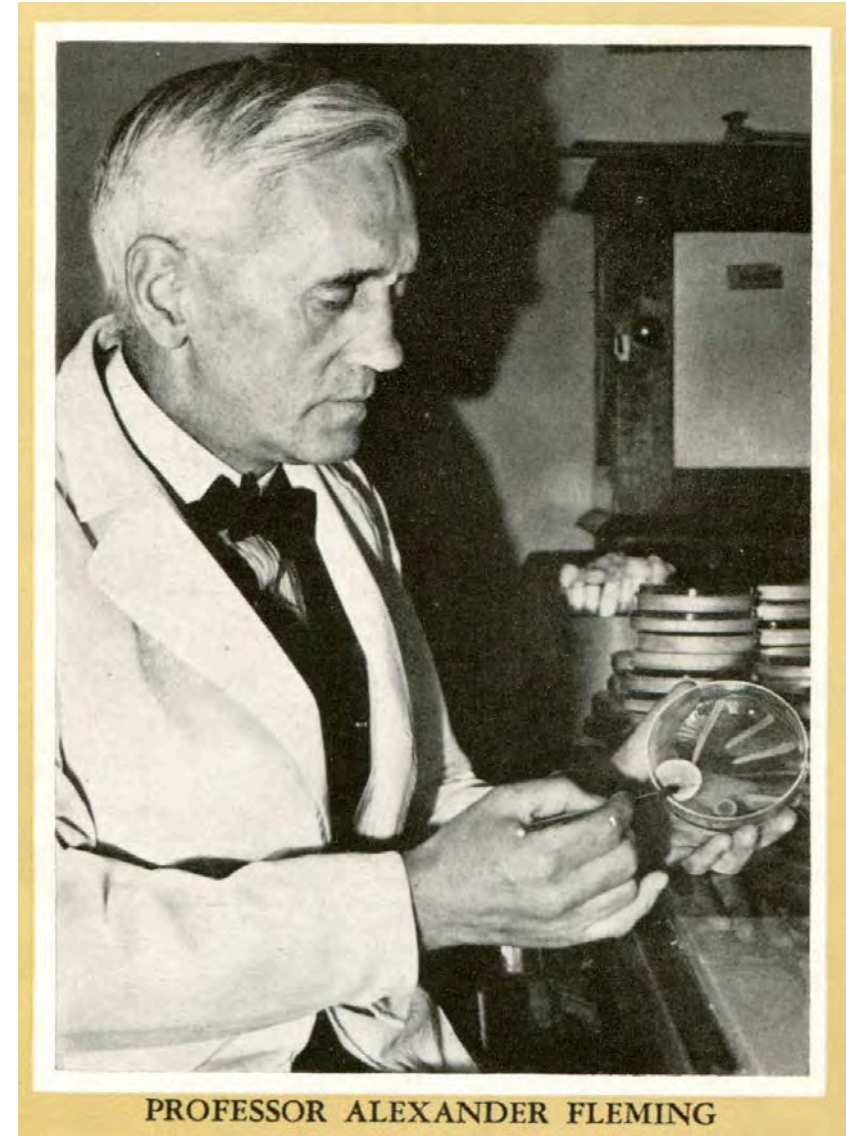


Alexander Fleming and Early Microbial Art

Illustrated by materials from the collections of
the Center for the History of Microbiology / ASM
Archives (CHOMA)



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Though most well-known as the discoverer of penicillin, Sir Alexander Fleming (1881-1955) was also a pioneer in the field of microbial art.

Six of his “paintings” were reproduced on the endpapers of Andre Maurois’s 1959 biography of the scientist.



Fleming attended the Second International Congress for Microbiology in London in 1936.

In addition to papers on “Selective Bacteriostasis” and “The Use of Stapylococcus Antitoxin, Toxin, Toxoid and Vaccine,” he offered this brief technical paper.



APPENDIX 1

Prof. A. Fleming (London): *The Growth of Micro-organisms on Paper.*

If a paper disc is placed on the surface of an agar plate the nutrient material diffuses through the paper sufficiently to maintain the growth of many micro-organisms implanted on the surface of the paper. At any stage growth can be stopped by the introduction of formalin. Finally the paper disc, with the culture on its surface, can be removed, dried and suitably mounted.

The nature of the paper makes a considerable difference to the result obtained. On filter-paper good growth takes place, but the extreme porosity of the paper makes the growth diffuse and, in the case of many moulds, the organism grows through the paper and adheres so firmly to the surface of the culture medium that the paper is torn in attempting to remove it from the surface of the medium. For most purposes a good stiff note-paper is suitable. The colonies remain discrete and the paper disc can be easily removed. If the colony is white a black paper disc can be used, but with chromogenic bacteria white paper is preferable.

