

Luminescence Detection of Latent Fingermarks on Non-porous Surfaces with Heavy-Metal-Free Quantum Dots

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Abstract: Current and proposed nanoparticle-based techniques for development of latent fingermarks suffer a number of drawbacks such as complicated, multi-step and time-consuming procedures, batch-to-batch variability, expensive reagents, large background noise and toxicity. Here, we introduce a promising green development technique based on heavy-metal-free quantum dots for the detection of latent fingermarks on non-porous surfaces. Red-near infrared luminescent Cu-In-S/ZnS core-shell QDs solution was produced in large scales using a water-based, simple and fast method using N-acetylcysteine as a biocompatible surfactant to coat the particles. The coated QDs were applied to the successful development of latent fingermarks deposited on a variety of surfaces, including highly patterned polymer banknotes and the sticky side of adhesive tape.

1.0 Introduction

The detection of latent (not visible to the naked eye) fingermarks has been carried out for more than a century to enable associations between individuals and objects found at crime scenes to be evaluated [1, 2]. In more recent years innovative methods and materials based on nanotechnology have been applied to the visualisation of latent fingermarks with noticeable progress from specifically targeting fingermark components [3-5].

Luminescent quantum dots (QDs) manifesting large cross sections of the UV/visible spectral range, narrow and size-dependent emission wavelengths, high quantum efficiencies, and enhanced photostabilities, have been applied to the development of latent fingermarks on non porous surfaces over the last two decades [6, 7]. However even though cadmium-based QDs such as CdS [6], CdSe [8] and CdTe [9, 10] have successfully visualised latent fingermarks, issues relating to the toxicity associated with heavy metals remain a concern [11]. In order to increase their biocompatibility some reported methods have included covering the QDs with a surfactant or over coating them with a non-toxic inorganic shell [12], however, even with these techniques, the underlying issue of toxicity is unsolved. Heavy-metal-free QDs such as carbon [13], Cu- and Mn-doped ZnS [14, 15] quantum dots have been proposed as alternatives, however, the latent fingermarks visualised by these QDs, which luminesce in the visible part of the spectrum, exhibit reduced sensitivity of the detection due to background noise [13-15].

Luminescence increases the contrast between latent fingermark ridges and the underlying substrate by moving the means of detection outside of the ambient illumination [16, 17]. Materials luminescing at longer wavelengths, (red to near infrared (NIR) region), have the potential to improve detectability of fingermarks

since there is less interference in this region. However, only a limited number of NIR-based techniques have been reported for the detection of fingermarks and these techniques suffered drawbacks including toxicity, being a post treatment rather than a standalone process and lack of contrast [18-20].

To fill these technology gaps, three main objectives for this work were identified: first, preparation of a red-NIR luminescent heavy-metal-free QDs water-based solution; second, modification of the QDs to boost their affinity to interact with the fingermark residue and third, making the procedure simple and practical for operational use.

Ternary metal complex chalcogenide, Cu-In-S/ZnS (CIS-ZS) core-shell quantum dots are of low toxicity and exhibit strong and broad absorption of ultraviolet and visible light, intense photoemission and reasonable stability under photoexcitation [21, 22]. CIS-ZS QDs have been used in a variety of light-emitting applications such as solar cells [23], electro-luminescence [24], photocatalysts [25] and bio-imaging [26-28]. However, to the best of our knowledge, ternary core-shell QDs with photoluminescence (PL) in red/NIR range have not been used for fingermark visualisation. This study provides a proof of concept for the use of CIS-ZS quantum dots in aqueous solution, modified with surfactant coatings, for the detection of latent fingermarks on a range of non-porous surfaces.

2.0 Materials and Methods

2.1 Chemicals

Copper acetate (Cu(Ac)₂, 99.99%), indium acetate (In(Ac)₃, 99.99%), zinc acetate (Zn(Ac)₂, 99.99%), 1-dodecanethiol (DDT, 98%), 1-octadecene (ODE, 90%), oleic acid (OA, 90%), oleylamine (OAm, 97%), 11-Mercaptoundecanoic acid (MUA, 98%), potassium hydroxide pellets (>85.0 %) 3-mercaptopropionic acid (MPA, ≥99 %), copper (II) chloride dihydrate (≥99.99%), ammonia solution (30%), indium (III) chloride (99.99%), sodium sulfide, zinc acetate dihydrate (≥99%) and N-acetyl-cysteine (NAC, ≥99%) were purchased from Sigma-Aldrich. All chemicals were used without further purification.

2.2 Nanoparticle Synthesis

CIS-ZS core-shell QDs synthesised in organic solvent: The Zn stock solution was prepared before reaction by dissolving 0.367 g of Zn(Ac)₂ (2 mmol) in 5 ml of ODE/OAm (4:1 volume ratio) at 160°C under N₂ flow. Colloidal CIS core QDs were synthesised based on a

literature method.[29] 0.0584 g of $\text{In}(\text{Ac})_3$ (0.2 mmol) and 0.0363 g of $\text{Cu}(\text{Ac})_2$ (0.2 mmol) were added into a 50 ml three-neck flask with 2.0 mL of DDT, 3 mL of ODE and 0.3 mL of OA. The reaction mixture was heated to 100 °C for 10 min under N_2 gas flow to form a clear solution. Then, the temperature was raised to 260 °C in 3 min. The CIS QDs were allowed to grow. The colour of the solution changed from slightly yellow to yellow, red, and finally black. Once the solution turned black, indicating the bigger core of CIS QDs, the temperature was dropped to 100 °C. For ZnS shell growth, the synthesised CIS QD crude solution was directly used without purification. In a typical synthesis, the Zn stock solution was injected into the reaction solution. The temperature was raised to 230 °C and kept for 30 min. Then the solution was quenched. The particles were precipitated and washed by hexane/ethanol. The obtained CIS-ZS QDs were redispersed in chloroform. Ligand exchange was carried out by following a previously reported method [30], 0.5 g of MUA in 7 ml of chloroform was added to the 2ml of CIS-ZS core-shell QDs solution. 5ml of 0.1 M KOH solution was added and stirred overnight to let the QDs transfer from organic solution to water. Then the aqueous phase including the QDs was separated using a pipette and centrifuged at 6000 rpm for 2 min. Supernatant was decanted and the QDS were re-dispersed in water for further use.

