

INVASIVE GROUP A STREPTOCOCCAL INFECTION IN NEW ZEALAND, 2023

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PREPARED FOR: Ministry of Health

CLIENT REPORT No: FW24014

PREPARED BY: Health Intelligence and Surveillance Group

PUBLISHED: 8 August 2024

This report is available on the ESR Digital Library
Published: 8 August 2024
Suggested citation:
Institute of Environmental Science and Research Ltd (ESR). Invasive group A streptococcal infection in New Zealand, 2023. ESR; 2024. Porirua, New Zealand
Client Report: FW24014
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ACKNOWLEDGEMENTS

This report was prepared as part of a Ministry of Health contract for scientific services.

Thanks to the following people for their contributions to this report:

- Diagnostic microbiology laboratories throughout New Zealand who participate in the national laboratory-based surveillance of invasive group A streptococcal infection by referring isolates to ESR;
- Audrey Tiong and Dr Juliet Elvy, ESR, for peer review.
- Associate Professor Nikki Moreland, University of Auckland, for external peer review.

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ABBREVIATIONS

Abbreviation	Description
CI	Confidence interval
DNA	Deoxyribonucleic acid
emm	M protein gene
ESR	Institute of Environmental Science and Research Ltd
GAS	Group A streptococcus
HISO	Health Information Standards Organisation
ICD-10-AM	International Statistical Classification of Diseases and Related Health Problems, Tenth Revision, Australian Modification
iGAS	Invasive group A streptococcal (infection)
IPL	ESR Invasive Pathogens Laboratory
NHI	National health index
NMDS	National Minimum Dataset (hospital discharges)
NZDep2018	New Zealand index of deprivation 2018
PCR	Polymerase chain reaction
RSV	Respiratory syncytial virus

SUMMARY

This report presents an overview of the epidemiology of invasive group A streptococcal (iGAS) infection in New Zealand in 2023. Surveillance of iGAS infection is undertaken by ESR, based on isolates sent to the ESR Invasive Pathogens Laboratory (IPL) from diagnostic laboratories in New Zealand.

Key findings

- Following the 2022/2023 iGAS increase in numerous high-income countries, there was also an increase in the incidence of iGAS in New Zealand. In 2023, iGAS infections rose to 10.9 cases per 100,000 population, compared with 4.8 per 100,000 in 2022 and 8.3 per 100,000 in 2019 (prior to the COVID-19 pandemic). This is the highest reported rate since at least 2002.
- In 2023, as in previous years, the highest incidence rates were among the youngest (less than 1 year) and oldest (80 years and over) age groups, with rates of 25.0 and 35.9 per 100,000 population, respectively. However, the most notable increase in incidence rates was among children aged 1–9 years, with a four-fold increase in 2023 compared with 2022 (from 2.5 to 9.9 per 100,000) and a three-fold increase compared with the pre-COVID-19 pandemic year 2019 (from 3.0 to 9.9 per 100,000). This increase among children is consistent with the 2022/2023 iGAS increase seen overseas.
- Pacific peoples and Māori were inequitably affected by iGAS infections with rates five and three times higher than the European or Other ethnic group, respectively. The higher incidence among Pacific peoples and Māori was seen for most age groups and all NZDep2018 quintiles.
- iGAS infection was also strongly associated with socioeconomic deprivation; almost two-thirds of iGAS cases lived in more deprived areas (NZDep2018 quintiles 4 and 5).
- In 2023, 62 unique *emm* types were characterised among iGAS isolates. *emm*1, *emm*12, *emm*92, and *emm*89 were the most common, with *emm*1 and *emm*12 increasing sharply between 2022 and 2023. The same *emm* types were also common in the 2022/2023 iGAS increase seen overseas.
- Ongoing surveillance of iGAS infection and molecular typing of isolates will be important to continue to monitor trends in New Zealand.

INTRODUCTION

Invasive group A streptococcal (iGAS) infections are part of the spectrum of disease that can be caused by the bacterium group A streptococcus (GAS), which includes necrotising fasciitis, cellulitis, bacteraemia, pneumonia, puerperal sepsis, and toxic shock syndrome. GAS also causes non-invasive diseases, such as pharyngitis, skin infections, and scarlet fever. Additional sequelae of GAS infection include acute rheumatic fever and post streptococcal glomerulonephritis. Together, invasive and non-invasive GAS diseases account for considerable mortality and morbidity in New Zealand [1, 2].

