Epidemiology of dengue reported in the World Health Organization's Western Pacific Region, 2013–2019

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The global burden of dengue, an emerging and re-emerging mosquito-borne disease, increased during the 20-year period ending in 2019, with approximately 70% of cases estimated to have been in Asia. This report describes the epidemiology of dengue in the World Health Organization's Western Pacific Region during 2013-2019 using regional surveillance data reported from indicator-based surveillance systems from countries and areas in the Region, supplemented by publicly available dengue outbreak situation reports. The total reported annual number of dengue cases in the Region increased from 430 023 in 2013 to 1 050 285 in 2019, surpassing 1 million cases for the first time in 2019. The reported case-fatality ratio ranged from 0.19% (724/376 972 in 2014 and 2030/1 050 285 in 2019) to 0.30% (1380/458 843 in 2016). The introduction or reintroduction of serotypes to specific areas caused several outbreaks and rare occurrences of local transmission in places where dengue was not previously reported. This report reinforces the increased importance of dengue surveillance systems in monitoring dengue across the Region.

he global burden of dengue, an emerging and reemerging mosquito-borne disease, increased from 2000 to 2019.¹ An estimated 70% of dengue virus infections are thought to occur in Asia.² It has previously been reported that in the World Health Organization's (WHO's) Western Pacific Region, the number of dengue cases increased from approximately 200 000 in 2008 to more than 450 000 in 2015.³ During this period, several countries and areas in the Region experienced large-scale outbreaks.4-6

Dengue is a public health threat because it is associated with large outbreaks in communities, severe disease and mortality.¹ Host immunity factors, such as serotype interaction, antibody-dependent enhancement and cross-immunity, complicate the clinical course, which leads to challenges in managing severe cases.^{1,7} Additionally, socioeconomic and environmental factors,

including climate change, drive disease transmission and complicate prevention and control activities.

In response to these challenges, a revised Western Pacific Regional Action Plan for Dengue Prevention and Control was developed and endorsed at the 67th meeting of the Regional Committee for the Western Pacific in October 2016.³ The Plan has guided countries and areas in the Region on improving the laboratory diagnosis of dengue, and the clinical management, surveillance and sustainable vector management for the disease to reduce morbidity and mortality, and decrease impacts on health systems.

Sharing information and data about dengue helps countries and areas better understand transmission patterns and supports the implementation of dengue prevention and control measures.² As a continuation of

Published: 22 March 2023 doi: 10.5365/wpsar.2023.14.1.973

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previous Regional dengue epidemiology updates in 2010, 2011 and 2012,⁸⁻¹⁰ this analysis reports data collated by the WHO Regional Office for the Western Pacific to describe the epidemiology of dengue in the Region from 2013 to 2019 using regional surveillance data. Data from 2020 to 2021 were excluded due to changes in reporting practices, population movement and people's behaviours as a result of the COVID-19 pandemic.

METHODS

Regional dengue data from 2013 to 2019 were collated from indicator-based surveillance systems from countries and areas in the Region. Information was also collected about laboratory sampling schemes and the confirmation methods used by each country and area. Data were either sent to WHO by ministries of health or collected from official websites where they were publicly available. Additional data - including serotype information, case definitions, and the numbers of clinically confirmed cases, laboratoryconfirmed cases and imported cases and deaths - were provided by Australia, Cambodia, Japan, the Republic of Korea, Malaysia, New Zealand, Pacific Island countries and areas (PICs), the Philippines, Singapore and Viet Nam. Information was reported based on the standard dengue case definitions used in each country or area (Table 1). Missing data were supplemented by using official dengue outbreak situation reports published on ReliefWeb (https:// reliefweb.int/), manuscripts identified through PubMed using keywords ["dengue" AND "outbreak" AND "(country/ area name)"], yearly aggregated data collected from all countries and areas in the Region through International Health Regulations (2005) channels, and WHO Regional biweekly dengue reports.¹¹

Table 1 summarizes the dengue surveillance systems, case definitions, laboratory sampling methods and serotype data. It was not possible to compare trends between countries and areas due to the differences in surveillance methods and reporting practices. The crude regional case notification rate per 100 000 population per year was calculated using the number of cases and deaths reported to WHO and standard calculation methods:

Case notification rate per 100 000 population per year = $(c/p) \times 100 000$ and

> 95% confidence interval = (100 000/p) (c \pm 1.96 \times \sqrt{c}),

where *c* is the total dengue notification case count in a given year and *p* is the population estimate for the Region in a given year. United Nations population estimate data were used for calculations. Population data for the Pitcairn Islands were not included in the United Nations population database.¹² Therefore, we used the closest population estimates based on the Pitcairn Islands' government website. In this report, an outbreak is defined as the "occurrence of cases of disease in excess of what would normally be expected in a defined community, geographical area or season".¹³

RESULTS

In the Region, the total number of annual dengue cases reported increased from 430 023 cases from 22 countries and areas in 2013 to 1 050 285 cases from 18 countries and areas in 2019 (data not shown). The lowest annual number of cases during these 7 years was reported in 2014, with 376 972 cases. In 2019, the total number of reported dengue cases surpassed 1 million for the first time. From 2013 to 2019, the case-fatality ratio (CFR) fluctuated between 0.19% (724/376 972 reported in 2014 and 2030/1 050 285 in 2019) and 0.30% (1380/458 843 reported in 2016) (Fig. 1). The number of cases reported from the PICs did not show a clear trend, with more cases reported in 2013 and 2014 compared with 2015 and 2016 (Fig. 2). There were challenges in calculating the CFRs for some countries due to limited reporting on dengue cases or deaths associated with dengue, or both.

From 2013 to 2018, the crude annual case notification rates in the Region ranged from a low of 19.82/100 000 population per year (95% confidence interval [CI]: 19.76–19.89) in 2014 to a high of 26.84/100 000 population per year (95% CI: 26.77–26.92) in 2015. In 2019, the case notification rate increased two-fold to 53.71/100 000 population per year (95% CI: 53.61–53.81) (Table 2).

