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One-step green approach for functional printing and finishing of textiles using silver and gold NPs

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In this study, we present a successful simple method for printing and finishing of polyester and cotton fabrics using gold and silver nanoparticles (Au-NPs and Ag-NPs, respectively) as stable, fast colorants and functional components. The surface plasmon resonance (SPR) bands of the colloidal gold and silver NPs were observed at λ_{max} 520 nm and 450 nm, respectively, indicating the presence of spherical Au-NPs and Ag-NPs, which was further confirmed by TEM analysis. The printed samples were subjected to SEM, XRD and EDX analyses. The SEM images and EDX spectra unequivocally confirmed the existence of embedded NPs on the fabric surfaces. Both the cotton and polyester samples possessed excellent color fastness, as indicated from the color fastness test. The functional properties of the printed fabrics indicated that the incorporation of Au-NPs and Ag-NPs into the fabrics simultaneously imparted multifunctional properties such as stable brilliant colors, highly durable antimicrobial activity and very good UV-protection properties.

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Introduction

Textile printing is the most versatile and important method used to introduce color and design in textile fabrics. It is also known as localized application of dye or pigment in a thickened form to a substrate to create an attractive design with welldefined boundaries. In the textile industry, screen-printing is the most frequently applied method for textile prints, and it accounts for nearly 50% of the printing production worldwide. The main factors for its widespread use are the quality of the prints, applicability to almost every type of fiber or mixture, ability to withstand any washing processes after fixation, its simplicity, low cost, and minimum requirements for wet-processing.^{1,2} However, the synthetic dyes or pigments and their auxiliaries used in the printing process may be toxic and produce harmful effects on humans and the environment. In addition, several synthetic and natural dyes have low durability to washing and UV light, leading to color fading.3

On the other hand, textile functionalization with noble metal (gold and silver) nanoparticles (NPs) has increased in recent years. Many attempts have been made to investigate different

methods for their preparation⁴⁻¹⁰ and to enhance the functions of the textiles such as flame retarding ability, hydrophobicity, self-cleaning, and resistance to wrinkles as well as antistatic, antimicrobial and UV-protective properties.¹¹⁻¹³ In addition, these NPs are endowed with unique localized surface plasmon resonance properties (SPR) and thus exhibit brilliant and different colors. With such excellent properties, novel shades of elegant hues and excellent multifunctional activities on different textile materials have been achieved.^{14,15}

Nevertheless, many dyes are not stable under irradiation of sunlight and have low resistance to washing and rubbing, causing color fading. Metal NPs are different from traditional dyes: it is not the chromophore of traditional dyes but the shape and size of NPs that define the colors. This property makes all the colors stable towards UV light, and they do not change provided that there is no change in the particle size through the growth or reduction of discrete nanoparticles or agglomeration. According to the previous literatures, most printing methods are based on the utilization of pigments or dyes as colorants incorporated with metal NPs as finishing agents to produce multifunctional printed textiles. However, these methods consume a lot of chemicals, time, and energy.

Recently, several attempts have been carried out to use gold and silver NPs to dye fabrics with colors and simultaneously implement functionality to the dyed fabric, *e.g.*, antibacterial activity^{22,23} and/or UV-protection.²⁴ Based on the literature, the incorporation of gold and silver NPs into fabric or fiber materials is applied using three different procedures: impregnation of fabrics in the prepared metal NP colloidal solution,^{25–27} preparing metal NPs *in situ* in fabrics,^{14,28–32} and synthesizing

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polymer-NP composites followed by a spinning process to form colored fibers. 33

To the best of our knowledge, textile printing with metal nanoparticles as a colorant has not been reported to date. Therefore, the main task of this study is to investigate a novel, simple, one-step, green approach for printing natural and synthetic fabrics, namely, cotton and polyester fabrics using gold and silver NPs as stable and fast colorants. This is an environmentally friendly solution for the current harmful printing, coloration and finishing processes.

2. Experimental

2.1. Materials

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Scoured and bleached plain weave polyester (100%, 109 g m $^{-2}$) and mill-scoured, bleached knitted cotton (200 g m $^{-2}$) fabrics were used.

