DNA which the organism contains (the most extreme ATtype DNA $\left(\frac{A+T}{G+C}=2.70\right)$ found so far in bacteria is that

occurring in Clostridium perfringens var. Freds), because it would be expected that the DNA from a bacteria and its L form would have an almost identical composition.

We thank Prof. M. Stacey for his interest, Dr. D. G. ff Edward for discussions, Mr. E. T. J. Chelton for assistance and the Salters' Institute of Industrial Chemistry for a research fellowship (to R. T. W.).

A. S. JONES R. T. WALKER

Department of Chemistry,

University of Birmingham.

¹ Lynn, R. J., and Smith, P. F., J. Bact., 74, 811 (1957).

² Schmidt, G., and Thannhauser, S. J., J. Biol. Chem., 161, 83 (1945). ³ Jones, A. S., and Walker, R. T., J. Gen. Microbiol., **31**, 333 (1963).

⁶ Wyatt, G. R., Biochem. J., 48, 581, 584 (1951).
 ⁵ Watson, J. D., and Crick, F. H. C., Nature, 171, 737 (1953).
 ⁶ Ann. New York Acad. Sci., 79, Art 10, 465 (1960).
 ⁷ Ann. New York Acad. Sci., 79, Art. 10, 481 (1960).
 ⁸ Ki Yong Lee et al., Ann. Inst. Pasteur, 91, 212 (1956).

A New Transferrin in New Guinea

IN 1957 Smithies1, using the technique of starch-gel electrophoresis, described inherited variants of human 3-globulins. Smithies and Hiller² established the identity of these β -globulins with the iron-binding protein, transferrin, and this was afterwards confirmed with iron-59 and autoradiography³. During the past few years widespread sampling of human populations has demonstrated the existence so far of 14 transferrins. In order of decreasing mobility in starch-gel electrophoresis these are4: Bo, $B_{0^{-1}}$, B_1 , $B_{1^{-2}}$, B_2 , B_3 , C, D_0 , D_4 ($D_{0^{-1}}$), $D_{Montreal}$, D_{Chi} , D_1 , D_2 , D_3 . The present communication reports another transferrin of the B series.

The new protein was readily detected in one-dimensional vertical starch-gel electrophoresis with borate buffer using conditions described previously5. Comparative investigations with different transferrin types showed that it migrated faster than B_0 (Fig. 1), and is thus the most rapidly migrating transferrin thus far described. From the intensity of staining with naphthalene black the protein appeared to have a concentration slightly higher than that of transferrin C. Autoradiographs of gels made after adding iron-59 to the serum before electrophoresis revealed two bands corresponding in position to the two protein bands. The faster-migrating band can thus be regarded as a transferrin. Since this new transferrin was first discovered in the serum of a native from a village near Lae in New Guinea it is proposed that this variant be called BLae.

The example of B_{Lae} already referred to was the only one in a series of 136 samples of serum from inhabitants



Fig. 1. The β -globulin are of a vertical starch-gel comparing B_{Lae} with other transferrin variants. *a*, Transferrin B; *b*, transferrin C

Table 1. TRANSFERRIN FREQUENCIES IN SOME POPULATIONS IN NEW GUINEA

Locality of New Guinea natives	No. tested	BC		CC		CD1		D_1D_1	
		No.	%	No.	%	No.	%	No.	%
Madang Megier Moroba:	16	_		16	100.0	_	-	-	
Watut Lae Panua	28 17	1	5.9	27 12	96·4 70·6	$\frac{1}{3}$	3.6 17.6	1	5-9
Port Moresby Kerema Orokolo	19 33 23			14 27 14	73·7 81·8 60·9	5 6 8	26·3 18·2 34·8	$\frac{-}{1}$	

in various places in New Guinea investigated in the present survey. Transferrin D₁ was observed in all population samples except that from Megier (Table 1). Careful comparison of the mobility of the D_1 in New Guinea failed to discriminate it from the D_1 of African Negroes or the D₁ of Australian Aborigines. Since the population samples were small no attempt has been made to calculate the gene frequencies of the transferrin alleles; but it is obvious that D_1 is relatively common in New Guinea. This result is in agreement with other investigations of transferrins in New Guinea⁶.

Although only one example of BLae was found in the survey reported here we have been successful in obtaining fresh samples of serum from the propositus and members of his family. This has confirmed the presence of B_{Lae} in the person originally sampled and has shown his brother and sister both to be heterozygous BLacC and mother to be homozygous $B_{Lae}B_{Lae}$. Further investigations on the distribution of B_{Lae} in New Guinea are at present in progress.

This work was supported by the University of Western Australia Research Grants Committee and by the Commonwealth Scholarship and Fellowship Scheme under the Colombo Plan. I thank Dr. W. R. Pitney for supplying the original New Guinea samples, and Dr. J. L. Jameson and Dr. J. C. Muirden for collecting the samples from family members, Dr. E. Ezekiel and Mr. J. Firman for carrying out the autoradiography and Dr. R. L. Kirk for his interest in the work. I also thank Dr. A. G. Bearn and his colleagues at the Rockefeller Institute for confirming the observations on BLae.

L. Y. C. LAI*

Department of Zoology,

University of Western Australia.

* Present address: Biological Laboratory, Western Reserve University, Cleveland 6, Ohio.

¹ Smithies, O., Nature, 180, 1482 (1957); 181, 1203 (1958).

² Smithies, O., and Hiller, O., Biochem. J., 72, 121 (1959).

- ³ Giblett, E. R., Hickman, C. G., and Smithies, O., Nature, 183, 1589 (1959).
 ⁴ Parker, W. C., and Bearn, A. G., Science, 137, 854 (1962).

- ¹ Lal, L. Y. C., Austral. J. Sci., 23, 228 (1961).
 ⁴ Barnicot, N. A., and Kariks, J., Med. J. Austral., 2, 859 (1960). Bennett, J. H., Auricht, C. O., Gray, A. J., Kirk, R. L., and Lai, L. Y. C., Nature, 189, 68 (1961).

Structure and Plant Growth-regulating Activity of some 2-Benzothiazolyloxyacetic Acids and 2-Oxobenzothiazolin-3-ylacetic Acids

THE preparation¹ and plant growth activity² of a compound stated to be 2-benzothiazolyloxyacetic acid (I) have been described in the patent literature. It became apparent, during the course of work on this and related compounds, that this structure was incorrect and that the compound is actually 2-oxobenzothiazolin-3-ylacetic acid (II).