From 2013 to 2019, large-scale outbreaks with notable increases in the number of cases were reported in multiple countries. Outbreaks were reported from the PICs every year from 2013 to 2019. There were two notable years, 2017 and 2019, when multiple outbreaks were reported across the Region, including in the PICs, with seven countries reporting outbreaks. All dengue serotypes (DENV-1, DENV-2, DENV-3 and DENV-4)

Table 1. Dengue clinical case definitions, and laboratory sampling and testing methods used for surveillance in countries, WHO Western Pacific Region, 2019

	Case definition ^a	· · · · · · · · · · · · · · · · · · ·			
Country	Clinically confirmed case	Laboratory confirmation required	Laboratory sampling and testing method	Surveillance and case reporting	
Australia ⁴⁶	Fever, headache, arthralgia, myalgia, rash, nausea or vomiting	Yes	NS1, IgG seroconversion, IgM detection, nucleic acid or virus isolation. All clinically diagnosed cases have laboratory testing.	All confirmed cases require both laboratory- definitive evidence and clinical evidence.	
				Both confirmed and probable cases are nationally notifiable.	
Cambodia ⁴⁷	Suspected dengue: high fever (39–40 °C) for 2–7 days (usually 3–4 days), with two or more of the following signs: flushed face, headache, retro- orbital pain, myalgia or arthralgia, cutaneous rash, haemorrhagic signs (e.g. petechiae, positive tourniquet test) and leucopoenia	No	Data are collected for the Cambodia Laboratory Information System, composed of 32 hospital laboratories where NS1 detection is conducted. Laboratory testing: Antibody haemagglutination inhibition ≥1/1280 or IgM- or IgG-positive by ELISA in convalescent serum.	Suspected cases are reported from all national hospitals and all provincial hospitals, but not from private clinics.	
	Probable dengue: signs of suspected dengue plus laboratory test results (right column) or a case that occurred in an area where a dengue case has been confirmed				
China⁵	More than two of the following symptoms: acute onset fever, severe headache, orbital pain, myalgia, arthralgia, fatigue, a history of travel in a dengue-endemic area during the 15 days before symptom onset or cohabitation with an individual with confirmed dengue, or no travel history but with a rash or positive tourniquet test AND leucopoenia or thrombocytopenia or serum IgM positivity	No	Real-time PCR, NS1 in acute-phase serum or virus isolation from an acutely infected patient's serum.	Both clinically confirmed and laboratory-confirmed cases are notified as an infectious disease.	
Japan ⁴⁸	Symptoms including acute onset of fever lasting for 2–7 days (commonly biphasic), headache, retro-orbital pain, arthralgia, myalgia, fatigue, conjunctivitis or rash AND laboratory confirmation (right column)	Yes	All clinically diagnosed cases have laboratory testing. Laboratory confirmation requires at least one of the following: a positive PCR test, NS1 detection, serology (IgM, seroconversion) and/or virus isolation.	All reported cases have laboratory testing.	
Lao People's Democratic Republic ²²	WHO 2009 dengue case classification ^c	No	Laboratory testing is conducted by RDT and PCR on a subset of specimens referred to the laboratories. Serotyping is also conducted on a subset of specimens.	Clinically confirmed cases (dengue with and without warning signs and severe dengue cases) are reported.	
Malaysia ⁴⁹	WHO 2009 dengue case classification ^c	Yes	All suspected cases are tested by the rapid combo test for NS1, IgM and IgG; ELISA for the dengue antigen and serology, real-time PCR for detecting viral RNA, or by viral isolation.	All reported cases have laboratory testing.	
New Zealand ^{31,50}	Acute onset of fever; headache, particularly retro-orbital; myalgia and arthralgia; and a fine rash, which may be itchy and usually begins on the extremities but spares the palms and soles. Other symptoms include weakness, depression, anorexia, abnormal taste, sore throat, coughing, vomiting and abdominal pain.	No	At least one of the following tests is required for laboratory confirmation: viral isolation, dengue virus (DENV) nucleic acid amplification, IgM or IgG seroconversion, a significant increase in antibodies (four-fold or greater) by serological test.	Both clinically confirmed and laboratory-confirmed cases are reported.	

Philippines ^{51–53}	WHO 2009 dengue case classification ^c	No	A subset of suspected cases have laboratory testing.	Suspected cases are reported.
	In addition, suspected cases are those who were previously well but have acute febrile illness for 2–7 days with clinical signs and symptoms of dengue.		Confirmed dengue is defined as a suspected case with positive viral culture isolation and/or PCR.	
			Probable dengue cases are NS1- or IgM-positive.	
Republic of Korea ⁵⁴	Acute onset of fever, headache, arthralgia, myalgia, leucopoenia, thrombocytopenia or bleeding AND laboratory confirmation (right column)	Yes	All clinically diagnosed cases have laboratory testing by real-time PCR or ELISA (IgM).	All reported cases have laboratory testing.
Singapore ⁵⁵	A clinical case meets the criteria of fever, headache, backache, myalgia, rash, abdominal discomfort and thrombocytopenia.	Yes	Samples are tested by the laboratory as ordered by the physician. Laboratory confirmation is done by dengue NS1 antigen testing, IgM or PCR.	All reported cases have laboratory testing.
Viet Nam ⁵⁶	^{m⁵⁶} Acute onset of fever lasting 2–7 days No AND at least two of the following: haemorrhagic manifestation or presentation, headache, loss of appetite, nausea, vomiting, rash, muscle pain, joint pain, orbital pain, lethargy, abdominal pain		MAC-ELISA is conducted for at least 7% of clinical cases and virus isolation is conducted for at least 3% of clinical cases. In an outbreak, at least 5–10 suspected cases are tested.	Both clinically confirmed and laboratory-confirmed cases are reported.

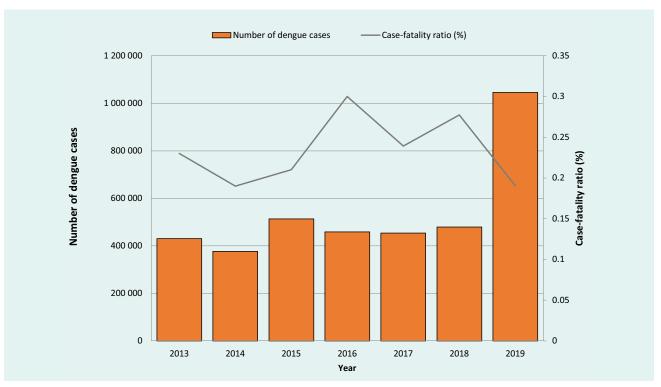
ELISA: enzyme-linked immunosorbent assay; IgG: immunoglobulin G; IgM: immunoglobulin M; MAC-ELISA: dengue IgM capture ELISA; NS1: rapid antigen diagnostic test to detect dengue virus non-structural protein; PCR: polymerase chain reaction; RDT: rapid diagnostic test.

