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# Insights into the nonlinear optical (NLO) response of pure $Au_m$ ( $2 \geq m \leq 7$ ) and copper-doped $Au_m-xCu_x$ clusters†

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A series of small pure  $Au_m$  ( $2 \geq m \leq 7$ ) and copper-doped  $Au_{m-x}Cu_x$  clusters was evaluated by density functional theory (DFT) at the CAM-B3LYP/LANL2DZ level for their geometric, electronic, and nonlinear optical (NLO) properties. The charge transfer for the Au cluster significantly improved by reducing the HOMO–LUMO energy gap from 3.67 eV to 0.91 eV after doping with Cu atoms. The doping of Cu also showed noteworthy impacts on other optical and NLO properties, including a decrease in the excitation energy and increase in the dipole moment and oscillator strength. Furthermore, changes in the linear isotropic and anisotropic polarizabilities ( $\alpha_{iso}$  and  $\alpha_{aniso}$ ) and first and second NLO hyperpolarizabilities ( $\beta_{static}$ ,  $\gamma_{static}$ ) were also observed in the pure and Cu-doped clusters, which enhanced the NLO response. The nonlinear optical properties of the clusters were evaluated by calculating the static and frequency dependent second- and third-order NLO polarizabilities at 1064 nm wavelength. Among all the doped structures, the  $Au_3Cu_1$  cluster showed the largest static first hyperpolarizability of  $\beta_{(total)} = 4.73 \times 10^3$  au, while the  $Au_1Cu_6$  cluster showed frequency dependent first hyperpolarizability of  $\beta_{(-2\omega;\omega,\omega)} = 1.26 \times 10^6$  au. Besides this, large static and frequency-dependent second hyperpolarizability values of  $6.30 \times 10^5$  au and  $1.05 \times 10$  au were exhibited by  $Cu_7$  and  $Au_1Cu_6$ , respectively. This study offers an effective approach to design high-performance NLO materials utilizing mixed metal clusters which might have broad applications in the fields of optoelectronics and electronics.

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## Introduction

Nonlinear optics (NLO) is one of the sub-disciplines of modern optics, with an important academic status at par with laser physics. The study of nonlinear optics involves a variety of nonlinear effects that occur in the interaction between laser and matter.<sup>1,2</sup> The purpose of nonlinear optics research, in particular, is for promoting new developments and new impacts on the co-operative process of powerful laser beams and materials, including gaining an in-depth understanding of the causes and common processes as well as evaluating potential applications in current or future fields. Laser interaction with a NLO material causes a modification in the optical properties of the material

system, so that the next incoming photon sees something different.<sup>3</sup> Generally, only laser light is strong enough to produce NLO phenomena; therefore, the beginning of this field of study is often regarded as the result of the discovery of second-generation harmonics by Franken *et al.* in 1961,<sup>4</sup> one year after the first laser was constructed by Maiman.<sup>5</sup> Although the observation of most NLO phenomena requires laser radiation, some classes of nonlinear optical effects were known long before the invention of laser. The most important examples of these phenomena include Pockels and Kerr electro-optic effects.<sup>6</sup>

Inorganic NLO objects associated with nonlinear light distribution from the central region play an important role in optical science. Nowadays, NLO materials are used in many technological applications, including photonic devices, optoelectronics, second-harmonic generation (SHG), frequency doublers, fiber-optic telecommunication, and laser technology.<sup>7,8</sup> The material requirement to reflect the NLO response is not centrosymmetric. In this regard, other non-centrosymmetric crystals exhibiting an indirect optical response are  $K_3B_6O_{10}Cl$  ref. 9 and  $SrB_5O_7F_3$ .<sup>10</sup> Continuous efforts have been devoted to design and integrate new NLO materials due to rapid advances in the fields of nanoscience and

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nanotechnology.<sup>11</sup> Materials used in the nano-scale have revealed an excellent NLO response that greatly enhances their growth in the current era.<sup>12</sup> Gold nanoparticles (Au-NPs) play a vital role in the field of photonics.<sup>13,14</sup>

In addition, several design strategies for NLO materials have been proposed, including, the significant diradical character,<sup>15</sup> designing octupolar molecules,<sup>16</sup> the push-pull effect,<sup>17</sup> and excess electron system.<sup>18</sup> Among the above-mentioned approaches for designing NLO materials, the introduction of excess electrons is a well-known and widely used strategy.<sup>7</sup> Gold copper bimetallic clusters are a good source of excess electrons and can be doped in many systems to generate excess electron compounds.<sup>19</sup> Therefore, a number of theoretical studies and experiments have been presented evaluating the electronic stability and chemical application of transitional metal clusters.<sup>20</sup>

Metal clusters are obtained from the bonding of a few to hundreds of atoms and are called molecular entities.<sup>21</sup> Gold clusters have declared as emerging materials in the 21st century due to their outstanding catalytic performances and their various NLO applications. Furthermore, gold clusters have proven to be key materials in electronics, optics<sup>22</sup> and also in biomedical fields.<sup>23,24</sup> Kokabi and Salehiyoun, studied the NLO and electronic properties of small neutral and singly charged iron-doped bismuth clusters.<sup>25</sup> Recently, Knoppe *et al.* studied the NLO properties of thiolate-protected and phosphine-protected Au<sub>20</sub> clusters.<sup>26,27</sup>

From the review of the valuable literature, it is evident that nano-scale materials possess excellent optical and nonlinear optical responses. Alloys of two clusters have also proved to be a better system with an excellent NLO response. Motivated by previous reports, we designed new alloys based on Cu and Au doping. The doping of two clusters can be an efficient approach for designing better NLO materials. In this report, we studied the changes produced in the electronic structure and geometry of Cu-doped Au-clusters by considering the dipole moment, HOMO-LUMO energy gap, NLO, average static  $\beta_{(0;0,0,0)}$  and frequency-dependent  $\beta_{(-2w;w,w)}$  second-order polarizability, and third-order static  $\gamma_{(0;0,0,0)}$  and frequency-dependent  $\gamma_{(-2w;w,0,0)}$  polarizability. The results of all the analyses suggested that our designed clusters are effective candidates as superb nonlinear optical materials.

### Computational methodology

The Gaussian 09 program was used for all the computational calculations.<sup>28</sup> The structures of all the studied gold and copper clusters were visualized and designed using the Gauss View 5.0 Program.<sup>29</sup> Density functional theory (DFT) methods were used to compute the molecular geometries of the neutral gold clusters. This theoretical methodology was selected due to the presence of a great number of electrons in the metal clusters and required the introduction of electron correlation effects. The CAM-B3LYP functional was used with LANL2DZ (Los Alamos National Laboratory 2 Double-Zeta), which is a widely used ECP (effective core potentials) type basis set to model the metal atoms and can reduce the computational cost and energy

required.<sup>30</sup> The HOMO-LUMO energy gap and NLO properties were calculated by using the same method as used for the optimization.

In our work, we focus on the comparison of the total electronic dipole moment  $\mu_{\text{tot}}$ , static and the frequency-dependent linear isotropic  $\alpha_{\text{iso}}$  and anisotropic  $\alpha_{\text{aniso}}$  polarizabilities, total second-order polarizability  $\beta_{(\text{total})}$ , average static  $\beta_{(0;0,0,0)}$  and frequency-dependent  $\beta_{(-2w;w,w)}$  first hyperpolarizabilities, and average static  $\gamma_{(0;0,0,0)}$  and frequency-dependent  $\gamma_{(-2w;w,0,0)}$  second hyperpolarizabilities [66, 67]. Mathematically, the anisotropic and isotropic polarizabilities can be stated as follows:<sup>31,32</sup>

$$\alpha_{\text{iso}} = \frac{1}{3}(\alpha_{xx} + \alpha_{yy} + \alpha_{zz}) \quad (1)$$

$$\alpha_{\text{aniso}} = \sqrt{(\alpha_{xx} - \alpha_{yy})^2 + (\alpha_{xx} - \alpha_{zz})^2 + (\alpha_{yy} - \alpha_{zz})^2} \quad (2)$$

Estimation of the non-linearities, particularly the first hyperpolarizabilities, were calculated as follows.<sup>33,34</sup>

$$\beta_{\text{total}} = (\beta_x^2 + \beta_y^2 + \beta_z^2)^{1/2} \quad (3)$$

$$\beta_x = \frac{3}{5}(\beta_{xxx} + \beta_{xyy} + \beta_{xzz}) \quad (4)$$

$$\beta_y = \frac{3}{5}(\beta_{yyy} + \beta_{yzz} + \beta_{xyy}) \quad (5)$$

$$\beta_z = \frac{3}{5}(\beta_{xzz} + \beta_{yyz} + \beta_{zzz}) \quad (6)$$

whereas the average third-order polarizability or second hyperpolarizability can be calculated as:

$$\gamma_o = \frac{1}{15} \sum_{ij=x,y,z} (\gamma_{ijij} + \gamma_{ijji} + \gamma_{ijji}) \quad (7)$$

The following equation was used to calculate the H-L energy gap.<sup>35,36</sup>

$$E_{\text{H-L}} = E_{\text{L}} - E_{\text{H}} \quad (8)$$

where  $E_{\text{H}}$  is the energy of the HOMO,  $E_{\text{L}}$  is the energy of the LUMO, and  $E_{\text{H-L}}$  represents the HOMO-LUMO energy gap.

