Cite this: *RSC Adv.*, 2021, 11, 28818

Bimetallic gold and palladium nanoparticles supported on copper oxide nanorods for enhanced H_2O_2 catalytic reduction and sensing†

Simbongile Sicwetsha,^a Omotayo Adeniyi^a and Philani Mashazi^{id}*^{ab}

The emergence of nanoscience and nanotechnology has revitalised research interest in using copper and its derived nanostructures to find exciting and novel applications. In this work, mono- and bimetallic gold and palladium nanoparticles supported on copper oxide nanorods (CuONRs) were prepared and their catalytic performance towards the reduction of H_2O_2 to form reactive oxygen radical species (ROS) was evaluated. The characterisation using microscopy and spectroscopic techniques confirms the successful synthesis of CuONRs, CuONRs@Au₆NPs, CuONRs@Pd₆NPs and CuONRs@Au₃Pd₃NPs. The efficient generation of ROS was confirmed using UV-vis spectroscopy and 1,3-diphenylisobenzofuran (DPBF) as a radical scavenger. The CuONRs possess excellent catalytic reduction activity for H_2O_2 by generating ROS. However, CuONRs also have lattice oxygens which do not participate in the catalytic reduction step. The lattice oxygens however allowed for the adsorption of gold and palladium nanoparticles (Au₆NPs, Pd₆NPs and Au₃Pd₃NPs) and thus enhanced catalytic reduction of H_2O_2 to produce ROS. The produced ROS was subsequently involved in the catalytic oxidation of a chromogenic substrate (TMB), resulting in blue coloured diimine (TMBDI) complex which was monitored using UV-vis and could also be observed using the naked eye. The catalyst dependence on pH, temperature, and H_2O_2 concentration towards efficient ROS generation was investigated. The gold and palladium-supported CuONRs nanocatalysts were evaluated for their potential applications in the fabrication of colorimetric biosensors to detect glucose oxidation by glucose oxidase (GOx). Glucose was used as a model analyte. The enzymatic reaction between GOx and β -D-glucose produces H_2O_2 as a by-product, which is then catalytically converted to ROS by the nanoparticles.

Received 7th July 2021
Accepted 20th August 2021

DOI: 10.1039/d1ra05247k

rsc.li/rsc-advances

Introduction

Bimetallic noble metal nanoparticles have received extensive research attention due to their intriguing synergistic properties and various applications in nanoscience and nanotechnology. They have been found to possess wide applications and of interest to this work is the applications as nanocatalysts in heterogeneous catalysis.^{1,2} In addition to their unique catalytic properties, nanocatalysts have found uses in various fields, including electronics, electrocatalysis, photocatalysis, and over the past two decades in nanozymology and biosensing. Integrating noble metal nanoparticles and inexpensive transition metal oxide nanoparticles into a single nanostructure produces cheaper and highly stable nanocatalysts with synergistic

properties.³ The morphology, internal structure, and catalytic efficiency of metal nanoparticles supported onto metal oxide nanoparticles (NM-MOs) can be achieved by altering the molar ratios of the precursors and the method used for preparation.^{4–6} The composition, structure, particle size and other auxiliary variables dictate the applications of nanostructured catalysts.^{7–9} Noble metal nanoparticles supported onto metal oxide nanoparticles exhibit better catalytic performance than their mono-metallic analogues.^{10–12} The improved catalytic performance can be ascribed to the synergistic effect induced by altering the electronic structure between the support material (metal oxide) and interfacial noble metals.^{10,13} The size of the noble metal nanoparticles and the nature of the metal oxide support was found to be essential in determining the catalytic performance.^{14,15} One dimensional (1D) nanomaterials such as nanowires, nanotubes and nanorods have attracted significant research attention as supports for noble metal nanocatalysts. They have high surface-to-volume ratio and sometimes offer fast electron transfer process in their synergistic applications.¹⁶ Therefore, small size noble metal nanoparticles such as gold nanoparticles supported on 1D metal oxide nanoparticles would be efficient as nanocatalysts.^{11,15}

^aChemistry Department, Rhodes University, PO Box 94, Makhanda 6140, South Africa. E-mail: p.mashazi@ru.ac.za

^bInstitute for Nanotechnology Innovation, Rhodes University, PO Box 94, Makhanda 6140, South Africa

† Electronic supplementary information (ESI) available: The EDX spectra and SEM-EDS elemental mapping results and discussion of the synthesised gold and palladium-supported nanoparticles. See DOI: 10.1039/d1ra05247k

The evolution of nanozymology has afforded scientists an insight into the enzyme-like properties of heterogeneous catalysts and their ability to mimic natural enzymes, thus their applications as artificial enzymes.^{17,18} Natural enzymes possess exceptional catalytic efficiency, substrate specificity, and high selectivity, resulting in their applications in biosensor developments.¹⁹ However, natural enzymes have some limitations that impede their wide industrial applications. These include low stability in harsh conditions, relatively high cost for preparation, purification and storage, narrow pH windows for optimal operation, and short shelf life.^{17,19} Artificial enzymes show superiority to natural enzymes due to low cost, high temperature and long-time storage stability.¹⁹ Therefore, artificial enzymes have been developed as potential substitutes for natural enzymes. Iron oxide nanoparticles were the first inorganic nanomaterials to display enzyme-like (peroxidase) activity.^{20,21} Various noble metals such as gold nanoparticles (AuNPs), platinum nanoparticles (PtNPs) and palladium nanoparticles (PdNPs), and metal oxide nanoparticles such as CeO₂, LaNiO₃ and V₂O₅ (ref. 22) have also demonstrated the various enzyme-like activity.²³ Therefore, there is a continued need to find nanomaterials with enhanced enzyme-like activity and investigate their mechanism of action.

In this work, we report the synthesis of CuONRs as support for noble metal nanocatalyst (Au₆NPs, Pd₆NPs and bimetallic Au₃Pd₃NPs) and evaluated their peroxidase-like activity. The CuONRs possess excellent catalytic reduction properties towards H₂O₂ generating ROS. However, CuONRs also has lattice oxygen which do not participate in the catalytic reduction. The lattice oxygens are suitable for the adsorption and deposition noble metal nanocatalysts of gold and palladium such as Au₆NPs, Pd₆NPs and Au₃Pd₃NPs. Hence enhancing the catalytic reduction of CuONRs towards H₂O₂ to produce ROS. The subscript number on the noble metal (Au₆NPs, Pd₆NPs and Au₃Pd₃NPs) nanocatalysts refers to the mole ratio of the gold or palladium salt to 20 mg of CuONRs used. The preparation and characterization of CuONRs using chemical reduction method and deposition of nanocatalysts is to our knowledge reported here for the first time. The synthesized CuONRs and the adsorbed or deposited noble metal nanocatalysts were evaluated for their H₂O₂ catalytic reduction and production of reactive oxygen radical species (ROS). Owing to the superior catalytic activity of small size Au₆NPs, Pd₆NPs and the bimetallic Au₃-Pd₃NPs and the catalytic activity of CuONRs, the CuONRs@Au₆NPs, CuONRs@Pd₆NPs, and CuONRs@Au₃Pd₃-NPs could possess synergistic catalytic properties. The facile synthesis and low cost of CuONRs make them suitable catalyst support. Also, the novelty of this work is not limited to the synthesis of the nanoparticles alone. The evaluation of their peroxidase mimetic activity and application in glucose detection is also reported.

