
Increasing hospitalizations for serious skin infections in New Zealand children, 1990–2007

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SUMMARY

The incidence of serious skin infections in New Zealand children is significantly higher than in comparative countries. This study aimed to describe the epidemiology of these infections and identify changes in disease distribution over time. Discharge data were analysed for all children admitted to a New Zealand public hospital with a serious skin infection during the period 1990–2007. Patient and admission variables were compared between 1990–1999 and 2000–2007. The incidence of serious skin infections almost doubled from 298·0/100 000 in 1990 to 547·3/100 000 in 2007. The highest rates were observed in boys, preschool-aged children, Māori and Pacific children, those living in deprived neighbourhoods, urban areas and northern regions. Over time there were disproportionate increases in infection rates in Māori and Pacific children and children from highly deprived areas. Serious skin infections are an increasing problem for New Zealand children. Worsening ethnic and socioeconomic health inequalities may be contributing to increasing rates.

Key words: Infectious disease epidemiology, skin infections, soft tissue infections, *Staphylococcus aureus*, *Streptococcus pyogenes*.

INTRODUCTION

Skin and subcutaneous tissue infections are a heterogeneous group of superficial bacterial infections, most frequently caused by opportunistic skin pathogens: *Staphylococcus aureus* and *Streptococcus pyogenes* [1]. These infections are common and are usually adequately treated within the community; however, in a number of cases serious skin infections develop which require hospitalization for often invasive treatments. An increase in the incidence of serious skin infections in children has been recognized worldwide [2–4]. In New Zealand (NZ) this trend has been particularly

marked with the rate of paediatric cellulitis double that of Australia and the USA [5]. The reasons for this high and increasing disease burden are not understood.

In 2007 a comprehensive report on the health and wellbeing of NZ children and young people was published by Craig *et al.* [6]. It found the incidence of serious skin infections in children had doubled during the last two decades, with the highest rates occurring in preschool-aged children, Māori and Pacific children, boys and children living in areas of greatest deprivation. There have been a number of regional reports with similar observations [5, 7–10]; however, the only study in this field to have been published in the peer-reviewed medical literature is a 1-year retrospective audit of paediatric skin infection admissions to a South Auckland hospital [11]. None of these

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studies have investigated whether changing epidemiology over time could be contributing to the increasing incidence of infections.

Furthermore, recent work has shown the case definition for serious skin infection currently in use is deficient in several areas, with a sensitivity of just 61% when tested against a set of clinically defined serious skin infections [12]. As this narrower case definition was used in the work by Craig *et al.* [6] currently reported national incidence rates are likely to underestimate the true burden of infection, and epidemiological trends are potentially different to those described.

This study aimed to use a newly developed and validated case definition of serious skin infection to describe the incidence of these conditions in NZ children during the period 1990–2007. It also aimed to investigate if there have been changes in the distribution of disease over time that could help explain the increasing infection rates.

METHODS

This study was based on hospital discharge data obtained from the NZ Ministry of Health. It selected children aged 0–14 years, admitted overnight to a NZ public hospital between 1 January 1990 and 31 December 2007, with a principal or additional discharge diagnosis from a defined list of serious skin infection International Classification of Disease (ICD) codes (see Appendix).

This case definition was developed in recent work which found its validity and clinical relevance was markedly improved by including categories of infection previously overlooked [12]. With the addition of skin infections of atypical anatomical sites, those secondary to either primary skin disease or trauma, and those recorded as additional diagnoses (see Appendix), the sensitivity of the case definition increased from 61.0% to 98.9%, with little loss in specificity.

Each discharge record included a unique patient identifier (encrypted National Health Index number) enabling transfers and readmissions within 30 days with the same principal diagnosis code to be removed. To ensure a better match with the census population overseas visitors were removed. Day cases were excluded due to inconsistencies in data recording over time.

Cases were assigned rurality and deprivation levels based on their home domicile census area units

(CAUs). Rurality assignment used a Statistics NZ classification which defines seven grades of rurality on the basis of population size and employment status. Assigning levels of socioeconomic deprivation used the New Zealand Deprivation Index (NZDep) which is based on nine variables extracted from census data [13]. NZDep 1 indicates least deprivation and 10 indicates highest deprivation. In 2.21% of cases domicile codes could not be linked to CAUs due to retired codes and addresses outside classification. To reduce the impact of these ‘missing CAUs’, retired domicile codes were linked to new codes using files from the Ministry of Health and Statistics NZ (R. Bishop, Statistics New Zealand, personal communication; CAU changes 1991–2006, Wellington, 2009; C. Lewis, New Zealand Health Information Service, personal communication; Domicile code mapping, Wellington, 2009).

The data were analysed using Microsoft Excel[®]. Denominators in rate calculations were derived from usually resident population counts from the 1991, 1996, 2001, and 2006 censuses. Counts from each census were used to approximate the population in the preceding and subsequent two years. Trends over time and between populations were explored by the calculation of rate ratios (RRs) with 95% confidence intervals (95% CIs) calculated using the log-transformation method [14]. Changes in the distribution of disease over time were measured by the difference in RRs for each variable between 1990–1999 and 2000–2007, with statistical significance indicated by a two-tailed *P* value <0.01.

