

CORRESPONDENCE

Avery in Retrospect

SIR,—Dr H. V. Wyatt has drawn attention to the muted manner in which Avery, MacLeod and McCarty expressed themselves in 1944 on the significance for genetics of their work on the transforming principle of pneumococcus¹. He has exposed the tendency of later commentators to read more “knowledge” into the statements of “information” than is perhaps justified. At the same time it is only fair to the three Rockefeller scientists to state Avery’s reasons for the narrow delimitation of this work and for the non-committal discussion of its significance. These features can be highlighted by a comparison of the 1944 paper with papers written by André Boivin and his collaborators who worked first at the Pasteur Institute then at the University of Strasbourg.

In May 1943 O. T. Avery wrote to his brother Roy a famous letter from which Dr Wyatt quotes the phrase “Sounds like a virus—may be a gene”. Avery then added, as if hastily: “But with mechanisms I am not now concerned. One step at a time and the first step is, what is the chemical nature of the transforming principle? Someone else can work out the rest. Of course, the problem bristles with implications”². He went on to assure his brother that a lot of well documented evidence was needed before anyone could be convinced that protein-free DNA had the properties he claimed. This was the task he undertook. Just how the DNA acted was a separate question, the answer to which would clarify the biological significance of transformation. In this connexion it should be borne in mind that although Robinow had demonstrated nuclear structures in rod-shaped bacteria in 1942³, no case of conjugation in bacteria had been reported in 1944, and it was not until 1946⁴ that Lederberg and Tatum had good evidence of bacterial recombination. Furthermore Avery had to contend with the traditional interpretation of bacterial transformation as given by Neufeld and Levinthal in 1928⁵, Dawson in 1930⁶, Alloway in 1933⁷—that the recipient cells have retained the power to elaborate the capsular polysaccharides of several types of pneumococci and need only the stimulus of the transforming principle, this being specific for the development of only the donor type coat. The 1944 interpretation was an advance in this position.

It was the enzyme studies which formed the bulwark of Avery’s case and accordingly his efforts with McCarty, after the 1944 paper, were directed at improving this evidence⁸. Here it must be conceded that trypsin and chymotrypsin alone are inadequate as agents to remove all possible types of protein from the transforming substance. Unfortunately they could not use pepsin because the DNA was damaged at the pH required for its action⁹. Here pronase, had it been known at that time, could have filled the gap. Also none of these enzymes effectively digests protein until it is denatured, hence it was possible for Mirsky to find a weak point here when he attacked the evidence from enzymology in 1947¹⁰.

In contrast to Avery, André Boivin gave all too few details of his work with Vendrely and Lehout at the Pasteur Institute on the transformation of *Escherichia coli* types. At a meeting of the Academie des Sciences in November 1945 these workers claimed to have obtained results like Avery’s but using *E. coli*¹¹. In 1942, stimulated by the work of Dawson, Sia and Alloway, they had tried to effect transformation of types in *E. coli* and before seeing the 1944 paper of the Rockefeller scientists they had come to the conclusion that the transforming principle was a nucleoprotein. When they learnt of this work they removed the protein from their nucleoprotein autolysates of *E. coli* and found the nucleic acid residues still capable of transformation. At the time this work was regarded as an extension of the pneumococcus case to another bacterial species. Only later was doubt thrown on Boivin’s work after subsequent attempts to reproduce it failed¹².

To the historian it is of interest that Boivin was prepared to go much further than Avery in his interpretation of the work. The title of his paper in *Experientia* contains the phrase “Significance for the Biochemistry of Heredity”. In the conclusion the discovery of the identity of the transforming principle is described as opening “new horizons”, promising for the biochemistry of heredity. “In particular, it is on the side of the nucleic acid and not at all on that of the protein of the nucleoprotein macromolecule constituting a gene that one must find the basis for the inductive properties belonging to the gene”¹³. When Boivin attended the Cold Spring Harbor Symposium on “Nucleic Acids and Nucleoproteins” in June 1947, he

gave a remarkable paper¹⁴ in which he related the work on bacterial transformation to Beadle and Tatum’s work on biochemical genetics, described Tulasne’s confirmation of Robinow’s work on the bacterial nucleus¹⁵, and the chemical mechanism involved (Vendrely and Lipardy¹⁶), and gave Tulasne’s and Vendrely’s cytochemical evidence, using RNase and DNase, for the localization of RNA in the bacterial cytoplasm and DNA in the nucleus of *E. coli*¹⁷.