Experimental

Reagents and apparatus

Copper acetate Cu(Ac)₂, hydrogen tetrachloroaurate trihydrate (HAuCl₄·3H₂O, 99%), palladium chloride (H₂PdCl₄), trisodium

citrate dihydrate (Na₃cit·2H₂O), sodium borohydride (NaBH₄), 3,3',5,5'-tetramethylbenzidine (TMB), glacial acetic acid (CH₃-COOH), 1,3-diphenylisobenzofuran (DPBF), D-glucose powder, glucose oxidase (GOx, EC 1.1.3.4. from *Aspergillus niger*, Type VII) were purchased from Sigma-Aldrich. Potassium hydroxide pellets (KOH), sodium acetate anhydrous (NaAc), and potassium dihydrogen orthophosphate (KH₂PO₄) were purchased from Merck. Sodium hydroxide pellets (NaOH), absolute ethanol (EtOH), methanol (MeOH), 50% hydrogen peroxide (H₂O₂), 32% hydrochloric acid (HCl) and dimethyl sulfoxide (DMSO) were purchased from B&M Scientific. All the reagents were of analytical grade and were used as received from the suppliers. Ultrapure water with the resistivity of 18 MΩ cm obtained from a Milli-Q water system (Millipore Corp, Bedford, MA, USA) was used to prepare all aqueous solutions.

UV-visible measurements were conducted on a Thermo-Scientific, Multiskan Sky w Cuvette & Touch Screen, 100–240 V, Belgium. Zeta potential measurements were carried out on a Malvern Zetasizer Nano-ZS90 series equipped with a 633 nm He/Ne laser. A disposable folded capillary cell (DTS 1060) was used for data collection. TEM images were taken from a Zeiss Libra 120 TEM operating at 80 kV. The nanoparticles were dissolved in water and dropped onto a carbon-coated copper grid, and allowed to dry at room temperature overnight before data collection. The EDS spectra were obtained from INCA PENTA FET coupled with VGA TASCAM at 20 kV acceleration voltage. X-ray powder diffraction (XRD) patterns were recorded on a Bruker D8 Discover equipped with a Lynx Eye detector, using Cu-Kα radiation (1.5405 Å, nickel filter). The samples were placed on a silicon wafer slide. The data was collected within the 2θ range of 10° to 100°.

Preparation of mono and bimetallic nanoparticles supported on CuONRs

Preparation of copper oxide nanorods (CuONRs). The copper oxide nanorods were prepared following a reported sol-gel method²⁴ with slight modifications. Briefly, 75 ml of 80 mM (6.0 mmol) copper acetate solution was blended with 1 ml glacial acetic acid in a round-bottom flask. The reaction was heated to reflux with continuous stirring. A 10 ml solution of sodium hydroxide (6 M, 60 mmol) was quickly injected into the solution. A black precipitate formed immediately. The reaction was allowed to stir for 30 minutes. The precipitate was centrifuged, washed three times with ethanol and air-dried at room temperature to yield copper oxide nanorods, CuONRs.

Preparation of AuNPs supported on CuONRs, CuONRs@Au₆NPs. The preparation of CuONRs@Au₆NPs was achieved using a method that has been previously reported.²⁰ Briefly, 20 mg of copper oxide nanorods (CuONRs) was dispersed in 60 ml of Millipore water and sonicated for 30 minutes. Then 10 ml of HAuCl₄·3H₂O (0.60 mM, 6.0 mmol) and 1.5 ml of trisodium citrate dihydrate (3.0 mg, 0.010 mmol) were added to the CuONRs. After 15 minutes, 1.5 ml of sodium borohydride (3.0 mg, 0.080 mmol) was added to the solution. The colour changed immediately from dark brown to black. The reaction was stirred vigorously for another 30 minutes at room

temperature. The black precipitate was centrifuged, washed three times with ethanol and air-dried at room temperature to yield gold nanoparticles-supported copper oxide nanorods, CuONRs@Au₆NPs. The subscript number 6 refers to 6.0 mmol of HAuCl₄·3H₂O used.

Preparation of PdNPs supported on CuONRs, CuONRs@Pd₆NPs. A similar procedure for CuONRs@Au₆NPs was used to prepare CuONRs@Pd₆NPs. Instead of HAuCl₄·3H₂O, H₂PdCl₄ salt was used. Briefly, 20 mg of copper oxide nanorods (CuONRs) was dispersed in 60 ml of Millipore water and sonicated for 30 minutes, and into this solution, 10 ml of H₂PdCl₄ (0.60 mM, 6.0 mmol) was added. About 1.5 ml of trisodium citrate dihydrate (3.0 mg, 0.010 mmol) and after 15 minutes, 1.5 ml of sodium borohydride (3.0 mg, 0.080 mmol) was added all at once. The colour changed immediately from dark brown to black. The reaction was stirred vigorously for another 30 minutes at room temperature. The black precipitate was centrifuged, washed three times with ethanol and air-dried at room temperature to yield palladium nanoparticles-supported CuONRs, CuONRs@Pd₆NPs. The subscript number 6 refers to 6.0 mmol of H₂PdCl₄ used.

Preparation of AuPdNPs supported on CuONRs, CuONRs@Au₃Pd₃NPs. The preparation of CuONRs@Au₃Pd₃NPs was accomplished by adding 20 mg of copper oxide nanorods (CuONRs) in 60 ml of water and sonicated for 30 minutes. Into this solution, HAuCl₄·3H₂O (5 ml, 3.0 mmol) and H₂PdCl₄ (5 ml, 3.0 mmol) aqueous solutions were added simultaneously at room temperature. Trisodium citrate dihydrate (3.0 mg, 0.010 mmol) was added to the mixture. After 15 minutes, sodium borohydride (3.0 mg, 0.080 mmol) was added rapidly into the solution. The colour changed immediately from dark brown to black. The reaction was stirred vigorously for 30 minutes at room temperature. The precipitate was centrifuged, washed three times with ethanol and air-dried at room temperature to yield bimetallic gold–palladium-supported copper oxide nanorods, CuONRs@Au₃Pd₃NPs.

Monitoring catalytic reduction of H₂O₂

The catalytic reduction of H₂O₂ by gold and palladium mono- and bimetallic nanoparticles supported on CuONRs was conducted at room temperature using 3,3',5,5'-tetramethyl benzidine (TMB) using UV-vis spectrophotometer. H₂O₂ (50 μL, 0.10 mol L⁻¹), TMB (50 μL, 25 mmol L⁻¹), and nanoparticle solution (50 μL of 2.0 mg mL⁻¹) were added to 0.20 mol L⁻¹ acetate buffer solution at the pH where IEP of the nanoparticles was zero. The blue colour developed with time, and the UV-vis spectra were measured.