Sera® Binder M-CPB liquid (acrylate-based copolymer, anionic, Clariant) and Sera® print M-CPK thickener 160 EG liquid (synthetic thickening agent based on ammonium polyacrylate, Clariant) were used.

Chloroauric acid (HAuCl₄·3H₂O, Sigma-Aldrich) was purchased from Skyspring Nanomaterials, Inc. USA.

All other chemicals used during this study such as AgNO $_3$ (Sigma) and tri-sodium citrate Na $_3$ C₆H $_5$ O $_7 \cdot$ 2H $_2$ O (99.5%) were of commercial grade.

2.2. Methods

2.2.1. Preparation of gold nanoparticles (Au-NPs). Au-NPs were prepared by adding 5 ml of 5 mM of gold(III) chloride hydrate, HAuCl₄ (2 mg ml⁻¹), to 85 ml of filtered deionized water, which was then refluxed in a 250 ml flask over a hot plate and heated to boiling point. Five milliliters of 25 mM sodium citrate solution was quickly added to the boiling solution and stirred for 30 min until the color became wine-red.³⁴

2.2.2. Preparation of silver nanoparticles (Ag-NPs)

- 2.2.2.1. Preparation of extract. Pluchea dioscoridis leaves were collected from Damietta, Egypt. The leaves were cleaned well and washed using tap water followed by distilled water. The clean leaves were dried for a week at room temperature and then was ground to powder form. Five g of the leaf powder was introduced into boiling distilled water (200 ml) in an appropriate flask for 5 min. Then, the mixture was filtered, and the filtrate was stored at $40\,^{\circ}\mathrm{C}$ for further use.
- 2.2.2.2. Synthesis of silver nanoparticles (Ag-NPs). Various volumes (1 and 12 ml) of Pluchea dioscoridis extracts (Pd) were placed in 100 mL Erlenmeyer flasks. Then, 45 ml AgNO₃ with different concentrations (1–5 mM) was titrated drop by drop in each flask for 10 min. The formation of Ag-NPs was confirmed by the change in colour from colorless to reddish-brown.³⁵
- **2.2.3. Printing paste method.** The formulation for aqueous printing using the flat screen technique is shown in Table 1

Printed fabric samples were then simultaneously dried and fixed in a high temperature steamer at 150 $^{\circ}$ C for 4 min followed by washing using 2 g l⁻¹ non-ionic detergent at a liquor ratio of 1:50 at 60 $^{\circ}$ C for 10 min.

Table 1 Formulation for aqueous printing using flat screen technique

Printing paste components	$\rm g~kg^{-1}$ paste
Thickener	20
Binder	100
Nano-material	20
Water	840
Total	1000

2.3. Characterization

- **2.3.1. TEM analysis.** Transmission electron microscopy (TEM) analysis of the cell-free extract was used to determine the size and morphology of the synthesized nanoparticles using a JOEL model 1200EX election microscope operated at an accelerating voltage of 120 kV. The specimen for TEM measurement was prepared by dissolving a drop of colloidal solution on a 400-mesh copper grid coated with an amorphous carbon film and evaporating the solvent in air at room temperature.
- **2.3.2. UV-vis spectroscopy.** UV-vis spectroscopy on Shimadzu UV-2450 was used to qualitatively confirm the presence of Au-NPs and Ag-NPs in the reaction medium. The reaction solutions exhibited intense absorption peaks due to surface plasmon resonance (SPR).
- **2.3.3. SEM and EDX analysis.** Scanning electron microscopy (SEM) images and EDX analysis of the printed metal NP samples were obtained using JEOL JSM-6510LB Japan-Tokyo.
- **2.3.4. X-ray diffraction (XRD).** The crystalline natures of the prepared NPs and printed samples were determined using an X-ray diffractometer (XRD) (Bruker D8 ADVANCE, Karlsruhe, Germany).

2.4. Testing

- **2.4.1.** Colorimetric analysis. The colorimetric parameters L^* (lightness), a^* (redness-greenness), b^* (yellowness-blueness) and color strength (K/S) of the printed cotton and polyester fabrics were measured using a Konica Minolta CM-3600 d spectrophotometer (Minolta, Tokyo, Japan).
- **2.4.2.** Color fastness. The color fastness properties of the obtained prints to washing, rubbing and light were determined according to the AATCC Test Methods (61-1972), (8-1972), and (16A-1972), respectively.
- **2.4.3. Durability test.** Durability to washing was evaluated according to the AATCC Test Method 61(2A)-1996 after 10 washing cycles.

