



Emerging Azithromycin Resistance among the *Neisseria gonorrhoeae* Strains Isolated in Danyang, China

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ABSTRACT

Neisseria gonorrhoeae is a human pathogen. Strain identification and effective antibiotic treatment are the main ways to prevent and control gonorrhea, but *N. gonorrhoeae* is now resistant to most antibiotics, and there is no vaccine available. The main purpose of this study was to understand the molecular characteristics of clinical isolates of *N. gonorrhoeae* and their resistance to azithromycin (AZM) in Danyang, China. Firstly, the clinical isolates of *N. gonorrhoeae* from January 2016 to December 2020 were collected by Matrix-Assisted Laser Desorption Ionization-Time of Flight Mass Spectrometry (MALDI-TOF MS). Secondly, the drug sensitivity of all clinical isolates was analyzed. Thirdly, polymerase chain reaction (PCR) and DNA sequencing were used to analyze the genes related to AZM resistance (AZM-R), including 23s rRNA allele mutation, *mtrR* promoter and coding region mutation, *rplD* and *rplV* mutation. Finally, the clinical isolates resistant to AZM were typed by *N. gonorrhoeae* multiantigen sequence typing (NG-MAST). 388 clinical isolates of *N. gonorrhoeae* were identified, of which 373, 329, 298, 5, 11 and 5 strains were resistant to ciprofloxacin, penicillin, tetracycline, AZM, spectinomycin and ceftriaxone, respectively. The mutation detection of AZM-R related gene showed that there were single (A) nucleotide deletion mutation in *mtrR* promoter region, G45D mutation in *mtrR* coding region, G70 mutation in *rplD* and A2047G mutation in 23s rRNA allele, but no mutation was found in *rplV*. A total of 8 different STs were identified in 5 AZM-R strains, of which two ST1866 isolates showed a high level of AZM-R. The clinical isolates of *N. gonorrhoeae* in Danyang have high genetic diversity. ST1866 isolates showed a high level of AZM-R. Therefore, measures should be taken to monitor the spread of ST1866 *N. gonorrhoeae* clones in eastern China.

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Authors' Contribution

JY and WZ prepared the study design, performed statistical analyses, and drafted the manuscript. QZ performed bactericidal assays and experiments. LC advised continuously in experiments and projects. ZC performed experiments with phagocytosis assay. XZ aided in study design and advising in experiments. WW, SZ and LW assisted in experiment and design of phagocytosis assay.

Key words

Neisseria gonorrhoeae, Azithromycin, Antimicrobial resistance, NG-MAST, Gonorrhoea

INTRODUCTION

Neisseria gonorrhoeae is a human-specific pathogen, causing 78 million new gonorrhea infections worldwide every year (Zhu *et al.*, 2016; Ma *et al.*, 2017; Lovett and Duncan, 2018; Quillin and Seifert, 2018). Gonorrhea can lead to serious complications, such as epididymitis in men and pelvic inflammation in women. Pelvic inflammation can result in involuntary infertility and ectopic pregnancy (Mitchell and Prabhu, 2013;

Wiesenfeld and Manhart, 2017). *N. gonorrhoeae* can also infect the eyes of newborns as they pass through the birth canal of an infected mother, which can lead to blindness (Rivacoba *et al.*, 2017; Jin, 2019). More importantly, gonorrhea is associated with other sexually transmitted infections and human immunodeficiency virus (HIV) infections (Xu *et al.*, 2018; Sanyal *et al.*, 2019; Dave *et al.*, 2020; Kato *et al.*, 2020). Gonorrhea is considered to be a non-ulcerative sexually transmitted infection, just like chlamydia and trichomoniasis, and like other non-ulcerative sexually transmitted diseases, people with gonorrhea have a higher risk of transmitting HIV to their partners (Peters *et al.*, 2021; Pottorff *et al.*, 2021). This is because the loss of genitals in patients with HIV infection increases the virus.

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Identification of strains and effective antibiotic treatment are the main ways to prevent and control gonorrhoea, but *N. gonorrhoeae* are now resistant to most antibiotics and no vaccine is available (Abbasi, 2017; Baarda *et al.*, 2018; Vincent and Jerse, 2019). Now, the double antimicrobial therapy of ceftriaxone and azithromycin (AZM) has been widely accepted worldwide, as an empirical first-line treatment for gonorrhoea (Maldonado and Takhar, 2013; Unemo *et al.*, 2021), including China (Chen *et al.*, 2014). In most countries, 2 g AZM single therapy is used for treatment of pairs β -Gonorrhoea in patients with lactam allergy. The results show that *N. gonorrhoeae* will produce resistance after introducing a new antimicrobial agent and replace the sensitive bacterial population within 20 years (Unemo and Dillon, 2011). Therefore, it is necessary to strengthen the monitoring of *N. gonorrhoeae* resistance.

In the 1990s, AZM became the first choice drug for many infectious diseases, but it was reported that AZM had drug resistance in the past decade (Steingrimsson *et al.*, 1990). AZM was recommended to Chinese patients with mixed infection of gonorrhoea and Chlamydia trachomatis around 2000 (Duan *et al.*, 2019), and was widely used because of its wide availability and easy management. However, the widespread use of AZM may lead to resistance of *N. gonorrhoeae*. AZM resistance (AZM-R) *N. gonorrhoeae* was first found in China from 2001 to 2003, and AZM-R isolates were first identified in Guangzhou in 2009 (Liang *et al.*, 2016). In the following years, AZM-R isolates and multi drug resistant isolates were reported in Nanjing, Hangzhou and Changsha (Ni *et al.*, 2016; Wan *et al.*, 2018; Yan *et al.*, 2019; Yuan *et al.*, 2019). Until 2013, little was known about the types of AZM-R *N. gonorrhoeae* prevalent in China. Therefore, the level of AZM-R and the molecular characteristics of AZM-R *N. gonorrhoeae* are still unclear (Jiang *et al.*, 2017).