## Results and discussion

### Geometry of the pure and Cu-doped gold clusters

The structural geometry of the clusters plays an important role in modulating the NLO and optoelectronic properties of advanced functional materials. So, the effect of variations in the geometrical structures on their NLO properties by doping Cu atom in pure Au-clusters was investigated. In the present study, small Au-clusters Au<sub>m</sub> ( $2 \leq m \leq 7$ ), namely Au<sub>2</sub>, Au<sub>3</sub>, Au<sub>4</sub>, Au<sub>5</sub>, Au<sub>6</sub>, and Au<sub>7</sub>, clusters were optimized through DFT calculations at the CAM-B3LYP/LANL2DZ level of theory, which is an efficient and frequently used method for doped clusters. In the literature, different reports effectively used the CAM-B3LYP/LanL2DZ level of theory for studying the electronic behavior



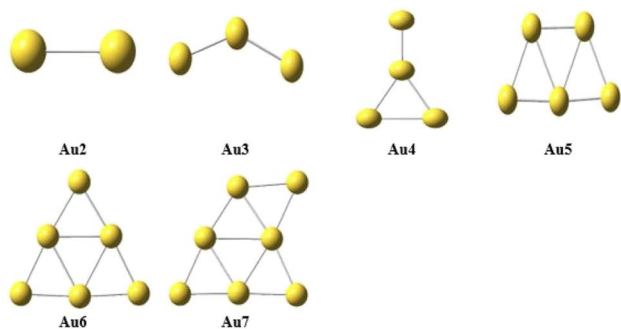


Fig. 1 Optimized  $Au_m$  ( $2 \leq m \leq 7$ ) clusters.

of clusters.<sup>37</sup> The average bond lengths and HOMO–LUMO energy gaps were calculated and compared with the reported Au-clusters, as it is very significant to relate the properties of the studied materials with others to ensure their uniqueness. As shown in Fig. 2, the reported bond lengths of the pure Au-clusters and our calculated values for pure Au-clusters were comparable.<sup>37</sup> After careful analysis of all the designed clusters, the following lowest energy geometry structures of pure gold  $Au_m$  ( $2 \leq m \leq 7$ ) clusters were selected (based on stability), as shown in Fig. 1.

Furthermore, by the replacement of both Au atoms of the  $Au_2$  cluster with a Cu atom the  $Au_1Cu_1$ ,  $Cu_2$  lowest energy geometry structures were obtained. Similarly, for  $Au_3$  ( $Au_2Cu_1$ ,  $Au_1Cu_2$ ,  $Cu_3$ ),  $Au_4$  ( $Au_3Cu_1$ ,  $Au_2Cu_2$ ,  $Au_1Cu_3$ ,  $Cu_4$ ),  $Au_5$  ( $Au_4Cu_1$ ,  $Au_3Cu_2$ ,  $Au_2Cu_3$ ,  $Au_1Cu_4$ ,  $Cu_5$ ),  $Au_6$  ( $Au_5Cu_1$ ,  $Au_4Cu_2$ ,  $Au_3Cu_3$ ,  $Au_2Cu_4$ ,  $Au_1Cu_5$ ,  $Cu_6$ ), and  $Au_7$  ( $Au_6Cu_1$ ,  $Au_5Cu_2$ ,  $Au_4Cu_3$ ,  $Au_3Cu_4$ ,  $Au_2Cu_5$ ,  $Au_1Cu_6$ ,  $Cu_7$ ) structures were obtained, as shown in Fig. 2.

Due to having applications in optoelectronics, gold clusters are considered as promising NLO materials. To enhance the conducting properties of pure gold clusters, these are doped with Cu atoms. The average bond length of the pure gold clusters  $Au_2$ ,  $Au_3$ ,  $Au_4$ ,  $Au_5$ ,  $Au_6$ , and  $Au_7$  were 2.57, 2.64, 2.71, 2.77, 2.77, and 2.79 Å, respectively. While these lengths decreased by doping with Cu atom as  $Au_1Cu_1$  2.57 Å 2.41,  $Cu_2$  2.26 Å. The average bond length for the pure Cu clusters  $Cu_2$ ,  $Cu_3$ ,  $Cu_4$ ,  $Cu_5$ ,  $Cu_6$ , and  $Cu_7$  were 2.26, 2.33, 2.40, 2.44, 2.44, and 2.45 Å respectively. The average bond lengths were considerably decreased by doping Cu in the pure gold clusters, as shown in Table 1. The  $E_g$  values for the pure gold cluster structures  $Au_2$ ,  $Au_3$ ,  $Au_4$ ,  $Au_5$ ,  $Au_6$ , and  $Au_7$  were 3.25, 3.14, 1.93, 2.49, 3.48, and 3.25 eV, respectively. A mixed trend for the energy bandgap was obtained for  $Cu_2$ ,  $Cu_3$ ,  $Cu_4$ ,  $Cu_5$ ,  $Cu_6$ , and  $Cu_7$ , as shown in Table 1. These results show that the HOMO–LUMO energy gap was altered upon doping with Cu atom. The band gap decreased from the pure to the doped system in the range of 3.67–0.91 eV. The lowest energy bandgap was observed for  $Au_2Cu_2$ ,  $Au_3Cu_1$ ,  $Cu_3$ ,  $Cu_4$ , and  $Au_4$ , as shown in Table 1.

This decreases in bond length and energy gap were due to deformation in the electronic structures of the clusters and increases in the dipole moment and the conducting properties of the clusters. All the doped clusters had a small ABL (average bond length) and low energy gap, which showed that all the doped clusters were stable.

## Dipole moment

The computed values of the dipole moment ( $\mu$ ) and change in the transition dipole moment ( $\delta\mu$ ) of the pure and Cu-doped gold clusters are given in Table 1. The doping of gold clusters significantly affected their electro-optical properties, such as the dipole moment and transition energy. The doping of metal unbalances the electron density of the pure clusters, which results in charge separation. The charge separation causes an increase in the dipole moment. A greater value of dipole moment in the doped system allows a higher NLO response. Therefore, the NLO polarizabilities were also calculated, which strongly depend on the dipole moment, oscillator strength, and orbital transition energy.

The dipole moment was almost zero due to the centrosymmetric nature of the pure clusters. The doping of a transition metal (Cu) could significantly change the dipole moments as compared to pure clusters. The total and transition dipole moment for the pure gold and pure copper clusters  $Au_2$ ,  $Au_3$ ,  $Au_4$ ,  $Au_5$ ,  $Au_6$ , and  $Au_7$  were almost 0.0 D. The doping of Cu into the pure Au-clusters imbalanced the electronic density, which resulted in charge separation. This charge separation caused an increase in the dipole moment. The calculated total dipole moment  $\mu_D$  and transition dipole moment  $\delta\mu$  for the optimized geometries of Au and Au–Cu doped clusters are presented in Table 1.

Large dipole moments  $\mu_D$  for the Cu-doped clusters  $Au_2Cu_1$ ,  $Au_4$ , and  $Au_3Cu_1$  were noted as 3.27, 3.63, and 5.03 D, respectively. The highest transition dipole moments  $\delta\mu$  were 0.5691, 0.7955, 0.8135, and 1.2070 D for  $Au_4Cu_2$ ,  $Au_3Cu_3$ ,  $Au_1Cu_5$ , and  $Au_2Cu_4$ , respectively, due to the highly imbalanced structures of the molecules. The dipole moments of the pure Au and pure Cu clusters were less than the mixed Au–Cu clusters. Interestingly, the dipole moment of the pure clusters increased upon doping with Cu atoms. The  $Au_2$  clusters showed a lower value of dipole moment for pure Au and pure Cu cluster compared to the others. The doped clusters, like  $Au_1$ – $Cu_1$  clusters, exhibited a higher value of dipole moment, which was due to the change in charge separation. The dipole moment for a symmetric structure is always less than for nonsymmetric structures, like in the  $Au_4$  cluster, where the  $\mu_D$  for the  $Au_2Cu_2$  symmetric structure was less than for the  $Au_3Cu_1$  nonsymmetric structure. The same trend of dipole moment was shown in  $Au_6$  ( $Au_3Cu_3$ ) and  $Au_7$  ( $Au_4Cu_4$ ) clusters. The dipole moment of the cluster structures containing a single Au atom was higher than the others with more than one Au atom clusters. The dipole moment of clusters containing a single Cu atom was also higher than the others with more than one Cu atom clusters, like in  $Au_5$  ( $Au_4Cu_1$ ),  $Au_6$  ( $Au_5Cu_1$ ) and  $Au_7$  ( $Au_6Cu_1$ ) clusters. Thus, one can expect better electronic and optoelectronic properties from these clusters.

