

# Ultraviolet fluorescence phototography of the Shroud of Turin

V. D. Miller and S. F. Pellicori

*Members, Shroud of Turin Research Project  
Santa Barbara, California*

One of the nondestructive techniques used to investigate the Shroud of Turin was ultraviolet fluorescence (UV) photography. This technique is able to detect organic and inorganic compounds by their integrated emission spectra and it is the complement of the more common technique of reflectance photography. Photographic data collection was one of many information resources designated by the Shroud of Turin Research Project<sup>1</sup> (STURP) team for the investigation of the body image and blood stains. The goal of the team was to determine the nature of the body image and its cause.

## Background

The investigation in October 1978 followed the conclusion of the public exhibition held in celebration of the 400th anniversary of the Shroud's housing in Turin, Italy. That was the first public exhibition in 45 years and the first full-scale multidisciplinary study in the Shroud's history.<sup>2,3</sup>

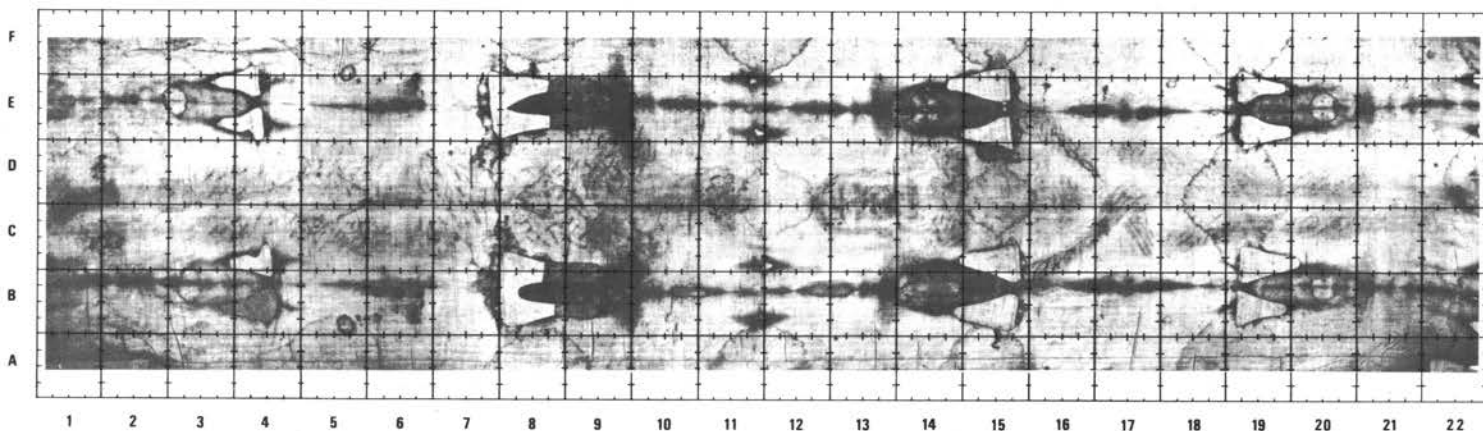
The Shroud is a 4.4 by 1.1 meter piece of age-yellowed linen which displays the life-sized dorsal and frontal images of a man. The appearance of the visible image and the locations of the blood stains suggest parallels with the descriptions of Christ's crucifixion in the Scriptures. For this reason, the legend associated with the Shroud of Turin is that it is the burial cloth of Jesus of Nazareth. It has also been suggested that the image was painted in the 14th century. Since the historical record is complete back only to the 1350's, the veracity of the legend cannot be directly addressed. Obvious scorch and water marks can be pinpointed to a fire in 1532. Hypotheses such as artistic painting, for example, can be tested for agreement with the observations. The team was thus charged with the task of collecting multifaceted data of necessary and sufficient quality and quantity for testing various hypotheses.

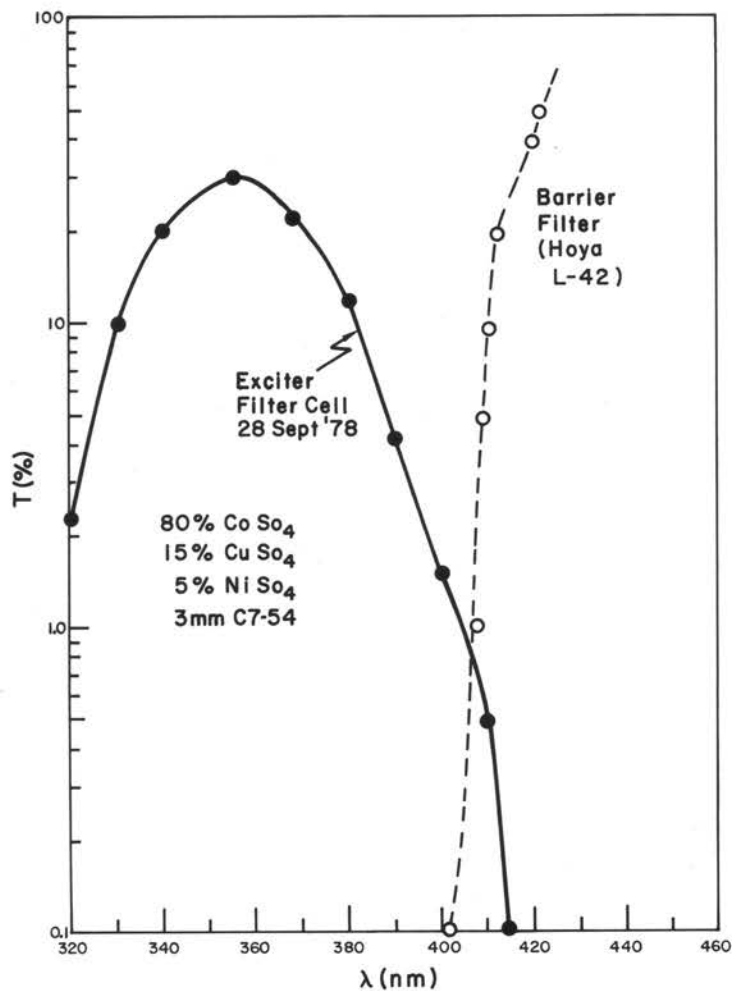
The United States team, an independent group of 32 scientists and assis-

tants, was equipped with instrumentation<sup>4</sup> capable of detecting work produced by known artistic techniques and adequate to provide a broadly-based foundation of information. The instrumentation included x-ray transmission and fluorescence<sup>5</sup> to detect high atomic number elements expected for inorganic pigments, microscopy<sup>6</sup> for visual and photographic examination of details, photoelectric spectrophotometry<sup>7,8</sup> for measurements of reflectance and fluorescence, photography<sup>9</sup> through bandpass filters, infrared spectrometry<sup>10</sup> and UV fluorescence photography documented here for the first time.