^a Only the minimum criteria required for fulfilling a clinical definition of dengue are included here; any additional signs and symptoms required for more severe forms are not listed.

^b Data sourced from WHO internal communications.

^c In the WHO 2009 dengue classification system, a probable case is any case with fever and two or more of the following: nausea, vomiting, rash, aches and pains, positive tourniquet test, leucopoenia or any warning sign. A case with warning signs is defined as a clinically diagnosed case if they have any of the following: abdominal pain or tenderness, persistent vomiting, clinical fluid accumulation, mucosal bleeding, lethargy, restlessness, liver enlargement >2 cm, or increase in haematocrit concurrent with rapid decrease in platelet count. Severe dengue is defined as severe plasma leakage leading to any of the following: shock, fluid accumulation with respiratory distress OR severe bleeding as evaluated by clinician OR severe organ involvement of the liver (i.e. aspartate amino transferase or alanine amino transferase \geq 1000 units/L), central nervous system (i.e. impaired consciousness), heart or other organs.

Fig. 1. Number of dengue cases and case-fatality ratios reported to WHO from the Western Pacific Region, 2013–2019



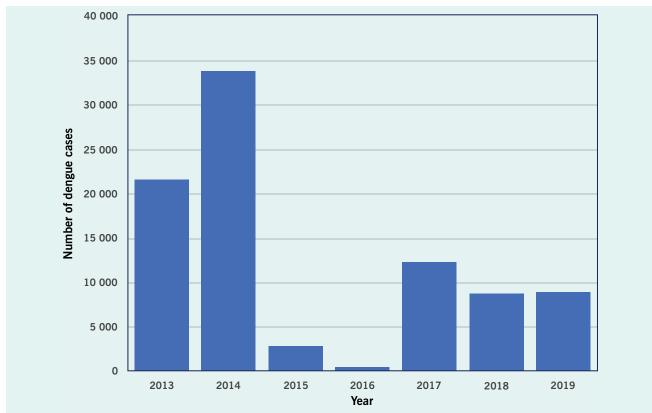


Fig. 2. Number of dengue cases reported to WHO from Pacific Island countries and areas, Western Pacific Region, 2013–2019

The data included in this figure are a subset of the data presented in Fig. 1.

were reported in the Region during the review period. Concurrent infections with two serotypes were reported in some countries. While some countries reported the same predominant serotype from 2016 to 2018, others reported changes in the predominant serotype. Additionally, there were reports of the introduction of a new serotype or switch in the predominant serotype, which was subsequently followed by outbreaks. Rare cases of autochthonous transmission were reported in countries where most previously reported cases had been imported.

Laboratory sampling schemes and confirmation methods varied by country and area. Some countries in this report were using the 2009 WHO dengue case classification system:¹⁴ (i) dengue without warning signs; (ii) dengue with warning signs that include abdominal pain or tenderness, persistent vomiting, clinical fluid accumulation, mucosal bleeding, lethargy or restlessness,

liver enlargement and increase in haematocrit with a rapid decrease in platelet count; and (iii) severe dengue, which is characterized by severe plasma leakage, severe haemorrhage and severe organ impairment. Other countries used other case definitions (**Table 1**). Some countries and areas in the Region report all identified cases of dengue, whereas others report only dengue cases at sentinel sites. In addition, some countries and areas conduct active surveillance or vector surveillance, or both (**Table 1**).

Reporting by country and area

Data for dengue cases were available from 35 countries and areas during this study period, including eight with complete case and death data for all years of this study (**Table 3**). Data were not available for three countries and areas: the Northern Mariana Islands, the Pitcairn Islands and Tokelau.

Table 2. Crude regional case notification rates for dengue reported to WHO from the Western Pacific Region, 2013–2019

Year	Number of cases reported in the Region	Population in the Region ^a	Case notification / 100 000 population per year ^b	95% confidence interval
2013	430 023	1 889 727 401	22.76	22.69–22.82
2014	376 972	1 901 609 413	19.82	19.76-19.89
2015	513 574	1 913 189 733	26.84	26.77–26.92
2016	458 843	1 924 437 124	23.84	23.77-23.91
2017	454 231	1 935 317 876	23.47	23.40-23.54
2018	479 263	1 945 715 729	24.63	24.56-24.70
2019	1 050 285	1 955 495 216	53.71	53.61-53.81

^a Population data were extracted from United Nations population estimates.¹²

^b Crude notification rates in the Region should be interpreted with caution, considering that the risks of disease and population sizes vary substantially across the Region, as well as the surveillance systems used to determine cases of dengue.

Asia subregion

Brunei Darussalam

Brunei Darussalam reported to WHO 2025 cases in 2013 and 436 cases and 2 deaths (CFR: 0.46%) in 2014. Reports for other years were not available.

Cambodia

During 2013–2019, Cambodia annually reported from 6372 to 68 597 suspected cases and from 3 to 59 deaths. The highest number of cases was reported during an outbreak in 2019 that peaked between June and August, with more than 5000 cases reported in epidemiological week 26.^{15,16} The highest number of deaths (59) was reported in 2013 (CFR: 0.34%).

In Cambodia, serotyping was conducted from sentinel laboratory surveillance at five sentinel sites. The predominant serotype reported from 2013 to 2015 was DENV-1, and in 2016, it switched to DENV-2. From the end of 2017 to the end of 2019, the predominant serotype switched back to DENV-1. This latter serotype switch preceded the large-scale outbreak in 2019, during which 73% (details on numerators and denominators are not available) of all serotyped samples between January and July 2019 were DENV-1, and the next most common serotype was DENV-2 (25%), followed by DENV-4 (2.2%) and DENV-3 (0.3%).

China

During 2013–2019, China annually reported from 2050 to 46 864 cases (including both clinically and laboratory-confirmed cases) and from 0 to 6 deaths. The highest number of cases and deaths were reported in 2014, with 46 864 cases and 6 deaths (CFR: 0.01%).