## Linear isotropic and anisotropic polarizabilities

In isotropic media, the polarizability  $\alpha_{iso}$  is defined as the ratio of the induced dipole moment of an atom to the electric field generating dipole moment, while in general, the polarizability cannot be expressed as a scalar quantity for anisotropic  $\alpha_{aniso}$  or



non-spherical media. In scalar terms, the anisotropic polarizability  $\alpha_{\text{aniso}}$  can be referred to as the applied electric fields can only cause field parallel components to become polarized and

as the  $x, y, z$  directions react in the same way to an applied electric field. The characterization of anisotropic media is done by the two tensor polarizability rank. Isotropic polarizability in

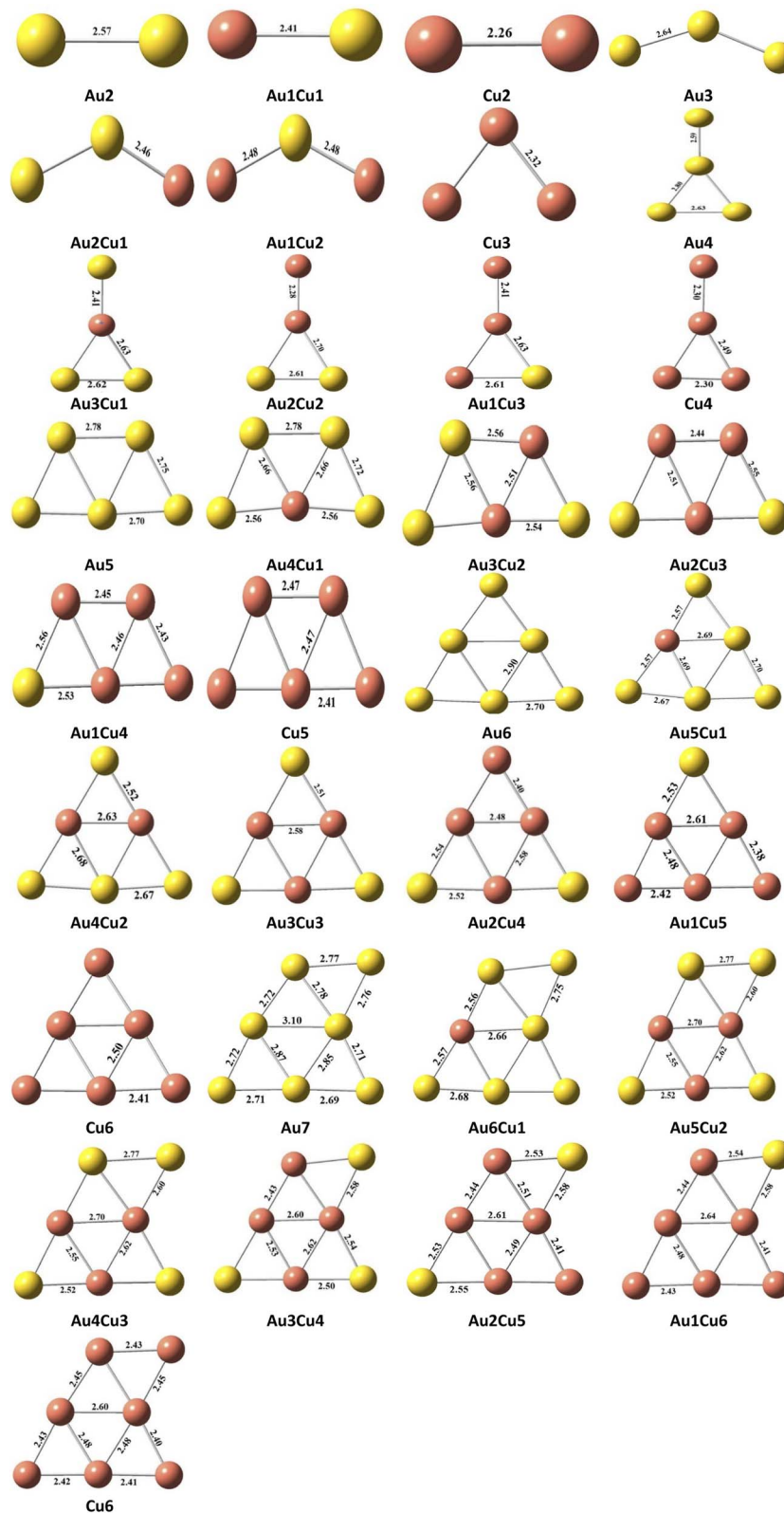


Fig. 2 Optimized structures and bond lengths (Å) of the pure and Cu-doped clusters.



**Table 1** Calculated values of the dipole moment  $\mu$  (D), transition dipole moment  $\delta\mu$ , first excited state transition energy (eV), oscillator strength ( $f_o$ ), HOMO–LUMO, energy gap  $E_g$  in eV, and average bond length ( $\text{\AA}$ )

Molecule	First transition dipole ( $\delta\mu$ )	$f_o$	Transition (%C·I)	$\mu$ (D)	HOMO (eV)	LUMO (eV)	$E_g$ (eV)	$\Delta E$ (eV)	ABL ( $\text{\AA}$ )
Au <sub>2</sub>	0.0000	0.0000	H–1 to L+1 (71)	0.00	–7.19	–3.93	3.25	2.7346	2.57
Au <sub>1</sub> Cu <sub>1</sub>	0.2152	0.0030	H–1 to L+1 (70)	2.63	–6.40	–3.02	3.38	2.6029	2.41
Cu <sub>2</sub>	0.0000	0.0000	H–1 to L+1 (70)	0.00	–5.59	–2.34	3.25	2.5317	2.26
Au <sub>3</sub>	0.0595	0.0001	H+3 to L+1 (99)	0.37	–6.59	–3.45	3.14	1.2269	2.64
Au <sub>2</sub> Cu <sub>1</sub>	0.0740	0.0003	H+2 to L–1 (73)	3.27	–5.72	–2.96	2.75	1.4927	2.56
Au <sub>1</sub> Cu <sub>2</sub>	0.4680	0.0085	H to L+2 (47)	1.40	–4.97	–2.53	2.44	1.5765	2.48
Cu <sub>3</sub>	0.4958	0.0034	H to L+1 (100)	0.42	–4.05	–2.66	1.39	0.5600	2.33
Au <sub>4</sub>	0.3176	0.0033	H to L+1 (70)	3.63	–6.34	–4.41	1.93	1.3526	2.71
Au <sub>3</sub> Cu <sub>1</sub>	0.1746	0.0005	H to L+1 (71)	5.03	–5.86	–4.58	1.28	0.6436	2.58
Au <sub>2</sub> Cu <sub>2</sub>	0.1961	0.0002	H to L+1 (72)	1.63	–5.15	–4.22	0.93	0.2505	2.56
Au <sub>1</sub> Cu <sub>3</sub>	0.2813	0.0035	H to L+1 (70)	2.51	–5.45	–2.92	2.53	1.8158	2.58
Cu <sub>4</sub>	0.2906	0.0025	H to L+1 (71)	3.02	–4.76	–2.91	1.85	1.2200	2.40
Au <sub>5</sub>	0.2302	0.0001	H to L+1 (97)	0.02	–5.91	–3.42	2.49	1.2608	2.77
Au <sub>4</sub> Cu <sub>1</sub>	0.1610	0.0008	H to L–2 (91)	0.22	–5.80	–3.32	2.48	1.2801	2.67
Au <sub>3</sub> Cu <sub>2</sub>	0.4196	0.0066	H to L–2 (96)	1.19	–5.66	–2.70	2.95	1.4614	2.59
Au <sub>2</sub> Cu <sub>3</sub>	0.3651	0.0054	H to L–2 (91)	1.71	–5.29	–2.57	2.72	1.6456	2.52
Au <sub>1</sub> Cu <sub>4</sub>	0.2076	0.0025	H to L–1 (24)	2.47	–4.88	–2.42	2.46	1.5801	2.48
Cu <sub>5</sub>	0.2558	0.0022	H to L+1 (97)	0.01	–4.55	–2.17	2.38	1.3774	2.44
Au <sub>6</sub>	0.1572	0.0017	H to L–1 (52)	0.00	–6.92	–3.43	3.48	2.8508	2.77
Au <sub>5</sub> Cu <sub>1</sub>	0.3057	0.0063	H to L–2 (59)	0.46	–6.79	–3.39	3.39	2.7549	2.69
Au <sub>4</sub> Cu <sub>2</sub>	0.5691	0.0218	H to L–1 (69)	0.25	–6.77	–3.36	3.40	2.7487	2.60
Au <sub>3</sub> Cu <sub>3</sub>	0.7955	0.0619	H to L–1 (63)	0.01	–6.74	–3.07	3.67	3.0298	2.54
Au <sub>2</sub> Cu <sub>4</sub>	1.2070	0.1024	H to L–1 (66)	2.47	–6.18	–2.79	3.39	2.8677	2.50
Au <sub>1</sub> Cu <sub>5</sub>	0.8135	0.0428	H to L–1 (65)	2.51	–5.74	–2.52	3.22	2.6353	2.47
Cu <sub>6</sub>	0.0882	0.0489	H to L–1 (61)	0.00	–5.50	–2.25	3.25	2.6802	2.44
Au <sub>7</sub>	0.0475	0.0001	H to L–1 (87)	0.39	–5.82	–3.58	2.24	1.3915	2.79
Au <sub>6</sub> Cu <sub>1</sub>	0.2629	0.0023	H to L–2 (95)	1.17	–5.61	–3.60	2.01	1.2903	2.72
Au <sub>5</sub> Cu <sub>2</sub>	0.2814	0.003	H to L–2 (97)	0.88	–5.61	–3.59	2.02	1.2706	2.65
Au <sub>4</sub> Cu <sub>3</sub>	0.1160	0.0009	H to L (91)	0.81	–5.70	–3.22	2.49	1.4583	2.59
Au <sub>3</sub> Cu <sub>4</sub>	0.1772	0.0016	H to L (91)	1.57	–5.51	–2.98	2.53	1.4386	2.54
Au <sub>2</sub> Cu <sub>5</sub>	0.2171	0.0015	H to L (98)	0.90	–5.34	–2.61	2.68	1.1374	2.51
Au <sub>1</sub> Cu <sub>6</sub>	0.2110	0.0021	H to L (97)	2.37	–4.93	–2.47	2.46	1.1618	2.48
Cu <sub>7</sub>	0.2449	0.0021	H to L–2 (92)	0.17	–4.40	–2.34	2.06	1.2803	2.45