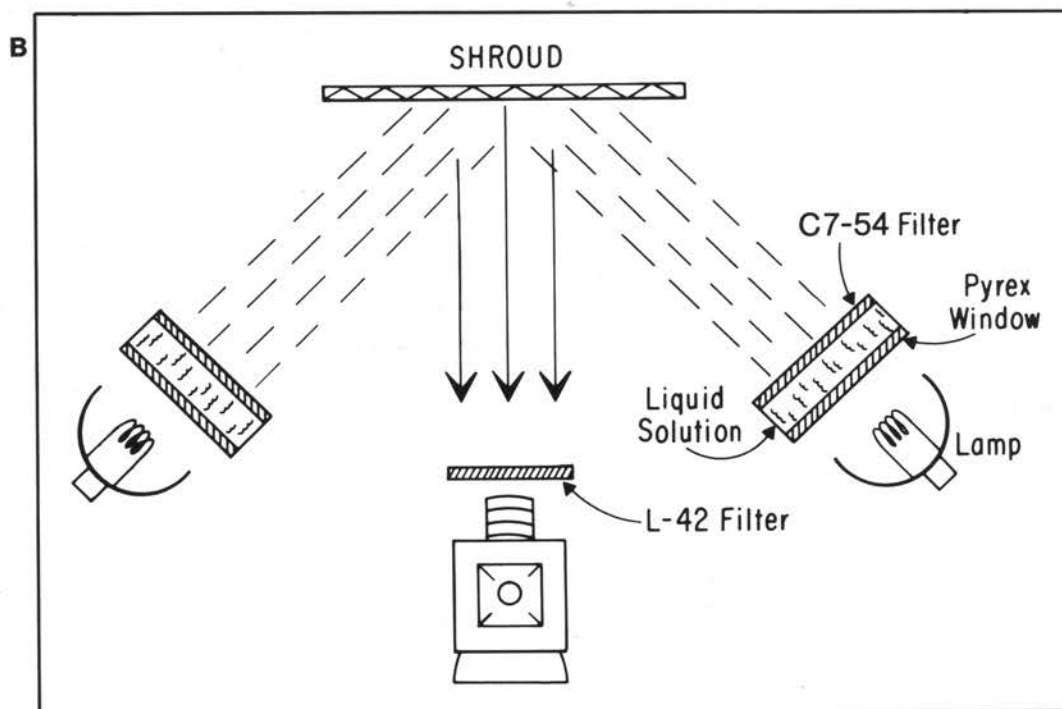
## Photographic procedure

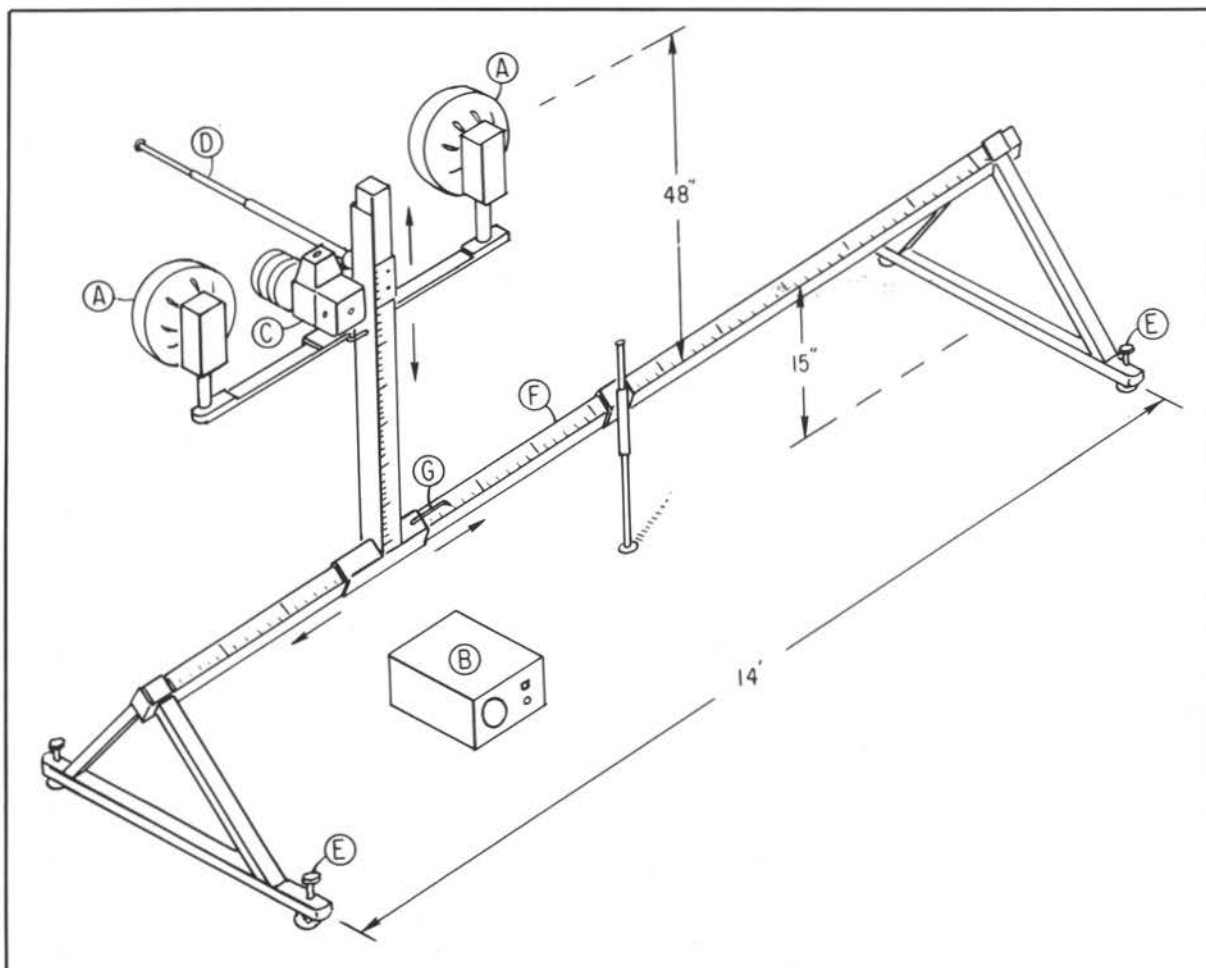
To record the emission excited by ultraviolet, a special light source was assembled. For our experiment a filter-window bandpass enabled wavelengths of 335 to 375 nm to be isolated from the sources for the exciting radiation. The two sources aimed at 45-degree angles from





**Figure 1A**—Transmittance curves for the ultraviolet (UV) exciting filter cell used in front of the xenon source and for the UV barrier filter used over the camera lens. **B**—Configuration of exciter and barrier filters for ultraviolet fluorescence photography. The exciter filter consisted of a Pyrex window and a filter cell containing a 1 cm path of the solution shown and a Corning 7-54 glass filter for absorption of the residual red leak. The Hoya L-42 filter served as a barrier filter transmitting only those wavelengths above 420 nm. **C**—Exciter filter cell attached to flash reflector.





**Figure 2**—Camera and illumination positioned on support rail for indexing along the length of the Shroud and thus creating a complete mosaic of photographs according to the reference layout in Figure 3. **A**—Lamp heads (Norman 200 B) with liquid cell filter. **B**—AC power pack. **C**—EL Hasselblad camera with 70 mm back. **D**—Telescoping extension rod for locating rail equi-distant from Shroud frame. **E**—Leveling screws. **F**—Centimeter scale with indexing marked for predetermined positions. **G**—Adjustable sliding index pointer for starting sequence.

the camera-subject axis were 200 watt-second xenon strobes with 15-cm reflectors. The film was to record only radiation emitted in the visible region of the spectrum, and none of the reflected, exciting UV. Consequently, the light source had to be free of visible radiation. To achieve these conditions special excitation filters were constructed to fit on the 15-cm diameter strobe sources.

