

Prevalence and Co-Infection of *Mycoplasma Genitalium* and *Gardnerella vaginalis*, Among Moroccan Women: Implications for Sexual and Reproductive Health
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INTRODUCTION & AIM

Women infertility affects around 50% of couples who have difficulty conceiving, impacting their emotional and relational well-being. While hormonal profiles and endometriosis are frequently assessed in reproductive health, bacterial vaginosis may act as a gateway indicator for detecting underlying infections, including undiagnosed Sexually transmitted infections STI. STI are emerging urogenital pathogens and represent a growing global sexual health challenge. Their coexistence with *Gardnerella vaginalis* a key contributor to bacterial vaginosis, raises clinical concerns due to potential reproductive health complications.

This study investigates the prevalence and co-infection of *Mycoplasma Genitalium* as a STI and *G. vaginalis* in a cohort of Moroccan women, evaluating potential impacts on sexual health and fertility. The aim of this study is to highlight that STIs may play a more significant role in fertility disorders than previously recognized, especially within the Moroccan context.

RESULTS & DISCUSSION

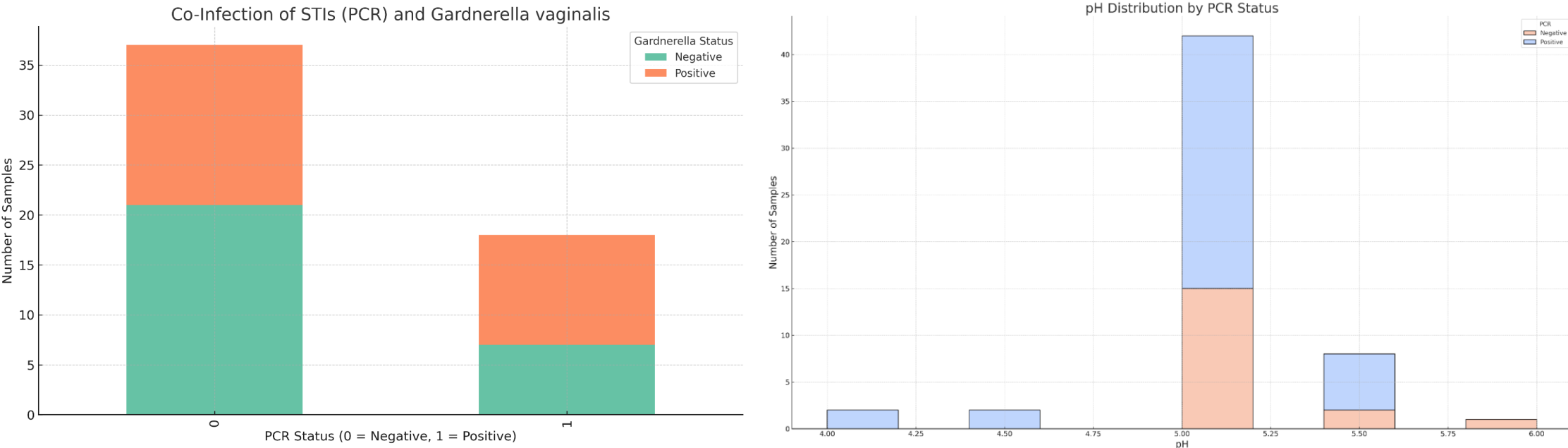


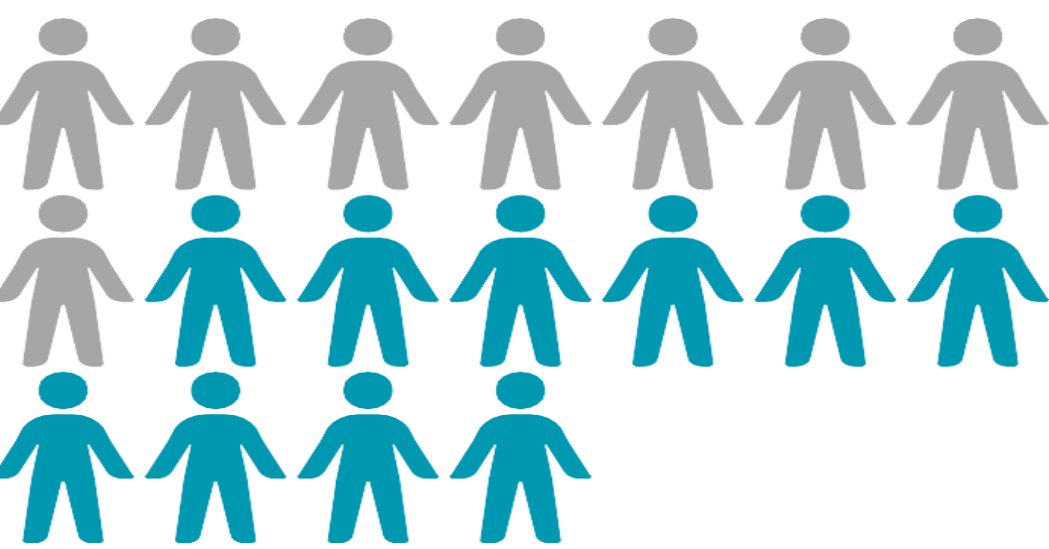
Figure 1: Distribution of *Gardnerella vaginalis* presence among PCR-positive and PCR-negative samples. While co-infection with *Mycoplasma* was frequently observed in *G. vaginalis*-positive women, a substantial number of *G. vaginalis*-only cases were PCR-negative ($p = 0,339$).