The multidrug resistance of *N. gonorrhoeae* is related to the overexpression of efflux pump. The most important efflux mechanism is the MtrC-MtrD-MtrE system, which is encoded by the *mtr* operon, in which *mtrR* is the regulatory gene and *mtrCDE* is the structural gene (Lucas *et al.*, 1997). Another efflux pump encoded by *mef* gene was first found in some gram-positive bacteria and then in clinical strains of gonorrhoea (Luna *et al.*, 2000).

In the *mtrR* gene, specific mutations in the promoter or coding region can lead to decrease the MtrCDE efflux pump repression and subsequently increase export of the antimicrobial. Mutations in the promoter or coding sequence of *mtrR* gene of macrolide resistant *N. gonorrhoeae* can reduce the expression of *mtrR* repressor and up regulate mtrCDE efflux pump (Chen *et al.*, 2019). In *N.*

gonorrhoeae, there was a single base pair (A) deletion in the 13 bp reverse repeat of *mtrR* promoter region, the expression of *mtrR* was cancelled, and the level of mtrCDE increased, most likely because the binding affinity of RNA polymerase to mtrCDE increased (Handing *et al.*, 2018). Missense mutations in the *mtrR*, such as G45D mutation in the helix trans helix motif in the *mtrR* repressor, can reduce the binding of the repressor to the mtrCDE promoter. The increased expression of mtrCDE efflux pump also increased AZM.

Further resistance to AZM is the result of 23S rRNA loop V mutation, which is a specific target of AZM. AZM exerts its bacteriostatic effect by directly interacting with the central ring of *rrl* gene domain V encoding 23S rRNA, resulting in obstruction of protein synthesis (Pham *et al.*, 2021). Specific point mutations in this region may lead to drug resistance by reducing the affinity of AZM to its target (Zhang and van der Veen, 2019). 23S rRNA point mutations have been described, including C2611T (numbering refers to the *E. coli* genome), given low to medium levels of AZM-R (minimum inhibitory concentration MIC = 2 to 32 μ g/mL) or A2059G (*E. coli* numbering), awarded high level AZM-R (MICs \geq 256 μ g/mL) (Demczuk *et al.*, 2016).

Ribosomal proteins L4 (encoded by *rplD*) and L22 (encoded by *rplV*) bind to domain I of 23S rRNA and act as channels for macrolide antibiotics to enter ribosomes (Reinert and Al-Lahham, 2005; Chisholm *et al.*, 2010). Point mutations of *rplD* and *rplV* in *E. coli* and *Streptococcus pneumoniae* lead to resistance to macrolides; However, this mutation is rarely detected in patients with gonorrhoea.

In the absence of new antibiotics for the treatment of gonorrhoea, it is important to classify the emergence and dynamics of AZM-R *N. gonorrhoeae* on a regional and national basis for the successful updating of treatment recommendations. The resistance level of *N. gonorrhoeae* to azithromycin in Danyang area of eastern China is still unclear. The purpose of this study was to investigate the prevalence and molecular typing of *N. gonorrhoeae* in Danyang City.

MATERIALS AND METHODS

N. gonorrhoeae isolation and species verification

The patients with *N. gonorrhoeae* infection were identified by consulting the microbiological laboratory records of Danyang people's Hospital of Jiangsu Province from January 2016 to December 2020. Clinical isolates are anonymous, so this study does not require ethical approval. There is no standard for selecting strains. The clinical data, such as gender and age and infection site, were

recorded. Species verification is performed biochemically using Matrix-Assisted Laser Desorption Ionization-Time of Flight Mass Spectrometry (MALDI-TOF MS) using VITEK-MS (bioMérieux).

Antimicrobial susceptibility testing

The susceptibility of *N. gonorrhoeae* to penicillin, tetracycline, ciprofloxacin, spectinomycin, ceftriaxone and AZM was determined by E-test. The results of the E-test were interpreted according to the guidelines for all antibiotics of the Institute of clinical and laboratory standards (Palmeira and Ferreira, 2019). All the strains were stored in glycerine broth and stored at -80°C until use. The quality control strain of drug sensitivity test was *N. gonorrhoeae* ATCC 49226, which was provided by the clinical laboratory of the Ministry of Health and kept in our laboratory. The strain was stored in liquid medium of glycerine broth.

Genetic determinants associated with resistance to AZM

Genomic DNA, was extracted using DNA rapid extraction kit (Shanghai Shenggong) and stored at -20°C. To identify site-specific mutations, we sequenced the genes associated with AZM-R, including 23s rRNA allele, *rplD* and *rplV* (encoding ribosomal proteins L4 and L22, respectively), as well as *mtrR* promoter and coding region. The primers and conditions for PCR have been previously published (Allen *et al.*, 2011; Zheng *et al.*, 2019). PCR products were bidirectionally sequenced by Applied Biosystems 3730xl DNA automatic sequencer. DNA sequences were compared with BLAST and GenBank programs (<http://www.ncbi.nlm.nih.gov/blast/>) was used to identify gene mutations.

N. gonorrhoeae multiantigen sequence typing (NG-MAST)

NG-MAST was used to analyze the molecular epidemiology of AZM-R *N. gonorrhoeae* isolates. Briefly, the internal regions of genes encoding two variable outer membrane proteins, *porB* and *tbpB* subunit B, were sequenced to generate a two allele map of a strain. Sequence type (ST) is assigned through NG-MAST website (www.ng-mast.net). Besides, the NG-MAST genome based on the sequence similarity of *porB* and *tbpB* alleles is defined as described above (Martin *et al.*, 2004). The phylogenetic tree of AZM-R *N. gonorrhoeae* was established by linking *porB* and *tbpB* alleles using MEGA 7.0 software and maximum likelihood method.

