

Impact of Alternative Growth Supplements on Antimicrobial Susceptibility Testing of *Neisseria gonorrhoeae*

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Purpose: The accurate detection of antibiotic susceptibility of *Neisseria gonorrhoeae* (*N. gonorrhoeae*) is of great importance for the treatment of patients with gonorrhea as well as to hinder the progress of drug resistance. To promote the application of gonococcal antibiotic susceptibility monitoring in primary hospitals and remote medical institutions, this study evaluated the effect of alternative growth supplements on the antibiotic susceptibility testing of *N. gonorrhoeae* isolates.

Methods: We divided the antimicrobial-containing media into three groups by adding different growth supplements (sterile defibrinated sheep blood, bovine hemoglobin, and Vitox). We tested the antimicrobial susceptibility of 80 *N. gonorrhoeae* isolates in different groups against eight antibiotics. Nonparametric signed-rank tests were utilized to compare the minimum inhibitory concentration (MIC) results of each group. Taking the MIC results of Vitox group as expected, the essential agreement (EA) and category agreement (CA) of the other two groups were calculated.

Results: For the group using sheep blood as growth supplements, the EA values and CA values of each antibiotic were above 90.00% and minor error rates were less than 7.00%. No very major error and major error were observed. For the group using hemoglobin as growth supplements, the EA values of the susceptibility results of zoliflodacin, penicillin, and ceftriaxone were lower than 90.00%. The overall MIC results of using hemoglobin as a growth supplement were higher than those of sheep blood and Vitox in the susceptibility testing of these three antibiotics.

Conclusion: Compared with the expected results, sheep blood may be considered for the use as an alternative material for *N. gonorrhoeae* antibiotics susceptibility surveillance, while hemoglobin may not be suitable for supplement to antimicrobial-containing medium.

Keywords: *Neisseria gonorrhoeae*, antibiotic susceptibility testing, growth supplements, agar dilution method

Introduction

Neisseria gonorrhoeae (*N. gonorrhoeae*) is the etiologic agent of gonorrhea, a sexually transmitted infection (STI) that remains a major global public health concern.¹ According to the WHO estimation, there were 87 million global incidents of gonorrhea annually.² In 2021, China reported 127,803 new cases of gonorrhea.³ Currently, the treatment of *N. gonorrhoeae* infection mainly depends on a variety of antibiotics.⁴ Notably, as the susceptibility of *N. gonorrhoeae* to various antibiotics decreases with time, multidrug-resistant (MDR) *N. gonorrhoeae* cases have been reported intermittently.⁵ Hence, the detection of antibiotic susceptibility of *N. gonorrhoeae* is of great importance for the accurate treatment of patients with gonorrhea as well as to hinder the progress of drug resistance.⁶

In 1990, the WHO Global Gonococcal Antimicrobial Surveillance Program (WHO GASP) was established to monitor gonococcal antimicrobial resistance (AMR) worldwide.⁷ China-GASP was initiated in 1987 by the National Center for Sexually Transmitted Disease Control (NCSTD).⁸ According to the laboratory diagnosis guidelines of WHO and EUCAST,^{9,10} the methods currently used for determining MICs include the agar dilution method and E-test method.

In the CLSI guidelines, disk diffusion or agar dilution MIC tests were used for routine clinical testing.¹¹ Disk diffusion only enabled qualitative testing of antimicrobial resistance.¹² Agar dilution is the most widely used method and the “gold standard” method for the determination of the MIC of gonococcal isolates against antimicrobial drugs.¹³

In the agar dilution method, the MIC determination is conducted in an antimicrobial-containing medium, which is a mix of GC agar base supplemented with a 1% defined growth supplement and antimicrobial solutions.^{8,11} From the data uploaded by various monitoring points in China, methodological differences among local laboratories and research institutions were summarized. In the standard operation of the agar dilution method, 1% Vitox was recommended as a growth supplement, while sterile defibrinated sheep blood and bovine hemoglobin were used as substitutes for Vitox in some hospitals and research institutions.^{14,15} For those institutions, the possible reasons for choosing sheep blood or hemoglobin as growth supplements were that these are more accessible and less costly. Hemoglobin was also used as a nutrient supplement in the Thayer Martin Agar used to isolate and culture *N. gonorrhoeae*.¹⁶

To assess possible systematic errors among the monitoring points and to find suitable growth supplement alternatives, we conducted this study to evaluate the accuracy for various experimental materials of antibiotic susceptibility testing of *N. gonorrhoeae*.

