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# Mechanistic insights of the reduction of gold salts in the Turkevich protocol†

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This paper presents fundamental understanding of the mechanism of the Turkevich protocol, the method recommended by the National Institute of Standards and Technology for the synthesis of gold nanoparticles using sodium citrate as reducing agent. Herein, we reveal that the Turkevich mechanism consists of two consecutive reduction steps ( $\text{Au}^{3+} \rightarrow \text{Au}^+ \rightarrow \text{Au}^0$ ) rather than a reduction followed by the disproportionation reaction as conventionally believed. This new understanding has profound implications: i. the second reduction step ( $\text{Au}^+ \rightarrow \text{Au}^0$ ), rather than the previously postulated first reduction step, is the rate-limiting reduction step and ii. the formation of acetone dicarboxylate ( $\text{DC}^{2-}$ ) as an intermediate product through the oxidation of citrate has a key role as stabilizer and as a reducing agent (stronger than sodium citrate). This knowledge enables the synthesis of monodispersed gold nanoparticles with sizes ranging from  $5.2 \pm 1.7$  nm to  $21.4 \pm 3.4$  nm, with the lower end considerably smaller than previously reported through the Turkevich route. This work provides fundamental guidance for the controllable synthesis of nanoparticles using  $\text{DC}^{2-}$  as a reducing agent directly applicable to other precious metals.

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## 1. Introduction

Gold nanoparticles (Au NPs) have potential applications in medicine, catalysis, electronics and photonics due to their unique chemical and physical properties compared with their bulk counterpart.<sup>1–6</sup> Au NPs are usually synthesized by the reduction of chloroauric acid ( $\text{HAuCl}_4$ ) with a reducing agent such as sodium borohydride,<sup>7</sup> sodium citrate,<sup>8</sup> and ascorbic acid.<sup>9</sup> The Turkevich protocol is the most reliable and popular method for the synthesis of Au NPs,<sup>10</sup> and consequently the recommended method by the National Institute of Standards and Technology for reference material preparation.<sup>11</sup> It consists of the use of sodium citrate to reduce chloroauric acid at elevated temperatures in an aqueous medium under various conditions. The citrate ion also acts as a stabilizer, which can be easily replaced post-synthesis by other ligands, in favor of the functionalization of Au NPs.<sup>12</sup>

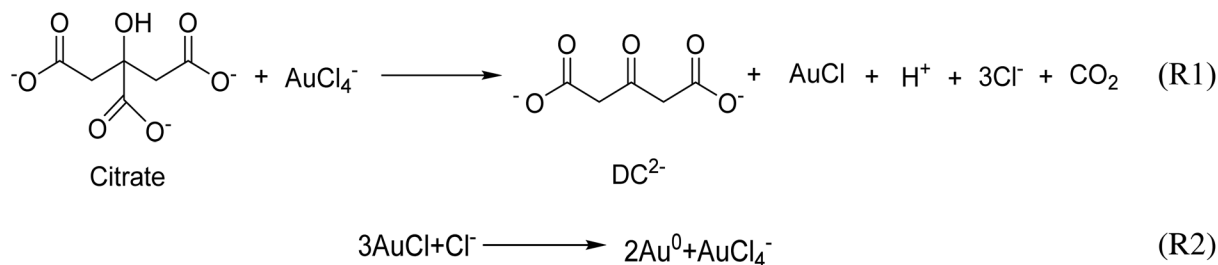
The accepted mechanism of the Turkevich method for the synthesis of gold nanoparticles consists of the initial redox reaction (R1) (Scheme 1),<sup>10,13,14</sup> where trivalent gold gets reduced to monovalent gold by citrate which in turn gets oxidised to acetone dicarboxylate ( $\text{DC}^{2-}$ ), the conjugated base of dicarboxyacetone (DCA). This first redox step is deemed as the

rate-determining step.<sup>15</sup> Consecutively, the disproportionation reaction (R2) (Scheme 1) takes place where metallic gold and trivalent gold are produced. In the original paper by Gammons *et al.* investigating the disproportionation reaction,<sup>16</sup> the reaction was conducted in the absence of any reducing agent and stabilizer. It is thus expected that in the presence of reducing agents, such as sodium citrate and  $\text{DC}^{2-}$  present in the conventional Turkevich protocol, a competitive parallel reduction of  $\text{Au}^+$  to metallic gold might take place, however, to the best of our knowledge, this parallel reduction reaction has not been investigated.

Much research has been done to unravel the mechanism of Turkevich protocol and optimize its synthesis conditions in order to obtain monodispersed Au NPs with controllable sizes.<sup>8,10,13–15,17–19</sup> The sodium citrate :  $\text{HAuCl}_4$  molar ratio has shown to have a great effect on the particle size,<sup>20</sup> while the temperature does not play a considerable role from 50 °C to 100 °C.<sup>21</sup> Increasing the sodium citrate :  $\text{HAuCl}_4$  molar ratio leads to smaller Au NPs due to higher nucleation rate and consequently higher concentration of generated nuclei.<sup>22–24</sup> The pH value of the reaction medium has also been shown to have an effect on both the particle size and kinetics.<sup>10,14,17,18,25,26</sup> This pH effect is related to the hydrolysis of the citrate and chloroauric species,<sup>14</sup> as  $\text{HCit}^{2-}$  was found to be the strongest reducing species whose concentration is maximised at pH 5.6.<sup>10,17,18,26</sup> Similarly,  $\text{AuCl}_4^-$  is the most reactive gold precursor compared with its hydrolysed species at high pH values,  $\text{AuCl}_3(\text{OH})^-$ ,  $\text{AuCl}_2(\text{OH})_2^-$ ,  $\text{AuCl}(\text{OH})_3^-$  and  $\text{Au}(\text{OH})_4^-$ . As a result, the reaction, especially the nucleation stage, is much

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**Scheme 1** The accepted mechanism for Turkevich protocol, consisting of the initial redox reaction (R1) and the disproportionation reaction (R2).

faster at pH values between  $\sim 3.7$ – $6.5$ , compared with pH of  $\sim 6.5$ – $7.7$ .<sup>17</sup> The Au NPs size consequently decreases from 22 nm to 11 nm when pH increases from 3.0 to 5.4, then increases to 28 nm when pH further increases from 5.4 to 7.5. The pH has also a dominating effect on the polydispersity of the Au NPs, being around 10% at pH values between 5.4 and 6.8, increasing to  $\sim 20\%$  when pH is below 4.5 or higher than 7.5.<sup>18</sup> The optimization of the reaction (1) by adjusting the pH range of chloroauric acid solution to 3–3.6, and the pH of the mixed solution between 5.4–6.0, at boiling temperature, enabled the synthesis of the smallest Au particles using the conventional Turkevich method to date ( $11.0 \pm 0.4$  nm).<sup>18,27</sup>

The order of addition of chloroauric acid and sodium citrate during the synthesis has a great effect on the particle size distribution.<sup>11,28</sup> In the standard Turkevich method, a concentrated sodium citrate solution is added to a diluted HAuCl<sub>4</sub> solution. However, in the so-called reversed Turkevich method, where the concentrated acidic HAuCl<sub>4</sub> solution is injected to a diluted sodium citrate solution, localised low pH values promote AuCl<sub>4</sub><sup>-</sup> species which contributes to the generation of more seeds and leads to smaller average particle sizes.<sup>11</sup>