Figure 2: Distribution of vaginal pH values among PCR-positive and PCR-negative samples. A slight elevation in pH tends to be more frequent in PCR-positive cases, although no statistically significant difference was found ($p = 0,197$).

METHOD

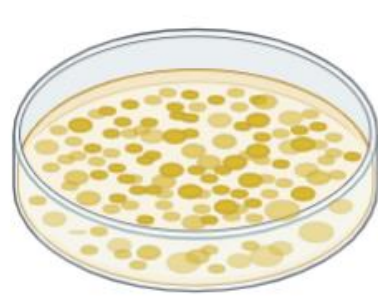
Vaginal swabs were collected from women presenting with symptoms of vaginal infection at gynecology and sexual health clinics in Tangier, Morocco. *G. vaginalis* detection was performed using standard bacteriological techniques: culture, Gram staining, and wet mount microscopy. *Mycoplasma Genitalium* analysis involved DNA extraction using a magnetic bead-based method, followed by multiplex PCR targeting *Mycoplasma Genitalium* genome.

Study population :

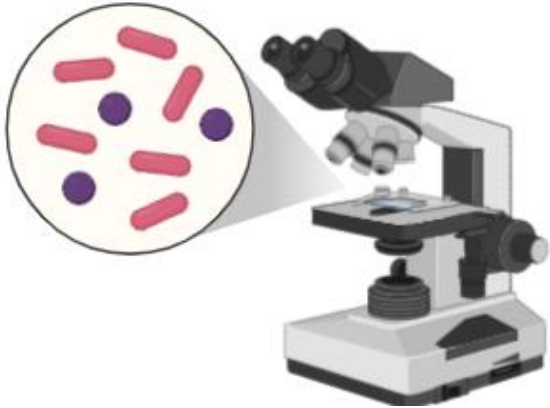


50 Women presenting vaginal infection symptostoms and discomfort

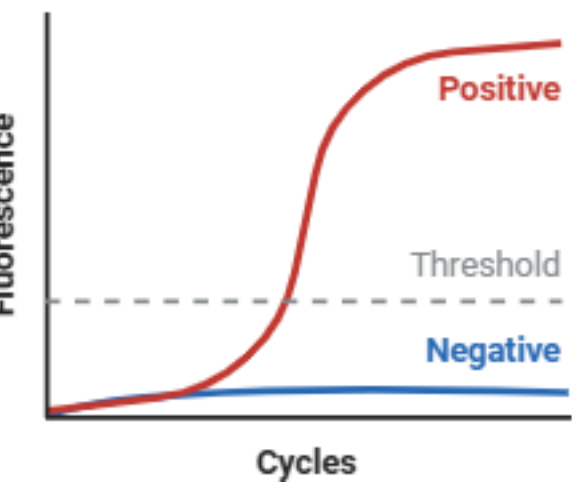
Laboratory techniques



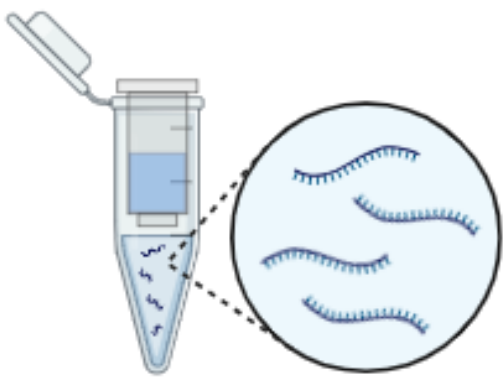
Culture



Wet amount microscopy and gram staining



PCR Amplification



Magnetic bead extraction DNA

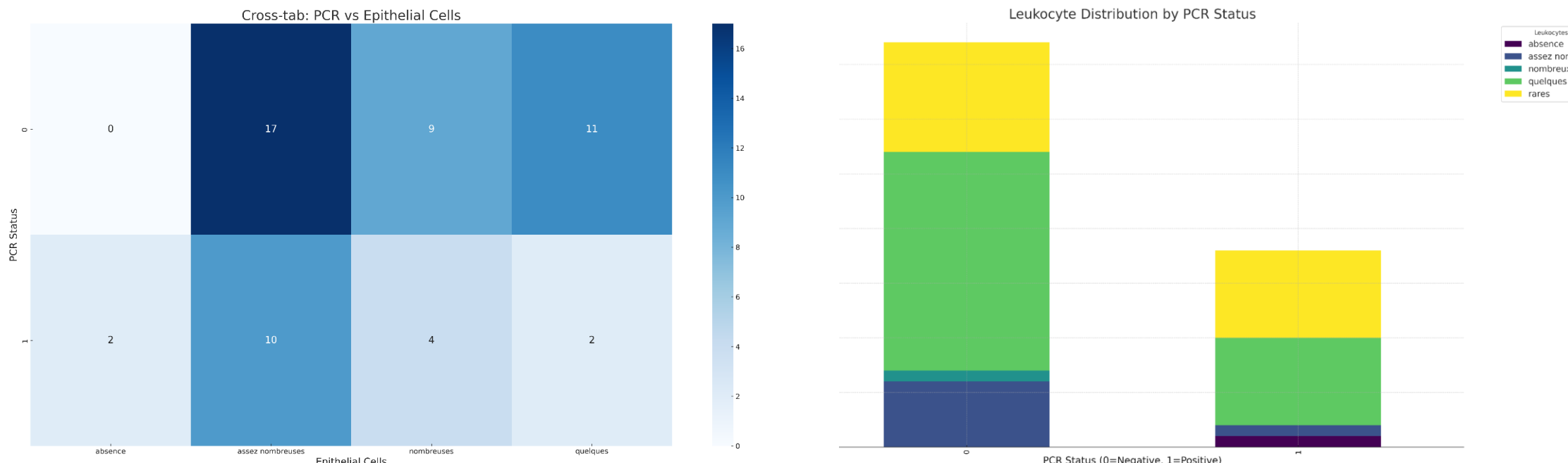