Methods

Bacterial Isolates

The *N. gonorrhoeae* isolates we used were 80 clinical gonococcal isolates from China Gonococcal Resistance Surveillance Program (China-GRSP). WHO *N. gonorrhoeae* reference isolates and ATCC 49226 were used for quality assurance. All the strains were previously determined to be *N. gonorrhoeae* through standardized methodologies.⁸ The Medical Ethics Committee at the Institute of Dermatology, the Chinese Academy of Medical Sciences & Peking Union Medical College and the National Center for Sexually Transmitted Disease Control all gave their approval to this project (2014-LS-026). This declaration of Helsinki complied in this study. Study participants signed an informed consent form before inclusion in the study. Before the antimicrobial agent susceptibility testing, all of the strains were preserved in skim milk at -80°C .

Resuscitation and Antimicrobial Susceptibility Testing

The effect of different growth supplements on the MIC of eight antibiotics (zofludacin, penicillin, gentamicin, tetracycline, spectinomycin, azithromycin, ciprofloxacin, and ceftriaxone) on *N. gonorrhoeae* isolates was determined. For this GC agar base (OXOID, USA) supplemented with 10% sterile defibrinated sheep blood (Lezhen, China) or 2% bovine hemoglobin (BD, USA) or 1% Vitox (OXOID, USA) were added separately to the media containing different concentrations of the antibiotics. Thereafter, 1 μL of bacterial suspension was transferred onto the agar surface using a replicator. Eventually, each medium was cultured for 18–24 hours at 36°C in a 5% CO_2 -enriched atmosphere. The growth of *N. gonorrhoeae* in different groups was observed and recorded. All steps strictly followed the WHO standard operation of the agar dilution method except for the other two additional growth supplements (sheep blood and hemoglobin).⁹ Each test was repeated for each bacteria strain to confirm the results.

Statistical Analysis

The MIC results of *N. gonorrhoeae* to the eight antibiotics were recorded. Graphpad Prism ver.6 Software (GraphPad Software, San Diego, CA, USA) was utilized to analyze all statistics. The phenotype characteristics of reference strains that were applied as the quality assessment of this experiment were gathered in the WHO global GASP.¹⁰ The percentage of MIC results of each antibiotic in different growth supplement groups was calculated. To assess the accuracy of each group, the MIC results of Vitox group were used to calculate the essential agreement (EA) and category agreement (CA). EA was defined as the percentage of strains with MIC results within plus or minus one doubling dilution step from the expected MIC result. CA was defined as the agreement of susceptible, intermediate and resistant results between MIC test and reference MIC result. The reproducibility was addressed with EA values and the performance of each group with CA values.¹¹ Very major errors were resistant strains being misclassified as susceptible. Major errors were susceptible strains

being misclassified as resistant. Minor errors occurred when one method categorized an isolate as intermediate while the other method defined it as susceptible or resistant.¹² Breakpoints from Clinical and Laboratory Standards Institute (CLSI) Antimicrobial Susceptibility Testing Guidelines¹³ were used to classify the *N. gonorrhoeae* isolates as sensitive, intermediately resistant, or resistant to antibiotics. The non-parametric Mann–Whitney *U*-test was implemented to examine the differences between MICs of eight antibiotics in different growth supplement.¹⁴

Results

A total of 80 *N. gonorrhoeae* isolates were tested for antibiotic susceptibility on antimicrobial-containing media supplemented with different growth supplements. During each batch of drug susceptibility testing, the MIC values of the reference strains were within 1 doubling dilution of those previously reported. The overall performances for eight antimicrobials are shown in Table 1. For the group using sheep blood as growth supplements, the EA values and CA values of each antibiotic were above 90.00% (EA values ranged from 90.00% to 100.00%, CA values ranged from 93.75% to 100.00%). No very major error and major error was observed, and minor error rates were less than 7.00% for the sheep blood group (minor error rates ranged from 0.00% to 6.25%). The EA values and CA values of gentamicin and spectinomycin were 100.0%. For the group using hemoglobin as growth supplements, the essential agreement values of the susceptibility results of zoliflodacin, penicillin, and ceftriaxone were 38.75%, 81.25%, and 60.00%, all lower than 90.00%. A major error was observed in the susceptibility results of ciprofloxacin.