Despite the numerous mechanistic studies of the Turkevich method, most of the researchers have focused on the optimisation of the first Au<sup>3+</sup> to Au<sup>+</sup> reduction step (R1).<sup>29,30</sup> However, some authors, including Turkevich, identified the potential role of the intermediate product DC<sup>2-</sup> as a reducing agent.<sup>8</sup> Doyen *et al.* confirmed the formation of DCA (at 75 °C) and its derived products acetoacetic acid, acetic acid and acetone by NMR.<sup>31</sup> The substitution of 10% sodium citrate by DCA was found to accelerate the reaction and decrease the size and Au NPs in the standard Turkevich protocol. It was postulated that DCA forms a multidentate complex with Au<sup>+</sup> which favors the disproportionation reaction, associated to the smaller Au NPs with narrower size distribution obtained in the reversed Turkevich method.<sup>28</sup>

The morphology of the gold nanoparticles can be controlled by the external addition of chemicals, for example, the presence of L-tyrosine leads to the formation of gold nanoflowers.<sup>32</sup> Gold nanowires, which self-assemble to gold foams, can be obtained by halting the Turkevich reaction or controlling low sodium citrate concentration.<sup>33–35</sup> Similarly, the addition of trace amounts of tannic acid in the presence of poly-vinyl-pyrrolidone (PVP) can lead to sub-10 nm particles.<sup>36,37</sup>

This paper presents a systematic study to shed light upon the mechanism of Turkevich for the synthesis of Au NPs by revealing the actual role of DC<sup>2-</sup> intermediate as a reducing agent and confirming the reduction of Au<sup>+</sup> over the previously believed disproportionation reaction. This new knowledge enables the design of the synthetic Turkevich conditions to tune the gold particle size while obtaining narrow size dispersions.

## 2. Experimental procedures

### Materials

Gold(i) chloride AuCl (99.9% trace metal basis), DCA (technical grade), sodium citrate dehydrate (>99%), chloroauric acid (99.99% trace metal basis, 30 wt%), citric acid monohydrate (>99%) and bovine serum albumin (BSA) are purchased from Sigma Aldrich. Sodium hydroxide (0.1 M) is purchased from Fisher Chemical. All chemicals are used without further purifications. 18.2 MΩ cm pure water is from the MilliQ ultra-purification system.

### Synthesis of Au NPs

The synthesis of Au NPs is carried out following the reversed Turkevich protocol in a 100 mL glass round-bottom flask reactor equipped with a glass condenser. The water bath temperature is controlled by an ETS-D5, IKA hotplate. In a typical Turkevich synthesis, 48.0 mL of water are pre-heated to 80 °C in a stirred round-bottom flask where a thermometer is introduced. Once the temperature is stable, 672 μL 0.125 M sodium citrate solution and 328 μL 0.125 M citric acid solution are added to the reaction vessel. The reaction starts when 1.0 mL 0.0125 M HAuCl<sub>4</sub> solution is added to reaction medium. In the reduction of HAuCl<sub>4</sub> by DC<sup>2-</sup> at 80 °C, 46.0 mL of water are pre-heated in a stirred round bottom flask, once the temperature is stable, 2.5 mL 0.1 M NaOH are added to adjust the pH at 4.8 (for comparison purposes with other experiments). 1 mL 0.0125 M HAuCl<sub>4</sub> is added to start the reaction about 3 seconds after the addition of 0.5 mL 0.25 M DCA. 1.5 mL samples are collected periodically (every one minute), cooled down in an ice water bath to quench the reaction and analyzed by UV-vis spectroscopy. The reaction is considered completed when the UV-vis spectrum does not change with time. High reproducible

results are obtained in all cases, confirmed by doing the experiments in duplicates and comparing their respective final UV-vis spectra.

### Characterization

The UV-vis spectrum of the Au NP colloidal suspensions is measured using an Agilent Cary-60 UV-Vis spectrophotometer. pH is measured using a HI98100 Checker Hanna pH indicator, with an accuracy of 0.2. Transmission electron microscopy (TEM) images are obtained using a FEI Tecnai 20 equipment. ImageJ software is used to analyze the area-equivalent diameter of more than 200 particles. TEM samples are prepared according to Michen *et al.*'s protocol,<sup>38</sup> which adopts BSA to prevent post-synthesis agglomeration. 1 mL of colloidal solution is mixed with 1.0 mL 0.3 mg mL<sup>-1</sup> BSA solution before kept at 5 °C for 2 h. Then, 5 µL mixed sample is drop on a 400 mesh carbon coated copper TEM grid and left to dry at room temperature. Scanning electron microscopy is carried out using a FEI Philips XL30 SFEG equipped with energy dispersive X-ray microscopy (EDS). X-ray photoelectron spectroscopy (XPS) is used to analyse the oxidation state of gold precursor using a Thermo Fischer Escalab 250 Xi with Al K alpha as source with 1486.68 eV energy and 50 eV pass energy. Dwell time is set as 50 ms and step size is 0.1 eV. X-ray spot size is 200 micrometres. The binding energy scale is calibrated by setting the Au<sub>4f7/2</sub> binding energy to 84.0 eV.<sup>39,40</sup> XPS samples are prepared dropping 5 µL of the Au NP colloidal solution into a silicon sample wafer, dried in a vacuum tank and then stored under argon atmosphere before characterization. Electrochemical experiments are performed to measure the oxidation potentials of sodium citrate and DC<sup>2-</sup> utilising a standard three-electrode setup connected to a potentiostat (VSP, Bio-Logic). A glassy carbon rod (Alfa Aesar, 2 mm diameter) is utilised as working electrode, with an exposed geometric surface area of 0.0314 cm<sup>2</sup>. Pt wire (Alfa Aesar, >99.997% purity, 0.5 mm diameter) is utilised as counter electrode, Hg/HgSO<sub>4</sub> (SI Analytics) is utilised as reference electrode. Prior to each experiment, glassy carbon and Pt electrodes are thoroughly rinsed in sequence with isopropanol and ultrapure water. All cyclic voltammetry experiments are performed at a scan speed of 10 mV s<sup>-1</sup>.

## 3. Results and discussion

To understand the mechanism of the Turkevich method for the synthesis of Au NPs, a first series of reactions at 25 and 80 °C is carried out following the reversed Turkevich method to investigate the actual role of citrate and its *in situ* formed oxidation form, dycarboxyacetone (DCA). Chloroauric acid (HAuCl<sub>4</sub>) is used as gold precursor. The Au:reducing agent molar ratio is kept constant at 1:10 at an initial pH 4.8–4.9 adjusted by addition of sodium citrate and citric acid mixtures or by addition of NaOH (Table 1).

Although the Au NPs synthesis is very slow at room temperature using citrate as reducing agent (A), with negligible formation of Au NPs after 20–25 min, changes in absorbance in the UV-region are observed (Fig. S1†), which are believed to be related to the fast substitution of one of the chloride ions by citrate in the Au environment as previously suggested by Ojea-Jiménez *et al.*<sup>14</sup> Increasing the reaction time to 24 h leads to the slow formation of Au NPs (26.0 ± 6.3 nm). On the other hand, the faster nucleation at 80 °C (B) leads to smaller Au NPs (15.2 ± 4.2 nm). It is important to note that the UV-vis spectra of the particles synthesized at room temperature show a significant absorbance at ~700 nm (Fig. S1†), previously associated to near-surface nucleation events during particle growth<sup>41</sup> a phenomena probably responsible of the irregular Au NPs shape in Fig. 3a (see below) and not visible during the synthesis at 80 °C. It is likely that autocatalysed secondary nucleation on the surface of existing gold surface dominates the process under slow reduction conditions at room temperature.<sup>41</sup> This effect of temperature on size is associated to the kinetics of reduction of the gold precursor from Au<sup>3+</sup> → Au<sup>+</sup> → Au<sup>0</sup> and consequently the generation of nuclei and/or seeds. A number of synthesis mechanisms have been reported to date from the original LaMer's nucleation theory<sup>42</sup> to the more comprehensive seed-mediated growth mechanism.<sup>18,25,43</sup> In both cases, when colloidal stability is guaranteed, the resulting size of the particles is directly related to kinetics of the reduction stage which determines not only the concentration of nuclei/seeds where growth take place, but also the amount of remaining precursor. Thus, faster reduction of the gold precursor at higher temperatures leads to a higher rate of nuclei/