Figure 3: Cross-tabulation between epithelial cell abundance and *Mycoplasma* PCR result.

Figure 4: Distribution of leukocyte levels across PCR-positive and PCR-negative samples ($p = 0.291$).

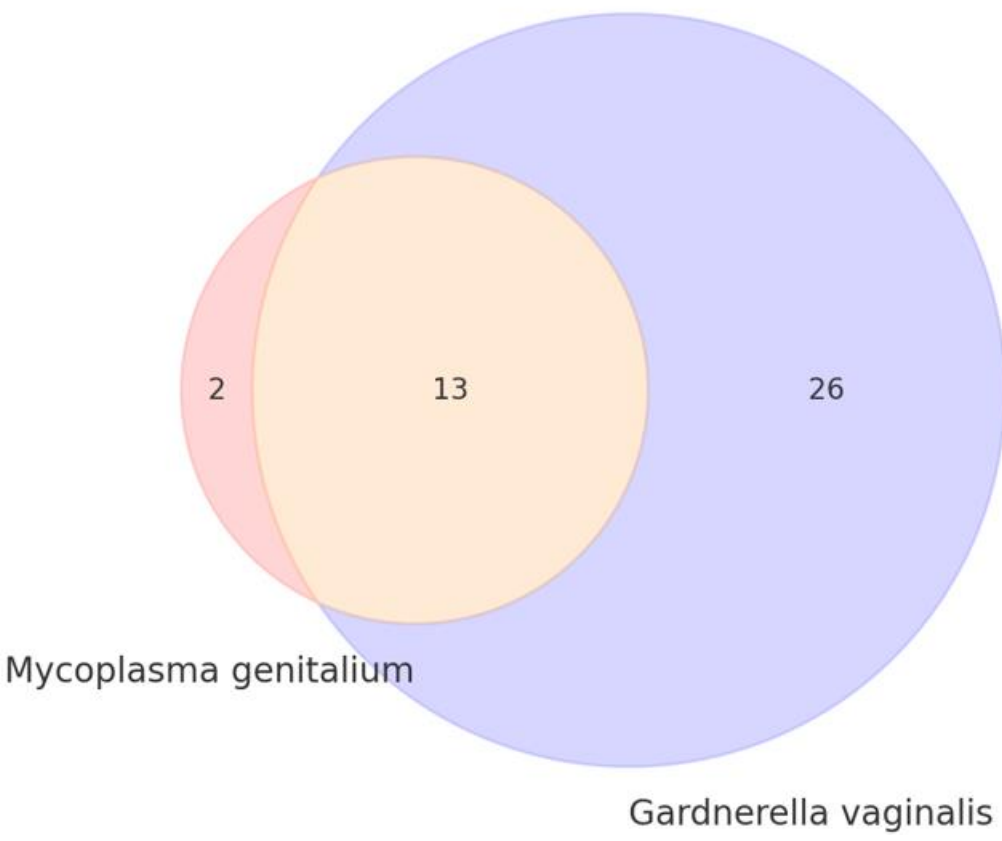


Figure 5: Distribution of patients according to the presence of *Mycoplasma Genitalium* and/or *Gardnerella vaginalis*.

CONCLUSION

Our study found that conventional cytological indicators, including pH, leukocyte count, and *Gardnerella vaginalis* presence, were **not significantly associated** with *Mycoplasma Genitalium* detection ($p > 0.05$). Despite this, the **clinical relevance of co-infections** remains notable, highlighting the **limitations of traditional diagnostic methods** that may fail to detect asymptomatic or subclinical infections. Given the strong link between *Mycoplasma* and infertility, integrating **molecular diagnostics such as PCR** into routine fertility assessments is crucial. Larger, long-term studies are needed to confirm these findings and better understand the **reproductive impact** of STI-related co-infections.