Comparing the results of the three drugs (zoliflodacin, penicillin, and ceftriaxone) with large differences between the two supplements in Table 2, the overall MIC results of using hemoglobin as a growth supplement were higher than those of sheep blood and Vitox in the drug susceptibility testing of zoliflodacin, penicillin, and ceftriaxone ($P \leq 0.05$). For example, in the susceptibility results of zoliflodacin, the percentage of ≥ 0.5 mg/L in the hemoglobin group was 42.50%,

Table 1 Categorical and Essential Agreements of Two Different Growth Supplements for 8 Antibiotics

	Spectinomycin		Azithromycin		Ciprofloxacin		Ceftriaxone	
	Sheep blood	Hemo-globin	Sheep blood	Hemo-globin	Sheep blood	Hemo-globin	Sheep blood	Hemo-globin
Number tested	80	80	80	80	80	80	80	80
Essential agreement	80	80	76	72	78	79	72	48
Essential agreement rate	100.00%	100.00%	95.00%	90.00%	97.50%	98.75%	90.00%	60.00%
Categorical agreement	80	80	76	78	80	79	79	73
Categorical agreement rate	100.00%	100.00%	95.00%	97.50%	100.00%	98.75%	98.75%	91.25%
Very major errors	0	0	0	0	0	0	0	0
Major errors	0	0	0	0	0	1	0	0
Minor errors	0	0	4	2	0	0	1	7
Minor error rate	0.00%	0.00%	5.00%	2.50%	0.00%	0.00%	1.25%	8.75%
	Zoliflodacin		Gentamicin		Penicillin		Tetracycline	
	Sheep blood	Hemo-globin	Sheep blood	Hemo-globin	Sheep blood	Hemo-globin	Sheep blood	Hemo-globin
Number tested	80	80	80	80	80	80	80	80
Essential agreement	72	31	80	80	75	65	78	73
Essential agreement rate	90.00%	38.75%	100%	100%	93.75%	81.25%	97.50%	91.25%
Categorical agreement	76	47	80	80	75	66	78	77
Categorical agreement rate	95.00%	58.75%	100%	100%	93.75%	82.50%	97.50%	96.25%
Very major errors	0	0	0	0	0	0	0	0
Major errors	0	0	0	0	0	0	0	0
Minor errors	4	33	0	0	5	14	2	3
Minor error rate	5.00%	41.25%	0.00%	0.00%	6.25%	17.50%	2.50%	3.75%

Table 2 The Percentage (%) of *N. Gonorrhoeae* Isolates with Different MICs (Mg/L) for Eight Antibiotics in Different Groups

MIC (mg/L)	SPE (%)			GEN (%)			PEN (%)			TET (%)		
	V	S	H	V	S	H	V	S	H	V	S	H
0.25									2.50			
0.5							5.00	5.00	1.30			
1							11.30	15.00	12.50	7.50	5.00	3.80
2							23.80	22.50	11.30	26.30	15.00	7.50
4							21.30	22.50	21.30	61.30	75.00	51.30
8				56.30	68.80	26.30	13.80	10.00	23.80	0.00	0.00	32.50
16	20.00	13.80	5.00	43.80	31.30	73.80	0.00	0.00	2.50	0.00	0.00	0.00
32	75.00	81.30	90.00				25.00	25.00	25.00	5.00	5.00	5.00
≥64	5.00	5.00	5.00									

MIC (mg/L)	ZOLI (%)			AZM (%)			CIP (%)			CRO (%)		
	V	S	H	V	S	H	V	S	H	V	S	H
≤0.008										12.50	8.80	7.50
0.015										18.80	18.80	0.00
0.03	7.50						20.00	20.00	18.80	0.00	10.00	3.80
0.06	21.30	5.00					0.00	0.00	0.00	8.80	0.00	27.50
0.125	65.00	48.80	6.30	2.50			5.00	5.00	5.00	18.80	12.50	16.30
0.25	6.30	41.30	51.30	3.80	1.30	18.80	0.00	0.00	0.00	26.30	27.50	2.50
0.5		5.00	40.00	57.50	61.30	56.30	0.00	0.00	0.00	10.00	12.50	28.80
1			2.50	13.80	15.00	5.00	0.00	0.00	0.00	5.00	10.00	13.80
2				2.50	2.50	0.00	5.00	2.50	5.00			
4				0.00	0.00	7.50	6.30	2.50	0.00			
8				20.00	20.00	12.50	0.00	7.50	5.00			
16							63.80	62.50	66.30			