**Table 1** Summary of reaction conditions for the investigation of the role of citrate and DCA in the synthesis of Au NPs

No	[HAuCl <sub>4</sub> ]/mM	[AuCl]/mM	[DCA]/mM	[Sodium citrate]/mM	[Citric acid]/mM	[NaOH]/mM	PVP/mM	T/°C	Initial pH <sup>a</sup>	Final pH	Diameter <sup>b</sup> /nm
A	0.25	—	—	1.68	0.82	—	—	25	4.9	5.0	26.0 ± 6.3
B	0.25	—	—	1.68	0.82	—	—	80	4.9	4.9	15.2 ± 4.2
C	0.25	—	2.5	—	—	5	—	25	4.9	7.8	21.1 ± 6.9
D	0.25	—	2.5	—	—	5	—	80	4.9	7.5	10.7 ± 1.4
E	0.25	—	2.5	2.5	—	1.62	—	25	4.8	6.0	23.0 ± 5.6
F	0.25	—	2.5	2.5	—	1.62	—	80	4.8	6.7	13.9 ± 3.0
G	—	0.25	—	—	—	0.01	5	80	4.8	3.9	11.9 ± 4.2
H	—	0.25	—	1.35	1.15	—	—	80	4.8	4.9	14.8 ± 4.3
I	—	0.25	2.5	—	—	4	—	25	4.8	7.0	9.8 ± 3.2
J	—	0.25	2.5	—	—	4	—	80	4.8	7.8	6.8 ± 1.5

<sup>a</sup>The initial pH was always measured at 25 °C for comparison purposes. <sup>b</sup> Measured by TEM.

seeds formation and a smaller particle size in agreement with our observations. In addition, the resulting dispersity of the particles will depend on the degree of separation between the nucleation/seed formation and growth stages.<sup>44</sup>

Reduction of gold precursor by citrate leads to its oxidation and thus the formation of  $\text{DC}^{2-}$ . Although the concentration of the latter is normally comparatively small to that of citrate ions during the conventional Turkevich method, for comparison purposes, gold nanoparticles are also synthesized by using directly  $\text{DC}^{2-}$  as reducing agent rather than sodium citrate (C-D, note that initial pH has been kept constant by adding NaOH into the reaction medium). In this case, the formation of Au NPs can take place even at room temperature (e.g. 25 °C) while increasing the temperature to 80 °C leads to faster kinetics (as expected). Due to the faster nucleation at 80 °C considerably smaller particles ( $10.7 \pm 1.4$  nm) are obtained compared with 25 °C ( $21.1 \pm 6.9$  nm).<sup>18,25</sup>

Fig. 1 shows the time-evolution of the formation of Au NPs using the absorption data. At 25 °C, while the reduction of gold is considerably quicker during the first ~30 min of reaction when  $\text{DC}^{2-}$  is used as reducing agent compared with citrate, the kinetics of the reduction greatly slows down after this time. Indeed, it requires ~20 days to achieve full conversion. On the other hand, full reduction of the gold precursor is

achieved in ~24 h in the presence of citrate despite the slower initial kinetics. The decrease of reaction kinetics using  $\text{DC}^{2-}$  as the reducing agent is likely to its adsorption on the surface of the formed Au NPs, resulting in a decrease of its concentration in the solution.<sup>12</sup> In addition, changes in pH (~0.1) in the first ~30 min could also affect the hydrolysis of the gold precursor from  $\text{AuCl}_4^-$  at pH of 4.9 to  $\text{AuCl}_3(\text{OH})^-$ , which is less reactive.<sup>26</sup> Indeed, after 20 days, pH increases from the initial value of 4.9 to 7.8 when  $\text{DC}^{2-}$  is used as the reducing agent, a rise caused by the hydroxide ions resulting from the decomposition of  $\text{DC}^{2-}$ , as shown in reaction (3), Scheme 2.<sup>13</sup> In contrast, sodium citrate acts as a buffer keeping the pH constant during the reaction. In any case, since the initial seed particle formation is faster when  $\text{DC}^{2-}$  is used as the reducing agent, the resulting Au NPs are smaller,  $21.1 \pm 6.9$  nm, compared with  $26.0 \pm 6.3$  nm when sodium citrate is the reducing agent.

The differences in the kinetics of the reduction are also observed at 80 °C, however, in this case, full conversion of the Au NPs is observed in both cases within ~15 min (Fig. 1b). Similarly than before, when  $\text{DC}^{2-}$  is used as reducing agent, smaller and more monodispersed Au NPs ( $10.7 \pm 1.4$  nm,  $\pm 13\%$ , Fig. 3d) are obtained compared with sodium citrate ( $15.2 \pm 4.2$  nm,  $\pm 28\%$ , Fig. 3b). Both colloidal stability and

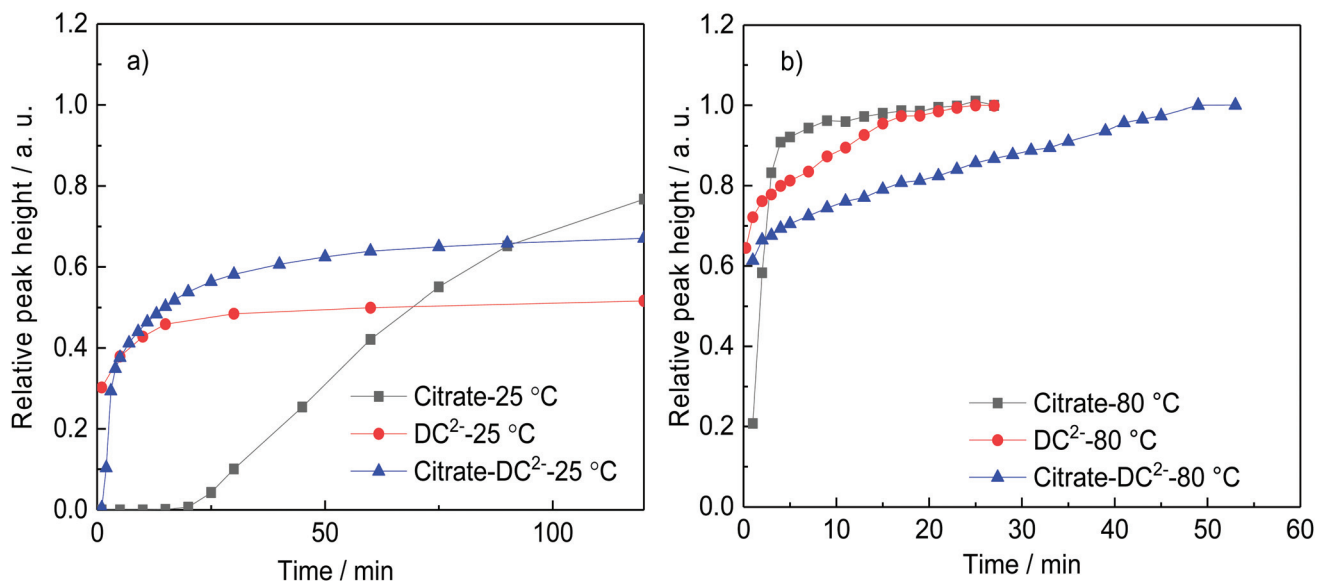
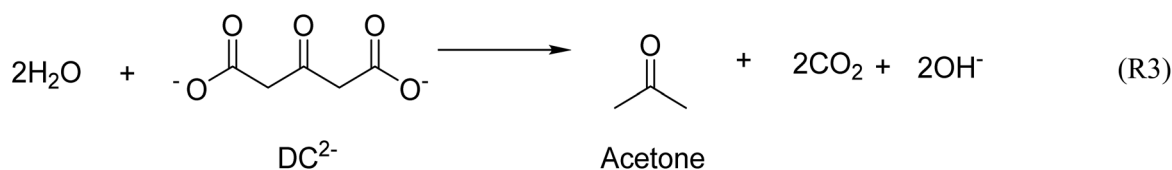
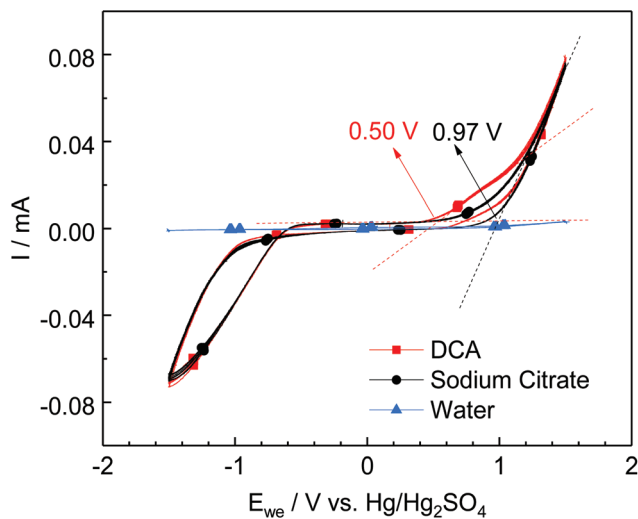


