

# Detection of latent fingermarks on thermal printer paper by dry contact with 1,2-indanedione

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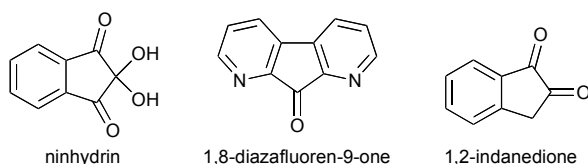
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A new method for the detection of latent fingermarks on thermal paper by dry contact with 1,2-indanedione, which reacts with the amino acids present to give coloured, photoluminescent prints is described. The simple and reliable method is based on contact between the thermal paper sample and reagent impregnated treatment papers. Different treatment paper reagent formulations and treatment times were investigated. The conditions which provided the best performance used treatment papers prepared from an acid-free fluoruous solution containing 1,2-indanedione and zinc chloride, with a contact time of 48 hrs. The dry contact approach was compared to current methods used by law enforcement agencies within Australia and was found to give similar or better performance.

**Key words:** latent fingermarks, fingerprint, 1,2-indanedione, thermal paper, dry contact

## Introduction

The impressions left by the ridge skin present on the grasping surfaces of the hands are extremely important in establishing contact and personal identification in criminal investigations.<sup>1,2</sup> Latent fingermarks, primarily composed of skin secretions (e.g. sweat), are invisible to the naked eye and thus require a development process to be visualised.<sup>1-4</sup> The principle approach for revealing latent fingermarks on porous surfaces (e.g. paper or cardboard) is chemical development with amino acid sensitive reagents.<sup>5</sup> This is essentially the trace detection of the amino acids present in skin secretions that have been deposited on a surface, where the position of the analyte within the matrix is important. Widely used reagents for this purpose include ninhydrin, 1,8-diazafluoren-9-one (DFO) or 1,2-indanedione (IND) (Figure 1), to form coloured products. DFO and IND have been termed “dual reagents” as their reaction products are both coloured and photoluminescent.<sup>6</sup> Dual reagents afford greater detection sensitivity and contrast due to their photoluminescent properties.<sup>5,6</sup>



**Fig.1** Chemical structures of amino acid sensitive reagents for the detection of latent fingermarks on porous surfaces.

Thermal paper is a heat sensitive paper most commonly used for sales receipts and fax papers, and can be of interest in cases of fraud and identity theft. It is different to most porous surfaces that are encountered by fingerprint examiners in that it is made up of several layers, each with a functional purpose.<sup>7</sup> The most important layers are the active coat, which

contains the chemicals for the printed image, and the top and back coats that seal the paper on which latent fingermarks may be deposited.<sup>7</sup>

Since its introduction, thermal paper has presented a challenge when using amino acid sensitive reagents to detect latent fingermarks. Samples are dipped into an organic solution containing the reagent (e.g. ninhydrin, IND), and depending on the procedure, additional components such as acid or metal salts.<sup>5,8</sup> After being air dried the sample may then be exposed to heat and/or humidity to develop the fingermarks as coloured impressions. Organic solvents, particularly polar solvents, react with the chemicals in the active coat of thermal paper causing irreversible damage to the sample and can obscure any printed text or developed fingermarks.<sup>7,9,10</sup> Additionally, the heating step commonly used in latent fingerprint development also results in discolouration of the paper.

Alternative methods have been proposed to overcome these problems. These include modification of a ninhydrin formulation to use a less polar solvent,<sup>7</sup> or the preparation of ninhydrin derivatives such as ninhydrin hemiketals, that minimise the concentration of polar solvents required for dissolution of the reagent.<sup>9,11</sup> Other methods have focused on whitening the background using cellophane/sticky tape (after a ninhydrin-based treatment),<sup>12</sup> or washing with acetone or solutions containing nitrogenous organic compounds (after treatments with either ninhydrin or DFO).<sup>9</sup>

Approaches which eliminate the requirement for a solvent have also been investigated. Treating samples by fuming with ninhydrin<sup>13</sup> or reagents such as dimethylaminocinnamaldehyde (DMAC),<sup>14</sup> iodine,<sup>15</sup> hydrochloric acid,<sup>16</sup> acetic acid,<sup>10</sup> and steam<sup>17</sup> have been reported. There has also been some research into the use of vacuum metal deposition<sup>17</sup> and controlled heating<sup>18</sup> as methods for latent fingerprint development.

