viewpoint that cells containing actively growing or dividing nuclei are more susceptible to the mutagenic action of various agents.

Regardless of the interpretation of the above facts, the results serve to point out the necessity of controlling the age and degree of hydration of conidia in studies concerned with the production of mutations by chemical agents.

References

- BEADLE, G. W., and TATUM, E. L. Amer. J. Bot., 1945, 32, 678.
- MCELROY, W. D., CUSHING, J. E., and MILLER, H. J. cell. comp. Physiol., in press.
- MITCHELL, H. K., and HOULAHAN, M. B. Amer. J. Bot., 1946, 33, 31.

The Chemical Nature and Origin of *Phaseolus* Virus 2 Crystalline Inclusions

SAUL RICH1

Department of Botany, Oregon State College

Isometric, crystalline inclusions were described by Mc-Whorter (\mathcal{S}) as occurring in plants infected with *Phaseolus* virus 2. In the present study, qualitative chemical tests were performed on epidermal peelings from broad bean (*Vicia faba*) plants infected with this virus. The fresh integumentary strips were examined under the microscope to find a field containing well-developed crystalline inclusions. The reagent solutions were allowed to flow under one edge of the cover slip as the original mounting material was withdrawn from the opposite edge with a blotter. Fixed material was also tested in some cases. The observations are summarized in Table 1.

In broad bean, there exists a melanin-producing system which may be initiated by wounding. This is made apparent by the discoloration of wounded tissue, which changes in color from normal green to red, and finally to black. Dark areas appear on detached broad bean leaves when they are gradually killed by soaking in physiological saline solution for 24 hrs or longer. The only microscopically visible cytological change in affected epidermal cells was the dark pigmentation of the nuclei. Mosaic-diseased broad bean leaves subjected to this treatment showed the discoloration not only in the nuclei, but also in the crystalline inclusions. The nuclei and inclusions appeared as if treated with hematoxylin. These results suggest the presence of tyrosine in both the nuclei and inclusions, since melanin is presumed to be the endproduct of a tyrosine-tyrosinase system (1).

The solubility of the crystalline inclusions in both acid and alkali indicates an amphoteric substance. Naturallyoccurring amphoteric compounds which are insoluble in fat solvents are limited largely to proteins. This fact,

¹Now with the Department of Plant Pathology, Connecticut Agricultural Experiment Station, New Haven, Connecticut. combined with the results obtained with picric acid, the biuret test, Millon's reagent, and the observation of melanin production, shows that the isometric crystalline inclusions of *Phaseolus* virus 2 are proteinaceous.

The crystalline inclusions are found within the nucleolus and cytoplasm only. The primary production of the inclusions within the nucleoli suggests that the crystals are either partly or entirely composed of nucleolar material. The presence of the crystals within the cytoplasm would argue against this theory, were it not for the work of Lenoir (2), who reported the secretion of nucleolar fragments from the dividing nucleus into the cytoplasm. It seems reasonable to conclude that the isometric crystalline inclusions of *Phaseolus* virus 2 may be the insoluble end-product of the interaction of virus material and nucleolar material. This theory is further supported by the lack of visible crystals in infected *Melilotus alba*. In this plant the nucleoli are extremely small. The small

TABLE 1

SUMMARY OF CRYSTALLINE INCLUSION REACTIONS

Chemical	Effect
Nitric acid (conc.) Sodium hydroxide (10%)	Crystalline inclusions dissolved """"""
Picric acid (sat. aq.)	Nuclei and crystalline inclusions stained light yellow
Biuret test	Inclusions positive for polypeptide linkage
Millon test	Inclusions positive for phenolic group (tyrosine?)
Formaldehyde (5%)	Slight yellowing of inclusions*
Water (100° C)	No visible effect
Alcohol (95%)	No visible effect ; tested on fixed material
Tertiary butyl alco- hol	No visible effect ; tested on fixed material
Xylol	No visible effect ; tested on fixed material
Dioxan	No visible effect ; tested on fixed material
Ether (10%)	No visible effect ; tested on fixed material
Feulgen's reagent	Crystalline inclusions remained un- stained; nuclear chromatin stained red
Acetocarmine (in 45% HAc)	Crystalline inclusions dissolved; nuclear chromatin stained red
Iodine green (10%)	Crystalline inclusions stained dull pink; nuclei stained green†

* McWhorter (4) found that inclusions and nuclei fixed in 5% formalin would not stain with trypan blue without previous peptonization in 10% citric acid.

 \dagger Purdy (5), using iodine green to stain tobacco infected with tobacco mosaic, found the amorphous inclusions stained dull pink, while the nuclei stained green.

amount of one of the interacting substances (nucleolar material) would, in this case, restrict the development of the end-product (crystalline inclusions).

References

- 1. GORTNER, R. A. J. biol. Chem., 1911, 10, 89-94.
- 2. LENOIR, M. C. R. Acad. Sci., Paris, 1925, 180, 160-163.
- 3. MCWHORTER, F. P. Phytopathology, 1941, 31, 760-761.
- 4. MCWHORTER, F. P. Stain Technology, 1941, 16, 143-149.
- 5. PURDY, H. A. Amer. J. Bot., 1928, 15, 94-99.