The first part of the analysis describes the incidence of serious skin infections for the entire period 1990–2007. To compare trends over time, the more detailed descriptive analyses have split the data into two periods, corresponding to the changeover from ICD-9 to ICD-10 in mid-1999.

RESULTS

Selection of cases, incidence and impact

A total of 82 408 hospitalizations met the case definition. From this we excluded 213 private hospital admissions, 955 overseas visitors, 3109 transfers, 12 353 day cases, and 1210 readmissions.

Of the remaining 64 568 cases, 12 were reported to have been discharged dead from hospital (case fatality 0.04%) from 1990 to 1999 and 17 (0.05%) from 2000 to 2007. Hospitalization data recorded a total of

Table 1. The incidence of serious skin infections in children aged 0–14 years in New Zealand, 1990–1999 and 2000–2007, disaggregated by category and level of diagnosis

Category	Level of diagnosis	1990–1999		2000–2007	
		<i>f</i>	Incidence† (per 100 000)	<i>f</i>	Incidence† (per 100 000)
Serious skin infections of typical sites (previously used case definition based on skin infection sub-chapter of ICD-10)	Principal	13 541	166.3	18 177	264.9
	All level	17 074	209.7	24 086	351.0
Serious skin infections of atypical anatomical sites	Principal	3170	38.9	1866	27.2
	All level	5233	64.3	2270	33.1
Serious skin infections secondary to primary skin disease	Principal	1406	17.3	1909	27.8
	All level	5364	65.9	6170	89.9
Serious skin infections secondary to external trauma	Principal	635	7.8	420	6.1
	All level	1270	15.6	3101	45.2
Total serious skin infections	Principal	18 752	230.3	22 372	326.0
	All level	28 941	355.4	35 627	519.2

f, Frequency is number of cases in 1990–1999 and 2000–2007.

† Average annual incidence per 100 000 based on usually resident population (from NZ Census).

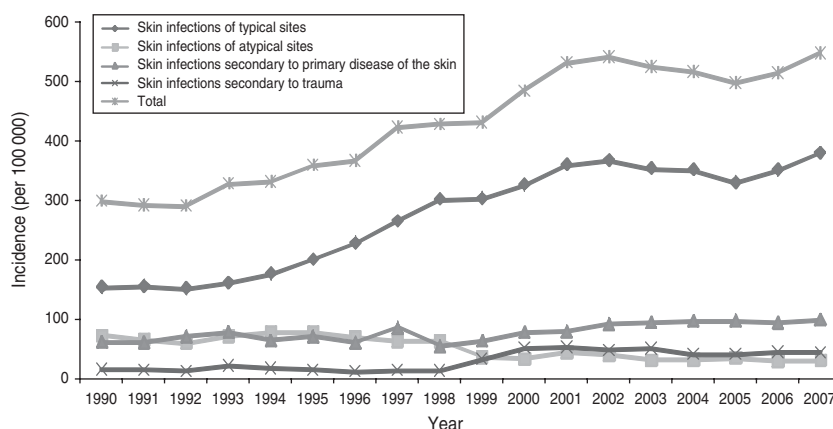


Fig. 1. Incidence of serious skin infection in children aged 0–14 years in New Zealand by ICD-10 code category 1990–2007.

213 141 hospital days over the study period. The median stay was 2 days and mean was 3.3 days in both 1990–1999 and 2000–2007.

Based on a 2003 estimate of hospitalization costs per case of NZ\$2180 [5], the direct cost of these infections for District Health Boards (DHBs) in 2007 alone was almost NZ\$15 million (based on an inflation-adjusted cost per case of NZ\$2434.21).

Table 1 shows the incidence of serious skin infections over 1990–1999 and 2000–2007. As recommended by the previous work developing the case definition [12], and to provide an indication of the level of certainty of these estimates, these data are disaggregated by category and level of diagnosis (see Appendix for details). This analysis shows that the distribution of disease between principal and

additional diagnosis categories changed little over this period (64.8% principal diagnosis in 1990–1999 and 62.8% in 2000–2007).