The filters consisted of a 1-cm path length containing a mixture of inorganic salts of transition metals<sup>11</sup> to absorb radiation above about 400 nm. One window of the liquid filter cell was Corning 9863 visible-absorbing UV-transmitting glass.

This 9863 filter was required to attenuate a small red leak in the exciter filter. The other window was pyrex. The isolated passband and out-of-band attenuation measured for the filter are shown in Figure 1. Since the fluorescences we sought were expected to be in the visible, the filter needed to attenuate to a level of  $10^{-4}$  since visible light from the xenon tubes would completely swamp the weak fluorescent signals. The reasons for using a liquid filter system were that no other filters such as multilayer coatings or ultraviolet transmission glass alone are able to define the required passband while possessing adequate rejection. Further,

none of the standard filter materials could survive the heat of the xenon flash. Finally, the 15-cm size was beyond standard UV accessories.

The exciting UV energy which was partially reflected by the Shroud was prevented from reaching the film by a nonfluorescing, long-wavelength pass, UV absorbing glass placed in front of the lens (Figure 1). This barrier filter passed wavelengths greater than 410 nanometers.

The camera and light source assembly was moved along a rail parallel to the long dimension of the Shroud, which was mounted with its short dimension vertical (Figure 2). The long dimension was di-

vided into eight 53.3-cm square areas for photography and later full-size reconstruction. The areas were intended to be coincident with the black-and-white color separation series taken at another time. The sections were numbered from left to right, beginning at the dorsal feet end of the Shroud. The number-letter coordinate set corresponded with a master reference mosaic and is shown in Figure 3.

The camera was a Hassleblad EL with a 80 mm lens. The film used was Kodacolor 400 film, a color negative film with an exposure index of 400. Prints were then made on Kodak Ektacolor no. 78 paper.

Differences between the fluorescent record and white light photos were noted. Assisted by laboratory data, interpretation of these differences provides insight

into the nature and causes of the various markings on the Shroud, namely the body image, blood stains, and scourge, water and scorch marks. These interpretations can be compared with laboratory-produced simulations.

### Detailed observations

Some general comparisons from color photographs of fluorescing colors and natural light colors are listed in Table I. Differences are detailed in the following pages for each section of the Shroud under discussion. Section B through E by 1 through 4, for example, defines a rectangle according to the coordinates of the reference overlay on the Shroud. (See Figure 3 and the reduced size overlay in the text.)

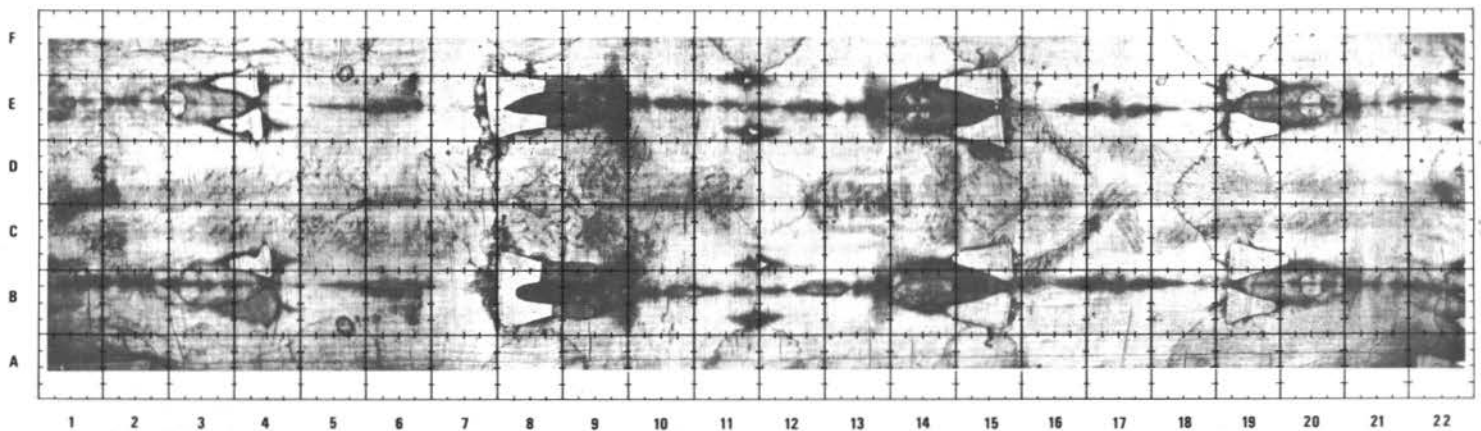
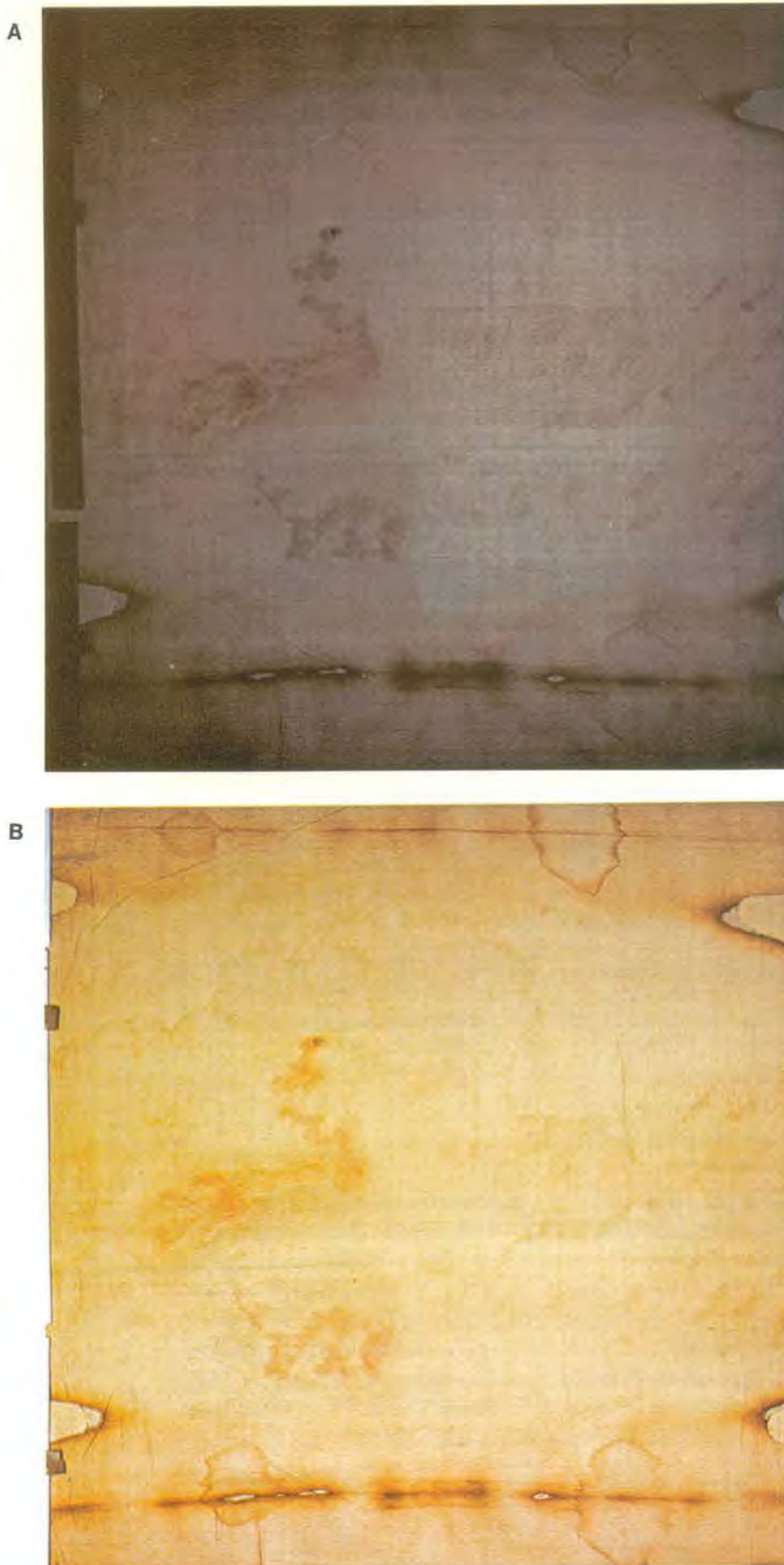


