Melioidosis at Royal Darwin Hospital in the big 2009–2010 wet season: comparison with the preceding 20 years

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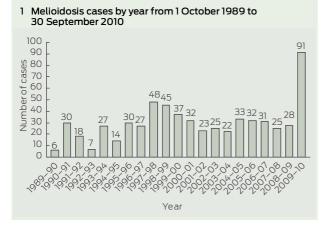
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MJA 2012; 196: 345-348 doi: 10.5694/mja11.11170 elioidosis is an infection caused by the soil and water bacterium *Burkholderia* pseudomallei, and was first documented in Australia in 1949 in Winton, Queensland, where sheep were infected.¹ It was first reported in the Northern Territory in 1960.² Since then, melioidosis has been increasingly recognised as an important cause of sepsis in northern Australia,³⁻⁶ with the vast majority of cases occurring during the monsoonal rainy season during October to April.

The Darwin Prospective Melioidosis Study commenced on 1 October 1989.7 Over the 20 years until 30 September 2009, 540 culture-confirmed cases of melioidosis were documented from the Top End of the NT.6 This represented a median of 27 (range, 6-48) cases per year. From 1 October 2009 until 30 September 2010, an unprecedented 91 patients with culture-confirmed cases of melioidosis were managed by Royal Darwin Hospital (RDH). In this study, we compare these 91 cases from that most recent 12-month period with the 540 cases from the preceding 20 years.

Methods

Clinical focus p 322 Case report p 350 The Top End of the NT has an estimated population of 177 500 people, with 124 500 living in Darwin.⁸ Indi-



Abstract

Objective: To compare the unprecedented 91 cases of melioidosis in the Top End of the Northern Territory of Australia from 1 October 2009 to 30 September 2010 with the 540 cases in the preceding 20 years and postulate reasons for this year of very high melioidosis incidence.

Design, setting and participants: Review of prospectively collected data on all patients with culture-confirmed melioidosis at Royal Darwin Hospital, the Top End's tertiary referral centre, since 1 October 1989.

Main outcome measures: Population-based annual incidence of melioidosis; differences in epidemiology, clinical presentations and outcomes for 2009–2010 compared with the preceding 20 years.

Results: In 2009–2010, the estimated population-based incidence of melioidosis was 50.2 cases per 100 000 in the Top End population overall, and 102.4 cases per 100 000 in the Top End Indigenous population. The proportion of patients acquiring melioidosis in the Darwin urban area increased from 49% in 1989–2009 to 65% in 2009–2010 (OR, 1.96; 95% CI, 1.20–3.19). Among the 49 Indigenous Australian patients with melioidosis in 2009–2010, 63% acquired the infection in Darwin, compared with 35% of Indigenous patients in the previous 20 years (OR, 3.17; 95% CI, 1.62–6.24).

Conclusions: In 2009–2010, the Top End had the highest annual incidence of melioidosis documented from anywhere to date. The prominent increase in cases in Darwin was associated with above average rainfall in Darwin during December 2009 to February 2010. The increase in the proportion of Indigenous Australians who acquired melioidosis in Darwin may reflect movement of some Indigenous people into Darwin from remote communities.

genous Australians account for 26% of the Top End population, and this proportion includes most of the people living in remote communities. All patients in the Top End with culture-confirmed melioidosis are managed by, or in consultation with, the infectious diseases department at RDH.

The Darwin Prospective Melioidosis Study records clinical and laboratory data for all patients with melioidosis, and defines risk factors and septic shock as has been described previously.^{6,7} We follow all patients until death or completion of treatment.

For the purposes of the study, the Darwin urban region includes the city limits of Darwin and Palmerston; and the Darwin rural region extends from outside Palmerston to include the rural blocks accessed along the Stuart and Arnhem highways as far as Adelaide River, 115 km south of Darwin.

Statistical analysis

Patient details were analysed using Stata, version 10 (StataCorp, College

Station, Tex, USA). Cases of melioidosis from 1 October 2009 until 30 September 2010 (12 months) were compared with cases from 1 October 1989 until 30 September 2009 (20 years). The χ^2 test was used to compare intergroup differences in categorical demographic and clinical parameters, with P < 0.05 considered significant. Odds ratios and 95% CIs were then calculated. We calculated population-based incidence rates for melioidosis in the Top End for 2009–2010 using the 2009 estimated population.

Ethics approval

This study was approved by the Human Research Ethics Committee of the NT Department of Health and Menzies School of Health Research (approval no. 02/38).

