

Does pristinamycin or sitafloxacin do the trick?

Treatment outcomes in people with macrolide-resistant mycoplasma genitalium who have failed first-line therapy

Anderson I¹, Varma R^{1,2}

1. Sydney Sexual Health Centre, South-Eastern Sydney Local Health District 2. The Kirby Institute, Sexual Health Program, University of New South Wales

Background

Treatment of Mycoplasma genitalium (Mgen) is complicated by resistance to macrolides and fluoroquinolones.

In the absence of fluoroquinolone resistance assays, a significant minority of clients fail first-line sequential treatment with doxycycline and moxifloxacin. Guidelines recommend sequential doxycycline and pristinamycin or sitafloxacin as second line. Elsewhere, pristinamycin-based regimens have achieved 75% cure rate (95% CI 66%–82%)¹.

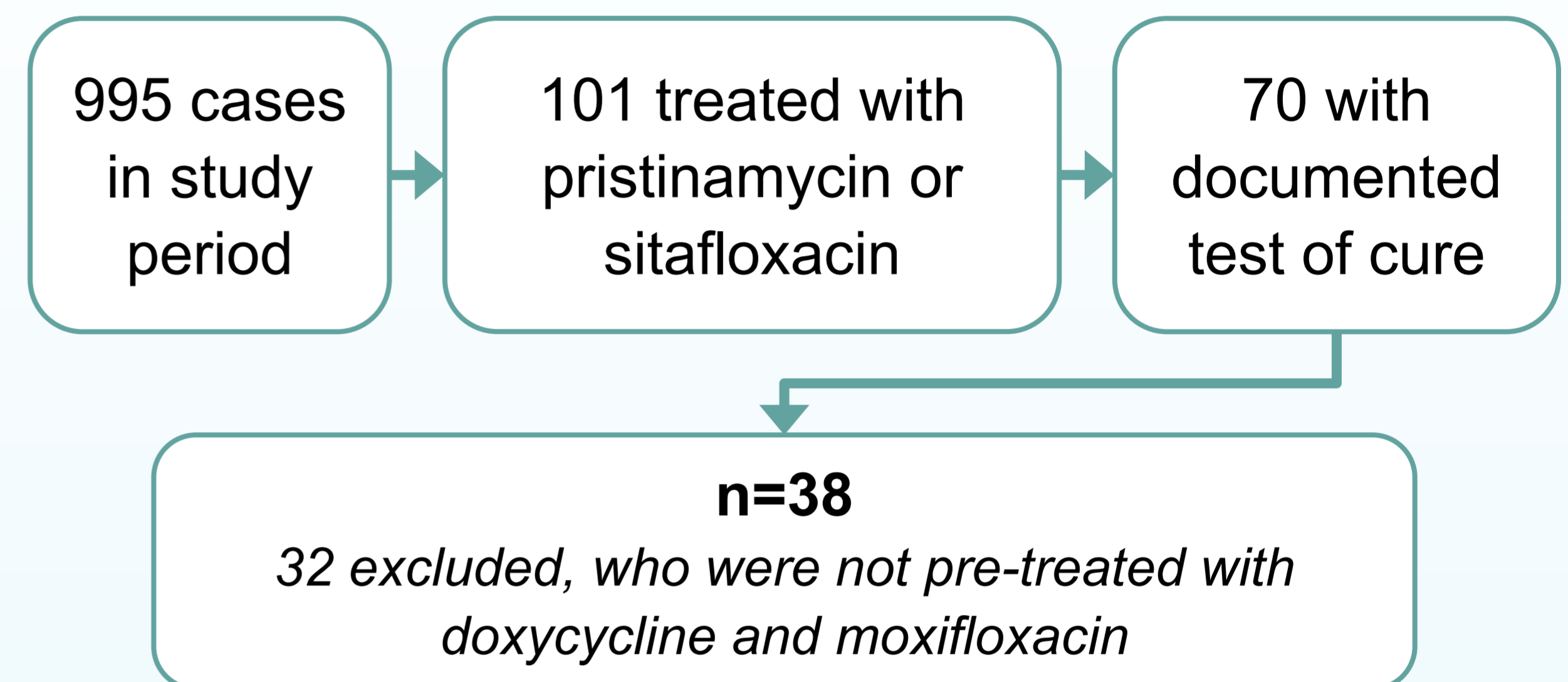
We looked at treatment outcomes of second-line therapy of Mgen in a sexual health clinic in Sydney, which receives community referrals and where testing was indicated for clients with STI syndromes and sexual contacts.