From early to mid-2022, several high-income countries (including the United Kingdom, France, Ireland, the Netherlands, Sweden, Canada, the United States, and Australia) reported a marked increase in GAS and iGAS infections, especially among children aged less than 10 years [3-7].

This report provides an overview of the epidemiology of iGAS infections in New Zealand in 2023.

METHODS

In New Zealand, diagnostic laboratories refer iGAS isolates to the ESR Invasive Pathogens Laboratory (IPL) for confirmation.

The following data was extracted from the ESR laboratory information system on 12 February 2024 for isolates collected between 1 January and 31 December 2023:

- national health index (NHI);
- age and sex;
- sample collection date;
- sample type (for example, blood, wound, tissue);
- symptoms, if available.

The NHI was used to obtain further information on cases, including ethnicity, New Zealand Index of Deprivation 2018 (NZDep2018), and date of death (if applicable). In addition, NHIs were used to obtain hospitalisation and diagnosis data (based on ICD-10-AM coding) from the National Minimum Dataset (NMDS). Diagnosis codes were also used to identify cases of invasive infection from non-sterile sites. The NMDS data was extracted on 4 March 2024.

Defining iGAS infections

Isolates that were included in this analysis were those collected by diagnostic laboratories in New Zealand in 2023 which met the case definition of iGAS infection.

The iGAS infection case definition that we used is:

1. isolation of GAS from a normally sterile site (e.g., blood, cerebrospinal fluid, joint, pleural, pericardial fluid) with or without evidence of severity;

OR

2. isolation of GAS from a non-sterile site with evidence of severity (e.g., streptococcal toxic-shock syndrome, necrotising fasciitis, meningitis or death).

The following ICD-10-AM codes were used to confirm whether GAS isolated from a nonsterile site met condition 2:

- A40.0 (sepsis due to streptococcus, group A)
- A48.3 (toxic shock syndrome)
- M72.6 (necrotising fasciitis)
- O85 (puerperal sepsis)

Population rates

All rates presented in this report are crude rates.

The 2023 mid-year population estimates from Statistics New Zealand were used to calculate the incidence rates.

All rates are presented as the number of cases per 100,000 population. Rates have not been reported where there were fewer than five cases in any category as this produces unstable rates.

Ethnicity

Prioritised ethnicity is used in this report. Level 1 ethnicity is used with ethnicity prioritised in the following order: Māori, Pacific peoples, Asian, and European or Other. For more detail on classification refer to the HISO Ethnicity Data Protocols [8].

Level 2 prioritised ethnicity is used to further describe the distribution of ethnicities in the Pacific peoples ethnic group. These are prioritised in the following order: Tokelauan, Fijian, Niuean, Tongan, Cook Island Māori, Samoan, and Other Pacific Peoples.

The incidence rates for level 1 ethnic groups were calculated by applying the ethnic proportions for the 2018 estimated resident population to the 2023 mid-year population estimates.

Socio-economic deprivation

The NZDep2018 is used to measure socio-economic deprivation. NZDep2018 is derived from a weighted combination of nine variables from the 2018 census, each reflecting a different aspect of material and socioeconomic deprivation [9]. The deprivation score is calculated for each geographical mesh block in New Zealand.

This report presents NZDep2018 by quintiles, where quintile 1 represents the least deprived areas and quintile 5 the most deprived areas.

The denominator data used to determine disease rates for NZDep2018 categories is based on the proportion of people in each NZDep2018 category from the usually resident 2018 census population.

30-day mortality rate

Mortality due to iGAS infection was estimated by reviewing deaths that occurred within 30 days of sample collection.

The 30-day mortality rate was calculated by dividing the number of cases who died within 30 days of sample collection by the total number of cases in a given year. If a sample collection date was not provided, the date the sample was received at ESR was used. This measure is used as a proxy for iGAS mortality. Since the associated death may not have been due to the iGAS infection, these results should be interpreted with caution.

emm typing

Diagnostic laboratories identify GAS isolates by culture following receipt of clinical samples. The diagnostic laboratories send isolates that meet the iGAS criteria to the ESR IPL for emm typing. Molecular typing is performed by polymerase chain reaction (PCR) and deoxyribonucleic acid (DNA) sequencing of the emm gene (method described by Beall et al [10]).

emm type diversity

The Simpson's reciprocal index was used to estimate iGAS strain diversity [11, 12].