Methods based on direct contact between thermal paper and

paper impregnated with reagent have been reported using ninhydrin and DMAC.<sup>19,21</sup> One technique, that has been termed ‘Nin-dry’, involves placing the sample between two pieces of dry ninhydrin-impregnated paper for a number of days in a sealed plastic bag.<sup>20,21</sup> This method requires no heat treatment and can afford good development of latent fingerprints with minimal damage to the document.<sup>21</sup> The DMAC contact method for thermal paper uses reagent impregnated sheets placed in a press for 24 hours.<sup>19</sup> This process developed fingerprints that were luminescent and left the printed text intact, but was deemed not as effective as treatment with DFO or ninhydrin.<sup>19</sup>

Of the aforementioned methods, only a few can produce photoluminescent fingerprints, primarily those treating with DFO solutions<sup>9,19</sup> and the DMAC contact method.<sup>19</sup> There has been little investigation into the more recently introduced reagent 1,2-indanedione (IND), with only two reported methods for the treatment of thermal papers,<sup>22,23</sup> both of which use wet, dipping treatments. The most recent method, recommended by the Australian Federal Police (AFP), involves dipping thermal paper samples into an acid-free IND and zinc chloride solution and then drying with no subsequent heat treatment.<sup>23</sup>

Here we present an investigation into a simple technique for the development of latent fingerprints on thermal papers using a dry contact method based on 1,2-indanedione, that produces photoluminescent fingerprints.

## Materials and Methods

### Chemicals

1,2-Indanedione (CASALI/Optimum Technology, Australia), anhydrous zinc chloride (BDH), dichloromethane (Mallingckrodt chemicals), ethyl acetate (Univar analytical, Australia), glacial acetic acid (CSR chemicals, Australia), absolute ethanol (CSR chemicals, Australia), HFE-7100™ (1-methoxynonafluorobutane, 3M Novec, Australia) and Petroleum spirit 60-80°C (APS chemicals, Australia) were used as received and were all analytical grade unless stated otherwise.

### Preparation of reagent solutions and treatment papers

Reagent solutions containing 1,2-indanedione and zinc chloride were prepared as recommended by the Australian Federal Police (AFP).<sup>23</sup> The two primary stock solutions were **IND/ZnCl<sub>2</sub>** (0.5 g 1,2-indanedione, 20 mg ZnCl<sub>2</sub>, 5 mL acetic acid, 0.5 mL ethanol, 15 mL dichloromethane and 30 mL ethyl acetate) and **acid-free/IND/ZnCl<sub>2</sub>** (0.75 g 1,2-indanedione, 20 mg ZnCl<sub>2</sub>, 0.5 mL ethanol, 15 mL dichloromethane and 35 mL ethyl acetate). Working solutions were prepared by dilution of the stock solution with either HFE-7100 or petroleum spirits, as per Table 1.

Four different dry contact treatment papers were prepared by dipping chromatography paper (Whatman No.1) in one of the four working solutions (Table 1) and allowed to air dry. These papers were used within 24 hours of preparation. For convenience we will refer to the treatment papers based on the working solutions used to prepare them.