Incidence by year and season, 1990–2007

Over the 18-year period analysed the average annual incidence rate of serious skin infections in NZ children almost doubled, from 298.0/100 000 in 1990 to 547.3/100 000 in 2007 (see Fig. 1). In the first two years of the study there was a largely stable incidence around 300/100 000, then from 1992–2002 infection rates steadily rose to over 500/100 000. Since 2002 the incidence has been relatively steady again. These trends were a direct reflection of changes in the incidence of serious skin infections of typical sites, with the rates of

Table 2. Childhood serious skin infection frequency, incidence and rate ratios by season, gender, age, ethnicity, rurality and deprivation, 1990–1999 and 2000–2007

Variable	1990–1999			2000–2007			Difference in RRs of each variable between time periods† <i>P</i>
	<i>f</i>	Incid.*	RR (95% CI)	<i>f</i>	Incid.*	RR (95% CI)	
Season‡							
Summer	7508	368.8	1.08 (1.05–1.11)	9594	559.3	1.15 (1.12–1.18)	0.13
Autumn	7722	379.3	1.11 (1.08–1.14)	9454	551.2	1.14 (1.11–1.17)	0.51
Winter	6963	342.0	1.00	8327	485.4	1.00	
Spring	6748	331.5	0.97 (0.94–1.00)	8252	481.1	0.99 (0.97–1.02)	0.63
Gender							
Male	16909	405.2	1.34 (1.31–1.37)	20440	581.5	1.28 (1.25–1.31)	0.21
Female	12031	303.3	1.00	15187	453.8	1.00	
Unknown	1						
Age (yr)							
0–4	16454	592.5	2.20 (2.15–2.25)	19922	912.4	2.45 (2.40–2.50)	0.03
5–9	7348	268.9	1.00	8525	372.1	1.00	
10–14	5139	195.1	0.73 (0.71–0.75)	7180	300.8	0.81 (0.79–0.83)	0.02
Ethnicity							
Māori	10002	546.3	2.28 (2.22–2.34)	13734	866.2	2.90 (2.84–2.96)	<0.001
Pacific	5251	885.7	3.70 (3.59–3.81)	7849	1351.9	4.52 (4.41–4.63)	<0.001
Other	13688	239.3	1.00	14044	299.1	1.00	
NZDep§							
1–2	2376	179.3	1.00	2937	226.2	1.00	
3–4	3361	236.7	1.32 (1.27–1.37)	3829	298.3	1.32 (1.28–1.37)	1.00
5–6	4494	295.5	1.65 (1.60–1.70)	4968	390.3	1.73 (1.68–1.78)	0.34
7–8	6300	374.4	2.09 (2.03–2.15)	7745	569.1	2.52 (2.46–2.58)	<0.001
9–10	12136	637.8	3.56 (3.48–3.65)	15966	972.0	4.30 (4.21–4.39)	<0.001
Missing¶	274			182			
Rurality							
Urban	25757	390.0	1.00	32177	555.3	1.00	
Rural	2925	235.3	0.60 (0.58–0.62)	3267	306.4	0.55 (0.53–0.57)	0.16
Missing¶	259			183			
Total	28941			35627			

RR, Rate ratio; CI, confidence interval.

f, Frequency is number of cases in 1990–1999 and 2000–2007.

* Average annual incidence per 100 000 based on usually resident population (from NZ Census).

† The *P* value is comparing the RRs of each variable across the two time periods. *P* < 0.01 indicates a statistically significant difference in the RR of the variable between 1990–1999 and 2000–2007.

‡ Autumn (March, April, May); winter (June, July, August); spring (September, October, November); summer (December, January, February).

§ The New Zealand Deprivation Index (NZDep) is a measure of socioeconomic deprivation based on nine variables extracted from census data [13]. NZDep 1 indicates least deprivation and 10 indicates highest deprivation.

|| Arbitrary reference category.

¶ Missing refers to cases with domicile codes that could not be linked to CAUs.

infections of atypical sites and those secondary to primary skin disease and trauma fairly stable over time.

Table 2 shows the incidence of serious skin infections by season of admission. Infections were significantly more frequent during summer and autumn

compared to winter (summer: RR 1.08 in 1990–1999 and 1.15 in 2000–2007; autumn: RR 1.11 in 1990–1999 and 1.14 in 2000–2007). There was no change in this seasonal trend over time (*P* > 0.01 for all seasons).

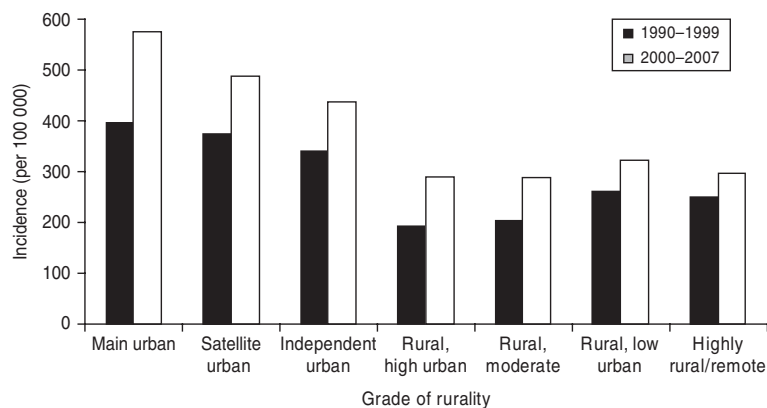


Fig. 2. Incidence of serious skin infections in children aged 0–14 years in New Zealand by rurality, 1990–2007.

Incidence by gender, age, ethnicity, deprivation level, and rurality, 1990–2007

Table 2 details serious skin infections in NZ children by a range of patient characteristics, across two time periods 1990–1999 and 2000–2007.