RESULT

Patient data

There were 388 strains of *N. gonorrhoeae*, including

75 strains in 2016, 78 strains in 2017, 77 strains in 2018, 78 strains in 2019, 80 strains in 2020. There were 340 strains in male and 48 strains in female. There were 41 strains in obstetrics and gynecology, 62 strains in dermatology and 285 strains in urology, and the age range was 16-78 years old, including 16 strains under 20 years old, 107 strains aged 20-30 years old, 103 strains aged 30-40 years old, 60 strains aged 40-50 years old, 59 strains aged 50-60 years old and 43 strains over 60 years old.

Antimicrobial susceptibility

Of the 388 strains of *N. gonorrhoeae*, 373 strains (96.13%), 329 strains (84.80%), 298 strains (26.80%), 5 strains (1.29%), 11 strains (2.83%) and 5 strains (1.29%) were resistant, respectively to ciprofloxacin, penicillin, tetracycline, AZM, spectinomycin and ceftriaxone. Among them, 88 strains were β -lactamase positive, with a positive rate of 22.68%. The MIC of AZM ranges from 0.064 to > 16 $\mu\text{g/mL}$, MIC₅₀ = 0.5 $\mu\text{g/mL}$, MIC₉₀ = 1.0 $\mu\text{g/mL}$. Among the 5 AZM-R strains, 2 strains with MIC = 1 $\mu\text{g/mL}$ showed low level resistance, 2 strains with MIC = 4 $\mu\text{g/mL}$ showed moderate resistance, and 1 strain with MIC > 16 $\mu\text{g/mL}$ showed high level resistance.

Detection of mutations in genes associated with AZM-R

The following mutations were detected in 5 AZM-R isolates: single nucleotide (A) deletion in the promoter region of *mtrR*, G70D in the *mtrR* coding region, and A2047G in the 23s rRNA allele (*N. gonorrhoeae* number, GenBank accession number: X67293.1) (Table I). Four allele mutations in 23s rRNA were detected in a gonorrhea isolate with high level of AZM-R. No mutation was detected in *rplV*.

Five azithromycin sensitive (AZM-S) strains were randomly selected as control group. The mutations detected in these controls included single nucleotide (A) deletion in the *mtrR* promoter region and G45D in the *mtrR* coding region in the three isolates (Table I). No mutation was found in *rplD* and *rplV* genes and 23s rRNA alleles in AZM-S group.

Molecular epidemiologic typing

Five AZM-R *N. gonorrhoeae* isolates were genotyped by NG-MAST, and 8 different sequence types of (STs) were identified. ST1866 were the most common isolates of ST, followed by ST4007, ST1407, ST12746, ST3287, ST1731, ST2286, ST2318 and ST12660. Both ST1866 isolates (which were not associated with each other in epidemiology) displayed high levels of AZM-R. All the 8 STs found in this study have been reported in NG-MAST database.

Based on phylogenetic analysis, there are four

groups. Group A strains include two different STs (ST2286

Table I. Characterization and molecular typing results of clinical isolates of *N. gonorrhoeae*.

Group	Strain no. (Years)	AZM MIC ($\mu\text{g/ml}$)	<i>mtrR</i> ^a promoter	<i>mtrR</i> ^b coding region	<i>rplD</i> ^c mutation	<i>rplV</i> ^d mutation	23S rRNA mutation ^e at position 2047	NG-MAST
AZM-R (n=5)	70(2016)	1	A deletion	No mutation	No mutation	No mutation	No mutation	1407
	13(2017)	1	A deletion	G45D	No mutation	No mutation	A→G, alleles 1, and 2	12746
	09(2018)	4	A deletion	G45D	No mutation	No mutation	A→G, alleles 1,2 and 3	1866
	23(2019)	4	A deletion	No mutation	No mutation	No mutation	A→G, alleles 1, and 4	3287
	45(2020)	18	A deletion	G45D	G70D	No mutation	A→G, alleles 1,2,3 and 4	1866
AZM-S (n=5)	04(2016)	0.25	No mutation	No mutation	No mutation	No mutation	No mutation	1731
	25(2017)	0.125	No mutation	No mutation	No mutation	No mutation	No mutation	2286
	55(2018)	0.064	A deletion	No mutation	No mutation	No mutation	No mutation	4007
	63(2019)	0.25	No mutation	G45D	No mutation	No mutation	No mutation	2318
	17(2020)	0.64	No mutation	No mutation	No mutation	No mutation	No mutation	12660

Notes: ^a The *mtrR* promoter mutation is the single-base-pair (A) deletion in the 13-bp inverted repeat in the promoter region. ^b GenBank accession number: Z25797.1. ^c GenBank accession number: YP-208871.1. ^d GenBank accession number: YP-208867.1. ^e GenBank accession number: X67293.1.

and ST2318). Group B strains include two different STs (ST12746 and ST1866). Group C strains are composed of three different STs (ST4007, ST1407, ST1731 and ST3287). Finally, the AZM-S cluster D contains a different STs (ST12660).

