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# THE OCCURRENCE OF METHEMOGLOBINEMIA DURING SULFANILAMIDE THERAPY

BY CHARLES L. FOX, JR.,<sup>1</sup> AND JAMES E. CLINE

(From the Department of Bacteriology and Immunology, Harvard Medical School, and the  
Chemical Laboratory of Harvard University, Cambridge)

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The presence of methemoglobin, more rarely sulfhemoglobin, in the blood of patients treated with sulfanilamide has been reported by many investigators (1 a-f). Evelyn and Malloy (2), using their photoelectric colorimeter, found both these pigments but felt they were not responsible for the "cyanosis." Wendel (3), using his visual spectroscopic method (4), found methemoglobin and (occasionally) sulfhemoglobin. On the other hand, Marshall (5), Posner (6a) in many instances, Chesley (6b), and some occasional observers have failed to find the characteristic band of methemoglobin.

A little spectroscopic experimentation with pure methemoglobin will speedily convince anyone that the light absorption characteristics of methemoglobin renders visual detection of the band at 630  $m\mu$  rather difficult in the presence of a large excess of hemoglobin. Even Wendel's ingenious method (4) does not eliminate the difficulty of finding this band. If the personal visual factor might be eliminated, incontrovertible evidence might be obtained.

This is possible with the Hardy recording spectrophotometer (7) which accurately draws a curve showing the light transmission of a sample from 400 to 700  $m\mu$ . Typical curves of the blood of patients receiving sulfanilamide are reproduced (Figure 1) with normal controls. The prominent depression at 630  $m\mu$  is readily seen. In one case a depression at 620  $m\mu$  is also seen indicating the presence of sulfhemoglobin.

In these patients from 10 to 18 per cent methemoglobin was estimated to be present by using Beer's and Bouguer's laws and absorption coefficients obtained on pure solutions by the Hardy recording spectrophotometer (Table I).

Inasmuch as sulfanilamide *in vitro* does not produce methemoglobin, the mechanism of its formation *in vivo* remains unexplained. Further ex-

<sup>1</sup> Moritz Rosenthal Fellow, Mount Sinai Hospital, New York City.

TABLE I  
*Absorption coefficients*

$\lambda m\mu$	Hemoglobin	Methemoglobin	Colored residue	Blue substance
700	0.0825	0.114	8.64	24.1
660	0.0862	0.472	10.8	56.8
630	0.133	2.32	17.6	75.0
620	0.179	2.18	19.1	80.2
600	0.566	1.98	21.9	89.1
580	7.14	2.40	24.2	93.9
560	5.83	2.52	25.9	92.9
540	8.60	3.70	26.7	87.2
520	4.12	4.75	27.1	78.1
500	3.30	5.57	27.8	68.5
480	4.15	5.00	29.4	59.8
460	6.82	5.69	32.1	53.6
440	16.12	11.5	36.5	49.6

Transmission curves were obtained with solutions of each substance.

These coefficients were calculated after using the equation

$$\log \frac{100}{\% \text{ transmission}} = kcd.$$

$k$  = absorption coefficient

$c$  = concentration of the solution—1 gram per 100 cc. for hemoglobin and methemoglobin

$d$  = thickness of cell in cm.

periments showed that when the blue derivative (8) (a reversible oxidation-reduction system formed by the aerobic ultra-violet irradiation of dilute aqueous sulfanilamide (9)) is added to hemoglobin, methemoglobin is formed<sup>2</sup> (Figure 2). Using the accurate Hardy recording spectrophotometer, and applying Beer's and Bouguer's laws, we analyzed the resulting solution. Assuming that hemoglobin and methemoglobin were the only colored components, the transmissions at 12 different wavelengths from 460 to 700  $m\mu$  were calculated (Table II, Column B). These do not agree with the experimental values (Table II, Column A), and indicate the presence of another colored component. When a third component, the brown solution resulting from the spontaneous

<sup>2</sup> "It is interesting to note, however, that the colored derivatives of sulfanilamide described (by Ottenberg and Fox (8)) convert hemoglobin to methemoglobin *in vitro*." Wendel (3).