When we look back over the mass of literature in the 1940s it seems scarcely possible that André Boivin could have so accurately predicted the structure which the nascent subject of molecular genetics was to take. Consider the following statement¹⁴: “We may, at the most, catch a glimpse of a series of catalytic actions which set out from primary directing centres (the deoxyribonucleic genes) proceed through secondary directing centres (the ribonucleic microsomes-plasma-genes) and thence through tertiary directing centres (the enzymes), to determine finally the nature of the metabolic chains involved, and to condition by this very means, all the characters of the cell in consideration . . .”.

Although the Avery, MacLeod and McCarty paper was published in a journal with a fairly limited readership, the subsequent papers in New York, Atlantic City, Hershey (Pennsylvania), and Cold Spring Harbor in 1946, and in the latter again in 1947, brought bacterial transformation to the attention of a wide audience. In addition Luria, Dobzhansky and Burnet visited Avery personally in the 1940s. In war-torn Europe conditions were not conducive to the public discussion of Avery’s work, yet in Paris André Lwoff and Boris Ephrussi held a colloquium with support from the Rockefeller Foundation at which the new work which had had its genesis in Avery’s discovery was reported.

This new work concerned the demonstration that bacterial transformation was not confined to one hereditary characteristic and that DNA did have the properties required of the hereditary substance. When we see Hotchkiss’s and Chargaff’s evidence against the tetranucleotide hypothesis, Chargaff’s demonstration of the species specific base composition of DNA, and the Boivin Vendrely Rule governing the DNA content of diploid and haploid cells as the fruit of work initiated by the

Avery, MacLeod, McCarty discovery, it is no longer possible to maintain that their paper was either ignored or unknown. When we see how little was known about genetic processes in bacteria and the chemistry of DNA in 1944 compared with 1950, Avery's caution can be seen as justified. As for the geneticists, it is clear that what caught their imagination was not the identity of the transforming principle—whether it was nucleic acid or nucleoprotein did not mean a great deal to them—but the possibility, at last, to bring about a given hereditary change by a specific treatment. Hence the reason for the widespread habit of referring to bacterial transformation as “directed mutation”.

H. J. Muller was exceptional among geneticists in being concerned about the chemical identity of Avery's transforming principle, but was impressed by Mirsky's opinion. To Darlington he wrote: “. . . Mirsky gave reasons for believing that Avery's so-called nucleic acid is probably nucleoprotein after all . . .”¹⁸. Yet again, what attracted Muller was the possibility of fitting transformation into the grand scheme of cytogenetics. In the transforming substance, he suggested, there were chromosomal fragments consisting of nucleoprotein, which were incorporated into the genetic apparatus of the recipient bacterium in transformation.

As in the case of Mendel's paper, the scientists of a given period found in Avery's paper what they were looking for, but unlike that earlier case, the 1944 paper was not ignored or unknown. It posed questions about DNA which by 1950 could be answered. What Avery failed to say, Boivin said, but his brave words did not profoundly alter the climate of opinion until the Boivin Vendrely Rule was established¹⁹. Only with the advantage of hindsight can we see the significance of the 1944 paper as obvious. Only by confining our

attention to the published record and the citation statistics on Avery's paper can we arrive at the view that it was little known or undervalued.