IGAS INFECTION IN NEW ZEALAND

There were 568 iGAS isolates referred to ESR in 2023. Table 1 summarises the number of isolates by sample site. Similar to previous years, almost all of the isolates (541/568, 95.2%) were from a sterile site, and the majority of these (508/541, 93.9%) were from blood cultures. Only a small proportion (4.8%) of isolates from iGAS infection cases were from non-sterile or unknown body sites.

Table 1. Invasive GAS isolates by sample site, 2023

Isolate site		Number of isolates	Percent
	Blood	508	89.4
Sterile site	Bone and Joints	23	4.0
Sterile site	Body fluids	10	1.8
	Total	541	95.2
Non-sterile or	unknown site	27	4.8
Total number	of isolates	568	100

Incidence of iGAS infection in New Zealand

There was an increase in the incidence of iGAS infections in 2023, with a rate of 10.9 per 100,000 (568 cases), compared with 4.8 per 100,000 (244 cases) in 2022 (Figure 1). This is the highest annual rate reported since at least 2002. Between 2014 and 2019, rates of iGAS infection fluctuated and ranged between 7.0 and 9.4 per 100,000 in 2011. No changes in isolate referral criteria have occurred between 2002 and 2023. A decrease in iGAS infections occurred in 2020–2022 coinciding with the COVID-19 pandemic years.

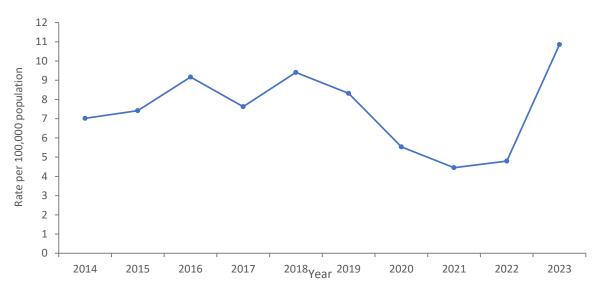


Figure 1. Incidence of iGAS infections in New Zealand, 2014–2023

Figure 2 shows that there is no clear seasonal pattern for iGAS infection in New Zealand. In 2023, the highest number of cases was reported in August (59 cases).

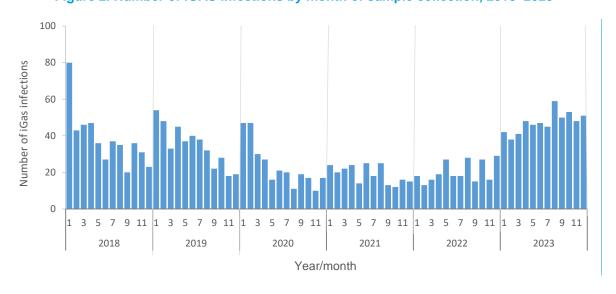


Figure 2. Number of iGAS infections by month of sample collection, 2018–2023

Incidence of iGAS infection by age group

Infants aged less than 1 year, and adults aged 80 years and over had the highest rates of iGAS infection in 2023 (25.0 and 35.9 per 100,000 respectively).

Among children aged 1–9 years, there was a four-fold increase in the incidence of iGAS infections in 2023, compared with 2022 (from 2.5 to 9.9 per 100,000), and a three-fold increase compared with 2019 (from 3.0 to 9.9 per 100,000) (Figure 3). In comparison, iGAS infection rates for infants aged less than 1 year and adults aged 70 years and over more than doubled compared with 2022 (11.7 to 25.0 per 100,000, and 12.9 to 27.5 per 100,000 respectively), but were similar to rates in 2019 (25.2 and 26.5 per 100,000 respectively).

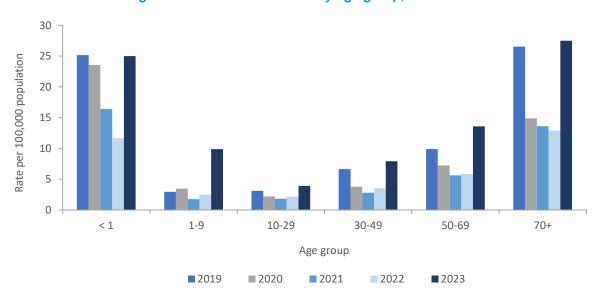


Figure 3. iGAS infection rates by age group, 2018–2023

As in previous years, males had a higher rate of iGAS infection than females (11.8 and 9.9 per 100,000 respectively).