Several outbreaks were reported from the southern and central regions of China. Yunnan Province in 2013 reported 1245 cases with 136 that were laboratoryconfirmed, no deaths, and a predominant serotype of DENV-3;¹⁷ Henan Province in 2013 reported 106 suspected cases, 73 confirmed cases and no deaths, with the predominant serotype being DENV-3;¹⁸ Guangdong Province in 2014 accounted for more than 40 000 cases, including 1942 cases that were laboratory-confirmed and hospitalized and 2 deaths, where the predominant serotype among cases was DENV-1.⁶

The introduction of a new serotype in China in 2017 caused an outbreak of 1138 autochthonous cases after multiple clades of DENV-2 were introduced to Hangzhou, Zhejiang Province, in a short period.¹⁹ During 2013–2019, Hong Kong Special Administrative Region, China, annually reported between 102 and 163 cases. During 2013–2019, Taiwan, China, annually reported between 10 and 43 467 cases, with the highest number of cases reported in 2015. During 2013–2018,

Table 3.

Year 2013 2014 2015 2016 2017 Country or area No. of CFR (%) CFR (%) CFR (%) CFR (%) CFR (%) cases deaths cases deaths cases deaths cases deaths cases deaths Asia subregion 2025 436 0.46 Brunei Darussalam _ 2 _ _ _ _ _ _ _ _ _ _ 12 843 6372 Cambodia 17 533 59 0.34 3684 21 0.57 15 412 3 0.05 24 China 4663 0 0.00 46 864 6 0.01 3858 0 0.00 2050 0 0.00 5893 2 0.03 5 China, Hong Kong Special Administrative Region 103 0 114 124 102 0 0.00 112 0.00 0 0.00 9 China, Macao Special Administrative Region 17 0 0.00 3 0 0.00 11 17 _ _ _ _ _ _ 596 381 10 China, Taiwan 15 509 0 0.00 43 467 0 0.00 Japan 249 0 0.00 341 0 0.00 293 0 0.00 338 1 0.30 245 0 0.00 Lao People's Democratic Republic 44 250 95 1716 0 1959 10 0.21 0.00 0 0.00 5618 0.18 11 067 14 0.13 6 Malaysia 43 346 92 0.21 108 698 215 0.20 120 836 336 0.28 109 037 237 0.22 89 487 177 0.20 81 0 Mongolia 0 NA 0 0 NA 0 0 NA Philippines 204 906 660 0.32 113 485 425 0.37 213 930 647 0.30 220 518 1092 0.50 152 224 811 0.53 216 Republic of Korea 251 0 0.00 164 0 0.00 259 0 0.00 313 0 0.00 171 0 0.00 Singapore 22 170 8 0.04 18 326 6 0.03 11 294 6 0.05 13 085 12 0.09 2767 2 0.07 3 Viet Nam 66 322 42 0.06 31 848 30 0.09 97 484 62 0.06 91 609 28 0.03 172 232 40 0.02 131 Total for subregion 406 423 956 0.24 341 200 705 0.21 508 909 1051 0.21 455 927 1380 0.30 440 587 1049 0.24 469 Pacific subregion American Samoa 479 4 0.84 0 0 0.00 _ _ _ _ _ _ _ _ _ Australia 1841 0 0.00 1721 0 0.00 1716 0 0.00 2237 0 0.00 1132 1 0.09 Cook Islands 946 0 0.00 765 0 0.00 0 0 _ _ Fiji 352 0 0.00 16 0.06 398 0 0.00 2699 9 0.33 26 595 4 1523 2155 0 0.00 French Polynesia 0 0.00 _ _ _ _ _ _ _ Guam Kiribati 0 0 _ _ _ _ _ _ _ _ _ _ Marshall Islands 217 0 0.00 0.00 0.00 90 0.00 0 Micronesia (Federated States of) 14 0 1 0 0 _ _ Nauru 251 0 964 3 0.31 New Caledonia 9958 0.04 4200 11 4 0.26 1 _ _ _ _ _ _ _ _ _ 179 125 191 New Zealand 106 0 0.00 0 0.00 0.00 0.00 161 0.00 0 0 0 Niue 0 2 _ _ _ _ _ _ _ _ _ _ Northern Mariana Islands 9 0.00 13 2 20 440 Palau 0 15.38 0 0.00 5 1.14 _ _ _ Papua New Guinea 6 Pitcairn Islands _ _ _ _ _ _ _ _ _ _ _ _ _ Samoa 2724 5 0.18 9500 1872 Solomon Islands 8 0.08 1 0.05 _ _ _ _ _ _ _ Tokelau _ _ _ 51 0 0.00 1559 0.00 100 0 0.00 Tonga 0 _ _ _ _ _ _ Tuvalu 408 _ Vanuatu 1561 >1000 _ _ _ _ _ _ _ _ _ _ _ _ _ Wallis and Futuna 94 222 0.00 0 _ _ 23 600 12 35 772 4665 2916 0.00 13 644 0.23 Total for subregion 0.05 19 0.05 0.09 34 0 4

Number of dengue cases (including imported cases), number of dengue-attributed deaths and case-fatality ratios reported to WHO from the Western Pacific Region, 2013–2019^a

CFR: case-fatality ratio; NA: cannot be calculated

430 023

968

0.23

376 972

724

0.19

513 574

1055

0.21

458 843

1380

0.30

454 231

1083

0.24

47

^a The – symbol indicates that no data were available.

TOTAL

	2018		2019			
lo. of ases	No. of deaths	CFR (%)	No. of cases	No. of deaths	CFR (%)	
-	-	-	-	-	-	
1 684	23	0.09	68 597	48	0.07	
136	1	0.02	22 188	3	0.01	
163	-	-	198	-	-	
18	-	-	-	-	-	
183	-	-	100	-	-	
201	0	0.00	463	0	0.00	
446	19	0.29	39 091	76	0.19	
l 360	147	0.18	130 101	147	0.11	
_	-	-	-	-	_	
6 190	1083	0.50	437 563	1689	0.39	
159	0	0.00	273	0	0.00	
283	6	0.18	15 999	3	0.02	
1 447	27	0.02	320 702	54	0.02	
9 270	1306	0.28	1 035 275	2020	0.20	
-	-	-	-	-	_	
917	0	0.00	1463	1	0.07	
0	-	-	126	0	0.00	
000	9	0.23	2500	0	0.00	
_	-	-	2400	0	0.00	
-	-	-	23	-	_	
899	2	0.11	-	-	_	
-	-	_	1635	1	0.06	
0	-	-	1464	1	0.07	
114	0	0.00	-	-	_	
997	0	0.00	3916	2	0.05	
294	0	0.00	224	0	0.00	
_	-	_	-	_	_	
_	_	_	-	_	_	
570	2	0.35	737	3	0.41	
-	_	_	_	_	_	
_	_	_	_	-	_	
	_	_	-	-	_	
-	_	_	-	-	_	
_	_	-	_	-	_	
_	_	_	_	_	_	
_	_	_	522	2	0.38	
	_	_	522	2	0.00	
202	-	-	_	-	_	
993	13	0.13	15 010	10	0.05	
	13	0.13	1.0.010	10	0.05	

Macao Special Administrative Region, China, annually \mathcal{N} reported between 3 and 18 cases.