scalar terms can be defined as the ratio of the induced atomic dipole moment to the dipole moment rising through an electric field.<sup>38</sup> Here, we specifically investigated the static and frequency-dependent  $\alpha_{\text{iso}}$  and  $\alpha_{\text{aniso}}$  of pure and Cu-doped Au-clusters at the CAM-B3LYP/LANL2DZ level of DFT and their values are shown in Table 2. The doping of Cu in pure Au-clusters meaningfully affected the electro-optical characteristics, such as transition energy and dipole moment. The maximum dipole moment was seen in doped systems, which may be due to their high NLO response.

The linear static isotropic polarizabilities for the pure Au-clusters Au<sub>2</sub>, Au<sub>3</sub>, Au<sub>4</sub>, Au<sub>5</sub>, Au<sub>6</sub>, and Au<sub>7</sub> were 78.86, 142.89, 167.34, 204.80, 234.61, and 298.49 au, respectively. The isotropic polarizabilities of pure Cu clusters were in the range of 76.74–281.49 au. The frequency-dependent isotropic polarizabilities for pure gold clusters were in the range of 60.16–322.77 au, and 45.97 = –248.24 au for pure Cu clusters. High values of static and frequency-dependent  $\alpha_{\text{iso}}$  were shown by Cu<sub>7</sub> and Au<sub>7</sub> clusters due to their large size and deformation structure.

The static anisotropic polarizabilities for pure Au-clusters were 69.91, 177.24, 156.36, 176.23, 170.68, and 270.15 au for Au<sub>2</sub>, Au<sub>3</sub>, Au<sub>4</sub>, Au<sub>5</sub>, Au<sub>6</sub>, and Au<sub>7</sub>, respectively. Frequency-

dependent anisotropic polarizabilities for the pure Au (2–7) clusters were in the range of 69.91–311.97 au. These show that the frequency-dependent anisotropic polarizability is always higher than the static. Therefore, we can say that the anisotropic polarizabilities for the pure Au and Cu clusters were very close to each other. High values of  $\alpha_{\text{aniso}}$  static and frequency-dependent polarizabilities were also shown by the Cu-doped Au<sub>7</sub>(Au<sub>6</sub>Cu<sub>1</sub>, Au<sub>5</sub>Cu<sub>2</sub>, Au<sub>4</sub>Cu<sub>3</sub>, Au<sub>3</sub>Cu<sub>4</sub>, Au<sub>2</sub>Cu<sub>5</sub>, Au<sub>1</sub>Cu<sub>6</sub>, Cu<sub>7</sub>) clusters due to deformation in their structure by the doping of copper and also due to their high dipole moment. A comparison of the linear and frequency-dependent isotropic and anisotropic polarizabilities is shown in Fig. 3.

Graphical comparison of the linear and frequency-dependent isotropic and anisotropic polarizabilities showed that the anisotropic polarizability was always less than isotropic polarizability. These results confirmed that the frequency-dependent isotropic and anisotropic polarizabilities were higher than the static one. The isotropic and anisotropic polarizability values were high for single Au and Cu containing cluster structures, like Au<sub>1</sub>Cu<sub>2</sub> and Au<sub>1</sub>Cu<sub>3</sub>, Au<sub>1</sub>Cu<sub>4</sub>, Au<sub>1</sub>Cu<sub>5</sub>, Au<sub>1</sub>Cu<sub>6</sub>, Au<sub>6</sub>Cu<sub>1</sub>, Au<sub>5</sub>Cu<sub>1</sub>, Au<sub>4</sub>Cu<sub>1</sub>, Au<sub>3</sub>Cu<sub>1</sub>, Au<sub>2</sub>Cu. The graphical comparison showed that by the doping of Cu, the values of the



**Table 2** Computed values of the static and frequency-dependent  $\alpha_{\text{iso}}$ ,  $\alpha_{\text{aniso}}$ , average second-order polarizability  $\beta_o$ ,  $\beta_{\text{total}}$ , and average third-order polarizability  $\gamma_o$  for the pure and Cu-doped gold clusters

