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## Gold nanoclusters for biomedical applications: toward *in vivo* studies

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In parallel with the rapidly growing and widespread use of nanomedicine in the clinic, we are also witnessing the development of so-called theranostic agents that combine diagnostic and therapeutic properties. Among them, ultra-small gold nanoclusters (Au NCs) show promising potential due to their optical properties and activatable therapeutic activities under irradiation. Furthermore, due to their hydrodynamic diameter of smaller than 6 nm and unique biophysical properties, they also present intriguing behaviors in biological and physio-pathological environments. In this review, we aim to present the latest research studies published on such nanoparticles in animals. We also propose guidelines to identify the main physico-chemical parameters that govern the behaviour of Au NCs after administration in small animals, notably concerning their renal elimination and their ability to accumulate in tumors. Then, we present recent advances in their use as theranostic agents putting them in parallel with other contrast agents.

### 1 Introduction

In the past few years, a new class of photoluminescent ultra-small size metal nanoparticles (NPs) with a core size between 0.2 and 3 nm and usually referred to in the literature as nanoclusters (NCs)<sup>1</sup> have gained growing interest in biomedical applications. They are composed of an assembly of tens to

hundreds of metallic atoms (gold, silver, platinum, zinc, copper, *etc.*)<sup>2–6</sup> stabilized by ligands (organic thiolate molecules,<sup>7–15</sup> dendrimers,<sup>16–18</sup> DNA,<sup>19,20</sup> amino acids,<sup>21–23</sup> peptides,<sup>24–26</sup> or proteins<sup>27–29</sup>). NCs can be viewed as the missing link between metal–ligand complexes and plasmonic metal NPs.

Their optical and physicochemical properties are suitable for biomedical applications since:

- Their ultra-small size should favor renal elimination, but increase passive accumulation in the tumor micro-environment compared to small molecules.<sup>30</sup>

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• They can be detected *in vivo* by multimodal imaging techniques owing to their tunable photoluminescence (PL), from the ultra-violet (UV) to near-infrared (NIR) region<sup>17</sup> and by X-ray CT<sup>31,32</sup> or, as recently detected, by photoacoustic imaging<sup>33</sup> owing to their metallic composition.

• They can be used as radio-sensitizers<sup>34–36</sup> due to the electronic properties of metal NCs.

• They can be used as delivery systems.<sup>37–39</sup>

Gold is often preferred over other metals for biomedical applications because of its biocompatibility and inertness. Therefore, we mainly focus this review on the *in vivo* studies of gold NCs (Au NCs).

## 2 Design of Au NCs

### 2.1 Synthesis parameters

Au NCs can be synthesized in solution by the classical “bottom-up” approach using metal precursors. Depending on the nature of the ligands and the synthesis parameters such as the metal:ligand:reducing agent proportion, pH, temperature, or strength of reducing agents, a large library of Au NCs have been synthesized and used for *in vivo* studies.<sup>40</sup> They can be divided into two main categories.

The first one corresponds to Au NCs with a discrete composition at the atomic level synthesized with thiolated molecules. These clusters are represented by their formula  $M_n(L)_m$ , where  $n$  and  $m$  are respectively the number of metal atoms (M) and ligands ( $L$  = thiolate ligand  $SR^{31}$  or poly(amino-amide) PAMAM<sup>17</sup>).

Using this “bottom-up” approach, it was difficult to control the growth of single-size stable Au NCs and the final products

were often composed of mixtures of Au NCs. Indeed, in 2004, the work of Tsukuda *et al.*<sup>24,41</sup> on the characterization of gold NCs stabilized with glutathione (SG) ( $Au_nSG_m$ ), by polyacrylamide gel electrophoresis and electrospray ionization mass spectrometry, revealed the presence of nine species ( $Au_{10}SG_{10}$ ,  $Au_{15}SG_{13}$ ,  $Au_{18}SG_{14}$ ,  $Au_{22}SG_{16}$ ,  $Au_{22}SG_{17}$ ,  $Au_{25}SG_{18}$ ,  $Au_{29}SG_{20}$ ,  $Au_{33–35}SG_{22}$ , and  $Au_{38–39}SG_{22}$ ). Furthermore, it was challenging to synthesize gram-scale monodispersed Au NCs. In 2007, a so-called etching method was then proposed<sup>42</sup> based on excess ligands with increasing temperature to obtain stable and single-size Au NCs. This process was first used to produce  $Au_{25}SG_{18}$  on a large scale,<sup>43</sup> prior to adopting a size-focusing methodology with the aim of extending the obtention of large-scale monodispersed Au NCs to other sizes such as  $Au_{10}SG_{10}$ ,  $Au_{15}SG_{13}$ ,  $Au_{18}SR_{13}$  ( $SR = SG, S-c-C_6H_{11}$ ),  $Au_{38}SR_{24}$  and  $Au_{144}SR_{60}$ , ( $SR = SC_2H_4Ph, SC_{12}H_{25}$ ).<sup>1,44–46</sup> Recent reviews have summarized the advances in synthesis of Au NCs and their properties.<sup>44,47–49</sup> Size focusing still has some limitations to obtain atomically precise Au NCs with some ligands. Au NCs stabilized with polyethylene glycol (PEG),<sup>15,50</sup> zwitterions (Zw)<sup>8,33</sup> or proteins (“bovine serum albumin” BSA or transferrin)<sup>27,29</sup> as ligands exhibit some polydispersity and correspond to the second category of Au NCs: subnanometer-scaled NPs with a core size between 1 and 3 nm.<sup>51</sup>

Metal NPs tend to aggregate in solutions over time due to ionic and electrostatic interactions (van der Waals forces).<sup>52</sup> To reduce this effect, NPs can be stabilized with ligands in order to generate electrostatic or steric repulsions between the NPs. Thiolate ligands are usually used due to the strong bond between the sulfur group and the gold surface (40 kcal mol<sup>−1</sup>).<sup>53</sup> A large library of biocompatible ligands such as proteins,<sup>27</sup> DNA,<sup>19</sup> peptides,<sup>24</sup> dendrimers,<sup>17</sup> polymers<sup>16</sup> or small organic molecules following the previous criteria are suitable for stabilizing NCs.<sup>7</sup>

Lipoic acid (LA)<sup>10,11,37,54</sup> is an example of a bidentate thiol ligand with strong anchors onto the metal surface, and its carboxylic groups that are negatively charged at physiological pH are known to generate electrostatic repulsion between NCs. Following this, a large panel of zwitterions based on lipoic acid sulfobetaine (LA-sulfobetaine) for stabilizing Au NCs have been designed, which exhibited remarkable colloidal stability in various media with high antifouling effects.<sup>12,13,55</sup> Indeed, F. Aldeek *et al.*<sup>8</sup> demonstrated that Au NCs stabilized by LA-sulfobetaine with a zwitterion moiety remained stable for at least 3 months even in acidic phosphate buffer (pH 2) or in the presence of 0.25 M antioxidant biomolecules. In contrast, Au NCs stabilized only by LA aggregated with time, which confirms the enhanced colloidal stability of zwitterionic functions. Another zwitterionic ligand widely used in NC synthesis is SG.<sup>26</sup> This tripeptide is naturally present in eukaryotic cells to maintain their intracellular redox homeostasis.<sup>3</sup> SG was used to synthesize atomically precise NC  $Au_nSG_m$ , negatively charged at physiological pH, initiating the metal growth from the cysteine end of the peptide.<sup>41,42</sup> Single amino acids, such as histidine<sup>23</sup> and cysteine,<sup>21,22</sup> which are positively charged and neutral at physiological pH, respectively, have also been used to stabilize NCs. The use of longer sequences of amino acid will induce



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steric repulsion between the NCs. The first protein employed to grow NCs is the BSA.<sup>27</sup> This 66.5 kDa protein is composed of tyrosines that could reduce gold atoms under alkaline conditions (pH ~ 12),<sup>56</sup> and then be trapped by the thiol group of cysteines. Several studies reported the *in vivo* applications of AuBSA NCs.<sup>34,39,57–62</sup> Following the same process, it was possible to grow Au NCs within transferrin,<sup>63</sup> a protein that transports iron with receptors overexpressed at the surface of some cancer cells.<sup>64</sup> NCs stabilized by transferrin and conjugated with graphene oxide have already been used *in vivo*.<sup>65</sup> Finally, PEG is often used as a biocompatible capping agent to form a hydrating layer and generate steric hindrance between the NCs.<sup>8,13–15</sup>

## 2.2 Specificities of the Au NCs for *in vivo* applications

**2.2.1 Ultra-small size.** The two main elimination pathways of NPs from the body are the urinary and hepatic systems. The first one is often preferred because it is a quick filtration process from the blood through the kidneys to the bladder. More precisely, according to their different pore sizes, the NPs pass through the glomerular capillary walls into the endothelium (70–90 nm), the glomerular basement membrane (2–6 nm), and the podocyte (4–11 nm).<sup>35,45,46</sup> The combined effects of these three layers have a fixed kidney filtration threshold (KFT) of around 6–8 nm for spherical NPs (Fig. 1).<sup>66,67</sup> Spherical metal NPs with HD below 6–8 nm, around 40 kDa,<sup>68</sup> are thus expected to be eliminated by the kidneys. NPs with a larger HD are generally retained in the reticuloendothelial system (RES) through nonspecific uptake by specific macrophages of the liver (Kupffer cells) and the spleen due to the endothelial leakiness effect. Indeed, the liver is composed of non-continuous endothelial cells with vascular fenestration between 50 and 100 nm and the inter-endothelial cell slit of the spleen is between 200 and 500 nm.<sup>69</sup> There are thus typical cut-off sizes that control the elimination and nonspecific storage of the NPs in the body.

