Feasibility of Screening for Penicillinase-Producing Neisseria gonorrhoeae from Primary Culture Plates by Using a Rapid Microacidometric Test

ALICE SCHAUER WEISSFELD,1 GAIL D. SANNER,2 JACK R. CHILDERESS,1 JOHN D. DYCKMAN,1 THOMAS W. HUBER,2 AND ROBERT P. WILLIAMS1

Houston City Health Department Laboratory2 and Department of Microbiology and Immunology, Baylor College of Medicine,1 Houston, Texas 77030

Received for publication 13 September 1977

Penicillin is the drug of choice in the treatment of gonorrhea (3). Although strains of Neisseria gonorrhoeae that are relatively resistant to penicillin have been known for a number of years, the mechanism of this resistance was not thought to involve production of an enzyme capable of degrading the antibiotic (2, 8). Thus, penicillinase-producing N. gonorrhoeae (PPNG) isolated from infections contracted in the Far East and Liverpool, England, were unusual in that these organisms elaborated a beta-lactamase (β-lac) that conferred resistance to penicillin, ampicillin, and cephaloridine (1, 7).

As of 2 May 1977, 150 cases of disease caused by PPNG have been recognized in the United States (4). Determined efforts to locate and investigate the patients’ sexual contacts apparently have successfully controlled the spread of these gonococci. However, because of the special threat posed by these bacteria, especially in terms of increased treatment costs, rapid and accurate susceptibility tests are important. Testing of growth from primary plates would save time, but current recommendations are to test reisolated organisms to prevent false-positive β-lactam tests resulting from possible contaminants. Jorgensen et al. (5) recently reported a modification of an iodometric procedure for detecting β-lactam-producing N. gonorrhoeae directly from colonies growing on the primary culture plate. Their procedure is more rapid than other tests currently available, since no further growth is required after the initial isolation of the bacteria on Thayer-Martin medium. The possibility remains, however, that other β-lactam-positive (β-lac+) bacteria present in the specimen taken for the diagnosis of gonorrhea could cause false-positive results.

In this communication, we analyze the feasibility of screening for PPNG from primary culture plates by examining the role of possible β-lac+ contaminants in causing spurious reactions. We also describe an adaptation of a microacidometric method (9) for the assay of β-lac production in which the suspect gonococcal colony is removed for testing either before or after the addition of oxidase reagent.

MATERIALS AND METHODS

Media. Modified Thayer-Martin medium (MTM) contained GC medium base (Difco), 1% hemoglobin (Difco), 1% defined supplement (IsoVitaleX; BBL), vancomycin (3.0 µg/ml), colistin (7.5 µg/ml), nystatin (12.5 U/ml), and trimethoprim lactate (5.0 µg/ml). Chocolate agar contained the same constituents as MTM, except that the antimicrobial inhibitors were omitted. Heart infusion broth (Difco) was prepared according to the manufacturer’s directions.

Test organisms. A β-lac-negative (β-lac−) strain of N. gonorrhoeae was obtained from the Texas Department of Health Resources in Austin. This organism exhibited a zone of inhibition measuring 36 mm by the penicillin disk diffusion test. Two β-lac+ strains were kindly made available by Clyde Thornberry of the Center for Disease Control, Atlanta, Ga.; they were designated CDC3389 and CDC1782 by his laboratory. Both of these organisms showed no zone of inhibition by the penicillin disk diffusion test. Penicillinase-producing strains of
Escherichia coli and Proteus rettgeri were isolated from clinical specimens at the Hermann Hospital Microbiology Laboratory; \( \beta \)-lac+ strains of Staphylococcus aureus and Flavobacterium meningosepticum were isolated from clinical specimens at the Houston City Health Department Laboratory.

Preparation of \( \beta \)-lac test reagent and performance of the test. We used the microtiter modification of the acidometric test (9). If \( \beta \)-lac is present, penicillin is degraded to penicilloic acid that acidifies the medium and changes the color of the pH indicator. The indicator solution containing penicillin and phenol red was prepared as described previously (9). The test solution was either used immediately or dispensed in 2-ml samples in 1 dram (ca. 1.177 g) vials and frozen at \(-70^\circ\)C. It was used as needed and could be refrozen and rethawed until it turned yellow.

The test was performed by dispensing 0.1 ml of penicillin-phenol red solution into wells of a microtiter plate (Cooke Engineering), one well for each culture to be tested. \( \beta \)-Lac+ and \( \beta \)-lac- control organisms were included with each run. If the organism to be tested had been treated with oxidase reagent (N,N-dimethyl-p-phenylenediamine monohydrochloride; Eastman Kodak), control cultures likewise treated with oxidase reagent were included. Growth from one to five colonies of the bacteria was removed from the plates with a small bacteriological loop and suspended in the penicillin-phenol red solution. If the cultures produced \( \beta \)-lac, the solution usually turned yellow within 5 min; if the culture did not produce \( \beta \)-lac, the test solution remained dark red. The plates were allowed to stand for at least 30 min before a reaction was considered negative.