Fig. 1 Comparison of the rate of formation of gold nanoparticles (represented by the relative height of the absorption peak) using sodium citrate or  $\text{DC}^{2-}$  as a reducing agent at a. 25 °C and b. 80 °C. Conditions:  $[\text{HAuCl}_4]$ : 0.25 mM, pH: 4.8–4.9, reducing agent: [sodium citrate]: 2.5 mM,  $[\text{DC}^{2-}]$ : 2.5 mM; or [sodium citrate] 2.5 mM +  $[\text{DC}^{2-}]$  2.5 mM, as in experiments A–F (Table 1).



Scheme 2 Decomposition of  $\text{DC}^{2-}$ .<sup>13</sup>



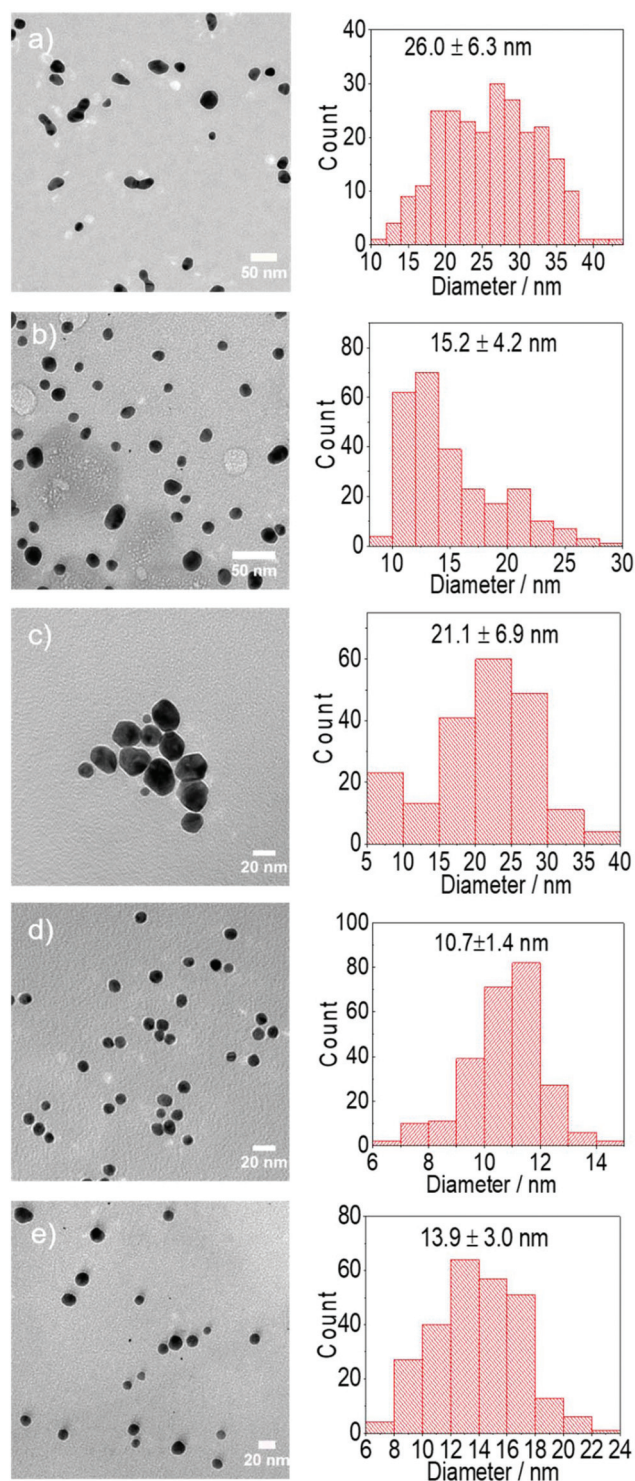


**Fig. 2** Current vs. voltage response for sodium citrate ([sodium citrate]: 1.35 mM, [citric acid]: 1.15 mM),  $\text{DC}^{2-}$  ([DCA]: 2.5 mM, [NaOH]: 4 mM), and diluted HCl solution. Conditions: 25 °C, pH 4.9.

chemistry of reduction should be taken into consideration to explain the difference.<sup>25</sup> As mentioned above,  $\text{DC}^{2-}$  promotes a faster reduction kinetics of the gold precursor leading to a higher concentration of initial seeds. In addition,  $\text{DC}^{2-}$  has a stronger stabilization effect compared with citrate according to Grasseschi *et al.*,<sup>12</sup> which may stabilize smaller stable seeds compared with citrate, leading to more seeds even when the same amount of gold precursor is initially reduced.<sup>18,25</sup> Both factors simultaneously lead to a higher concentration of seeds and thus, they lead to smaller Au NPs for the same precursor concentration.

These results agree with the measured oxidation potential of the two reducing agents as shown in Fig. 2, where it is shown that  $\text{DC}^{2-}$  is a stronger reducing agent with an oxidation potential of 0.50 V, compared to sodium citrate at 0.97 V at the studied conditions.