Water-synthesis of CIS-ZS core-shell QDs: In a typical synthesis of the QDs, a mixture of 95 mL of deionised water, 2 mL of MPA (1.0 M), 1 mL of CuCl_2 (0.1 M), 0.4 mL of NH_4OH (30%), 0.5 mL of InCl_3 (1.0 M) and 1 mL of Na_2S (1.0 M) aqueous solutions were heated to 90-95 °C while aggressively stirred in a flask for 20 min. Then different amounts of $\text{Zn}(\text{Ac})_2$ (1.0 M) were added dropwise. The mixture was stirred for another 15 min and then left to cool down to the room temperature. In a typical post-synthesis treatment, where applicable, 0.3 ml MPA or 1.08 g of NAC was added to the as-synthesised solution.

2.3 Transmission electron microscopy

Transmission electron microscopy (TEM) was performed on a JEOL 2100 transmission electron microscope with a tungsten filament running at an accelerating voltage of 120 keV. TEM specimens were prepared by depositing one drop of the QDs solution on carbon-coated copper grids.

2.4 Optical spectroscopy

A PerkinElmer Lambda 35 ultraviolet-visible (UV-Vis) absorption spectrophotometer with a wavelength range of 190-1000nm (± 0.1 nm wavelength accuracy) was used to collect optical density of the solutions. The instrument was equipped with deuterium and tungsten halogen lamps and a Seya Namokia monochromator. PerkinElmer UV WinLab software (version 6.0.2.0723) was used for analysing the data. Luminescent characteristics of the prepared QDs were investigated using a Cary Eclipse fluorescence spectrophotometer equipped with two Czerny-Turner monochromators and a 15 W Xenon Flash Lamp, Schwartzchild collection optics and two photomultiplier detection tubes. The spectral range of this spectrometer is from the deep ultraviolet (UV) at 190 nm to near infrared (NIR) at 900 nm with an accuracy of ± 1.0 nm. The data were analysed with Cary Eclipse Scan Application software (version 1.2).

2.5 Collection and treatment of latent fingermarks

Substrates including glass slide, new \$5 Australian polymer banknote and aluminium drink can were washed with detergent

and deionised water and other substrates including black adhesive insulating tape, transparent sticky tape, aluminium foil and zip-lock bag were used in their new state. Donors both male and female aged between 20-50 years were asked not to wash their hands for at least half an hour before fingerprint collection. For collecting uncharged (natural) latent fingerprints, donors only rubbed their hands together and touched the substrates for approximately 5 seconds. For collecting charged latent fingerprints, donors were asked to first rub their hands on their face and hair and then rub them together. Fingermarks were kept in an office environment away from sunlight without any light protection. Fresh fingermarks were treated immediately after collection. The temperature and the relative humidity were monitored using a Digitech QP-6013 datalogger over the period of storage, producing recorded temperature and relative humidity ranges of 20-23 °C and 25-70% RH, respectively.

The collected latent fingerprints were immersed in the QDs working solutions. The working solutions were orbitally shaken on a PathTech Basic Orbital Mixer for different duration. The developed fingermarks were rinsed and shaken in a deionised water bath to remove excess QDs adhered to the substrate. Finally, the developed fingermarks were dried in an oven at 30°C for a maximum of 10 min.

2.6 Photography

Following development, developed latent fingerprints were photographed in a dark room using a Nikon D300 camera with an AF-S Micro NIKKOR 60 mm lens, a Tamron 62 mm Skylight 1A filter and an Anti-Glare Schott RG630 AG 1%(nom) deep red filter (long-pass 613 nm). The Nikon camera was connected to a desktop computer running Camera Control Pro 2 software (version 2.8.0) to use the live view and remote-control functions of the camera. The camera was mounted overhead on a Firenze mini repro stand in a darkened room. The photography specifications of the fingermarks photos were chosen to enhance contrast and visibility of the treated marks, and are specified in Table S1. Developed fingermarks on various substrates were illuminated with a forensic light source (Polilight PL500, Rofine, Australia). Photos were processed and adjusted in brightness using Adobe Photoshop CC (version 19.1.16) to make fingermarks details more visible. However, the same modifications were applied on the split fingermarks.

3.0 Results and Discussion

A preliminary investigation was carried out using CIS-ZS QDs synthesised in organic solvent which were then transferred to the aqueous phase by ligand exchange. The surfactant coating has a dual role in that it allows dispersion of the QDs in aqueous solution, as well as providing the capability of being attracted to the latent fingerprint residue. The aqueous solution of QDs gave promising results demonstrating that it is possible to develop fingermarks with CIS-ZS QDs on non-porous surfaces (see Figures S1 and S2 in ESI).

Synthesis of QDs in an organic phase produces uniform and brightly luminescent products. However, an organic synthesis method is complicated, time-consuming and expensive due to the tedious preparation steps, inert atmosphere condition, and high reaction temperature ($>200^\circ\text{C}$) [26, 31]. Also, the yield of QDs is very low (\sim tens of mg) and the required organic substances raise significant safety issues [32, 33]. Furthermore, as a post-synthesis treatment, the QDs must be transferred from organic to aqueous phase by ligand exchange to render them biocompatible and applicable to

fingermark development, which is a long process and may result in agglomeration and loss of luminescence [9]. We experienced all the aforementioned issues during the organic-synthesis and ligand exchange of the CIS-ZS QDs as well as their application to fingermarks development (Figure S1 and Figure S2). These drawbacks limit the usage of this method for application in an operational environment.