Determination of the feasibility of screening for PGNG from primary culture plates. A \( \beta \)-lac+ strain of S. aureus, E. coli, F. meningosepticum, or P. rettgeri and a \( \beta \)-lac- strain of N. gonorrhoeae were suspended in heart infusion broth at a concentration equivalent to a standard McFarland I. The gonococci were further diluted 1:10, so that when mixed together, the \( \beta \)-lac+ bacteria were present at 10 times the concentration of the \( \beta \)-lac- gonococci. These mixtures, as well as pure cultures of the test organisms, then were plated on MTM and chocolate agar, and the resultant growth was tested for \( \beta \)-lac reactivity.

RESULTS

Presumptive identification of \( N. \) gonorrhoeae at the Houston City Health Department Laboratory is made on the basis of colonial morphology and oxidase and Gram reactions. Since it is convenient to flood suspect colonies with oxidase reagent, we tested whether this reagent would interfere with the performance of the \( \beta \)-lac test (Table 1). A positive test was obtained with the \( \beta \)-lac+ \( N. \) gonorrhoeae both before and after exposure to oxidase reagent. Moreover, no false-positive results were obtained when \( \beta \)-lac-, oxidase-positive control cultures were tested.

<table>
<thead>
<tr>
<th>Strain of ( N. ) gonorrhoeae</th>
<th>Exposure to oxidase reagent</th>
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<tbody>
<tr>
<td>Before addition</td>
<td>After addition</td>
</tr>
<tr>
<td>( \beta )-Lac+</td>
<td>+</td>
</tr>
<tr>
<td>( \beta )-Lac-</td>
<td>-</td>
</tr>
</tbody>
</table>

\( \beta \)-Lac reactivity was defected up to 9 h after oxidase was added to a 24-h culture; up to 5.5 h after oxidase was added to a 48-h culture; and up to 2.5 h after oxidase was added to a 72-h culture.

We next determined whether a false-positive \( \beta \)-lac test would result if \( \beta \)-lac- gonococci were grown in the presence of a variety of \( \beta \)-lac+ bacteria. These organisms were mixed with the gonococcus as described in Materials and Methods. Either a loopful of bacteria was removed from the primary streak area (swipe) or an isolated colony (colony) was removed for testing (Table 2). In no case did exposure to oxidase reagent interfere with the performance of the test, although it did facilitate the identification of the gonococci, particularly in mixed cultures.

Whether plated on chocolate agar or MTM, the pure culture of \( \beta \)-lac- \( N. \) gonorrhoeae gave a negative \( \beta \)-lac test. The pure cultures of S. aureus, P. rettgeri, E. coli, and F. meningosepticum gave positive tests from chocolate agar. The S. aureus and E. coli did not grow on MTM, so the \( \beta \)-lac test could not be done. Only one colony of P. rettgeri grew on MTM; this colony was \( \beta \)-lac+. Growth of the F. meningosepticum was greatly reduced on MTM, but colonies removed from this plate were \( \beta \)-lac+.

As expected, when the mixtures of organisms were plated, the S. aureus, E. coli, F. meningosepticum, and P. rettgeri overgrew the \( N. \) gonorrhoeae on chocolate agar. When a sweep was made of these plates, the \( \beta \)-lac test was invariably positive. However, when care was taken to pick only colonies resembling \( N. \) gonorrhoeae, a negative \( \beta \)-lac test was obtained. Small, gray, entire, mucoid colonies characteristic of \( N. \) gonorrhoeae could be picked without trouble, but the identification of gonococcal colonies was made particularly easy by the addition of oxidase reagent. When the mixtures of S. aureus, E. coli, or P. rettgeri and \( N. \) gonorrhoeae were plated on MTM, only colonies resembling \( N. \) gonorrhoeae were seen, and all the tests for \( \beta \)-lac were negative. There was a slight change in the color of the indicator solution when a sweep was made of the MTM plate containing F. meningosepticum and \( N. \) gonorrhoeae. This was attributed to the small number of gram-negative rods (i.e., \( \beta \)-lac+ F. meningosepticum).
P. rettgeri

meningosepticum

+ E. coli +

S. aureus +

Of the was false-positive not meningosepticum (gonorrhoeae.

P-lac-

nies case the test ange-yellow but MTM indicates that bacterium lac+ bacteria.

However, we the picking lac+ and 8-lac- control cultures

2. N. b Sweep indicates that a d this well never turned frankly yellow; a Gram stain of the area on the plate from which the inoculum was removed showed a few gram-negative rods mixed with mostly gram-negative diplococci.