Molecule	$\alpha_{(0,0)}$ (au)		$\alpha_{(-w,w)}$ (au)		2nd polarizability $\beta_o$ (au)			$\gamma_o$ (au)	
	$(\alpha_{\text{iso}})$	$(\alpha_{\text{aniso}})$	$(\alpha_{\text{iso}})$	$(\alpha_{\text{aniso}})$	$\beta_{(0;0,0)}$	$\beta_{(-2w;w,w)}$	$\beta_{\text{total}}$	$\gamma_{o(0;0,0,0)}$	$\gamma_{o(-2w;w,w,0)}$
Au <sub>2</sub>	78.86	69.91	60.16	84.03	$0.00 \times 10^1$	$0.00 \times 10^0$	$1.15 \times 10^1$	$6.01 \times 10^3$	$8.03 \times 10^3$
Au <sub>1</sub> Cu <sub>1</sub>	70.54	47.37	75.37	55.24	$1.01 \times 10^3$	$1.27 \times 10^3$	$4.70 \times 10^2$	$5.67 \times 10^5$	$6.74 \times 10^5$
Cu <sub>2</sub>	76.74	45.97	83.24	55.01	$0.00 \times 10^0$	$0.00 \times 10^0$	$6.74 \times 10^1$	$4.56 \times 10^4$	$1.42 \times 10^5$
Au <sub>3</sub>	142.89	177.24	186.02	285.96	$2.41 \times 10^2$	$1.26 \times 10^3$	$1.26 \times 10^3$	$1.42 \times 10^3$	$1.45 \times 10^5$
Au <sub>2</sub> Cu <sub>1</sub>	135.37	161.31	154.15	205.61	$2.21 \times 10^3$	$2.28 \times 10^4$	$3.64 \times 10^3$	$1.41 \times 10^5$	$1.69 \times 10^6$
Au <sub>1</sub> Cu <sub>2</sub>	146.45	189.12	175.11	262.17	$1.08 \times 10^3$	$3.26 \times 10^3$	$1.79 \times 10^3$	$1.08 \times 10^5$	$5.03 \times 10^5$
Cu <sub>3</sub>	131.57	106.17	133.82	943.76	$1.43 \times 10^3$	$6.21 \times 10^2$	$2.39 \times 10^3$	$2.47 \times 10^5$	$9.80 \times 10^5$
Au <sub>4</sub>	167.34	156.36	182.22	174.18	$1.28 \times 10^3$	$2.84 \times 10^3$	$2.84 \times 10^3$	$6.52 \times 10^4$	$1.16 \times 10^5$
Au <sub>3</sub> Cu <sub>1</sub>	152.36	121.33	162.80	143.06	$2.73 \times 10^3$	$4.46 \times 10^3$	$4.73 \times 10^3$	$1.44 \times 10^5$	$2.99 \times 10^5$
Au <sub>2</sub> Cu <sub>2</sub>	168.42	142.17	181.57	172.96	$1.78 \times 10^3$	$1.91 \times 10^3$	$3.36 \times 10^3$	$4.04 \times 10^5$	$1.56 \times 10^6$
Au <sub>1</sub> Cu <sub>3</sub>	138.90	119.24	148.71	134.41	$1.19 \times 10^3$	$2.62 \times 10^3$	$1.98 \times 10^3$	$1.79 \times 10^5$	$3.83 \times 10^5$
Cu <sub>4</sub>	145	147.35	132.26	139.74	$3.78 \times 10^2$	$5.21 \times 10^2$	$3.66 \times 10^2$	$3.17 \times 10^5$	$9.83 \times 10^5$
Au <sub>5</sub>	204.80	176.23	226.60	219.58	$3.51 \times 10^2$	$3.52 \times 10^2$	$2.63 \times 10^2$	$8.67 \times 10^4$	$4.24 \times 10^4$
Au <sub>4</sub> Cu <sub>1</sub>	194.87	160.67	217.13	205.13	$9.10 \times 10^1$	$1.28 \times 10^4$	$7.07 \times 10^1$	$2.02 \times 10^5$	$1.52 \times 10^7$
Au <sub>3</sub> Cu <sub>2</sub>	178.72	134.84	194.70	162.94	$3.05 \times 10^2$	$6.06 \times 10^3$	$4.42 \times 10^2$	$1.49 \times 10^4$	$1.09 \times 10^6$
Au <sub>2</sub> Cu <sub>3</sub>	177.16	136.25	192.44	159.26	$4.17 \times 10^2$	$1.82 \times 10^4$	$7.15 \times 10^2$	$2.15 \times 10^5$	$4.26 \times 10^7$
Au <sub>1</sub> Cu <sub>4</sub>	185.39	144.20	203.75	170.81	$1.11 \times 10^3$	$2.10 \times 10^3$	$1.86 \times 10^3$	$3.27 \times 10^5$	$2.10 \times 10^5$
Cu <sub>5</sub>	195.65	154.01	218.85	187.47	$2.95 \times 10^2$	$2.23 \times 10^2$	$4.99 \times 10^2$	$3.57 \times 10^5$	$5.15 \times 10^5$
Au <sub>6</sub>	234.61	170.68	247.08	186.02	$1.74 \times 10^0$	$4.35 \times 10^1$	$1.94 \times 10^1$	$9.84 \times 10^4$	$2.10 \times 10^5$
Au <sub>5</sub> Cu <sub>1</sub>	222.74	161.23	234.90	176.46	$1.21 \times 10^2$	$3.42 \times 10^2$	$9.31 \times 10^1$	$1.73 \times 10^4$	$2.10 \times 10^4$
Au <sub>4</sub> Cu <sub>2</sub>	209.73	146.72	221.02	160.91	$1.92 \times 10^2$	$7.56 \times 10^2$	$3.97 \times 10^2$	$1.19 \times 10^5$	$2.65 \times 10^5$
Au <sub>3</sub> Cu <sub>3</sub>	195.39	124.79	205.43	136.18	$1.03 \times 10^2$	$1.82 \times 10^2$	$1.06 \times 10^2$	$1.89 \times 10^5$	$3.44 \times 10^5$
Au <sub>2</sub> Cu <sub>4</sub>	201.60	132.80	213.71	146.69	$1.08 \times 10^3$	$2.31 \times 10^3$	$1.79 \times 10^3$	$9.13 \times 10^4$	$1.84 \times 10^5$
Au <sub>1</sub> Cu <sub>5</sub>	210.24	142.33	224.94	159.32	$1.11 \times 10^3$	$2.35 \times 10^3$	$1.86 \times 10^3$	$3.80 \times 10^5$	$8.57 \times 10^5$
Cu <sub>6</sub>	221.25	153.97	239.15	174.58	$8.36 \times 10^2$	$5.11 \times 10^1$	$8.69 \times 10^1$	$3.46 \times 10^5$	$8.81 \times 10^5$
Au <sub>7</sub>	298.49	270.15	322.77	311.97	$3.95 \times 10^2$	$6.20 \times 10^3$	$6.60 \times 10^2$	$1.83 \times 10^5$	$1.15 \times 10^6$
Au <sub>6</sub> Cu <sub>1</sub>	286.96	254.14	310.79	293.15	$4.52 \times 10^2$	$6.06 \times 10^2$	$6.67 \times 10^2$	$5.06 \times 10^4$	$3.79 \times 10^5$
Au <sub>5</sub> Cu <sub>2</sub>	269.18	220.05	295.90	254.51	$3.89 \times 10^2$	$1.13 \times 10^4$	$6.55 \times 10^2$	$3.53 \times 10^5$	$2.42 \times 10^7$
Au <sub>4</sub> Cu <sub>3</sub>	251.88	204.48	271.71	236.66	$3.11 \times 10^2$	$4.09 \times 10^3$	$3.56 \times 10^2$	$2.03 \times 10^5$	$2.96 \times 10^6$
Au <sub>3</sub> Cu <sub>4</sub>	251.05	205.78	272.79	238.00	$2.94 \times 10^2$	$2.80 \times 10^3$	$4.98 \times 10^2$	$2.00 \times 10^5$	$2.88 \times 10^5$
Au <sub>2</sub> Cu <sub>5</sub>	255.72	209.45	258.74	228.01	$6.61 \times 10^2$	$8.23 \times 10^3$	$1.14 \times 10^3$	$3.27 \times 10^5$	$5.77 \times 10^6$
Au <sub>1</sub> Cu <sub>6</sub>	267.89	226.57	287.96	438.78	$1.61 \times 10^3$	$1.26 \times 10^6$	$2.68 \times 10^3$	$4.22 \times 10^5$	$1.05 \times 10^8$
Cu <sub>7</sub>	281.49	248.24	316.76	303.68	$5.35 \times 10^2$	$1.31 \times 10^4$	$9.00 \times 10^2$	$6.30 \times 10^5$	$5.15 \times 10^6$

$\alpha_{\text{iso}}$  and  $\alpha_{\text{aniso}}$  first decreased, like in Au<sub>5</sub> clusters, where the  $\alpha_{\text{iso}}$  and  $\alpha_{\text{aniso}}$  first decreased from Au<sub>5</sub> to Au<sub>2</sub>Cu<sub>3</sub> cluster and then increases by further doping up to the pure Cu<sub>5</sub> cluster. The Au<sub>6</sub> and Au<sub>7</sub> clusters showed a similar trend for the isotropic and anisotropic polarizabilities. It could also be noted that the clusters containing equal numbers of gold and copper atoms showed low polarizability; for example, the  $\alpha_{\text{iso}}$  and  $\alpha_{\text{aniso}}$  for the Au<sub>1</sub>Cu<sub>1</sub>, Au<sub>2</sub>Cu<sub>2</sub>, and Au<sub>3</sub>Cu<sub>3</sub> structures were less than for Au<sub>2</sub>Cu<sub>3</sub>, Au<sub>3</sub>Cu<sub>2</sub>, and Au<sub>4</sub>Cu<sub>5</sub>, etc. When the anisotropic polarizability  $\alpha_{\text{aniso}}$  values of the pure and Cu-doped clusters were compared, the values of the Cu-doped clusters were less than for the pure gold and pure copper clusters. The polarizability of the studied clusters gradually increased with increased size (atomic size) and the number of metals.

### Second-order NLO polarizability

Density functional theory (DFT) at the CAM-B3LYP/LANL2DZ level was used to theoretically calculate the total second-order polarizability  $\beta_{\text{(total)}}$ , average static second-order polarizability  $\beta_{(0;0,0)}$ , and average frequency-dependent second-order polarizability  $\beta_{(-2w;w,w)}$  of the pure and Cu-doped gold nano clusters.

The computed values of  $\beta_{(0;0,0)}$ ,  $\beta_{(-2w;w,w)}$ , and  $\beta_{\text{(total)}}$  of the pure and doped Au–Cu clusters are shown in Table 2.