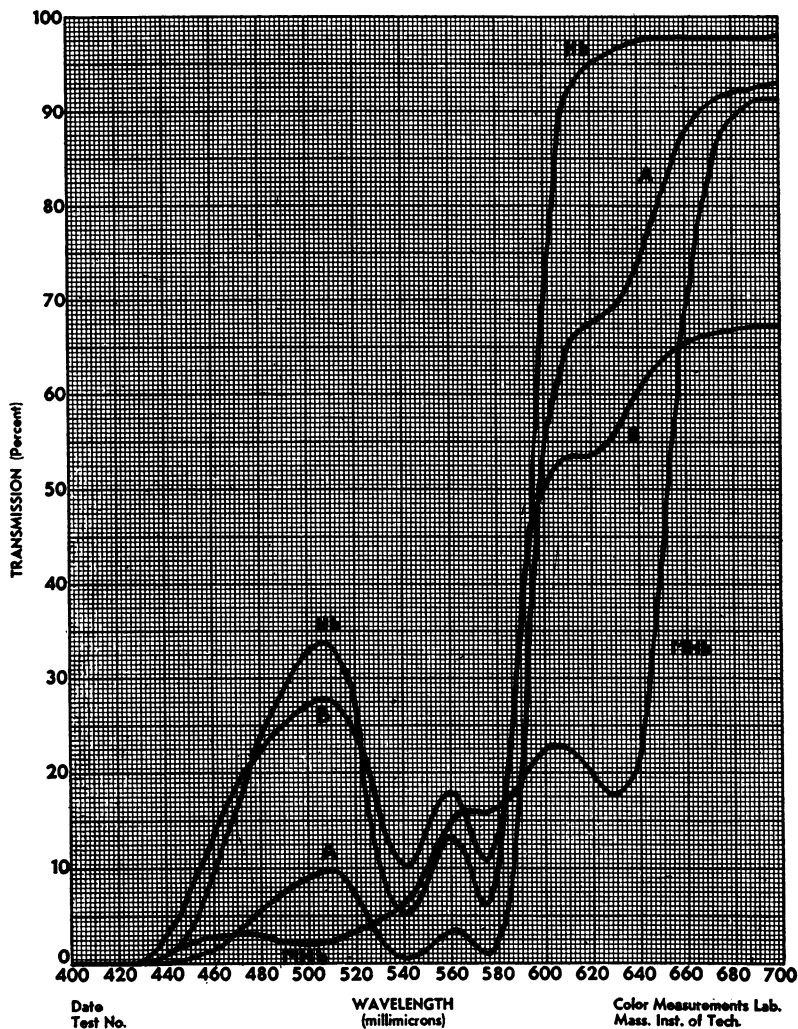


FIG. 1. CURVES OF WASHED, LAKED HUMAN ERYTHROCYTES DRAWN BY HARDY RECORDING SPECTROPHOTOMETER (pH—7.1)

Hb is the curve of 1/100 dilution of blood of normal subject—unaltered hemoglobin solution with characteristic bands at 541 and 573  $m\mu$ .

MHb is the curve of 100 per cent methemoglobin (same concentration as Hb) with prominent band at 630  $m\mu$ .

A is typical curve of patients' blood during sulfanilamide therapy (diluted 1/50) showing unmistakable prominence at 630  $m\mu$  caused by methemoglobin and general depression from 660 to 700  $m\mu$  (possibly due to a colored oxide of sulfanilamide).

B is curve of patients' blood during sulfanilamide therapy (diluted 1/100) showing prominence at 620  $m\mu$  caused by sulfhemoglobin and general depression from 660 to 700  $m\mu$  (possibly due to a colored oxide of sulfanilamide).

reduction in air of the blue oxidizing solution, is introduced into the calculations, the predicted transmission values (Table II, Column C) show significantly better agreement with the experimental values (Table II, Column A).