Preparation of QDs in water has the potential to be more rapid, simpler, more biocompatible and inexpensive. However, the surfaces of the as-prepared QDs have dangling bonds which are tentatively passivated by organic surfactants. Subsequent exposure of the synthesised QDs to an aqueous environment results in surface oxidation and a decrease in luminescence efficiency [31]. Previous studies have used nitrogen gas purging to attempt to solve this problem but this method cannot completely suppress the quenching effect [14, 32]. A more effective strategy, namely passivation of QDs with an inorganic shell made of other semiconductor materials, noticeably minimises the trap-state defects and increases the luminescent quantum yield and stability [34, 35]. Aqueous-synthesis of CIS-ZS QDs has been achieved using microwave-assisted [26] and hydrothermal methods [36], however these approaches suffer drawbacks including the necessity of special facilities and being a time consuming procedure. More recently, a convenient, fast and inexpensive water-based synthesis method with a possibility of upscaling has been established to produce CIS-ZS QDs in a safer way [37, 38]. In this approach, a mixture of copper, indium and sulfur precursors in water is heated at low temperatures (<100°C) to produce the Cu-In-S core, and then a zinc precursor is added to form the shell around the cores. We used this approach with modifications to produce CIS-ZS working solutions that could be used directly for the effective visualisation of latent fingermarks on different non-porous substrates without any further changes such as purification or pH and temperature adjustments.

Colloidal CIS-ZS core-shell QDs were synthesised in water in a simple one-step technique. First, aqueous solutions of CuCl_2 , InCl_3 , Na_2S were mixed in water in presence of a surfactant and ammonium solution. The reaction mixture was heated and kept at 90-95°C. Cu and In salts participate in a precipitation reaction with sodium sulfide and produce the Cu-In-S spherical nanoparticles [31]. Zn (II) precursor was added to the reaction mixture without any purification and the ZnS layers epitaxially grew on the cores as a shell producing a luminescent CIS-ZS QDs solution (Figures 1a-1c). This QD solution can be prepared within 90 min on a large scale (at least 400 mL of 0.96 M solution per batch). Figure 1d and Figures S3a-S3d show the TEM images of the synthesised CIS-ZS core-shell QDs. The size of the QDs was less than 5 nm. Absorption (Ab) and PL spectra of the CIS core and CIS-ZS core-shell QDs are illustrated in Figure 1e.

The synthesised QDs fluoresced at deep red to NIR (emission maximum at 660 nm) when excited in a wide range of wavelengths from UV to red region. The growth of ZnS shells around the CIS cores noticeably increased the PL intensity (approximately 15 times). Increasing the amount of Zn^{2+} precursor resulted in significant enhancement of the luminosity due to an increase of the thickness of the shell (Inset in Figure 1e); however, it led to a slightly blue-shift from 680 nm for the core to 660 nm for the final product. This is most probably related to the surface reconstruction rather than the diffusion of Zn into the CIS core taking into account the relatively low temperature of 90-95°C [27].

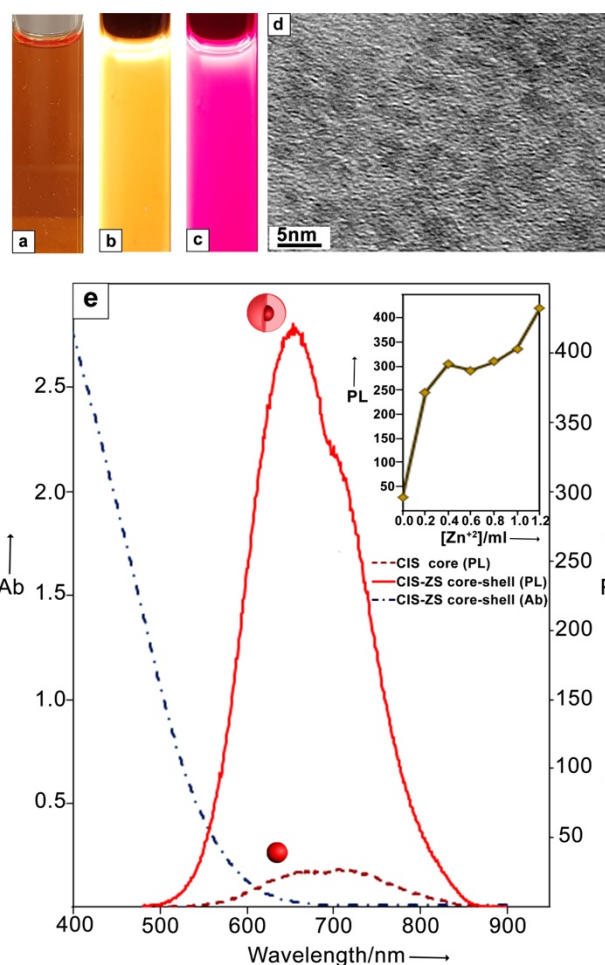


Figure 1. (a-c) CIS-ZS QDs solution photographed under (a) no illumination and (b and c) 350 nm illumination and through (a and b) no filter and (c) deep red filter (long-pass 613 nm). (d) TEM image of CIS-ZS QDs. (e) PL (excited at 350 nm) and Ab spectra of CIS core and CIS-ZS core-shell QDs. Inset in (e): maximum (excited at 350 nm) PL of CIS-ZS QDs with different amounts of Zn^{2+} precursor ($[\text{Zn}^{2+}]$).

Aging the prepared QDs solution under light for about one month, noticeably boosted the PL intensity (Figure S3e) due to the photodegradation of the surfactant leading to thickening of the shell, increasing the crystallinity and reducing the surface defects, as reported previously [32, 39]. Addition of more than 0.6 mL of Zn precursor led to the formation of a cloudy solution due to agglomeration and precipitation of the QDs (Figures S4). This unstable solution was undesirable as it hindered fingermark development, developing less ridge detail.

Initially 3-Mercaptopropionic acid (MPA) surfactant (the same surfactant used during the synthesis) was added to the prepared solution to stabilize the quantum dots as a post-synthesis step to prevent sedimentation by providing an electrostatic shield against inter-particle interactions. The long carbon chain and carboxylic group in the chemical structure of MPA was also proposed to increase both polar and non-polar interaction affinity with the components within the fingermark residue. Although only used in small amounts, MPA is a toxic and malodorous chemical. To increase the safety of the working solution alternative molecules were explored as capping ligands, including L-cysteine and N-acetylcysteine (NAC), to replace MPA in the post-synthesis step. Both are non-hazardous chemicals with NAC used for medical

purposes such treatment of paracetamol poisoning [40]. Xu et al. has previously demonstrated that NAC-capped QDs developed more ridge details of latent fingermarks compared to other ligands including MPA [15], fingermarks developed on transparent tape in our studies with NAC as the endcapping ligand consistently producing defined ridge detail and contrast when compared to MPA, with L-cysteine yielding no development of latent fingermarks (See Figure S5). This is suggested to be due to the variety of functional groups present in NAC, such as thiols, carbonyl and amide groups which enable interactions with lipids and amino acids present in fingermark secretions. Further studies are required to investigate the nature of the interaction of the quantum dots with the latent fingermark, as well as exploring other potential end-capping ligands.

