PHOTOGRAPHIC NATURE OF TANNING OF THE HUMAN SKIN AS SHOWN BY STUDIES OF MALE HORMONE THERAPY¹

THE condition of the skin is known to be influenced by the endocrine system, as for examples, the bronzing in Addison's disease of the cortex of the adrenal gland, the acne commonly associated with puberty and the pigmentation changes during pregnancy. Changes in the skin of a hypogonadal patient following the use of synthetic male sex substance testosterone propionate² have been reported.³ A further series of three surgically castrated⁴ and four hypogonadal males has been studied. Before treatment the skin in all but two cases was of a characteristic pasty, sallow color, gray and lacking in pink tinge. This was most pronounced in the castrated men. After treatment with testosterone propionate there was a rapid flushing, followed by increased oiliness of the skin and growth of hair on the face, chest, abdomen, arms and legs. All patients presented a more tanned appearance, particularly of the face, neck, hands and exposed parts of the skin. Part of this increased pigmentation is due to "developing" of pigment from previous exposure as indicated by the following history:

Case 1. Orchidectomy was performed in May of 1937, after which time the patient tanned but poorly and burned easily upon exposure to the sun. In August of 1937 he spent part of every morning and afternoon for a week lying on the beach clad in an abbreviated bathing suit of a peculiar cut. Only slight coloration resulted. The patient was examined by the authors several times during December and January, at which time the body skin was of a pasty sallow color. Treatment with male hormone substance was begun in the dead of winter, January 17, 1938. Within three weeks there appeared, along with the bronzing of the face, a tanning of the body save where it had been protected by a bathing suit. The patient had not worn the bathing suit, whose peculiar pattern the tan fitted, or any other bathing suit for five months. Neither had he sunned himself or used a sun-lamp in similar fashion.

Upon withdrawal of the hormone treatment in February of 1938 the flush disappeared from the skin and the tanned areas gradually faded. Subsequent injection and withdrawal periods induced, respectively, coloration and a fading of these areas which had not been exposed to sunlight since the previous August. These phenomena might be accounted for in part by assuming either that tanning consists of a continued production of pigment over a long

¹Supported in part by the International Cancer Research Foundation.

² Furnished by the Ciba Company under the trade-name Perandren.

³ James B. Hamilton, Endocrinology, 21: 649, 1937.

⁴ Neal E. Miller, Gilbert Hubert and J. B. Hamilton, Proc. Soc. Exp. Biol. and Med., 38: 538, 1938. period of time or that the melanin becomes colorless unless an adequate hormone supply be present.

Graded exposures to a sun-lamp have been given to this and to other patients and a similar series of pigmentation changes have been observed during periods of treatment and withdrawal.

From studies of men with low amounts of testicular secretion it appears that male hormone substance exerts a "developing" action upon the rather colorless material which is laid down in the skin following exposure to the sun or sun-lamp. This "developing" action may be exerted as late as five months after exposure. This indicates that tanning may be a "photographic-like process" of "exposure" and "development," with the sex hormone acting to "develop" color-lacking material laid down in the skin by exposure. Further, the pigmentation is not established continuously but will fade upon cessation of hormone treatment and reappear upon later resumption of treatment.

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ACTION OF VANADIUM ON TISSUE OXIDATIONS