2.5. Functional properties of metal NP-printed fabrics

2.5.1. Antimicrobial activity. The antimicrobial activities against G+ve bacteria (*Staphylococcus aureus* and *Bacillus cereus*) and G–ve bacteria (*Escherichia coli* and *Candida utilis*) were evaluated qualitatively according to the AATCC Test Method (147-1988) and were expressed as the zone of growth inhibition (mm).

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2.5.2. UV-protection properties. UV-protection functionality, expressed as UV-protection factor (UPF), was evaluated according to AS/NZS 4399: 1996, and protection was rated as good, very good or excellent if the UPF values ranged from 15 to 24, 25 to 39, or above 40, respectively.

3. Results and discussion

The main task of the current study is to obtain a facile novel one step green procedure for printing cotton and polyester fabrics using colloidal gold (Au-NPs) and silver (Ag-NPs) as stable and fast colorants to attain multifunctional prints based on nanomaterials. The results obtained and the appropriate discussions are presented below.

3.1. Characterization of the synthesized nanoparticles

TEM analysis. The morphology, shape, and size of the synthesized Au-NPs and Ag-NPs were characterized using transmission electron microscopy. Fig. 1 shows the TEM images of Au-NPs and Ag-NPs. The size of the NPs was determined by measuring the diameter of the entire particle in the TEM images. As depicted from Fig. 1a, the average diameter of the gold NPs was found to range between 13 and 20 nm in a monodispersed state due to the negative charge of the citrate ions,

which repel each other.36 Moreover, the TEM image demonstrates the spherical shape of the synthesized Au-NPs. Fig. 1b presents the TEM image of the synthesized Ag-NPs. It was clear that the particles exhibited a spherical-like shape and were uniformly distributed and well dispersed. The average size of the prepared Ag-NPs was in the range of 15-25 nm, which indicated sufficient homogeneity and well dispersion of the silver particles. Ag-NPs were observed to be sufficiently close to each other, which attributed to the lack of reducing or stabilizing agent in the solution.37

The size of numerous particles were measured using the TEM images, and histograms providing different size distributions of the well-dispersed suspensions of Au and Ag NPs were given in Fig. 1c and d, respectively. The width of the Au-NP bins was 3 nm, as illustrated in Fig. 1c, and they were centered at 12, 15, 18 and 21 nm. Au-NPs around 18 nm in size exhibited the highest percentage value (47.79%) in comparison with other Au-NPs present in the colloid. From Fig. 1d, for Ag-NPs, the bins of the histogram were 5 nm wide, and they were centered at 5, 10, 15, 20 and 25 nm. The majority of particles (30-33%) were located at around 5 nm, which may be the main reason for the high antimicrobial activity of Ag-NPs.

3.1.2. UV-vis spectroscopy. UV-vis spectroscopy is one of the most common techniques used to verify the formation and stability of metal NPs in aqueous solutions. Gold and silver NPs

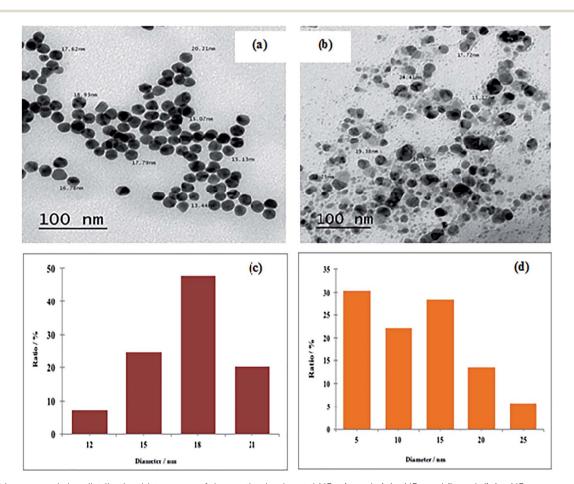


Fig. 1 TEM images and size distribution histograms of the synthesized metal NPs: (a and c) Au NPs and (b and d) Aq-NPs.