Boys had a significantly greater risk of infection than girls, with an incidence rate of 405.2/100 000 compared to 303.3/100 000 (RR 1.34) in 1990–1999 and 581.5/100 000 compared to 453.8/100 000 (RR 1.28) in 2000–2007. There was no significant difference in the RRs between the two time periods ($P=0.21$).

The incidence of skin infections decreased with increasing age. Children aged 0–4 years had more than double the risk of infection than those aged 5–9 years (RR 2.20 in 1990–1999 and 2.45 in 2000–2007) and the 10–14 years age group had the lowest rates of infection overall. Between the two time periods there were increases in the proportion of cases in both the youngest and oldest age groups relative to the reference 5–9 years age group; this difference in RRs approached but did not reach statistical significance ($P=0.03$ and $P=0.02$).

The rate of serious skin infections was significantly higher in Māori and Pacific children than those in other ethnic groups. In 1990–1999 the incidence rate was 2.28 times higher in Māori children, and 3.70 times higher in Pacific children, compared to those of other ethnicities. By 2000–2007 that difference had increased to 2.90 times higher in Māori children and 4.52 times higher in Pacific children. The difference in RRs over time was statistically significant ($P<0.001$).

The incidence of serious skin infections was lowest in areas of least deprivation and increased markedly with rising deprivation levels. During the period 1990–1999, the rate of infection in children from

NZDep 9–10 areas was 3.56 times greater than for children from NZDep 1–2 areas (179.3/100 000 and 637.8/100 000, respectively). By 2000–2007 this difference had increased significantly to 4.30 times higher ($P<0.001$).

Serious skin infection rates were more than 1.5 times higher in children from urban areas compared to children from rural areas (see Fig. 2). Of the three urban classifications, the incidence of infection steadily decreased as areas became increasingly rural. The gradient of this trend appeared steeper in 2000–2007 compared to 1990–1999, but there was no statistically significant difference found ($P=0.16$). The incidence of infection in the four rural areas was similar, with no change over time.

Incidence by DHB

Figure 3 shows the incidence of serious skin infections across all 21 geographically assigned NZ DHBs. There was a rough north–south gradient with higher rates generally observed in North Island DHBs (Northland–Wairarapa) compared to South Island DHBs (Nelson–Malborough–Southland). The incidence of infection increased over time in all DHBs except West Coast, where a small decrease was observed. Tairāwhiti DHB had the highest incidence rate during both time periods studied, approaching double the national rate with an incidence of 644.9/100 000 in 1990–1999 rising to 961.4/100 000 in 2000–2007.

DISCUSSION

Serious skin infections are an important and increasing challenge to the health of NZ children, and are

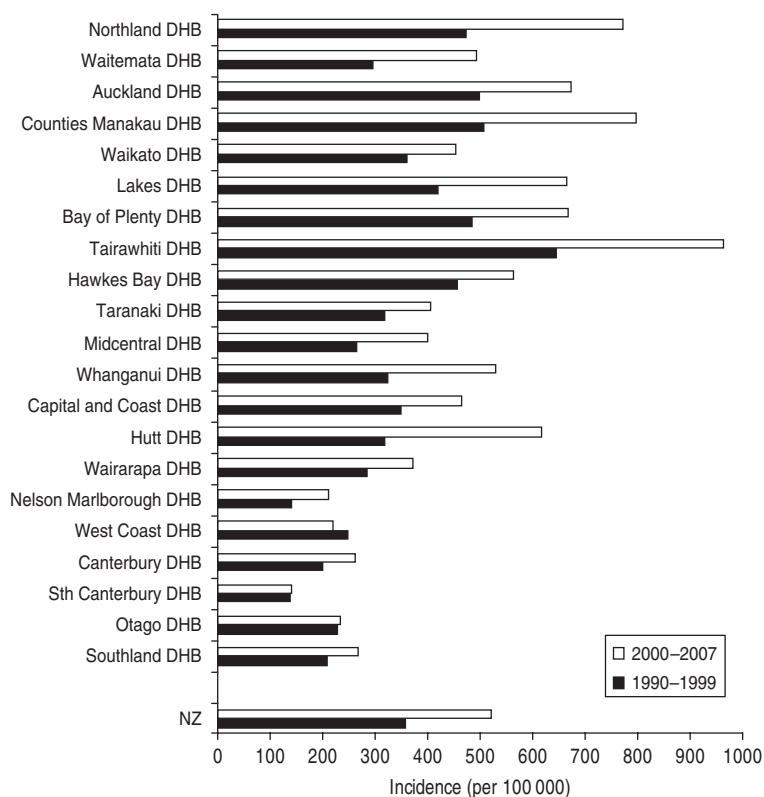


Fig. 3. Incidence of serious skin infections in children aged 0–14 years in New Zealand by District Health Board (DHB), 1990–2007.

making a growing contribution to ethnic and socio-economic health inequalities.

