Convenient Method for Detecting $^{14}$CO$_2$ in Multiple Samples: Application to Rapid Screening for Mutants

HERBERT TABOR,* CELIA WHITE TABOR, AND EDMUND W. HAFNER

Laboratory of Biochemical Pharmacology, National Institute of Arthritis, Metabolism, and Digestive Diseases, Bethesda, Maryland 20014

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A procedure is presented for the rapid screening of bacterial colonies to detect mutants unable to produce $^{14}$CO$_2$ from a labeled precursor. The method is especially useful for mass screening for mutants that cannot be easily detected by their phenotypic characteristics.

The method described herein permits the rapid screening of many individual bacterial cultures for mutations in pathways producing $^{14}$CO$_2$ by trapping the $^{14}$CO$_2$ from each culture on paper saturated with barium hydroxide. This technique, combined with the procedures of Weiss and Milcarek (5) for the rapid isolation and cultivation of separate bacterial colonies in depression plates, permits the rapid screening of large numbers of separate colonies for the presence or absence of a given metabolic pathway producing CO$_2$. The use of Ba(OH)$_2$ paper takes advantage of the well-known insolubility of Ba$^{14}$CO$_3$, resulting in the fixation of $^{14}$CO$_2$ immediately above the colony in which it was formed. All of our studies have been with Escherichia coli, but the same method could be used for other bacteria, yeast, or mammalian cells in culture.

The method is as follows. (i) In each well of a Falcon 96-well plate (catalog no. 3040), place 100 μl of an appropriate minimal medium (with the use of a peristaltic pump) containing any necessary supplements and the appropriate $^{14}$C-labeled substrate. (ii) Inoculate the plate with organisms grown in another Falcon plate, with the help of the 48-prong stamping device described by Weiss and Milcarek (5). (iii) Immediately cover the plate with filter paper that has been impregnated with Ba(OH)$_2$ and cover this with a Falcon plastic lid (catalog no. 3041). (iv) The paper is prepared by wetting pieces of Whatman 3MM paper (11.7 by 7.5 cm) with saturated Ba(OH)$_2$, drying the papers by blotting with filter paper, and storing the papers in a desiccator over a KOH solution until use. Care must be taken to avoid prolonged contact of the paper with atmospheric CO$_2$. NaOH or KOH solutions cannot be used to wet the filter paper, as soluble carbonates are formed, which diffuse. (v) Secure the lid firmly on the dish with rubber bands. (iv) Wrap the plate in aluminum foil to limit evaporation, and incubate the cultures overnight at the desired temperature. (v) Remove the paper and immerse in acetone; remove the acetone by air-drying, followed by heating for 5 min at 110°C. (vi) Place the paper on Kodak RP/R-54 X-ray film for 1 to 7 days, depending on the number of counts used. (For a 24-h exposure, the specific activity of the $^{14}$C-labeled substrate added should be sufficient to produce 1,000 dpm of Ba$^{14}$CO$_3$ above each well.)

A typical radioautogram showing the use of the above method is presented in Fig. 1. In this experiment L-[U-$^{14}$C]methionine was added to the growth medium to detect mutants deficient in adenosylmethionine decarboxylase. The black spots are above the wild-type colonies, and the gray spots are above the mutants.

To screen for temperature-sensitive mutants, the following modification is suggested. The organisms are grown overnight without labeled substrate at the permissive temperature with limiting glucose (0.02%). Additional glucose is then added (it can be added with the stamping device used above, transferring ca. 3 μl of 20% glucose per well). The plate is then incubated at the permissive temperature for 1 h and at the nonpermissive temperature (e.g., 43°C) for 30 to 60 min; radioactive substrate (5 μl) is then added to each well with a repeating Hamilton syringe, and the plate is immediately covered with a piece of Ba(OH)$_2$-impregnated paper. After further incubation for 2 h at the nonpermissive temperature, a radioautogram is prepared as described above.

This method can also be used to assay a reaction that does not produce $^{14}$CO$_2$, provided that the product of the reaction can be coupled to another reaction that does produce $^{14}$CO$_2$. For example, during growth on [guanido-$^{14}$C]arginine, [$^{14}$C]urea is produced by the sequential action of arginine decarboxylase and
agmatine ureohydrolase, and the [14C]urea is excreted into the medium (2, 3). If urease is added to the growth medium, 14CO₂ is formed (2). We have used this method for detection of strains containing known mutations in agmatine ureohydrolase and have obtained patterns similar to those seen in Fig. 1.

The method has also been used for the qualitative assay of a reaction producing 14CO₂ in vitro. For this purpose, extracts are prepared in each well by either the lysozyme-ethylene diaminetetraacetic acid method of Weiss and Milcarek (5) or by treating the cells with toluene (4). With the latter technique, we have assayed for ornithine decarboxylase in ornithine decarboxylase-deficient mutants (1) in transduction experiments, with results similar to those in Fig. 1. This method may also be convenient for monitoring column eluants during the purification of any enzyme producing 14CO₂ either directly or in coupled reactions.

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LITERATURE CITED