Methods

- Retrospective case series of pristinamycin and sitafloxacin use for macrolide-resistant Mgen
- Study period: May 2017 – May 2021
- Analysed using descriptive statistics and Fisher's exact test with $p \leq 0.05$

Inclusion Criteria

- diagnosis with macrolide-resistant Mgen infection
- treatment with pristinamycin or sitafloxacin at SSHC
- documented test of cure
- initial treatment with doxycycline and moxifloxacin

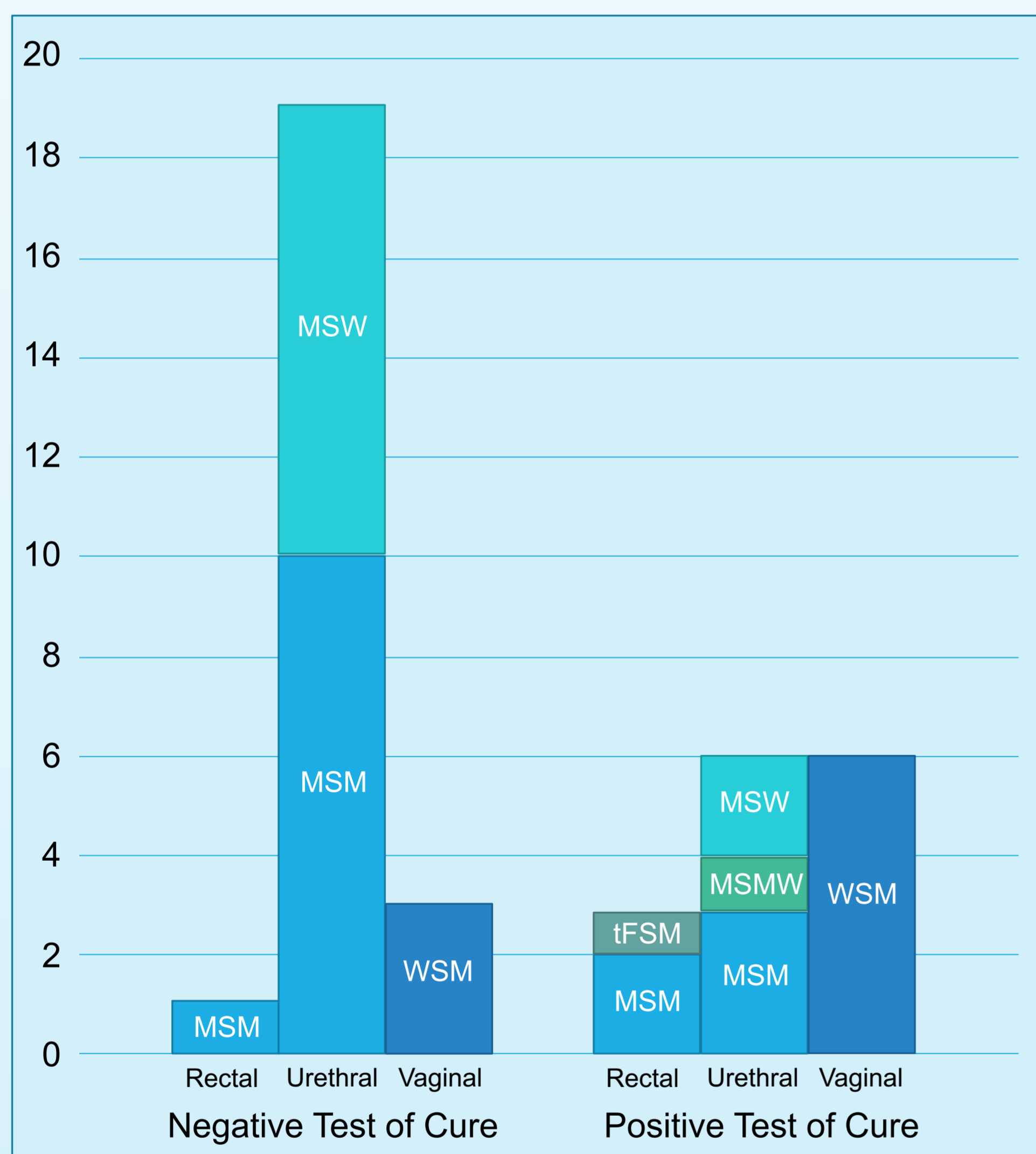


Why is looking at Mgen treatment important?

- Mgen is the culprit in NGU in up to 35% of cases^{2,3}
- Australian MSM have more macrolide-resistance than in the general population (76% vs 39%)⁴
- 16% of Australian Mgen have dual-class resistance (macrolide [azithromycin] + fluoroquinolone [moxifloxacin])⁵

Results

Test of cure by site and sexual behaviour



	n=38		Test of cure result (% of n)			
	n	%	Negative		Positive	
Total	38	100%	23	61%	15	39%
Treatment Drug						
Pristinamycin	31	82%	20	65%	11	35%
Sitafloxacin	7	18%	3	43%	4	57%
Gender						
Female	9	24%	3	33%	6	67%
Trans Female	1	3%	0	0%	1	100%
Male	28	74%	20	71%	8	29%
Site (Symptoms at TOC)						
Rectal	4	11%	1	25%	3	75%
No	3	8%	1	33%	2	67%
Yes	1	3%	0	0%	1	100%
Urethral	25	66%	19	76%	6	24%
No	20	53%	16	80%	4	20%
Unknown	1	3%	0	0%	1	100%
Yes	4	11%	3	75%	1	25%
Vaginal	9	24%	3	33%	6	67%
No	4	11%	3	75%	1	25%
Yes	5	13%	0	0%	5	100%

Significance	
	p-value
Pristinamycin vs sitafloxacin	0.401
Cis male vs Non-cis male	0.030
Urethral vs non-urethral	0.013
MSM vs non-MSM	0.506
Symptomatic vs non-symptomatic at TOC	0.023

Conclusion