The reason of the higher monodispersity of the Au NPs formed using  $\text{DC}^{2-}$  as reducing agent at 80 °C is threefold – firstly,  $\text{DC}^{2-}$  serves as the stabilizer for Au NPs. Although  $\text{DC}^{2-}$  is not stable in the aqueous solution, its enol form coordinates with Au NPs forming a complex and consequently stabilising DCA *versus* its decomposition.<sup>12</sup> On the other hand, the enol-Au NPs complex is negatively charged, stabilising the particles electrostatically. Secondly, a fast nucleation takes place due to the high reducing potential of  $\text{DC}^{2-}$  as discussed above, leading to the formation of seeds to which the remaining  $\text{DC}^{2-}$  coordinates.<sup>12</sup> Both aspects consequently lead to a lower concentration of  $\text{DC}^{2-}$  in the solution, decreasing the reduction kinetics and thus the formation of new nuclei. Thirdly, an increase in pH due to  $\text{DC}^{2-}$  decomposition (as discussed above) leads to the hydrolysis of the remaining gold precursor to less reactive forms. These new conditions reduce the seed formation rate to a minimum, separating the seed formation and their growth stages.<sup>18,44</sup>

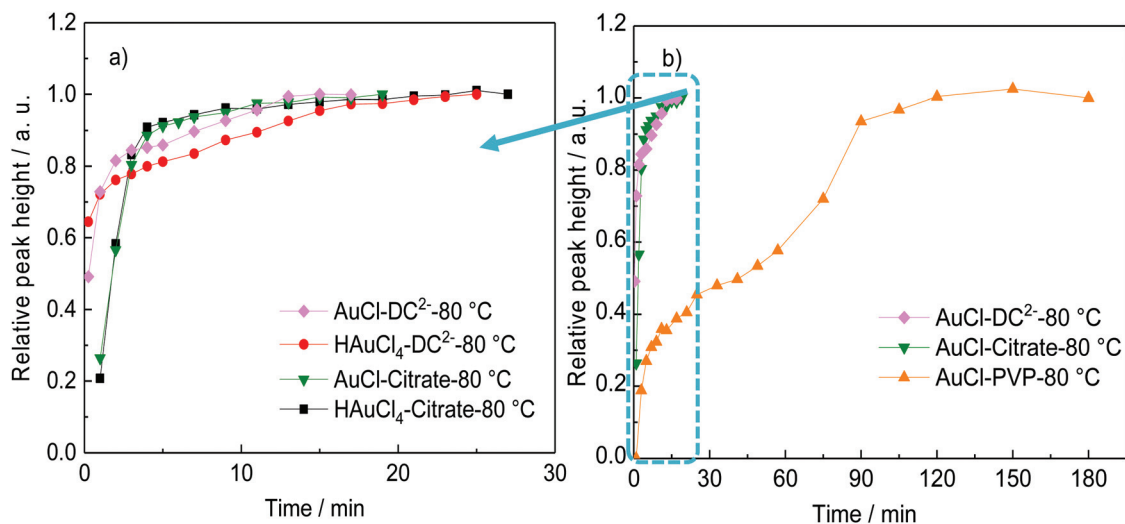


**Fig. 3** Representative TEM images and particle size distribution of Au NPs synthesised using chloroauric acid as the precursor a. [sodium citrate] 2.5 mM at 25 °C, b. [sodium citrate] 2.5 mM at 80 °C, c. [ $\text{DC}^{2-}$ ] 2.5 mM at 25 °C, d. [ $\text{DC}^{2-}$ ] 2.5 mM at 80 °C, e. [sodium citrate] 2.5 mM and [ $\text{DC}^{2-}$ ] 2.5 mM at 80 °C. Conditions: [ $\text{HAuCl}_4$ ]: 0.25 mM, pH: 4.8–4.9, samples are taken after reaction completion.

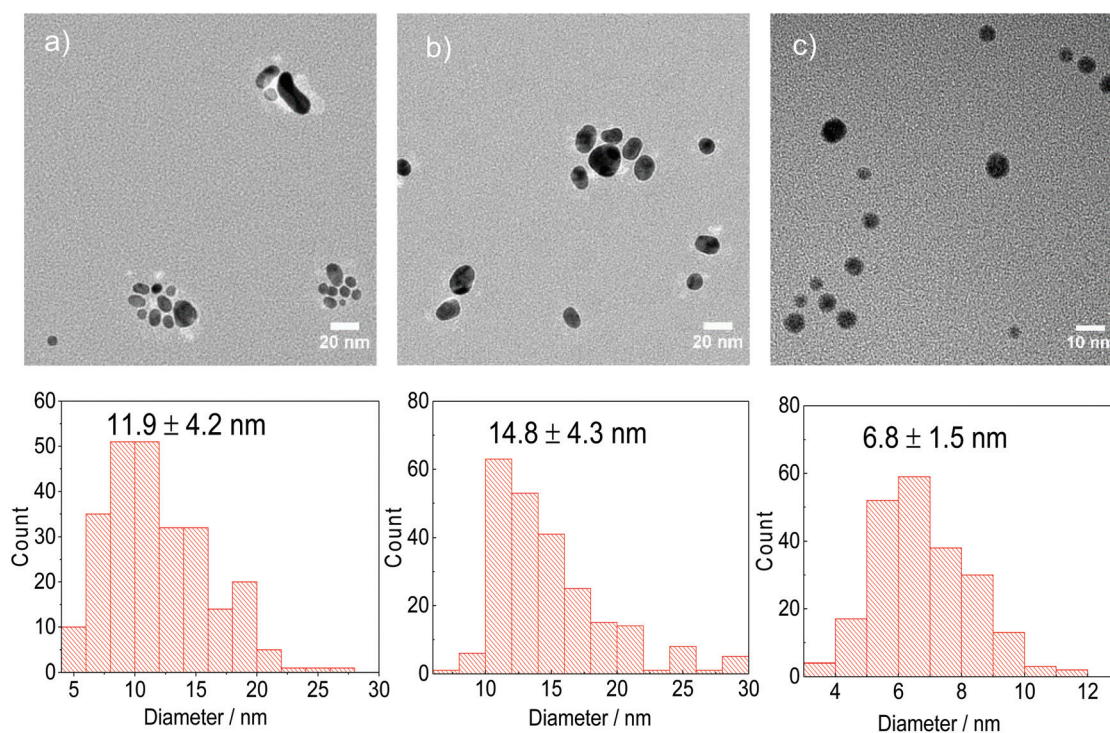
Both Au NPs stabilized by  $\text{DC}^{2-}$  and citrate are stable for at least 5 months as confirmed by UV-vis analysis. Grasseschi *et al.*<sup>12</sup> have previously demonstrated that  $\text{DC}^{2-}$  have a stronger interaction with the Au surface than citrate ions however, both can be easily replaced by molecules with stronger affinity to

gold atoms, *e.g.* 4-mercaptopyridine, which enables the convenient post-synthesis functionalization of Au NPs.

It is important to note that this high monodispersity of Au NPs achieved with  $\text{DC}^{2-}$  as reducing agent is only feasible at pH values  $>\sim 4.8$  when the DCA fully deprotonates into  $\text{DC}^{2-}$



**Fig. 4** Comparison of the rate of formation of gold nanoparticles (shown by the relative height of the absorption peak) (a) using different gold precursors and reducing agents at 80 °C. Conditions:  $[\text{HAuCl}_4]$ : 0.25 mM,  $[\text{AuCl}]$ : 0.25 mM, [sodium citrate]: 2.5 mM,  $[\text{DC}^{2-}]$ : 2.5 mM, pH: 4.8–4.9. Original data are shown in Fig. S3.† (b) By the reduction or the disproportionation reaction of AuCl with  $\text{DC}^{2-}$  or sodium citrate as a reducing agent (same with (a)), or PVP as stabilizer in the absence of sodium citrate and  $\text{DC}^{2-}$ . Conditions:  $[\text{AuCl}]$ : 0.25 mM, [PVP monomer]: 5 mM, pH: 4.8, 80 °C.



**Fig. 5** Representative TEM images and particle size distribution of Au NPs synthesised using AuCl as the precursor a. in the absence of reducing agent using PVP as stabilizer, b. sodium citrate and c.  $\text{DC}^{2-}$ . Conditions:  $[\text{AuCl}]$ : 0.25 mM, [reducing agent]: 2.5 mM (when applicable), 80 °C pH: 4.8, time: 15–180 min.