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have been reported to exhibit different colors in aqueous solutions, depending on their size and intensity in relation to their SPR.38 The synthesis of stable Au-NPs was conducted through a reduction process using sodium citrate as the reductant and stabilizer. The interest in this synthetic process is due to the resultant versatile citrate layer on the Au-NP surface, which allowed easy multifunctionalization of the obtained particles.³⁹

Fig. 2a shows the UV-vis spectrum of the synthesized Au-NPs, and a maximum absorbance peak of the colloidal Au-NPs occurred at about 520 nm. Generally, Au-NPs displayed a single absorption peak in the visible range between 510 and 550 nm due to its SPR and exhibited heavy absorption of visible light at 520 nm. This was resulted in a brilliant wine-red color for the gold NPs, which varied according to its size and shape. Alagad et al. 40 also observed the absorbance spectrum of Au-NPs at around 520 nm, whereas HAuCl4 did not exhibit any absorbance at the same wavelength.

Fig. 2b shows the UV-vis spectrum of the synthesized Ag-NPs; it can be clarified that due to the yellowish color of the colloidal Ag-NPs, the maximum absorption peak was at about 450 nm, which was specific to SPR of spherical Ag-NPs.37 This result confirmed the proposed formation according to transmission electron microscopy (TEM). Both the UV-vis absorption (Fig. 2b) and TEM (Fig. 1b) results verified that the yellow color of the colloidal Ag-NPs was a result of the presence of spherical Ag-NPs.

The observed broad peak illustrates that the synthesized Ag-NPs were different in size and shape, which was in agreement with the TEM analysis (Fig. 1b). Krishnaraj et al.41 reported that a single SPR band indicates spherical nanoparticles, whereas two or more SPR bands indicate anisotropic particles. Furthermore, it was observed that another strong UV absorption peak appeared at about 350 nm, which may be due to the residual plant extract used in the synthesis of Ag-NPs or trace iron impurities, which appeared in the ppm level.42

3.1.3. SEM and EDX analysis. Fig. 3 shows the SEM images of the polyester and cotton fabrics printed with Au-NPs and Ag-NPs. It is clear that there is a change in their surface morphology with the deposition of NPs onto their surfaces. Furthermore, the EDX spectra confirmed the existence of metal

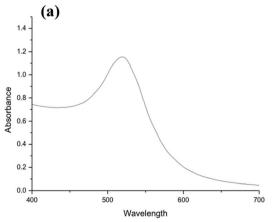
nanoparticles (Au-NPs and Ag-NPs) on the fabric surfaces. Therefore, the results indicated that Ag-NPs and Au-NPs were successfully distributed on all the examined printed fabric surfaces.

3.1.4. X-ray diffraction (XRD). The formation of metal NPs was also confirmed by X-ray diffraction (XRD) analysis. The XRD patterns of the synthesized Au-NPs in Fig. 4 revealed that Au is crystalline with a face-centered cubic (fcc) structure corresponding to clear peaks at 38.25 (111), 44.46 (200), 64.64 (220), and 77.20 (311), respectively. The obtained results were in good agreement with the data for standard gold in JCPDS file no. 04-0784. On the other hand, the XRD patterns of the synthesized Ag-NPs in Fig. 4 exhibited a face centered cubic structure according to the four reflection peaks at 2θ values of 38.11° , 44.29°, 64.45° and 77.39°, which were indexed to the (111), (200), (220), and (311) crystal planes, respectively (JCPDS card no. 89-3722). The XRD results provided strong evidence for the presence of gold and silver particles and thus confirmed the UVvis and TEM results.43,44

Color measurements

The color of printed fabrics is of key significance since it may impact the product demand and consumer inclination. To investigate the color properties of nano-printed fabric surfaces, their colorimetric data were measured for Au-NP- and Ag-NPprinted fabrics using the CIELAB system in terms of L*, a*, and *b** to study the effect of gold and silver NPs as new colorants for cotton and polyester fabrics. The color coordinates are listed in Tables 2 and 3. After printing fabrics using colloidal solutions of Au-NPs and Ag-NPs, red-wine and yellow printed polyester and cotton fabrics were obtained with different shades and very good homogeneity, as shown in Fig. 5.