Japan

During 2013–2019, Japan annually reported between 201 and 463 laboratory-confirmed cases, with 1 death reported in 2016. In 2014, an outbreak of 162 autoch-thonous dengue cases was reported for the first time in nearly 70 years, of which more than 90% (148/160, from available data) had visited or worked near parks in central Tokyo, and the dominant serotype was DENV-1.^{5,20,21}

All cases reported from 2016 to 2018 were imported. The predominant serotype was DENV-2 (36% [61/172] of cases in 2016, 35% [39/113] in 2017, 42% [34/81] in 2018), followed by DENV-3 (23% [39/172] of cases in 2016, 27% [31/113] in 2017, 31% [25/81] in 2018), DENV-1 (31% [54/172] of cases in 2016, 27% [31/113] in 2017, 24% [19/81] in 2018) and DENV-4 (11% [18/172] of cases in 2016, 11% [12/113] in 2017, 4% [3/81] in 2018). In 2019, 17% (78/463) of serotyped cases were DENV-1, 16% (74/463) were DENV-2, 9% (40/463) were DENV-3 and 3% (16/463) were DENV-4.

Lao People's Democratic Republic

During 2013–2019, the Lao People's Democratic Republic annually reported between 1716 and 44 250 clinically confirmed cases and 0 to 95 deaths. In 2013, the country reported the largest dengue outbreak in its history,²² with 44 250 cases and 95 deaths reported nationwide. In the southern part of the country alone, 4638 cases and 32 deaths were reported, among which DENV-2, DENV-3 and chikungunya virus were detected, as were concurrent infections with more than one serotype of DENV, or DENV and chikungunya virus.²³ More than 90% (numerator not available) of 537 samples serotyped in 2013 were DENV-3.24 In 2015, an outbreak was reported as predominantly due to DENV-1.24 In 2019, there was a dengue outbreak with 39 091 cases reported and 76 deaths (CFR: 0.19%), and 65% (numerator not available) of 1178 specimens collected and serotyped were found to be DENV-2.24 The predominant serotypes during outbreaks in 2013, 2015 and 2019 were attributed to three different serotypes, indicating two serotype switches.²⁴

Malaysia

During 2013–2019, Malaysia annually reported between 43 346 and 130 101 laboratory-confirmed cases and 92 to 336 deaths. No imported cases were reported from 2016 to 2018. Malaysia launched the web-based e-Notification system and e-Dengue system in 2014, and all registered dengue cases since January 2014 have been laboratory-confirmed. More than 100 000 cases were reported in 2014, 2015, 2016 and 2019.

All four serotypes were reported in Malaysia, with the predominant serotype differing each year from 2016 to 2018, with significant cocirculation. In 2016, the predominant serotype was DENV-1 (40%, 2211/5482), followed by DENV-3 (32%, 1745/5482), DENV-2 (25%, 1381/5482) and DENV-4 (3%, 145/5482). In 2017, the predominant serotype was DENV-3 (41%, 2200/5420), followed by DENV-2 (35%, 1887/5420), DENV-1 (23%, 1245/5420) and DENV-4 (2%, 88/5420). In 2018, the predominant serotype was DENV-2 (47%, 2608/5544), followed by DENV-3 (33%, 1833/5544), DENV-1 (19%, 1055/5544) and DENV-4 (1%, 48/5544).

Mongolia

During 2013–2015, Mongolia reported no dengue cases and no deaths. Data for 2016–2019 were not available.

The Philippines

During 2013–2019, the Philippines annually reported between 113 485 and 437 563 suspected dengue cases and 425 to 1689 deaths. Among these suspected cases, 1488 cases in 2016, 1333 cases in 2017 and 998 cases in 2018 were laboratory-confirmed. The highest number of cases and deaths were reported during a large-scale outbreak in 2019, with 437 563 cases and 1689 deaths (CFR: 0.39%).

All four serotypes were reported from the Philippines. In 2016, the predominant serotype among 1488 cases tested was DENV-1 (44%, 659/1488), followed by DENV-3 (26%, 384/1488), DENV-2 (24%, 349/1488) and DENV-4 (6%, 95/1488); 1 case tested positive for both DENV-1 and DENV-2 (0.1%, 1/1488). In 2017, the predominant serotype among 1333 cases tested was DENV-3 (60%, 795/1333), followed by DENV-1 (24%, 318/1333), DENV-2 (12%, 164/1333) and DENV-4 (4%, 47/1333); 2 cases tested positive for DENV-1 and DENV-2 (0.2%, 2/1333), 5 cases tested positive for DENV-1 and DENV-3 (0.4%, 5/1333) and 2 cases tested positive for DENV-2 and DENV-3 (0.2%, 2/1333). In 2018, the predominant serotype among 998 cases tested was DENV-3 (60%, 598/998), followed by DENV-1 (22%, 223/998), DENV-2 (15%, 149/998) and DENV-4 (3%, 25/988); 2 cases tested positive for DENV-1 and DENV-3 (0.2%, 2/988) and 1 case tested positive for DENV-2 and DENV-3 (0.1%, 1/988). In 2019, the predominant serotype among the 100 cases with serotype data available was DENV-3 (64%), followed by DENV-2 (18%), DENV-1 (15%) and DENV-4 (3%).²⁵

Republic of Korea

During 2013–2019, the Republic of Korea annually reported between 164 and 313 laboratory-confirmed cases and no deaths. The highest number of cases was reported in 2016. All cases reported from 2016 to 2018 were imported, comprising all four serotypes. In 2016, the predominant serotype was DENV-1 (38%, 57/149), followed by DENV-2 (35%, 52/149), DENV-3 (20%, 30/149) and DENV-4 (7%, 10/149). In 2017, the predominant serotype among imported cases was DENV-1 (44%, 38/86), followed by DENV-3 (23%, 20/86), DENV-2 (19%, 16/86) and DENV-4 (14%, 12/86). In 2018, the predominant serotype among imported cases was DENV-2 (37%, 35/96), followed by DENV-1 (33%, 32/96), DENV-3 (28%, 27/96) and DENV-4 (2%, 2/96).