Yours faithfully,

R. OLBY

Department of Philosophy,
University of Leeds,
Leeds LS2 9JT

- ¹ Wyatt, H. V., *Nature*, **235**, 86 (1972).
- ² Hotchkiss, R. D., in *Phage and the Origins of Molecular Biology* (edit. by Cairns, J., Stent, G. S., and Watson, J. D.) (Cold Spring Harbor Laboratory, 1966).
- ³ Robinow, C. F., *Proc. Roy. Soc., B*, **130**, 299 (1942).
- ⁴ Lederberg, J., and Tatum, E. L., *Nature*, **158**, 558 (1946).
- ⁵ Neufeld, F., and Levinthal, W., *Z. Immunitätsforsch. Exp. Ther.*, **55**, 324 (1928).
- ⁶ Dawson, M. H., *J. Exp. Med.*, **51**, 143 (1930).
- ⁷ Alloway, J. L., *J. Exp. Med.*, **57**, 265 (1933).
- ⁸ McCarty, M., and Avery, O. T., *J. Exp. Med.*, **83**, 89 (1946).
- ⁹ Avery, O. T., MacLeod, C., and McCarty, M., *J. Exp. Med.*, **79**, 137 (1944).
- ¹⁰ Mirsky, A. E., *Cold Spring Harbor Symp. Quant. Biol.*, **12**, 16 (1947).
- ¹¹ Boivin, A., Vendrely, R., and Lehoul, Y., *C. R. Acad. Sci., Paris*, **221**, 646 (1945).
- ¹² Avadhani, N.-G., Mehta, B. M., and Rege, D. V., *J. Mol. Biol.*, **42**, 413 (1969).
- ¹³ Boivin, A., Delaunay, A., Vendrely, R., and Lehoul, Y., *Experientia*, **1**, 334 (1945).
- ¹⁴ Boivin, A., *Cold Spring Harbor Symp. Quant. Biol.*, **12**, 7 (1947).
- ¹⁵ Tulasne, R., *CR Seanc. Soc. Biol.*, **141**, 411 (1947).
- ¹⁶ Vendrely, R., and Lipardy, J., *C. R. Acad. Sci., Paris*, **223**, 342 (1946).
- ¹⁷ Tulasne, R., and Vendrely, R., *C. R. Seanc. Soc. Biol.*, **141**, 674 (1947).
- ¹⁸ Letter from H. J. Muller to C. D. Darlington dated March 2, 1946.
- ¹⁹ Boivin, A., Vendrely, R., and Vendrely, C., *C. R. Acad. Sci.*, **226**, 1061 (1948).

Peregrines and Propaganda

STR.—Dr Cramp writes that “there is more than propaganda to justify the

belief that the persistent pesticides led to striking declines in peregrines” (*Nature*, **238**, 475; 1972). He then refers to evidence, as if the existence of evidence sufficed to make the case, irrespective of contrary evidence or unsoundness. But rarely is there absolutely no evidence behind propaganda. In this case, the evidence was examined by the Wilson Committee¹ and the Mrak Commission² and found to be inadequate.

Dr Cramps chides me for not quoting the Wilson Report for the suggestion that dieldrin was responsible; but that was just a suggestion and not altogether convincing. In this country, for most purposes the small tonnages of dieldrin and DDT used could be replaced by other insecticides, as recommended by the Wilson Committee, because there were few disadvantages in doing so. But the balance of advantage would be quite different in some countries, where the lives and happiness of many millions of human beings would be put at risk by abandoning DDT and dieldrin. In those countries, much more rigorous examination of and search for evidence would be essential and mere suggestions ought not to be lightly accepted. In particular, in this country and in North America, suggestions and insufficient evidence are converted by propaganda into beliefs³; such beliefs will be accepted by malarious countries at their peril.

Yours faithfully,

D. L. GUNN

Chilham,
Kent

¹ *Advisory Committee on Pesticides and other Toxic Chemicals*, 148 (HMSO, 1969).

² *Report of the Secretary's Commission on Pesticides and their Relationship to Environmental Health*, 677 (US Department of Health, Education and Welfare, 1969).

³ Gunn, D. L., *Ann. Appl. Biol.*, **72** (in the press).

HOW TO BUY NATURE

Volumes start in January, March, May, July, September and November, but subscriptions may begin at any time.

The direct postal price per subscription is:

12 MONTHS* (52 issues per title)

	Surface mail UK and worldwide	U.S.A. and Canada
Nature (Friday)	£14	\$48
Nature + Nature Physical Science	£24	\$83
Nature + Nature New Biology	£24	\$83
All three editions	£29.50	\$108
Annual Index	£1	\$3

*Rates for shorter periods *pro rata* (minimum three months)
(Charge for delivery by air mail on application)

Editorial and Publishing Offices of NATURE

MACMILLAN JOURNALS LIMITED
4 LITTLE ESSEX STREET, LONDON WC2R 3LF
Telephone Number: 01-836 6633. Telegrams: Phusis London WC2R 3LF
Telex 262024

711 NATIONAL PRESS BUILDING
WASHINGTON DC 20004
Telephone Number: 202-737 2355. Telex 64280

Subscription Department
MACMILLAN JOURNALS LIMITED
BRUNEL ROAD, BASINGSTOKE, HANTS
Telephone Number: Basingstoke 29242

American display advertisements
NATURE SCIENTIFIC PUBLICATIONS INC
711 NATIONAL PRESS BUILDING
WASHINGTON DC 20004

All other advertisements
T. G. SCOTT & SON, LIMITED
1 CLEMENT'S INN, LONDON WC2A 2ED
Telephone: 01-242 6264/01-405 4743
Telegrams: Textualist London WC2A 2ED
Registered as a newspaper at the Post Office

Copyright © Macmillan Journals Limited, September 29 1972