theory of DNA structure<sup>12</sup> is provided by the constancy of molecular weight throughout the structural changes which we have described. In addition, however, we are led to wonder whether the observed phenomena do not point to the participation of water in stabilizing the structure of DNA. With this end in view, we are investigating the behavior of DNA in a variety of organic solvents with different hydrogen-bonding characteristics.

(12) C. A. Dekker and H. K. Schachman, Proc. Nat. Acad. Sci., 40, 89 (1954).

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## SEGREGATION COEFFICIENTS OF VARIOUS IM-PURITIES IN A SILICON TETRAIODIDE MATRIX Sir:

The demand for transistors and other semiconductor devices has stimulated considerable research in the preparation of ultra-pure silicon. It was found that zone-refining techniques<sup>1</sup> were inadequate for the removal of certain impurities, e.g., boron, in a silicon matrix,<sup>2</sup> and it was decided to approach the problem by preparing a suitable compound of silicon, subjecting this to such purification techniques as recrystallization, sublimation, and zone purification and ultimately decomposing it to elemental silicon. Thermodynamic calculations indicated that of the four tetrahalides, silicon tetraiodide decomposed most readily and that it lent itself best to zone-melting techniques because of its relatively high melting point (121.5-122.5°). Furthermore, it could be expected that the segregation coefficients for the various impurities in silicon tetraiodide would differ from those found in silicon.

Silicon tetraiodide was prepared by passing iodine vapor over Coleman and Bell Company ninety-eight per cent. pure silicon at 800° and the product was then crystallized from toluene. This material was used to fill Pyrex ampoules nine millimeters in diameter and thirty centimeters long. It was then densified, the tube sealed, and zonepurification was effected by vertical passage at the rate of five centimeters an hour. The zone width was two and one-half centimeters, and the temperature of the molten zone was about  $135^{\circ}$ . A small molten zone was passed through the charge only once in order to maintain impurities at spectrographically detectable levels. Spectrographic analyses of the successive two and one-half centimeter zones permitted preliminary calculations to be made based on Pfann's original equation.<sup>1</sup> The results of these calculations gave plots of  $C/C_0$ versus x/l (where C is the concentration of an impurity in a solid frozen from a mother zone,  $C_0$  is the mean concentration of the impurity before zone

(1) W. G. Pfann, Trans. AIME, J. Metals, 194, 747 (July, 1952).

refining, x is the distance that the zone has traveled along the tube, and l is the zone length) which indicated that certain impurities can be efficiently removed by this technique. In the case of boron where the concentration level was below the limit of spectrographic detectability, a mathematical extrapolation was employed based on the minimum spectrographically detectable concentration found. Using this value in conjunction with the expression:

$$C_{z} = K[C_{0} + (C_{z-1}/K - C_{z-1})]$$

where  $C_z$  is the concentration of an impurity frozen into the zth zone,  $C_{z-1}$  is the concentration of the impurity frozen into the previously frozen zone, k is the segregation coefficient of the impurity, and  $C_0$  is as described above, the maximum value for the k of boron was determined. The values of the k's determined for various metallic impurity species in the SiI<sub>4</sub> matrix are: boron, 0.16  $\pm$  0.07, aluminum, 0.88  $\pm$  0.04, sodium, 0.07  $\pm$  0.01, magnesium, 0.58  $\pm$  0.06, copper 0.63  $\pm$  0.05.

It is apparent from the above that the impurities listed can be removed to levels below the one part per million range by a suitable number of passes. Work on this project, and the method and/or methods of preparing pure silicon tetraiodide, as well as the preparation of the elemental silicon, is continuing.

AIR FORCE CAMBRIDGE RESEARCH CENTER ELECTRONICS RESEARCH DIRECTORATE LAURENCE G. HANSCOM FIELD BERNARD RUBIN BEDFORD, MASSACHUSETTS GUY H. MOATES RECEIVED JANUARY 9, 1956

## GLUCOSIDURONIC ACID SYNTHESIS BY $\beta\text{-}GLUCU\text{-}$ RONIDASE IN A TRANSFER REACTION

Sir:

Certain biological phenomena have been correlated with the activity of the enzyme  $\beta$ -glucuronidase. Among these are glucuronidogenesis,<sup>1</sup> action of gonadal hormones,<sup>2,3,4,5</sup> human cancer,<sup>6,7,8</sup> genetic control in mouse tissues,<sup>3,9,10</sup> and effects of pituitary interstitial cell stimulating hormone.<sup>11</sup> In attempts to arrive at an interpretation of the function of the enzyme *in vivo*, we have found it difficult to explain the findings on the basis of a purely hydrolytic action of the enzyme or its simple reversal. It was postulated that  $\beta$ -glucuronidase participates as a member of a multicomponent system concerned with glucosiduronic

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(2) W. H. Fishman, ibid., 159, 7 (1947).

(3) W. H. Fishman and M. H. Farmelant, Endocrinology, 52, 536 (1953).

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(5) W. H. Fishman, in 'Vitamins and Hormones,' Vol. IX, Academic Press, New York, N. Y., 1951, p. 213.
(6) W. H. Fishman, A. J. Anlyan and E. Gordon, Cancer Research, 7,

(b) W. H. Fishman, A. J. Aniyan and E. Gordon, Cancer Research, 1, 808 (1947).
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(7) W. H. Fishman and R. Bigelow, J. Natl. Cancer Inst., 10, 1115 (1950).

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(9) A. G. Morrow, E. M. Greenspan and D. M. Carroll, J. Natl. Cancer Inst., 10, 657 (1949).

(10) L. W. Law, A. G. Morrow and E. M. Greenspan, *ibid.*, 12, 909 (1952).

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<sup>(2)</sup> J. A. Burton, Physica, 20, 845 (Nov., 1954).