Incidence of iGAS infection by ethnicity

Ethnicity was recorded for all 568 iGAS infections in 2023. Pacific peoples experienced the highest rate of iGAS infection (37.5 per 100,000), followed by Māori (20.1 per 100,000). Compared with 2022, iGAS infection rates increased across all ethnic groups, and this increase was highest among Pacific peoples and Māori (Figure 4). Table 2 in the appendix shows iGAS infections by ethnicity for 2018–2023.

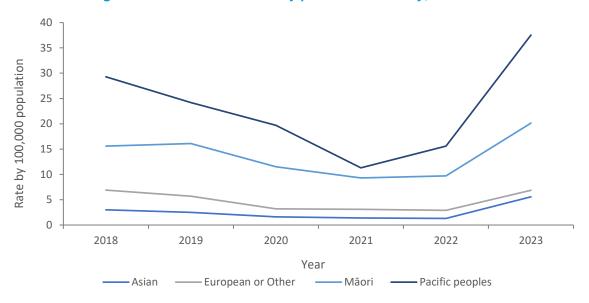


Figure 4. iGAS infection rates by prioritised ethnicity, 2018–2023

Samoan and Tongan are the largest Pacific ethnic groups in New Zealand. Of the 130 iGAS infections among Pacific peoples in 2023, most were Samoan (60 cases), followed by Tongan (32 cases), Cook Islands Māori and Fijian (12 cases each) and Niuean (7 cases). The remaining cases were identified as Other Pacific peoples (5 cases), or Pacific not further defined (1 case).

Figure 5 shows iGAS infection rates by age group and ethnicity in 2023. Pacific peoples had the highest rates among all age groups. Pacific peoples aged 60 years and over had the highest rate overall (132.6 per 100,000).

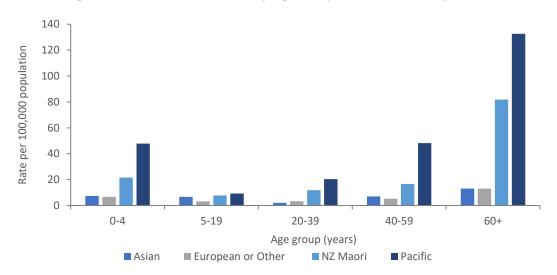


Figure 5. iGAS infection rates by age and prioritised ethnicity, 2023

Incidence of iGAS infection by socioeconomic deprivation

An NZDep2018 quintile could be assigned for 98.4% (559/568) of iGAS infections. The rate of iGAS infection increased with increasing socioeconomic deprivation, from 6.4 per 100,000 in quintile 1 to 21.2 per 100,000 in quintile 5; a more than three-fold increase (Table 3). Almost two-thirds (62.7%) of iGAS infections were in people living in the most deprived quintiles (NZDep2018 quintiles 4 and 5).

An increase in iGAS infection with increasing deprivation was seen among the Māori and European or Other ethnic groups. However, the Pacific peoples and Asian ethnic groups saw the highest rates in NZDep2018 quintile 4 (Figure 6). Across all NZDep2018 quintiles, Pacific peoples had the highest rate, followed by Māori.

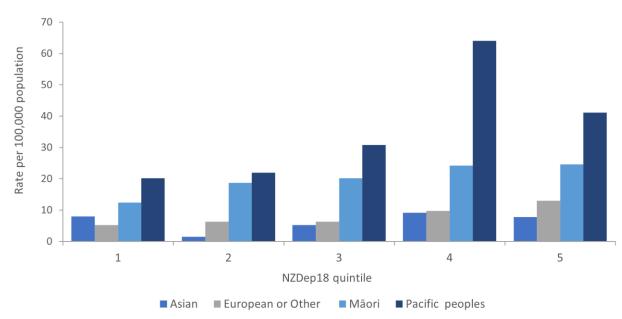


Figure 6. iGAS infection rates by prioritised ethnicity and NZDep2018 quintile, 2023

Incidence of iGAS infection by health district

The health district was known for 98.2% (558/568) of iGAS infections in 2023. Lakes (19.2 per 100,000), Tairāwhiti (19.0 per 100,000), Counties Manukau (18.9 per 100,000) and Taranaki (15.5 per 100,000) districts had the highest rates of iGAS infection (Figure 7). Table 4 shows iGAS infection rates by health district for 2018–2023.