Table 2 shows the colorimetric data for the printed fabrics using Au-NPs and Ag-NPs. At low concentrations of NPs, the colors of the printed cotton and polyester fabrics were light redwine for Au-NPs and light-yellow for Ag-NPs, as shown by the low positive a^* and b^* values. With an increase in the concentration of Au-NPs, the positive a^* value progressively increased giving a reddish tone to the printed fabrics. The color of the Ag-



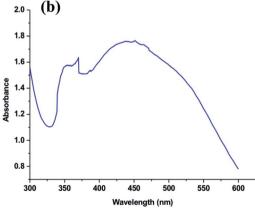


Fig. 2 UV-visible absorption spectra of the synthesized NPs: (a) Au-NPs and (b) Aq-NPs.

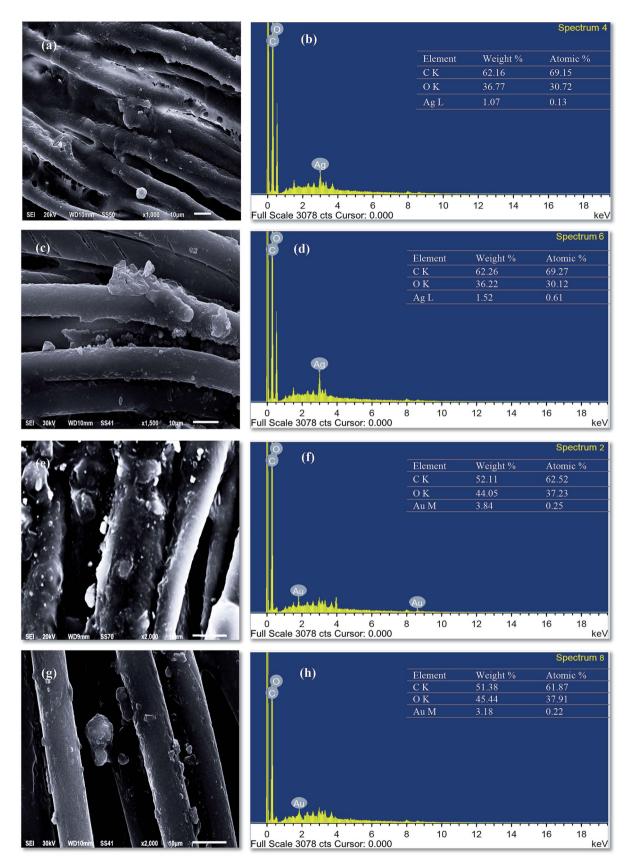
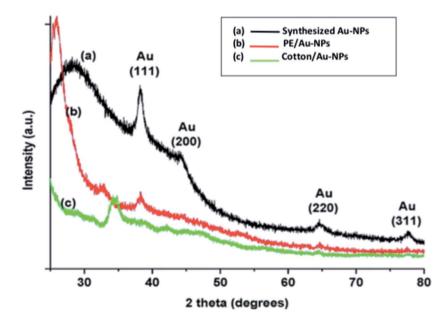


Fig. 3 SEM images and EDX spectra of the metal NP-printed samples: Ag-NPs cotton (a and b) and polyester (c and d). Au-NPs cotton (e and f) and polyester (g and h).



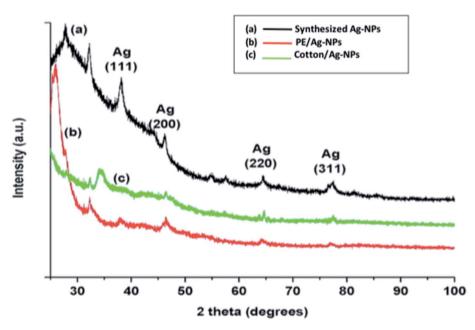


Fig. 4 XRD patterns of the synthesized metal NPs and printed fabrics.

