radiotherapy treatments is proposed. As a test of concept, experimental acquisitions and Monte Carlo simulations were performed to determine the performance that a XFCT detector has to fulfil.

INTRODUCTION

Contrast agents with high atomic number, such as gold nanoparticles (GNPs), have the potential to enhance the radiation dose locally in external radiotherapy with kV X-rays. In the kilovoltage region, gold has an attenuation coefficient two orders of magnitude greater than that of soft tissue. So, even a small amount of GNPs allows a dose enhancement inside the tumour volume by sparing surrounding normal tissue provided that the GNP concentration is lower there. The greatest ratio between the absorption coefficient of gold and that of soft tissue is found near 40–50 keV. However, the use of such energies for treatment would be complicated by the high absorption in tissue. Therefore, X-rays of higher energies, near the gold K-edge (80.7 keV), are usually used. Several developments have already been performed with monoenergetic beams around the gold K-edge, using synchrotron radiation, but that technique is not useful in a clinical context. Instead, more traditional X-ray production methods are considered with some refinements in the filtration to optimise the photon energy distribution of the beam.

One of the key tasks during preclinical studies of such a treatment method is to determine the biodistribution of GNPs injected into animals. Indeed, the Kα fluorescent X-ray energies of gold (66.99 and 68.80 keV) are both able to escape a body when produced by polychromatic X-rays. However, it has been noted that the following three technical challenges are yet to be met satisfactorily to perform in vivo quantification of GNPs:

- Goal 1: imaging of GNPs distributed within the tumour and critical organs,
- Goal 2: quantification of the amount of GNPs present,
- Goal 3: achieving system sensitivity capable of detecting GNP concentration typically found in tumours.

X-ray fluorescence computed tomography (XFCT) has shown the potential to overcome these challenges by providing GNP images as well as concentration information. XFCT imaging of GNP-loaded phantoms has been performed using K X-ray fluorescence (K-XRF) in gold. However, relatively high GNP concentrations were tested by the system. Simulation studies comparing X-ray fluorescence CT and K-edge CT show that fluorescence imaging can provide a better sensitivity for very low tracer concentration.

The development of a new experimental device based on the XFCT technique and allowing in vivo control of GNP radiotherapy treatments is now proposed. As a test of concept, in a first step the X-ray tube and the detector used at the authors’ institute for non-destructive testing were used. In parallel, Monte Carlo (MC) simulations performed with PENELOPE were used; on one hand, as an optimisation tool to enhance the experimental set-up, and on the other hand, as a planning system to estimate the GNP concentration needed to treat tumours.

MATERIALS AND METHODS

Experimental set-up

The proposed system (Figure 1) was configured according to the experimental set-up reported by MacMahon et al. and consisted of a polychromatic X-ray source (125 kVp, 300 μA), a 6 mm diameter collimator; a spectrometer system (XR 100T-CdTe, Amptek, Inc., USA), a multichannel analyser (Novelec SM-1024) and a phantom.
The XR 100T-CdTe was a thermoelectrically cooled $5 \times 5 \times 1 \text{ mm}^3$ CdTe diode X-ray detector and preamplifier. Its energy resolution was $\sim 1.5$ keV (FWHM) at 150 keV and 0.6 keV at 59.54 keV ($^{241}$Am source). It had high detection efficiency up to 100 keV, which made it ideal for XRF applications. Its calibration was performed with a metallic sheet of pure gold (99%).

The phantom was composed of two PMMA parallelepipeds of 5 cm thickness mimicking the healthy tissue and a cylindrical ‘tumour’ volume of 5 cm diameter, positioned between the two PMMA parallelepipeds and in the intersection of the incident and detected beams. The tumour volume was made of PMMA. It had a small hole into which a sheet of pure gold of 1 mm height and 25 mm diameter was inserted. The dimensions of the sheet were chosen to obtain a concentration of gold in the tumour volume equivalent to 10 mg Au g$^{-1}$ of tumour.

Several strategies were applied to effectively minimise the unwanted background in the registered pulse-height distributions: (i) two 0.6-mm-thick copper filters were placed at the exit of the collimated beam to reduce the unnecessary X-rays with energies below gold K-edge energy (80.7 keV), (ii) the spectrometer was positioned at 135° to the incident X-ray beam to minimise the impact of the tungsten peaks and of the Compton scattered X-rays in the region of the Au peaks, and (iii) a brass collimator with an opening of 6 mm diameter was placed to cover the sensitive element of the spectrometer in order to receive the directional beam while improving the signal-to-noise ratio.