A two-level model was applied to understand the factors affecting the hyperpolarizability.<sup>39–41</sup> This model explains the dependency of the first hyperpolarizability on the oscillator strength ( $f_o$ ) and transition energy ( $\Delta E$ ) of the crucial excited state, and the difference in the dipole moment between the ground and crucial excited states ( $\Delta\mu$ ). The two-level model can be represented as follows:

$$\beta_o \propto \frac{\Delta\mu f_o}{\Delta E^3} \quad (9)$$

This equation shows that the first hyperpolarizability ( $\beta_o$ ) is directly related to the oscillator strength ( $f_o$ ) and transition dipole moment ( $\Delta\mu$ ), but inversely proportional to the third power of the transition energy ( $\Delta E$ ).

The first hyperpolarizabilities  $\beta_{\text{(total)}}$  for the pure Au-clusters Au<sub>2</sub>, Au<sub>3</sub>, Au<sub>4</sub>, Au<sub>5</sub>, Au<sub>6</sub>, and Au<sub>7</sub> were  $1.15 \times 10^1$ ,  $1.26 \times 10^3$ ,  $2.84 \times 10^3$ ,  $2.63 \times 10^2$ ,  $1.94 \times 10^1$ , and  $6.60 \times 10^2$  au, respectively. The high first hyperpolarizabilities  $\beta_{\text{(total)}}$  for the doped



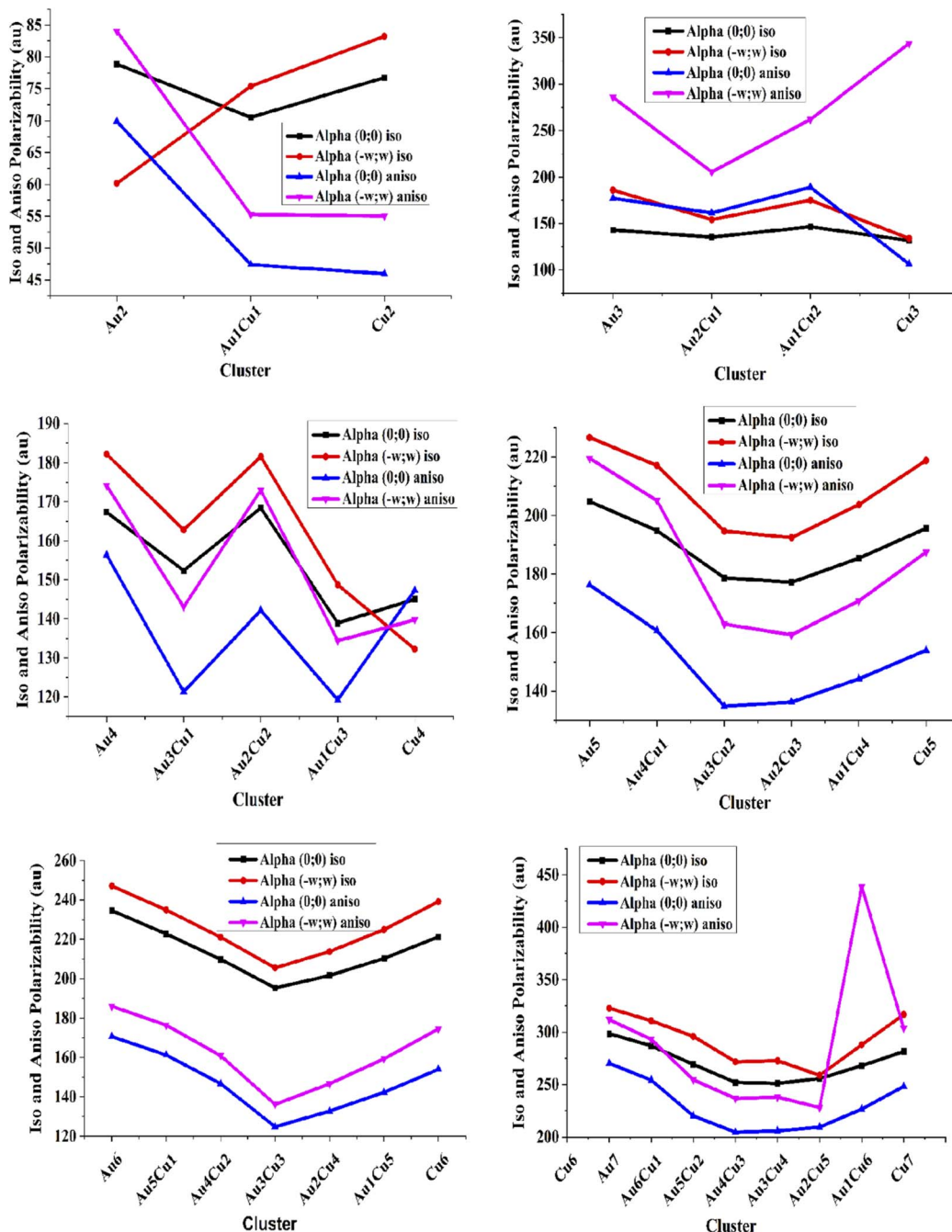


Fig. 3 Linear isotropic and anisotropic polarizabilities of the pure and Cu-doped gold clusters.

clusters  $\text{Au}_1\text{Cu}_6$ ,  $\text{Au}_4$ ,  $\text{Cu}_4$ ,  $\text{Au}_2\text{Cu}_2$ ,  $\text{Au}_2\text{Cu}_1$ , and  $\text{Au}_3\text{Cu}_1$  were  $2.68 \times 10^3$ ,  $2.84 \times 10^3$ ,  $3.21 \times 10^3$ ,  $3.36 \times 10^3$ ,  $3.64 \times 10^3$ , and  $4.73 \times 10^3$  au, respectively. The static second-order polarizabilities  $\beta_{(0;0,0)}$  for the pure Au (2–7) clusters were in the range of  $0$ – $2.73 \times 10^3$  au. The highest static first hyperpolarizability for the doped clusters  $\text{Au}_4$ ,  $\text{Cu}_3$ ,  $\text{Au}_1\text{Cu}_6$ ,  $\text{Au}_2\text{Cu}_2$ ,  $\text{Au}_2\text{Cu}_1$ , and  $\text{Au}_3\text{Cu}_1$  were  $1.28 \times 10^3$ ,  $1.43 \times 10^3$ ,  $1.61 \times 10^3$ ,  $1.78 \times 10^3$ ,  $2.21 \times 10^3$  and  $2.73 \times 10^3$  au, respectively.

The frequency-dependent  $\beta_{(-2w;w,w)}$  polarizabilities for the pure  $\text{Au}_m$  ( $m = 2$ – $7$ ) clusters were in the range of  $0$ – $1.26 \times 10^6$  au. These showed that the frequency-dependent first

hyperpolarizability  $\beta_{(-2w;w,w)}$  was always higher than the static one. The first hyperpolarizability values of the pure Au and Cu clusters were very close to each other. While, the average frequency-dependent  $\beta_{(-2w;w,w)}$  values were very high for the mixed Au–Cu clusters  $\text{Au}_5\text{Cu}_2$ ,  $\text{Au}_4\text{Cu}_1$ ,  $\text{Cu}_7$ ,  $\text{Au}_2\text{Cu}_3$ ,  $\text{Au}_2\text{Cu}_1$ , and  $\text{Au}_1\text{Cu}_6$  as  $1.13 \times 10^4$ ,  $1.28 \times 10^4$ ,  $1.31 \times 10^4$ ,  $1.82 \times 10^4$ ,  $2.28 \times 10^4$ , and  $1.26 \times 10^6$ , which were in accordance with the two-level model. These resulting values of the second hyperpolarizability for the doped clusters were higher than those of previously reported NLO materials, ranging from  $2.55 \times 10^4$ – $3.63 \times 10^4$  au.<sup>42</sup> For further investigation of the consistency of