Results

Box 1 shows the yearly cases of melioidosis from 1 October 1989. The baseline characteristics, risk factors and clinical presentations of the 91

2 Demographic characteristics and location of residence for patients with melioidosis managed at Royal Darwin Hospital from 1 October 1989 to 30 September 2010

Demographic characteristics	1989–2009	2009-2010	Odds ratio (95% CI)*	P
All patients				
Number of cases	540	91		
Median number of cases/year	27	91		< 0.001
Median age (years)	49	53		
No. of children aged < 15 years	26 (5%)	3 (3%)		0.71
Male	372 (69%)	61 (67%)		0.82
Indigenous	281 (52%)	49 (54%)		0.80
Darwin urban [†]	262 (49%)	59 (65%)	1.96 (1.20–3.19)	0.006
Indigenous	99 (38% [‡])	31 (53% [‡])	1.82 (0.99–3.35)	0.05
Darwin rural [†]	65 (12%)	12 (13%)		0.89
Remote Indigenous communities	169 (31%)	15 (16%)	0.43 (0.23-0.80)	0.006
Other ⁶	44 (8%)	5 (5%)		0.51
Indigenous patients				
Number of cases	281	49		
Median age (years)	45	49		
Male	169 (60%)	28 (57%)		0.81
Darwin urban [†]	99 (35%)	31 (63%)	3.17 (1.62-6.24)	< 0.001
Darwin rural [†]	8 (3%)	0		0.49
Remote Indigenous communities	153 (54%)	14 (29%)	0.34(0.16-0.68)	0.001
Other [§]	21 (7%)	4 (8%)		1.00

^{*}Odds ratios and 95% confidence intervals were calculated, comparing the odds of the parameter occurring in melioidosis cases in 2009–2010 versus 1989–2009. † See Methods for definitions. ‡ Percentage of Darwin urban. § Including Katherine, Gove and central Australia.

3 Host risk factors and associated deaths for patients with melioidosis managed at Royal Darwin Hospital from 1 October 1989 to 30 September 2010

Risk factors	1989–2009	2009–2010	P	Deaths 1989–2009*	Deaths 2009–2010*
Total patients	540	91			
Diabetes	213 (39%)	44 (48%)	0.14	33 (15%)	4 (9%)
Hazardous alcohol intake	211 (39%)	40 (44%)	0.45	33 (16%)	7 (18%)
Chronic lung disease	140 (26%)	26 (29%)	0.69	27 (19%)	5 (19%)
Chronic renal disease	65 (12%)	7(8%)	0.30	13 (20%)	1 (14%)
Rheumatic heart disease and/or cardiac failure	39 (7%)	11 (12%)	0.17	9 (23%)	3 (27%)
Malignancy	31 (6%)	9 (10%)	0.20	8 (26%)	1 (11%)
Immunosuppression	31 (6%)	10 (11%)	0.10	6 (19%)	1 (10%)
No host risk factor	106 (20%)	16 (18%)	0.75	2 (2%)	0

^{*} Percentage with risk factor who died.

patients in 2009–2010 were similar to those of the 540 patients in 1989–2009 (Box 2, Box 3, Box 4). The proportion of patients acquiring melioidosis in the Darwin urban area increased significantly from 49% in 1989–2009 to 65% in 2009–2010 (OR, 1.96; 95% CI, 1.20–3.19). Of the patients who acquired melioidosis in the Darwin urban area, the proportion who were Indigenous Australians increased from 38% in 1989–2009 to 53% in 2009–2010 (OR, 1.82; 95% CI, 0.99–3.35). Of the 49 Indigenous Australian patients who acquired melioidosis in

2009–2010, 63% were in the Darwin urban area; this compares with 35% of all Indigenous patients with melioidosis in the previous 20 years (OR, 3.17; 95% CI, 1.62–6.24). Among non-Indigenous patients, there was no significant change, with 28 of 42 (67%) in 2009–2010 and 163 of 259 (63%) in 1989–2009 acquiring melioidosis in the Darwin urban area.

The population-based annual incidence of melioidosis in the Top End in 2009–2010 was 50.2 cases per 100 000, compared with a prior mean annual incidence of 19.6 per 100 000.9 The

annual incidence of melioidosis in the Top End Indigenous population in 2009–2010 was 102.4 cases per 100,000.

Box 5 shows some clinical examples of melioidosis. No patient without a recognised risk factor for melioidosis died from the disease during 2009-2010; 10 of the 11 patients who died had either diabetes or hazardous alcohol intake or both. Of the seven Indigenous patients who died from melioidosis in 2009-2010, five were initially from remote Indigenous communities but were living in the Darwin urban area when they acquired melioidosis (four of these were homeless); all five had hazardous alcohol intake. Eleven other Indigenous patients living in urban Darwin at diagnosis were originally from remote NT Indigenous communities. Over half had moved to Darwin within the preceding year. Nine of these 11 patients had hazardous alcohol intake as a risk factor, including three who were homeless.

Box 6 shows the monthly Top End cases of melioidosis for 2009–2010 in comparison to rainfall recorded at Darwin airport. In the Darwin urban area, there was substantially above average rainfall in December 2009 and January 2010. Total Darwin rainfall for the period 1 October 2009 to 30 September 2010 was 2122.4 mm, compared with the mean yearly rainfall of 1712.7 mm.