A typical procedure for the treatment of latent fingermarks with CIS-ZS QDs solution involved immersing the sample in the solution and agitating for varying durations, with most substrates requiring no longer than 20 minutes. International Fingerprint Research Group (IFRG) guidelines relating to initial proof of concept (Phase 1) studies were followed to conduct the experiment [41]. A range of substrates including transparent, black and blue adhesive insulating tapes, glass slides, aluminium foil, aluminium drink cans and zip-lock bags were tested and developed fingermark ridge detail (see Figure 2). The sticky side of adhesive tapes demonstrated particularly promising results, Figure 2g shows excellent ridge detail including the sweat pores of a natural latent fingermark on a transparent sticky tape developed with the NAC-coated CIS-ZS core-shell QDs. To further investigate potential of the technique, the natural split fingermarks of 6 donors on transparent tape, aluminium foil and glass slides were developed. The results demonstrated full or partial development of the latent fingermarks from all 6 donors (Figure S6).

As a proof of concept, charged latent fingermarks on new Australian polymer banknotes as a nonporous multi-coloured substrate were treated with MPA capped CIS-ZS QDs, development produced marks well enough to distinguish high levels of ridge detail (Figure 3 b,d).

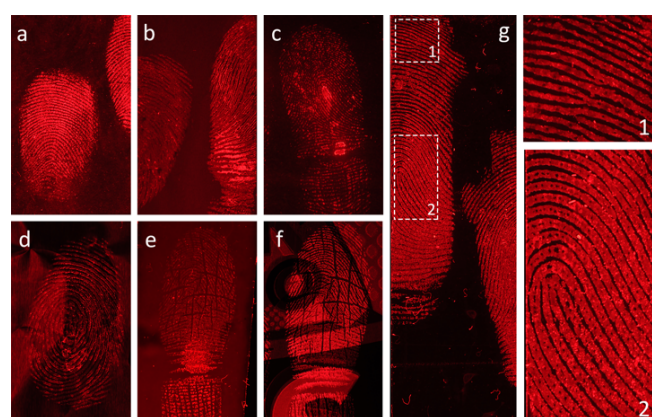


Figure 2. Fresh developed (a-d,g) natural and (e,f) charged latent fingermarks on (a) glass slide, (b) black adhesive insulating tape, (c) zip lock bag, (d) aluminium foil, (e) blue adhesive insulating tape, (f) aluminium drink can and (g) transparent adhesive tape. The marks were illuminated at 350 nm and pictured through the deep red filter. The marks were shaken at ((a-d and f-g) 20 rpm for (b)10 min, (a,c) 20 min, (d)15 min, (f) 2 hrs and (g) 5 min in CIS-ZS QDs solution, (e) the mark was treated with no shaking for 20 min,. The marks were illuminated at 350 nm and pictured through the deep red filter. See Table S1 for the photography conditions.

Being luminescent in deep red to NIR region made it possible to visualise the latent fingermarks on these multi-coloured and multi-patterned substrates. The ClZ-Zn QDs can be excited with a broad range of wavelengths. This is advantageous when working with inherently luminescent surfaces. Figure 3f shows that the security feature of a UV-fluorescent bird interfered with the fingermark on top of it. To visualise the hidden part of the fingermark, it was illuminated at 450 nm instead of 350 nm leading to the elimination of fluorescence from the bird feature (Figure 3g).

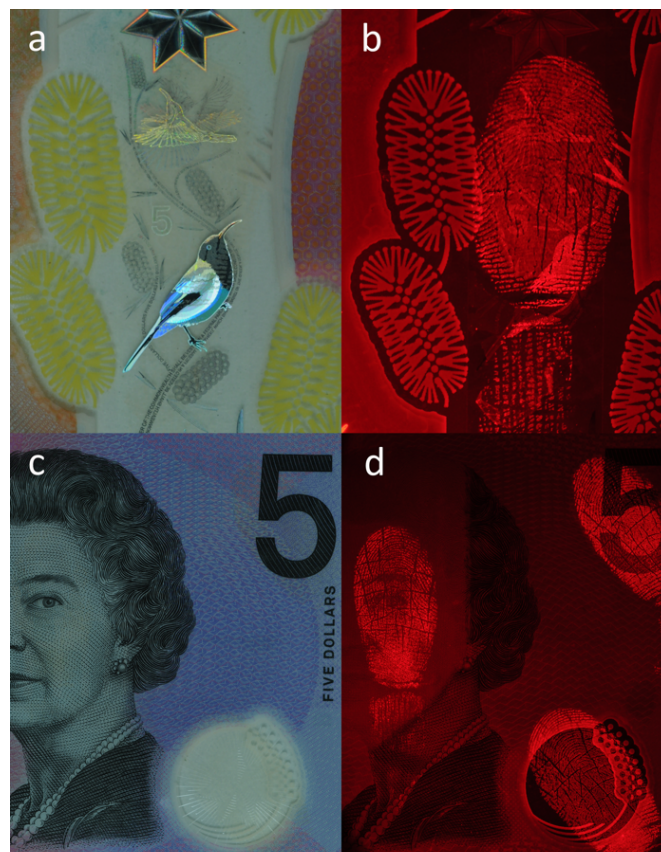


Figure 3. (a,c,e) Unilluminated and (b,d,f) illuminated at 350 nm and (g) 450 nm charged latent fingermarks (one female donor) deposited on Australian polymer banknote. The illuminated marks were pictured through a deep red filter. The marks were shaken at 20 rpm for 20 min in a MPA-coated CIS-ZS QDs solution. See ES1+ and Table S1 for the photography details.