Recent work has shown the case definition for serious skin infections used in previous publications has been deficient in a number of areas, lacking sensitivity, standardization and clinical relevance. By including several categories of skin infections previously overlooked, the sensitivity of this definition could be increased from 61.0% to 98.9% with minimal loss of specificity [12]. This is the first study to apply this standardized case definition to the NZ paediatric population. Findings showed that between 1990 and 2007 the incidence of serious skin infections almost doubled, from 298.0/100 000 to 547.3/100 000. Besides the considerable medical and social impact, this incidence equates to a sizable public expenditure on health services, even without accounting for other direct and indirect economic costs.

Furthermore, this is the only study we are aware of that has investigated whether there have been changes in the distribution of serious skin infections over time which could account for the increasing incidence of disease. Between 1990 and 2007 the highest infection rates were observed in boys, children aged <5 years,

Māori and Pacific children, those living in deprived neighbourhoods, in urban areas and in northern districts of the country. While these high-risk groups have remained the same between 1990–1999 and 2000–2007, ethnic and deprivation-related disparities have significantly increased. This change in the distribution of serious skin infections over time is contributing to the increasing incidence of disease; however, it cannot explain the whole increase as rates have risen across all population groups.

The incidence of infections showed marked and consistent seasonality. High disease rates in summer and autumn have been observed in other settings and are believed to result from warmer air temperatures leading to more frequent insect bites, deficiencies in hygienic precautions, and increased skin exposure resulting in greater skin-to-skin contact and minor trauma [2, 3, 5, 15–20].

High rates of infection in preschool-aged children and boys have important implications for directing community prevention efforts. In both groups this increased risk could be attributable to more frequent injuries, poorer general hygiene, or longer delays in seeking medical attention. It is not known whether

these trends reflect similar patterns in the community, or in the case of younger children, of lower hospital admission thresholds. Previous NZ reports have observed similar trends [5–7, 11], but interestingly studies in the international literature report no gender predominance [2, 16, 20–23]. As these studies mainly comprise primary-care and population surveys, it is possible that simple skin infections are experienced equally by both genders but boys are more likely to suffer progression to a serious skin infection.

Māori and Pacific children had higher rates of serious skin infections than children of other ethnicities. This finding is consistent with the wider observation that Māori and Pacific peoples generally experience high rates of infectious diseases [24]. The reasons for this pattern are complex and multifactorial; they include household crowding and a range of socio-economic factors [25, 26]. It is not known whether the high incidence of serious skin infections directly reflects higher community rates of disease, but it is known that Māori and Pacific families experience greater barriers to accessing primary healthcare including cost, cultural differences and longer travel distances [27–29]. In addition, hospitalization data have been found to often undercount Māori [30, 31] which would result in an underestimation of ethnic inequalities. The significant increase in ethnic disparities between the two time periods is of particular concern. While some of this increase could be due to inconsistencies in how census (denominator) data has been collected over time, this factor is unlikely to completely account for the difference.

Socioeconomic deprivation was seen to be an important risk factor for serious skin infection with a steady increase in infection incidence with increasing neighbourhood deprivation. While this association is well established and thought to be mediated by hygiene, nutrition, household crowding, and the ability to afford timely medical treatment [5, 6, 17, 21], the evidence of increasing inequality has not been previously recognized. As there has been little change to the allocation and recording of deprivation status, these findings suggest truly worsening deprivation disparities over time.

The association between serious skin infection incidence and level of rurality has not been reported previously in NZ. The high incidence of infection in children from urban areas could be due to socio-economic deprivation, household crowding and a higher frequency of skin contact with other children in more densely populated cities. It is possible that

lower infection rates in rural areas could reflect more frequent treatment in the community due to reduced access to hospitals. However, as the management of serious skin infections in children frequently involves hospital-based treatments such as intravenous antibiotics and surgical debridement, it is unlikely that many true serious skin infections would be managed in an outpatient setting.

Within NZ, Tairāwhiti DHB had the highest incidence of childhood serious skin infections. This finding could be a result of the large Māori population and high deprivation of the region, but requires greater investigation. The observed north–south gradient may in part reflect the distribution of population groups who experience higher disease rates, but could also relate to climatic differences, with northern districts of NZ experiencing relatively warmer weather compared to those in the south.

The descriptive data presented in this paper have limited potential for identifying factors that have caused the rise in serious skin infections and increases in ethnic and socio-economic inequalities. Other work in New Zealand over a similar time period has shown a marked increase in rates of infectious diseases generally, along with rising ethnic inequalities [24]. In addition, rheumatic fever incidence has shown worsening ethnic inequalities from 1996 to 2005 with this disease increasingly concentrated in Māori and Pacific children [32]. The causes for these changes are not known.

There are several programmes monitoring health determinants and outcomes over time for NZ children and households [6, 33, 34]. This monitoring showed a mixed pattern over the last two decades. Mean income levels have been rising for New Zealanders but inequalities have increased, particularly during the 1990s [34]. While average levels of household crowding have generally decreased, there have been comparatively small declines in the proportion of children living in crowded conditions and persisting large ethnic inequalities [33].