IF 10-20 γ of vanadium in the form of sodium meta vanadate is added to rat or guinea pig liver suspension at pH 6.7 a large extra oxygen uptake occurs. This effect is much less in kidney and absent in brain. Concentration curves indicated that some substance in the liver was oxidized in the presence of vanadium. To prove this the following experiment was done. Rat liver was ground with buffer pH 6.7, squeezed through muslin and centrifuged. The solid was resuspended in 40 cc of water to which 10 cc buffer pH 6.7 was added and centrifuged again. This process was repeated four times and the resultant solid material which was free of hemoglobin and light yellow in color was finally suspended in 15 cc buffer pH 6.7. This constituted the enzyme preparation. The substrate was prepared as follows. A guinea pig liver was chopped and ground in buffer pH 6.7. Enough alcohol was added to make the final concentration 70 per cent. The precipitate was filtered and the alcohol evaporated off in vacuo at 40° C. The resulting suspension was extracted with ether three times, leaving a clear light yellow solution. This solution was then treated with a small amount of Lloyd's reagent and filtered. The filtrate was now almost colorless. It could now be evaporated down to dryness and extracted with boiling 95 per cent. alcohol. The alcohol extracts the substrate, which is redissolved in water after the alcohol is evaporated off. Table 1 gives the oxygen uptake in c.mm. of the various combinations of enzyme, substrate and vanadium after half an hour at pH 6.7 and 37° C. 0.5 ce enzyme suspension and about 10 per cent. of the amount of substrate present in one guinea pig liver was used in a total volume of 2.0 cc in the Warburg vessels.

TABLE 1

	O2 uptake c.mm.
Enzyme	0
Enzyme + vanadate	Ó
Vanadate + substrate	0
Enzyme + substrate	12
Enzyme + vanadate + substrate	119

Work on the chemical identification of the substrate is now proceeding. Experiments have shown that it is probably not an amino acid, amine, simple alcohol or aldehyde, purine, low fatty acid, choline, succinate, cholic acid, citrate, lactate, pyruvate, glucose or ascorbic acid. It is probably a phospholipid.

As vanadium is found in small traces in all tissues these results raise the question whether it has a normal catalytic function in the body and whether it is an essential element.

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A COMPARATIVE STUDY OF THE SUBTER-RANEAN MEMBERS OF THREE FIELD GRASSES

A COMPARATIVE study was made of the roots and root hairs in upper soil levels for oats, winter-rye and Kentucky bluegrass. Soil samples 3 inches in diameter and 6 inches deep (42 cubic inches) were taken from the fields by means of a cutting tube, and measurements made of the included subterranean plant parts. Total lengths of both roots and root hairs were used in computing the extent and surface exposed by the underground members. In Tables 1 and 2 the values given are the average of the three soil samples surveyed for each species.

TABLE	1
Roots*	

	Total	Total	Total root
	number of	length of	surface
	roots	roots (ft.)	(sq. in.)
Oats Rye Bluegrass	$4,700 \\ 6,400 \\ 84,500$	$150 \\ 210 \\ 1,260$	50 78 330

* Per soil sample (42 cubic inches).

In a comparison of the cultivated rye plants grown in competition with a non-competing greenhouse rye plant, previously surveyed,¹ it was found that the field

¹ H. J. Dittmer, Am. Jour. Bot., 24: 417-420, 1937.

TABLE 2

ROOT	HAIRS*
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,	Total num- ber of root hairs (in millions)	Total length of root hairs (miles)	Total root hair surface (sq. ft.)
Oats Rye Bluegrass	$\begin{array}{c} 6.3 \\ 12.5 \\ 51.6 \end{array}$	$\begin{array}{r} 4.9\\10.0\\32.0\end{array}$	3.7 8.2 16.9

* Per soil sample (42 cubic inches).

rye had approximately 5 times the number of root hairs per unit of root length as the non-competing greenhouse plant. However, the indoor plant had far more and longer roots, and consequently a greater total number of root hairs.

Assuming that roots and root hairs were evenly distributed throughout the samples, one cubic inch of soil from this oats field would have approximately 110 roots and 150,000 root hairs, with a combined length of about 630 feet and a surface area of 15 square inches. A similar cube of soil from a field of winter rye would have approximately 150 roots and 300,000 root hairs with a combined length of 1,300 feet and a surface of about 30 square inches. Kentucky bluegrass would have, per cubic inch of soil, approximately 2,000 roots and 1,000,000 root hairs, with a combined length of over 4,000 feet and a surface area of about 65 square inches. When it is considered that these grasses have from 150,000 to 1,000,000 root hairs per cubic inch of soil their importance in the physics of the soil is obvious. From the standpoint of their usefulness as soil binders oats would be least efficient, rye intermediate and bluegrass far superior to either of the others in retarding erosion.

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