In addition to the TMB, ROS generation was further monitored using 1,3-diphenyliso-benzofuran (DPBF). H₂O₂ (75 μL, 0.10 mmol L⁻¹) was mixed with nanoparticles (50 μL, 2.0 mg mL⁻¹). DPBF (50 μL, 0.25 mmol L⁻¹) was added, and immediately the UV-vis spectra were measured. The absorption band at 430 nm was monitored. DPBF reacts with the produced ROS resulting in the oxidation of DPBF and a decrease in its signal intensity at 430 nm. The absorption spectra at 430 nm were obtained at a predetermined time interval.

Detection of glucose at gold and palladium nanocatalysts supported on CuONRs

The ROS generation and optimal properties were evaluated towards the detection of glucose. The enzymatic reaction between glucose and glucose oxidase (GOx) to yield gluconolactone and H₂O₂ (by-product) was used. GOx (50 μL, 1.0 mg mL⁻¹) was added into a 75 μL phosphate buffer saline (pH 7.4) solution of varied concentrations of glucose and allowed to react at 35 °C for 45 minutes. This results in the oxidation of glucose to produce gluconolactone and H₂O₂. After 45 minutes, 50 μL solution (2.0 mg mL⁻¹) of gold and palladium-supported nanorods was added. This was followed by the addition of TMB (100 μL, 2.0 mmol L⁻¹) in acetate buffer (1.0 ml, 0.20 mol L⁻¹) at optimum pH. The temperature of the solution was kept at 35 °C. The colour development was monitored by the naked eye (blue colour development) and spectrophotometrically using UV-vis.

Results and discussion

Characterisation of gold- and palladium-supported CuONRs

CuONRs nanoparticles were prepared using the chemical reduction method.²⁰ The conventional sol-gel method was optimised to yield CuO nanorods (CuONRs). The rod-like morphology of CuONRs provided a suitable support surface for anchoring gold and palladium nanoparticles. A seed-mediated method was used to deposit gold and palladium nanoparticles and played an essential role in the formation CuONRs@Au₆NPs, CuONRs@Pd₆NPs and CuONRs@Au₃Pd₃NPs. The preparation of gold and palladium nanoparticles-supported on CuONRs was conducted in three separate reaction vessels.

Into each reaction vessel, CuONRs was dispersed in solution, and noble metal salt HAuCl₄·3H₂O or H₂PdCl₄ or a mixture of both was added. After 15 minutes of homogeneously mixing the reaction, a strong reducing agent NaBH₄ was added. Trisodium citrate was added as the stabilising agent. The concentrations of metal salts and NaBH₄ were such that small noble metal nanoparticles were deposited onto CuONRs. The reaction was carried out in an ice-water bath. The low temperatures were used to facilitate the formation of smaller size gold and palladium nanoparticles onto the CuONRs surface. For metal salts, 0.60 mmol L⁻¹ concentration was used, and for mixed metal salts, a concentration of 0.30 mmol L⁻¹ for each metal salt was used. The formation of CuONRs@Au₆NPs and CuONRs@Pd₆NPs occurred and represented monometallic supported nanoparticles. Bimetallic gold–palladium nanoparticles involved simultaneous co-reduction of HAuCl₄·3H₂O (0.30 mM) and H₂PdCl₄ (0.30 mM) to form CuONRs@Au₃Pd₃NPs. The morphology and structural properties were studied to understand the synergistic effect between metallic systems.²⁵

X-ray diffraction (XRD) was used to analyse the successful formation of CuONRs and mono and bimetallic gold and palladium nanoparticles supported on CuONRs. Fig. 1 shows the (a) X-ray diffraction patterns and (b) zeta-potentials of (i) CuONRs, (ii) CuONRs@Au₆NPs, (iii) CuONRs@Pd₆NPs, and (iv) CuONRs@Au₃Pd₃NPs. The X-ray diffractogram for CuONRs in

Fig. 1 (a) X-ray diffraction (XRD) patterns and (b) zeta potential vs. pH of (i) CuONRs, (ii) CuONRs@Au₆NPs, (iii) CuONRs@Pd₆NPs, and (iv) CuONRs@Au₃Pd₃NPs.

Fig. 1(a)(i) exhibited peaks at $2\theta = 32.6^\circ, 35.6^\circ, 38.8^\circ, 48.0^\circ, 53.3^\circ, 57.9^\circ, 61.5^\circ, 65.9^\circ, 67.5^\circ, 71.9^\circ, 74.5^\circ$ corresponding to (110), (-111), (111), (-202), (020), (202), (-113), (022), (311), (004) Miller indices. The X-ray diffraction patterns showed single phase copper oxide nanoparticles with a monoclinic structure with lattice constants, $a = 4.67970$, $b = 3.43140$, $c = 5.13620$, $\alpha = 90.00$, $\beta = 99.26$ and $\gamma = 90.00$. The peaks were broad in shape, confirming the formation of the nanoparticles. The CuONRs@Au₆NPs in Fig. 1(a)(ii) showed the diffraction patterns due to CuONRs as observed above and additional peaks at $2\theta = 37.4^\circ, 44.4^\circ, 64.5^\circ, 77.7^\circ$ corresponding to (111), (200), (220), (311) Miller indices of gold nanoparticles similar to a face-centred cubic crystal structure of gold (JCPDS: 65-8601). The CuONRs@Pd₆NPs in Fig. 1(a)(iii) showed the diffraction patterns due to CuONRs and PdNPs at $2\theta = 40.1^\circ, 46.3^\circ, 68.5^\circ, 82.1^\circ, 86.0^\circ$ corresponding to (111), (200), (220), (311), (222) Miller indices of PdNPs similar to the structure of palladium (JCPDS: 05-0681). The CuONRs@Au₃Pd₃NPs in Fig. 1(a)(iv) showed the mixture of diffraction patterns of CuONRs, CuONRs@Au₆NPs and CuONRs@Pd₆NPs. There were also few additional peaks at $2\theta = 34.1, 38.2, 63.8$ and 82.7° , corresponding to (111), (200), (220), and (311) for the nanoalloy of Au@PdNPs. The distinct diffraction patterns emanating from CuONRs@Au₃Pd₃NPs confirmed co-reduction of $\text{HAuCl}_4 \cdot 3\text{H}_2\text{O}$ and H_2PdCl_4 salts occurred, and the formation of Au@PdNPs nanoalloy formed successful.⁴ Fig. 1(b) shows the zeta potential (mV) plot of the synthesised nanomaterials at different pH conditions. The zeta potential graph showed similar trends with the zeta potential values positive at pH 2 for CuONRs and CuONRs@Au₆NPs. For CuONRs@Pd₆NPs and CuONRs@Au₃Pd₃NPs, the values were positive at pH 2 and pH 3. The zeta potential values were negative and continued to increase negatively as the pH increased. This is due to hydroxide (OH^-) from the sodium hydroxide salt used to prepare CuONRs. In addition to the OH^- , also carboxylic acid and hydride surface functional groups due to adsorbed species which act as stabilisers of gold and palladium nanoparticles. The negative increase is due to the deprotonation of the surface carboxyl