Figure 3—Reference layout of the Shroud showing the numbered and lettered sections discussed in detail in the text.

**Table I—Comparison between UV Fluorescence and Visible Reflectance Photographs**

Feature	UV Fluorescent Characteristics	Visual Appearance
Background (Non-Image Areas)	Generally yellow-green, with blue flecks thought to be modern lint contamination. Patches show a variety of colors; blue and brown. Creases and wrinkles not easily discerned.	Yellow as expected for aged linen. Patches are lighter in color. Creases and wrinkles are obvious because of shadowing.
Blood	Highly absorbing. No color. Fluorescing borders apparent around some areas.	Has color of dried blood; i.e., red to reddish-brown.
Scourges	Highly absorbing and many resolve into scratch-like lines in parallel grouping of 3 or 4.	Reddish, diffuse in structure. Some dark spots seen within.
Body Image	No color; prevents or absorbs the background fluorescence. Greater contrast with background here than in reflected visible; less gradation in shading across a body feature than contained in visible image. Variations in image density can be abrupt, associated with weave pattern.	Straw colored, low contrast. Gradual shading into background density toward outer areas of a body feature. Weave has a smaller influence on image clarity.
Water Marks	Some edges are more highly accentuated, others nearly disappear when compared with reflected light borders just outside the absorbing edges. Colors grade as in unresolved liquid chromatography from yellow/green to brown.	Generally brown with sharp edges and an unresolved liquid chromatographic appearance.
Scorches	The visually dark brown burns fluoresce brownish-red. The color reddens as the scorch density decreases. Comparable to pyrolysis products, produced under limited oxygen combustion, such as furfurals. The open burn holes of regions 6 and 16 show total absorption; i.e., no color.	Burns are dark brown to black. Those associated with the 1532 fire are dark brown and grade to light brown with distance. Weak, light-toned scorches resemble the body image color. The open burn holes of regions 6 and 16 have black borders (carbonized).



### Dorsal feet area—B through E by 1 through 4 (Figure 4)

Some cloth fluorescent characteristics stand out as different from the white light appearance, namely creases, thread, and shadowing are less apparent in UV. The striation in the weave pattern (warp versus weft) are enhanced in fluorescence. There is a general decrease in cloth emission at the corners. The teflon coated magnets, quite visible in white light, are nonfluorescent. This signifies the rejection level of reflected light. The visual water marks are not detectable fluorescently, while the scorches around the burns emit reddish fluorescence.

### Left foot—C through D by 1 through 2

Blood is red visually, but neutral to black (absorbing) fluorescently. Detailed shading in absorption in the blood in center of left foot. A triangular-shaped pattern is seen at the ball of the foot. Note the very dense absorption in the blood area at the left end of the rivulet from the heel: scorching might be associated. A fluorescing border in the blood flow off the body image areas is seen.

### Right foot

A more distinct light border area is seen. The body image appears different from the blood in fluorescence, that is, it absorbs less intensely. The scourge marks have greater contrast in fluorescence than in white light and on the calf appear as lines rather than as dumbbell shapes as they do further up on the leg. Some scourges are not perceptible in the white light photo. The scourge lines run parallel and diagonally from top left to bottom right. The background cloth fluorescence is prominent between the leg images (C-D area). The absorbing water marks at 3 and B through E have light border areas.

**Figure 4**—Section B through E by 1 through 4 of the Shroud. **A**—UV fluorescence. **B**—white light reflectance photograph.

### General dorsal areas—4 through 6 (Figure 5)

Red fluorescence is seen around patches at left and right boundaries. There is a series of holes through the Shroud cloth (through which the reinforcing backing cloth sewn to the Shroud after the fire can be seen). The densely burned borders of these burned holes show no emission. Where the more lightly scorched material matches the appearance of other scorched areas (around added patches, for example), the fluorescent appearance is quite different, that is, no emission is seen. The different appearances might relate to different generation mechanisms.

### Calf areas of right and left leg

Scourges favor orthogonal orientation in this area. As the thigh is approached, the scourges become scratch-like. On the right leg at C-5 and D-6, the scourges appear to be off the body image area. Weave striations are obvious in fluorescence. In D-6 and C-6, we note small areas of brightness between center burns in both emission and reflection photos. The circular mark in the thighs (D-5 and C-5) resembling a scourge might be a scorch mark. Similarly, in D-7 and C-7, there is no fluorescence emission, but perhaps the marks have the same origin as the circular burn holes in this section. At the boundaries, B and E, and around the patches, the scorches display shading in red emission. These scorches appear different from the burn holes. The water stain, C-5 and D-5, did not disturb the scourge marks. In areas B through E by 6-7, faint red-brown bands are apparent: they traverse the burn holes. Notice at the expected knee joint, 6, that there is no body image apparent and limited scourging.