A study was conducted to follow the biodistribution of Au and Ag NPs with sizes ranging between 1.4 and 250 nm after intravenous administration in mice. It confirmed the size-dependent toxicity of NPs. If small NPs were widely spread into the organs, larger ones were mainly found in the liver and spleen.<sup>70</sup> The accumulation of metal NPs in different organs is considered a major issue due to their poor degradation that could induce acute toxicity.<sup>69,71</sup> This size effect could be extended to other inorganic nanomaterials such as quantum dots,<sup>72</sup> lanthanide NPs,<sup>73</sup> and carbon nanotubes.<sup>74</sup>

A deeper toxicology study<sup>75</sup> was carried out on Au NPs with sizes between 3 and 100 nm after their intraperitoneal injection in mice. Surprisingly, the smallest and largest NPs (3, 5, 50, and 100 nm) did not show any harmful effects in mice, while those with intermediate sizes (8, 12, 17, and 37 nm) induced severe sickness that can lead to premature death in mice. These studies revealed the potential toxicity of Au NPs larger than 6 nm and advantages of using ultra-small Au NCs.

**2.2.2 Optical properties.** Owing to their sub-nanometer diameter, Au NCs exhibit molecular-like properties such as discrete electronic states (HOMO–LUMO, highest occupied molecular orbital and lowest unoccupied molecular orbital), leading to photoluminescence (PL) properties tunable from the UV to the NIR.<sup>17</sup> The PL has been attributed to complex and multiple energy transfers taking place in the metal core *via* metal-to-metal charge transfer (MMCT), and between the metal and the ligands *via* ligand-to-metal charge transfer (LMCT) or ligand-to-metal–metal charge transfer (LMMCT). The correlation between the structure of NCs and their optical properties is still not completely understood and we invite readers to refer to fully comprehensive reviews addressing this topic.<sup>51,76</sup>

Several Au NCs exhibit PL in the NIR window between 600 and 850 nm.<sup>14,21,77</sup> Very recently, the spectral window has been extended to the shortwave infrared (SWIR) region (900–1700 nm).<sup>78</sup> Developing biocompatible optical probes in the NIR/SWIR



**Fig. 1** Layer composition of the glomerular filtration barrier. The filtration takes place through the endothelial fenestrae, across the glomerular basement membrane and the pores between podocytes. The combination of these three layers defines the size of the compounds that could cross the glomerular filtration barrier. Reproduced from ref. 66 with permission from Springer Nature publishing group.



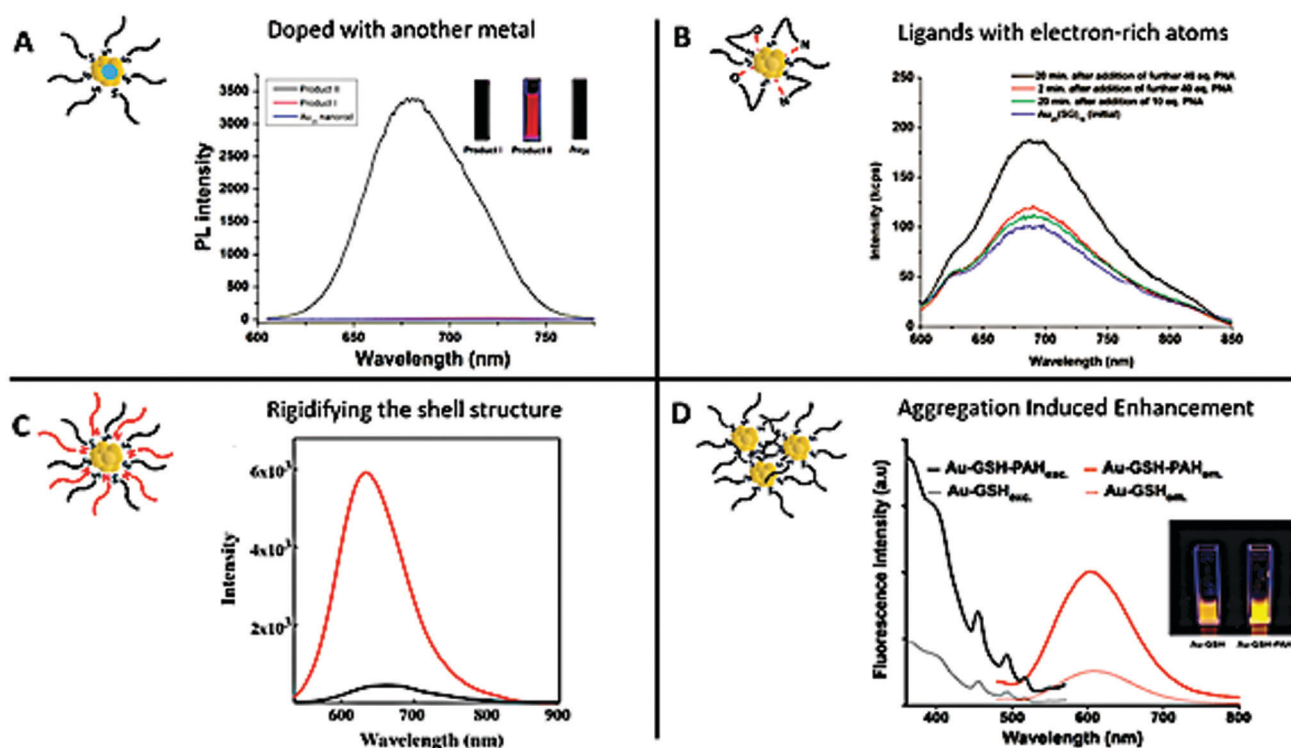


optical window presents an advantage for *in vivo* imaging in deep tissues due to reduced auto-fluorescence, exponential decrease in light scattering and improved penetration of light into the tissues, offering potentially a higher spatial resolution.<sup>79</sup> Au NCs also present large Stokes shift in comparison to most of the organic fluorophores ( $<20\text{--}30\text{ nm}^{80}$ ), which can exceed  $100\text{ nm}^{81,82}$ . This reduces the scattering and absorption of light and improves fluorescence detection.

The main limitation of atomically precise Au NCs as optical probes is related to their low quantum yield (QY) one order of magnitude lower than those of quantum dots,<sup>83</sup> lanthanide NPs,<sup>84</sup> and organic dyes<sup>85</sup> especially in the NIR/SWIR windows. Most of the Au NCs exhibit a QY  $< 1\%$  with the well-characterized  $\text{Au}_{25}\text{SG}_{18}$  exhibiting a QY  $\sim 0.3\%$ ,<sup>86</sup> which is  $10^7$  times higher than that of bulk gold.<sup>87</sup> Fortunately, a few  $\text{Au}_n\text{SG}_m$  exhibit higher QYs, such as  $\text{Au}_{18}\text{SG}_{14}$  ( $\lambda_{\text{exc.}/\text{em.}} = 590\text{ nm}/745\text{ nm}$ , QY  $\sim 5.3\%$ )<sup>88</sup> or  $\text{Au}_{22}\text{SG}_{18}$  ( $\lambda_{\text{exc.}/\text{em.}} = 520\text{ nm}/665\text{ nm}$ , QY  $\sim 8\%$ ).<sup>89</sup> Different strategies have been reported to improve the brightness of Au NCs based on (i) doping with another metal,<sup>90–92</sup> (ii) using ligands that can delocalize their electronic density to the gold core,<sup>93,94</sup> or (iii) rigidification of the ligand shell surrounding the metal core.<sup>77,95,96</sup>

Therefore, it is crucial to develop new strategies to improve their QY. A first approach to enhance the brightness involves doping Au NCs with another metal. As an example, Rongchao Jin *et al.* obtained a QY of around 40% by doping Au NCs with 13 silver atoms<sup>90</sup> (Fig. 2A). We also obtained a QY of around 15% by doping AuSG with silver.<sup>91</sup> Similar results were obtained when doping Au NCs with 2% of different metals (Ag, Cu, Pt, Zn, and Cd).<sup>68</sup> A second approach is based on the use of ligands that can delocalize their electronic density to the gold core. Replacing hexyl ( $\text{C}_6\text{H}_{13}$ ) by dodecyl ( $\text{C}_{12}\text{H}_{25}$ ) to stabilize  $\text{Au}_{25}$ , or mixing  $\text{Au}_{25}\text{SG}_{18}$  with a peptide nucleic acid (PNA) containing electron-rich atoms (*e.g.* O, N) induced a 2–3 fold enhancement of the PL<sup>93</sup> (Fig. 2B). The exchange of some ligands of  $\text{Au}_{38}(\text{SPhC}_2)_{24}$  by  $\text{HSPhNO}_2$  rather  $\text{SPhOCH}_3$  also induced an increase of the PL.<sup>94</sup>

A third strategy is based on blocking the loss of energy caused by intramolecular rotation of the ligand, by rigidifying the shell structure. Very bright fluorescent  $\text{Au}_{22}\text{SG}_{18}$  (QY  $\sim 62\%$  in toluene,  $\lambda_{\text{em.}} = 630\text{ nm}$ ) was obtained after adding tetraoctylammonium cations while the initial NCs presented a QY  $\sim 7\%$  (in water,  $\lambda_{\text{em.}} = 665\text{ nm}$ )<sup>95</sup> (Fig. 2C). A rigidification