groups as the pH increases. The zeta potential values increased to -29.1 mV for CuONRs, -28.3 mV for CuONRs@Au₆NPs, -27.4 mV for CuONRs@Pd₆NPs and -25.1 mV for CuONRs@Au₃Pd₃NPs at pH 10. The negative zeta potentials for CuONRs@Au₆NPs, CuONRs@Pd₆NPs and CuONRs@Au₃Pd₃NPs were ascribed to the citrate stabiliser used to minimise aggregation of Au and Pd nanoparticles on the surface of CuONRs. The negative zeta potential was also due to the exposed CuONRs surface not covered by Au and Pd nanoparticles. The CuONRs exhibited the isoelectric point (IEP) or point-of-zero charge (PZC) at pH 2.5. The IEP was slightly shifted for CuONRs@Au₆NPs to pH 3.2, CuONRs@Pd₆NPs to pH 3.3 and CuONRs@Au₃Pd₃NPs to pH 2.7. High positive ($>+20$ mV) or negative (>-20 mV) zeta potential values are related to the stability of the nanoparticles at that pH as these induce particle-particle separation due to charge repulsion.²⁶ The zeta potential values more than -20 mV for stable nanoparticles were at pH ≥ 5 for CuONRs, pH ≥ 6 for CuONRs@Au₆NPs, pH ≥ 6 for CuONRs@Pd₆NPs and pH ≥ 7 for CuONRs@Au₃Pd₃NPs. The negative zeta potential shows that the mono and bimetallic nanoparticles supported on CuONRs are negatively charged and stable at physiological pH.

Fig. 2 shows TEM micrographs with their corresponding size distribution histograms for (a) CuONRs, (b) CuONRs@Au₆NPs, (c) CuONRs@Pd₆NPs, and (d) CuONRs@Au₃Pd₃NPs. The TEM image of CuONRs in Fig. 2(a) exhibited rod-like nanostructures with the length of 40 ± 5.6 nm and the width (ϕ) of 11 ± 4.5 nm. The CuONRs TEM image also showed some aggregation, which is typical of copper oxide nanoparticles.²⁴ The bigger-sized nanoparticles were due to the overlapping CuONRs, and the longer CuONRs were head-to-tail or head-to-head or tail-to-tail connected. The CuONRs that overlapped or head-to-tail or head-to-head or tail-to-tail connected were not included in measuring the length (40 ± 5.6 nm) and width (11 ± 4.5 nm). The TEM image of CuONRs@Au₆NPs in Fig. 2(b) showed that the rod-like morphology of CuONRs was retained, but these were clustered. Spherical gold nanoparticles were observed as dark spots deposited onto CuONRs. The deposited AuNPs onto CuONRs resulted in the rough surfaces of CuONRs. The average particle size of AuNPs on the surface of CuONRs was 2.8 ± 0.8 nm. The TEM image of CuONRs@Pd₆NPs exhibited spherical palladium nanoparticles as deposits on the surface of CuONRs. The average size of palladium nanoparticles was 2.9 ± 0.8 nm. The gold and palladium nanoparticles gave similar particle sizes, and this was due to similar concentrations (0.60 mmol L^{-1}) of $\text{HAuCl}_4 \cdot 3\text{H}_2\text{O}$ and H_2PdCl_4 used and similar reaction conditions. The average size of AuPd bimetallic nanoparticles on the surface of CuONRs was 2.5 ± 0.6 nm. The relatively smaller size nanoparticles of AuPdNPs was due to the concentration (0.30 mmol L^{-1}) used for each metal salt. The total metal salt content was 0.60 mmol L^{-1} and equivalent to the CuONRs@Au₆NPs and CuONRs@Pd₆NPs. The SEM-EDS was used to characterise the nanoparticles, and the results are shown in Fig. S2 to S4 of the (ESI†). The SEM-EDS elemental mapping showed the homogeneous and uniform distribution of copper, oxygen, gold and palladium in the prepared nanoparticles.

Fig. 2 TEM micrographs with their corresponding size distribution histograms and their corresponding for (a) CuONRs, (b) CuONRs@Au₆NPs, (c) CuONRs@Pd₆NPs, and (d) CuONRs@Au₃Pd₃NPs.

Catalytic reduction of H₂O₂ at Au₆NPs-, Pd₆NPs- and Au₃Pd₃NPs-supported on CuONRs

Gold and palladium nanoparticles supported on CuONRs were investigated for their catalytic reduction of H₂O₂ to form ROS (HO[•], HO₂[•] and O₂[•]). The reaction was monitored following the oxidation of a chromogenic substrate, 3,3',5,5'-tetramethylbenzidine (TMB). TMB is a colourless chromogenic substrate. Upon oxidation by ROS, it changes colour to blue, resulting in a 3,3',5,5'-tetramethylbenzidinediimine (TMBDI) charge transfer complex with a maximum absorption bands at 652 nm and 370 nm. Fig. 3 shows the UV-vis absorption spectra of (a)(i)

nanomaterials (Au₆NPs-, Pd₆NPs- and Au₃Pd₃NPs-supported on CuONRs) + H₂O₂, (ii) nanomaterials + TMB, and (iii) CuONRs + H₂O₂ + TMB (inset: images of corresponding solutions), and (b)(i) CuONRs@Au₆NPs + H₂O₂ + TMB, (ii) CuONRs@Pd₆NPs + H₂O₂ + TMB, (iii) CuONRs@Au₃Pd₃NPs + H₂O₂ + TMB in 0.20 M acetate buffer solution (at the pH where IEP is zero for different nanoparticles). No colour development and absorption in the UV-vis spectra were observed in Fig. 3(a)(i) and (ii). Fig. 3(a)(iii) shows the UV-vis spectrum with absorption maxima at 370 nm and 652 nm due to the presence of H₂O₂ + TMB + CuONRs. The images of the different solutions in the inset of Fig. 3(a)(iii) showed that blue colour developed. The UV-vis spectra in Fig. 3(b)(i) CuONRs@Au₆NPs, (ii) CuONRs@Pd₆NPs and (iii) CuONRs@Au₃Pd₃NPs showed absorption with maxima at 370 nm and 652 nm. The observed absorption bands at 370 nm and 652 nm confirmed ROS generation from the reduction of H₂O₂ and oxidation of TMB. The oxidation of TMB only occurs when the nanoparticles are in the presence of H₂O₂.

Fig. 3 UV-vis absorption spectra of (a)(i) NRs + H₂O₂, (ii) NRs + TMB, (iii) CuONRs + H₂O₂ + TMB and (b)(i) CuONRs@Au₆NPs + H₂O₂ + TMB, (ii) CuONRs@Pd₆NPs + H₂O₂ + TMB, (iii) CuONRs@Au₃Pd₃NPs + H₂O₂ + TMB in 0.20 M acetate buffer solution.