The experiment involving cocultivation of β-lac+ bacteria and β-lac- N. gonorrhoeae on MTM indicates that suppressed β-lac+ bacteria will not give false-positive results. Even in the case where the inoculum contained a few colonies of F. meningosepticum along with the N. gonorrhoeae, the indicator solution turned orange-yellow but never became yellow. Thus, the test would have been read as negative. However, we recommend that in instances where colonial morphology and Gram stain show mixed cultures, only well-isolated colonies be assayed for β-lac production. The addition of oxidase reagent to the plate simplifies the picking of colonies of gonococci for testing.

Further, it is important to include known β-lac+ and β-lac- control cultures with every run to ensure the proper function of the test.

The finding that β-lac activity can be assessed from primary culture plates will facilitate screening large populations for PPNG and quickly identifying patients in whom alternate therapy should be considered. During a 1-month period, for example, 1,641 primary cultures showing oxidase-positive colonies were screened at the Houston City Health Department Laboratory, using the microacidimetric method described here. A presumptive identification of N. gonorrhoeae was made in 1,619 of the cultures; the remaining 22 cultures were found to be gram-negative bacilli or coclobacilli. None of the gonococci tested displayed β-lac activity. These included isolates of N. gonorrhoeae from eight patients with positive cultures after routine penicillin administration, all of which exhibited zones of inhibition of ≥36 mm by the penicillin disk diffusion test. One of the nongonococcal cultures, subsequently identified as F. meningosepticum, gave a positive reaction.

We prefer the microacidimetric method for β-lac detection to the paper strip modificatio of the iodometric procedure (5), because the endpoint is more easily read by the laboratorian, a large number of specimens can be processed with greater speed, and a smaller inoculum is required. During routine use, we have found that it is possible to test as little as one oxidase-positive colony of gram-negative diplococci, although the time required for the test to become positive is increased with smaller inocula. However, more than one isolated colony should be tested before calling a culture negative, because enzyme-negative colonies can represent 1 to 30% of the population obtained from primary clinical specimens of patients infected with β-lac- N. gonorrhoeae (6).

We presently are using this procedure to test positive cultures of recently treated individuals who return for test of cure evaluation, and we plan to employ this protocol for routine screening of all N. gonorrhoeae isolates. In addition, we are using the test to examine the β-lac reactivity of other primary isolates, including Haemophilus influenzae and Nocardia sp.

ACKNOWLEDGMENTS

A. S. W. was supported by Public Health Service training grant ST01 AI00462 from the National Institute of Allergy and Infectious Diseases.

<table>
<thead>
<tr>
<th>N. gonorrhoeae plus β-lac- inoculum*</th>
<th>Chocolate agar</th>
<th>β-Lac reaction</th>
<th>MTM</th>
</tr>
</thead>
<tbody>
<tr>
<td>Without oxidase</td>
<td>With oxidase</td>
<td>Without oxidase</td>
<td>With oxidase</td>
</tr>
<tr>
<td>Sweep*</td>
<td>Colony*</td>
<td>Sweep</td>
<td>Colony</td>
</tr>
<tr>
<td>S. aureus</td>
<td>+</td>
<td>–</td>
<td>+</td>
</tr>
<tr>
<td>F. meningosepticum</td>
<td>+</td>
<td>–</td>
<td>+</td>
</tr>
<tr>
<td>E. coli</td>
<td>+</td>
<td>–</td>
<td>+</td>
</tr>
<tr>
<td>P. rettgeri</td>
<td>+</td>
<td>–</td>
<td>+</td>
</tr>
</tbody>
</table>

* The mixture was prepared so that the β-lac- N. gonorrhoeae was present at one-tenth the concentration of the β-lac+ inoculum.

† Sweep indicates that the inoculum for the test included a sample of all the growth on the plate.

‡ One typical, well-isolated colony of N. gonorrhoeae was tested.

§ This well never turned frankly yellow; a Gram stain of the area on the plate from which the inoculum was removed showed a few gram-negative rods mixed with mostly gram-negative diplococci.

DISCUSSION

The experiment involving cocultivation of β-lac+ bacteria and β-lac- N. gonorrhoeae on MTM indicates that suppressed β-lac+ bacteria will not give false-positive results. Even in the case where the inoculum contained a few colonies of F. meningosepticum along with the N. gonorrhoeae, the indicator solution turned orange-yellow but never became yellow. Thus, the test would have been read as negative. However, we recommend that in instances where colonial morphology and Gram stain show mixed cultures, only well-isolated colonies be assayed for β-lac production. The addition of oxidase reagent to the plate simplifies the picking of colonies of gonococci for testing.

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LITERATURE CITED