**Table 1** Composition of working solutions.

Working solution	Stock solution	Diluent
<b>Fluorous IND/ZnCl<sub>2</sub></b>	5 mL <b>IND/ZnCl<sub>2</sub></b>	45 mL HFE-7100
<b>Non-fluorous IND/ZnCl<sub>2</sub></b>	5 mL <b>IND/ZnCl<sub>2</sub></b>	45 mL Pet. spirits
<b>Fluorous acid-free/IND/ZnCl<sub>2</sub></b>	5 mL <b>acid-free/IND/ZnCl<sub>2</sub></b>	45 mL HFE-7100
<b>Non-fluorous acid-free/IND/ZnCl<sub>2</sub></b>	5 mL <b>acid-free/IND/ZnCl<sub>2</sub></b>	45 mL Pet. spirits

### Collection of latent fingerprint samples

Latent fingerprint impressions were taken from a single donor, unless otherwise stated. When replicate latent impressions were required from a single donor, they were collected periodically over a 7 hour period, except for the depletion series. Donors were asked not to wash their hands immediately before collecting impressions, or to ‘charge’ their fingers by rubbing their fingers on their faces or hair. The donors were asked to rub their hands together prior to deposition of latent fingerprints and to touch the paper lightly. Latent fingerprint samples were treated between 1 and 7 days after collection.

For the initial investigations, whole latent fingerprints were collected on unprinted Thermal Register Rolls (Officeworks, Australia) and printed receipts from a number of supermarkets. These samples were then divided in two, and each half was developed with a different method. Investigations into dipping and air drying development methods were carried out on whole latent fingerprints. For investigations into the use of treatment papers in dry contact mode, whole latent fingerprints were collected on unprinted Thermal Register Rolls (Officeworks, Australia) and printed receipts. These fingerprints were then divided into quarters and each quarter was developed with a different treatment paper, or for a different length of time.

### Development of thermal paper samples using wet methods

Samples developed by dipping and heat treatment were dipped into a working solution, allowed to air dry, then heat treated in an Elna laundry press (initial experiments: *ca.* 160 °C, 10 s; later experiments: *ca.* 50 °C, 5 s). Samples developed by dipping without heat treatment were dipped into a working solution and then left in the fume cupboard until dry. These two methods represent the AFP methods for non-thermal and thermal papers respectively.

### Development of thermal paper samples using the dry contact method

Development of samples by the dry contact method using prepared treatment papers involved placing a sample between two pieces of dry treatment paper, sealing in a plastic zip-lock bag and leaving in the dark, unweighted, for a specific period of time. Samples were removed from the bag and photographed in absorbance and luminescence modes at 24 h intervals for up to 96 h unless otherwise stated.

### Testing the dry contact method for normal paper

Depletion series<sup>24</sup> on white photocopy paper (FUJI Xerox Professional, Australia) were collected from one male and one female donor. The depletion series were halved and one half was developed using the AFP method for non-thermal paper (dipping in **fluorous IND/ZnCl<sub>2</sub>** working solution followed by heat treatment) and the other half was developed with a modified dry contact method (the sample was placed between **fluorous IND/ZnCl<sub>2</sub>** treatment papers and subjected to *ca.* 160 °C for 10 s in an Elna laundry press).

### Comparison of the dry contact method to other development methods

A blind trial comparison was conducted, using four fingerprint donors (3 male and 1 female). Over the course of 6 hours the donors deposited a number of prints randomly across 20 unprinted thermal paper samples using varying amounts of pressure. A list of number and position of the deposits on each piece of paper was recorded and remained unsighted by the analyst until the samples had been developed and analysed. One set of 10 thermal paper samples were developed using the AFP method for thermal paper (dipping in **fluorous acid-free/IND/ZnCl<sub>2</sub>** working solution followed by air drying and no heat treatment); and the other set using the dry contact method described in this paper (using **fluorous acid-free/IND/ZnCl<sub>2</sub>** treatment papers), with a 48 h development time.