DISCUSSION

In this study, the drug sensitivity test of *N. gonorrhoeae* in Danyang area from 2016 to 2020 was combined with AZM-R molecular typing. This study found for the first time that AZM-R began to appear in Danyang area, which was consistent with the previous reports from China (Yuan *et al.*, 2011; Chen *et al.*, 2013; Yuan *et al.*, 2019). World Health Organization (WHO) suggested that once 5 per cent of locally acquired *N. gonorrhoeae* isolates develop drug resistance, the empirical use of antibiotics should be stopped. Therefore, AZM is not recommended as a monotherapy for gonococcal urethritis or cervicitis in China and many other countries in the world. To improve treatment effectiveness and delay further selection of cephalosporin-resistant *N. gonorrhoeae*, most current guidelines recommend a dual treatment regimen of ceftriaxone (250mg or 500mg intramuscular injection) or cefixime (400mg, if no ceftriaxone option) combined with AZM (1g or 2g oral) in the treatment of gonorrhoea. However, in recent years, the decreased sensitivity of AZM-R *N. gonorrhoeae* to extended-spectrum cephalosporins has been reported, which seriously threatens the future efficacy of current treatment recommendations.

In recent years, there have been studies on AZM-R *N.*

gonorrhoeae in China. For example, 32% of the isolates showed AZM-R, and 10% showed high level of drug resistance (Zheng *et al.*, 2019). The analysis of 126 strains isolated from Hefei from 2014 to 2015 showed that 29% of the strains were AZM-R, and 10% of them showed high level of AZM-R.

Mutations in the promoter or coding sequence of *mtrR* gene in macrolide resistant *N. gonorrhoeae* strains can reduce the expression of *mtrR* repressor, resulting in the up regulation of *mtrCDE* efflux pump (Ohneck *et al.*, 2015). In *N. gonorrhoeae*, there was A-deletion in the 13 bp reverse repeat of *mtrR* promoter, which overlaps the *mtrCDE* promoter at the -35 region, *mtrR* expression was cancelled, and *mtrCDE* expression was increased, which was probably because RNA polymerase had greater binding affinity for *mtrCDE* (Johnson and Shafer, 2015). *N. gonorrhoeae* resistant to macrolides can reduce the expression of MtrR repressor and up-regulate MtrCDE efflux pump (Ng *et al.*, 2002; Cousin *et al.*, 2003; Fernandez-Huerta and Espasa, 2019; Hall *et al.*, 2019; Ma *et al.*, 2020). In *N. gonorrhoeae* strains with deletion of 13-bp reverse repeat sequence in MtrR promoter region, the expression of MtrR was cancelled and the level of MtrCDE increased, which was probably due to the increase of binding affinity of RNA polymerase to *mtrCDE* (Rouquette *et al.*, 1999). Missense mutations in the *mtrR* (Handing *et al.*, 2018; Beggs *et al.*, 2021), such as the G45D mutation in the helix-to-helix motif of the *mtrR* repressor, can reduce the binding of the repressor to the *mtrCDE* promoter. In this study, the mutation rates of *mtrR* promoter and coding region of AZM-R strain were

100.0% and 60%, respectively. The mutation rate of *mtrR* promoter region in AZM-R group was significantly higher than that in AZM-S group, but there was no significant difference in *mtrR* coding region. This is consistent with the previously reported results that the mutation in the *mtrR* promoter region plays a more important role in the resistance of *N. gonorrhoeae* to azithromycin than the mutation in the *mtrR* coding region.

AZM exerts its bacteriostatic effect by directly interacting with the central ring of RRL gene domain V encoding 23S rRNA, resulting in obstruction of protein synthesis (Wu *et al.*, 2011; Trembizki *et al.*, 2015). Specific point mutations in this region may lead to drug resistance by reducing the affinity of azithromycin to its target (Ng *et al.*, 2002; Chisholm *et al.*, 2010; Galarza *et al.*, 2010). The HL-AZM-R isolate had A2143G mutation in at least three of the four alleles. Including A2143G (the number in *E. coli* corresponds to A2059) or C2599T (the number in *E. coli* corresponds to C2611T) (Ng *et al.*, 2002). Previous studies have reported that one or more of the four alleles of *rplD* gene in 23s rRNA domain V are associated with AZM-R, including mutation, A2059G and C2611T (corresponding to A2059G and C2611T in *E. coli*, respectively) (Belkacem *et al.*, 2016). In this study, A2047G mutation was detected in the V region of AZM-R gonorrhea virus, but not in AZM-S group. The 23s rRNA alleles of A2047G mutation were alleles 1 and 2, alleles 1 and 2 and 3, alleles 1 and 4, alleles 1 and 2 and 3 and 4, respectively, and the corresponding MIC values were 1 µg/mL, 4 µg/mL, 4 µg/mL and 16 µg/mL, respectively. It is worth noting that A2047G mutation was detected in all strains with MIC > 16 µg/mL. Therefore, we believe that A2047G allele is the main determinant of AZM-R.

The *rplD* mutation (G70D) of a *N. gonorrhoeae* isolate to AZM was 16 µg/mL, which was consistent with the previously reported results (Jacobsson *et al.*, 2016). G68D and G70D were previously described by (Zheng *et al.*, 2019), so the mutation at position 68-70 of the *rplD* gene of *N. gonorrhoeae* seems to be associated with a high level of AZM resistance. The mutant of *rplD* gene belongs to group B, and the NG-MAST classification of ST1866, indicates that the mutant has genetic diversity.

Gonorrhea typing methods, such as NG-MAST, are helpful to understand the spread of gonorrhea. In this study, phylogenetic analysis showed that AZM-R strains had wide differences and did not belong to any specific group. Different from the reports of other countries, the isolates with high level of AZM-R in this study belong to NG-MAST-ST1866. It is worth mentioning that the high-level *N. gonorrhoeae* AZM-R found in this study has the same NA-MAST type as the high-level *N. gonorrhoeae* AZM-R found in the Liu-YH report, which indicates that

the transmission of NG-MAST ST1866 *N. gonorrhoeae* has occurred across the Taiwan Strait. In addition, NG-MAST ST1866 clones have been reported in Nanjing, Hangzhou and Hefei (Ni *et al.*, 2016; Jiang *et al.*, 2017; Wan *et al.*, 2018), which indicates that the clone has a high level of AZM-R, has spread in eastern China, which is a matter of concern, which must be paid attention to.