Table 2 Colour characteristics and K/S values of the Au-NP-printed fabrics

Sample Au-NP Conc.	Cotton				Polyester				
	L^*	a*	<i>b</i> *	K/S	L^*	a*	<i>b</i> *	K/S	
0.3 mµ	69.58	20.17	0.71	$1.4 (1.2)^a$	65.39	27.71	4.36	1.3 (1.2)	
0.5 mμ	60.09	29.05	1.14	3.6 (3.2)	57.24	28.54	9.1	2.6 (2.5)	
0.8 mμ	50.16	36.74	5.15	5.7 (5.3)	48.95	34.62	10.7	4.6 (4.5)	
2.5 mµ	47.69	36.45	9.6	6.8 (6.1)	43.92	34.82	11.53	6.7 (6.6)	
4 mμ	43.89	37.79	6.07	8.2 (8.01)	38.63	38.11	12.83	9.3 (9.1)	

 $[^]a$ Values in parentheses indicate K/S after 10 washing cycles.

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Table 3 Colour characteristics and K/S values of the Aq-NP-printed fabrics

Sample Ag-NP Conc.	Cotton				Polyester			
	L^*	a*	<i>b</i> *	K/S	L^*	a*	<i>b</i> *	K/S
5 mμ	85.72	6.52	6.69	$2.7 (2.5)^a$	81.87	4.68	20.76	1.2 (1.2)
6 mμ	82.91	4.54	21.54	4.6 (4.6)	69.55	13.93	33.6	2.5 (2.4)
7 mμ	78.94	5.95	30.67	5.7 (5.6)	61.97	18.8	37.38	3.7 (3.5)
8 mµ	75.7	11.1	39.74	7.4 (7.2)	60.35	23.43	46.59	6.1 (6)
9 mμ	58.91	21.57	43.44	9.4 (9.3)	46.59	24.59	52.01	8.9 (8.7)

^a Values in parentheses indicate K/S after 10 washing cycles.

NP-printed fabrics gradually increased, giving a dark yellowish tone with an increase in the concentration of Ag-NPs, as shown by the positive high b^* values. Furthermore, the lightness (L^*) values gradually decreased for all the printed fabrics. This decrease in L^* was a result of the coloration of the fabrics, which became deeper by increasing the concentrations of Au and Ag metal NPs.

The color strength (K/S) data were measured at the wavelength of 425 nm, which is related to the yellow color of Ag-NPs, whereas the wine-red color of Au-NPs was measured at the wavelength of 520 nm. Fig. 6 shows that increasing the concentration of the synthesized Au-NPs and Ag-NPs resulted in a significant improvement in the color depth of the obtained metal Au and Ag NP prints expressed as the K/S values. The observed data indicated that the darker yellow and red-wine colors of the printed fabrics were achieved by using higher concentrations of Au-NPs and Ag-NPs. This can be explained by the fact that a high number of spherical NPs were formed in the fabrics under investigation with the use of more concentrated Au-NP and Ag-NP colloidal solutions.

Moreover, the fabrics printed with Au-NPs had a red-wine color, and the fabrics printed with Ag-NPs had a yellow color with different shades [from faint (red-wine/yellow) to dark (redwine/yellow)], but the nature of the color did not change regardless of the concentration of NPs. This may be regarded to the fact that the change in SPR of Au-NPs and Ag-NPs with different shapes and sizes on the polyester and cotton samples led to bright colors of the printed fabrics.

These results support the suggestion that cotton and polyester fabrics can be printed by synthesized silver and gold colloidal NPs as stable and fast colorants, and the color strength (K/S) of these fabrics can be controlled by changing the concentration of synthesized Au-NPs and Ag-NPs.

3.3. Fastness properties

The coloration requirements are not limited to imparting color to the fabrics, but the acquired color should withstand the action of certain tests. Thus, different color fastness properties (washing, rubbing and light) of all the printed cotton and polyester fabrics were measured according to standard AATCC Test Methods (61-1972) and (8-1972), and (16A-1972), respectively, and the results are reported in Table 4. The results showed that all the printed fabrics exhibited excellent fastness