For a direct comparison between the various development methods, latent fingerprints from two donors on unprinted thermal paper samples were cut into quarters and treated with one of four visualisation methods: (i) the AFP method for non-thermal papers (dipping in **fluorous IND/ZnCl<sub>2</sub>** working solution followed by heat treatment); (ii) the AFP method for thermal paper (dipping in **fluorous acid-free/IND/ZnCl<sub>2</sub>** working solution followed by air drying and no heat treatment); (iii) the dry contact method described in this paper (using **fluorous acid-free/IND/ZnCl<sub>2</sub>** treatment papers); and (iv) a dry contact method using ninhydrin impregnated treatment papers (the 'Nin-dry' method), used by the Western Australian Police (WAPOL). The 'Nin-dry' method is as follows: a ninhydrin solution was prepared as detailed in the AFP workshop manual<sup>23</sup> and sprayed onto white photocopy paper (Green Wrap, Fuji Xerox, Australia) which was allowed to air dry. Samples were then placed between 'Nin-dry' treatment papers in a zip-lock plastic bag in a dark cupboard, unweighted. Samples were checked for fingerprint development at 24, 48 and 72 h.

### Photography of Samples

Samples were photographed in both absorbance (white-light) mode and luminescence mode using a Nikon D100 or D300 digital cameras. Illumination in absorbance mode was achieved using incandescent light bulbs with no camera filter attachments. Illumination in luminescence mode was achieved using a Rofin Polilight® PL500 (Rofin, Australia), with an excitation wavelength of 505 nm and an orange camera filter attachment (550 nm barrier filter). The camera settings for all photographs were as shown in Table 2.

**Table 2** Photographic conditions for absorbance and luminescence mode photographs, unless otherwise stated.

	Absorbance mode	Luminescence mode
Focal Length (mm)	60	85
Exposure Mode	Manual	Manual
White Balance	Auto	Auto
Shutter Speed (s)	1/25	4
Aperture	f/11	f/11
Sensitivity	ISO 200	ISO 200

### Visual analysis of developed fingerprints

In this study, a system based on that used by the Home Office Police Scientific Development Branch (HOPSDB) UK,<sup>24</sup> consisting of four levels was used to grade the development of fingerprints (Table 3).

**Table 3** Latent fingerprint development grading system used in this study.

Grade	Description
0 No development	No visible ridge detail in colour or luminescence
1 Weak development	Patchy ridge detail development with less visible colour or luminescence
2 Medium development	Mostly continuous ridge detail development with visible colour or luminescence
3 Strong development	Continuous ridge detail development with readily visible colour or luminescence

## Results and Discussion

### Preliminary considerations

Fingerprints are not absolutely reproducible and so it is important to establish a system of assessing the quality of developed fingerprints that takes into account their variable nature. Variability in the quality of developed fingerprints can arise due to: different skin secretion compositions, the presence of contaminants on the hands; and variation in the pressure applied when leaving the deposit.

Two complementary methods can be used to directly compare the performance of fingerprint development reagents. These are the 'split print' and depletion series approaches. The 'split print' approach is where a single latent fingerprint is divided into two or more sections, so that each section can be treated differently. The depletion series approach is where a series of latent fingerprints are deposited one after the other, without the donor's fingers touching anything between each deposition, so as to give a series of prints that have decreasing amounts of deposited material.<sup>24,25</sup>

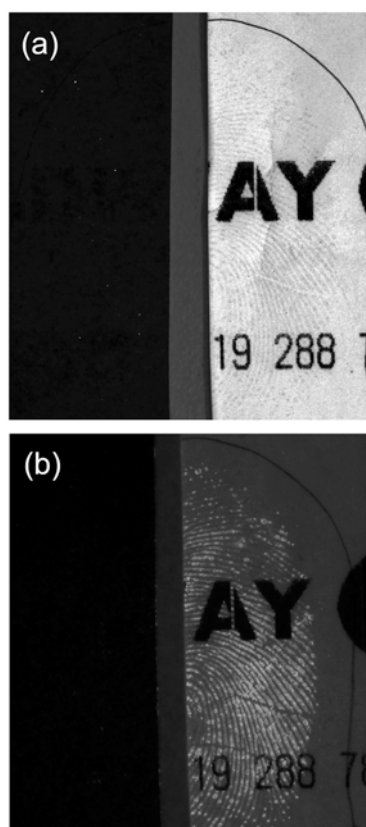
Both approaches have been used in this study.