Subsequent testing explored natural fingermarks on these substrates with NAC capped CIS-ZS QDs, Figure 4 demonstrates the level of ridge detail developed from 8 day old natural fingermarks. Whilst a larger number of donors showed poor development, the use of aged, natural marks gives a more realistic representation of fingerprint evidence, further investigation will continue to explore the use of CIS-ZS QD's on these surfaces, particularly on the role of the surfactant.

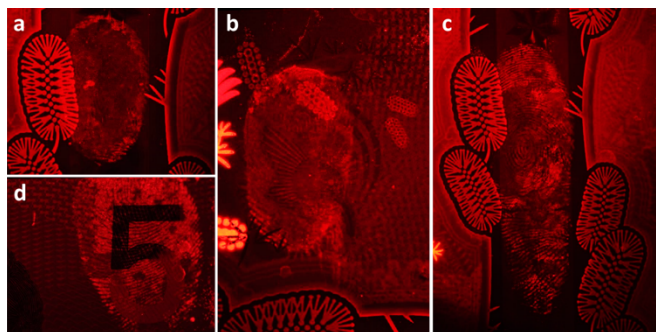


Figure 4. Treated (shaken at 20 rpm for 10 min in CIS-ZS QDs solution) 8-days-old latent fingerprints on Australian polymer banknote. The marks were illuminated at 350 nm and pictured through the deep red filter. See Table S1 for the photography conditions.

4.0 Conclusions

In conclusion, a safe innovative and straightforward procedure was developed for preparation and application of an aqueous working solution containing surface modified CIS-ZS core-shell QDs for visualisation of latent fingerprints. The surface modification consists of a surfactant coating which allows dispersion of the QDs in aqueous solution, as well as providing the capability of being attracted to the latent fingerprint residue. Latent fingerprints on various non-porous substrates were rapidly developed with high levels of ridge detail being observed. Being luminescent at deep red to NIR wavelength made it possible to eliminate the problem of dark, multi-coloured and multi-patterned backgrounds. However, issues remain in application to natural fingerprints on some challenging substrates such as polymer banknotes and cans. Further studies with a wider range of donors and substrates are required, as well as optimisation of the technique, including factors such as additional surface modification of the QDs, pH adjustment of the working solution, working solution stability/shelf-life and combination with other methods (such as cyanoacrylate fuming).

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Keywords: Latent fingerprints • quantum dots • photoluminescence • forensic science

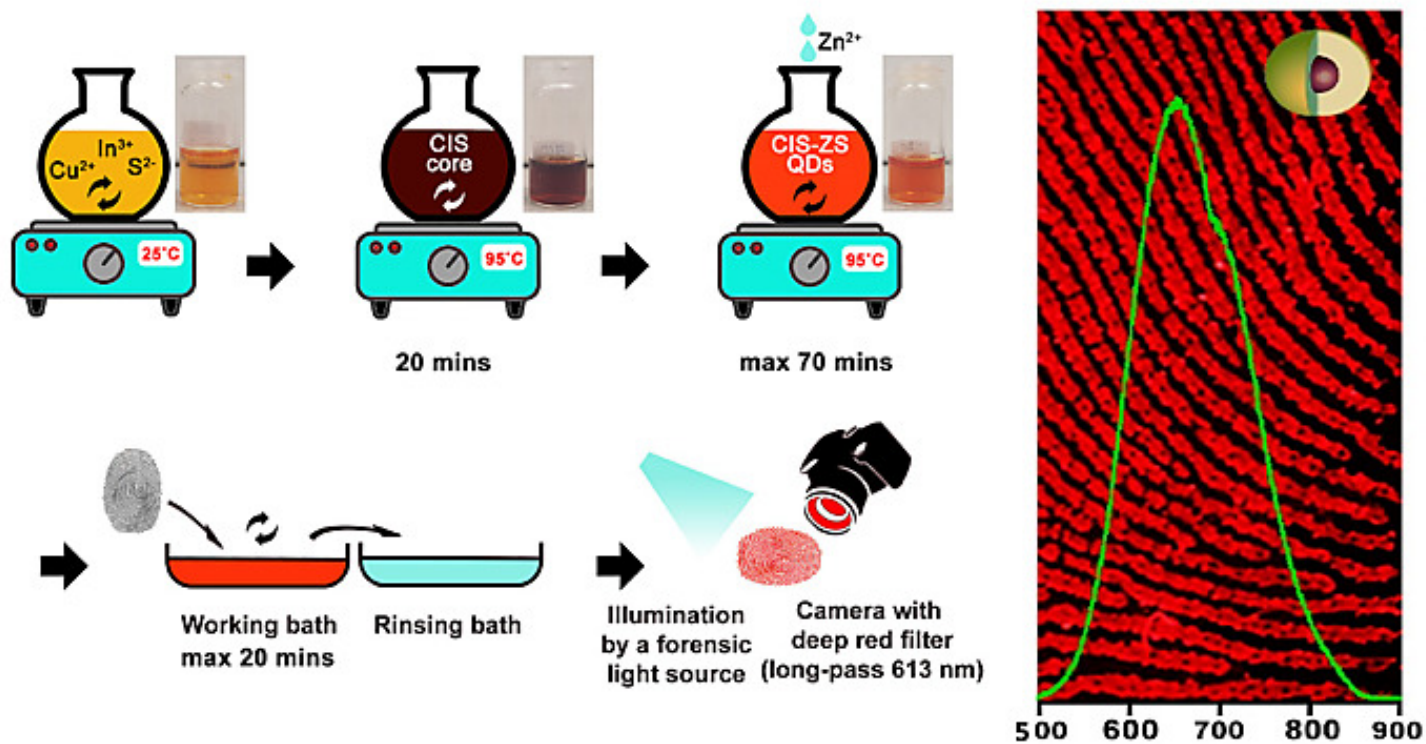
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Supplementary Information: Luminescence Detection of Latent Fingermarks on Non-porous surfaces with Heavy-Metal-Free Quantum Dots

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Table S1. Photography specifications of the photos taken from the latent fingermarks