Optimum conditions for H₂O₂ reduction and ROS generation

The effect of environmental conditions such as pH, H₂O₂ concentration and temperature on the ROS generation efficiency were investigated. Fig. 4 shows the effect of (a) reaction time, (b) pH, (c) varying H₂O₂ concentration, and (d) temperature on ROS generation for (i) CuONRs, (ii) CuONRs@Au₆NPs

(red), (iii) CuONRs@Pd₆NPs (green), and (iv) CuONRs@Au₃-Pd₃NPs (purple). In Fig. 4(a), the effect of reaction time was investigated by increasing the reaction time from 0–18 minutes. The CuONRs@Au₃Pd₃NPs in Fig. 4(a)(iv) showed the highest rate of ROS generation compared to other nanoparticles. CuONRs and CuONRs@Au₆NPs showed the lowest ROS generation in Fig. 4(a)(i) and (ii), respectively. CuONRs@Pd₆NPs in Fig. 4(a)(iii) showed higher ROS generation when compared with CuONRs and CuONRs@Au₆NPs. In Fig. 4(b), the effect of pH was investigated by changing the pH from pH 3–9 of the substrate solution. ROS generation was highest at pH 4.0 for CuONRs@Au₆NPs and CuONRs@Au₃Pd₃NPs, pH 5.0 for CuONRs, and pH 6.0 for CuONRs@Pd₆NPs. The differences in optimum pH conditions were attributed to variation in the composition of the prepared nanomaterials. It could be inferred that CuONRs@Au₆NPs and CuONRs@Au₃Pd₃NPs favoured ROS generation in pH 4.0, whilst CuONRs favoured ROS generation at pH 5.0 with CuONRs@Pd₆NPs favouring ROS generation at pH 6.0. As the pH increased, a decrease in ROS generation was observed due to the instability and decomposition of H₂O₂ in alkaline conditions. In Fig. 4(c), the effect of varying H₂O₂ concentration from 0–60 mM was investigated. The concentration of TMB was kept constant (25 mM). When the H₂O₂ concentration increased from 0–60 mM, ROS generation increased for CuONRs, CuONRs@Au₆NPs, CuONRs@Pd₆NPs and CuONRs@Au₃Pd₃NPs. A linear relationship between the absorbance at 652 nm and TMB or H₂O₂ concentration was established. CuONRs@Au₃Pd₃NPs had the highest slope of 0.0084 a.u. mM⁻¹ followed by CuONRs@Pd₆NPs (0.0080 a.u. mM⁻¹) and CuONRs@Au₆NPs (0.0043 a.u. mM⁻¹) and CuONRs (0.0033 a.u. mM⁻¹). The high slope for CuONRs@Au₃Pd₃NPs was due to the excellent catalysis of the bimetallic system. In Fig. 4(d), the effect of increasing the temperature from 20–70 °C was investigated. The studied nanomaterials showed an increase in ROS generation as the temperature increased up to 40 °C and remained stable afterwards. It was interesting to see

that ROS generation for the nanomaterials was not affected by the temperature up to 70 °C. The stability of the ROS generation even at increased temperatures demonstrates the applicability of the nanomaterials in temperature conditions unfavourable to natural enzymes. In terms of the nanomaterials used, CuONRs@Au₃Pd₃NPs showed higher absorption intensity as the reaction time increased, and this was followed by the CuONRs@Pd₆NPs. This could be attributed to the excellent catalytic properties of PdNPs in CuONRs@Pd₆NPs. The bimetallic nanoparticle system, CuONRs@Au₃Pd₃NPs, containing AuNPs and PdNPs as nanocatalysts resulted in the synergistic effect and higher rate of ROS generation as confirmed by an enhanced increase in absorption at 652 nm.

Steady-state kinetics of Au₆NPs-, Pd₆NPs- and Au₃Pd₃NPs-supported on CuONRs

The steady-state kinetic parameters were evaluated for CuONRs and Au₆NPs-, Pd₆NPs-, and Au₃Pd₃NPs supported on CuONRs. The Michaelis–Menten and Lineweaver–Burk (double reciprocal) models were used to obtain the kinetic parameters derived from varying one substrate concentration and keeping the other substrate concentration constant. H₂O₂ and TMB were used as substrates. The Michaelis–Menten parameters such as Michaelis–Menten constant (K_m) and maximum velocity (V_{max})

Fig. 4 Effect of (a) reaction time, (b) pH, (c) changing H₂O₂ concentration, and (d) temperature on peroxidase-like activity of (i) CuONRs (black), (ii) CuONRs@Au₆NPs (red), (iii) CuONRs@Pd₆NPs (green), and (iv) CuONRs@Au₃Pd₃NPs (purple).

Fig. 5 The steady-state kinetic plots of varying [H₂O₂] at a fixed 25.0 mM TMB concentration and their corresponding double reciprocal plots for (a) CuONRs, (b) CuONRs@Au₆NPs, (c) CuONRs@Pd₆NPs and (d) CuONRs@Au₃Pd₃NPs.

were obtained by fitting the data obtained from the plot of initial velocity (V_0) versus substrate concentration $[S]$ in the Lineweaver–Burk double reciprocal plot. Fig. 5 shows a plot of initial velocity (V_0) versus varied $[H_2O_2]$ at a fixed 25.0 mM of $[TMB]$, and the corresponding double reciprocal plots for (a) CuONRs, (b) CuONRs@Au₆NPs, (c) CuONRs@Pd₆NPs and (d) CuONRs@Au₃Pd₃NPs. The linear relationship between initial velocity (V_0) versus $[H_2O_2]$ was observed at low $[H_2O_2]$ concentrations and a plateau at higher $[H_2O_2]$ concentrations confirming Michaelis–Menten kinetics. The Lineweaver–Burk (double reciprocal plots) were used to calculate V_{max} values from the intercepts and the slopes of the graph for calculating K_m values. The K_m and V_{max} values obtained are summarised in Table 1 for H_2O_2 and TMB as substrates. The K_m value indicates the binding affinity of the nanoparticles towards the substrate.^{27,28} Low K_m values confirm a stronger binding affinity, while higher K_m values signify a weak binding affinity towards a particular substrate. The CuONRs@Pd₆NPs showed a lowest K_m value of 2.94 mM, whilst CuONRs@Au₆NPs showed the highest K_m value of 3.11 mM when H_2O_2 was used as a substrate. The results indicate that the CuONRs@Pd₆NPs had a stronger binding affinity for H_2O_2 . CuONRs@Au₃Pd₃NPs exhibited a high K_m value of 10.68 mM compared to CuONRs@Au₆NPs (3.11 mM) and CuONRs@Pd₆NPs (2.94 mM). However, the K_m value for CuONRs@Au₃Pd₃NPs of 10.68 mM was lower than 39.97 mM of CuONRs, confirming that Au₃-Pd₃NPs had an enhancing effect. The K_m values for CuONRs@Au₆NPs and CuONRs@Pd₆NPs were within the K_m values for the HRP enzyme, ranging between 0.214–3.72 mM. The following trend of increasing K_m values was observed; CuONRs@Pd₆NPs (2.94 mM) < CuONRs@Au₆NPs (3.11 mM) < CuONRs@Au₃Pd₃NPs (10.68 mM) < CuONRs (39.97 mM). The excellent K_m values for CuONRs@Pd₆NPs confirms the observed trend on the enhancement of ROS generation and due to electron transfer from gold to palladium as observed above.