Similar calculations with the data of the curves of patients' blood showed good agreement (Table III) between the observed curve and one predicted on the basis of two components: hemoglobin and methemoglobin in the region from

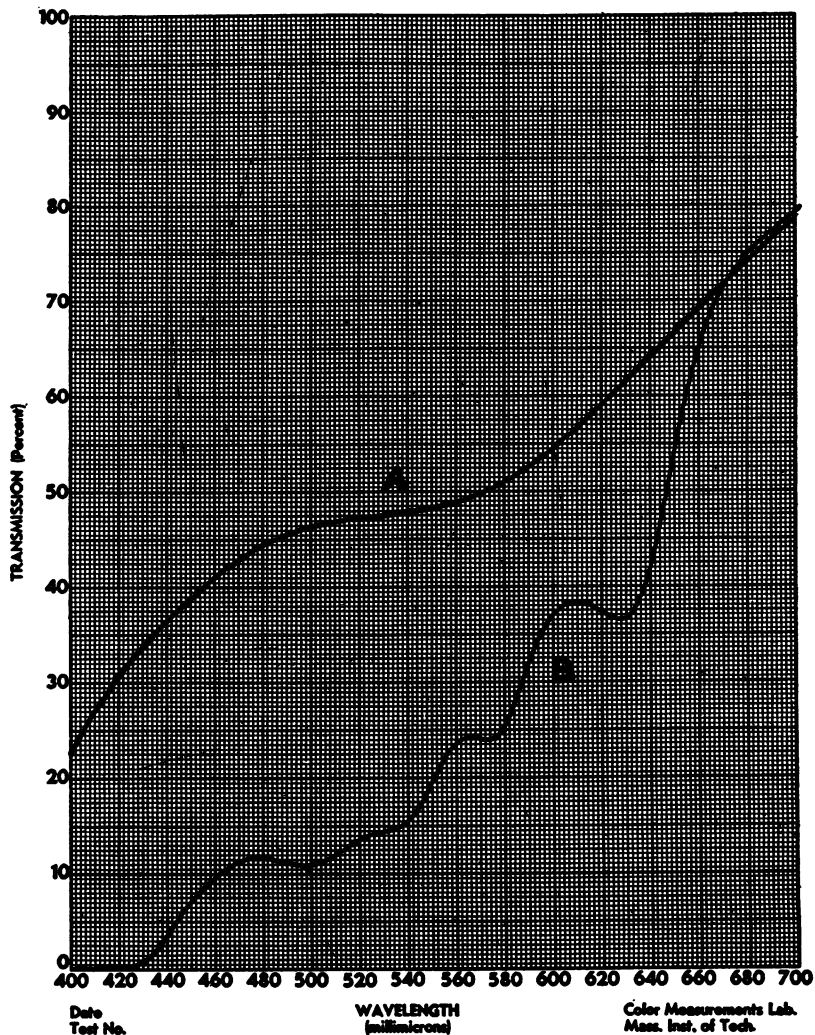


FIG. 2. FORMATION OF METHEMOGLOBIN BY PHOTO-OXIDATION PRODUCT OF SULFANILAMIDE

B is curve after reaction between the blue product of irradiation of sulfanilamide and human hemoglobin.

A is curve of brown solution resulting from spontaneous reduction in air of the blue oxidizing solution.

460 to 600  $m\mu$ . In this region, light absorption by hemoglobin is very great, obscuring small deviations. But in the red region beyond, where hemoglobin absorption is minimal, there was demonstrated a small residual absorption. This might be due to any brown substance<sup>8</sup> such as the brown reduced form of the blue oxidation product (9).

<sup>8</sup> An unidentified "foreign pigment" is just now reported in normal rats receiving sulfanilamide (10) and Harris and Michel (11) have also observed an extraneous absorption at 670  $m\mu$ .

This effect, however, was too slight to influence the gross appearance of blood. These observations suggest that *in vivo* sulfanilamide is oxidized to the blue derivative which in turn oxidizes hemoglobin to methemoglobin and is itself reversibly reduced. Although this residual absorption indicates in addition to hemoglobin and methemoglobin the existence of a third colored component, and while our spectrophotometric findings are consistent with our interpretation, equally satisfactory fit of the curves might per-

TABLE II  
Hemoglobin—blue substance reaction

$\lambda m\mu$	Experimental A	Calculated B	Calculated C
	$\log \frac{1}{T}$	$\log \frac{1}{T}$	$\log \frac{1}{T}$
700	0.118	0.024	0.118
660	0.211	0.094	0.182
630	0.455	0.454	0.455
620	0.460	0.428	0.460
600	0.478	0.396	0.478
580	0.699	0.603	0.638
560	0.693	0.702	0.656
540	0.886	0.885	0.827
520	0.886	1.00	0.876
500	0.959	1.15	0.959
480	0.943	1.05	0.931
460	1.04	1.23	1.07

Experimental A—obtained from spectrophotometer curve. Calculated B—on basis of 91.0 per cent methemoglobin and 9.0 per cent hemoglobin.