**Fig. 2** Strategies to enhance the PL of the metal NCs. (A) Silver doping: Product I was obtained by reacting Au NPs with an  $\text{Ag}^{\text{I}}$  thiolate complex, which gave rise to  $\text{Ag}_x\text{Au}_{25-x}$  NCs with a maximum of 12 Ag atoms. Product II was obtained by reacting  $\text{Au}_{11}$  NCs with the  $\text{Ag}^{\text{I}}$  thiolate complex, which gave rise to  $\text{Ag}_{13}\text{Au}_{12}$  NCs. The PL spectra of  $\text{Au}_{25}$  NCs (bottom, blue line), Product I (middle, red line), and Product II (top, black line) show that the number of silver atoms has a strong impact on the optical properties of the NCs. The inset presents the fluorescence emission of the solution of the corresponding products under UV illumination ( $\lambda_{\text{exc.}} = 365\text{ nm}$ ). (B) LMCT: luminescence spectra of  $\text{Au}_{25}\text{SG}_{18}$  (blue) and after the addition of different equivalents of an electron-rich atom, PNA (initial concentration of  $\text{Au}_{25}\text{SG}_{18} : 1.1\text{ }\mu\text{M}$ ,  $\text{OD}_{614} \sim 0.025$ ,  $\lambda_{\text{exc.}} = 514\text{ nm}$ ). (C) Rigidification of the shell: luminescence spectra of  $\text{Au}_{22}\text{SG}_{18}$  (black) and after the addition of tetraoctylammonium cations (red) into toluene to rigidify the shell of ligands ( $\text{OD}_{514} \sim 0.025$ ,  $\lambda_{\text{exc.}} = 514\text{ nm}$ ). (D) AIE effect: excitation and emission spectra at similar Au NC concentrations of AuSG (dashed lines) and after the addition of PAH to promote electrostatic cross-linking between Au NCs (PAH-AuSG, solid lines). The inset presents the fluorescence emission of the solution of the corresponding products under UV illumination ( $\lambda_{\text{exc.}} = 366\text{ nm}$ ). Reproduced with permission from John Wiley and Sons for (A) ref. 90, and American Chemical Society for (B) ref. 93 and 94, (C) ref. 95, and (D) ref. 97.



of the shell structure of Au NCs stabilized with 6-aza-2thiothymine (ATT) was also obtained by adding L-arginine (QY  $\sim$  65%, versus QY  $\sim$  1.8% for the NCs alone;  $\lambda_{\text{em.}}$  = 530 nm).<sup>96</sup> It is also possible to rigidify the shell by creating a second layer and a PL enhancement of 300 was observed when the ratio Au:Zw passed from 1:1 to 1:40.<sup>77</sup>

Finally, the aggregation-induced enhancement effect<sup>98</sup> can also be seen as a strategy to boost Au NC PL. Indeed, aggregation will create stronger intra- and inter-complex aurophilic Au(I)  $\cdots$  Au(I) interactions that could enhance the PL signal. At the same time, the intramolecular rotation and vibration of the ligands are reduced and the Au core is better protected from solvent molecules, thereby reducing the probability of nonradiative relaxation of the excited states.<sup>47,99</sup> These two phenomena induce an enhancement of the emission of the NCs. By generating self-assembled AuSG using cationic polyelectrolytes such as poly(allylamine hydrochloride) (PAH), it was possible to control the distance between Au NCs by adjusting the pH and to increase the QY from 7 to almost 25% using the AIE effect<sup>97</sup> (Fig. 2D).

**2.2.3 Surface properties.** Surface functionalization of Au NCs has been largely investigated and an extensive library of biocompatible ligands able to generate electrostatic repulsion such as small organic molecules,<sup>7–11</sup> amino acids,<sup>21,22</sup> and peptides<sup>24–26</sup> and/or steric repulsion such as proteins,<sup>27–29</sup> DNA,<sup>19,20</sup> and dendrimers,<sup>16–18</sup> between NCs have been used.

For example, zwitterions based on lipoic acid sulfobetaine (LA-sulfobetaine), a bidentate thiol ligand with strong anchors to the metal surface, present remarkable colloidal stability in various media with high antifouling effects,<sup>8,13,100</sup> while PEGylated ligands are often used as a biocompatible capping agent to form a hydrating layer and generate steric hindrance between the NCs.<sup>8,13–15</sup>

Other small organic molecules and biomolecules are also commonly used for functionalizing the NC surface:

— fluorophores to shift the optical properties of Au NCs toward the NIR region,<sup>23,62,101</sup> drugs, photo or radiosensitizers for cancer therapy;

— targeting molecules to specifically interact with receptors overexpressed at the surface of tumor cells, such as:

- Folic acid (FA) that can recognize the folate receptors (FRs), overexpressed in many human cancer cells (ovary, breast, colon, kidneys, liver, testes, brain, lungs, and blood),<sup>54,61,101–103</sup>

- Luteinizing hormone-releasing hormone (LHRH) overexpressed in several types of cancer cells (ovary, breast, prostate, lungs, and liver),<sup>104</sup>

- Hyaluronic acid (HA) that can bind to CD44 receptors overexpressed at the surface of different cancer cells, in particular cancer stem cells,<sup>61</sup>

- AMD3100 that can interact with CXCR4, an up-regulating receptor, in particular in leukemia or breast cancer cells,<sup>105</sup>

- Cyclic RGD (cRGD), a zwitterionic compound capable of interacting with  $\alpha_v\beta_3$ ,  $\alpha_5\beta_1$ , and  $\alpha_v\beta_5$  integrins highly present in neoangiogenic endothelial cells and several solid tumors.<sup>106–108</sup>

Two main techniques can be used to functionalize the Au NCs with specific molecules. An efficient approach involves directly synthesizing Au NCs with the molecule of interest usually with a terminal thiol group that could bind to the metal surface.<sup>8,9</sup> A second approach is based on the post-functionalization of Au NCs. Click chemistry<sup>15</sup> and succinimidyl ester<sup>105,109,110</sup> reactions have been used to covalently bind molecules of interest to the ligand stabilizing the Au NCs,<sup>15,54,101,107</sup> or ligand exchange could be used if the molecule of interest has a thiol group.<sup>110</sup>

### 3. Physico-chemical parameters affecting the biodistribution of Au NCs

C. Zhou *et al.*<sup>111</sup> studied the biodistribution of AuSG with NIR emission. The highest uptake of the AuSG in the kidneys (21% of injected dose per gram of tissue, ID per g) was obtained at 5 min pi (post injection) and decreased almost by 3 fold (7.5 ID per g) after urinary excretion. These observations were consistent with their short circulation time in blood (half-life,  $t_{1/2\alpha} \sim$  5 min) and indicated that AuSG were mainly eliminated *via* the urinary system. The accumulation of AuSG in the liver and spleen was lower than that in the kidneys (respectively 4 and 2.4% ID per g) and remained constant over time. After 48 h, more than 50% of the injected AuSG were eliminated.

We obtained comparable results with AuZw, which presented a short blood circulation time ( $t_{1/2\alpha} \sim$  6 min).<sup>112</sup> Using laser-induced breakdown spectroscopy (LIBS), we detected a strong Au signal in the medulla of the kidneys 30 min pi that was almost undetectable at 1 h (Fig. 3A). In contrast, a low but

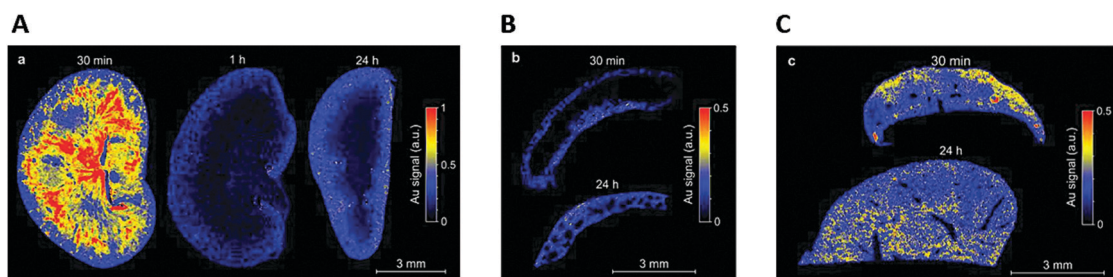
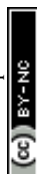


Fig. 3 After the intravenous injection of AuZwMe<sub>2</sub> (600  $\mu$ M; 200  $\mu$ L) into NMRI (Naval Medical Research Institute) nude mice, LIBS detection was carried out for: (A) kidney slices 30 min, 1 h, and 24 h pi, (B) spleen slices 30 min and 24 h, and (C) liver slices 30 min and 24 h. Reproduced from ref. 112 with permission from The Royal Society of Chemistry publishing group.



constant Au signal was observed in the liver and spleen during 24 h (Fig. 3B and C, respectively). These results were confirmed *ex vivo* by fluorescence and inductively coupled plasma mass spectrometry measurements.

These two studies demonstrated that ultra-small size Au NCs have a very rapid renal clearance, but it is important to better understand the different physicochemical parameters that may affect the kinetics and completeness of this renal elimination.