Figure No.	F-stop	Exposure time (sec)	Figure No.	F-stop	Exposure time (sec)
2a	f/3.2	5	S2	f/13	5
2b	f/3.2	10	S5	f/3.2	6
2c	f/3.2	20	S6a	f/3.2	5
2d	f/3.2	10	S6b	f/3.2	10
2e	f/3.2	2	S6c-e	f/3.2	30
2f	f/3.2	1	S6f	f/3.2	20
2g	f/4	15	S6g-r	f/4	15
3a,c,e	f/11	3/5	S7a	f/3	8
3b	f/3.2	2	S7b	f/3.2	1
3d	f/3.2	8/5	S8a-b	f/3.5	5/2
3f	f/3.2	1	S8c	f/3.5	4
3g	f/3.2	1/8	S8d	f/3.5	6

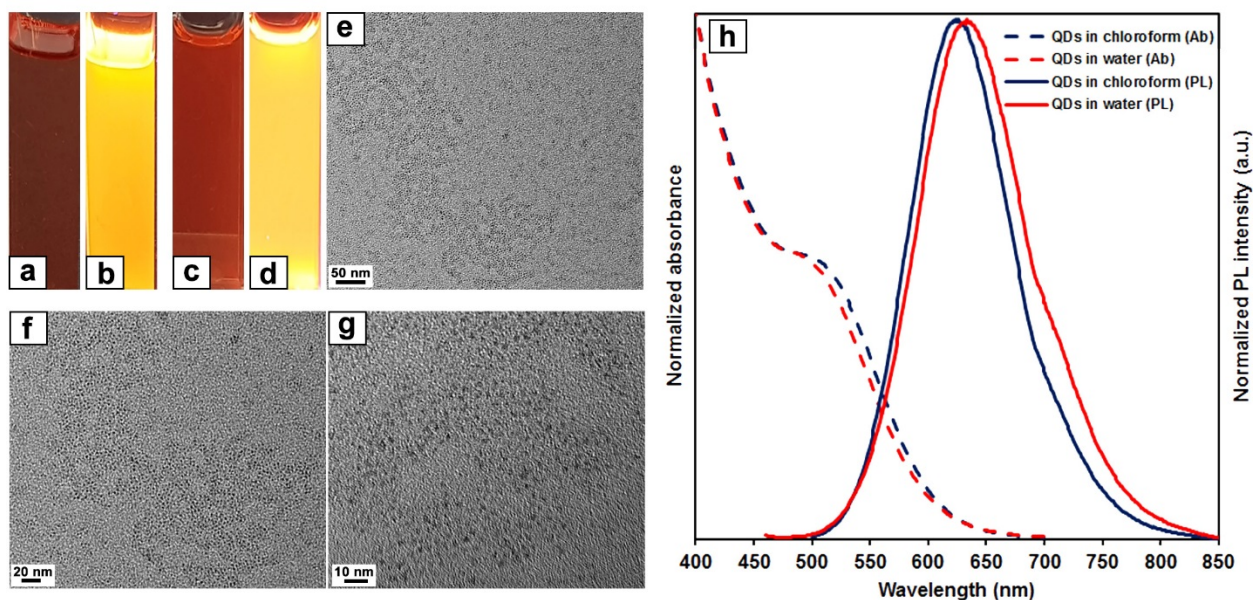


Figure S1. (a-d) CIS-ZS QDs dispersed in (a and b) chloroform and (c and d) in water and photographed under (a and c) no illumination and (b and d) 350 nm illumination. (e-g) TEM images of the CIS-ZS QDs in different magnifications. (h) PL (excited at 450 nm) and Ab spectra of the QDs solutions.

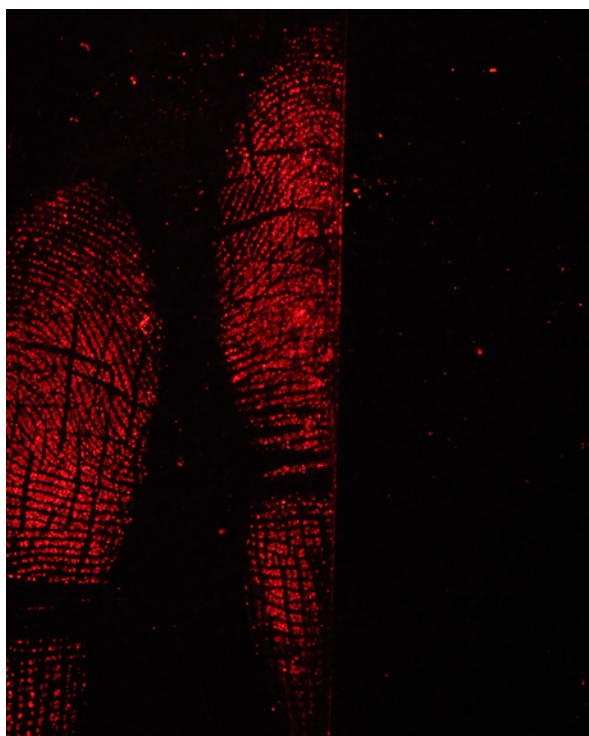


Figure S2. (a) Untreated (right side) and CIS-ZS QDs treated (left side) fresh charged split latent fingerprints (one female donor) on glass slides. The mark was illuminated at 350 nm and pictured through the deep red filter. See Table S1 for the photography conditions.