Calculated C—on basis of 10.8 per cent hemoglobin, 79.7 per cent methemoglobin and 9.5 per cent colored residue.

Italicized figures are reference points used in computation.

TABLE III  
Values of  $\log 1/T$  for sulfanilamide-treated patients

$\lambda m\mu$	Patient 1			Patient 2			Patient 3		
	A exper- imental	B cal- cu- lated	C cal- cu- lated	D exper- imental	E cal- cu- lated	F cal- cu- lated	G exper- imental	H cal- cu- lated	I cal- culated
700	0.032	0.025	0.088	0.024	0.021	0.084	0.0355	0.023	0.0854
660	0.052	0.046	0.052	0.033	0.032	0.033	0.056	0.047	0.059
630	0.159	0.169	0.169	0.090	0.090	0.090	0.177	0.177	0.177
620	0.169	0.161	0.164	0.100	0.098	0.098	0.180	0.176	0.180
600	0.268	0.237	0.245	0.197	0.184	0.185	0.260	0.236	0.250
580	1.68	1.74	1.80	1.64	1.75	1.75	1.59	1.50	1.56
560	1.47	1.45	1.40	1.444	1.440	1.446	1.29	1.26	1.32
540	2.16	2.14	2.17	2.15	1.92	2.12	1.93	1.86	1.92
520	1.16	1.19	1.20	1.08	1.093	1.095	1.09	1.09	1.10
500	1.05	1.05	1.05	0.92	0.92	0.92	0.99	0.99	0.99
480	1.24	1.21	1.21	1.12	1.11	1.11	1.14	1.11	1.13
460	1.80	1.85	1.87	1.70	1.75	1.76	1.69	1.66	1.69

Column A—Values of  $\log 1/T$  from experimental curve

Column B—Values calculated on basis of 80.2 per cent hemoglobin and 19.8 per cent methemoglobin

Column C—Values calculated on basis of 82.3 per cent hemoglobin, 17.4 per cent methemoglobin and 0.314 per cent colored residue

Column D—Values from experimental curve

Column E—Values calculated on basis of 90.3 per cent hemoglobin and 9.7 per cent methemoglobin

Column F—Values calculated on basis of 90.6 per cent hemoglobin, 9.3 per cent methemoglobin and 0.096 per cent colored residue

Column G—Values of  $\log 1/T$  from experimental curve

Column H—Values calculated on basis of 74.3 per cent hemoglobin and 25.7 per cent methemoglobin

Column I—Values calculated on basis of 78.0 per cent hemoglobin, 21.4 per cent methemoglobin and 0.573 per cent colored residue.

Italicized figures are reference points used in computation.

haps be obtained on the basis of different assumptions as to the nature of the pigments present. Our interpretation is consistent with the findings but cannot be considered as established by our data.

Further evidence that *in vivo* sulfanilamide is oxidized is provided by Dr. Sanford Rosenthal (12) who reported indications of hydroxylamine sulphonamide, an oxidation product, in the urine of sulfanilamide-treated animals and patients. Preliminary experiments by one of us (C. L. F.) indicate that the Rosenthal test (13) is positive in the methemoglobin forming colored irradiation products and negative in the unadsorbed colorless residual solution which does not form methemoglobin. The data are correlated with the mode of action of sulfanilamide (14).

#### SUMMARY

1. Objective evidence in the form of curves drawn by the Hardy recording spectrophotometer is presented showing the occurrence of methemoglobin and sulfhemoglobin in the blood of sulfanilamide-treated patients.

2. An explanation for the occurrence of methemoglobin in the blood of sulfanilamide-treated patients is provided by the demonstration *in vitro* of the conversion of hemoglobin to methemoglobin by certain oxidation products of sulfanilamide.

3. Spectrophotometric evidence for the occurrence *in vivo* of this reaction is adduced.

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