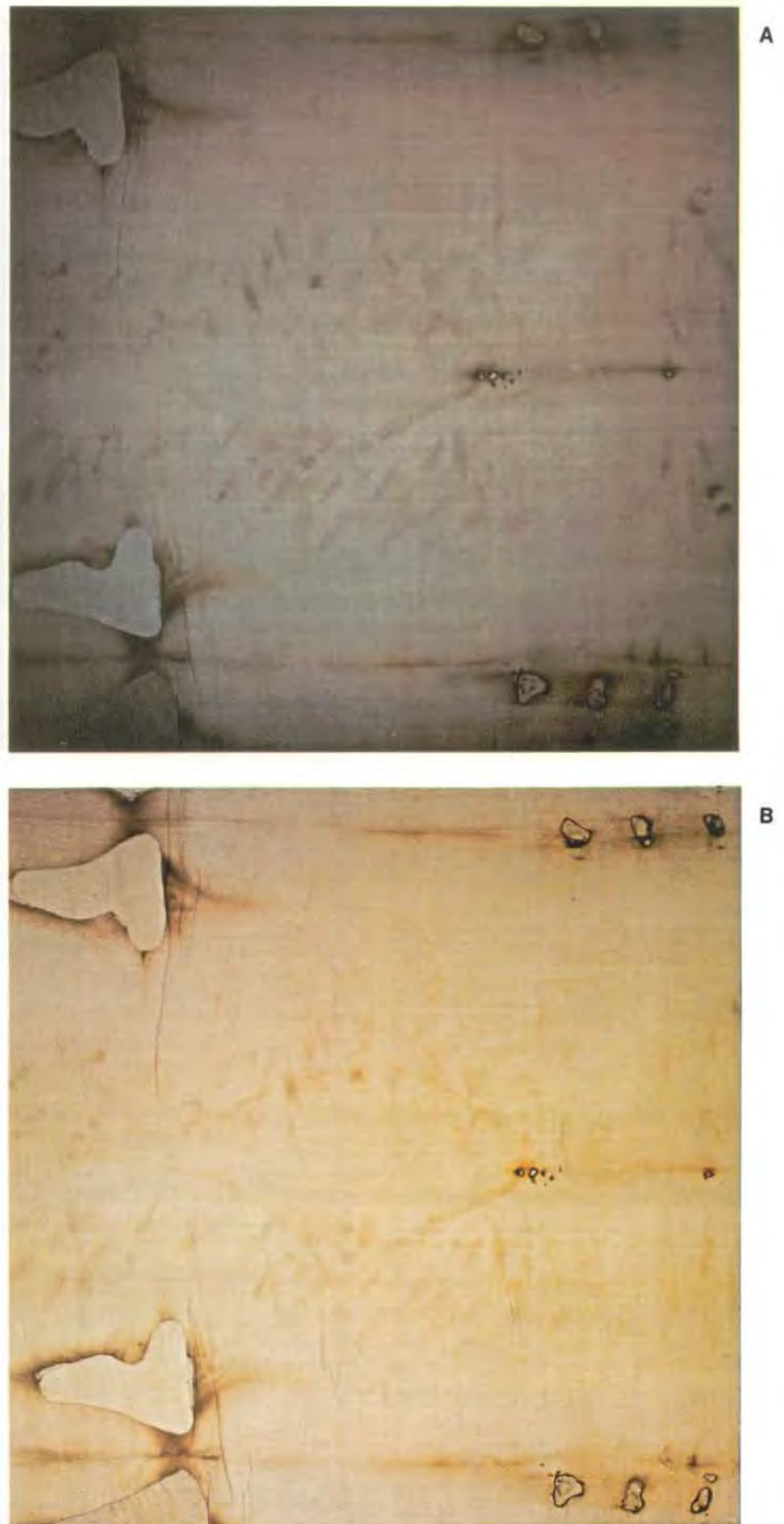


Figure 5—Section B through E by 4 through 6. A—UV fluorescence. B—white light reflectance photograph.

**A**



**B**

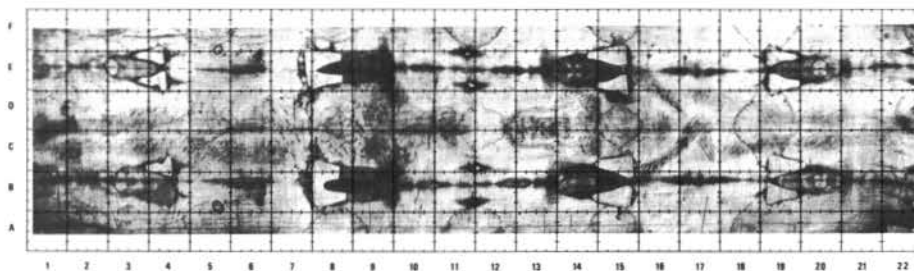




**Figure 6**—Section B through E by 7 through 9. **A**—UV fluorescence. **B**—white light reflectance photograph.

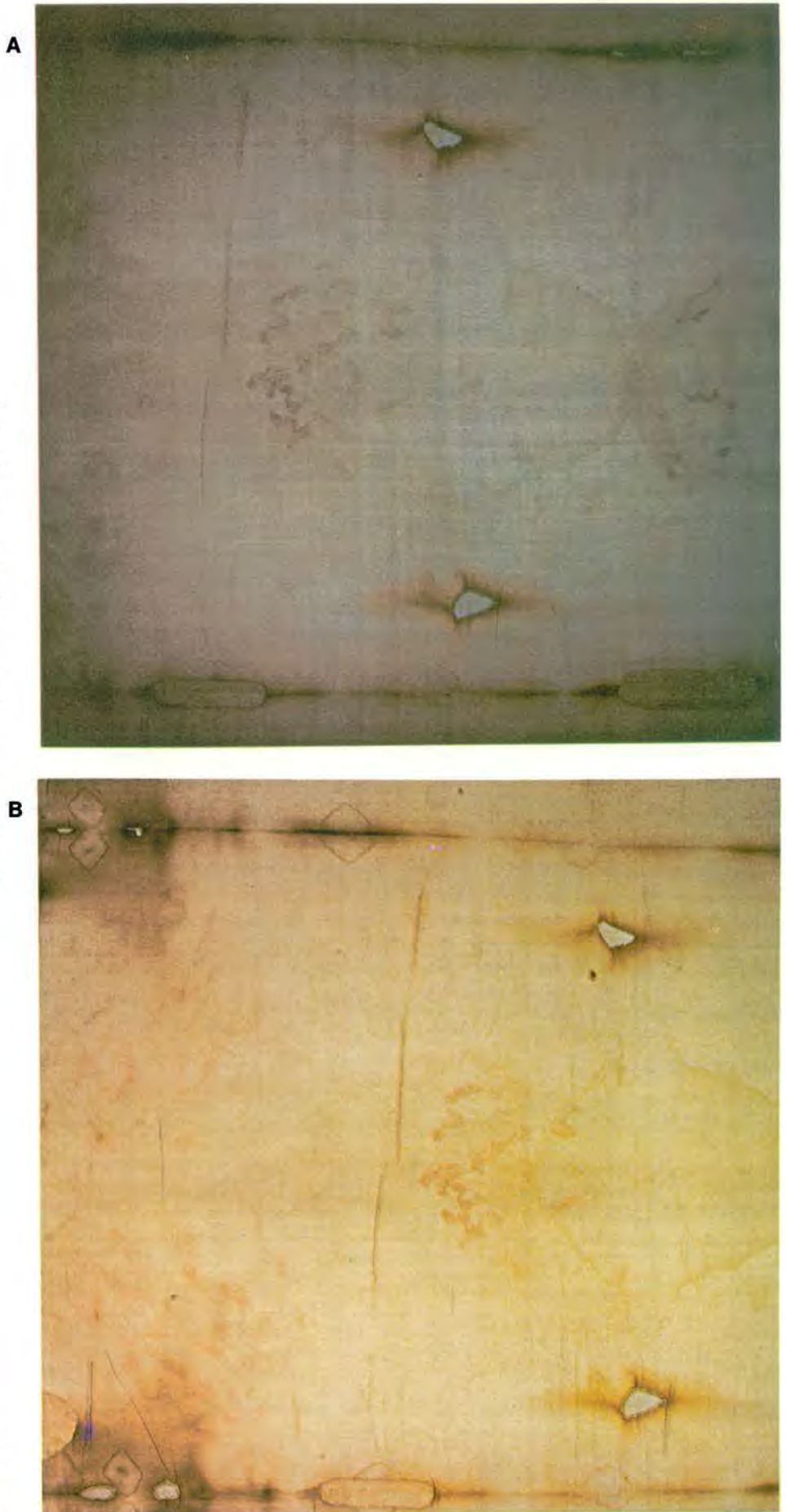
### Dorsal midsection area—7 through 9 (Figure 6)

