Spectroscopic and Photophysical Studies of Fingerprint Dusting Compositions

Keywords: Fingerprints; Fluorescence; Fourier transform infrared spectroscopy; Photophysical parameters; UV-visible spectra

Abstract

The two fingerprint dusting compositions considered here were based on fluorescent brilliant blue R and acridine orange stains. The bonding material in each case was a combination of silica gel and hydrated magnesium silicate. Spectral and photochemical studies revealed that the compositions were not merely physical mixtures of bonding material and colorant but the constituents interact chemically with each other, profoundly affecting the quality of developed prints.

Introduction

Regular fingerprint powders consist of coloured particles, each of which is bonded with a resinous polymer; the coloured particle provides contrast and the polymer coating provides adhesion [1]. It is commonly perceived that in a fingerprint dusting compositions, the two components act independently of each other. The adhesive gets adhered to the moisture and oily component of sweat residue by the mechanism of which are treated by G.L. Thomas, while the colorant gets adsorbed on the adhesive [2].

Luminescent powders are advantageous as these can develop prints with a good contrast on multicoloured surfaces [3,4]. Moreover, these assist detection of weak, accidental finger impressions that are often found at crime scenes.

The application of finely divided material and subsequent removal of the excess powder by brushing, blowing or tapping has been the universal method of developing fingerprints on an array of surfaces [5]. The technique relies on the chemical adherence of fingerprint powder to the moisture and oily components of skin ridge deposit [6].

In continuation to our interest in investigating the application of fluorescent dusting formulations to detection of latent fingerprints [7,8], we reported in this communication, two powder compositions incorporating a bonding material mixture of silica gel (major) and hydrated magnesium silicate (minor). Brilliant blue R is the colorant in one composition, acridine orange in the other. On the basis of spectral and photophysical studies, we concluded that the mixture and the colorant were not independent entities. These were interacted with each other, as a result of which the quality of developed fingerprints was profoundly affected.

Experimental Methods

Brilliant blue R, acridine orange and silica gel (mesh size 100-200) were procured from Sigma-Aldrich and were used without further purification. Commercial grade hydrated magnesium silicate was used.

The following instruments were used: SPECTRUM RX I (Perkin Elmer), collecting eight scans from 400-4000 cm⁻¹, with a resolution of 4 cm⁻¹ for Fourier Transform Infrared studies; Shimadzu Model UV 1600 spectrophotometer for recording Ultraviolet Visible spectra; Jobon Yvon Horiba Fluorolog Fluorimeter for fluorescence studies.

The curve for fluorescence spectra was obtained by setting the excitation wavelength to be the absorption maximum. Corrected fluorescence spectra of the two soluble dyes were obtained using a high pressure Xenon flash lamp as excitation source. The light path was 1 cm in all spectral recordings. Rhodamine B in ethanol (Quantum yield for fluorescence, Φ = 0.65) was used as a standard. All spectra were recorded in ethanol at a concentration of about 10⁻⁴ mol/L.

Twenty five students and faculty members of our group, in the age group of 20 to 50 years, with variable donor capabilities were asked to impinge latent fingerprints on different porous and non porous surfaces by mild pressure, without prior washing and cleaning of hands. This ensured practical variability in the print samples. The entire project was conducted in winter season when the temperature was 18 – 20 °C and relative humidity 30 – 75%.

Surfaces considered

The compositions were tested on both absorbent (porous) and non-absorbent (non-porous) items. The absorbent surfaces used were xerox paper, newspaper and glossy paper. The non-absorbent surfaces were glass, lamination sheet and metal.

The quality of developed fingerprints was gauged on the basis of ridge clarity and degree of fluorescence, on a scale ranging from zero to 10, according to the criteria generally followed in India [9], as summarized in Table 1.

Procedure: To a mixture of silica gel (4.0 g) and hydrated...
Depletion studies were carried out on glossy paper.

**Results and Discussions**

Using the novel formulations, latent fingerprints were developed on a broad spectrum of absorbent and non-absorbent items. Representative cases on Photostat paper and metallic surface were depicted in Figures 1 and 2 respectively. Figure 2 showed a representative fingerprint developed on a metallic surface by Brilliant blue R and acridine orange based compositions. Figure 3 showed the latent fingerprints under fluorescence light by Brilliant blue R based composition.

Depletion studies to test the validity of brilliant blue R based composition were carried out on a plastic item. The finger was wiped on the forehead and four impressions were impinged in succession. The developed fingerprints were illuminated with radiation having wavelength of 505-550 nm. When observed through red goggles, the fingerprints showed fluorescence.

The quality of developed fingerprints was observed on the basis of ridge clarity and degree of fluorescence, on a scale ranging from 0 to 10 (Table 1). On an average the print develop by brilliant blue R based composition were assigned a value of 8 while those developed by acridine orange based composition were assigned a grading of 6. Graph 1 showed an assessment of the effectiveness of two fluorescent powder compositions for latent fingerprint development on different substrates.