### Method of applying development reagents

Preliminary experiments were carried out to determine whether 1,2-indanedione would develop latent fingerprints on thermal paper through a dry contact process. Latent fingerprints were deposited on thermal paper which was placed between sheets of 1,2-indanedione impregnated paper and sealing in a plastic zip-lock bag and left overnight in the dark. Coloured, photoluminescent fingerprints were successfully developed with all samples.

Following these successful preliminary studies the dry contact method was compared to other approaches for application of 1,2-indanedione/zinc chloride including dipping and heating (as per the AFP recommendations for non-thermal papers<sup>23</sup>), and dipping and air drying (as per the AFP recommendations for thermal paper<sup>23</sup>). The techniques recommended by the AFP are widely used throughout law enforcement in Australia as standard methods for the development of latent fingerprints.

Initial investigations into the different methods of applying 1,2-indanedione/zinc chloride reagents focused on the comparison between the dry contact method (using **fluorous IND/ZnCl<sub>2</sub>** treatment papers) and the AFP method for non-thermal papers (dipping in **fluorous IND/ZnCl<sub>2</sub>** working solution and subsequent heat treatment). These experiments showed that when a thermal paper sample, printed or unprinted, is dipped and heated at *ca.* 160 °C significant blackening of the background occurs obscuring both latent fingerprints and printed evidence. However, the dry contact treatment developed fingerprints that were visible in absorbance and photoluminescence imaging modes (Figure 2). Decreasing the heating to *ca.* 50 °C for 5 seconds still resulted in discolouration of the background.



**Fig.2** Latent fingerprints on printed thermal paper, where halves were treated with fluoruous IND/ZnCl<sub>2</sub> using (on the left) the AFP method for non-thermal papers (dipping with heating) or (on the right) the dry contact method. Photographs taken with a Nikon D300 Camera in (a) absorbance mode or (b) luminescence mode with settings; Focal length: 60 mm, Shutter Speed: 2.5 s and Aperture: f/14.

When samples were dipped into working solutions and air dried, without heat treatment, discolouration also occurred. The extent of this varied depending on the time the sample was left in solution and the formulation of the development solution. In terms of discolouration of the thermal paper, the non-fluorous solutions (petroleum spirits based) were more detrimental than fluorous development solutions (HFE-7100 based). This was also the case with solutions containing acid. It is important to note that the discolouration caused by dipping without heating was significantly less than when the samples were dipped and heated.

In comparison the thermal paper samples developed by the dry contact method developed slowly but without any darkening of the background.

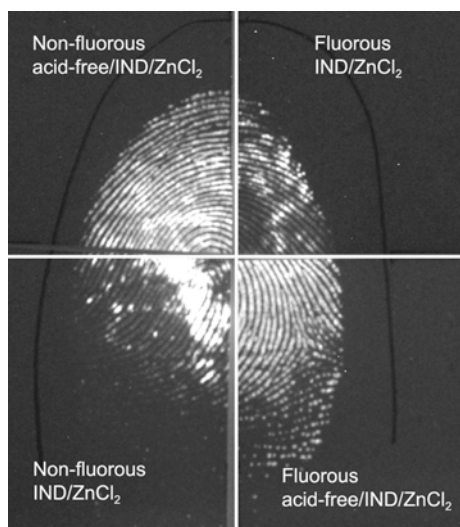
#### Investigations into reagent formulation for dry contact treatment papers

Since acidic components of development solutions had a detrimental effect on the active layer of thermal paper in the wet dipping methods, it was postulated that residual acid in the treatment papers may also be problematic. Working solutions, used to prepare treatment papers, containing no acid were investigated (e.g. **fluorous acid-free/IND/ZnCl<sub>2</sub>**, Table 1). While the AFP recommend the use of the fluorous solvent HFE-7100 for development solutions, petroleum spirits was also investigated as it is a less expensive alternative to HFE-7100.