All the patched burn areas fluoresce reddish brown, including the water stain front. Other white light water boundaries are indistinguishable in fluorescence. The scourges orient orthogonally and are distinctly absorbent against the fluorescent background linen. The emission of the linen is not uniform. Across the small of the back the rivulets of blood are absorbing; see also D-9 and C-9. There are some lighter border areas associated with the blood flow. Some of the densest marks within the blood flow area might be scorches. In area D-8, the blood boundaries fade in fluorescence. At E-8 and C-8 the stains at the inner boundary of the large patches appear to be mirror images. Two small circular stains at C-7 and D-7 are mirror images and visually resemble weak scorches. In fluorescence these stains are absorbing. Note also lighter boundary areas associated with water marks. The body image (nonfluorescing) is very dense at the scapula and rib cage. Scourges are apparent through the scorched shoulder areas above the patches. Many scourges have fluorescing bordering areas.



**Dorsal head and neck area  
and ventral head area—10  
through 12 by B through E  
(Figure 7)**

At E and B and at patches 11 and 12, red fluorescing scorched areas are obvious. The brightly fluorescing yellow-green donut-shaped areas to the right of patch 11-12 at E and B appear to be scorches. They are mirrored elsewhere at symmetrically located fold sections. The faint water stain between the head images has light blue boundaries in fluorescence. The cloth weave striation is an apparent nonuniformity. The blood stains on the dorsal head area are bounded by brighter areas. A smudge or scourge appears to the right of the general blood flow on the dorsal head. Faint scorch pattern is visible in the light blood stains. At C-D by 11-12 a smudge resembling blood is visible between the head images. The dorsal body image is more distinct than the ventral image. At the center of the dorsal head, a blue fluorescence is noted. This has a different color than the body image.



**Figure 7**—Section B through E by 10 through 12.  
**A**—UV fluorescence. **B**—white light reflectance  
photograph.

**A**

### Head and chest—13 through 15 by D through C (Figure 8)

Scorches near the patches emit reddish brown fluorescence. The water stain boundary below patches at C-15 and E-15 appear to contain reddish fluorescent material as from scorches, while those at C and D at 15 (abdomen) do not contain fluorescing material. The lower left arm blood stains, B and C at 16, have light border areas. Through the scorches of B at 14 and 15, upper left arm, can be seen blood or scourges. The upper area of the left shoulder shows faint blood, scourge and body detail. The diffusely appearing scourge marks below the central water stain, C and D at 16, become more distinct (scratch-like) near C-15 and D-15. Scourges are diagonal and orthogonal, with a scratch characteristic predominant on the left. The face is more strongly bounded by the weave pattern on its left side than on its right, C and D at 13. Going beyond this feature, no image is discernible. The blood streaks in the hair are denser on the right side and have fluorescing boundaries, C and D at 13. In comparison with the general body image, the beard and mustache are denser. In fact, the density is greatest on the left portion of the beard and mustache and more diffuse on the right. A distinct boundary is present at the lower left. The lower lip appears to have a fluorescent boundary. On the right shoulder, the blood stains are in very sharp detail, with the lower stain broken into dots. Compare this area with some of the scorches on the right side. Circles of yellow-green fluorescence are associated with these wounds.

**B**

**Figure 8**—Section B through E by 13 through 15. **A**—UV fluorescence. **B**—white light reflectance photograph.

### Ventral hands and thighs— area 16 through 19 (Figure 9)

Scorches show red fluorescence except for those associated with the sets of three holes which have heavily burned borders. Plumes of pyrolyzed materials pointing toward the feet are seen associated with the burned holes. At 19 the plume pointing toward the hands fluoresces red. Some shading from red to yellow fluorescence can be seen. The water mark above the knees at 18 has an absorbant edge with density gradations. Some fluorescing bordering can be seen also. In white light, however, this water stain is not prominent. The fluorescent color is brown as opposed to grey. The water stain situated above the series of holes to the right side has very little emission. Some of the water stains are better defined in fluorescence, others are not. The blood and body image are similar in the fluorescence photos; i.e., grey and non-emitting. Notice the clear fluorescing borders around the hand wound blood stains. As is true of scourges elsewhere, the scourge wounds are more distinguishable in the fluorescence photos. As the upper thigh area is approached, however (see 19), the scourges become more diffuse. Often scratches are contained within the diffuse areas. Some scourge marks appear only in the fluorescence photos: examples are noted between the hands and forearm areas. Around the knuckle of the left hand a slight blood flow can be seen. The body image absorption, especially arm and thigh, is greater on the left side of the body. There is a detail located where the genitals would be expected below the intersection of the hands. The dark patch below the wrist wound appears distinct from the body image. It is not understood. The fingers possess more contrast by fluorescence photography. The differences among water stain appearances might be due to differing material content, with some containing mobile pyrolysis products.