Gender, site of infection, and symptoms could guide clinicians when counselling clients on therapeutic effectiveness when they present for second-line M. gen treatment. Larger, prospective multi-centre studies that specifically examine treatment differences between urethritis and vaginitis/cervicitis are required.

Key Messages

- No difference between pristinamycin and sitafloxacin as second-line drug
- Cis males with urethral infection cured more readily
- Sexual behaviour does not matter, in a cohort of people with macrolide-resistant Mgen
- Symptoms at TOC are likely to indicate ongoing infection

References

1. Durukan D, Doyle M, Murray G, Bodiababu K, Vodstrcil L, Chow EPF, et al. Doxycycline and Sitafloxacin Combination Therapy for Treating Highly Resistant Mycoplasma genitalium. Emerg Infect Dis. 2020 Aug;26(8):1870–4. 2. Gnanadurai R, Fifer H 2020. Mycoplasma genitalium: A Review. Microbiology. 166(1):21–9. 3. McIver R, Jalocon D, McNulty A, Jeffreys NJ, Bourne C, Varma R, et al. Men who have sex with men with Mycoplasma genitalium-positive non-gonococcal urethritis are more likely to have macrolide resistant strains than men with only female partners. SSHC Manuscript. 2018. 4. Read TRH, Murray GL, Danielewski JA, Fairley CK, Doyle M, Worthington K, et al. Symptoms, Sites, and Significance of Mycoplasma genitalium in Men Who Have Sex with Men. Emerg Infect Dis. 2019 Apr;25(4):719–27. 5. Sethi S, Zaman K, Jain N. Mycoplasma genitalium infections: current treatment options and resistance issues. Infection and Drug Resistance. 2017;10:283–92.