### 3.1 Length of the ligand

Due to the high surface reactivity of NCs, small changes in the ligand have huge impacts on their biodistribution. For example, the liver-to-blood and the kidney-to-blood ratios are decreased 22 and 1.9 times, respectively, when a glycine moiety is added to a cysteine ligand.<sup>21</sup>

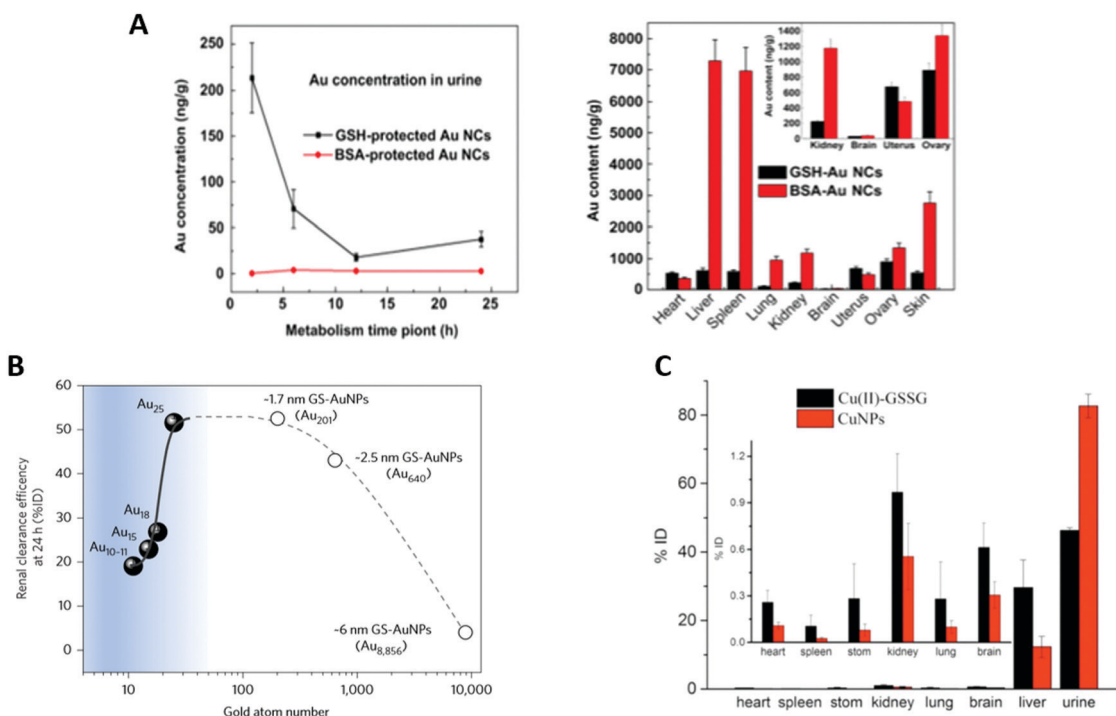
X. D. Zhang *et al.*<sup>59</sup> synthesized Au NCs with the same core size, but stabilized by different ligands such as SG or BSA. AuSG showed a very efficient renal clearance with 36% of the NCs excreted into the urine 24 h pi and only 6% were remaining in the mouse 28 days pi. In contrast, more than 95% of AuBSA were retained 28 days pi, with elevated concentrations in the liver and spleen (Fig. 4A). This difference in terms of pharmacokinetics and biodistribution was attributed to the size of the ligand that changed the HD of the Au NCs. Indeed,

the AuSG with its HD of 2 nm could be taken up by the kidneys, but not the HD of 7 nm of AuBSA.<sup>34,113</sup>

### 3.2 Size of the metal core

AuSG NPs with increasing metallic core sizes, leading to an increase in their HD (2, 6, and 13 nm) were administered intravenously<sup>114</sup> and, as could be expected, the results indicated that the larger the size, the stronger the liver uptake and the lower the kidney elimination. After one day pi, 50% of the injected dose (ID) of AuSG with an HD of 2 nm was eliminated in the urine, while only 4% or 0.5% ID for the 6 or 13 nm ones was observed. On the other hand, 3.7% ID, 27.1% ID, and 40.5% ID were respectively blocked in the liver. This tendency was also confirmed in a study that compared the clearance efficiency of 3.4 and 18.4 nm large Au NPs stabilized by SG.<sup>115</sup> However, a reverse size dependency has been observed recently when a sub-nanometer size was reached. Au<sub>25</sub>SG<sub>18</sub> were filtered faster than Au<sub>18</sub>SG<sub>14</sub> or Au<sub>10–11</sub>SG<sub>10–11</sub><sup>46</sup> (Fig. 4B). In the latter case, a very small reduction in the number of gold atoms (from 18 to 10–11), and thus a very small reduction in their HD dramatically reduced their renal elimination because of an augmented binding of the clusters on the glomerular glycocalyx.

Accordingly, biodegradable NCs that dissociate into small fragments should also be cleared from the body. CuSG are 2 nm large NCs (core diameter and 2.7 nm HD) that gradually



**Fig. 4** (A) Renal elimination and biodistribution in mice intravenously injected ( $7550 \mu\text{g kg}^{-1}$ ,  $151 \mu\text{g mL}^{-1}$ ) with AuSG and AuBSA NCs: renal elimination 24 h pi (top left) and biodistribution in the main organs 28 days pi (top right). AuBSA NCs have 10 times higher distribution in the liver and spleen than AuSG NCs. (B) Effect of the sub-nanometer size of the NCs on their renal elimination. Renal elimination efficiencies of different sizes of Au NCs, intravenously injected ( $\sim 100 \mu\text{M}$ ,  $100 \mu\text{L}$ ) in BALB/c mice, versus the number of gold atoms at 24 h pi. The renal elimination increased with the number of gold atoms up to 25 (Au<sub>25</sub>SG<sub>18</sub>) and reached a plateau followed by a decrease for larger sizes. (C) Biodistribution in the main organs 24 h after intravenous injections of Cu(II)-GSSG complexes ( $300 \mu\text{L}$ ,  $0.33 \text{ mg mL}^{-1}$ ) or luminescent CuSG NCs ( $300 \mu\text{L}$ ,  $0.83 \text{ mg mL}^{-1}$ ) in BALB/c mice ( $N = 6$ ). Reproduced with permission from Elsevier for (A) ref. 59, Springer Nature for (B) ref. 46 and American Chemical Society for (C) ref. 6.





degrade in Cu(II)-SG and disulfide (Cu(II)-GSSG) under physiological conditions.<sup>6</sup> Surprisingly, 90% of the Cu collected in the urine 2 h pi was still in the form of CuSG, while 60% of Cu in the liver was in the form of Cu(II)-GSSG-complexes bound to serum proteins. Indeed, CuSG had a higher resistance to serum protein adsorption than Cu(II)-GSSG. This explains why these NCs were quickly eliminated and less accumulated in the liver than the degradation product (Fig. 4C).

### 3.3 Protein corona formation

Proteins can be adsorbed onto the surface of NPs forming the protein corona.<sup>113,116</sup> The protein coating is a dynamic system with competition between different proteins until an equilibrium is reached. The first layer of proteins is rapidly adsorbed onto the surface of the particles to form the soft corona. In the second step, proteins that have a stronger affinity to the surface of the particles slowly remove the first layer of proteins to generate a stable layer: the hard corona. This means that the composition and thickness of the protein corona evolve with time.<sup>117</sup> The affinity and exchange rate of the protein corona depend on the size, charge, composition, and shape of the NPs, the incubation conditions (temperature, concentration, and time), the type of proteins, and their stability.<sup>116,117</sup>

The formation of this protein layer will have an impact on cell internalization, bio-distribution, and toxicity of the NPs,<sup>118,119</sup> and NCs.<sup>120</sup>

Concerning NCs, it was established that incubation of Au NCs with an increasing concentration of human serum albumin (HSA) improved their fluorescence intensity by 6 at high protein concentrations (3  $\mu$ M).<sup>121</sup> This was probably due to the adsorption of proteins that decreased the non-radiative recombination by forming a rigid protection layer or inducing new metal-ligand interactions.

The surface charge strongly affects the amount and composition of the protein corona.<sup>109</sup> Indeed, negative NCs interact preferentially with Apolipoprotein (Pi > 5.5), whereas positive NCs prefer albumin (Pi < 5.5), and as a result, positive NCs accumulate in the spleen, lungs, heart, and kidneys, while negatively charged NCs will rather end up in the liver and testis.<sup>109</sup>

Neutral surfaces obtained with PEGylated ligands or with zwitterionic molecules (SG or BSA) do not bind proteins. However, the non-fouling properties observed with zwitterionic systems are pH and particle curvature dependent.<sup>14,55,115</sup>

### 3.4 Density

The size of a nanoparticle is important, but its density (*i.e.* mass/volume) may also have a strong impact. Increasing the density of a NC will augment its interaction with the surface of the blood vessels. NCs stabilized by SG but with different metal cores (Au, Ag, and Au/Ag alloy) were synthesized.<sup>122</sup> As compared to high density AuSG, the low-density AgSG presented low affinity for the blood vessels, faster clearance, and shorter retention time.

### 3.5 Electrical charge

The charge of the NCs also determines the affinity of the NCs for the blood vessels. AuSG NCs functionalized with ethanediamine

or ethanedioic acid to generate positively and negatively charged surfaces were more slowly excreted than neutral NCs over a 90 day period.<sup>109</sup> Just 24 h after intraperitoneal administration, the amount of positively charged NCs in the kidneys was two times higher than those of neutral NCs. Various studies have demonstrated that, after glomerular filtration, positively charged compounds bind through their cationic sites to megalin receptors overexpressed on negatively charged proximal tubule epithelial cells.<sup>66,67,123</sup> This re-absorption phenomenon could explain that small positively charged compounds were slowly excreted by the kidneys.

### 3.6 Influence of the concentration of injected Au NCs

Finally, the concentration of the injected Au NCs also influences their transport by the blood flow.<sup>124</sup> An interesting two-compartment kidney elimination process was observed for AuSG intravenously injected into mice at nine different concentrations (from 0.15 to 1 059 mg kg<sup>-1</sup> of body weight). At low concentrations (<15 mg kg<sup>-1</sup>), kidney elimination was constant at 35% ID. Then, the renal elimination linearly increased with the concentration of NCs. One explanation is that, for small doses, the AuSG could easily cross the blood vessel wall and enter the extravascular space. The reduction in their blood concentration slows down the renal process. At higher concentrations, the Au NCs are strictly confined to the blood vessels, more rapidly transported by the blood flow, and more efficiently eliminated through the kidneys.