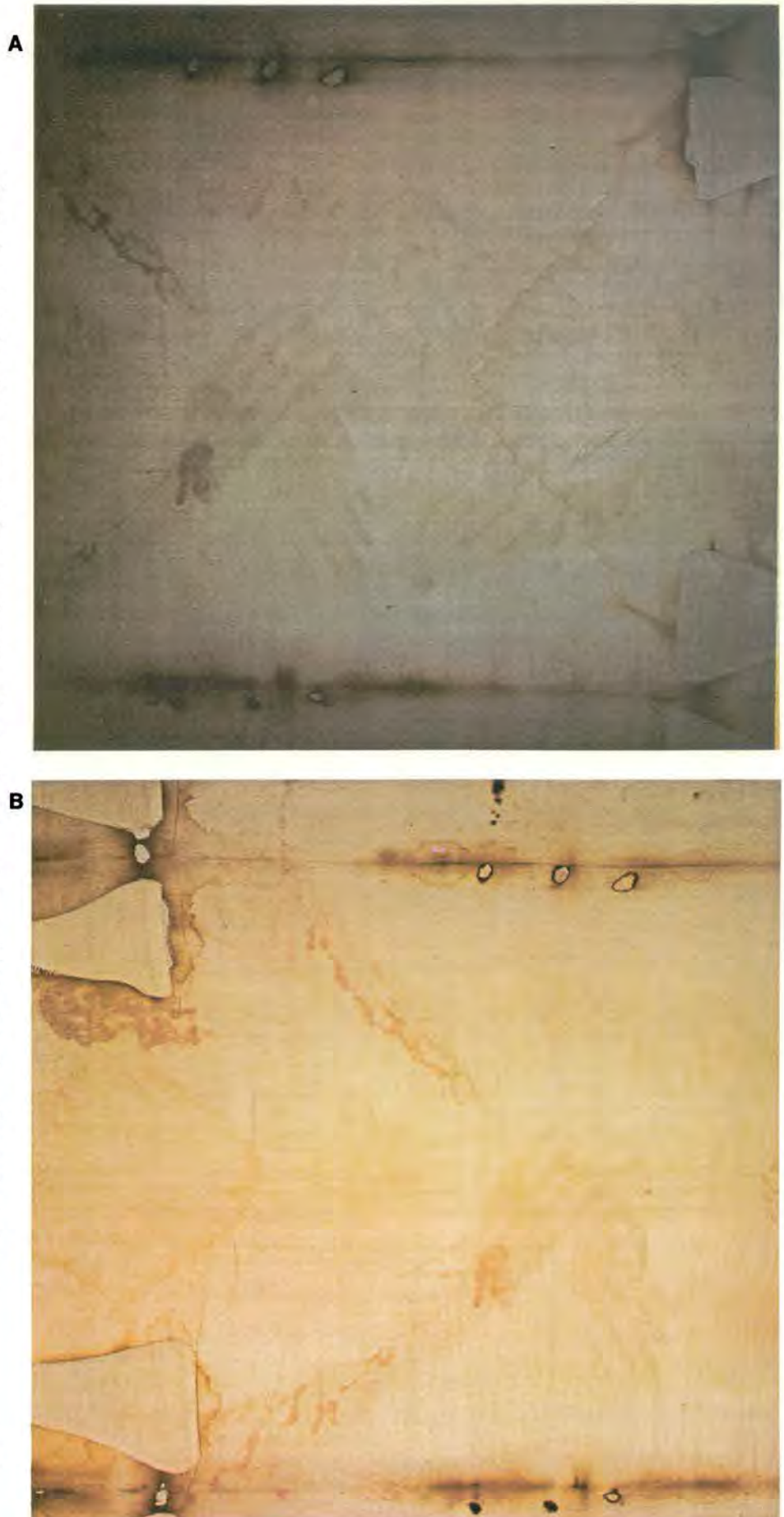


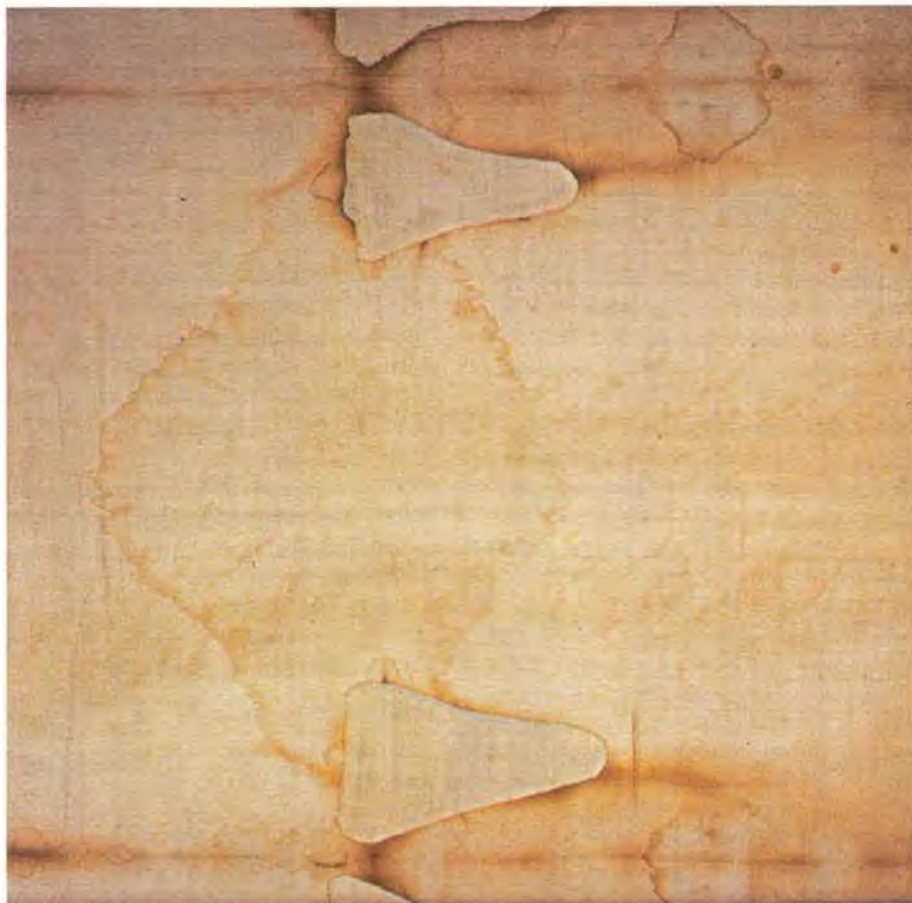
Figure 9—Section B through E by 16 through 19.  
A—UV fluorescence. B—white light reflectance photograph.

**A**

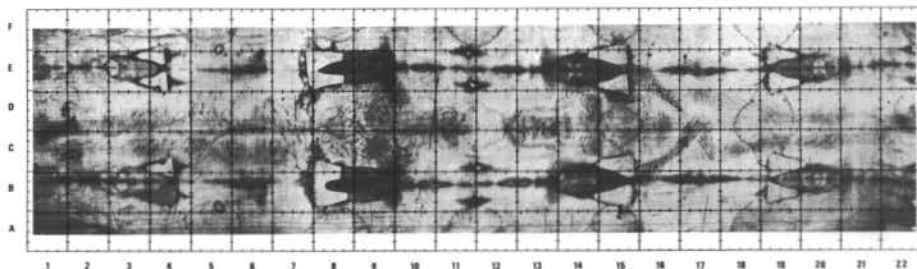
### Ventral feet, knees and thighs—19 through 22 (Figure 10)

A very faint body outline is discernible in this section, even in fluorescence photos. The feet are not defined. The leg outline and scourge markings are limited by a weave line appearing blue in fluorescent emission where the weave direction changes. This is an area of “no-print.” The central water stain at B and C by 18 and 19 is contrast-enhanced by fluorescence: it absorbs strongly against the emitting background. While the water stain at D-20 has fluorescing border areas, the opposing one at A-20 does not. Also, the one at D-22 has an unusually wide absorbing boundary. As in the other scourged areas, the scourges run diagonally left to lower right, and both diffuse and scratch-like marks, some perpendicular to the leg, are visible. See the left leg at B-20 and 21. Some scourges appear to be bounded by the central water stain. No scourges can be found on the ankles.

On the feet, two blood stain areas are distinguishable on the right foot; the smaller one is denser, the larger has a fluorescing border area. Curious less densely absorbing flows without definite boundary trail from the blood spots. They do not resemble the usual blood characteristics. The small circular mark above the water boundary at D-22/21 has the fluorescent colors of a scorch: red grading to a yellow border. To the upper right a 2-inch diameter stain containing three dark lines appears by fluorescence contrast, but not by white light. Wax drops seen in this area are bright fluorescers colored yellow, or green. At the lower edge, D-22, two sharp-edged marks visible as dark brown emit bright white in fluorescence.