A first measurement was acquired with gold and without any PMMA phantom to detect the gold peaks and to calibrate the spectrometer’s energy scale. A second measurement was performed with the PMMA inserts and the tumour volume without gold. A third measurement was performed with the PMMA inserts and the tumour cylindrical volume with gold. The three measurements lasted 10 min at 300 $\mu$A.

MC simulations

Simulations were performed with the 2006 version of PENELOPE$^{(11)}$. An RQR10 spectrum$^{(12, 13)}$ was simulated with a 0.95° aperture. The simulated geometry corresponded to the experimental set-up. A brass collimator with a 6 mm diameter opening was placed on the beam axis, and all the photons emerging it were registered to build an energy spectrum. The same brass collimator was also positioned at 45° from the beam axis, and the photons emerging it were also registered.

A second simulation was performed with the same parameters, except that the tumour cylinder was replaced by water.

RESULTS

Experimental results

Figure 2 shows the spectra obtained with and without any gold sheet in the tumour volume. Both spectra present a Compton scattering front between 62 and 66 keV. The $K_\alpha$ peaks of tungsten are visible at 67.2 and 69.1 keV for the ‘PMMA spectrum’ and the Au $K_\alpha$ peaks can be detected at 67.0 and 68.8 keV for the ‘Gold + PMMA spectrum’. The difference between both spectra is plotted in the second panel of Figure 2. Au peaks are hardly detected due to the high background level.

Simulated results

Spectra simulated with and without gold in the tumour volume are reported in Figure 3. The difference between the two simulated spectra is shown in
the second panel of Figure 3. It shows a very good discrimination of the Au $K_{\alpha 2}$ peak, due to the better energy resolution used in the MC simulation and to the absence of background induced by the detector. However, the Au $K_{\alpha 1}$ peak is hardly detectable, probably being inferior due to the MC uncertainty.

DISCUSSION
This pilot study has been performed for one experimental and one simulated configuration. The results show that gold peaks are not easily detected because their contribution is mixed with tungsten $K_{\alpha}$ peaks,
phantom Compton scattering and interactions in the detector. It has been shown\(^{10}\) that an optimisation of the beam filtration can be performed to optimise the spectral shape of the excited source and increase the sensitivity. Moreover, the detector position has also to be tested in order to find the most suitable detection set-up. Scattered X-rays can be minimised by using a backscatter detector orientation\(^{14}\). Further simulations will be performed to determine the experimental set-up that is most suitable for application in the present study. To improve the simulation, the detector model will be integrated into the simulation to analyse its impact on the Au peaks detection.

Moreover, in a first step, the authors’ laboratory detector usually used for applications in non-destructive testing was used to check the feasibility of the method. However, the energy resolution of this one is too bad (1.5 keV at 150 keV) to allow a good discrimination of the gold peaks in the Compton background. A better energy resolution is needed to allow a good discrimination of the Au peaks.

In the future, the authors plan to test the performance of a detector initially developed in their institute for Astrophysics\(^{15, 16}\). This is an imaging spectrometer (CALISTE HD), which uses a 1-mm-thick Schottky CdTe detector divided into 256 pixels disposed in a 16 × 16 matrix with a pixel pitch of 625 μm. It has been tested with a \(^{241}\)Am source, and excellent spectroscopic performance was obtained, in particular between 10 and 100 keV (0.56 and 0.67 keV FWHM at 13.9 and 59.5 keV, equivalent to 4 and 1.1 % energy resolution, respectively). A better energy resolution (~0.52 keV at 59.5 keV) can be attained for the present application after a finer calibration step and a reduction of the electronic noise.

**CONCLUSION**

Measurements performed during this study prove the feasibility to detect Au peaks in GNP radiotherapy treatments. The MC simulations show that a detector with a better energy resolution than the one used during this study is needed to get a good discrimination of the peaks. Other parameters of the detector have also to be taken into account, such as the calibration process, the scattering interactions into the elements of the detector, the pile-up effect and the dead-time to minimise the background visible in Figure 2. A more detailed MC simulation taking into account the geometry of the detector could provide useful information to characterise the ideal detector. Experimental acquisitions will thus be conducted with a better energy resolution detector developed initially for astrophysical applications. In parallel, MC simulations will be performed to optimise the experimental set-up and quantify the performance of the detector.

**REFERENCES**

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