Discussion

The year beginning with the wet season of 2009-2010 had the highest recorded number of cases of melioidosis in the Top End for the past 21 years. Over these 12 months, our estimated population-based annual incidence of 50.2 cases per 100 000 is the highest recorded anywhere in the melioidosis-endemic region of South-East Asia and northern Australia. The previous highest annual incidence in the Top End was 41.7 cases per 100 000 population in the calendar year of 1998, which included case clusters following Tropical Cyclone Les, which flooded the town of Katherine, and Tropical Cyclone Thelma, which passed over the north-west end of the Tiwi Islands. The prior highest recorded annual incidence anywhere

4 Clinical presentation and outcomes for patients with melioidosis managed at Royal Darwin Hospital from 1 October 1989 to 30 September 2010

Clinical presentation	1989–2009	2009–2010	P
Total patients	540	91	
Pneumonia	278 (51%)	47 (52%)	1.00
Bacteraemia, no focus	59 (11%)	14 (15%)	0.29
Skin ulcer or abscess	68 (13%)	10 (11%)	0.80
Genitourinary	76 (14%)	8 (9%)	0.29
Soft tissue abscess	19 (4%)	5 (5%)	0.54
Osteomyelitis/ septic arthritis	20 (4%)	3 (3%)	1.00
Neurological melioidosis	14 (3%)	1 (1%)	0.62
Other	6 (1%)	3 (3%)*	0.25
Severity and outcomes			
Bacteraemia	298 (55%)	57 (63%)	0.23
Deaths	61 (20%)	10 (18%)	0.75
Septic shock	116 (21%)	19 (21%)	1.00
Deaths	58 (50%)	9 (47%)	1.00
Ventilated in intensive care	97 (18%)	16 (18%)	1.00
Deaths	41 (42%)	9 (56%)	0.44
Total deaths	77 (14%)	11 (12%)	0.70

^{*} Others in 2009–2010 were ascitic fluid culture-positive melioidosis, liver abscess, mediastinal lymphadenopathy/abscess.

5 Clinical presentations of melioidosis during 2009–2010





A: Chest x-ray of a 66-year-old man from Darwin with chronic lung disease, diabetes and ischaemic heart disease; his was the first death from melioidosis of the wet season, December 2009. B: A 4-year-old boy from Darwin with cutaneous melioidosis; possibly the result of an inoculating event in his muddy backyard. C: A 66-year-old person from Darwin with diabetes, osteomyelitis and a soft tissue abscess of right middle finger.

else was 42.7 per 100 000 in the Torres Strait Islands during 1995–2000. 11

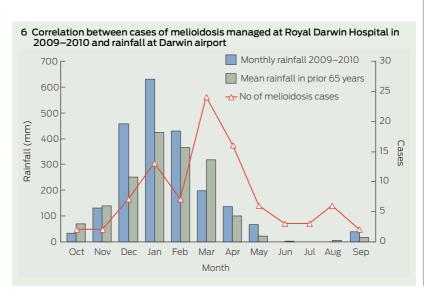
The association between melioidosis and rainfall and severe weather events is well recognised in Australia^{3,5,6} and South-East Asia.^{12,13} The 2009–2010 rainfall pattern is considered the major contributor to the increase in cases. There was consistently heavy rainfall in the Darwin area in December 2009 and January 2010, with two severe weather events leading to intense rainfall in December and February. Other Top End locations had variable rainfall patterns which were generally less intense than in Darwin.

In 2009–2010, the incidence of melioidosis of 102.4 cases per 100 000 for Indigenous Australians was three times higher than that for non-Indigenous Australians. Even though the overall proportion of Indigenous cases remained the same, there was a significant reversal of the previous predominance of infected Indigenous patients being from remote communities to most being from urban Darwin in 2009–2010. The proportion of urban Darwin cases that was Indigenous increased

from 38% to 53%. That four out of five of the Indigenous people from Darwin who died from melioidosis in 2009–2010 were homeless, previously from remote communities and with excessive alcohol intake as their major risk factor, supports concerns that the Northern Territory Emergency Response initiated in 2007 may have had possible unintended consequences resulting from displacement

of drinkers from remote communities to urban centres such as Darwin, where alcohol is available. 14-16

In conclusion, the dramatic increase in the number of cases of melioidosis in the Top End in 2009–2010 represented an estimated population-based annual incidence that is the highest on record. One in every 1000 Indigenous people living in the Top End had culture-confirmed



melioidosis over that 12-month period. The increase in cases was most prominent in the Darwin urban area, and was associated with increased rainfall. There was also an increased proportion of Indigenous Australians who acquired melioidosis in Darwin compared with the preceding 20 years, which may reflect movement of some Indigenous people into Darwin from remote communities.

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