Hospitalization data have strengths and weaknesses as a basis for surveillance of serious skin infections. The main limitation of these data is that, by definition, they only represent the ‘tip of the iceberg’ and cannot measure the incidence of mild to moderate skin infections in the community. This limitation is common to other areas of infectious disease epidemiology, such as acute gastroenteritis [35]. The advantages of this data source is that it is accessible and is likely to be relatively sensitive for serious skin

infections as few paediatric cases would be treated outside of the public hospital setting due to the close monitoring and invasive treatments required. On this basis, and as by definition serious skin infections are those skin infections which require hospitalization, we used the term 'incidence' to describe hospitalization rates.

It is possible that the sensitivity of such surveillance has changed over time, for example, changes to the recording of day patients as admissions. We have attempted to minimize such effects by the use of a fairly high threshold for inclusion. A further consideration is modifications to the disease coding system; despite using standardized mapping tables, translating diagnoses between ICD-9 and ICD-10 was problematic in some areas with the incidence of several diagnoses markedly varying over time (see Appendix). However, as there was a steady increase in the total infection incidence over the years when the ICD revision occurred, the variation is more likely to reflect inter-code and inter-category drift, and further justifies our use of a more inclusive case definition than that used previously.

This study demonstrates an urgent need for action to prevent serious skin infections in NZ children. It highlights population groups with disparate rates of disease to which these efforts should be particularly focused. Future work could include extending this simple univariate analysis to a multivariate model to identify the independent effects of each risk factor. A retrospective case-note review or a case-control study would better elucidate the aetiological processes contributing to the development of serious skin infections. In order to allow clear and accurate comparison, any future work on hospitalization data could utilize the standardized case definition of serious skin infection used in this study. In addition, the epidemiology of skin infections in primary care is largely unknown; future study in this area could improve our understanding of whether inequalities in serious skin infection rates directly reflect community trends. In combination with such ongoing work, the findings of this present study indicate some priority areas for directing interventions to reduce the morbidity of serious skin infections in NZ children and to narrow health inequalities.

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DECLARATION OF INTEREST

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REFERENCES

1. Sladden MJ, Johnston GA. Common skin infections in children. *British Medical Journal* 2004; **329**: 95–99.
2. Koning S, *et al.* Impetigo: incidence and treatment in Dutch general practice in 1987 and 2001 – Results from two national surveys. *British Journal of Dermatology* 2006; **154**: 239–243.
3. Loffeld A, *et al.* Seasonal occurrence of impetigo: a retrospective 8-year review (1996–2003). *Clinical and Experimental Dermatology* 2005; **30**: 512–514.
4. Hersh AL, *et al.* National trends in ambulatory visits and antibiotic prescribing for skin and soft-tissue infections. *Archives of Internal Medicine* 2008; **168**: 1585–1591.
5. Hunt D. Assessing and reducing the burden of serious skin infections in children and young people in the Greater Wellington Region. Wellington: Capital and Coast DHB, Hutt Valley DHB, Regional Public Health, 2004 (http://www.skininfections.co.nz/documents/Serious_Skin_Infections_Nov2004.pdf).
6. Craig E, Jackson C, Han DY, NZCYES Steering Committee. Monitoring the health of New Zealand children and young people: Indicator Handbook. Auckland: Paediatric Society of New Zealand, New Zealand Child and Youth Epidemiology Service, 2007. (<http://www.paediatrics.org.nz/files/Indicator%20Handbook%20Version%202008.3.pdf>).
7. Lawes C. Paediatric cellulitis hospital discharges in the Auckland Region. Auckland: Public Health Protection Service, Auckland Healthcare, 1998.
8. Morgan C, Selak V, Bullen C. Glen Innes Serious Skin Infection Prevention Project: Final Report 1 February 2003–31 January 2004. Auckland: Auckland Regional Public Health Services, 2004 (http://www.arphs.govt.nz/Publications_Reports/archive/GlenInnesSkinProject.pdf).
9. Leversha A. Starship Hospital cellulitis case-control study. Preliminary results reported in reference 8.
10. Leversha A, *et al.* Case series of children admitted to Starship Hospital with cellulitis. Preliminary results reported in reference 8.
11. Finger F, *et al.* Skin infections of the limbs of Polynesian children. *New Zealand Medical Journal* 2004; **117**: U847.
12. O'Sullivan C, Baker M. A proposed epidemiological case definition for serious skin infection in children. *Journal of Paediatrics and Child Health* 2010; **46**: 176–183.
13. Salmond C, Crampton P, Atkinson J. NZDep 2006 Index of Deprivation: User's Manual. Wellington: Ministry of