**B**

**Figure 10**—Section B through E by 19 through 22. **A**—UV fluorescence. **B**—white light reflectance photograph.



### Discussion—Laboratory experiments

In fluorescence photographs most of the details are visible in contrast with the Shroud linen, which itself fluoresces. With the exception of lightly scorched areas and some water stained areas, the Shroud features absorb UV energy without visible emission. Compared with modern linen, the Shroud linen fluoresces less brightly. It emits a yellow-green color. Modern linen can be artificially aged by baking at high temperatures ( $125^{\circ}$ – $150^{\circ}$  C) to the point where its reflected color and fluorescent emission approach those of the Shroud.<sup>7</sup> When foreign materials are applied to the linen, a reaction which results in locally visible darkening of the linen can be stimulated by air baking as above. These experiments<sup>7</sup> have led to the interpretation that the body image is the result of locally accelerated dehydration/oxidation and conjugation of the cellulose molecular structure. Images produced in hours at high temperature are comparable in both reflectance and UV stimulated emittance to images produced at normal temperatures during longer time intervals (years?). Laboratory-produced images were photographed using the same equipment as was used in Turin. Exposure to sunlight of wavelengths less than 340 nm influences the rate and degree of cellulose degradation.

Laboratory data for whole blood displayed total absorption, which is in agreement with the Shroud data.

Weak scorches, which in white light have densities and colors similar to body image areas, show a significant difference with UV fluorescence photography. Scorches emit a reddish brown fluorescence while the body image is nonfluorescent. The significance of this difference is evident in relation to the suggestion that the body image was caused by con-

tact with a hot statue or was scorched by other means. Laboratory-produced scorches emit a bright greenish-yellow fluorescence if they were produced in air and reddish if produced under conditions of limited available oxygen. The scorches associated with the fire of 1532, during which the Shroud was involved, attest to the rapid consumption of the available oxygen.<sup>12</sup> Their reddish emission is probably due to furfurals, which can be produced under such conditions.

Linen lightly scorched by a soldering iron in air shows the green-yellow emission, often distributed in plumes of deposited pyrolysis products. We demonstrated in one experiment that the material of the plumes could be transported by water, but the underlying scorched cellulose retained a bright yellow-green fluorescence. This demonstration together with the observed absence of body image fluorescence is strong evidence against the cause for the body image being a scorch.

A quite frequent hypothesis is that the body image was painted. The binders used in paints as early as the 14th century would be made of proteinaceous materials, animal collagen being a favored material; egg white and gelatin are other examples. But these collagens would inherently be contaminated by fluorescing amino acids.<sup>13</sup> Random sampling of book bindings and illustrations from the 14th and 15th centuries were observed to emit bright fluorescence when excited by long-wave mercury vapor lamp excitation. Microchemical analyses on fibrils retrieved from the Shroud have shown the absence of paints, pigments, stains, dyes and protein in body image areas.<sup>14</sup>

### Conclusion

The sharp detail revealed for the first time, particularly in the scourges, suggests that intimate cloth-body contact occurred. The detail (and contrast) is only slightly less prominent on the front than

on the dorsal image, indicating that the large difference in weight for each side had only a minor influence on the imprinting of the scourges. This observation is contrary to what might be intuitively expected, and it might be a clue to some future understanding of the image production mechanism.

The occurrence of contact could also have transferred substances present on the skin to the cloth where they stimulated the cellulose alteration process and caused an image of the body to develop as local darkening of the linen.<sup>7</sup>

Hypotheses such as a scorch cause or paint are contradicted by the fluorescence photography results.

UV fluorescence photography has revealed some near-invisible details, many of which require explanation. For example, the pattern of distinct burn holes has characteristics unlike the burn damage attributed to the 1532 fire. Areas in the weave where the image density abruptly decreases (e.g., sides of the face) might actually contain very faint images which possibly could be retrieved by using stimulating radiation of shorter wavelengths. The property of the linen thread that didn't develop image density should also be discovered. The 8-cm side strip running the length of the Shroud shows weft bands that are continuous with the main body of the Shroud. Similar appearances result from backlighting and low-energy radiography. The suggestion is that this strip was not separated from the main body over its entire length. Many of these unexplained details might relate to the history of the Shroud.

Another feature requiring explanation is the lighter bordering area seen with many bloodstained areas. The interpretation is that blood serum is present. It might have acted to retard the image development reactions associated with the body image. Fibrils from many water stain fronts and from the area be-

tween the head images contain blood particles.<sup>14</sup> Further discussions are in preparation.<sup>15</sup>

### Acknowledgments

We appreciate the assistance given by D. Devan and other members of the STURP team, and also acknowledge the cooperation extended by the church officials in Turin. J. Druzik, Conservation Center, Los Angeles County Museum of Arts, critically reviewed the manuscript.

**About the authors**—Vernon Miller is Head of the Scientific and Industrial Photography Department at the Brooks Institute of Photography, Santa Barbara, California. He was the chief scientific photographer for the STURP team in Turin, and has employed a variety of image processing techniques to the photographic data obtained.

Samuel Pellicori is an optical physicist with the Santa Barbara Research Center. He has been evaluating and simulating in the laboratory some of the spectrophotometric, fluorometric and microscopic data he collected in Turin. Address correspondence to Samuel Pellicori, Santa Barbara Research Center, 75 Coromar Drive, Goleta, California 93017.

### References

1. Proceedings of the 1977 U.S. Conference of Research on the Shroud of Turin, Holy Shroud Guild, 294 E. 150 St., Bronx, NY 10451.
2. Kenneth F. Weaver, "The Mystery of the Shroud," *National Geographic* 157, 730 (1980).
3. B. J. Culliton, "Science Investigates the Shroud of Turin," *Science* 201, 235 (1978).
4. E. J. Jumper and R. W. Mottern, "Scientific Investigation of the Shroud of Turin," *Appl. Opt.* 19, 1909 (1980).
5. R. A. Morris, L. A. Schwalbe and J. R. London, "X-Ray Fluorescence Investigation of the Shroud of Turin," *X-Ray Spectrometry* 9, No. 2, 40 (1980).
6. Samuel Pellicori and Mark S. Evans, "The Shroud of Turin Through the Microscope," *Archaeology* 34, 34, Jan/Feb 1981.
7. S. F. Pellicori, "Spectral Properties of the Shroud of Turin," *Applied Optics* 19, 1913 (1980).
8. Roger Gilbert, Jr., and Marion M. Gilbert, "Ultraviolet-Visible Reflectance and Fluorescence Spectra of the Shroud of Turin," *Applied Optics* 19, 1930 (1980).
9. V. Miller and D. Lynn, "Photography of the Turin Shroud," *Science and Technology*, Feb. 1981 (in Dutch).
10. J. S. Accetta and J. S. Baumgart, "Infrared Reflectance Spectroscopy and Thermographic Investigations of the Shroud of Turin," *Appl. Opt.* 19, 1921 (1980).
11. S. F. Pellicori, "Transmittances of Some Optical Materials for Use Between 1900 and 3400 Å," *Appl. Opt.* 3, 361 (1964).
12. R. N. Rogers in Reference 1, p. 133.
13. Eric J. Jumper, John P. Jackson, John H. Heller, Alan D. Adler, Samuel F. Pellicori and Raymond N. Rogers, "A Comprehensive Examination of the Various Stains and Images on the Shroud of Turin," *American Chem. Soc. Proc. on Archaeological Chem.* (1982), in preparation.
14. J. H. Heller and A. D. Adler, "A Chemical Investigation of the Shroud of Turin," submitted to *Journal of Forensic Sciences*.
15. L. A. Schwalbe and R. N. Rogers, "Physics and Chemistry of the Shroud of Turin: Summary of the 1978 Investigation," in preparation.