

HHS Public Access

Author manuscript *Sex Transm Dis.* Author manuscript; available in PMC 2019 April 01.

Published in final edited form as:

Sex Transm Dis. 2018 April; 45(4): e18. doi:10.1097/OLQ.000000000000758.

Ciprofloxacin May be Efficacious in Treating Wild-Type Gyrase A Genotype *Neisseria gonorrhoeae* Infections

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Dear Editor

We read with great interest the article by Chesson et al. which illustrated the possible economic losses that may be averted by slowing the emergence of ceftriaxone resistant *Neisseria gonorrhoeae* infections [1]. As the threat of untreatable *Neisseria gonorrhoeae* infection increases [2], rapid molecular assays for the timely prediction of antimicrobial susceptibility are increasingly being utilized to target antimicrobial therapy [3]. A recent modeling study demonstrated the theoretical delay in the emergence of multidrug-resistant *Neisseria gonorrhoeae* with the use of such rapid molecular assays [4]. The University of California, Los Angeles Health System has implemented a rapid molecular assay for the determination of mutation in the gyrase A (*gyrA*) gene of *Neisseria gonorrhoeae* [5, 6]. The absence of mutation in that gene has been shown to reliably predict susceptibility to ciprofloxacin [7]. As suggested by the article by Chesson et al. the use of ciprofloxacin may spare ceftriaxone, and thus reduce the emergence of ceftriaxone resistant *Neisseria gonorrhoeae* [8].

Ciprofloxacin has been shown to be >99% effective among infections with phenotypic susceptibility [9], however one concern regarding the current utility of ciprofloxacin in the treatment of *Neisseria gonorrhoeae* infections is that there have been no studies evaluating patient outcomes among those with wild-type (non-mutated) *gyr*A genotype *Neisseria gonorrhoeae* infection. We have collected outcome data from participants with wild-type

The authors have no conflict of interest.

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Neisseria gonorrhoeae infections treated with ciprofloxacin at the University of California, Los Angeles Health System. We completed a retrospective review of 25 patient records between June 2016 and September 2017 with a repeat *Neisseria gonorrhoeae* test performed between 7-90 days after single dose ciprofloxacin 500 mg oral treatment of the initial infection. Of those 25 participants, 100% (95% CI 83% – 100%) had a negative test of cure. The anatomic sites of infection included rectal (n=7), pharyngeal (n=7), urethral (n=7), and unspecified genital swabs (n=4). There was no difference in repeat test results by time to repeat test 7–21 days (100%) vs. 22–90 days (100%) (*p*-value>0.1).

Those preliminary results provide promising evidence of the efficacy of ciprofloxacin for the treatment of wild-type *gyr*A genotype *Neisseria gonorrhoeae* infections. There are additional costs to *gyr*A genotyping and genotype-based therapy compared with recommended empiric two-drug therapy with ceftriaxone and azithromycin [10]; however, in the light of the potential costs incurred by the emergence of ceftriaxone resistant infections as suggested by Chesson et al. we advocate for further implementation of the *gyr*A genotypic assay in other health systems for the promotion of targeted *Neisseria gonorrhoeae* therapy.

Acknowledgments

This research was supported by the National Institutes of Health, grants: R21AI77256 and R21AI109005.

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