Fig. 6 shows the steady-state kinetic parameters, plot of initial velocity (V_0) versus $[TMB]$, and their corresponding double reciprocal plots for (a) CuONRs, (b) CuONRs@Au₆NPs, (c) CuONRs@Pd₆NPs and (d) CuONRs@Au₃Pd₃NPs. The TMB

concentration was varied from 0–7.0 mM whilst keeping the concentration of H_2O_2 (0.10 M) constant for the different nanomaterials. At lower TMB concentrations (<5.0 mM), the initial velocity (V_0) increased linearly with increasing concentration of TMB for CuONRs in Fig. 6(a), CuONRs@Au₆NPs

Fig. 6 The steady-state kinetic plots of varying TMB concentrations at a fixed 0.10 M H_2O_2 concentration and the corresponding double reciprocal plots for (a) CuONRs, (b) CuONRs@Au₆NPs, (c) CuONRs@Pd₆NPs, and (d) CuONRs@Au₃Pd₃NPs.

Table 1 Steady-state kinetic parameters, K_m and V_{max} for the AuNPs, PdNPs and AuPdNPs-supported CuONRs

Material	Substrate	K_m (mM)	V_{max} ($M s^{-1}$)
CuONRs ^{TW}	$[H_2O_2]$	39.97	4.24×10^{-8}
	$[TMB]$	2.79	1.55×10^{-8}
CuONRs@Au ₆ NPs ^{TW}	$[H_2O_2]$	3.11	1.27×10^{-8}
	$[TMB]$	6.34	2.44×10^{-8}
CuONRs@Pd ₆ NPs ^{TW}	$[H_2O_2]$	2.94	1.23×10^{-9}
	$[TMB]$	3.74	1.65×10^{-8}
CuONRs@Au ₃ Pd ₃ NPs ^{TW}	$[H_2O_2]$	10.68	1.90×10^{-8}
	$[TMB]$	4.55	7.25×10^{-9}
HRP ^{21,26}	$[H_2O_2]$	0.214–3.72	1.24×10^{-8} to 8.71×10^{-8}
	$[TMB]$	0.275–0.434	2.46×10^{-8} to 10.0×10^{-8}
CuO–Au nanoalloy ³⁰	$[H_2O_2]$	4.08	1.11×10^{-10}
	$[TMB]$	3.54	1.05×10^{-10}
Fe ₃ O ₄ @SiO ₂ –NH ₂ –Au@Pd _{0.30} NPs ³¹	$[H_2O_2]$	0.35	6.78×10^{-8}
	$[TMB]$	0.09	8.65×10^{-8}

Fig. 6(b), CuONRs@Pd₆NPs in Fig. 6(c) and CuONRs@Au₃Pd₃-NPs in Fig. 6(d) and reaching the plateau afterwards. The obtained results are representative of the Michaelis-Menten mechanism. The K_m and V_{max} values were calculated and are summarised in Table 1 for both H₂O₂ and TMB as substrates. The CuONRs showed a strong binding affinity towards TMB with K_m equals to 2.79 mM followed by CuONRs@Pd₆NPs with a K_m value of 3.74 mM. The K_m values for CuONRs@Au₆NPs and CuONRs@Au₃Pd₃ NPs were 4.55 mM and 6.34 mM, respectively. The trend of increasing K_m values was CuONRs (2.79 mM) < CuONRs@Pd₆NPs (3.74 mM) < CuONRs@Au₃Pd₃NPs (4.55 mM) < CuONRs@Pd₆NPs (6.34 mM). The maximum velocity (V_{max}) for TMB was 7.25×10^{-9} M s⁻¹ for CuONRs@Au₃Pd₃NPs, whilst that of other materials was 10^{-8} M s⁻¹. The K_m values of TMB for the prepared nanomaterials were higher than for HRP enzyme towards TMB with K_m values 0.275–0.434 mM (ref. 21 and 29) and for Au@Pd_{0.30}NPs supported on magnetic-silica nanoparticles.³⁰ The K_m values were comparable to the reported CuO–Au nanoalloys recently reported.³¹ CuONRs@Pd₆NPs exhibited better kinetic properties, especially towards H₂O₂, which is the substrate that interacts with the nanomaterial and generates ROS that oxidises the TMB substrate.

Confirmation of ROS generation using DPBF

The catalysis of CuONRs, CuONRs@Au₆NPs, CuONRs@Pd₆NPs and CuONRs@Au₃Pd₃NPs was proposed to be *via* the reduction of H₂O₂ to produce ROS (OH[•], HO₂[•] and O₂[•]). The generation of ROS was evaluated using 1,3-diphenylisobenzofuran (DPBF) as a radical scavenger. DPBF is a fluorescent dye that reacts in a specific manner with singlet oxygen (¹O₂), leading to its oxidation, monitored using UV-vis spectroscopy.³² In this work, DPBF was used to monitor the production of ROS and its oxidation. The Au₆NPs-, Pd₆NPs- and Au₃Pd₃NPs supported on CuONRs produced radicals (HO[•], HO₂[•], O₂[•]) upon interacting with H₂O₂. Spectroscopically, the produced radicals were scavenged by DPBF, and a decrease in its absorption peak was observed. The rate of oxidation of DPBF was monitored at 1 or 2 minutes time intervals for the prepared nanoparticles. Fig. 7 shows the rate of oxidation plots for (a)(i) DPBF + H₂O₂, (ii) DPBF + CuONRs and (iii) DPBF + H₂O₂ + CuONRs, (b) CuONRs@Au₆NPs (i) with DPBF alone and (ii) with DPBF + H₂O₂, (c) CuONRs@Pd₆NPs with (i) DPBF alone and (ii) DPBF + H₂O₂ and (d) CuONRs@Au₃Pd₃NPs with (i) DPBF alone and (ii) DPBF + H₂O₂. There was a slight change in the absorption intensity for DPBF + H₂O₂, and the rate of oxidation of DPBF was 7.23×10^{-4} a.u. min⁻¹. A slight decrease in DPBF absorption was observed in the presence of CuONRs, with the rate of oxidation increasing to 8.70×10^{-3} a.u. min⁻¹. The rate of oxidation of DPBF was increased further when H₂O₂ and nanoparticles were all present in the solution. The rate of oxidation of DPBF was obtained as a slope of the linear plot of absorbance (a.u.) vs. time (min). The rates of oxidation for CuONRs, CuONRs@Au₆NPs, CuONRs@Pd₆NPs and CuONRs@Au₃Pd₃NPs are summarised in Table 2. The rate of oxidation of DPBF was faster in the presence of H₂O₂ and CuONRs@Au₃Pd₃NPs (4.44×10^{-2} a.u. min⁻¹). The higher the

Fig. 7 The rate of oxidation plots for (a)(i) DPBF + H₂O₂, (ii) DPBF + CuONRs and (iii) DPBF + H₂O₂ + CuONRs, (b) CuONRs@Au₆NPs with (i) DPBF alone and (ii) DPBF + H₂O₂, (c) CuONRs@Pd₆NPs (i) DPBF alone and (ii) DPBF + H₂O₂, and (d) CuONRs@Au₃Pd₃NPs with (i) DPBF alone and (ii) DPBF + H₂O₂. DPBF and (ii) DPBF + H₂O₂.