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Ethics approval

Because the *N. gonorrhoeae* were part of the routine hospital laboratory procedure, ethics approval was not required.

Statement of conflict of interest

The authors have declared no conflicts of interest.

REFERENCES

- Abbasi, J., 2017. New hope for a gonorrhea vaccine. *J. Am. med. Assoc.*, **318**: 894-895. <https://doi.org/10.1001/jama.2017.11037>
- Allen, V.G., Farrell, D.J., Rebbapragada, A., Tan, J., Tijet, N., Perusini, S.J., Towns, L., Lo, S., Low D.E. and Melano, R.G., 2011. Molecular analysis of antimicrobial resistance mechanisms in neisseria gonorrhoeae isolates from ontario, canada. *Antimicrob. Agents Chemother.*, **55**: 703-712. <https://doi.org/10.1128/AAC.00788-10>
- Baarda, B.I., Martinez F.G. and Sikora, A.E., 2018. Proteomics, bioinformatics and structure-function antigen mining for gonorrhea vaccines. *Front. Immunol.*, **9**: 2793. <https://doi.org/10.3389/fimmu.2018.02793>
- Beggs, G.A., Ayala, J.C., Kavanaugh, L.G., Read, T.D., Hooks, G.M., Schumacher, M.A., Shafer, W.M. and Brennan, R.G., 2021. Structures of neisseria gonorrhoeae mtrr-operator complexes reveal molecular mechanisms of DNA recognition and antibiotic resistance-conferring clinical mutations. *Nucl. Acids Res.*, **49**: 4155-4170. <https://doi.org/10.1093/nar/gkab213>
- Belkacem, A., Jacquier, H., Goubard, A., Mougari, F., La Ruche, G., Patey, O., Micaelo, M., Semaille, C., Cambau, E. and Bercot, B., 2016. Molecular epidemiology and mechanisms of resistance of

- azithromycin-resistant neisseria gonorrhoeae isolated in france during 2013-14. *J. Antimicrob. Chemother.*, **71**: 2471-2478. <https://doi.org/10.1093/jac/dkw182>
- Chen, S., Connolly, K.L., Rouquette-Loughlin, C., D'Andrea, A., Jerse, A.E. and Shafer, W.M., 2019. Could dampening expression of the neisseria gonorrhoeae mtrcde-encoded efflux pump be a strategy to preserve currently or resurrect formerly used antibiotics to treat gonorrhea? *mBio*, **10**. <https://doi.org/10.1128/mBio.01576-19>
- Chen, S.C., Yin, Y.P., Dai, X.Q., Unemo, M. and Chen, X.S., 2014. Antimicrobial resistance, genetic resistance determinants for ceftriaxone and molecular epidemiology of neisseria gonorrhoeae isolates in nanjing, china. *J. Antimicrob. Chemother.*, **69**: 2959-2965. <https://doi.org/10.1093/jac/dku245>
- Chen, X.S., Yin, Y.P., Wei, W.H., Wang, H.C., Peng, R.R., Zheng, H.P., Zhang, J.P., Zhu, B.Y., Liu Q.Z. and Huang, S.J., 2013. High prevalence of azithromycin resistance to treponema pallidum in geographically different areas in china. *Clin. Microbiol. Infect. Off. Publ. Eur. Soc. Clin. Microbiol. Infect. Dis.*, **19**: 975-979. <https://doi.org/10.1111/1469-0691.12098>
- Chisholm, S.A., Dave, J. and Ison, C.A., 2010. High-level azithromycin resistance occurs in neisseria gonorrhoeae as a result of a single point mutation in the 23s rna genes. *Antimicrob. Agents Chemother.*, **54**: 3812-3816. <https://doi.org/10.1128/AAC.00309-10>
- Cousin, S.L., Jr., Whittington, W.L. and Roberts, M.C., 2003. Acquired macrolide resistance genes and the 1 bp deletion in the mtrr promoter in neisseria gonorrhoeae. *J. Antimicrob. Chemother.*, **51**: 131-133. <https://doi.org/10.1093/jac/dkg040>
- Dave, J., Paul, J., Pasvol, T.J., Williams, A., Warburton, F., Cole, K., Miari, V.F., Stabler, R. and Eyre, D.W., 2020. Ethnically diverse urban transmission networks of neisseria gonorrhoeae without evidence of hiv serosorting. *Sex. Transmitt. Infect.*, **96**: 106-109. <https://doi.org/10.1136/sextrans-2019-054025>
- Demczuk, W., Martin, I., Peterson, S., Bharat, A., Van Domselaar, G., Graham, M., Lefebvre, B., Allen, V., Hoang, L., Tyrrell, G., Horsman, G., Wylie, J., Haldane, D., Archibald, C., Wong, T., Unemo, M. and Mulvey, M.R., 2016. Genomic epidemiology and molecular resistance mechanisms of azithromycin-resistant neisseria gonorrhoeae in canada from 1997 to 2014. *J. clin. Microbiol.*, **54**: 1304-1313. <https://doi.org/10.1128/JCM.03195-15>
- Duan, X., Wang, K., Wu, J., Zhang, D., Liu, X., Ni, M., Liu, S. and Meng, Z., 2019. Comparative efficacy of chinese herbal injections combined with azithromycin for mycoplasma pneumonia in children: A bayesian network meta-analysis of randomized controlled trials. *J. clin. Pharm. Therapeut.*, **44**: 675-684. <https://doi.org/10.1111/jcpt.12855>
- Fernandez-Huerta, M. and Espasa, M., 2019. Mycoplasma genitalium co-infection with chlamydia trachomatis and neisseria gonorrhoeae among asymptomatic patients: The silent wick for macrolide resistance spread. *Sex. Transmitt. Infect.*, **95**: 391. <https://doi.org/10.1136/sextrans-2018-053848>
- Galarza, P.G., Abad, R., Canigia, L.F., Buscemi, L., Pagano, I., Oviedo, C. and Vazquez, J.A., 2010. New mutation in 23s rna gene associated with high level of azithromycin resistance in neisseria gonorrhoeae. *Antimicrob. Agents Chemother.*, **54**: 1652-1653. <https://doi.org/10.1128/AAC.01506-09>
- Hall, C.L., Harrison, M.A., Pond, M.J., Chow, C., Harding-Esch, E.M. and Sadiq, S.T., 2019. Genotypic determinants of fluoroquinolone and macrolide resistance in neisseria gonorrhoeae. *Sex. Hlth.*, **16**: 479-487. <https://doi.org/10.1071/SH18225>
- Handing, J.W., Ragland, S.A., Bharathan, U.V. and Criss, A.K., 2018. The mtrcde efflux pump contributes to survival of neisseria gonorrhoeae from human neutrophils and their antimicrobial components. *Front. Microbiol.*, **9**: 2688. <https://doi.org/10.3389/fmicb.2018.02688>
- Jacobsson, S., Golparian, D., Cole, M., Spiteri, G., Martin, I., Bergheim, T., Borrego, M.J., Crowley, B., Crucitti, T., Van Dam, A.P., Hoffmann, S., Jeverica, S., Kohl, P., Mlynarczyk-Bonikowska, B., Pakarna, G., Stary, A., Stefanelli, P., Pavlik, P., Tzelepi, E., Abad, R., Harris, S.R. and Unemo, M., 2016. Wgs analysis and molecular resistance mechanisms of azithromycin-resistant (mic >2 mg/l) neisseria gonorrhoeae isolates in europe from 2009 to 2014. *J. Antimicrob. Chemother.*, **71**: 3109-3116. <https://doi.org/10.1093/jac/dkw279>
- Jiang, F.X., Lan, Q., Le, W.J. and Su, X.H., 2017. Antimicrobial susceptibility of neisseria gonorrhoeae isolates from hefei (2014-2015): Genetic characteristics of antimicrobial resistance. *BMC Infect. Dis.*, **17**: 366. <https://doi.org/10.1186/s12879-017-2472-z>
- Jin, J., 2019. Prevention of gonococcal eye infection in newborns. *J. Am. med. Assoc.*, **321**: 414. <https://doi.org/10.1001/jama.2018.21434>
- Johnson, P.J. and Shafer, W.M., 2015. The transcriptional repressor, mtrr, of the mtrcde efflux pump operon