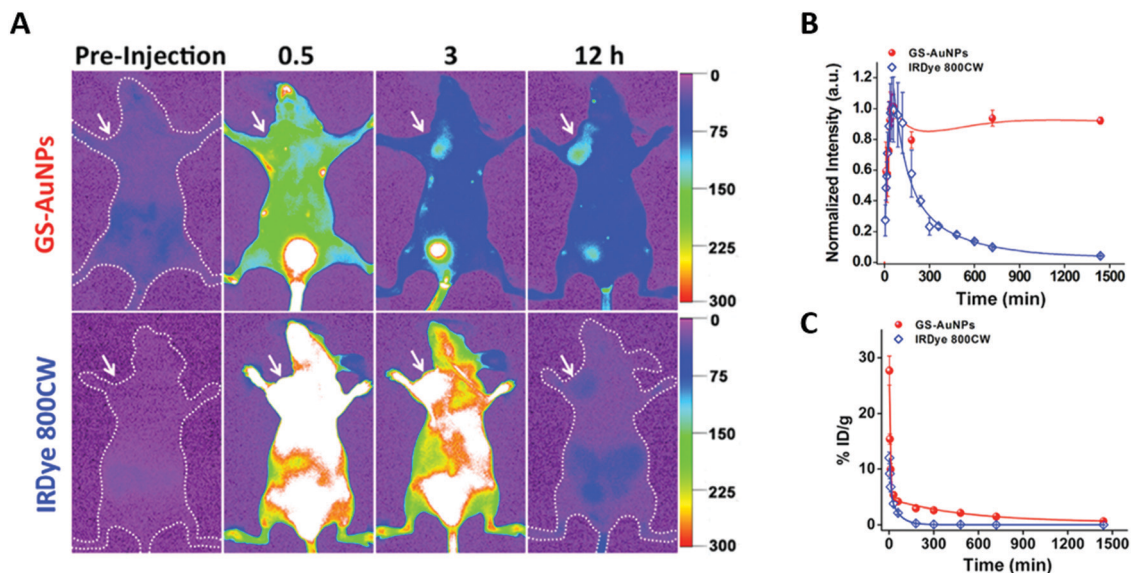
## 4 Tumor targeting of Au NCs

If the renal clearance of the NCs is reckoned as highly relevant to reduce their toxicity, these species should still have enough time to circulate in the body in order to accumulate passively and/or specifically in the tumor microenvironment. The growth of a tumor generally induces the formation of new blood vessels with pore sizes between 300 and 1200 nm and a perturbation of the lymphatic drainage. This allows NP accumulation in cancer tissues at higher concentrations and for a longer time than in normal tissues. This phenomenon is called the Enhanced Permeability and Retention (EPR) effect. Maeda *et al.*<sup>125</sup> were one of the first groups who studied the EPR effect in 1986. They discovered that NPs with a size larger than the KFT can circulate in the blood at high concentrations and for longer times (at least 6 h) to allow a passive uptake.

### 4.1 Passive accumulation of renal-clearable Au NCs by the EPR effect

Au NCs combine a high renal clearance with surprisingly elevated passive uptake in MCF-7 tumors.<sup>30</sup> As can be seen in Fig. 5A, AuSG and IRDye 800CW present a rapid whole-body distribution and a maximum tumor accumulation 40 min pi. However, 24 h pi, the concentration of NCs in the tumor was 10 times higher than that of IRDye 800CW (Fig. 5B). This augmented EPR effect probably occurred because AuSG NCs have a longer blood elimination half-life ( $t_{1/2\beta}$ ) than IRDye (Fig. 5C).





**Fig. 5** Tumor uptake and renal elimination in mice intravenously injected with AuSG NCs and IRDye 800CW. (A) *In vivo* NIR fluorescence images of MCF-7 tumor-bearing mice 0.5, 3, and 12 h after intravenous injection of AuSG NCs and IRDye 800CW (200  $\mu$ L at 20 mg mL<sup>-1</sup> and 10  $\mu$ M, respectively). The tumor areas are indicated by arrows. (B) MCF-7 tumor uptake of AuSG NCs and IRDye 800CW 24 h pi. (C) Renal elimination of AuSG and IRDye 800CW, intravenously injected (200  $\mu$ L at 7 mg mL<sup>-1</sup> and 10 mM, respectively) at 24 h pi. The curves were fitted to a biexponential function with  $R^2$  values of 0.9711 and 0.9838, respectively. The distribution half-life ( $t_{1/2\alpha}$ ) values are  $5.4 \pm 1.2$  and  $6.3 \pm 2.5$  min, respectively, and the  $t_{1/2\beta}$  values are  $8.5 \pm 2.1$  and  $0.98 \pm 0.08$  h, respectively, for AuSG NCs and IRDye 800CW ( $n = 3$ ). Reproduced from ref. 30 with permission from the American Chemical Society.

Compared to larger Au NPs that are not cleared by the kidneys, AuSG NCs present a two times higher tumor accumulation. This could be attributed to the 10 times lower accumulation of AuSG in the RES compared with Au NPs, which also maintained the highest NC concentration in the blood during the first hours. This implies that NCs escaping RES uptake and remaining long enough in circulation in the blood stream will have a better chance to accumulate in tumors *via* the EPR effect.

**4.1.1 Increasing the circulation time by reducing Au NC elimination by the kidneys.** Au NCs stabilized with SG or PEG ligands with close HD (3.3 and 5.5 nm, respectively) were tested<sup>14</sup> in order to study the impact of the ligand on renal elimination and tumor targeting. Both NCs were excreted to a similar extent at 24 h pi in the urine, but their kinetics of elimination were different. The fluorescence intensity of the bladder reached a maximum at 1 h for the AuSG, while it took 5 h for the AuPEG. It was thus not surprising that AuPEG showed a tumor accumulation in MCF-7 tumors three times higher than that of AuSG at 12 h pi. Similar results were obtained with Au<sub>25</sub>SG<sub>18</sub> (~1.9 nm) and AuZwMe<sub>2</sub> (~2.36 nm)<sup>33,112</sup> as well as when Au<sub>25</sub>SG<sub>18</sub> were compared with smaller Au<sub>10-11</sub>SG<sub>10-11</sub>, Au<sub>15</sub>SG<sub>13</sub>, Au<sub>18</sub>SG<sub>14</sub><sup>46</sup> (Fig. 6). However, increasing the size of NCs with Au<sub>201</sub> and Au<sub>640</sub> had no impact on tumor uptake.

Overall, this confirms that increasing the circulation time of the NCs favors their passive uptake by the tumor.

**4.1.2 Increasing the cellular interaction.** Neutral, positively, or negatively functionalized AuSG will not accumulate equally in tumors.<sup>37</sup> One day after intra-peritoneal administration, 149, 222, and 320 ng g<sup>-1</sup> of Au NCs were measured in U14 tumors for neutral, positive, and negative NCs, respectively. The fact that charged Au NCs show higher tumor accumulation than

neutral NCs could be explained by different phenomena. The negatively charged cell membranes favor the adsorption of positively charged NCs, thus augmenting their binding to the cells and internalization.<sup>126</sup> For the negative ones, the enhancement of the EPR effect is more likely due to the lower protein corona adsorption.<sup>127</sup>

## 4.2 Active tumor targeting

The addition of a targeting agent, usually a small (bio)molecule that could specifically interact with receptors overexpressed at the surface of tumor cells is expected to augment the specific tumor accumulation. Cell surface proteins such as integrins, transferrin, and folate receptors (FRs) are commonly targeted because they are overexpressed in tumors.<sup>54,61,64,65,101-103,106,108,128</sup>

AuBSA functionalized with folic acid (FA)<sup>51</sup> provided maximum tumor/normal tissue ratios (T/N) in mice with FR-positive HCT116 tumors two times higher than for negative A549 tumor. The functionalization of AuBSA with hyaluronic acid (HA)<sup>61</sup> (Fig. 7) or Luteinizing hormone-releasing hormone (LHRH)<sup>83</sup> allowed increasing the tumor accumulation by three compared to the AuBSA.

Grafting an RGD-containing peptide on AuBSA can improve the targeting of human glioblastoma U87MG tumors engrafted in mice.<sup>86</sup> U87MG cells express integrin  $\alpha_v\beta_3$ , a cell surface receptor recognized by RGD. If, 1 h after the intravenous injection of AuBSA, the tumor was detected due to the EPR effect, the signal dramatically decreased after 2 h. In contrast, the NIR fluorescence of the RGD-Au in tumors was still visible after 24 h. Thus, functionalized Au NCs with a targeting molecule could improve the tumor accumulation of the Au NCs and prolong their retention time.







**Fig. 6** Effect of the subnanometer size of the NCs on tumor uptake: (A) MCF-7 tumor accumulation efficiency of different-sized Au NCs *versus* the number of gold atoms at 24 h p.i. For NCs smaller than Au<sub>25</sub>SG<sub>18</sub> (Au<sub>10-11</sub>SG<sub>10-11</sub>, Au<sub>15</sub>SG<sub>13</sub>, Au<sub>18</sub>SG<sub>14</sub>), the tumor accumulation decreased with increasing number of gold atoms. For NCs larger than Au<sub>25</sub>SG<sub>18</sub> (Au<sub>201</sub> and Au<sub>640</sub>), tumor uptake was constant. \**P* < 0.05, \*\**P* < 0.005, Student's *t*-test. (B) Blood pharmacokinetics of Au<sub>10-11</sub>SG<sub>10-11</sub>, Au<sub>15</sub>SG<sub>13</sub>, Au<sub>18</sub>SG<sub>14</sub>, and Au<sub>25</sub>SG<sub>18</sub> show two-compartment pharmacokinetics with different distributions and *t*<sub>1/2β</sub> between the NCs. BALB/c mice (*n* = 3) were intravenously injected (~100 μM, 100 μL). Reproduced from ref. 46 with permission from Springer Nature publishing group.