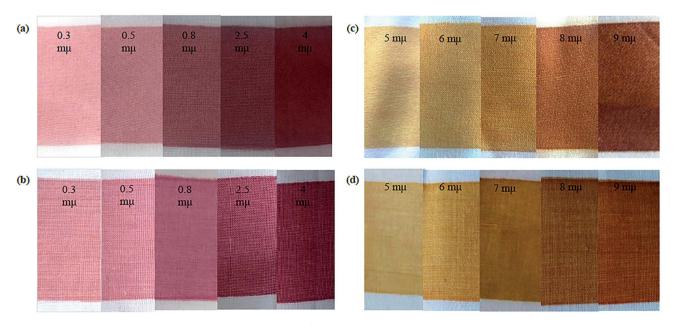
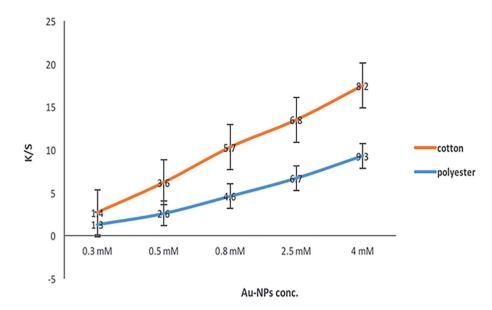


Fig. 5 Photographs of the metal NP-printed fabrics: (a) polyester/Au-NPs, (b) cotton fabrics with Au-NPs, (c) polyester fabrics with Ag-NPs, and (d) cotton fabrics with Aq-NPs.

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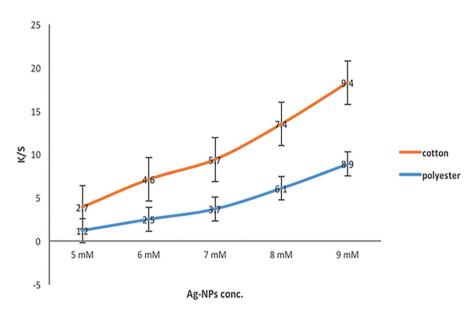


Fig. 6 Effect of metal NP concentration on K/S of the printed fabrics.

levels to washing and rubbing and a very good fastness level to light. The obtained results reflected the ability of the suggested technique to yield good printed fabrics with excellent fastness properties without the use of any other chemicals (*e.g.*, crosslinkers or coating materials).

3.4. Antimicrobial activity

The antimicrobial properties of the Au-NP- and Ag-NP-printed fabrics are shown in Table 5. The results revealed that the unprinted samples were not affected, and there were no inhibition areas. On the other hand, the cotton samples printed using Ag and Au-NPs as colorants exhibited excellent antimicrobial activities against *S. aureus*, *Bacillus cereus*, *E. coli* and *Candida utilis*. It is known that nano-materials have strong inhibiting effects towards a broad spectrum of bacteria and

yeasts, especially those Ag-NPs displaying good activity against all the indicator pathogens, which shows the potential broad spectrum antimicrobial activity. 45 The remarkable improvement

Table 4 Fastness properties of the metal NP-printed fabrics

		WF^a		RF^b		
Type of NPs	Type of fabric	St.	Alt.	Wet	Dry	LF^c
Control	Polyester	_	_	_	_	_
	Cotton	_	_	_	_	_
Au-NPs	Polyester	5	5	4/5	5	4
	Cotton	5	5	4/5	5	3/4
Ag-NPs	Polyester	5	5	4/5	5	3/4
	Cotton	5	5	4/5	5	4

^a Wash fastness. ^b Rubbing fastness. ^c Light fastness.

Table 5 Antimicrobial activity of the metal NP-printed fabrics

Type of NPs		Antimicrobial activity ZI ^a (mm)						
		G+ve		G-ve	Yeast			
	Type of fabric	S. aureus	Bacillus cereus	E. coli	Candida utilis			
Control	Polyester	0	0	0	0			
	Cotton	0	0	0	0			
Au-NPs	Polyester	16	20	22	18			
	Cotton	18	22	16	16			
Ag-NPs	Polyester	28	26	30	18			
	Cotton	24	28	18	20			

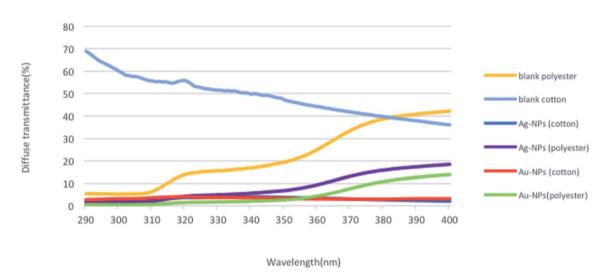


Fig. 7 Transmittance spectra of the blank and metal NP-printed fabrics.