Abbreviations: ZOLI, zoliflodacin; CRO, ceftriaxone; PEN, penicillin; GEN, gentamicin; TET, tetracycline; SPE, spectinomycin; AZM, azithromycin; CIP, ciprofloxacin; CRO, ceftriaxone; V, Vitox; S, sheep blood; H, hemoglobin.

while those in the Vitox and sheep blood group were 0.00% and 5.00%. An isolate with an MIC value of 1mg/L was observed in the results of the hemoglobin group. In the susceptibility results of ceftriaxone, the percentage of ≥0.5 mg/L in the hemoglobin group was 42.60%, while those in the Vitox and sheep blood group were 15.00% and 22.50%. Similar results were found for the results of penicillin (Figure 1).

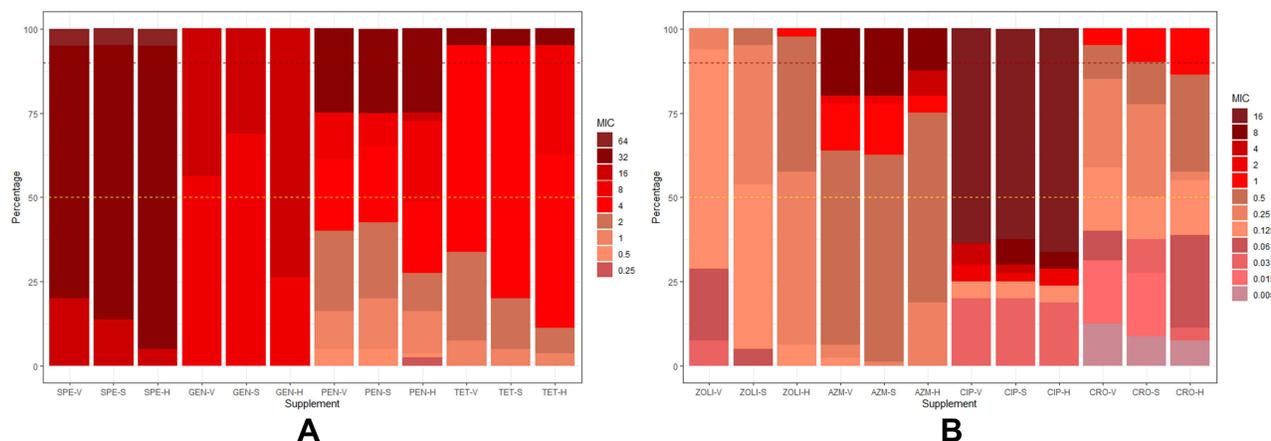


Figure 1 (A and B) The proportions of *N. gonorrhoeae* isolates with different MICs for eight antibiotics in different groups. The MIC50 (yellow-dotted line), MIC90 (red-dotted line) and number of isolates per province are shown. The block color represents the MIC value and the block length represents the percentage. The vertical coordinate represents the antibiotics of different growth supplement groups.
Abbreviations: ZOLI, zoliflodacin; CRO, ceftriaxone; PEN, penicillin; GEN, gentamicin; TET, tetracycline; SPE, spectinomycin; AZM, azithromycin; CIP, ciprofloxacin; CRO, ceftriaxone; V, Vitox; S, sheep blood; H, hemoglobin.

Discussion

As a result of the significant decline in the efficacy of available antimicrobials, *N. gonorrhoeae* has been identified as an emerging public health problem. Consequently, it is necessary to detect the trend of *N. gonorrhoeae* antimicrobial resistance (AMR) for controlling the gonorrhea epidemic.¹⁵ In China, the *N. gonorrhoeae* resistance surveillance program has been carried out since 1987. By comparing the susceptibility testing methods at various participating clinics in China, we found that there were differences among STD laboratories. Previous studies have proven that the results of the agar dilution method can be affected by many factors: the composition of the agar medium, pH-value, and culture parameters.¹⁶ Thus, despite the MICs estimated by different laboratories being comparable, random errors and systematic errors in the values due to technical nuances can also affect the clinical interpretation. Therefore, exploring the effect and the accuracy of different experimental materials on the MIC testing of *N. gonorrhoeae* isolates is of great meaning for the accurate detection of *N. gonorrhoeae* AMR.