**Fig. 7** *Ex vivo* fluorescence images of the tumor and of the major organs in mice 7 h p.i. of HA-AuBSA NCs (A) or free HA and AuBSA NCs (B) with the corresponding average fluorescence intensity analysis (C and D). The addition of the targeting ligand HA grafted onto the Au NC surface improves the tumor uptake by two times. Reproduced from ref. 61 with permission from The Royal Society of Chemistry publishing group.

In another study, a dual targeting was tested. A cRGD peptide and a 26-base G-rich DNA oligonucleotide that functions as a nucleolin-binding aptamer (AS1411) were added to Au NCs to induce a dual targeting.<sup>38</sup> The nucleolin receptors are overexpressed on the cell membrane of most of the cancer cells. The tumor signal disappeared 8 h pi when naked Au NCs were

used. In contrast, it was still detectable 48 h later when only one targeting molecule was added. This confirmed the interest of using targeted Au NCs to augment their retention time in the tumor. If both ligands were co-presented by the Au NCs, the T/N fluorescence ratio was increased to a greater extent (7.2) than when the Au NCs were labeled with only cRGD (5.4).



If the addition of a targeting agent on the Au NC surface favors higher and longer tumor accumulation,<sup>38,103–105,107</sup> it also increases the size and changes the charge of the Au NCs, which could be at the expense of their renal elimination.

## 5 Au NCs: promising theranostic agents

Au NCs are useful for multimodal cancer diagnosis and therapy.

### 5.1 Multimodal diagnostic applications

Many imaging techniques are used daily in hospitals for diagnosis, image-guided surgery, and follow-up of treatment efficacy. We will present only imaging techniques that have already been tested with Au NCs in animal models.

**5.1.1 Optical imaging.** Optical imaging (OI) is a multiscale (from molecules to cells and up to the patient), noninvasive technique<sup>129</sup> that facilitates real-time<sup>79</sup> *in vivo* monitoring with high sensitivity<sup>130,131</sup> and temporal resolution to visualize dynamic processes.<sup>111</sup>

However, OI also presents drawbacks and in particular suffers from a weak penetration of light in biological tissues, especially because hemoglobin absorbs light below 650 nm while water will absorb wavelengths above 900 nm.<sup>129,132</sup> In addition, in the visible range ( $\pm 400$ –700 nm), auto-fluorescence of some tissues (skin especially) leads to a high background noise that reduces the optical contrast. For all these reasons, it is particularly interesting to work in the NIR wavelength range between 700 and 900 nm, called the “transparent imaging window.”<sup>79,129</sup>

Au NCs, with their water solubility, biocompatibility, tunable PL from the UV to the NIR, and resistance to photobleaching in contrast to organic fluorophores, are potentially good optical probes. In the earliest works, the low emission wavelength at 480 nm of Au NCs stabilized with histidine (His)<sup>23</sup> limited their use to superficial tissues or cell imaging. A hydrophilic indocyanine green fluorophore needed to be added to have an emission at 800 nm that can be followed *in vivo*. In the case of AuBSA, even though the 680 nm emission<sup>58,61,133</sup> enabled their detection under few millimeters of tissues, the conjugation with an indocyanine green fluorophore also permitted a shift of the emission in the NIR and facilitated their detection.<sup>62</sup> As mentioned in the first part, several strategies exist to shift the emission to higher wavelengths and to increase the QY of the NCs for improving their limit of detection. In the meantime, other NCs such as AuSG<sup>14,30</sup> and AuPEG<sup>14</sup> with intrinsic NIR fluorescence were employed.

Seminal works by H. Dai<sup>79,132,134</sup> have demonstrated the benefit in terms of spatial and temporal resolutions to move from the NIR to a SWIR spectral window with the parallel development of new contrast agents and camera.<sup>79</sup> LA-sulfobetaine-capped Au NCs with relatively good fluorescence in the NIR/SWIR region (QY of 0.6% at 1000 nm and 3.8% at 900 nm) and broad emission were produced.<sup>78</sup> After their intravenous injection, images of the blood vessels were obtained with higher contrast and

spatial resolution in the SWIR using a long-pass filter at 1250 nm than in the NIR.

Au NCs thus appeared as new promising SWIR (Fig. 8A) and NIR (Fig. 8B) contrast agents.

**5.1.2 X-ray CT.** X-ray CT is also a noninvasive technique<sup>31</sup> with a high 3D resolution that offers anatomical information, but suffers from low sensitivity.<sup>31,131,136,137</sup> An additional contrast agent is often required to distinguish soft tissues.<sup>31,137</sup> Iodine and gold are used as X-ray contrast agents;<sup>31,52,137,138</sup> however, the short circulation time, nonspecific distribution, and potential renal toxicity of iodine limit its application. The idea is to find a compound slightly larger to increase the circulation time: Au NCs satisfy this criterion.<sup>114</sup>

AuBSA were thus tested in mice, allowing an improved visualization of the structure of major organs (heart, liver, kidneys, bladder, and intestine) and a clear delineation of the calyces, pelvis, ureters, and bladder (Fig. 8C) making AuBSA NC a promising candidate to study renal excretion.<sup>31</sup> Other NCs such as AuSG were also successfully tested.<sup>32</sup>

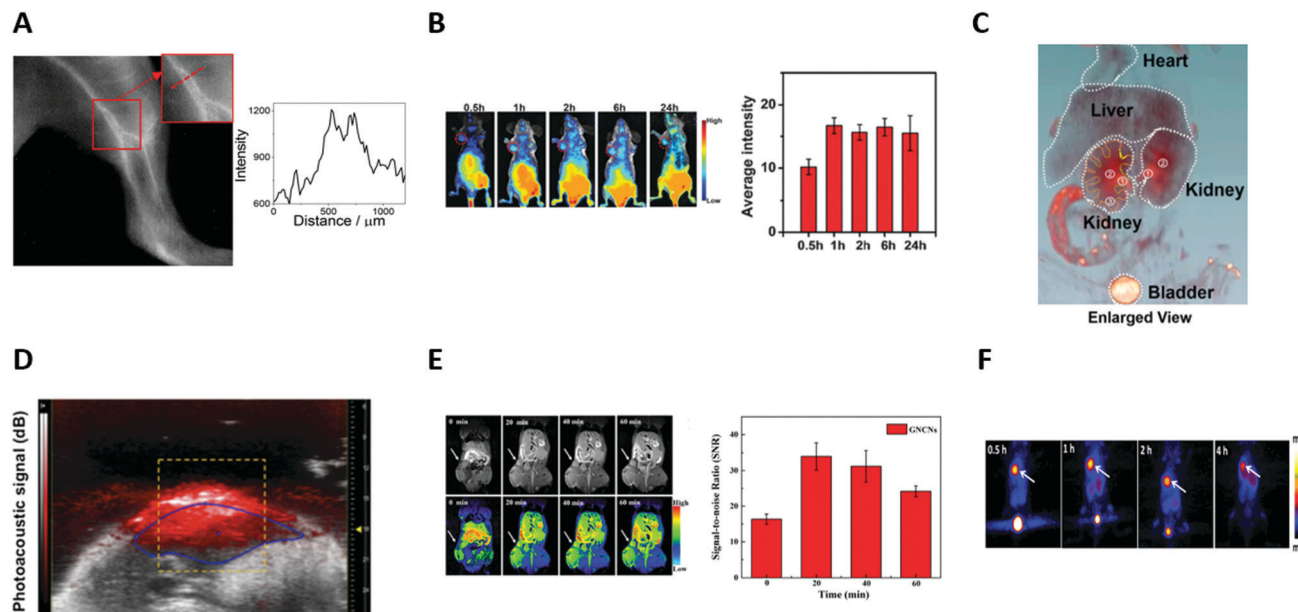
Because the Au NCs are also fluorescent, they can serve as bi-modal imaging agents.<sup>60</sup> X-ray CT images provide anatomical information, but can also be used to confirm the results obtained by optical imaging, such as tumor uptake<sup>139</sup> or renal elimination of AuSG.<sup>32,35,36,114</sup>

**5.1.3 Photoacoustic.** Au NCs have recently been employed in photoacoustic imaging<sup>140</sup> and OI<sup>33</sup> (Fig. 8D). AuZw with different metal core sizes and ligand thicknesses were produced, in order to investigate the influence of the metal/ligand ratio on their optical and photoacoustic properties. When the amount of ligand increased, the shell became more rigid, which decreased the nonradiative loss due to vibrations of the ligand. This was associated with an increased fluorescence but decreased photoacoustic signal. However, by decreasing the quantity of ligand, the core of the NCs became larger. Upon illumination, the metallic core produced more heat, which generated more intense vibrations and ultimately improved the photoacoustic signal. Because of this opposite evolution in the signal, Au NCs should be carefully chosen, depending on the technique that will be required for the experiment.

**5.1.4 Magnetic resonance imaging.** Magnetic resonance imaging (MRI) is a noninvasive technique<sup>141</sup> largely employed in hospital and clinical settings based on the observation of water's protons contained in the organism when an external magnetic field is applied. This technique allows 3D reconstruction<sup>131,141</sup> with high spatial resolution,<sup>137,141</sup> but suffers from low sensitivity.<sup>131,137,141</sup> Gadolinium (Gd<sup>3+</sup>) is often used as a contrast agent for MRI detection.<sup>107,135,142</sup>

The use of Au NCs decorated with Gd<sup>3+</sup> ions compared to traditional Gd chelates (diethylenetriamine penta-acetic acid, DTPA) improved the performance of Gd as a MRI contrast agent itself and prolonged its detection time three-fold<sup>135</sup> (Fig. 8E). These Au NCs were still visible by X-ray CT or optical imaging techniques which permit visualization of the organs, where MRI had a weak signal-to-noise ratio, making Gd-functionalized Au NCs promising tools for tri-modal imaging: MIR/X-ray CT/optical imaging.





