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FORTY YEARS OF PLANT PHYSIOLOGY

SOME GENERAL IMPRESSIONS¹

By Professor EDWIN C. MILLER

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FORTY years ago the work in plant physiology was changing from the old to the new. Those who were interested in the subject were concerned chiefly with the nature of the response of *Mimosa* or similar plants to stimuli of various sorts. In the main, they were not interested at all in any practical or applicable results that might accrue from their investigations. At about the time of my entrance into the field, the conflict between the purist and the practical man was at its height and was being waged bitterly. It is said that some purist when asked of what practical value his findings were in the field of science, replied, "None

¹ Address of the retiring president of the American Society of Plant Physiologists, December, 1942.

Contribution No. 443, Department of Botany, Kansas Agricultural Experiment Station.

whatsoever and if I had thought before undertaking the work that they would be of any practical value, I would never have undertaken the investigation." Such a happening may be somewhat exaggerated, but it, nevertheless, illustrates the state of mind of some of the individuals who waged this bitter conflict. This condition illustrates the same spirit that was expressed by the so-called "malefactors of great wealth" who are reputed to have said that "the public could be damned" as far as they were concerned.

The public, rightly or wrongly, may eventually reach the stage where the workers not only in plant physiology, but also in most other lines of scientific work, must show that the results of their labors will contribute to the happiness or advancement of man-

it, and finally centrifuging it at 5,000 r.p.m. in an angle-head centrifuge and discarding the sediment. These preparations were then rendered non-virulent by exposure to the rays of a mercury arc lamp for a determined period of time.²

Freezing and drying of these antigens are accom-

same in hyperimmune serum, whether virulent or frozen and dried antigen was employed.

Similar antigens with St. Louis and West Nile viruses have been prepared and tested. They have proved practically identical to the virulent antigens in antigenicity and specificity. Our observation of their

TABLE 1
EFFECT OF IRRADIATION AND FREEZING AND DRYING ON THE COMPLEMENT-FIXING ANTIGEN OF
WESTERN EQUINE ENCEPHALOMYELITIS VIRUS

(Antigen irradiated for 100 minutes; tested for virulence by intracerebral inoculation into Swiss mice. Out of 12 mice inoculated none died).

Antigen	Anti-complementary power*	Antigenicity†							Specificity and titer of serum‡							
		1:1	1:2	1:4	1:8	1:16	1:32	1:64	1:1	1:2	1:4	1:8	1:16	1:32	1:64	1:128
No antigen; saline control	0.13 cc								0	0	0					
Western equine encephalomyelitis, non-irradiated .	0.13 cc	4¶	4	4	4	0	0	0	4	4	4	4	4	4	1	0
Western equine encephalomyelitis, irradiated and lyophilized	0.13 cc	4	4	4	4	0	0	0	4	4	4	4	4	4	2	0
West Nile, irradiated and lyophilized	0.13 cc	0	0	0	0	0	0	0	0	0	0	0	0	0	0	0

* Amount of guinea pig serum in dilution of 1:30 equivalent to one unit.

† Dilution of antigen reacting with a constant amount of Western equine encephalomyelitis mouse immune serum.

‡ Dilution of Western equine encephalomyelitis mouse immune serum reacting with a constant amount of antigen.

¶ 4 = Complete fixation; 0 = complete hemolysis; 1, 2 = intermediate degree of hemolysis.

plished in the following manner. The irradiated antigen is freed of sediment by centrifugation in a horizontal centrifuge for 10 minutes at 2,500 r.p.m. To the clear supernatant, merthiolate in a dilution of 1:10,000 is added. The antigen is then pipetted in 2 or 5 cc quantities into glass ampoules, frozen quickly by immersion into a dry ice-alcohol mixture, and dried over a period of 20 hours in a Flosdorf-Mudd apparatus,³ after which the ampoules are sealed. The ampoules containing the desiccated antigen in an air-free space are stored at 2° C. When needed for use in tests the ampoules are opened and 2 or 5 cc of distilled water added to the desiccated material. A similar method was employed by Smadel and Wall for preserving spleen lymphocytic choriomeningitis antigen.⁴

In the following table the results of one test with an ampoule of irradiated frozen and dried Western equine encephalomyelitis antigen are shown. The figures in the first column indicate that the titer of the complement was the same with frozen and dried as with standard virulent antigen or saline control (0.13 cc), showing that the dried material was not anti-complementary. Column 2 indicates that the titer of dried and virulent antigens was the same, 1:8, showing that there was no loss of antigenicity from irradiation or freezing and drying. Finally, column 3 indicates that the fixation titer was the

keeping qualities is limited thus far to 6 weeks, but we believe that in view of universal experience with lyophilized materials, these antigens will maintain their specific properties for long periods of time.

J. CASALS

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BOOKS RECEIVED

- Carnegie Institution of Washington Publication 524. Papers from Tortugas Laboratory. Vol. xxxiii. Illustrated. Pp. 195. \$1.50, paper cover; \$2.00, cloth binding.*
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- GENTRY, HOWARD SCOTT. *Rio Mayo Plants, A Study of the Flora and Vegetation of the Valley of the Rio Mayo, Sonora. Illustrated. Pp. vii + 328. Carnegie Institution of Washington. \$2.25, paper cover; \$2.75, cloth binding.*
- GOLDBERG, LEO and LAWRENCE H. ALLER. *Atoms, Stars and Nebulae. Illustrated. Pp. v + 323. The Blakiston Company. \$2.50.*
- HAILE, FATHER BERARD. *Origin Legend of the Navaho Flintway. Pp. 319. University of Chicago Press. \$3.00.*
- HUDDLESON, I. FORREST and OTHERS. *Brucellosis in Man and Animals. Illustrated. Pp. xix + 377. The Commonwealth Fund. \$3.50.*
- PENNEY, A. C. *Everyday Electricity and Magnetism. Illustrated. Pp. vii + 248. Longmans, Green and Co. \$1.75.*
- SUTTON, RICHARD L. and RICHARD L. SUTTON, JR. *Synopsis of Diseases of the Skin. Illustrated. Pp. 481. C. V. Mosby Company.*
- ZOETHOUT, W. D. *Laboratory Experiments in Physiology. Illustrated. Pp. 256. C. V. Mosby Company.*

³ Earl W. Flosdorf and Stuart Mudd, *Jour. Immunol.*, 29: 389-425, 1935.

⁴ J. E. Smadel and M. J. Wall, *Jour. Bact.*, 41: 421-30, 1941.



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