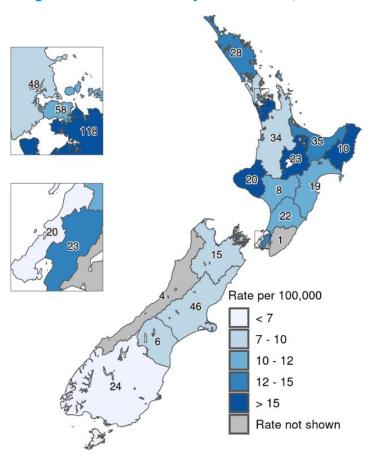


Figure 7. iGAS infections by health district, 2023

Rates not shown if count <5

30-day mortality

In 2023, there were 72 deaths that occurred within 30 days of collection of an iGAS sample. The 30-day mortality rate for iGAS infections in 2023 was 12.7%, compared with the 2017–2022 average of 10.5%.

The 30-day mortality rate increased with age, with the highest rate among those aged 80 years and over (29.7%), followed by those aged 70–79 years (18.7%). One death was reported for infants aged less than 1 year and two deaths for children between 1-9 within 30 days of sample collection (Figure 8).

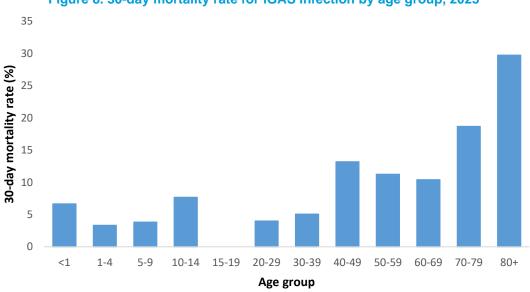


Figure 8. 30-day mortality rate for iGAS infection by age group, 2023

Minor differences were seen in the 30-day mortality rate by ethnicity. The mortality rate was 15.3% for the European or Other ethnic group, 13.4% for Māori, and 11.4% for Asian. Pacific peoples had the lowest mortality rate at 7.7%.

emm type distribution

emm typing was carried out on all 568 iGAS isolates received by the IPL in 2023. There were 62 *emm* types associated with iGAS infections in New Zealand in 2023. This is compared with 47 different *emm* types in 2022, 37 *emm* types in 2021, and 51 *emm* types in 2020.

The most common *emm* types were *emm*1 and *emm*12, accounting for 15% of isolates each. Both *emm*1 and *emm*12 increased sharply from 2022 to 2023. In contrast, *emm*114, which has been the dominant strain since 2019, decreased in 2023 (Figure 9). Table 5 in the Appendix shows the number of iGAS infections by *emm* type from 2018 to 2023.

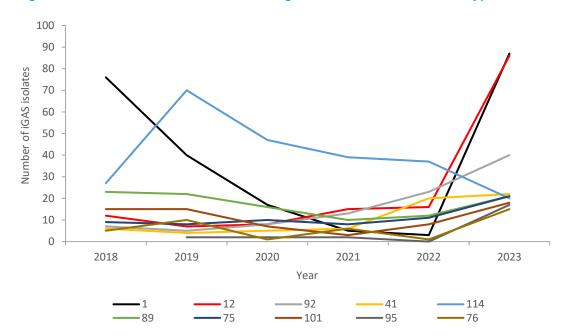


Figure 9. Number of iGAS infections among the 10 most common emm types, 2018–2023

In 2023, the Simpson's reciprocal index was 15.81 (95% Confidence Interval (CI): 15.80–15.81), suggesting a moderate diversity of *emm* types. Between 2018 and 2023, there has been an overall decreasing trend in the diversity of *emm* types (Table 6).

DISCUSSION

iGAS infections increased in New Zealand in 2023, mirroring the 2022/2023 increase overseas

There was a sharp increase in the incidence of iGAS infections between 2022 and 2023 (from 4.8 to 10.9 per 100,000), following a period of low incidence during the COVID-19 pandemic. The incidence of iGAS infections in 2023 is the highest reported in New Zealand in more than 20 years. Between 2002 and 2022, iGAS infection rates ranged from 3.8 per 100,000 in 2002 to 9.3 per 100,000 in 2011 [13].