**Fourier transform infrared spectra**

The infrared spectra obtained by grounding each formulation with KBr powder (Infrared grade by Merck) and made pellets with the help of hydraulic press, maintained five tonnes of pressure for few seconds. A comparison of FTIR spectra of the pure silica gel, SG-B and SG-A with the vibrational modes of the major peaks was shown in Figure 4. In the infrared spectra of pure silica gel, a prominent and intense absorption occurred at 1050 cm<sup>-1</sup> due to (Si-O-Si) antisymmetric vibrations. In the FT-IR spectrum, this absorption split up in a shoulder peak at 1083 cm<sup>-1</sup> and a sharp intense peak at 1037 cm<sup>-1</sup>. The former was assigned to the longitudinal optical mode of Si-O-Si vibrations, the latter to the transverse optical mode [10]. Both these vibrations undergo perturbations in the fingerprint compositions.
The longitudinal optical mode of Si-O-Si vibrations absorbed at 1053 cm\(^{-1}\), while transverse optical mode absorbed at 1018 cm\(^{-1}\). Moreover, the (Si-O-H) stretching vibration, which occurs at 965 cm\(^{-1}\), completely disappeared in the novel compositions, indicating deprotonation. The absorption band at 670 cm\(^{-1}\), observed in the formulations, was attributed to (Mg-O) stretching vibrations, indicating the presence of talc as one of the adhesive materials.

**Ultraviolet visible spectra**

All the powder formulations and their individual stains were dissolved in ethanol for ultraviolet visible studies. The brilliant blue R and acridine orange showed an intense absorption at 596 nm and 501 nm, respectively, due to π-π* transitions. In the compositions, these shift to 587 nm and 490 nm, respectively, although their intensity was not affected \(\log \varepsilon\) (molar extinction coefficient) calculated 4.25. The hypsochromic shift indicated less delocalized electronic structures of the stains and, in turn, implied the interaction of the colorants with a mixture.

**Fluorescence spectra**

The photophysical parameters of the fluorescent stains, as well as the fingerprints powder compositions were presented in Table 2. The emission band was observed at a longer wavelength than the absorption band. Thus, in accordance with Franck-Condon principle and thermal relaxation of vibrational modes, the fluorescence spectra was observed on the red side of the absorption band, in an approximately mirror image relationship [11]. The bathochromic shift, in turn, might be attributed to solvation and/or caging effects. The more pronounced stokes shift in the compositions, as compared to those in pure stains, suggested an excited state species that was structurally different from the ground state analog, implying, in turn, a considerable degree of conformational changes in the compositions vis-à-vis the pure stains.

The actual radiative lifetime of the excited state, \(\tau\), was smaller than the intrinsic radiative lifetime, \(\tau_n\), indicating the possibility of non-radiating energy dissipation processes depopulating the excited state. Thus fluorescence remained the dominant but certainly not the exclusive mode of emission.

The difference between the actual and intrinsic radiative lifetime was more pronounced in the compositions than in the stains. The acridine orange-based composition had a larger difference than the brilliant blue R based composition. This meant that in the former case the non-radiative processes that were, intersystem crossing and internal conversion compete with fluorescence to a larger extent than in the later. This was duly reflected in the larger value of Einstein’s probability of spontaneous absorption, \(B_{nm}\) than the corresponding probability of spontaneous emission, \(A_{nm}\) for the brilliant blue R based formulation. For acridine orange based composition, the value of spontaneous absorption and emission were comparable. The dominance of fluorescence in brilliant blue R based powder was responsible for better quality fingerprints developed by this composition than those developed by acridine orange based composition.

The quantum yield of fluorescence, intersystem crossing and internal conversion followed the order \(\Phi_f > \Phi_{isc} > \Phi_{ic}\). Their priorities were also established by their respective rate constants [12].

The oscillator strength, \(f\) had been calculated from the relation \(f = \frac{4.31 \times 10^{-9} \int \varepsilon(\nu) d\nu}{\Delta \nu}\) which was valid if we assumed a Lorenzian shape for the absorption band. The factor \(\int \varepsilon(\nu) d\nu\) was replaced by \(\varepsilon_{\max}\ \Delta \nu\) where \(\Delta \nu\) was the half band width of the absorption band [13]. The greater value of oscillator strength of brilliant blue R based composition as compared to the acridine orange based composition was also a factor contributing to the better quality of developed fingerprints.
fingerprints.

Conclusion

On the basis of this investigation, we endeavoured to suggest that while formulating the fingerprint dusting powders, the degree of chemical interaction between the bonding material and the colorant should be taken into consideration. The level of interaction between these two broad components of a fingerprint composition, which might be gauged on the basis of spectral and photophysical studies, profoundly affected the quality of developed impressions. Thus, in the present case, better results obtained with brilliant blue R based composition may be attributed to a greater degree of interaction between the mixture and colorant, leading to greater fluorescent characteristics and oscillator strength of this formulation as compared the acidine orange based composition.

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