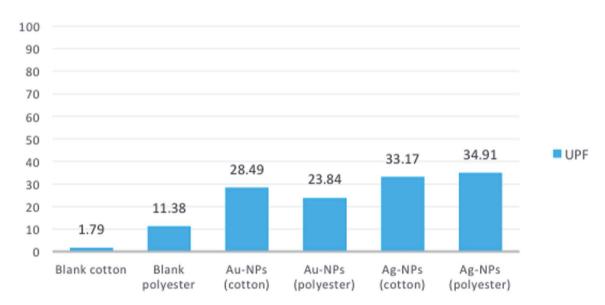


Fig. 8 UPF values of the blank and metal NP-printed fabrics.

in the antimicrobial functionality is ascribed to the interaction of the negatively charged cell walls of the pathogens with the positively charged cationic sites of the antimicrobial agent, which changes its chemical and physical properties. This action interrupts cell membrane functions and protein activity as well as the ability to multiply. In addition, the silver and gold Table 6 Durability of the imparted functional and color fastness properties after 10 washing cycles

		Fastness properties					Functional properties				
${ m WF}^a$		RF^b	RF^b			Zi ^d (mm)					
							G+ve		G-ve	Yeast	
Type of NPs	Type of fabric	Type of fabric St. Alt. We	Wet	Wet Dry LF ^c		S. aureus	Bacillus Cereus	E. coli	Candida utilis	UPF	
Au-NPs	Polyester	5	4/5	4/5	5	4/5	14	16	18	14	22.34
	Cotton	5	4/5	4/5	5	4	16	20	14	12	26.87
g-NPs	Polyester	5	4/5	4/5	5	4	23	21	25	14	32.88
	Cotton	5	4/5	4/5	5	4/5	21	25	18	16	30.52

^a Wash fastness. ^b Rubbing fastness. ^c Light fastness. ^d Zone of inhibition.

nanoparticles released silver and gold ions, which produced higher biocidal effect on the microorganisms.^{11,45}

3.5. UV-protection activity

To investigate the UV-protection ability of the blank and polyester and cotton fabrics printed with Au-NPs and Ag-NPs, their UV light transmittance and UPF values were measured, and the results were illustrated in Fig. 7 and 8.

As expected, the blank samples are not good UV filters and a high percentage of UV light can penetrate into the fabric. However, the average transmittance values in the UVA (315–400 nm) and UVB (280–315 nm) regions for the fabrics decreased clearly after printing the fabrics with Au-NPs and Ag-NPs, which indicated that the gold and silver NPs noticeably improved the UV-blocking abilities of the polyester and cotton samples.

The UPF values of Au-NP- and Ag-NP-printed fabrics exhibit very good UV-protection according to the Australian classification scheme compared with those of blank fabrics, as shown in Fig. 8. Therefore, these results confirmed the UV reflection ability of Au-NPs and Ag-NPs, which can effectively decrease aging and reduce human skin damage caused by harmful UV-radiation.

3.6. Durability to washing

The results illustrated in Table 6 reveal that the retention of the imparted functional properties, *i.e.*, antimicrobial and UV-protection together with the fastness properties as well as the color strength (K/S) of the obtained metal NPs prints was still high even after 10 washing cycles. This means that the metal NPs (Au-NPs and Ag-NPs) were still tightly loaded and fixed onto the simultaneously printed and finished fabric surfaces.

4. Conclusion

A facile and feasible strategy was devised to produce colorful and multifunctional printed polyester and cotton fabrics using synthesized NPs. This method featured the novelty of employing gold and silver NPs as eco-friendly stable colorants and functional components in a one-step process for printing and functional finishing of polyester and cotton fabrics. The results of the UV-visible, TEM, SEM and XRD measurements confirmed

the successful synthesis and uniform distribution of spherical Au-NPs and Ag-NPs on polyester and cotton fabrics. The brilliant colored appearance, excellent color fastness properties, highly durable antibacterial activity and very good UV-protective properties of the nano-printed fabrics indicated their ability to produce good printed/finished fabrics in a one-step process without using traditional dyes or auxiliary chemicals.

Conflicts of interest

There are no conflicts to declare.

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