On samples of unprinted thermal paper, fingerprint quarters developed using treatment papers prepared from acetic acid-containing working solutions (**fluorous IND/ZnCl<sub>2</sub>** and **non-fluorous IND/ZnCl<sub>2</sub>**) often showed a more intense colour and luminescence than those without acid (**fluorous acid-free/IND/ZnCl<sub>2</sub>** and **non-fluorous acid-free/IND/ZnCl<sub>2</sub>**). However, the development was patchy and uneven across the quarter, and in some cases there was no development at all, averaging a development grade from 0-1 (as per Table 3). While the fingerprints developed using treatment papers prepared from solutions containing no acid (**fluorous acid-free/IND/ZnCl<sub>2</sub>** and **non-fluorous acid-free/IND/ZnCl<sub>2</sub>**) often had less intense colour and luminescence, there was more consistent development of the ridge patterns of the fingerprint, with an average grade from 2-3 (see Figure 3). These results indicate that the inclusion of acid in visualisation solutions does have a detrimental effect on fingerprint development on thermal paper. There was no notable difference in development of fingerprints between treatment papers prepared from fluorous or non-fluorous working solutions.

On printed thermal paper samples similar observations were made. Fingerprint quarters developed with **fluorous acid-free/IND/ZnCl<sub>2</sub>**-based treatment papers often demonstrated brighter luminescence than quarters developed with the other treatment papers. Of the four different working solutions investigated, the treatment papers prepared with **fluorous acid-free/IND/ZnCl<sub>2</sub>** (Table 1) generally developed more fingerprint quarters of greater quality (grade 2 and 3) than the treatment papers prepared with the other working solutions. It was also found that petroleum spirit was a

suitable replacement for HFE-7100, however, the quality of the developed fingermark is decreased. It should be noted that when the printed text overlapped the fingermarks, the ridge detail was obscured in both absorbance and luminescence mode.



**Fig.3** Latent fingermark on unprinted thermal paper where quarters were treated with visualisation solutions using the dry contact method.

Photographs were taken using a Nikon D100 Camera in luminescence mode with settings; Focal length: 85 mm, Shutter Speed: 4 s and Aperture: f/11.

#### Investigations into development time

When treating non-thermal papers, heat is used to speed up the reaction between the amino acids in the latent fingermark and 1,2-indanedione. For thermal paper the application of heat to shorten development time is not feasible. Therefore the development time would be a compromise between quality of developed fingermarks and length of development period.

Latent fingermark quarters on unprinted and printed thermal papers were developed using **fluorous acid-free/IND/ZnCl<sub>2</sub>**-based treatment papers, with treatment times of either 24, 48, 72 or 96 hours (Figure 4).



**Fig.4** Latent fingermark on printed thermal paper where quarters were treated for different development times with fluorous acid-free/IND/ZnCl<sub>2</sub> by the dry contact method. Photographs were taken using a Nikon D100 Camera in luminescence mode with settings; Focal length: 85 mm, Shutter Speed: 2.5 s and Aperture: f/11.

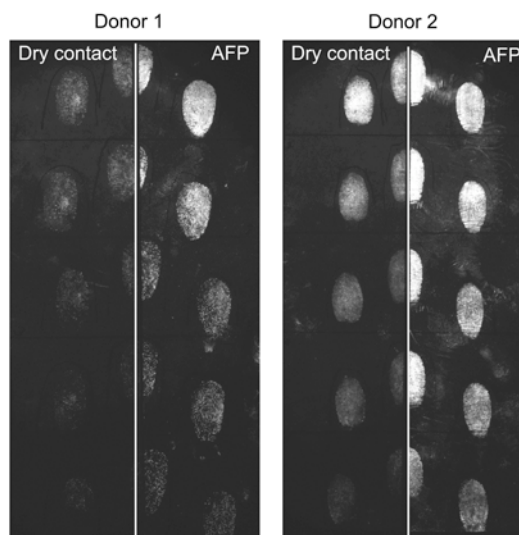
On both unprinted and printed thermal papers, strong prints showed sufficient colour and luminescence after 24 hours with no noticeable increase in development at longer treatment times. However, weak fingermarks required longer development times, with sufficient development observed after 48 hours.