Singapore

During 2013–2019, Singapore annually reported between 2767 and 22 170 laboratory-confirmed cases and 2 to 12 deaths. Large numbers of cases were reported during outbreaks in 2013, 2014, 2015 and 2019. The numbers of reported cases were low in 2017 and 2018. Among the 20 deaths reported during 2016-2018, 14 were autochthonous cases and the rest were imported cases. All four serotypes were reported from Singapore; however, denominators were not available, so the percentage for each serotype is reported along with the number of positive cases. The predominant serotypes from 2016 to 2018 were DENV-2 (51% [2257 positive cases] in 2016, 45% [361] in 2017 and 52% [637] in 2018), followed by in 2016 DENV-1 (29%, 278 positive cases) then DENV-3 (18%, 806), and in 2017 DENV-3 (24%, 192) then DENV-1 (21%, 171), and in 2018 DENV-3 (25%, 305)

and DENV-1 (20%, 240). DENV-4 was reported from 2% (n = 67) of cases in 2016, 10% (77) in 2017 and 4% (47) in 2018.

Viet Nam

During 2013–2019, Viet Nam annually reported between 66 322 and 320 702 cases (including both clinically and laboratory-confirmed cases) and 27 to 62 deaths. More than 100 000 cases were reported in 2017, 2018 and 2019; notably, 320 702 cases were reported in 2019. During the outbreak in 2017, more than 59 000 cases were reported in northern Viet Nam, eight times higher than the number of cases in 2016.²⁶

All four serotypes were reported from Viet Nam during 2016–2018. In 2016, the predominant serotype was DENV-1 (61%, 1104/1803), followed by DENV-4 (25%, 453/1803), DENV-2 (13%, 240/1803) and DENV-3 (0.3%, 6/1803). In 2017, the predominant serotype was DENV-1 (72%, 2057/2870), followed by DENV-2 (21%, 607/2870), DENV-4 (7%, 204/2870) and DENV-3 (0.1%, 2/2870). In 2018, the predominant serotype changed to DENV-2 (50%, 988/1980), followed by DENV-1 (33%, 661/1980), DENV-4 (17%, 328/1980) and DENV-3 (0.2%, 3/1980).

Pacific subregion

Australia

During 2013–2019, Australia annually reported between 917 and 2237 laboratory-confirmed cases and 0 to 1 death. More than 1700 cases were reported annually in 2013, 2014, 2015 and 2016; in 2016, 2237 cases were reported. During 2016–2018, more than 98% of reported cases were imported (2204/2237 in 2016, 1113/1132 in 2017 and 907/917 in 2018). In Australia, dengue cases occur each year in North Queensland, generally originating from an imported case, although in 2019 an outbreak associated with 13 locally acquired cases was reported for the first time in decades in the Rockhampton region, Queensland.^{27,28}

All four serotypes were reported from Australia, with the predominant serotype being DENV-2 (44% [468/1052 of known and serotyped cases] in 2016, 56% [246/436] in 2017, 43% [120/282] in 2018), followed in 2016 by DENV-3 (24%, 257/1052), DENV-1 (19%,

202/1052) and DENV-4 (12%, 125/1052); in 2017 by DENV-1 (20%, 88/436), DENV-3 (13%, 57/436) and DENV-4 (10%, 45/436); and in 2018 by DENV-1 (30%, 86/282), DENV-3 (20%, 55/282) and DENV-4 (7%, 21/282). In addition to these serotyped cases, concurrent infection with two serotypes was reported in 2016 and 2017. In 2016, concurrent infections were reported with DENV-1 and DENV-2 (1 case), DENV-2 and DENV-3 (1 case), and DENV-3 and DENV-4 (4 cases); in 2017, concurrent infection with DENV-1 and DENV-4 was reported in 1 case; in 2019, concurrent infection with DENV-3 and DENV-4 was reported in 1 case.

New Zealand

During 2013–2019, New Zealand annually reported between 106 and 294 cases (including both clinically confirmed and laboratory-confirmed cases, although most are laboratory-confirmed); during 2016–2019, no deaths were reported. Among reported cases, 98% (158/161) were laboratory-confirmed in 2017, 95% (280/294) in 2018 and 98% (219/224) in 2019.^{29–31} The largest number of cases was reported in 2018, at 294 cases. In 2016, two dengue fever outbreaks were reported that involved 12 cases. During 2013–2019, all cases reported in New Zealand were imported (information on travel history was not available for 1 case in 2015 and 2 cases in 2019).

All four serotypes were reported from New Zealand. In 2016, the predominant serotype was DENV-3 (63%, 59/93), followed by DENV-2 (20%, 19/93), DENV-1 (11%, 10/93) and DENV-4 (5%, 5/93). In 2017 and 2018, the predominant serotype was DENV-2 (83% [82/99] and 84% [167/200], respectively), followed by DENV-1 (10% [10/99] and 9% [18/200], respectively), DENV-3 (6% [6/99] and 5% [9/200], respectively) and DENV-4 (1% [1/99] and 3% [6/200], respectively).

American Samoa

American Samoa reported clinically confirmed cases to WHO using the 2009 WHO dengue case classification system. Laboratory confirmation is conducted to confirm outbreaks using reverse transcription polymerase chain reaction (RT-PCR) or an antigen rapid diagnostic test (NS1). In 2015, American Samoa reported 479 cases and 4 deaths (CFR: 0.84%). Outbreaks were also reported in 2017 and 2018, but the total numbers of cases are not available.