In this study, we aimed to compare the effect of different growth supplements in the antimicrobial-containing media to the MIC results of *N. gonorrhoeae* and to find suitable growth supplement alternatives. The results demonstrate that sheep blood as an alternative nutrient showed good categorical and essential agreements in multiple antibiotics' susceptibility testing results, and no very major error and major error was observed in the results of the sheep blood group. While in the hemoglobin group, there were three groups of antibiotics susceptibility results less than accepted percentages (defined as >90% for categorical agreements and <7% for minor errors).¹² After comparing the results of the three antibiotics with large differences between two supplements, we also found that the MIC results of the hemoglobin group were higher than those of the sheep blood group and the Vitox group. The results suggested that sheep blood as an alternative growth supplement showed good reproducibility in the susceptibility testing of *N. gonorrhoeae*, while hemoglobin may not be suitable for a supplement to antibiotics-containing media.

Such a discrepancy between hemoglobin and Vitox in these three antibiotics could arise because of two reasons. The first is the characteristics of the drugs. The main component of Vitox is vitamin B12, adenine, L-glutamine, guanine, aminobenzoic acid, cystine, cysteine, etc., while the main component of sheep blood is hemoglobin, serum protein, vitamin B, lysine, glutamic acid, folic acid, etc.¹⁷ In hemoglobin supplements, it is the only substance, which is mainly composed of aspartic acid, leucine and glutamic acid. Therefore, when sheep blood and Vitox are added, some of the components may interact with the drug. Another reason is the impact of resistance mechanisms. According to the guidelines of CLSI,¹⁸ the use of a cysteine-free growth supplement is required for agar dilution tests with carbapenems and clavulanate. The main reason is that cysteine is related to the drug resistance of these two kinds of antibiotics.¹⁹ Therefore, we suspect that there is a similar situation in this experiment.

During our retrieval of relevant literature, this is the first study comparing the accuracy of alternative growth supplements like sheep blood and hemoglobin with the standard material in the antibiotic susceptibility testing in *N. gonorrhoeae*. The results of this study provided a database for the use of sheep blood as an alternative growth supplement. Considering that sheep blood is less expensive and more easily available than Vitox, for some primary or remote institutions, the use of sheep blood may be suitable for use as an experimental material for *N. gonorrhoeae* antibiotics susceptibility surveillance. Under strict quality assurance, including the use of reference strains as quality control in each experiment and the MIC results of the reference strains within 1 doubling dilution of the panel, sheep blood could be used as an alternative growth supplement to Vitox.

In this study, we provided the first exploratory results. Nevertheless, there were also some limitations in our study. First, the sample size of this study is limited, a large panel of clinical isolates and more antibiotic-resistant isolates are needed to verify the trend we have found. Second, the function of specific components of each growth supplement is out of the scope of this study. We did not analyze the possible reasons for the results differences among the three growth supplements in terms of the composition and background. A possible cause of the higher results in hemoglobin medium may be the lower concentration of antibiotics.^{20,21} According to the CLSI M100 Performance Standards, cysteine-containing defined growth supplement will significantly alter dilution test results with carbapenems and clavulanate. Hence, the role of the specific components in different growth supplements by the control variable method will be our next focus.

Conclusion

In conclusion, this study highlights the effects of different growth supplements on antibiotic susceptibility testing of *N. gonorrhoeae*. Sheep blood may be considered for use as an alternative material for Vitox in the *N. gonorrhoeae* antibiotics susceptibility surveillance, while hemoglobin may not be suitable for a supplement to antimicrobial-containing medium.

Data Sharing Statement

The data that supports the findings of this study are available in the submitted article.

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Author Contributions

All authors made a significant contribution to the work reported, whether that is in the conception, study design, execution, acquisition of data, analysis and interpretation, or in all these areas; took part in drafting, revising or critically reviewing the article; gave final approval of the version to be published; have agreed on the journal to which the article has been submitted; and agree to be accountable for all aspects of the work.

Disclosure

The authors declare that there is no conflict of interest regarding the publication of this article.

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