This increase in iGAS infections follows the 2022/2023 iGAS increase seen in high income countries (including in the United Kingdom, France, Ireland, the Netherlands, Sweden, Canada, the United States and Australia [3-7]), which coincided with the easing of pandemic restrictions. While higher case numbers continue to be seen in a number of countries, including parts of Canada and Australia [14-16], in the United Kingdom, iGAS infections have decreased but remain higher than pre-pandemic levels [17].

The drivers of the increase in iGAS infections in New Zealand and overseas remain unclear. A possible explanation is that low exposure to GAS and predisposing illnesses due to COVID-19 pandemic measures may have reduced immunity to GAS, particularly among children. Increased population mixing and a rise in predisposing viral illnesses (e.g., respiratory viruses or varicella zoster) post-pandemic have also been theorised to contribute to this increase [6, 18]. The delay in the increase in iGAS infections in New Zealand may be due to the removal of COVID-19 restrictions in New Zealand occurring later than many other countries, and/or delayed introduction of iGAS strains that were common in the iGAS increase overseas.

No clear seasonal pattern was seen in iGAS infection 2023, similar to previous years. This is in contrast to some other temperate climates, where GAS and iGAS infections typically peak in winter and early spring, coinciding with increases in respiratory infections, including influenza and respiratory syncytial virus (RSV) [19, 20].

iGAS increases were most notable among children

In 2023, iGAS infections followed a U-shaped curve, with those aged less than 1 year and 80 years and over having the highest rates. The largest increases compared with the prepandemic era were among children aged 1–9 years; this is consistent with reports from a number of countries during the 2022/23 iGAS increase where an increase in paediatric infections was noted [18, 21-23].

Inequities in iGAS infections persist

As in previous years, ethnic inequities were seen in overall rates, with rates among Pacific peoples five times higher than European or Other, and rates among Māori three times higher. Pacific peoples and Māori had the highest rates across all NZDep2018 quintiles and most age groups. These disparities are similar to ethnic inequities reported internationally, with iGAS infections disproportionally affecting Indigenous peoples and other ethnic minorities [24-26].

Socioeconomic deprivation continues to be strongly associated with iGAS infection. Almost two-thirds of iGAS infections are reported among people living in areas with higher socioeconomic deprivation (quintiles 4 and 5).

emm1 and emm12 increased sharply to become the dominant emm types in 2023

In 2023, the most dominant emm types were emm1, emm12, and emm92, with the prevalence of emm1 and emm12 increasing sharply from 2022 to 2023. The 2022/23 iGAS increase overseas shows a similar distribution, with emm1 and emm12 identified as the dominant emm types in the United Kingdom [7], the Netherlands [21], Australia [22], Italy [27], the United States [23], Denmark [28], and France [6]. A number of studies highlight emm1 as a major contributor to the iGAS increase [28, 29]. In particular, the M1UK strain has gained considerable attention in the context of the 2022/2023 iGAS increase, given its increasing prevalence, higher virulence, and emm1's association with more severe infection than other genotypes [28, 30, 31]. M1UK was the most prevalent variant during the 2022/23 winter season in both England and Australia and accounted for 31% of all cases under 15 years in England [18, 29].

The Simpson's reciprocal index in 2023 was 15.8 and, considering the 62 emm types characterised in 2023, this suggests there were some dominant emm types combined with a reasonable level of strain diversity. The 2023 index is slightly lower than that reported in a 2018–2019 sample of school-aged children with GAS infections in Auckland (20.2; 95%CI: 16.6–23.8) [32], and falls within the range reported in a recent study that examined GAS emm type diversity in low- and high-income settings [33]. Overall, there has been a slight decline in emm type diversity since 2018.

The 30-day mortality rate increased in 2023, rising from an average rate of 10.5% in 2017– 2022 to 12.7% in 2023. Higher case fatality rates were reported in some countries during the 2022/2023 iGAS increase, including Canada [34], and Ireland [35]. It has been postulated that the dominance of emm1 GAS drove the rise in mortality observed in multiple high-income countries in 2022/2023 [33, 34]. The shift in emm types in New Zealand in 2023 may have contributed to the increase in the 30-day mortality rate seen amongst iGAS infection cases. While further research is needed to better understand the distribution of M1UK in New Zealand, M1UK was detected in New Zealand as early as 2018 [20].