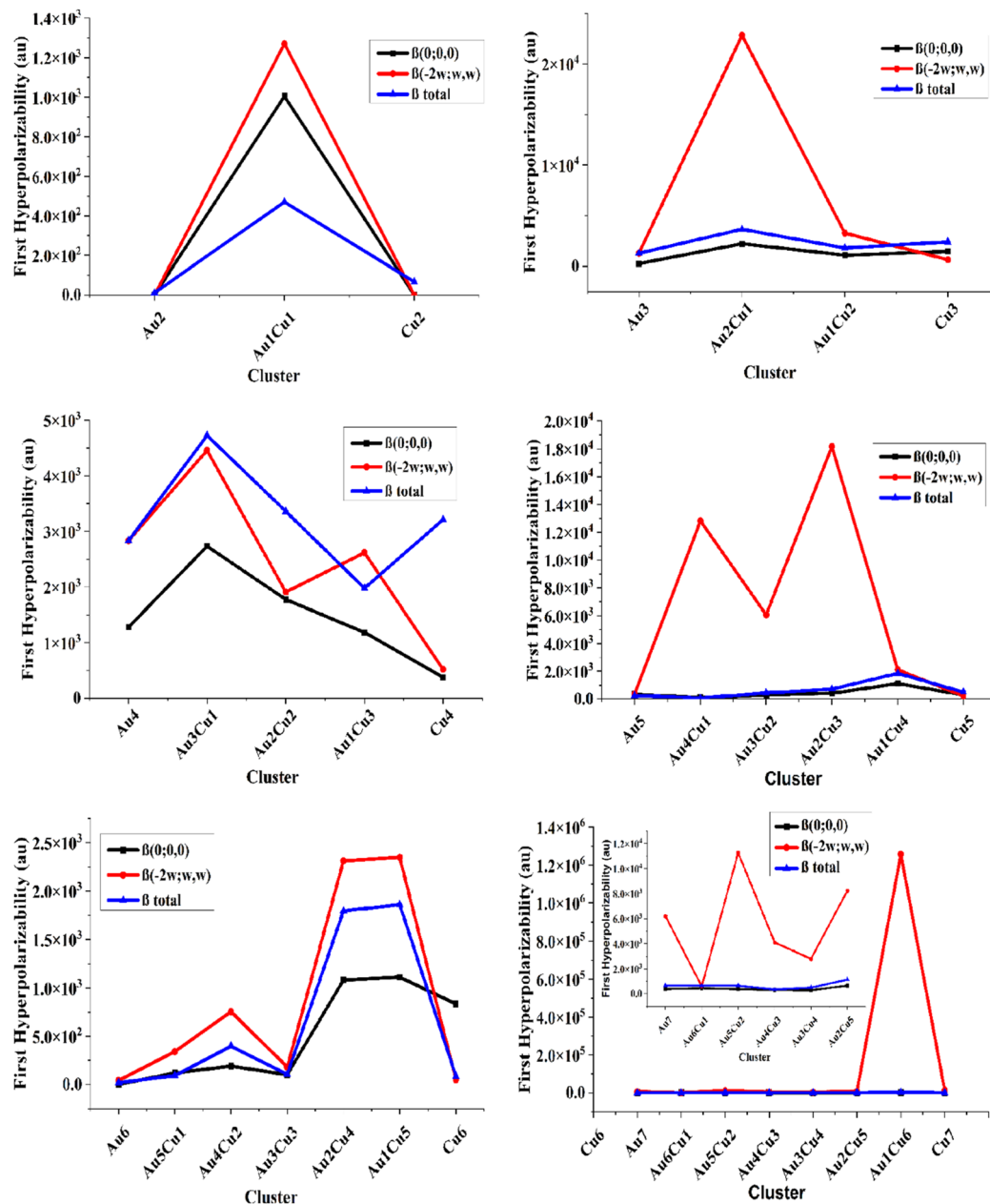


Fig. 4 First hyperpolarizability comparison for the pure and Cu-doped gold clusters.

the different pure and Cu-doped Au-clusters, a graphical comparison was made for the first hyperpolarizability or second-order polarizability, as shown in Fig. 4.

The  $\beta$  values evaluated by the two-level model nicely correlated with the computed values of the static first hyperpolarizability. They increased with the increase in size of the clusters and thus followed a monotonic behavior. The doping of Cu resulted in the highest hyperpolarizability values in each cluster. It was observed that the frequency-dependent first hyperpolarizability  $\beta_{(-2w;w,w)}$  was always higher than the static one and the pure Au-clusters showed the lowest values of the average first hyperpolarizability.

Comparing the pure Au and Cu-doped clusters, the  $\beta_{(0;0,0)}$  for the pure cluster was always less than for the mixed Au-Cu clusters. Fascinatingly, the  $\beta_{(-2w;w,w)}$  of the pure clusters was increased by doping with Cu atoms. For the 2A clusters, the value of the second-order polarizability for the pure Au and pure Cu clusters was less than for the mixed Au<sub>1</sub>-Cu<sub>1</sub> clusters due to the large change in the electric dipole moment according to the two levels. The Au<sub>3</sub>Cu<sub>1</sub> nonsymmetric structure showed notable values of hyperpolarizability, which signified their NLO behavior as compared to Cu<sub>2</sub>, Cu<sub>3</sub>, and Cu<sub>4</sub> symmetric structures. The average first hyperpolarizability of clusters containing a single Au atom (Au<sub>1</sub>Cu<sub>4</sub>, Au<sub>1</sub>Cu<sub>5</sub>, and Au<sub>1</sub>Cu<sub>6</sub>) cluster was higher than the mixed doped clusters due to the deformation





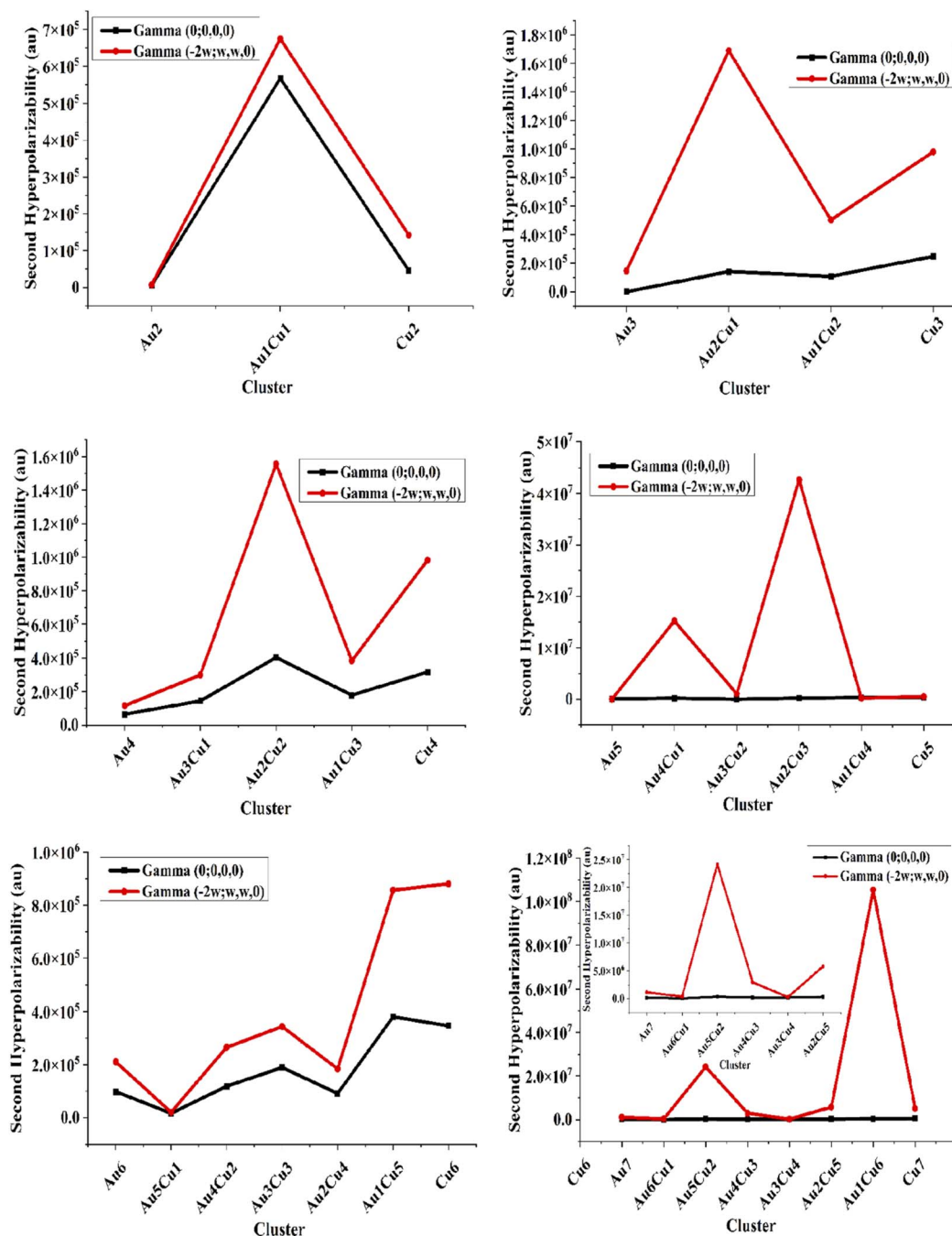


Fig. 5 Second hyperpolarizability comparison for the pure and Cu-doped gold clusters.

produced by doping, which enhances the dipole moment and changes the electronic structure of the clusters. The electronic dipole moment followed the same trend of variation as observed in the first and second hyperpolarizabilities.

