

# Invasive Urogenital *Neisseria meningitidis* Serogroup Y Multilocus Sequence Type 1466

C.R. Robert George, Hemalatha Varadhan, Syeda Naqvi, Brett Locker, Matthew Howes, Sebastiaan van Hal,<sup>1</sup> Monica M. Lahra<sup>1</sup>

Author affiliations: New South Wales Health Pathology, New Lambton Heights, New South Wales, Australia (C.R.R. George, H. Varadhan, S. Naqvi); John Hunter Hospital, New Lambton Heights (B. Locker, M. Howes); New South Wales Health Pathology, Camperdown, New South Wales, Australia (S. van Hal); The University of Sydney, Camperdown (S. van Hal); New South Wales Health Pathology, Randwick, New South Wales, Australia (M.M. Lahra); The University of New South Wales, Randwick (M.M. Lahra).

DOI: <https://doi.org/10.3201/eid3110.250902>

**To the Editor:** Coincident outbreaks of *Neisseria meningitidis* serogroup Y (MenY) multilocus sequence type (ST) 1466 were detected in 2024, manifesting as invasive meningococcal disease (IMD) in the United States and urogenital infections in Australia (1). We read with interest the genomic analysis of globally available MenY ST1466 sequences that found that isolates from outbreaks in the United States and Australia were closely related (2). Although the study provided evidence that the urogenital site could act as a reservoir for IMD cases, risk estimates of urogenital MenY ST1466 ability to cause IMD or the propensity for the invasive isolates to inhabit the urogenital niche were lacking (2). Here, we report the case of a previously healthy 29-year-old woman who sought care at an emergency department for postprocedural pelvic inflammatory syndrome 3 days after the insertion of an intrauterine device.

<sup>1</sup>These senior authors contributed equally to this article.

We collected microbiological samples as the initial case workup. Vaginal swab specimens were negative for *Chlamydia trachomatis* and *N. gonorrhoeae*. Oropharyngeal specimens were not collected. MenY was isolated from a positive blood culture, and subsequent investigation included additional vaginal samples for culture. Although culture results were negative, MenY was detected by PCR from 5 vaginal and intrauterine device fluid samples. We conducted whole-genome sequencing on the blood culture after DNA extraction by using Illumina iSeq (Illumina, <https://www.illumina.com>), according to manufacturer instructions. In silico typing confirmed MenY ST1466. The patient was treated with ceftriaxone and made a full recovery.

This case represents presumptive progression of urogenital carriage of MenY ST1466 to invasive disease, supporting the premise that urogenital carriage is a risk factor for IMD (2). On the basis of the link between upper airway inflammation associated with influenza and IMD, it is biologically plausible that meningococcal urethritis similarly poses an increased risk for IMD (3). Concerns remain in Australia regarding the risk posed by urogenital MenY cases being detected through enhanced laboratory practices.

## References

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Address for correspondence: C.R. Robert George, New South Wales Health Pathology, Level 5, 45 Watt St, Newcastle, NSW 2300, Australia; email: [rob.george@health.nsw.gov.au](mailto:rob.george@health.nsw.gov.au)