- of neisseria gonorrhoeae can also serve as an activator of “off target” gene (glne) expression. *Antibiotics*, **4**: 188-197. <https://doi.org/10.3390/antibiotics4020188>
- Kato, Y., Kawaguchi, S., Shigehara, K., Yaegashi, H., Nakashima, K., Nakagawa, T., Sakamoto, J., Itoda, I., Ueda, M., Izumi, K., Kadono, Y. and Mizokami, A., 2020. Prevalence of n. Gonorrhoeae, c. Trachomatis, m. Genitalium, m. Hominis and ureaplasma spp. In the anus and urine among japanese hiv-infected men who have sex with men. *J. Infect. Chemother. Off. J. Japan Soc. Chemother.*, **26**: 403-406. <https://doi.org/10.1016/j.jiac.2019.12.007>
- Liang, J.Y., Cao, W.L., Li, X.D., Bi, C., Yang, R.D., Liang, Y.H., Li, P., Ye, X.D., Chen, X.X. and Zhang, X.B., 2016. Azithromycin-resistant neisseria gonorrhoeae isolates in guangzhou, china (2009-2013): Coevolution with decreased susceptibilities to ceftriaxone and genetic characteristics. *BMC Infect. Dis.*, **16**: 152. <https://doi.org/10.1186/s12879-016-1469-3>
- Lovett, A. and Duncan, J.A., 2018. Human immune responses and the natural history of neisseria gonorrhoeae infection. *Front. Immunol.*, **9**: 3187. <https://doi.org/10.3389/fimmu.2018.03187>
- Lucas, C.E., Balthazar, J.T., Hagman, K.E. and Shafer, W.M., 1997. The mtrr repressor binds the DNA sequence between the mtrr and mtrc genes of neisseria gonorrhoeae. *J. Bact.*, **179**: 4123-4128. <https://doi.org/10.1128/jb.179.13.4123-4128.1997>
- Luna, V.A., Cousin, S., Jr., Whittington, W.L. and Roberts, M.C., 2000. Identification of the conjugative mef gene in clinical acinetobacter junii and neisseria gonorrhoeae isolates. *Antimicrob. Agents Chemother.*, **44**: 2503-2506. <https://doi.org/10.1128/AAC.44.9.2503-2506.2000>
- Ma, K.C., Mortimer, T.D., Duckett, M.A., Hicks, A.L., Wheeler, N.E., Sanchez-Buso, L. and Grad, Y.H., 2020. Increased power from conditional bacterial genome-wide association identifies macrolide resistance mutations in neisseria gonorrhoeae. *Nat. Commun.*, **11**: 5374. <https://doi.org/10.1038/s41467-020-19250-6>
- Ma, Q., Xu, X., Luo, M., Wang, J., Yang, C., Hu, X., Liang, B., Wu, F., Yang, X., Wang, J., Liu, H., Li, W., Zhong, Y., Li, P., Xie, J., Jia, L., Wang, L., Hao, R., Du, X., Qiu, S., Song, H. and Sun, Y., 2017. A waterborne outbreak of shigella sonnei with resistance to azithromycin and third-generation cephalosporins in china in 2015. *Antimicrob. Agents Chemother.*, **61**. <https://doi.org/10.1128/AAC.00308-17>
- Maldonado, N.G. and Takhar, S.S., 2013. Update on emerging infections: News from the centers for disease control and prevention. Update to the cdc’s sexually transmitted diseases treatment guidelines, 2010: Oral cephalosporins no longer a recommended treatment for gonococcal infections. *Annls Emerg. Med.*, **61**: 91-95. <https://doi.org/10.1016/j.annemergmed.2012.10.014>
- Martin, I.M., Ison, C.A., Aanensen, D.M., Fenton, K.A. and Spratt, B.G., 2004. Rapid sequence-based identification of gonococcal transmission clusters in a large metropolitan area. *J. Infect. Dis.*, **189**: 1497-1505. <https://doi.org/10.1086/383047>
- Mitchell, C. and Prabhu, M., 2013. Pelvic inflammatory disease: Current concepts in pathogenesis, diagnosis and treatment. *Infect. Dis. Clin. North Am.*, **27**: 793-809. <https://doi.org/10.1016/j.idc.2013.08.004>
- Ng, L.K., Martin, I., Liu, G. and Bryden, L., 2002. Mutation in 23s rRNA associated with macrolide resistance in neisseria gonorrhoeae. *Antimicrob. Agents Chemother.*, **46**: 3020-3025. <https://doi.org/10.1128/AAC.46.9.3020-3025.2002>
- Ni, C., Xue, J., Zhang, C., Zhou, H. and van der Veen, S., 2016. High prevalence of neisseria gonorrhoeae with high-level resistance to azithromycin in Hangzhou, China. *J. Antimicrob. Chemother.*, **71**: 2355-2357. <https://doi.org/10.1093/jac/dkw131>
- Ohneck, E.A., Goytia, M., Rouquette-Loughlin, C.E., Joseph, S.J., Read, T.D., Jerse, A.E. and Shafer, W.M., 2015. Overproduction of the mtrcde efflux pump in neisseria gonorrhoeae produces unexpected changes in cellular transcription patterns. *Antimicrob. Agents Chemother.*, **59**: 724-726. <https://doi.org/10.1128/AAC.04148-14>
- Palmeira, J.D. and Ferreira, H., 2019. CLSI - EUCAST: Comparison of antibiotic-susceptibility profile of Enterobacteriaceae of animal origin according to the standards. *Acta Microbiol. Immunol. Hung.*, **66**: 413-422. <https://doi.org/10.1556/030.66.2019.037>
- Peters, R.P., Feucht, U.D., de Vos, L., Ngwepe, P., McIntyre, J.A., Klausner, J.D. and Medina-Marino, A., 2021. Mother-to-child transmission of chlamydia trachomatis, neisseria gonorrhoeae, and trichomonas vaginalis in hiv-infected pregnant women in south africa. *Int. J. STD AIDS.*, 956462421990218. <https://doi.org/10.1177/0956462421990218>
- Pham, C.D., Nash, E., Liu, H., Schmerer, M.W., Sharpe, S., Woods, G., Roland, B., Schlanger, K., St Cyr, S.B., Carlson, J., Sellers, K., Olsen,

