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An Isotope Effect in Photosynthesis

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Authors

Weigl, J.W.
Calvin, M.

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AN ISOTOPE EFFECT IN PHOTOSYNTHESIS

by

John W. Weigl and M. Calvin

23 November 1948

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AN ISOTOPE EFFECT IN PHOTOSYNTHESIS

by

John W. Weigl and M. Calvin

Radiation Laboratory and Department of
Chemistry, University of California,
Berkeley, California.*

23 November 1948

In the course of some kinetic studies on photosynthesis of barley seedlings, it has been found that plants utilize $C^{12}O_2$ faster than $C^{14}O_2$. The plants were placed in a closed system containing an infra-red absorption-cell for the analysis of total CO_2 and an ionization chamber for the determination of $C^{14}O_2$ in the gas phase, both instruments recording continuously.

Carbon dioxide, containing about 2% $C^{14}O_2$, was introduced in the dark, and the specific activity at this point taken as unity. After a short dark period, the lights were turned on and photosynthesis was allowed to take place. Figure 1 shows the result of a typical experiment.

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During the initial dark period the specific activity fell because of dilution by inactive respired CO_2 . However, as photosynthesis proceeded, the specific activity of the residual CO_2 rose until, when only 1/6 of it remained, the specific activity reached a peak some 20% higher than it had been at the start of photosynthesis. At this point the steady respiratory dilution became an appreciable fraction of the total remaining CO_2 , and the specific activity dropped rapidly.

Two other possible explanations of the rise in specific activity were:

- 1) a difference in the response times between the instruments of the order of a minute or two.
- 2) the conversion of the CO_2 into some other volatile compound.

Independent determinations of the response time to changes in the system revealed no appreciable time lags. The interpretation as an isotope effect was confirmed by removing samples of CO_2 during the course of a run and determining its specific activity by counting as BaCO_3 , reconvertng to CO_2 , and measuring in a separate ionization chamber.

A similar selectivity favoring C^{12}O_2 over C^{13}O_2 has been observed by Nier (1) and Urey (2) in steady state systems, in which the net effect is considerably smaller than that reported here.

The authors wish to acknowledge the valuable aid of Mr. Paul M. Warrington in these experiments.

(1) A. O. Nier and E. A. Gulbransen, J. Am. Chem. Soc., 61, 697 (1939);
B. F. Murphey and A. O. Nier, Phys. Rev., 59, 771 (1941).

(2) H. C. Urey, Science, 108, 489 (1948).