- Health, 2007 ([http://www.moh.govt.nz/moh.nsf/Files/phi-users-manual/\\$file/phi-users-manual.pdf](http://www.moh.govt.nz/moh.nsf/Files/phi-users-manual/$file/phi-users-manual.pdf)).
14. **Clayton D, Hills M.** *Statistical Methods in Epidemiology*. Oxford: Oxford University Press, 1993, pp. 80–82.
 15. **Kristensen JK.** Scabies and pyoderma in Lilongwe, Malawi: prevalence and seasonal fluctuation. *International Journal of Dermatology* 1991; **30**: 699–702.
 16. **Elliot AJ, et al.** The association between impetigo, insect bites and air temperature: a retrospective 5-year study (1999–2003) using morbidity data collected from a sentinel general practice network database. *Family Practice* 2006; **23**: 490–496.
 17. **Kakar N, et al.** Clinico-bacteriological study of pyoderma in children. *Journal of Dermatology* 1999; **26**: 288–293.
 18. **Masawe A, Nsanzumuhire H, Mhalu F.** Bacterial skin infections in preschool and school children in costal Tanzania. *Archives of Dermatology* 1975; **111**: 1312–1316.
 19. **Taplin D, et al.** Prevalence of streptococcal pyoderma in relation to climate and hygiene. *Lancet* 1973; **1**: 501–503.
 20. **Rogers M, et al.** A three-year study of impetigo in Sydney. *Medical Journal of Australia* 1987; **147**: 63–65.
 21. **Bailie RS, et al.** Skin infection, housing and social circumstances in children living in remote Indigenous communities: testing conceptual and methodological approaches. *BMC Public Health* 2005; **5**: 128.
 22. **Lawrence D, et al.** Epidemiologic studies among Amerindian populations of Amazonia. I. Pyoderma: prevalence and associated pathogens. *American Journal of Tropical Medicine and Hygiene* 1979; **28**: 548–58.
 23. **Dajani A, Ferrieri P, Wannamaker L.** Endemic superficial pyoderma in children. *Archives of Dermatology* 1973; **108**: 517–522.
 24. **Baker M, et al.** Close-contact infectious diseases in New Zealand: Trends in ethnic inequalities in hospitalisations, 1989–2008. Wellington: University of Otago, 2010 (<http://www.healthyhousing.org.nz/wp-content/uploads/2010/06/Close-Contact-IDs-in-NZ-June-2010.pdf>).
 25. **Baker M, et al.** Household crowding a major risk factor for epidemic meningococcal disease in Auckland children. *Pediatric Infectious Disease Journal* 2000; **19**: 983–990.
 26. **Grant CC, et al.** Hospitalization for pneumonia in children in Auckland, New Zealand. *Journal of Paediatrics and Child Health* 1998; **34**: 355–359.
 27. **Malcolm L.** Inequities in access to and utilisation of primary medical care services for Maori and low income New Zealanders. *New Zealand Medical Journal* 1996; **109**: 356–358.
 28. **Tukuitonga CR, Bell S, Robinson E.** Hospital admission among Pacific children Auckland 1992–97. *New Zealand Medical Journal* 2000; **113**: 358–361.
 29. **Brabyn L, Barnett R.** Population need and geographical access to general practitioners in rural New Zealand. *New Zealand Medical Journal* 2004; **117**: U996.
 30. **Harris R, et al.** Estimating Maori hospitalisations and cancer registrations. In: Robson B, Harris R, eds. *Hauora: Maori Standards of Health*, 4th edn. Wellington: Te Ropu Rangahau Hauora a Eru Pomare, 2007, pp. 249–259.
 31. **Ministry of Health.** Tatau Kahukura: Maori health chart book. Public Health Intelligence Monitoring Report No. 5. Wellington: Ministry of Health, 2006.
 32. **Jaine R, Baker M, Venugopal K.** Epidemiology of acute rheumatic fever in New Zealand 1996–2005. *Journal of Paediatrics and Child Health* 2008; **44**: 564–571.
 33. **Ministry of Social Development.** Children and Young People: Indicators of Wellbeing in New Zealand 2008. Wellington: Ministry of Social Development, 2008 (<http://www.msd.govt.nz/documents/about-msd-and-our-work/publications-resources/monitoring/children-young-indicators-wellbeing/2008-report/cyi-report-2008.pdf>).
 34. **Ministry of Social Development.** The Social Report, 2009. Wellington: Ministry of Social Development, 2009 (<http://socialreport.msd.govt.nz/documents/social-report-2009.pdf>).
 35. **Lake R, et al.** The disease pyramid for acute gastrointestinal illness in New Zealand. *Epidemiology and Infection* 2010; **138**: 1468–1471.