Whole genome sequencing information was not available for our 2023 surveillance data, therefore the prevalence and distribution of M1UK in New Zealand has not been described in this report.

Data interpretation and limitations

There are a few notable limitations to the data used in this report. Firstly, iGAS is currently not a notifiable disease in New Zealand, and surveillance relies on voluntary referral of isolates to ESR. An audit of iGAS isolates referred to the ESR IPL in 2022 found that 84% of isolates identified by participating laboratories were referred to ESR and that the current surveillance system underestimates the burden of iGAS infection in New Zealand [13]. Following the audit, the findings were shared with laboratories with a reminder to refer samples to ESR. This may have changed referral patterns for isolates to ESR in 2023 and could affect trend data. Later in 2024 iGAS infection will become a notifiable disease, which will likely increase iGAS reporting.

The ethnic-specific population estimates were based on the 2018 census population and adjusted for any undercounting by Statistics New Zealand using information from multiple sources. Despite these efforts to adjust population estimates and impute missing ethnicity data, research suggests that undercounts of Pacific and Māori populations remain [36]. NHI data are also known to undercount both Māori and Pacific peoples. Together, these limitations affect the quality and reliability of our Māori and Pacific population data and therefore the estimates of iGAS incidence in these groups [37]. Additionally, reliable and up-to-date population denominators for L2 Pacific ethnicities were not available [36], precluding the ability to calculate incidence rates for these groups.

Finally, whole genome sequencing data was not available as part of our surveillance data. Therefore, we were unable to describe the distribution of the iGAS M1UK lineage.

CONCLUSION

There was an increase in iGAS infections in New Zealand in 2023, with the highest annual rate seen in more than 20 years, at 10.9 per 100,000. iGAS rates were highest among the youngest (less than 1 year) and oldest (80 years and over) age groups. While iGAS infections increased across all age groups, the increase was most notable among those aged 1-4 and 5-9 years compared with the pre-COVID-19 pandemic era. Additionally, an increase in the 30-day mortality rate was seen in 2023. In 2023, emm1 and emm12 were the most common emm types. These trends are consistent with those reported during the 2022/2023 iGAS increase overseas.

Inequities continue to be experienced by Māori and Pacific peoples. Given the high and increasing number of iGAS infections seen in 2023, continued surveillance of iGAS infections, alongside efforts to address systemic and health care access issues, remain important. The upcoming change to make iGAS notifiable will improve the completeness of reporting and data collection for iGAS infection surveillance in New Zealand.

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APPENDIX

Table 2. Number of cases and rate of iGAS infection by prioritised ethnicity, 2018-2023

	2018		2019		2020		2021		2022		2023	
Ethnic group	Cases	Rate ¹										
Māori	127	15.6	133	16.1	97	11.5	79	9.3	83	9.7	172	20.1
Pacific	97	29.3	81	24.2	67	19.7	39	11.3	54	15.6	130	37.5
Asian	22	3	19	2.5	12	1.6	11	1.4	10	1.3	44	5.6
European/Other	207	6.9	174	5.7	100	3.2	99	3.1	90	2.9	222	6.9
Unknown	8		7		6				7			
Total	461	9.4	414	8.3	282	5.5	228	4.5	244	4.8	568	10.8

¹ Rate per 100,000 population.

Table 3. Number of cases and rate of iGAS infection by prioritised ethnicity and NZDep2018 quintile, 2023

NZDep2018 quintile	Asian		European or Other		Māori		Pacific peoples		Total	
	Cases	Rate ¹	Cases	Rate ¹	Cases	Rate ¹	Cases	Rate ¹	Cases	Rate ¹
1 (least deprived)	9	8.0	37	5.3	8	12.4	3	20.2	57	6.4
2	2	1.5	42	6.4	16	18.7	5	21.9	65	7.2
3	8	5.3	39	6.3	23	20.1	11	30.8	81	8.8
4	14	9.2	54	9.7	42	24.2	39	64.1	149	15.8
5 (most deprived)	9	7.7	47	13.0	81	24.6	70	41.1	207	21.2

¹ Rate per 100,000 population.