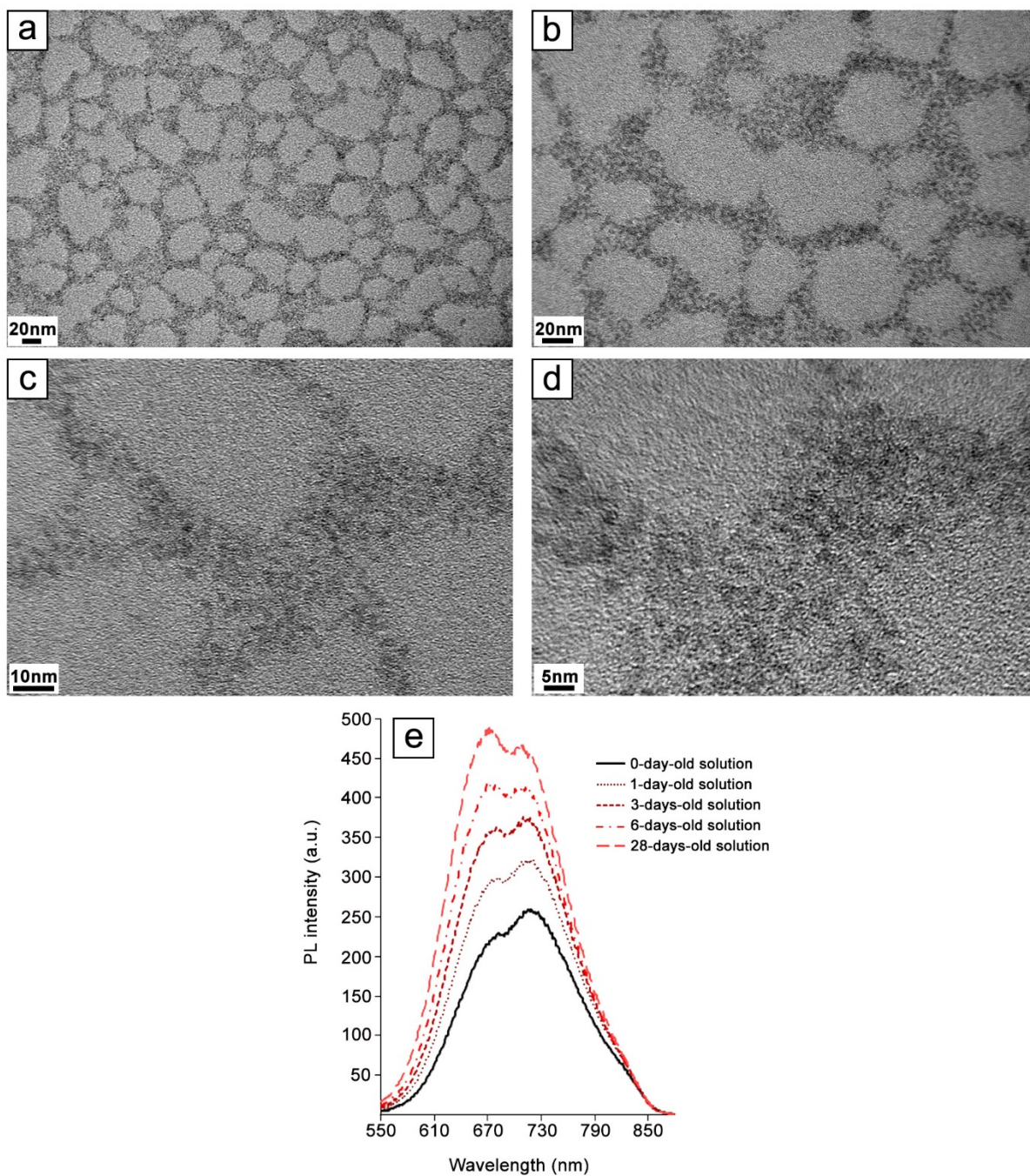


Figure S3. (a-d) TEM images of the CIS-ZS core-shell QDs synthesised in water in different magnifications. (e) PL spectra of fresh and old CIS-ZS QDs solutions. (f-g) CIS-ZS QDs solutions (f) before and (g) after addition of MPA. Inset in (f) is the precipitated QDs solution before the addition of MPA.

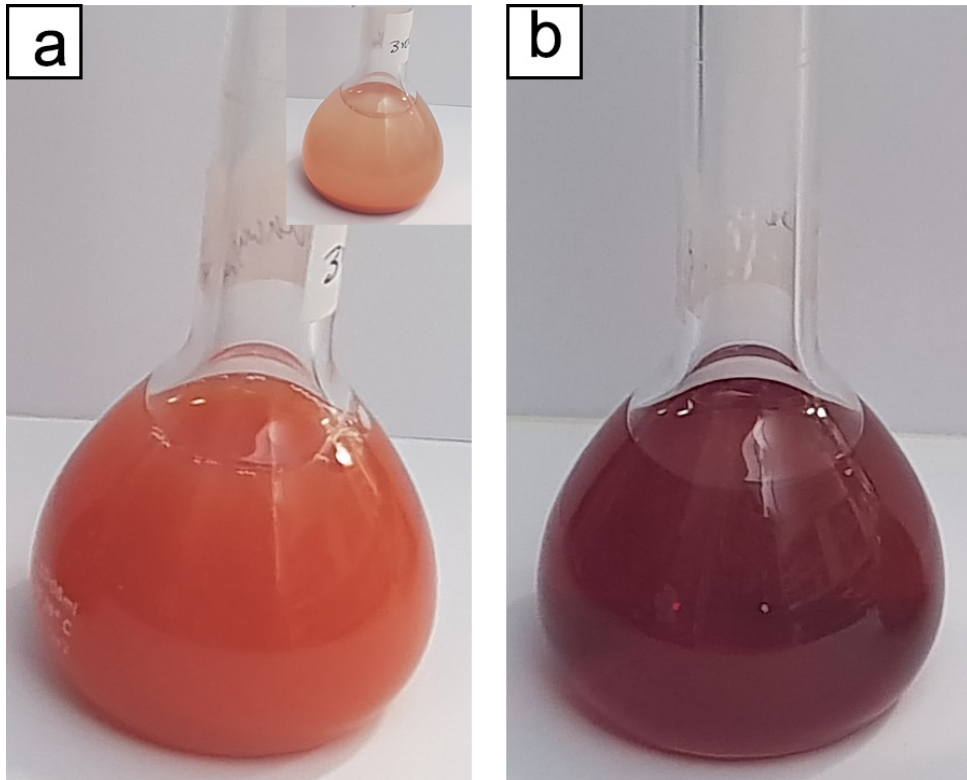


Figure S4. CIS-ZS QDs solutions (a) before and (b) after addition of MPA. Inset in (a) is the precipitated QDs solution before the addition of MPA.