APPENDIX. The incidence of serious skin infections in children aged 0–14 years in New Zealand, 1990–1999 and 2000–2007, disaggregated by ICD code, coding category and level of diagnosis

Skin infection	ICD-9†	ICD-10‡	Level of diagnosis	<i>f</i> 1990–1999	Incid. 1990–1999‡	<i>f</i> 2000–2007	Incid. 2000–2007‡
Serious skin infections of typical sites*							
Impetigo	684	L01.0-L01.1	Principal	643	7.9	694	10.1
			All level	1486	18.2	1609	23.5
Cutaneous abscess, furuncle and carbuncle	6800-6809	L02.0-L02.9	Principal	767	9.4	7368	107.4
			All level	900	11.1	8120	118.3
Cellulitis	68100-68102, 68110, 68111, 6819-6829	L03.01-L03.9	Principal	10 300	126.5	8411	122.6
			All level	11949	146.7	11002	160.3
Acute lymphadenitis	683	L0.40-L04.9	Principal	553	6.8	1026	15.0
			All level	632	7.8	1168	17.0
Pilonidal cyst with abscess	6850	L05.0	Principal	43	0.5	108	1.6
			All level	49	0.6	111	1.6
Pyoderma	6860	L08.0	Principal	35	0.4	45	0.7
			All level	94	1.2	452	6.6
Other infections of skin and subcutaneous tissue	390, 6868-6869, 9101-9179, 9191-9199	L08.1, L08.8, L08.9	Principal	1200	17.5	525	7.7
			All level	1964	24.1	1624	23.7
Total			Principal	13 541	166.3	18177	264.9
			All level	17 074	209.7	24 086	351.0
Serious skin infections of atypical anatomical sites							
Erysipelas	035	A46	Principal	42	0.5	22	0.3
			All level	47	0.6	25	0.4
Hordeolum/cellulitis/abscess eyelid	37 311-37 313	H00.0	Principal	308	3.8	449	6.5
			All level	371	4.6	555	8.1
Abscess/cellulitis external ear and infective otitis externa	38 010, 38 011, 38 013, 38 014	H60.0-H60.3, H62.0, H62.4	Principal	599	7.4	378	5.5
			All level	974	12.0	533	7.8
Abscess/cellulitis nose	4781	J34.0	Principal	695	8.5	111	1.6
			All level	2080	25.5	133	1.9
Anal abscess/cellulitis (excludes rectal, ischiorectal or intersphincteric regions)	566	K61.0	Principal	666	8.2	496	7.2
			All level	701	8.6	536	7.8
Acute inflammation/cellulitis/abscess of orbit	37600-37601	H05.0	Principal	584	7.2	143	2.1
			All level	733	9.0	155	2.3
Other inflammatory disorders of penis, scrotum and unspecified male genital organ (excludes deeper tissues)	6072, 6084	N48.2, N49.2, N49.9	Principal	149	1.8	97	1.4
			All level	194	2.4	150	2.2
Abscess/cellulitis of vulva	6164	N76.4	Principal	127	1.6	170	2.5
			All level	133	1.6	183	2.7
Total			Principal	3170	38.9	1866	27.2
			All level	5233	64.3	2270	33.1
Serious skin infections secondary to primary skin disease							
Varicella with other complications	0527-0528	B01.8	Principal	240	2.9	271	3.9
			All level	293	3.6	316	4.6
Scabies	1330	B86	Principal	269	3.3	92	1.3
			All level	1273	15.6	504	7.3

APPENDIX (cont.)

Skin infection	ICD-9†	ICD-10†	Level of diagnosis	<i>f</i> 1990–1999	Incid. 1990–1999‡	<i>f</i> 2000–2007	Incid. 2000–2007‡
Dermatitis unspecified and other specified (eczema) and infective eczema§	6908, 6929, 7028	L30.8, L30.9, L30.3 0	Principal	897	11.0	1546	22.5
			All level	3798	46.6	5350	78.0
Total			Principal	1406	17.3	1909	27.8
			All level	5364	65.9	6170	89.9
Serious skin infections secondary to external trauma							
Insect/spider bites	9104, 9114, 9124, 9134, 9144, 9154, 9164, 9174, 9192, 9194, 9198, 9248, 9895	S10.13, S10.8, S10.93, S20.13, S20.33, S20.43, S20.83, S30.83, 30.93, S40.83, S50.83, S60.83, S70.83, S80.83, S90.83, T00.9, T09.03, T11.08, T13.03, T14.03, T14.03, T63.3, 63.4	Principal	469	5.8	288	4.2
			All level	646	7.9	454	6.6
Post-traumatic wound infection not elsewhere classified	9583	T79.3	Principal	158	1.9	108	1.6
Open wound infection with foreign body (+infection) and open wound with infection	8799	T89.01, T89.02	Principal	442	5.4	457	6.7
			All level	8	0.1	24	0.3
			All level	182	2.2	2190	31.9
Total			Principal	635	7.8	420	6.1
			All level	1270	15.6	3101	45.2

f, Frequency is number of cases in 1990–1999 and 2000–2007.

* Case definition based on skin infection sub-chapter of ICD-10.

† Cases after 1999 were identified using ICD-10 diagnostic codes, while cases prior to this year were identified by ICD-9 codes which were forward and backward mapped from ICD-10.

‡ Average annual incidence per 100 000 based on usually resident population (from NZ Census).

§ The medical definition of infective eczema (a primarily inflammatory condition) is not in keeping with the clinical description of a serious skin infection; however, due to similarities in terminology, this code is incorrectly used for eczema with a superficial bacterial infection.