



Australian Government
Department of Health

2022 · Volume 46

Communicable Diseases Intelligence

Australian Gonococcal Surveillance Programme, 1 July to 30 September 2021

Monica M Lahra, Masoud Shoushtari, Tiffany R Hogan

<https://doi.org/10.33321/cdi.2022.46.19>

Electronic publication date: 26/4/2022

<http://health.gov.au/cdi>

Communicable Diseases Intelligence

ISSN: 2209-6051 Online

This journal is indexed by Index Medicus and Medline.

Creative Commons Licence - Attribution-NonCommercial-NoDerivatives CC BY-NC-ND

© 2022 Commonwealth of Australia as represented by the Department of Health

This publication is licensed under a Creative Commons Attribution-Non-Commercial NoDerivatives 4.0 International Licence from <https://creativecommons.org/licenses/by-nc-nd/4.0/legalcode> (Licence). You must read and understand the Licence before using any material from this publication.

Restrictions

The Licence does not cover, and there is no permission given for, use of any of the following material found in this publication (if any):

- the Commonwealth Coat of Arms (by way of information, the terms under which the Coat of Arms may be used can be found at www.itsanhonour.gov.au);
- any logos (including the Department of Health's logo) and trademarks;
- any photographs and images;
- any signatures; and
- any material belonging to third parties.

Disclaimer

Opinions expressed in Communicable Diseases Intelligence are those of the authors and not necessarily those of the Australian Government Department of Health or the Communicable Diseases Network Australia. Data may be subject to revision.

Enquiries

Enquiries regarding any other use of this publication should be addressed to the Communication Branch, Department of Health, GPO Box 9848, Canberra ACT 2601, or via e-mail to: copyright@health.gov.au

Communicable Diseases Network Australia

Communicable Diseases Intelligence contributes to the work of the Communicable Diseases Network Australia.
<http://www.health.gov.au/cdna>



Communicable Diseases Intelligence (CDI) is a peer-reviewed scientific journal published by the Office of Health Protection and Response, Department of Health. The journal aims to disseminate information on the epidemiology, surveillance, prevention and control of communicable diseases of relevance to Australia.

Editor

Jennie Hood and Noel Lally

Deputy Editor

Simon Petrie

Design and Production

Kasra Yousefi

Editorial Advisory Board

David Durrheim,
Mark Ferson, John Kaldor,
Martyn Kirk and Linda Selvey

Website

<http://www.health.gov.au/cdi>

Contacts

CDI is produced by the Office of Health Protection and Response, Australian Government Department of Health, GPO Box 9848, (MDP 6) CANBERRA ACT 2601

Email:

cdi.editor@health.gov.au

Submit an Article

You are invited to submit your next communicable disease related article to the Communicable Diseases Intelligence (CDI) for consideration. More information regarding CDI can be found at: <http://health.gov.au/cdi>.

Further enquiries should be directed to:
cdi.editor@health.gov.au.

Australian Gonococcal Surveillance Programme, 1 July to 30 September 2021

Monica M Lahra, Masoud Shoushtari, Tiffany R Hogan

Introduction

The National Neisseria Network (NNN), Australia, comprises reference laboratories in each state and territory established in 1979. The NNN has reported data on susceptibility profiles for all *Neisseria gonorrhoeae* isolated from each jurisdiction for an agreed group of antimicrobial agents for the Australian Gonococcal Surveillance Programme (AGSP) since 1981. The antibiotics reported represent current or potential agents used for the treatment of gonorrhoea and include ceftriaxone; azithromycin; ciprofloxacin; and penicillin. More recently, gentamicin susceptibilities are included in the AGSP Annual Report. Ceftriaxone, combined with azithromycin, is the recommended treatment regimen for gonorrhoea in the majority of Australia. However, there are substantial geographic differences in susceptibility patterns in Australia, with certain remote regions of the Northern Territory and Western Australia having low gonococcal antimicrobial resistance rates. In these regions, an oral treatment regimen comprising amoxicillin, probenecid, and azithromycin is recommended for the treatment of gonorrhoea.

Results

A summary of the proportion of *Neisseria gonorrhoeae* isolates with decreased susceptibility (DS) to ceftriaxone (minimum inhibitory concentration, MIC \geq 0.06 mg/L); and the proportions resistant to azithromycin (MIC \geq 1.0 mg/L), penicillin (MIC \geq 1.0 mg/L), and ciprofloxacin (MIC \geq 1.0 mg/L) for Quarter 3 2021, is shown in Table 1.

Ceftriaxone

N. gonorrhoeae isolates with ceftriaxone MIC values \geq 0.06 mg/L are categorised as Decreased Susceptible (DS) in the AGSP. For trend data, reporting is at MIC values of 0.06 mg/L and MIC \geq 0.125 mg/L. In the third quarter of 2021, 0.65% of *N. gonorrhoeae* isolates tested had ceftriaxone DS. This proportion was lower than that reported in the first two quarters of 2021 and 2020 annually (0.94%), as shown in Table 2.¹ Noting, however, that the number of isolates tested was lower in 2021, coinciding with the public health measures in place during the COVID-19 pandemic.