Cook Islands

The Cook Islands reported clinically confirmed cases to WHO using the 2009 WHO dengue case classification system. In 2014, the Cook Islands reported 946 cases and no deaths, and in 2015 the Islands reported 765 cases and no deaths (CFR: 0%). No cases were reported to WHO during 2016–2018. In 2019, a dengue outbreak was declared in February, with 41 confirmed cases and 85 probable, 48 hospitalizations and no deaths.³² The predominant serotype in 2019 was DENV-1, accounting for 93% (35/38) of cases with available serotype information. Additionally, 3 cases who were tourists with a history of travelling to French Polynesia were confirmed with DENV-2 in October 2019.³³

Fiji

During 2013–2018, Fiji annually reported between 352 and 26 595 cases and 0 to 16 deaths. Fiji reported clinically confirmed cases to WHO using the 2009 WHO dengue case classification system. Samples from different health divisions were tested using RT-PCR, an antigen rapid diagnostic test (NS1) and an enzymelinked immunosorbent assay (ELISA). An outbreak was reported in 2014 of at least 26 595 cases (more than 27 000 reported according to some sources) and 16 deaths (CFR: 0.06%). From 2017 to 2018, an outbreak was reported, for which the predominant serotype was DENV-2.³⁴

French Polynesia

French Polynesia used the 2009 WHO dengue case classification system, and the laboratory method for confirmation was RT-PCR. In 2013, French Polynesia reported 1523 dengue cases associated with an outbreak, with 258 being laboratory-confirmed; during the outbreak, 70% (170/244) of cases with the serotype identified had DENV-1 infections, 30% (73/244) had DENV-3 infections (genotype III) and 0.4% (1/244) had coinfection with both serotypes.³⁵ DENV-3 was reported to have been introduced from South America.35 In 2014, 2155 confirmed and 34 000 suspected cases were reported in French Polynesia, and outbreaks were also reported in 2016 and 2017. In 2016 and 2017, DENV-1 was reported, and in 2018, DENV-2 was reported. In April 2019, an outbreak of DENV-2 was declared, with 2400 autochthonous cases reported.³⁶

Guam

Guam reported clinically confirmed cases to WHO: 23 cases were reported in 2019, with no further information available.

Kiribati

Kiribati reported clinically confirmed cases to WHO using the 2009 WHO dengue case classification system. Laboratory testing to confirm outbreaks is conducted using RT-PCR or an antigen rapid diagnostic test (NS1). In Kiribati, outbreaks were reported in 2013 and 2014, and no cases were reported in 2016 and 2017. In 2018, 1899 cases and 2 deaths were reported, with DENV-2 detected.

Republic of the Marshall Islands

In the Republic of the Marshall Islands, outbreaks were reported in 2013 and 2014. In 2019, a DENV-3 outbreak was reported with at least 1395 cases of dengue-like illness, including 431 laboratory-confirmed cases and 1 death.³⁷ A health emergency was declared in relation to this event; internal movement restrictions were imposed between the affected and unaffected islands; and emergency medical teams were deployed to support the dengue response.

Federated States of Micronesia

The Federated States of Micronesia reported clinically confirmed cases to WHO using the 2009 WHO dengue case classification system. Laboratory methods used to confirm outbreaks include RT-PCR and an antigen rapid diagnostic test (NS1). There were 217 cases reported to WHO in 2013, associated with an outbreak of 729 suspected dengue cases and no deaths in Kosrae from September 2012 to March 2013. DENV-4 was detected from 3 specimens collected during this period; 11% (728/6600) of Kosrae residents met the case definition for suspected dengue, and almost 4% (242/6600) were hospitalized.³⁸ In 2018, DENV-4 was reported. In 2019, 1464 dengue cases including 1 death were reported from Yap state, and the predominant serotype was DENV-3. The dengue outbreak in 2019 coincided with a concurrent leptospirosis outbreak in Yap state, and an executive order determining a public health crisis was issued.

Nauru

Nauru reported clinically confirmed cases to WHO using the 2009 WHO dengue case classification system. Laboratory testing to confirm outbreaks uses RT-PCR or an antigen rapid diagnostic test (NS1). Nauru reported 251 cases in 2014, no cases in 2016, 964 cases and 3 deaths in 2017 and 114 cases and no deaths in 2018. In 2017, DENV-2 was reported and in 2018, DENV-1 was reported.

New Caledonia

New Caledonia reported cases to WHO using the 2009 WHO dengue case classification system and RT-PCR for laboratory confirmation. In 2013, New Caledonia reported 9958 cases including 4 deaths during an outbreak in which the predominant serotype was DENV-1.39 Based on available information, an outbreak was also reported in 2014. In 2017, 4200 cases and 11 deaths were reported, with DENV-1, DENV-2 and DENV-3 detected. From November 2018 to September 2019, a dengue outbreak was declared. From 1 January to 31 December 2019, 3916 cases, 368 hospitalizations and 2 deaths were reported. Among the 316 cases with serotype information available, the predominant serotype was DENV-2. Two cases of DENV-1 and 1 case of DENV-4 were imported from French Polynesia and Indonesia, respectively.40

Niue

Niue reported clinically confirmed cases to WHO. In Niue, 2 cases were reported in 2017. In 2018, DENV-2 was reported, but information on the number of cases was not available.

Palau

Palau reported cases to WHO using the 2009 WHO dengue case classification system and RT-PCR or an antigen rapid diagnostic test (NS1) for laboratory testing to confirm outbreaks. During 2013–2017, Palau annually reported between 9 and 737 cases and 0 to 5 deaths. Outbreaks were reported in 2016 and again in 2017, the latter comprising 440 cases and 5 deaths, with a predominant serotype of DENV-2. In 2018, 570 cases and 2 deaths were reported, and in 2019, there were

737 cases including 3 deaths. From December 2018 to September 2019, 160 cases were confirmed as DENV-3. Two serotypes were reported from Palau; DENV-2 was reported in 2016 and 2017, and DENV-3 was reported in 2018.

Papua New Guinea

In 2014, Papua New Guinea reported 6 cases. Further information was not available.

Samoa

Samoa reported cases to WHO using the 2009 WHO dengue case classification system and RT-PCR or an antigen rapid diagnostic test (NS1) to confirm outbreaks. In Samoa, outbreaks were reported in 2015 and 2016. In 2017, 2724 cases and 5 deaths were reported, with the predominant serotype being DENV-3. In 2018, DENV-2 was reported.

Solomon Islands

The Solomon Islands reported cases to WHO using the 2009 WHO dengue case classification system. In the Solomon Islands in 2013, 9500 cases and 8 deaths (CFR: 0.10%) were associated with an outbreak in Honiara. DENV-3 genotype I was isolated from specimens collected during this outbreak, suggesting introduction from south-east Asia after 18 years of dengue absence in the PICs.³⁵ In 2014, 1872 cases and 1 death (CFR: 0.05%) were reported. The introduction of DENV-2 to the Solomon Islands resulted in outbreaks in 2016 and 2017.41,42 From September 2016 to April 2017, an outbreak of DENV-2 was reported in 9 of 10 provinces in the Solomon Islands, with 12 329 suspected cases, including 1510 cases positive by dengue rapid diagnostic test, and 16 deaths.⁴² An outbreak was also reported in 2019.