- A., Sanon, R., Hardin, H., Soge, O.O., Raphael, B.H. and Kersh, E.N., 2021. Atypical mutation in neisseria gonorrhoeae 23s rna associated with high-level azithromycin resistance. *Antimicrob. Agents Chemother.*, **65**. <https://doi.org/10.1128/AAC.00885-20>
- Pottorff, A., Duarte, P., Chow, J., Luque, A. and Nijhawan, A.E., 2021. Extragenital testing for neisseria gonorrhoeae and chlamydia trachomatis in a large hiv clinic in the us south: Implementation and epidemiology. *Sex. Transmitt. Dis.*, **48**: e22-e26. <https://doi.org/10.1097/OLQ.0000000000001349>
- Quillin, S.J. and Seifert, H.S., 2018. Neisseria gonorrhoeae host adaptation and pathogenesis. *Nat. Rev. Microbiol.*, **16**: 226-240. <https://doi.org/10.1038/nrmicro.2017.169>
- Reinert, R.R. and Al-Lahham, A., 2005. Time-kill study of the activity of telithromycin against macrolide-resistant streptococcus pneumoniae isolates with 23s rna mutations and changes in ribosomal proteins l4 and l22. *Antimicrob. Agents Chemother.*, **49**: 3011-3013. <https://doi.org/10.1128/AAC.49.7.3011-3013.2005>
- Rivacoba, M.C., Izquierdo, G., Zenteno, N. and Porte, L., 2017. Neisseria subflava bacteremia in newborns: Case report and review of the literature. *Rev. Chile. Infect. Org. Off. Soci. Chilen. Infect.*, **34**: 389-392. <https://doi.org/10.4067/s0716-10182017000400389>
- Rouquette, C., Harmon, J.B. and Shafer, W.M., 1999. Induction of the mtrcde-encoded efflux pump system of neisseria gonorrhoeae requires mtra, an arac-like protein. *Mol. Microbiol.*, **33**: 651-658. <https://doi.org/10.1046/j.1365-2958.1999.01517.x>
- Sanyal, A., Shen, C., Ding, M., Reinhart, T.A., Chen, Y., Sankapal, S. and Gupta, P., 2019. Neisseria gonorrhoeae uses cellular proteins excl10 and il8 to enhance hiv-1 transmission across cervical mucosa. *Am. J. Reprod. Immunol.*, **81**: e13111. <https://doi.org/10.1111/aji.13111>
- Steingrimsson, O., Olafsson, J.H., Thorarinsson, H., Ryan, R.W., Johnson, R.B. and Tilton, R.C., 1990. Azithromycin in the treatment of sexually transmitted disease. *J. Antimicrob. Chemother.*, **25 (Suppl A)**: 109-114. https://doi.org/10.1093/jac/25.suppl_A.109
- Trembizki, E., Buckley, C., Donovan, B., Chen, M., Guy, R., Kaldor, J., Lahra, M.M., Regan, D.G., Smith, H., Ward, J. and Whiley, D.M., 2015. Direct real-time pcr-based detection of neisseria gonorrhoeae 23s rna mutations associated with azithromycin resistance. *J. Antimicrob. Chemother.*, **70**: 3244-3249. <https://doi.org/10.1093/jac/dkv274>
- Unemo, M. and Dillon, J.A., 2011. Review and international recommendation of methods for typing neisseria gonorrhoeae isolates and their implications for improved knowledge of gonococcal epidemiology, treatment, and biology. *Clin. Microbiol. Rev.*, **24**: 447-458. <https://doi.org/10.1128/CMR.00040-10>
- Unemo, M., Ross, J., Serwin, A.B., Gomberg, M., Cusini, M. and Jensen, J.S., 2021. Background review for the 2020 european guideline for the diagnosis and treatment of gonorrhoea in adults. *Int. J. STD AIDS*, **32**: 108-126. <https://doi.org/10.1177/0956462420948739>
- Vincent, L.R. and Jerse, A.E., 2019. Biological feasibility and importance of a gonorrhea vaccine for global public health. *Vaccine*, **37**: 7419-7426. <https://doi.org/10.1016/j.vaccine.2018.02.081>
- Wan, C., Li, Y., Le, W.J., Liu, Y.R., Li, S., Wang, B.X., Rice, P.A. and Su, X.H., 2018. Increasing resistance to azithromycin in neisseria gonorrhoeae in eastern chinese cities: Resistance mechanisms and genetic diversity among isolates from nanjing. *Antimicrob. Agents Chemother.*, **62**. <https://doi.org/10.1128/AAC.02499-17>
- Wiesenfeld, H.C. and Manhart, L.E., 2017. Mycoplasma genitalium in women: Current knowledge and research priorities for this recently emerged pathogen. *J. Infect. Dis.*, **216**: S389-S395. <https://doi.org/10.1093/infdis/jix198>
- Wu, A., Buono, S., Katz, K.A. and Pandori, M.W., 2011. Clinical neisseria gonorrhoeae isolates in the united states with resistance to azithromycin possess mutations in all 23s rna alleles and the mtrr coding region. *Microbial. Drug Resist.*, **17**: 425-427. <https://doi.org/10.1089/mdr.2010.0199>
- Xu, S.X., Leontyev, D., Kaul, R. and Gray-Owen, S.D., 2018. Neisseria gonorrhoeae co-infection exacerbates vaginal hiv shedding without affecting systemic viral loads in human cd34+ engrafted mice. *PLoS One*, **13**: e0191672. <https://doi.org/10.1371/journal.pone.0191672>
- Yan, J., Xue, J., Chen, Y., Chen, S., Wang, Q., Zhang, C., Wu, S., Lv, H., Yu, Y. and van der Veen, S., 2019. Increasing prevalence of neisseria gonorrhoeae with decreased susceptibility to ceftriaxone and resistance to azithromycin in hangzhou, china (2015-17). *J. Antimicrob. Chemother.*, **74**: 29-37. <https://doi.org/10.1093/jac/dky412>
- Yuan, L.F., Yin, Y.P., Dai, X.Q., Pearline, R.V., Xiang, Z., Unemo, M. and Chen, X.S., 2011. Resistance to azithromycin of neisseria gonorrhoeae isolates from