rate of DPBF oxidation, the higher the concentration of ROS produced. The rate of oxidation of DPBF is equivalent to the rate of ROS produced and it was found to decrease following this trend: CuONRs@Au₃Pd₃NPs (4.44×10^{-2} a.u. min⁻¹) > CuONRs@Pd₆NPs (1.74×10^{-2} a.u. min⁻¹) > CuONRs (1.62×10^{-2} a.u. min⁻¹) > CuONRs@Au₆NPs (1.23×10^{-2} a.u. min⁻¹). The DPBF studies confirmed ROS production from the interaction of H₂O₂ with the nanomaterials and the highest ROS generation efficiency of CuONRs@Au₃Pd₃NPs. ROS generation from the H₂O₂ was an important first step towards the oxidation of TMB and not observed in the absence of H₂O₂. The mechanistic steps involved in H₂O₂ catalytic reduction are as follows: step 1 adsorption of H₂O₂ *via* oxygen atoms onto the metal surfaces (Pd, Au and Cu) of the nanoparticle. Step 2 is the catalytic reduction which leads to bond breaking to produce ROS (OH[•], HO₂[•] and O₂[•]). Step 3, the generated ROS diffuses back into solution for further reaction. The noble metal nanoparticles (Au₆NPs, Pd₆NPs and Au₃Pd₃NPs) were adsorbed onto CuONRs *via* the lattice oxygen atoms. CuONRs@Pd₆NPs and CuONRs@Au₃Pd₃NPs exhibited enhanced catalytic activity due to the presence of Pd nanoparticles.

Glucose detection at Au₆NPs-, Pd₆NPs- and Au₃Pd₃NPs-supported on CuONRs

Glucose detection was investigated as the model analyte to demonstrate the applicability of the nanomaterials. Au₆NPs, Pd₆NPs and Au₃Pd₃NPs-supported on CuONRs were used in the colorimetric detection of glucose. H₂O₂ is the by-product of various oxidase enzymes with FADH₂ enzyme co-factor such as glucose oxidase (GOx), alcohol oxidase (AOx), uric oxidase (UOx), and cholesterol oxidase (ChOx).³³ GOx is used in the fabrication of biosensors for the detection of glucose. GOx enzymes as biocatalysts offer excellent selectivity and sensitivity towards the β-D-glucose substrates. After the production, H₂O₂ is catalytically oxidised at the nanoparticle surface and generate ROS, as confirmed above. In the presence of TMB as

Table 2 Comparison of the rate of oxidation of DPBF for the prepared Au₆NPs, Pd₆NPs and Au₃Pd₃NPs-supported on CuONRs

Nanomaterials	Rate of oxidation (min ⁻¹)		
	DPBF + H ₂ O ₂ (×10 ⁻⁴)	DPBF (×10 ⁻³)	DPBF + H ₂ O ₂ (×10 ⁻²)
CuONRs	7.23	8.70	1.62
CuONRs@Au ₆ NPs		4.10	1.23
CuONRs@Pd ₆ NPs		6.07	1.74
CuONRs@Au ₃ Pd ₃ NPs		5.62	4.44

a chromogenic substrate, the produced ROS oxidises TMB to form blue coloured products (TMBDI) characterised by the absorption band at 652 nm in the UV-vis spectrum. The mechanism of this biocatalysis is shown in Scheme 1.

Fig. 8 shows the UV-vis spectra and the corresponding plots of the absorbance at 652 nm *versus* concentration of glucose for (a) CuONRs, (b) CuONRs@Au₆NPs, (c) CuONRs@Pd₆NPs, and (d) CuONRs@Au₃Pd₃NPs. The concentration of glucose was varied from 0–60 μM, whilst the concentration of GOx, nano-materials, TMB and reaction time were kept constant. The UV-vis absorption at 652 nm increased linearly as the glucose concentration increases up to 40 μM for CuONRs and CuONRs@Au₆NPs. A wider linear curve was observed for CuONRs@Pd₆NPs and CuONRs@Au₃Pd₃NPs within the studied concentration range (0–60 μM) of glucose. The intensity of the absorption band at 652 nm was proportional to the concentration of glucose. Therefore, the amount or concentration of the produced H₂O₂ is directly proportional to the concentration of glucose. In the absence of glucose, no absorption signal was observed. The results confirmed that the presence of GOx in solution induces the biocatalytic oxidation of glucose to form gluconolactone. The enzyme was reduced to form GOx(FADH₂). GOx(FADH₂) reacts with oxygen to form GOx(FAD) and H₂O₂ as a by-product. The linear regression eqn (1)–(4) between the absorbance and varied glucose concentration were:

Fig. 8 UV-vis spectra and the corresponding plots of absorbance @ 652 nm *versus* [glucose] concentration for (a) CuONRs, (b) CuONRs@Au₆NPs, (c) CuONRs@Pd₆NPs and (d) CuONRs@Au₃Pd₃NPs.

$$\text{CuONRs: Abs} = 0.00376 [\text{glucose}] + 0.0663 \quad (1)$$

$$\text{CuONRs@Au}_6\text{NPs: Abs} = 0.00372 [\text{glucose}] + 0.0708 \quad (2)$$

$$\text{CuONRs@Pd}_6\text{NPs: Abs} = 0.00230 [\text{glucose}] + 0.102 \quad (3)$$

$$\text{CuONRs@Au}_3\text{Pd}_3\text{NPs: Abs} = 0.00305 [\text{glucose}] + 0.112 \quad (4)$$

Scheme 1 The mechanism of glucose detection using glucose oxidase (GOx) enzyme and CuONRs@Au₃Pd₃NPs as nanocatalyst for the catalytic reduction of H₂O₂ to produce ROS.

The limit of detection (LoD) and limit of quantification (LoQ) for the CuONRs supported catalysts were investigated. LoD and LoQ values were calculated using $S = 3\sigma$ for LoD and $S = 10\sigma$ for LoQ, where S is a signal and σ blank. The LoD and LoQ were calculated using the linear regression eqn (1) for CuONRs, and the LoD was 3.50 μM, and LoQ was 15.80 μM. Eqn (2) was used for LoD and LoQ for CuONRs@Au₆NPs and were found to be 3.80 μM and 17.10 μM. For CuONRs@Pd₆NPs, the LoD was 8.87 μM, and LoQ was 39.90 μM, using the linear regression eqn (3). For CuONRs@Au₃Pd₃NPs, the linear regression eqn (4) was used, the LoD was 7.34 μM, and the LoQ was found to be 33.10 μM. The selectivity and specificity for glucose detection was attributed to the selectivity of GOx enzyme towards β-D-glucose oxidation. The interference studies were therefore not investigated.

Conclusions

The preparation of Au₆NPs-, Pd₆NPs- and Au₃Pd₃NPs-supported on CuONRs was accomplished and confirmed by various methods. The prepared nanomaterials were investigated for their catalytic reduction of H₂O₂. The catalytic reduction of H₂O₂ by the nanomaterials was affected by different reaction conditions such as pH, reaction time, temperature and varied H₂O₂ concentrations. The investigation of the steady-state kinetic parameters showed that the prepared nanomaterials followed the Michaelis–Menten kinetics behaviour. The CuONRs@Pd₆NPs and CuONRs@Au₆NPs showed the lowest *K_m* values (2.94 mM and 3.11 mM) towards H₂O₂, and the values were within range for the HRP enzyme. TMB *K_m* values were higher for all our investigated nanomaterials when compared to the HRP enzyme. Nanomaterials demonstrated the effective production of ROS, with various experiments confirming this. The nanomaterials interact with H₂O₂ to produce reactive oxygen species (ROS). The prepared nanomaterials were successfully applied for the colorimetric detection of glucose with good linearity, limit of detection (LoD) and limit of quantification (LOQ) were determined.