Azithromycin

In the third quarter of 2021, the proportion of isolates resistant to azithromycin (MIC \geq 1.0 mg/L) in Australia was 4.5% (Table 2), similar to the first two quarters of 2021. The AGSP trend data for azithromycin resistance since 2010 is shown in Table 2. Globally, there have been reports of increased azithromycin resistance in *N. gonorrhoeae*, heightened since dual therapy was introduced.² In the third quarter of 2021, all states reported isolates with resistance to azithromycin, with the exception of South Australia and remote regions of Western Australia. No azithromycin-resistant isolates were reported in the territories.

Dual therapy using ceftriaxone plus azithromycin is the recommended treatment for gonorrhoea as a strategy to temper development of more widespread ceftriaxone resistance. Patients with infections in extragenital sites, where the isolate has decreased susceptibility to ceftriaxone, should have test of cure cultures collected.

Table 1: Gonococcal isolates showing decreased susceptibility to ceftriaxone, and resistance to azithromycin, ciprofloxacin and penicillin, Australia, 1 July to 30 September 2021, by state or territory

State or territory	Number of isolates tested		Decreased susceptibility		Resistance					
	Q3, 2021		Ceftriaxone MIC \geq 0.06 mg/L		Azithromycin MIC \geq 1.0 mg/L		Ciprofloxacin MIC \geq 1.0 mg/L		Penicillin ^a MIC \geq 1.0 mg/L	
	n	%	n	%	n	%	n	%	n	%
Australian Capital Territory	37	2.7	0	0.0	16	43.2	9	24.3		
New South Wales	388	0.5	25	6.4	284	73.2	156	40.2		
Queensland	276	0.4	3	1.1	112	40.6	74	26.8		
South Australia	68	0.0	0	0.0	14	20.6	28	41.2		
Tasmania	26	0.0	1	3.8	5	19.2	5	19.2		
Victoria	416	1.2	5	1.2	267	64.2	212	51.0		
Northern Territory non-remote	11	0.0	0	0.0	6	54.5	0	0.0		
Northern Territory remote	26	0.0	0	0.0	1	3.8	0	0.0		
Western Australia non-remote	125	0.0	7	5.6	47	37.6	36	28.8		
Western Australia remote	17	0.0	0	0.0	1	5.9	0	0.0		
Australia	1390	0.65	62	4.5	753	54.2	520	37.4		

a Penicillin resistance includes a MIC value of \geq 1.0 mg/L or penicillinase production.

Table 2: Percentage of gonococcal isolates with decreased susceptibility to ceftriaxone (MIC 0.06 and \geq 0.125 mg/L) and resistance to azithromycin (MIC \geq 1mg/L), Australia, 2010 to 2020, and 1 January to 31 March 2021, 1 April to 30 June 2021 and 1 July to 30 September 2021

YEAR	2010	2011	2012	2013	2014	2015	2016	2017	2018	2019	2020	2021 Q1	2021 Q2	2021 Q3
Ceftriaxone MIC 0.06 mg/L	4.80%	3.20%	4.10%	8.20%	4.80%	1.70%	1.65%	1.02%	1.67%	1.19%	0.87%	0.86%	0.90%	0.65%
Ceftriaxone MIC \geq 0.125 mg/L	0.10%	0.10%	0.30%	0.60%	0.60%	0.10%	0.05%	0.04%	0.06%	0.11%	0.07%	0.00%	0.00%	0.00%
Ceftriaxone DS Total	4.90%	3.30%	4.40%	8.80%	5.40%	1.80%	1.70%	1.06%	1.73%	1.30%	0.94%	0.86%	0.90%	0.65%
Azithromycin MIC \geq 1mg/L	n/a	1.1%	1.3%	2.1%	2.5%	2.6%	5.0%	9.3%	6.2%	4.6%	3.9%	4.8%	4.2%	4.5%

Continued surveillance to monitor *N. gonorrhoeae* with elevated MIC values, coupled with sentinel site surveillance in high-risk populations, remain essential to inform therapeutic strategies, identify incursion of resistant strains, and detect instances of treatment failure.

Author details

Monica M Lahra^{1,2}
Masoud Shoushtari¹
Tiffany R Hogan¹

1. The World Health Organization Collaborating Centre for STI and AMR, Sydney and Neisseria Reference Laboratory, NSW Health Pathology, Microbiology, The Prince of Wales Hospital, Randwick, NSW, 2031, Australia
2. School of Medical Sciences, Faculty of Medicine, the University of New South Wales, Kensington, NSW 2052 Australia.

Corresponding author

Professor Monica M Lahra

The World Health Organization Collaborating Centre for STI and AMR, Sydney and Neisseria Reference Laboratory, NSW Health Pathology Microbiology, The Prince of Wales Hospital, Randwick, NSW, 2031, Australia.

Telephone: +61 2 9382 9054

Facsimile: +61 2 9382 9098

Email: monica.lahra@health.nsw.gov.au

References

1. Lahra MM, Shoushtari M, George CRR, Armstrong BH, Hogan TR. Australian Gonococcal Surveillance Programme Annual Report 2020. *Commun Dis Intell (2018)*. 2021;45. doi: <https://doi.org/10.33321/cdi.2021.45.58>
2. Unemo M. Current and future antimicrobial treatment of gonorrhoea – the rapidly evolving *Neisseria gonorrhoeae* continues to challenge. *BMC Infect Dis*. 2015;15:364. doi: <https://doi.org/10.1186/s12879-015-1029-2>