While there was no noticeable increase in performance after 48 hours, samples observed after 78 days of development revealed fingermarks with significantly darker colour, more intense luminescence and ridge detail. While operationally, it is not feasible to leave exhibits in treatment for 78 days, it can be concluded that leaving thermal paper exhibits in treatment for extended periods of time does not have an adverse effect.

#### The dry contact method for non-thermal papers

An investigation into the effectiveness of the 1,2-indanedione dry contact method for non-thermal papers compared to the AFP method for non-thermal papers (dipping the sample into the **fluorous IND/ZnCl<sub>2</sub>** working solution followed by heat treatment) was carried out. A solvent free method for the treatment of fingermarks on these surfaces would be beneficial particularly in cases where police are required to travel internationally with forensic equipment as the air transportation of hazardous materials is restricted. A dry reagent method would also be useful for samples that are fragile and have the potential to be damaged by standard development techniques.

Results of this preliminary experiment showed that for a depletion series of fingermarks on white photocopy paper the dry contact method using **fluorous IND/ZnCl<sub>2</sub>** treatment papers, with additional heating, developed fingermarks that were visible in both absorbance and luminescence mode (Figure 5).



**Fig.5** A depletion series from two donors on white photocopy paper, where halves were treated with fluorous IND/ZnCl<sub>2</sub> using the dry contact method with additional heating or the AFP method for non-thermal papers (dipping and heat treatment). Photographs taken using a Nikon D300 Camera taken in luminescence mode with settings; Focal length: 60 mm, Shutter Speed: 1/2.5 s and Aperture: f/14. A full size version of Fig.5 is available in the Electronic Supplementary Information.

Halves treated with the AFP method (dipping, **fluorous IND/ZnCl<sub>2</sub>**, heat treatment) developed greater colour and luminescence than halves treated using the dry contact method. This is not unexpected as the fingerprints are exposed to direct contact with a greater concentration of the development reagents in the wet dipping methods, than in the dry contact method. The reduced luminescence of samples treated using the adapted dry contact method may be due to the physical properties of the paper itself. Paper has a fibrous surface, consisting of 'hills' and 'troughs', the uneven nature of the surface may limit the amount of contact between the sample and the treatment papers, thus reducing the methods ability to develop fingerprints. It should be noted that the dipping method results in noticeably higher background luminescence compared to the dry contact method.

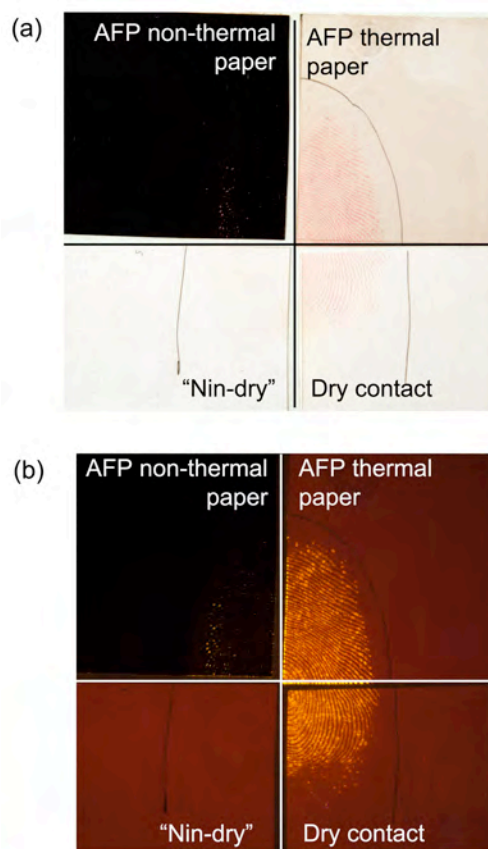
While the dry contact method could not develop fingerprints of equal luminescence and colour to the AFP method for non-thermal paper, it did develop fingerprints of reasonable quality when viewed in luminescence mode.