Tonga

Tonga reported cases to WHO using the 2009 WHO dengue case classification system. In Tonga, 51 cases and no deaths were reported in 2014; 1559 cases and no deaths were reported in 2015; and more than 100 cases were reported in 2017.

Tuvalu

Tuvalu reported cases to WHO using the 2009 WHO dengue case classification system. In Tuvalu, 408 cases were reported in 2014. In March 2019, a dengue outbreak was declared. In 2019, 522 cases were reported, including at least 21 hospitalizations and 2 deaths in children. The predominant serotype in the 2019 outbreak was DENV-1.

Vanuatu

Vanuatu reported clinically confirmed cases to WHO. In Vanuatu, 1561 cases were reported in 2014 and more than 1000 cases were reported in 2017; DENV-2 was reported in 2018.

Wallis and Futuna

Wallis and Futuna reported cases to WHO using the 2009 WHO dengue case classification system. In Wallis and Futuna, 94 cases were reported in 2013. In 2017, an outbreak was declared in November, with 222 cases and no deaths, and DENV-1 was identified from 2 samples. In 2018, 202 cases and DENV-1 were reported. In November 2019, an outbreak was declared in Wallis and Futuna, and 30 confirmed cases were reported from February to December 2019, with the predominant serotype being DENV-2.⁴³

DISCUSSION

Dengue continued to pose a health burden in the Region during 2013–2019, with the number of annually reported cases ranging from a little more than 430 000 to more than 1 million and with the annual number of reported deaths ranging from 724 to 2025. Outbreaks were reported from the Region every year during the study period. The introduction or reintroduction of serotypes to specific areas caused several outbreaks and rare occurrences of local transmission in places where dengue had not been previously reported. With support from countries and areas, WHO continued to share timely information during the study period through its biweekly dengue epidemiological reports for the Region¹¹ and conducted regional and country-specific risk assessments to inform dengue prevention and control efforts.

The increases in reported cases and regional case incidence may be attributed to several factors. First, a true increase in dengue incidence may have occurred due to expanding urbanization and increasing population size and density, particularly in settings with increased exposure to competent dengue vectors and mosquito breeding grounds.44 Shifts in ecological factors due to climate change, such as intensified rainy seasons and higher ambient temperatures, have expanded the geographical range of Aedes mosquitos globally during the past 50 years and led to intensified dengue transmission.⁴⁵ Second, increased international travel and trade have led to the importation of cases with different serotypes and the introduction of mosquito eggs through the importation of goods to areas where the population is susceptible and competent mosquitos exist.^{44,45} Third, reports to national health authorities likely increased due to strengthened surveillance systems and diagnostic capacities, including laboratory networks that supported confirmatory diagnosis in the PICs, as well as an emphasis on risk communication activities to improve the awareness of dengue among the public.³ The range of CFRs may be associated with differences in case reporting, the timing of the case presentation to health-care facilities and clinical management.

The number of cases reported in 2019 was higher than in the years from 2013 to 2018, and the CFR was relatively low. This increase in 2019 included at least 14 countries and areas that reported dengue outbreaks in the Region, including large-scale outbreaks; during 2019, four countries and areas in the Asia subregion and three in the Pacific subregion reported their highest number of cases of the 7-year study. It is possible that case detection and reporting increased due to improved awareness of dengue among health-care professionals and the public because of the large outbreaks. These outbreaks may have also increased health-care-seeking behaviour, leading to fewer deaths, thereby decreasing the CFR.

Our findings show that there is a substantial burden of dengue in the Region and that it continues to increase over time. However, dengue surveillance practices throughout the Region are inconsistent and require strengthening. To inform national and regional risk assessments and actions, information is required not only on the time, place and demographics of a case, such as age and sex, but also on the DENV serotype and whether the infection was locally acquired or imported. These details will also support risk assessments for and responses to events with new epidemiological patterns, such as outbreaks associated with the introduction or reintroduction of serotypes to specific areas, as well as rare occurrences of local transmission in places where it was not previously reported. Furthermore, in some settings, the capacities for surveillance, outbreak response, clinical management and diagnosis may be limited. Several approaches could fill these gaps, including strengthening laboratory capacity and laboratory networks, institutionalizing active surveillance to detect dengue cases who are self-managed and inapparent, and implementing integrated vector surveillance.

Although several countries and areas have adopted the 2009 WHO dengue case classification system,¹⁴ there are differences in countries and areas across the Region in surveillance methodology, including whether universal or sentinel reporting is used; laboratory sampling schemes and confirmation methods; and reporting practices. These differences are a limitation of this report, indicating why comparison across countries should be avoided and comparisons within one country should be informed by the local reporting practices, which may change over time. As a result of differences in case definitions and other factors, there is likely to be underreporting and, thus, an underestimation of the true regional burden in terms of the number of cases, CFRs and incidence.^{1,2} Despite these limitations, continued reporting of dengue in line with the Regional Action Plan is important to guide public health authorities in their national and subnational response efforts.

The burden of dengue, including the increased risks of dengue outbreaks, will continue amid other public health emergencies. Disaggregating data by age and sex at all levels will enable public health authorities to implement improved and targeted response measures. Additional information about cases, including their travel history and serotype, should also be routinely collected and reported. The Region's capacity to mitigate the impact of dengue can be strengthened by making a shift in its management, from a reactive, acute outbreak response to one that reduces fatalities through undertaking activities, including sustainable implementation of mosquito control measures, engaging communities to raise their awareness about the risk of dengue and to communicate relevant behavioural changes, and strengthening diagnostics and case management. Enhancing collaboration and coordination within and beyond the health sector is key to carrying out these activities successfully.

Acknowledgements

We thank the WHO Member States in the Western Pacific Region for collecting and sharing data, as well as the WHO Country Office staff in the Region for supporting surveillance activities for dengue. We also thank the WHO Regional Health Emergencies Programme Country Office focal points, and the Regional Division for Disease Control focal points for their contributions to this report.

Conflicts of interest

MC, LB, SO, CKL, RRA, SC and TM are associate editors and BO is executive editor of the Western Pacific Surveillance and Response journal. They were not involved in the editorial decision to publish the manuscript. All other authors have no conflicts of interest to declare.

Ethics statement

Ethical clearance was not required because this report used routinely available data and no personal identifying information was collected.

Funding

None.

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