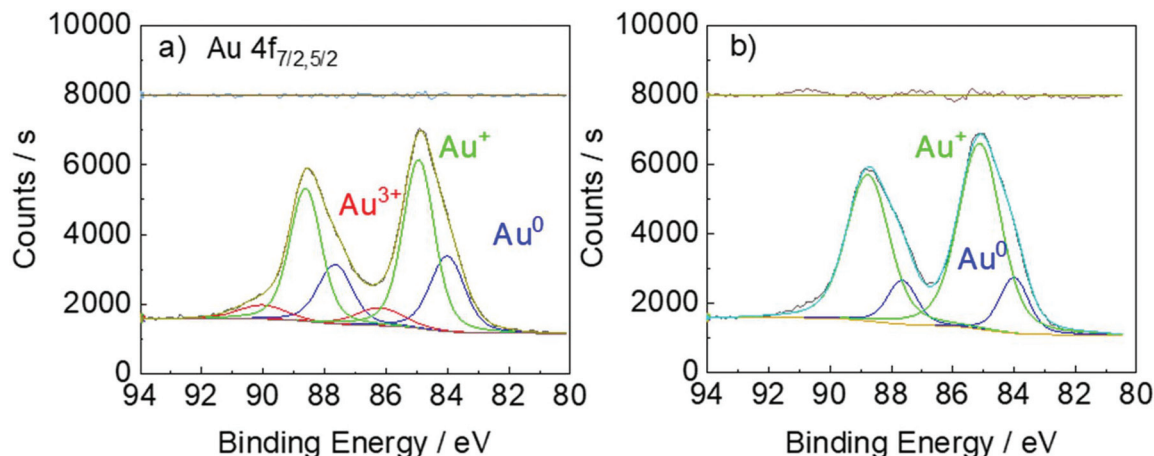


Fig. 6 XPS of a. [HAuCl<sub>4</sub>]: 0.25 mM, [DC<sup>2-</sup>]: 2.5 mM, sampling time: 5 min; b. [AuCl]: 0.25 mM, [DC<sup>2-</sup>]: 2.5 mM, sampling time: 15 min. Conditions: pH: 4.8, 25 °C.

Table 2 Summary of reaction conditions for the investigation of the Au<sup>+</sup> reduction step in the Turkevich method

No	[AuCl]/mM	[DCA]/mM	[Sodium citrate]/mM	[Citric acid]/mM	[NaOH]/mM	T/°C	Initial pH	Final pH	Diameter/nm
K	0.25	5			7.62	80	4.4	9.2	6.6 ± 2.1
L	0.25	5	5			80	4.4	6.5	6.9 ± 1.9
M	0.25	2.5	3.68	1.32		80	4.4	5.2	7.2 ± 1.6
N	0.25	0.25	2.53	2.47		80	4.4	4.4	10.8 ± 2.5
O	0.25	5		5		80	2.5	2.8	21.4 ± 3.4
P	0.25	5	3.21	1.79		80	3.6	5.1	12.6 ± 3.4
Q	0.25	5	5		7.72		6.2	7.6	5.2 ± 1.7
R	0.25	5	5		10	80	9.0	8.6	6.8 ± 2.5
S	0.25	5	5		10.1	80	9.9	9.0	7.8 ± 1.8

species. Indeed, in contrast to our observations, Wuitschick *et al.* concluded that the reduction of chloroauric acid with DCA results on the quick formation of polydispersed and non-reproducible Au NPs,<sup>25</sup> while this experiment is done three times to confirm the reproducibility as shown in Fig. S4.† The differences are related to the pH conditions. In their case, pH values <4.8, lead to the protonated form of DCA, which is known to have a low reducing and stabilisation power.<sup>11,28</sup> Similarly, Ojea-Jiménez *et al.* reported that no Au NPs formation is observed at pH 3.7 and 100 °C due to the low reactivity of the protonated form of DCA.<sup>28</sup> In addition, it is likely that high temperatures (*e.g.* boiling conditions) of these two studies lead to the fast decomposition of DCA.

In the presence of both reducing agents, citrate and DC<sup>2-</sup>, at 80 °C, the seed particle formation rate is faster in comparison with citrate-only but slower than with DC<sup>2-</sup>-only (Fig. 1b). This is reflected on the resulting Au NPs size with sizes of 10.7 ± 1.4 nm, 15.2 ± 4.2 and 13.9 ± 3.0 nm obtained with only-DC<sup>2-</sup>, only-citrate and a DC<sup>2-</sup>-citrate mixture (keeping the individual concentration constant) respectively. A possible explanation is the competitive substitution of chloride ligands in the gold precursor by both DC<sup>2-</sup> and citrate, which follows the substitution of chloride ligands in the gold precursor by hydroxycarboxylates proposed by Ojea-Jimenez *et al.* which was later extended to a range of α-hydroxycarboxylate ions with car-

boxyl and hydroxyl groups by Bartosewicz *et al.*<sup>14,27,45</sup> According to Grasseschi *et al.*,<sup>12</sup> DC<sup>2-</sup> can isomerize to form a carbon-carbon double bond and a hydroxyl group (enol form)