### Third-order NLO polarizability

To calculate the static and frequency-dependent second hyperpolarizability or third-order polarizability of the pure and Cu-doped gold clusters, DFT calculations at the CAM-B3LYP/LANL2DZ level were used, and the results for the average

second hyperpolarizabilities are summarized in Table 2. The static third-order static polarizabilities  $\gamma_{(0;0,0)}$  for pure Au-clusters Au<sub>2</sub>, Au<sub>3</sub>, Au<sub>4</sub>, Au<sub>5</sub>, Au<sub>6</sub>, and Au<sub>7</sub> were  $6.01 \times 10^3$ ,  $1.42 \times 10^3$ ,  $6.52 \times 10^4$ ,  $8.67 \times 10^4$ ,  $9.84 \times 10^4$ , and  $1.83 \times 10^5$  au, respectively. For the pure Cu (2–7) clusters, these were in the range of  $4.56 \times 10^4$ – $6.30 \times 10^5$  au. The highest static second hyperpolarizabilities for the doped clusters Au<sub>2</sub>Cu<sub>2</sub>, Au<sub>1</sub>Cu<sub>6</sub>, Au<sub>1</sub>Cu<sub>1</sub>, and Cu<sub>7</sub> were  $4.04 \times 10^5$ ,  $4.22 \times 10^5$ ,  $5.67 \times 10^5$ , and  $6.30 \times 10^5$ , respectively. The third-order frequency-dependent  $\gamma_{(-2w;w,w,0)}$  polarizabilities for the pure gold clusters Au<sub>2</sub>, Au<sub>3</sub>, Au<sub>4</sub>, Au<sub>5</sub>, Au<sub>6</sub>, and Au<sub>7</sub> were  $8.03 \times 10^3$ ,  $1.45 \times 10^5$ ,  $1.16 \times 10^5$ ,



$4.24 \times 10^4$ ,  $2.10 \times 10^5$ , and  $1.15 \times 10^6$  au, respectively. The frequency-dependent  $\gamma_{(-2w;w,w,0)}$  polarizabilities for the pure  $\text{Cu}_m$  ( $m = 2-7$ ) clusters were in the range of  $1.42 \times 10^5$ – $5.15 \times 10^6$  au. It was found that the second hyperpolarizability was much greater than the first hyperpolarizabilities and also that the frequency-dependent  $\gamma_{(-2w;w,w,0)}$  polarizability was always higher than the static one. The average third-order frequency-dependent  $\gamma_{(-2w;w,w,0)}$  polarizabilities were very high for the mixed Au–Cu doped  $\text{Cu}_7$ ,  $\text{Au}_2\text{Cu}_5$ ,  $\text{Au}_4\text{Cu}_1$ ,  $\text{Au}_5\text{Cu}_2$ ,  $\text{Au}_2\text{Cu}_3$ , and  $\text{Au}_1\text{Cu}_6$  clusters at  $5.15 \times 10^6$ ,  $5.67 \times 10^6$ ,  $1.52 \times 10^7$ ,  $2.42 \times 10^7$ ,  $4.26 \times 10^7$ , and  $1.05 \times 10^8$  au, respectively. These computed first and second hyperpolarizabilities for the pure and Cu-doped gold clusters were larger than the already reported NLO materials, e.g.,  $51.99 \times 10^4$  to  $138.25 \times 10^4$  au for superhalogen and superalkali doped phosphorenes,<sup>43</sup> and  $6.1 \times 10^7$  au for bimetallic superalkali clusters.<sup>44</sup> To highlight the difference between the pure and Cu-doped Au-clusters, a graphical comparison was made for the third-order polarizability or second hyperpolarizability, as shown in Fig. 5.

A graphical comparison between the static and frequency-dependent third-order polarizability is shown in Fig. 5. It was observed that the frequency-dependent second hyperpolarizability  $\gamma_{(-2w;w,w,0)}$  was always higher than the static one. As we moved from  $\text{Au}_2$  to  $\text{Au}_6$  clusters, the second hyperpolarizability of the pure gold clusters was less than for the pure Cu clusters. Comparing between the Cu-doped clusters and pure clusters, the  $\gamma_{(0;0,0)}$  of the pure clusters was less than for the mixed Au–Cu clusters. Interestingly, the  $\gamma_{(0;0,0)}$  of the pure clusters was increased by doping with Cu atoms. For the  $\text{Au}_2$  clusters, the value of  $\gamma_{(0;0,0)}$  and  $\gamma_{(-2w;w,w,0)}$  for the  $\text{Au}_1\text{Cu}_1$  cluster were higher than for the pure gold and copper clusters due to the change in the electronic structure. Also, it was found that the second hyperpolarizability for the symmetric structure was comparatively less than for the nonsymmetric structures, like the  $\text{Au}_2$  to  $\text{Au}_7$  clusters, where the second hyperpolarizability for the  $\text{Au}_2\text{Cu}_2$ ,  $\text{Au}_3\text{Cu}_3$ , and  $\text{Au}_4\text{Cu}_4$  symmetric structures was less than for the  $\text{Au}_3\text{Cu}_1$  nonsymmetric structure. The  $\gamma_{(0;0,0)}$  of clusters containing a single Au ( $\text{Au}_1\text{Cu}_4$ ,  $\text{Au}_1\text{Cu}_5$ ,  $\text{Au}_1\text{Cu}_6$ ) cluster was higher than for the others with more than one Au-doped clusters due to the deformation produced by the doping, which enhanced the dipole moment. The  $\text{Au}_4\text{Cu}_2$ ,  $\text{Au}_5\text{Cu}_1$ , and  $\text{Au}_5\text{Cu}_2$  clusters showed very high  $\gamma_{(-2w;w,w,0)}$  third-order polarizability due to their high dipole moment and complex structures. The excitation energy ( $\Delta E$  in eV) and oscillator strength ( $f_o$ ) for all the doped and pure Au-clusters were also investigated. The excitation energies ( $\Delta E$ ) were 2.73, 1.23, 1.35, 1.26, 2.85, and 2.68 eV for the  $\text{Au}_2$ ,  $\text{Au}_3$ ,  $\text{Au}_4$ ,  $\text{Au}_5$ ,  $\text{Au}_6$ , and  $\text{Au}_7$  pure Au-clusters, and decreased from 2.85–1.23 eV. These smaller values of  $\Delta E$  for the doped complexes led to higher values of hyperpolarizability. The oscillator strength ( $f_o$ ) for the pure Au-cluster was 0 and it changed upon Cu-doping, whereby the values of  $f_o$  were 0.0218, 0.0063, and 0.0030 for  $\text{Au}_4\text{Cu}_2$ ,  $\text{Au}_5\text{Cu}_1$ , and  $\text{Au}_5\text{Cu}_2$ , respectively. All the calculated results for the excitation energy ( $\Delta E$ ) and oscillator strength ( $f_o$ ) were in accordance with the two-level model and related with their hyperpolarizability results. These computed first and second hyperpolarizabilities for the pure

and Cu-doped gold clusters were much higher than those of the already reported NLO materials. Thus, the third-order NLO polarizability analysis showed that all the compounds in the present investigation are potential candidates for efficient third-order NLO applications.

## Conclusions

The theoretical study of the current investigated Au–Cu clusters showed that the doping of Cu atoms did not largely affect the structural framework of the Au-clusters, while doping with Cu significantly enhanced the NLO properties of the clusters. A significant decrease in the HOMO–LUMO energy gap from 3.67 to 0.91 eV was observed, which was reduced by 74% by doping with Cu. Similarly, the optical properties, including the linear isotropic and anisotropic polarizabilities, increased by 4 to 5 times. Among all the structures, the  $\text{Au}_3\text{Cu}_1$  and  $\text{Au}_1\text{Cu}_6$  clusters showed the largest static  $\beta_{(\text{total})} = 4.73 \times 10^3$  au and frequency-dependent  $\beta(-2w; w, w) = 1.26 \times 10^6$  au first hyperpolarizabilities, respectively. The large static and frequency-dependent second hyperpolarizabilities of  $6.30 \times 10^5$  au and  $1.05 \times 10^8$  au were exhibited by  $\text{Cu}_7$  and  $\text{Au}_1\text{Cu}_6$ , respectively. Theoretically, it was shown that a single-doped system  $\text{Au}_m$  ( $2 \geq m \leq 7$ ) is a better choice to get a high NLO response as compared to other combinations. These computed values of the second and third-order polarizabilities are larger than some already reported clusters. This methodology can be extended to explore the NLO response of gold nanometallic structures having different shapes, such as stars, cage, rods, and core-shell.

## Author contributions

All authors contributed to the study conception and design. Materials designing, data collection and analysis were performed by F. ·H, R. H, K. A, M. U. K. and J. Y. The initial draft was prepared by F. ·H, R. H, Z. I and M. A. The final draft was edited by S. M, K. A and M. A. All authors read and approved the final manuscript.

## Conflicts of interest

There are no conflicts to declare.

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