**Fig. 8** (A) Image of the left leg of a wild-type CS7BL/6 mouse taken using an InGaAs camera equipped with a 1250 LP filter after the intravenous injection of LA-sulfobetaine-capped Au NCs (0.5 mg of Au). Signal intensity across a line of interest drawn in the inset images showed the spatial resolution of the vessels. (B) *In vivo* fluorescence images of a HeLa tumor-bearing nude mouse at different time points (30 min, 1 h, 2 h, 6 h, and 24 h) after intravenous injection of cRGD-Au NCs (200  $\mu$ L, 8.0 mg mL<sup>-1</sup>). Red circles indicate the tumor. Quantification of the signal intensity of the tumor side ( $n = 3$ ). (C) *In vivo* 3D CT images of the heart, liver, kidneys, and bladder (excluding bone) in mice 2 h after intravenous injection of AuBSA NCs (200  $\mu$ L, 9.5 mgAu per mL). The ureter, renal pelvis, and the major calyx are marked in orange and yellow dashed curves. (D) Noninvasive *in vivo* photoacoustic imaging of the belly of a NMRI nude mouse, 3 h after intravenous injection of AuZw NCs (200  $\mu$ L, 600  $\mu$ M). (E) *In vivo* T1-weighted MR images of A549 tumor in a BALB/c-nude mouse at different time points (20, 30, and 40 min) after intratumoral injection of Gd-AuSG NCs (50  $\mu$ L, 0.11 M). White arrows indicate the position of the tumor. The graph represents the signal-to-noise ratio around the tumor ( $n = 3$ ). (F) Representative PET images of coronal single slices of an orthotopic A549 lung tumor-bearing mouse at different time points (0.5, 1, 2, and 4 h), after intravenous injection of 6.7 MBq of [64Cu]CuNC@BSA-LHRH. White arrows indicate the position of the tumor. Reproduced with permission from American Chemical Society for (A) ref. 78, (C) ref. 31 and (F) ref. 104; The Royal Society of Chemistry for (B) ref. 106, AIP for (D) ref. 33 and Elsevier for (E) ref. 135.

**5.1.5 Positron emission tomography.** The last noninvasive imaging method described with Au NCs is positron-enhanced tomography (PET). PET has the highest detection sensitivity in human<sup>39,143</sup> but a poor spatial resolution.<sup>131</sup>

<sup>64</sup>Cu, a well-known  $\beta^+$  radioisotope, is generally complexed with bifunctional chelates, such as DOTA (1,4,7,10-tetraazacyclododecane-1,4,7,10-tetraacetic acid) and TETA (1,4,8,11-tetraazacyclododecane-1,4,8,11-tetraacetic acid). However, these complexes suffer from low stability, and the released copper ions will be complexed with proteins to be finally stored in the liver. This could induce toxic effects.<sup>104</sup>

The direct incorporation of <sup>64</sup>Cu in NCs may overcome these limitations and the ligand shell could favor the radiolabeling stability (Fig. 8F). These NCs keep their initial properties, which means renal excretion and the ability to escape the RES.<sup>6,104</sup>

For example, CuSG NCs are progressively degraded into Cu(II)-GSSG.<sup>6</sup> Due to their interaction with serum proteins, they were then retained in the liver. After intravenous injection in mice, CuSG or Cu(II)-GSSG showed  $t_{1/2\beta}$  of 3.2 and 4.9 h, respectively. The successful addition of <sup>64</sup>Cu, with its half-time of decay of 12.7 h, helped in following the elimination of CuSG with precision. Indeed, the strong signal from the kidneys and the bladder just 1.5 min pi confirmed the fast renal elimination of [<sup>64</sup>Cu]CuSG. The signal from the bladder still

visible 4 h pi attested the constant elimination of the NCs from the body. Finally, at 4 h pi, the signal detected in the liver confirmed the dissociation of CuSG into Cu(II)-GSSG complexes.

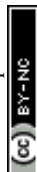
Radioactive NCs can thus be used for dual imaging. PET offers more sensitive and quantitative follow-up, which will be an addition to the benefits of optical imaging.<sup>104,144</sup>

## 5.2 Therapy

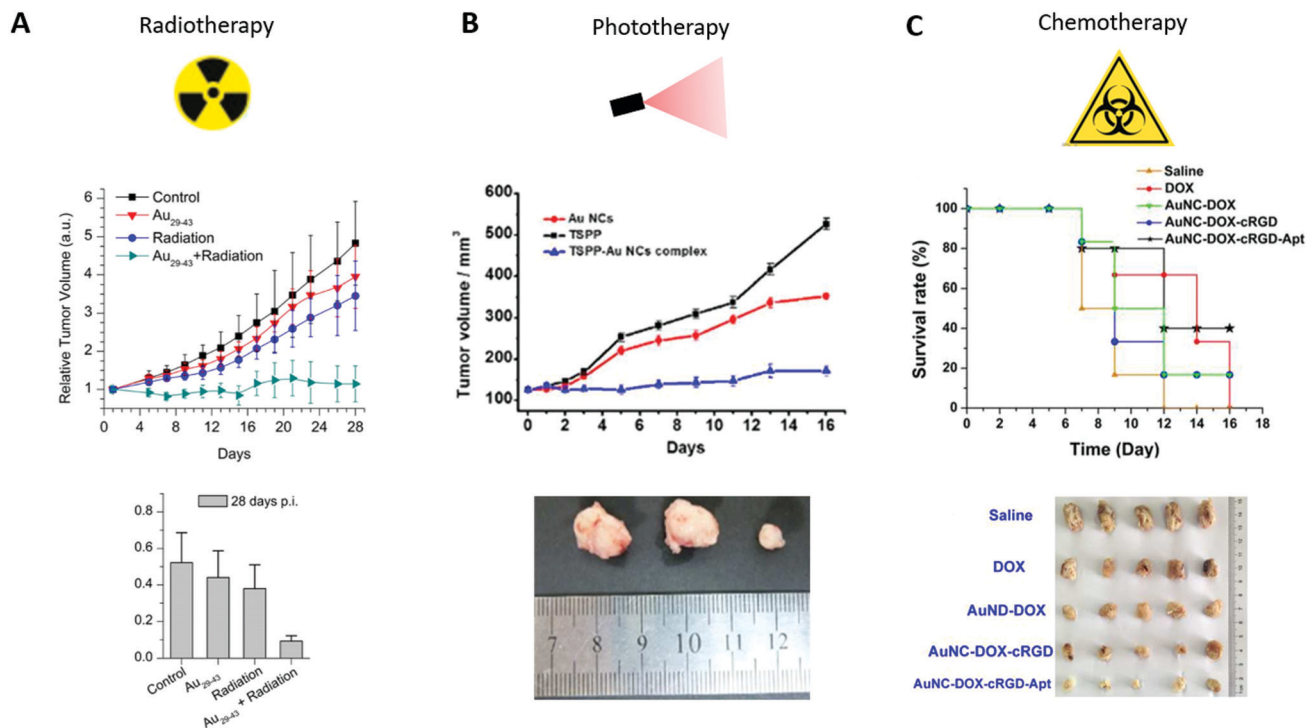
In addition to their use in diagnostics, Au NCs can be used as therapeutic agents and delivery systems.

**5.2.1 Radiotherapy.** Gold can serve as a radiosensitizer<sup>34–36</sup> if it is formulated to be biocompatible, to have an efficient renal clearance and good tumor accumulation capability.

Zhang *et al.*<sup>35</sup> first studied the potential of a Au<sub>10–12</sub>SG<sub>10–12</sub> or of a saline solution after intraperitoneal administration (0.2 mL, 3 mM) in nude mice engrafted with a human cervical U14 tumor. When the tumor uptake reached a maximum (24 pi), the mice were either irradiated (under <sup>137</sup>Cs gamma radiation of 3600 Ci at 5 Gy) or not. Regarding the controls, no decrease in tumor size was observed in mice treated with only the Au NCs over a period of 23 days. In contrast, the tumor volume decreased by 8% upon irradiation and 65% with the combination of Au NCs plus irradiation.







**Fig. 9** (A) Time-course studies of the U14 tumor volumes until 28 days pi (top) and measurement of the U14 tumor weights at 28 days pi (bottom) of mice intraperitoneally injected with saline solution (control) or with  $\text{Au}_{29-43}\text{SG}_{27-37}$  (5.9 mgAu per kg of body), nude mice treated with only radiation (under  $^{137}\text{Cs}$  gamma radiation of 3600 Ci at 5 Gy, 24 h pi), and nude mice injected with  $\text{Au}_{29-43}\text{SG}_{27-37}$  followed by irradiation ( $n = 8$ ). (B) Time-course studies of U87 tumor volumes until 16 days pi (top) of mice intraperitoneally injected with Au NCs (0.1 mL, 5 mM), TSPP (0.1 mL, 0.1 mM) and TSPP-Au NCs (equivalent 0.1 mL, 0.1 mM TSPP) ( $n = 3$ ) and corresponding images isolated from tumor-bearing mice after 16 days of treatment (Au NCs, TSPP and TSPP-Au NCs from left to right, bottom). (C) Survival rate curves of U87MG tumor-bearing mice intravenously injected with free DOX, DOX-Au NCs, cRGD-DOX-Au NCs, and Apt-cRGD-DOX-Au NCs (all at  $5.0 \mu\text{g kg}^{-1}$  equivalent DOX) and corresponding images isolated from tumor-bearing mice after 14 days of treatment (bottom). Reproduced from ref. 36, 38 and 133 with permission from Scientific Reports, The Royal Society of Chemistry and Elsevier publishing group.