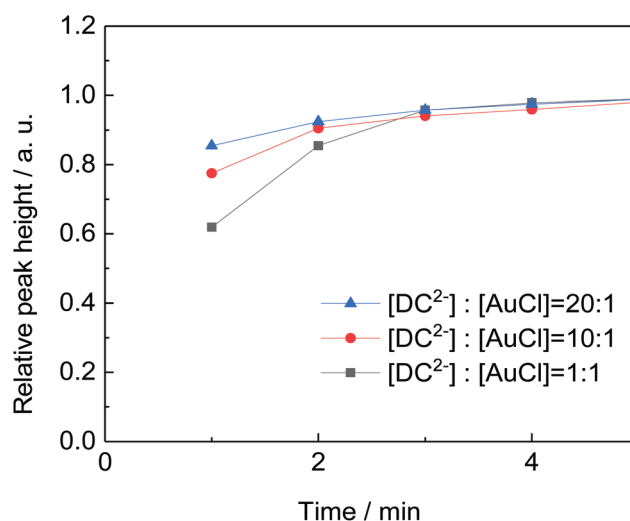


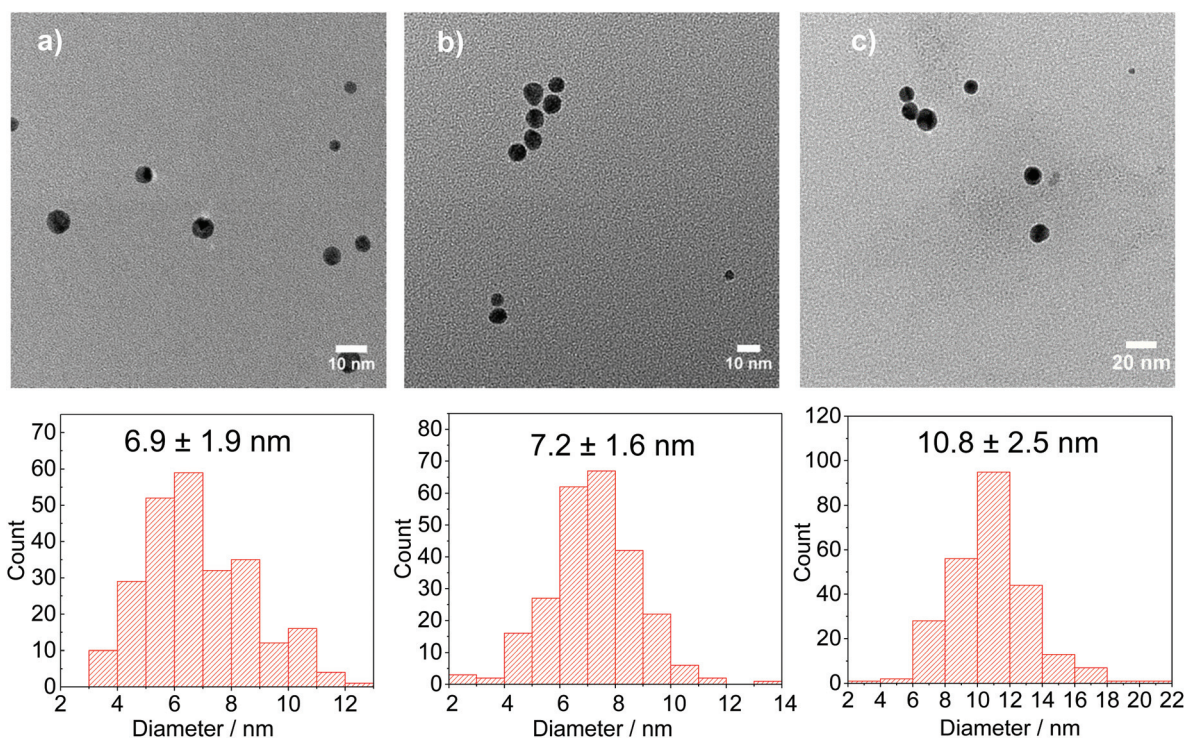
Fig. 7 Comparison of the rate of formation of gold nanoparticles (shown by the height of the absorption peak) by varying the [DC<sup>2-</sup>]:[AuCl] molar ratio at 80 °C. Conditions: [AuCl]: 0.25 mM, [citrate]: 5 mM, pH: 4.4, 80 °C.

which would be expected to have a similar ligand exchange capability although further mechanistic studies would be needed to verify this. Similar results are obtained at 25 °C; the nucleation rate is faster when only  $\text{DC}^{2-}$  is used as reducing agent, while the combination of citrate and  $\text{DC}^{2-}$  gives a faster nucleation than citrate only (Fig. 1a).

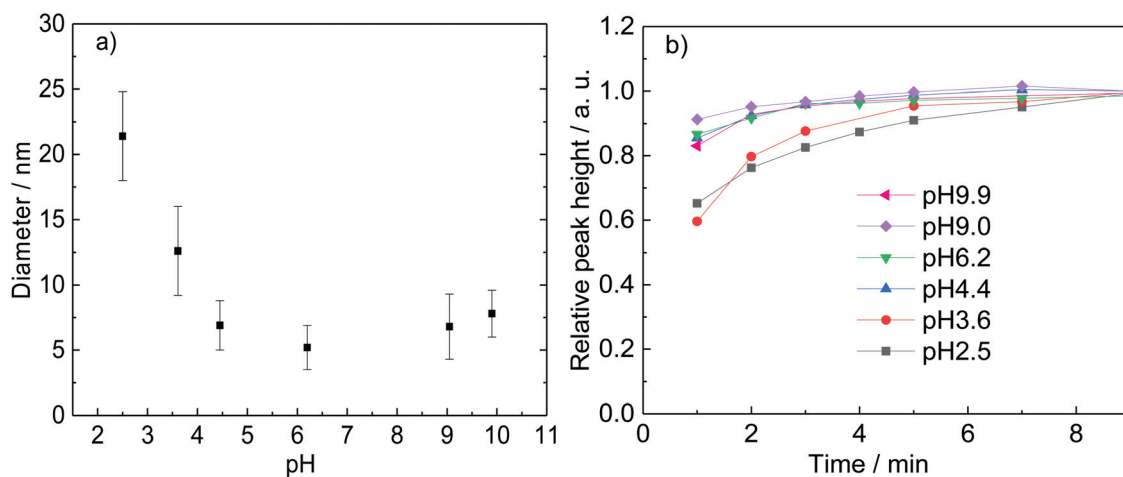
According to the accepted mechanism of the Turkevich method, the second reduction step ( $\text{Au}^+$  to  $\text{Au}^0$ ) and consequently the formation of NPs takes place through the disproportionation reaction (reaction (2)). To investigate the role of the reducing agents and specially the  $\text{DC}^{2-}$  formed as an intermediate species in this second reduction step, a similar series of reactions is carried out but in this case, intermediate  $\text{Au}^+$  species ( $\text{AuCl}$ ) is used as gold precursor rather than  $\text{Au}^{3+}$  present in chloroauric acid ( $\text{HAuCl}_4$ ).

Both citrate and  $\text{DC}^{2-}$  are capable of reducing  $\text{AuCl}$  to form Au NPs at 80 °C. When  $\text{AuCl}$  is the gold precursor,  $\text{DC}^{2-}$  shows a faster nucleation kinetics, (Fig. 4a), and thus, smaller Au

portation reaction (reaction (2)). To investigate the role of the reducing agents and specially the  $\text{DC}^{2-}$  formed as an intermediate species in this second reduction step, a similar series of reactions is carried out but in this case, intermediate  $\text{Au}^+$  species ( $\text{AuCl}$ ) is used as gold precursor rather than  $\text{Au}^{3+}$  present in chloroauric acid ( $\text{HAuCl}_4$ ).



**Fig. 8** Representative TEM images and particle size distribution of Au NPs synthesised using  $\text{AuCl}$  as precursor with a  $\text{DC}^{2-}$  : Au molar ratio of a. 20 : 1, b. 10 : 1 and c. 1 : 1. Conditions:  $[\text{AuCl}]$ : 0.25 mM,  $[\text{citrate}]$ : 5 mM, pH: 4.4, 80 °C, time: 7 min–11 min.



**Fig. 9** Effect of pH on a. Au NP sizes (the error bars represent the standard deviation) and b. rate of formation. Conditions:  $[\text{AuCl}]$ : 0.25 mM,  $[\text{DC}^{2-}]$ : 5 mM,  $[\text{citrate}]$ : 5 mM, 80 °C, time: 7 min–13 min.



NPs ( $6.8 \pm 1.5$  nm, Fig. 5c), compared with sodium citrate ( $14.8 \pm 4.3$  nm, Fig. 5b), at the same pH.

Similar results to the ones above are obtained when  $\text{HAuCl}_4$  is used as gold precursor. Indeed, same kinetics and similar Au NPs size distribution are observed when starting with  $\text{HAuCl}_4$  ( $15.2 \pm 4.2$  nm) or  $\text{AuCl}$  ( $14.8 \pm 4.3$  nm) using sodium citrate as reducing agent (Fig. 4a). Considering that the reduction of gold precursor consists of two reduction steps in series  $\text{Au}^{3+} \rightarrow \text{Au}^+ \rightarrow \text{Au}^0$  as demonstrated above, the similar rate of reaction shown in Fig. 4a indicate that under these conditions and at  $80^\circ\text{C}$ , the  $\text{Au}^+$  reduction is considerably slower than the  $\text{Au}^{3+}$  reduction to  $\text{Au}^+$  and the dissolution of  $\text{AuCl}$ .<sup>15</sup>

The solubility of  $\text{AuCl}$  in water is very low, however,  $\text{DC}^{2-}$  facilitates its dissolution by coordination,<sup>13,45</sup> which consequently facilitates its reduction to  $\text{Au}^0$  and the formation of Au NPs.  $\text{DC}^{2-}$  drives the reduction of  $\text{Au}^+$ , so indirectly further dissolves the  $\text{AuCl}$ .