Author contributions

S. S.: methodology, investigation, validation, data curation, visualisation, writing – original draft. O. A.: methodology, investigation, validation, data curation, visualisation, writing – review and editing. P. M.: methodology, validation, data curation, visualization, resources, funding acquisition; writing – review & editing, supervision, project administration, corresponding.

Conflicts of interest

The authors declare no known competing financial interests.

Acknowledgements

This research was funded by the National Research Foundation through Research Development Grant for Y-Rated Grant (UID 116331), NRF-STINT Bilateral (UID 118725), Rated Incentive funding (UID 96001) and Rhodes University through Sandisa Imbewu and Researcher Development Grant. SS thanks NRF for an MSc scholarship, and OA thanks Rhodes University for a postdoctoral fellowship.

References

- 1 C. Evangelisti, E. Schiavi, L. A. Aronica, A. M. Caporusso, G. Vitulli, L. Bertinetti, G. Martra, A. Balerna and S. Mobilio, *J. Catal.*, 2012, **286**, 224–236.
- 2 M. B. Gawande, A. Goswami, T. Asefa, H. Guo, A. V. Biradar, D. L. Peng, R. Zboril and R. S. Varma, *Chem. Soc. Rev.*, 2015, **44**, 7540.
- 3 S. Y. Tee, E. Ye, P. H. Pan, C. J. J. Lee, H. K. Hui, S. Y. Zhang, L. D. Koh, Z. Dong and M. Y. Han, *Nanoscale*, 2015, **7**, 11190.
- 4 W. Yao, F. L. Li, H. X. Li and J. P. Lang, *J. Mater. Chem. A*, 2015, **3**, 4578.
- 5 Y. G. Sun and Y. N. Xia, *J. Am. Chem. Soc.*, 2004, **126**, 3892.
- 6 C. M. Cobley, D. J. Campbell and Y. N. Xia, *Adv. Mater.*, 2008, **20**, 748.
- 7 X. Liu, J. Locozzia, Y. Wang, X. Cui, Y. Chen, S. Zhao, Z. Li and Z. Lin, *Energy Environ. Sci.*, 2017, **10**, 402.
- 8 K. C.-F. Leung, S. Xuan, X. Zhu, D. Wang, C.-P. Chak, S.-F. Lee, W. K.-W. Ho and B. C.-T. Chung, *Chem. Soc. Rev.*, 2012, **41**, 1911.
- 9 N. E. Motl, A. F. Smith, C. J. DeSantis and S. E. Skrabalak, *Chem. Soc. Rev.*, 2014, **43**, 3823.
- 10 J. Liu, W. Wang, T. Shen, Z. Zhao, H. Feng and F. Cui, *RSC Adv.*, 2014, **4**, 30624.
- 11 A. Ahmad, V. N. Kalevaru and A. Martin, *Catalysts*, 2016, **6**(7), 97.
- 12 W. He, H.-K. Kim, W. G. Wamer, D. Melka, J. H. Callahan and J.-J. Yin, *J. Am. Chem. Soc.*, 2013, **136**, 750.
- 13 C. Xu, J. Xie, D. Ho, C. Wang, N. Kohler, E. G. Walsh, J. R. Morgan, Y. E. Chin and S. Sun, *Angew. Chem., Int. Ed.*, 2008, **47**, 173.
- 14 S. Wang, Z. Wang and Z. Zha, *Dalton Trans.*, 2009, 9363.
- 15 G. J. Hutchings and C. J. Kiely, *Acc. Chem. Res.*, 2013, **46**(8), 1759–1772.
- 16 M. S. Chavali and M. P. Nikolova, *SN Appl. Sci.*, 2019, **1**, 607.
- 17 L. Long, J. Liu, K. Lu, T. Zhang, Y. Xie, Y. Ji and X. Wu, *J. Nanobiotechnol.*, 2018, **16**, 46.
- 18 L. Gao and X. Yan, Nanozymes: an emerging field bridging nanotechnology and biology, *Sci. China: Life Sci.*, 2016, **59**, 400, DOI: 10.1007/s11427-016-5044-3.
- 19 Y.-C. Yang, Y.-T. Wang and W.-L. Tseng, *ACS Appl. Mater. Interfaces*, 2017, **9**, 10069–10077.
- 20 X. Wang, Q. Han, S. Cai, T. Wang, C. Qi, R. Yang and C. Wang, *Analyst*, 2017, **142**, 2500.
- 21 L. Gao, J. Zhuang, L. Nie, J. Zhang, Y. Zhang, N. Gu, T. Wang, J. Feng, D. Yang, S. Perrett and X. Yang, *Nat. Nanotechnol.*, 2007, **2**, 577.
- 22 R. Andre, F. Natalio, M. Humanes, J. Leppin, K. Heinze, R. Wever, H.-C. Schroder, W. E. G. Muller and W. Tremel, *Adv. Funct. Mater.*, 2011, **21**, 501.
- 23 G.-W. Wu, S.-B. He, H.-P. Peng, H.-H. Deng, A.-L. Liu, X.-H. Lin, X.-H. Xia and W. Chen, *Anal. Chem.*, 2014, **86**, 10955.
- 24 W. Chen, J. Chen, Y.-B. Feng, L. Hong, Q.-Y. Chen, L.-F. Wu, X.-H. Lin and X.-H. Xia, *Analyst*, 2012, **137**, 1706.
- 25 M. Marelli, A. Jouve, A. Villa, R. Psaro, A. Barlena, L. Prati and C. Evangelisti, *J. Phys. Chem.*, 2009, **123**, 2864.
- 26 J. M. Berg, A. Romoser, N. Banerjee, R. Zedba and C. M. Sayes, *Nanotoxicology*, 2009, **3**, 276.
- 27 D. Zhou, K. Zeng and M. Yang, *Microchim. Acta*, 2019, **186**, 161.
- 28 M.-C. Kim, D. Lee, S. H. Jeong, S. Y. Lee and E. Kang, *ACS Appl. Mater. Interfaces*, 2016, **8**, 34317.
- 29 Y. Song, K. Qu, C. Zhao, J. Ren and X. Qu, *Adv. Mater.*, 2010, **22**, 2206.
- 30 O. Adeniyi, S. Sicwetsha and P. Mashazi, *ACS Appl. Mater. Interfaces*, 2020, **12**, 1973.

- 31 S. Mvango and P. Mashazi, *Mater. Sci. Eng., C*, 2019, **96**, 814.
- 32 K. Zamojc, M. Zdrowowicz, P. B. Rudnicki-Velasquez, K. Krzyminski, B. Zaborowski, P. Niedzialkowski, D. Jacewicz and L. Chmurzynski, *Free Radical Res.*, 2017, **51**, 38.
- 33 T. Li, K. Zhu, S. He, X. Xia, S. Liu, Z. Wang and X. Jiang, *Analyst*, 2011, **136**, 2893.