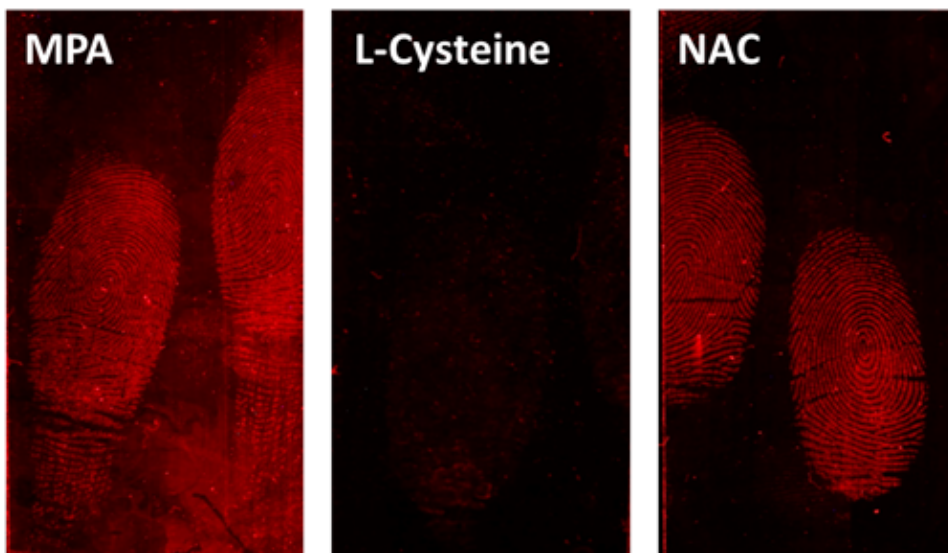


Figure S5 - Developed natural latent fingerprint (one female donor) deposited on a transparent sticky tape. Developed with MPA-coated CIS-ZS QD's (left), L-cysteine-coated CIS-ZS QD's (middle), NAC-coated CIS-ZS QD's (right) solutions. The marks were illuminated at 350 nm and pictured through the deep red filter.

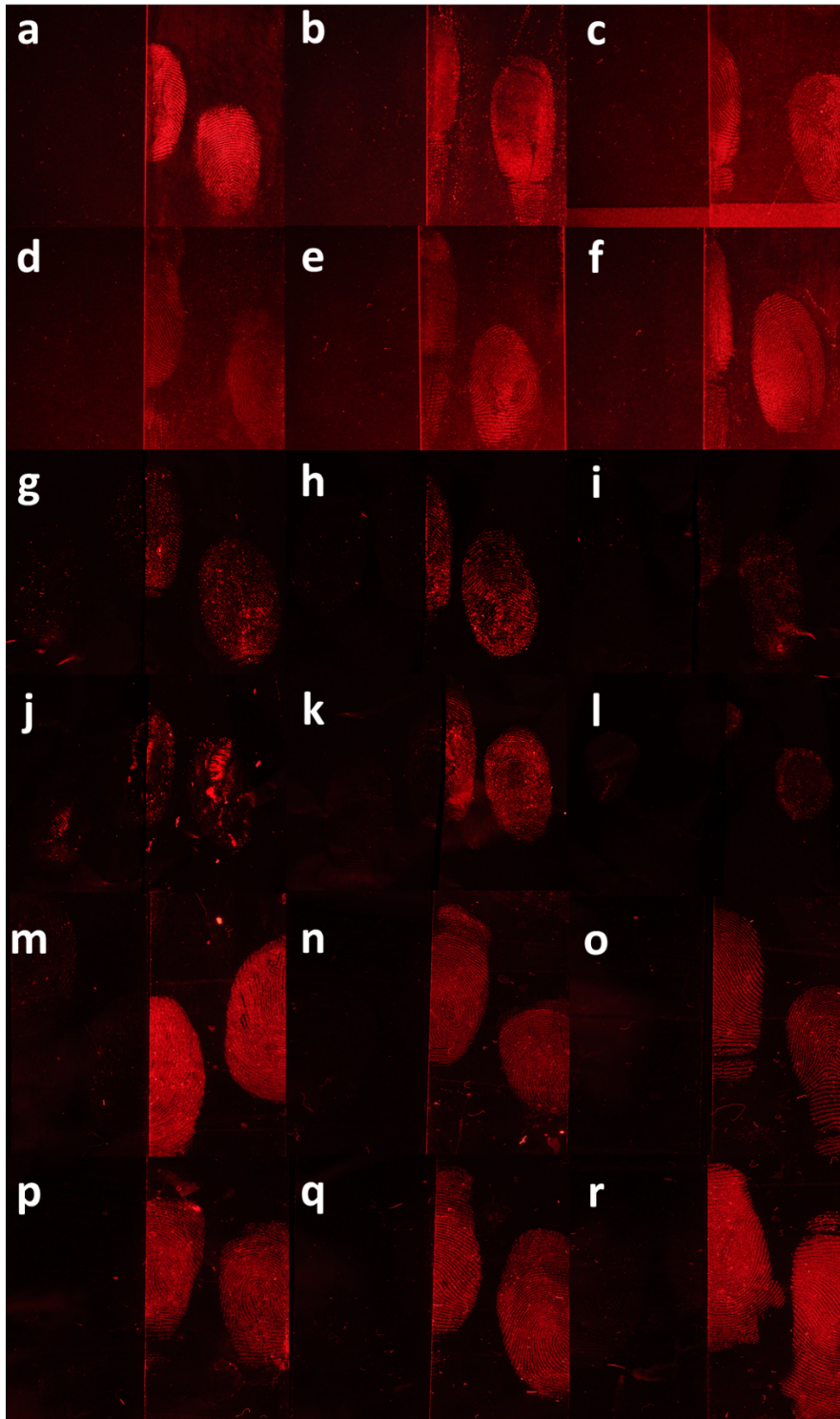


Figure S6. Untreated (left side) and treated (right side) natural split latent fingerprints (a-f) 1 day old on glass slides (g-l) d) 2 days old on aluminium foil (m-r) 3 days old on transparent adhesive tape, from a range of donors. The marks were illuminated at 350 nm and pictured through the deep red filter. See Table S1 for the photography conditions.

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