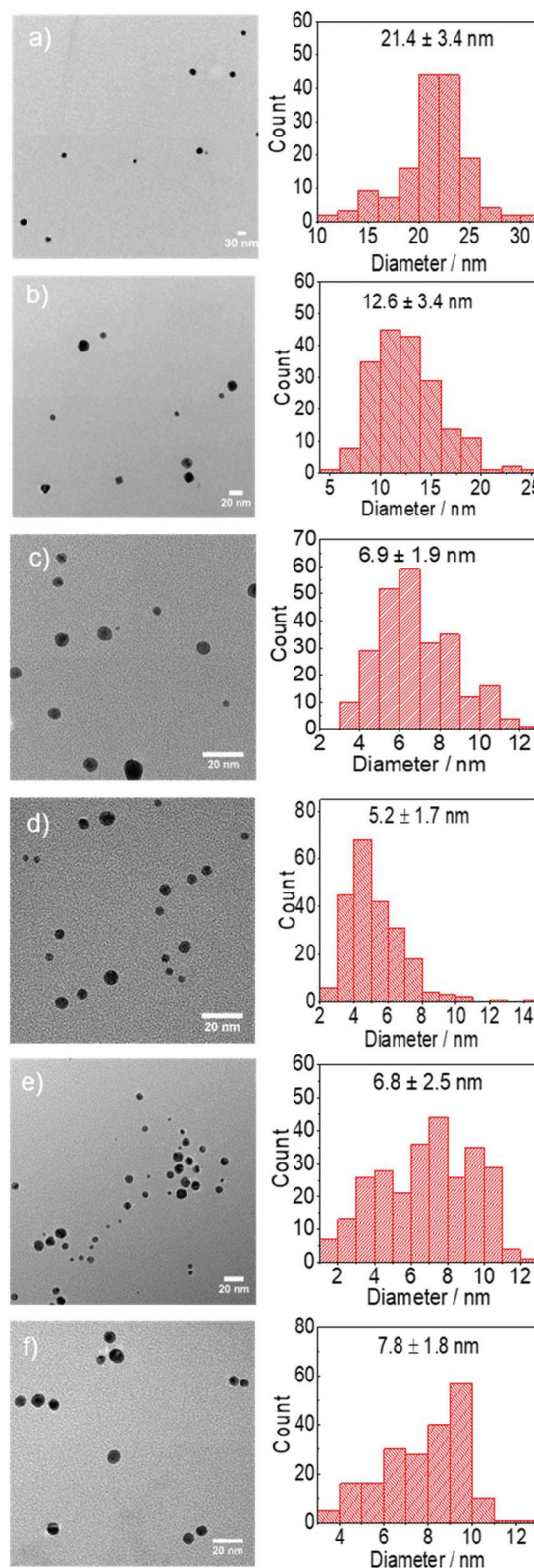
As mentioned above,  $\text{DC}^{2-}$  is not stable at high temperatures and it decomposes quickly at  $80^\circ\text{C}$  (according to thermal stability studies followed by the characteristic absorption peak of DCA at 240 nm, Fig. S2†). However, as shown in Fig. 4, the reduction of  $\text{AuCl}$  is quicker than the decomposition of  $\text{DC}^{2-}$ , successfully leading to the formation of Au NPs. In addition, as discussed before, the coordination of  $\text{DC}^{2-}$  to Au NPs enhances its stability.<sup>12</sup>

Despite the low solubility of  $\text{AuCl}$ , a disproportionation reaction starting with  $\text{Au}^+$  is carried out (*i.e.* in the absence of  $\text{DC}^{2-}$  and citrate) at  $80^\circ\text{C}$  and pH value of 4.8, using PVP as the stabilizer for Au NPs.<sup>37</sup> Au NPs with average sizes of  $11.9 \pm 4.2$  nm are synthesized, as shown in Fig. 5a after 150 min. It is important to note that the overall kinetics of the disproportionation reaction and the potential reduction of  $\text{AuCl}$  by PVP (with a weak reduction potential<sup>46</sup>) are considerably slower than the reduction of  $\text{AuCl}$  by both sodium citrate and  $\text{DC}^{2-}$  (Fig. 4b). These results suggest that in the presence of a reducing agent (sodium citrate or  $\text{DC}^{2-}$ ), the formation of the Au NPs consists of two reduction steps ( $\text{Au}^{3+} \rightarrow \text{Au}^+ \rightarrow \text{Au}^0$ ), the second one competing over the much slower disproportionation reaction.

Indeed, our XPS results (Fig. 6) confirm that the extent of the disproportionation reaction in the presence of  $\text{DC}^{2-}$  is almost negligible. When  $\text{HAuCl}_4$  is reduced by  $\text{DC}^{2-}$ ,  $\text{Au}^{3+}$ ,  $\text{Au}^+$  and  $\text{Au}^0$  are all detected (Fig. 6a). This is consistent with the XPS results of the reduction of  $\text{HAuCl}_4$  by sodium citrate.<sup>40</sup> However, when  $\text{AuCl}$  is reduced by  $\text{DC}^{2-}$  (Fig. 6b), only  $\text{Au}^+$  and  $\text{Au}^0$  species are present, confirming the direct reduction of  $\text{Au}^+$  to  $\text{Au}^0$ , with the disproportionation reaction negligible (*i.e.* absence of  $\text{Au}^{3+}$ ).

Once demonstrated that the second reduction step in the Turkevich method is dominated by the reduction of the  $\text{Au}^+$  by the reducing agent, further understanding of this step is needed to find out the relationship between reaction conditions and particle size and distribution.

During the conventional Turkevich method, the second reduction step takes place in the presence of both, citrate (present as reactant) and  $\text{DC}^{2-}$  (present as intermediate product in the first reduction step, reaction (1)). Thus, a new



**Fig. 10** Representative TEM images and particle size distribution of Au NPs synthesised using  $\text{AuCl}$  as precursor at pH of a. 2.5, b. 3.6, c. 4.4, d. 6.2, e. 9.0 and f. 9.9. Conditions:  $[\text{AuCl}]$ : 0.25 mM,  $[\text{DC}^{2-}]$ : 5 mM, [citrate]: 5 mM,  $80^\circ\text{C}$ , time: 7 min–13 min.



with a minimum (~0.5%) of Cl. Such precipitates are not observed when H<sub>2</sub>AuCl<sub>4</sub> is used as precursor suggesting that due to the low solubility of AuCl, its solid form serves as nucleation points during the reduction process, leading to such precipitation.

## 4. Conclusions

This study challenges the traditional Turkevich mechanism for the synthesis of gold nanoparticles using citrate as reducing agent demonstrating that it consists of two consecutive reduction steps rather than a reduction followed by the disproportionation reaction as commonly believed in the literature. DC<sup>2-</sup>, formed as intermediate product through the oxidation of citrate, plays a key role in the synthesis as it has a considerably higher reduction potential than citrate and enables the electrostatic stabilization of the Au nanoparticles through coordination of its enol form. As a consequence, high nucleation rates can be achieved in the presence of DC<sup>2-</sup> leading to monodispersed particles. Directly related, while previous research considered the first reduction (Au<sup>3+</sup> → Au<sup>+</sup>) as the rate-determining step to control the size of Au nanoparticles, in this paper, the second reduction reaction (Au<sup>+</sup> → Au<sup>0</sup>) is inferred as the rate-limiting step. In this way, the size of Au NPs can be tuned from 5.2 ± 1.7 nm to 21.4 ± 3.4 nm by tuning the conditions of the second reduction step.

## Conflicts of interest

There are no conflicts to declare.

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