The same experiment was carried out with  $\text{Au}_{29-43}\text{SG}_{27-37}$ .<sup>36</sup> Compared with the control group, the tumor volume in mice treated with only radiation and mice treated with both  $\text{Au}_{29-43}\text{SG}_{27-37}$  and radiation decreased by 10% and 76%, respectively. No effects were observed in mice injected only with Au NCs (Fig. 9A). In both cases, Au NCs thus generated a toxic effect only if they were irradiated. Similar results were obtained with AuBSA.<sup>34</sup>

**5.2.2 Phototherapy.** The second well-developed approach in cancer treatment is phototherapy, divided into photo-dynamic therapy (PDT) and photothermal therapy (PTT) but the potential of Au NCs as PTT agents in mice has not yet been proven.

Concerning PDT,  $\text{Au}_{25}\text{SG}_{18}$  can generate singlet oxygen species under 650 and 808 nm irradiation in solution<sup>145</sup> but most of the current studies use Au NCs combined with photosensitizer molecules such as chlorine6 (Ce6)<sup>37,39,146,147</sup> or porphyrin derivatives.<sup>37,133</sup>

Most of the PDT compounds present unwanted cutaneous photosensitivity, and inadequate selectivity that can be ameliorated when combined with NCs. A four-time decrease in the size of C6 tumors was obtained 8 days after the intravenous injection of Au NCs grafted with protoporphyrin IX, followed by laser treatment 3 h later (532 nm, 15 min,  $1.5 \text{ W cm}^{-2}$ )<sup>37</sup> compared to the control treated with only radiation. Moreover,

U87 tumor growth can be stopped, after intratumoral injection of Au NCs conjugated with TSPP (meso-tetra(4-sulfonatophenyl)-porphyrine dihydrochloride), a photosensitizer, in U87 tumors, and light excitation for 30 min per day with IR light indicating that Au NCs can enhance the PDT effect of the photosensitizer alone<sup>133</sup> (Fig. 9B).

In order to study whether this synergistic effect could be generalized, C. Zhang *et al.*<sup>39</sup> intravenously injected Ce6- $\text{Au}_{25}\text{SG}_{18}$ , free Ce6, or PBS into mice with human gastric MGC-803 tumors of 150 mm<sup>3</sup>. The tumors were illuminated (633 nm, 10 min,  $100 \text{ mW cm}^{-2}$ ) 8 h and 24 h later. After 21 days, the volumes of the tumors for the mice injected with PBS, Ce6, and Ce6- $\text{Au}_{25}\text{SG}_{18}$  without laser treatment were similar and close to 1200 mm<sup>3</sup>. However, the tumor growth slowed down and reached a volume of around 650 mm<sup>3</sup> or 350 mm<sup>3</sup>, in mice injected with free Ce6 or Ce6- $\text{Au}_{25}\text{SG}_{18}$  followed by illumination. This thus confirmed that Au NCs can enhance the radiotherapeutic activity of a co-injected radiosensitizer.

Based on this observation, it appeared interesting to generate a core-shell structure using silica<sup>146</sup> or liposomes<sup>147</sup> to increase the amount of Ce6 in contact with the Au NCs. The compounds were intravenously injected into mice with melanoma MDA-MB-435 or breast MDA-MB-361 tumors for compounds with silica or liposome shells, respectively.



The tumors were irradiated for 10 min using laser (671 nm, 100 mW cm<sup>-2</sup>) for the silica model and 3 times every 3 days for 25 min (660 nm, 150 mW cm<sup>-2</sup>) for the liposome model. In both tumor models, regarding the control, the injection of free Ce6 slowed down the growth of the tumors almost 2 times, while the core-shell structure composed of Ce6 and Au NCs totally inhibited the growth of the tumors. These studies confirmed that Au NCs can enhance the effect of the Ce6 radiosensitizer even after encapsulation in silica or liposomes.

**5.2.3 Chemotherapy.** Au NCs can also be used for drug delivery, as demonstrated with the delivery of doxorubicin (DOX), a DNA-intercalating agent.<sup>38</sup> After repeated intravenous administration every 48 h for 2 weeks, mice with U87MG tumors presented reduced tumor growth with increased necrosis as compared to controls (Fig. 9C).

More complex systems combining multimodal imaging and therapeutic compounds were generated<sup>146</sup> combining Au NCs functionalized with Ce6 carbon dots. These theranostic agents were usable for fluorescence, MRI and PDT cancer treatments.

To conclude, Au NCs are promising multimodal imaging probes. Due to their intrinsic properties, toxic effects can be generated in the microenvironment of the tumors upon activation by an external stimulus like X-ray radiation or illumination, with reduced damage to healthy tissues. They can be used as therapeutic compounds alone or to enhance the effects of sensitizer agents.

The multimodal properties of the Au NCs make these theranostic agents highly promising in oncology.

**5.2.4 Optical-guided surgery.** Biocompatible zwitterionic or pegylated ligands containing Au NCs present augmented optical properties and plasma half-life that can improve their passive accumulation in an orthotopic animal model of Head and Neck Squamous Cell Carcinoma (HNSCC). This can serve to detect the presence of tumors in the mouth using non-invasive optical imaging, and can be used intra-operatively by surgeons to perform optical-guided surgery with improved survival rates as compared to standard surgery. Indeed, the presence of fluorescent Au NC-labeled tumor margins allows the surgeon to better remove all tumor infiltrated surrounding tissues that are not visible by the naked eye, thus reducing the appearance of local relapse.<sup>100</sup>

## 6 Conclusions and perspectives

In this review, we described recent works performed on metal NCs, mainly gold, for *in vivo* applications. Progress in synthesis methods has enabled the development of effective strategies to produce photoluminescent atomically precise NCs on a large scale, with high purity and different functional groups.

The unique physicochemical properties and good biocompatibility of metal NCs have facilitated their use in biomedical applications. The PL and metallic composition of NCs allow their use as optical or X-ray probes, or radiosensitizers. By integrating Au NCs with other imaging or therapeutic agents, complex systems could be achieved. These combinations result in powerful image-guided therapy that takes advantage of each

technique. Finally, their fast renal elimination decreases their nonspecific accumulation in different organs, and thus, their toxicity. These optical features combined with their low toxicity and rather simple synthesis method make them very good candidates as contrast agents with respect to other nanomaterials such as Quantum Dots, organic fluorophores, lanthanide NPs, and carbon nanotubes.

Different parameters such as size, length, and charge of the ligands or density affect the formation of the protein corona, optical properties, biodistribution, targeting potential, renal elimination, tumor uptake, and biodegradability of metal NCs. Each of these parameters must be carefully addressed when designing NCs for specific biomedical applications and their possible clinical translation.

Au NCs have already been applied *in vivo* on mice and monkeys.<sup>124</sup> They have shown promising results as theranostic agents, even in parts of the body difficult to access. NCs are not limited to cancer treatments and can be used for diagnosis of other diseases such as renal dysfunctions,<sup>31,148,149</sup> and Alzheimer's disease.<sup>150,151</sup>

However, some of their properties such as their tumor accumulation and retention or their luminescent properties in the NIR and SWIR regions for *in vivo* imaging still need to be improved. Further studies are necessary to confirm their radio and photosensitizing effects and more broadly, to get a deeper understanding of the relationships among structure, stability, and luminescence of Au NCs.

Precise and early diagnosis of tumors and the development of targeted therapies are two major lines of research in oncology. In this context, we believe that photoluminescent Au NCs are promising candidates. Their biomedical potential will go beyond cancer research and toward a wide range of (biomedical) applications (such as fluorophores for fluorescence resonance energy transfer applications,<sup>152,153</sup> biosensors, and optoelectronics).

Once their pharmacokinetic and luminescence properties, combined with their proven theranostic potential under light or X-ray irradiation, are clearly demonstrated at the preclinical level, we can expect a rapid translation of such ultrasmall Au NCs in clinical trials because of their method of preparation, low toxicity and powerful elimination. In addition, they will benefit from the fact that the Food and Drug Administration has already approved different gold nanoparticles for diagnostics and therapy.

## Abbreviations

BSA	Bovine serum albumin
CI	Contrast index
cRGD	Cyclic RGD
DTPA	Diethylenetriamine penta-acetic acid
EPR	Enhancement permeability and retention
FA	Folic acid
FR	Folate receptor
HA	Hyaluronic acid



HD	Hydrodynamic diameter
H	Hour
ID	Injected dose
ID per g	Percentage of injected dose per gram of tissue
KFT	Kidney filtration threshold
LA	Lipoic acid
LHRH	Luteinizing hormone-releasing hormone
LIBS	Laser induced breakdown spectroscopy
LMCT	Ligand-to-metal charge transfer
LMMCT	Ligand-to-metal-metal charge transfer
MRI	Magnetic resonance imaging
NCs	Nanoclusters
NIR	Near-infrared
NPs	Nanoparticles
PDT	Photodynamic therapy
PEG	Polyethylene glycol
PET	Positron emission tomography
pi	Post injection
PL	Photoluminescence
PNA	Peptide nucleic acid
PTT	Photothermal therapy
QY	Quantum yield
ROS	Reactive oxygen species
RES	Reticuloendothelial system
SG	Glutathione
SWIR	Shortwave infrared
$t_{1/2\alpha}$	Distribution half-life
$t_{1/2\beta}$	Elimination half-life
UV	Ultraviolet
X-ray CT	X-ray computed tomography
Zw	Zwitterion molecules

## Conflicts of interest

There are no conflicts to declare.

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