### Comparison of the dry contact method to other development methods

A blind trial was used to compare the performance of the dry contact method with the standard method for thermal papers. Both the AFP thermal paper method (dipping in **fluorous acid-free/IND/ZnCl<sub>2</sub>** working solution followed by air drying and no heat treatment) and the dry contact method (using **fluorous acid-free/IND/ZnCl<sub>2</sub>** treatment papers) developed a similar proportion of fingerprints in luminescence and absorbance mode. However, in absorbance mode it was easier to see fingerprints developed using the dry contact method as there was no discolouration of the background. This experiment showed that both the AFP method for thermal paper and dry contact method were equally effective in developing latent fingerprints on thermal paper.

A direct comparison of the dry contact method with the AFP methods for non-thermal and thermal papers, and the 'Nin-dry' method was also performed using thermal paper (an example is shown in Figure 6). As expected, quarters treated with the AFP method for normal paper (dipping in **fluorous IND/ZnCl<sub>2</sub>** working solution followed by heat treatment) showed blackening of the paper with no development (grade 0) of the latent fingerprints visible in absorbance mode, and weak to medium development (grades 1-2) in luminescence mode. Quarters treated with the AFP method for thermal paper (dipping in **fluorous acid-free/IND/ZnCl<sub>2</sub>** working solution followed by air drying and no heat treatment) showed some discolouration of the background with the majority of latent fingerprints developing to grades 1 and 2 in absorbance mode and grade 3 in luminescence mode. The dry contact method (using **fluorous acid-free/IND/ZnCl<sub>2</sub>** treatment papers) consistently developed fingerprints that were grade 3 in both absorbance and luminescence viewing modes. The 'Nin-dry' treated quarters were checked for development at 24, 48 and 72 hours, with the majority of samples showing grade 2 development in absorbance mode only after 72 hours of treatment. There was no visible luminescence from 'Nin-

dry' treated quarters, as expected for a ninhydrin-based development. The development of quarters using the 'Nin-dry' method was compared to the 1,2-indanedione dry contact method at 24 hours, and it was observed that the 1,2-indanedione dry contact method had developed all fingerprints to a better standard than the 'Nin-dry' method.



**Fig.6** Latent fingerprints on unprinted thermal paper where quarters were treated using the AFP method for non-thermal paper (dipping in fluoruous IND/ZnCl<sub>2</sub> working solution followed by heat treatment), the AFP method for thermal paper (dipping in fluoruous acid-free/IND/ZnCl<sub>2</sub> working solution followed by air drying and no heat treatment), the 1,2-indanedione dry contact method (using fluoruous acid-free/IND/ZnCl<sub>2</sub> treatment papers) or 'Nin-dry' (a ninhydrin-based dry contact method). Photographs taken with a Nikon D300 Camera in (a) absorbance mode with settings; Aperture: f/16 and Shutter Speed: 1/10 s and (b) luminescence mode with settings; Focal length: 60 mm, Aperture: f/3.2 and Shutter Speed: 1/30 s.

### Conclusions

These studies indicate that the 1,2-indanedione dry contact method described here is a simple and reliable, non-destructive method for the development of latent fingerprints on thermal paper. When compared to the 'Nin-dry' method, the 1,2-indanedione dry contact method developed fingerprints to a darker colour, that were also luminescent, in a shorter period of time. The dry contact method appears to be comparable to the current AFP recommended (dipping without additional heat treatment) method for treating thermal paper with 1,2-indanedione (24 hours), without damaging the paper.

This method shows promise for non-thermal documents that are fragile or for cases where international travel with forensic equipment is required. A larger scale operational trial in collaboration with a number of police fingerprint bureaux throughout Australia is currently being planned. This will include investigations into the use of the 1,2-indanedione dry contact method with aged samples (older than one week).

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## Notes and references

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<sup>†</sup> Electronic Supplementary Information (ESI) available: A full size version of Fig.5. See DOI: 10.1039/b000000x/

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