- 2 cities in china. *Sex. Transmitt. Dis.*, **38**: 764-768. <https://doi.org/10.1097/OLQ.0b013e318219cdb5>
- Yuan, Q., Li, Y., Xiu, L., Zhang, C., Fu, Y., Jiang, C., Tang, L. and Peng, J., 2019. Identification of multidrug-resistant neisseria gonorrhoeae isolates with combined resistance to both ceftriaxone and azithromycin, china, 2017-2018. *Emerg. Microbes Infect.*, **8**: 1546-1549. <https://doi.org/10.1080/22221751.2019.1681242>
- Zhang, J. and van der Veen, S., 2019. Neisseria gonorrhoeae 23s rRNA a2059g mutation is the only determinant necessary for high-level azithromycin resistance and improves *in vivo* biological fitness. *J. Antimicrob. Chemother.*, **74**: 407-415. <https://doi.org/10.1093/jac/dky438>
- Zheng, Z., Liu, L., Shen, X., Yu, J., Chen, L., Zhan, L., Chen, H., Lin, C., Jiang, Y., Xia, H., Wang, L. and Yu, F., 2019. Antimicrobial resistance and molecular characteristics among neisseria gonorrhoeae clinical isolates in a chinese tertiary hospital. *Infect. Drug Resist.*, **12**: 3301-3309. <https://doi.org/10.2147/IDR.S221109>
- Zhu, B., Bu, J., Li, W., Zhang, J., Huang, G., Cao, J., Tang, Z., Gan, Q. and Wei, P., 2016. High resistance to azithromycin in clinical samples from patients with sexually transmitted diseases in guangxi zhuang autonomous region, china. *PLoS One*, **11**: e0159787. <https://doi.org/10.1371/journal.pone.0159787>

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