Table 4. Number of cases and rate of iGAS infection by health district, 2018–2023

Health district	20 ⁻	18	20 ⁻	19	20:	20	202	21	20:	22	202	23
Health district	Cases	Rate ¹	Cases	Rate ¹	Cases	Rate ¹	Cases	Rate ¹	Cases	Rate ¹	Cases	Rate ¹
Northland	41	22.1	35	18.5	15	7.7	19	9.6	20	9.9	28	13.7
Waitematā	40	6.5	44	7.0	31	4.9	17	2.7	22	3.5	48	7.4
Auckland	56	11.4	37	7.4	26	5.1	30	6.0	23	4.8	58	11.8
Counties Manukau	106	18.7	88	15.2	78	13.1	35	5.8	44	7.3	118	18.9
Waikato	35	8.3	36	8.4	21	4.8	22	4.9	19	4.2	34	7.4
Lakes	9	7.9	22	19.1	7	6.0	8	6.8	7	5.9	23	19.2
Bay of Plenty	11	4.4	26	10.1	18	6.8	23	8.5	24	8.7	35	12.5
Tairāwhiti	9	18.2	5	10.0	8	15.7	6	11.7	4	-	10	19.0
Hawke's Bay	17	9.9	16	9.2	20	11.1	13	7.2	12	6.6	19	10.3
Taranaki	15	12.4	15	12.2	9	7.2	9	7.1	0	-	20	15.5
MidCentral	14	7.7	8	4.3	7	3.7	9	4.8	3	-	22	11.4
Whanganui	8	12.0	6	8.9	3	-	2	-	9	12.9	8	11.4
Capital & Coast	15	4.7	15	4.7	7	2.2	4	-	12	3.7	20	6.1
Hutt Valley	5	3.2	10	6.4	6	3.8	2	-	4	-	23	14.2
Wairarapa	2	-	1	-	1	-	0	-	2	-	1	-
Nelson Marlborough	12	7.7	3	-	1	-	8	4.9	1	-	15	9.0
West Coast	0	-	1	-	0.	-	0	-	0	-	4	-
Canterbury	34	6.1	24	4.2	17	2.9	13	2.2	17	2.9	46	7.6
South Canterbury	10	16.4	5	8.2	1	-	0	-	0	-	6	9.5
Southern	19	5.6	11	3.2	5	1.4	8	2.3	10	2.9	24	6.7

¹ Rate per 100,000 population. Where there were fewer than five cases in any category, a rate has not been calculated.

Table 5. Number of iGAS infections by emm type, 2018–2023

emm type*	2018	2019	2020	2021	2022	2023
1	76	40	17	5	3	87
12	12	7	8	15	16	86
92	7	5	8	13	23	40
41	6	4	5	6	20	22
114	27	70	47	39	37	20
89	23	22	16	10	12	21
75	9	8	10	8	11	21
101	15	15	7	3	8	18
95	0	2	2	2	0	17
76	5	10	1	6	1	15
108	6	0	0	7	6	14
53	19	30	32	28	11	14
58	17	2	1	2	4	14
82	15	9	11	8	7	14
103	38	27	20	12	12	13
77	8	5	3	4	4	11
81	22	18	11	7	2	10
22	2	7	4	5	5	9
44	9	9	0	3	3	8
91	6	8	4	4	4	8
49	7	12	11	7	1	7
3	6	1	3		2	6
28	4	9	4	4	2	5
65	5	12	2	4	1	5
86	2	1	0	0	0	5
112	4	3	0	2	0	4
116	2	0	2	3	0	4
4	13	10	5	3	1	3
59	4	5	2	1	3	3
100	2	0	1	0	6	3
25	1	1	2	0	0	3
68	3	2	2	0	0	3
9	0	1	1	0	0	3
110	2	1	0	0	0	3
71	2	1	0	0	0	3
33	1	0	0	1	0	3
99	1	0	0	0	0	3
109	0	0	0	0	0	3
197	0	0	0	0	0	3

*emm types with less than two cases in 2023 have not been included in the table

Table 6. Simpson's reciprocal index, 2018–2023

Year	Simpson's reciprocal index (95% CI)
2018	20.54 (20.53–20.55)
2019	16.97 (16.96–16.98)
2020	15.60 (15.58–15.61)
2021	16.74 (16.73–16.76)
2022	17.68 (17.67–17.69)
2023	15.81 (15.80–15.81)



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