The method is especially useful for making specimens for museum and teaching purposes. Single colonies of moulds in all stages of development can be preserved dry for an indefinite period. Permanent specimens illustrating such phenomena as selective bacteriostasis, droplet infection, etc. can easily be prepared.

Dried cultures of chromogenic bacteria have been preserved in this way for two years and the colours have not faded except in cases where the specimens had been exposed to light, especially bright sunlight. The colour of *B. prodigiosum* was especially

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sensitive to light and faded very considerably in a few months when exposed to diffuse sunlight on a laboratory wall. In similar conditions there was no perceptible fading of the cultures of staphylococcus, sarcina, or *B. violaceum*.

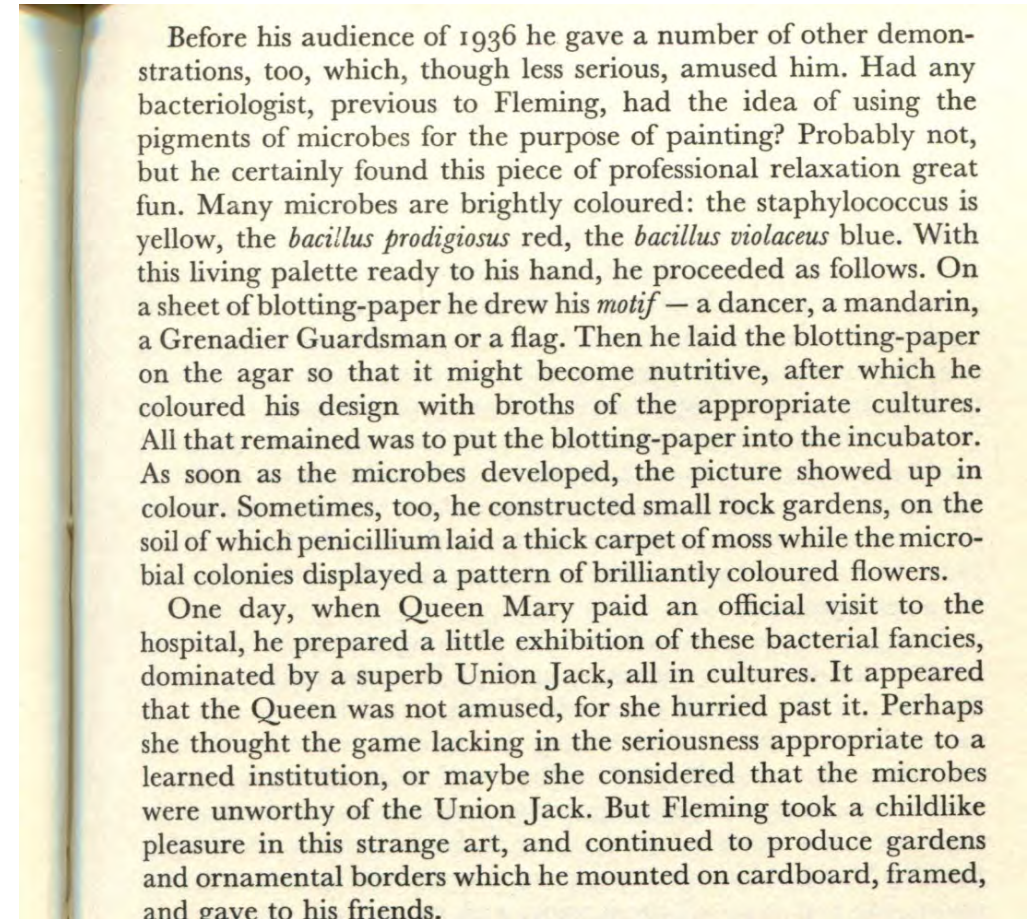
Here, Maurois discusses Fleming's technique

The Life of Sir Alexander Fleming. Translated from the French by Gerard Manley Hopkins

New York: E. P. Dutton & Co., Inc. 1959, p. 153



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This memento, prepared by Fleming using the technique described in the 1936 abstract, was presented to Dr. Kenneth Raper during Fleming's 1945 visit to America.

Raper (1908-1987) was a mycologist at the USDA's Northern Regional Research Laboratory in Peoria during World War II and directed the search for more productive strains of Fleming's *Penicillium notatum*. The result was the isolation of *Penicillium chrysogenum* NRRL 1951, which became the forerunner of all commercial strains.



“Functional Incubator – Daily Registration Totals Plated Ea[ch] Night”

This is from the May, 1955 Annual Meeting of the Society of American Bacteriologists (now ASM.) It may be that this practice, unique to this SAB meeting, was an homage to Fleming, who had died two months earlier.

