

# CLINICAL FEATURES OF *MYCOPLASMA GENITALIUM* ASSOCIATED PELVIC INFLAMMATORY DISEASE AND RESPONSE TO MOXIFLOXACIN: A CASE SERIES

## Authors:

Latimer RL<sup>1</sup>, Read TRH<sup>1-2</sup>, Vodstrcil LA<sup>1-2</sup>, Goller JL<sup>3</sup>, Fairley CK<sup>1-2</sup>, Ong JJ<sup>1,2,4</sup>, Hocking JS<sup>3</sup>, Bradshaw CS<sup>1-2</sup>

<sup>1</sup>Central Clinical School, Monash University, Melbourne, Australia

<sup>2</sup>Melbourne Sexual Health Centre, Alfred Health, Melbourne, Australia

<sup>3</sup>Centre for Epidemiology and Biostatistics, Melbourne School of Population & Global Health, University of Melbourne, Parkville, Victoria, 3010, Australia

<sup>4</sup>London School of Hygiene and Tropical Medicine, London, United Kingdom

## Introduction:

There are limited published data describing the clinical features and therapeutic response in women meeting criteria for presumptive treatment for pelvic inflammatory disease associated with *Mycoplasma genitalium* (henceforth MG-PID). MG-PID has been reported to respond poorly to standard PID treatment regimens and while moxifloxacin is recommended in several treatment guidelines, published data to support its use are scant.

## Methods:

We conducted a retrospective study of women at Melbourne Sexual Health Centre (MSHC) between 2006-2017, who met the CDC criteria for presumptive treatment of PID, and had MG detected as the sole pathogen. Clinical and laboratory characteristics of MG-PID were compared to cases of chlamydial PID (CT-PID) by multivariable analysis. Microbiological and clinical cure following moxifloxacin and standard PID treatment was determined for women with MG-PID who returned for test of cure between 14-120 days.

## Results:

Ninety-two patients with MG-PID were compared with 92 women with CT-PID. MG-PID was associated with increased lower abdominal tenderness [Adjusted Odds Ratio (AOR)=2.29 (95%CI 1.14-4.60)], but a lesser vaginal polymorphonuclear (PMN) response compared to CT-PID by multivariable analysis. Of the 92 women with MG-PID, 54/92 (59%) received moxifloxacin (duration 10-14 days) and 37/54 returned a test of cure between 14-120 days after treatment– 27/37(73%) cases had a median of 7 days of a standard PID regimen containing doxycycline and metronidazole +/- azithromycin prior to commencing moxifloxacin. Microbial and clinical cure following moxifloxacin was 95% (95%CI 82-99) and 89% (95% CI 75-97), respectively.

## Conclusion:

Women meeting CDC criteria for presumptive treatment of MG-PID did not significantly differ to those with CT-PID. MG-PID was associated with more abdominal tenderness but less of a vaginal PMN inflammatory response than CT-PID. Moxifloxacin, often commenced after standard PID treatment, achieved high

microbiological cure (95%), in the context of emergence of quinolone resistance and high level macrolide resistance in Australia.

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