ABSTRACT

Introduction New Zealand (NZ) has a persistently high rate of suicide, particularly among young people. Hospital presentation for self-harm (SH) is one of the strongest predictors of death by suicide. Improving the monitoring of SH and suicide is a key recommendation for suicide prevention by WHO. This study will establish the first ever sentinel surveillance for SH at several large hospitals and a monthly survey of all practicing paediatricians in NZ. The study will provide robust information about the epidemiology of SH, factors associated with SH and the types of interventions required for those presenting to hospital with SH.

Method and analysis This observational study will establish SH surveillance in the emergency departments of three public hospitals for the first time in NZ, where study population will include individuals of all ages who present with SH or suicidal ideation. The study methodology is in line with the WHO Best Practice guidelines and international collaborators in Australia and Europe. Electronic triage records will be reviewed manually by the research team to identify potential cases that meet inclusion criteria. For all eligible cases, variables of interest will be extracted from routine clinical records by the research team and recorded on a secure web-based survey application. Additionally, SH surveillance data for the national paediatric population (<15 years) will be obtained via the New Zealand Paediatric Surveillance Unit (NZPSU); paediatricians will report on included cases using the same variables using a secure survey application. A deidentified dataset will be produced for aggregated statistical analysis.

Ethics and dissemination The University of Otago Health Ethics Committee granted ethical approval for this study in addition to local ethics approval at participating hospital sites. The study findings will be disseminated to relevant stakeholders in NZ, in addition to international audiences through publications in peer-reviewed scientific journals and conference presentations.

INTRODUCTION

In 2016, the WHO estimated an annual global suicide rate of 10.5 per 100 000.1 Premature deaths by suicide are particularly tragic; they first occur early in the life course, resulting in significant disability-adjusted life-years2 and economic costs of around US$2.2 billion.3 Second, suicide is preventable. Third, the social and health cost of suicide borne by those affected by suicide death is significant.

People who have engaged in self-harm (SH) (intentional self-injury or self-poisoning regardless of degree of intention to die) are at much greater risk suicide than the general population4–6 with a risk of suicide around 30 thirty higher in the year following hospital presentation for SH. Males, older adolescents, those presenting with self-injury (compared with self-poisoning) and repeated SH are at particularly elevated risk of death by suicide.7 Although relatively common, suicidal ideation (having thoughts or plans for suicide but not acted on them) is also an important factor to target in suicide prevention. Around three in five individuals transition from ideation to a plan and suicide attempt within the first year of having suicidal ideation.8 Hospital presentation for SH is key predictor of suicide, so a comprehensive understanding

Strengths and limitations of this study

⇒ Using the WHO best practice guidelines, the surveillance system will provide real-time data monitoring of self-harm (SH) to establish good quality epidemiological information about SH presentations to hospitals in New Zealand (NZ), which have not been publicly reported since 2013.
⇒ The surveillance system will provide information about the characteristics of people presenting to hospital following SH, allowing robust service planning decisions and the ability to evaluate the impact of interventions on subsequent presentations.
⇒ Real-time surveillance allows the detection of novel methods of SH, and appropriate intervention to reduce access, where possible.
⇒ This study is based on presentations for SH to public hospitals so is not an accurate estimate of community cases of SH.
⇒ There is significant variation in the completeness of routinely collected data, so not all cases of SH may be identified.
of this high-risk population, and their treatment needs is essential as part of a multi-level approach to suicide prevention.¹⁰

New Zealand (NZ) is a high-income country spread across two large islands (North Island, South Island) with a population of 5 million people, from a range of ethnic groups including European/Pakeha (70%), indigenous Māori (17%), Asian (15%) and Pacific Peoples (8%).¹¹ The healthcare system in NZ provides free of charge universal care to eligible citizens funded via general taxes. Around one third of the population also carry private health insurance.¹² Currently 20 District Health Boards (DHB) operate around 40 public hospitals, of which 28 have 24/7 emergency departments. There are approximately 1 million presentations per annum with wait time and resource challenges similar to other high-income countries.¹³

In 2016, the most recent year for which there are finalised publicly available data, NZ reported a total population suicide rate of 11.5 deaths per 100,000 people, with higher rates among young people (16.8 per 100,000)¹⁴ and indigenous Māori having rates which are nearly double (RR¹ 1.85, 95% CI 1.64 to 2.10)¹⁵ those of non-Māori (17.1 per 100,000) compared with 8.1 per 100,000 for Pacific peoples and 11.3 per 100,000 for other ethnic groups. The burden of suicide is particularly evident for Māori males (31.3 per 100,000 compared with non-Māori males 14.4 per 100,000), however rates among Māori women have risen 40% between 1996 and 2001.¹⁶ The WHO estimates that NZ suicide rates in 2019 (provisional data) were 10.3 per 100,000, the same as Canada, higher than Ireland (8.9 per 100,000) and United Kingdom (6.9 per 100,000) and lower than Australia (11.3 per 100,000) and the USA (14.5 per 100,000).¹⁷

Despite high public interest, relatively little progress has been made in reducing suicide in NZ,¹⁴ partly due to lack of good quality data to inform interventions; current data reporting practices based on routinely collected data on presentations to public hospitals in undercount SH presentations in NZ by an estimated 50%–60% for several reasons including only episodes with a length of stay greater than 48 hours which misses a significant proportion of presentations. Government reported data also refers to episodes rather than individuals, making it hard to estimate repetition rates, and report on a limited number of variables¹⁴¹⁸ most of which are not amenable to change. In addition, mental health presentations appear particularly poorly coded by hospital staff compared with physical health complaints, which leads to further undercounting of this population.¹⁹

Improving the monitoring of SH and suicide is a key recommendation from the WHO and setting up a SH surveillance system is a key strategy.²⁰ Surveillance data can help establish the true rate of hospital presentations for SH and suicidal ideation, provide robust data about risk factors, give a much better understanding of who comes to hospital following SH/suicidal ideation, the types of hospital treatments they receive, and indications of which interventions may be likely to reduce representations to hospital, and ultimately deaths by suicide.¹²¹

Internationally there are a small number of dedicated surveillance systems in place.²² These include the Multi-centre Study of Self-Harm in England,²³ the Hunter Area Toxidology Service in Australia,²⁴ the Bristol Self-Harm Surveillance Register²⁵ and the National Self-Harm Registry Ireland.²⁶ These systems operate within hospital emergency department settings and collect data from either regional geographical area in the case of Hunter Area Toxidology Service and Bristol Self-Harm Surveillance Register or a broader geographical area in the case of the Multi-centre Study of Self-Harm in the UK, and nationally in the case of the National Self-Harm Registry Ireland. These surveillance systems contribute valuable epidemiological data about prevalence and patterns of SH in their respective geographical areas and are actively used to provide high-quality information to inform suicide prevention initiatives. This NZ surveillance system will use similar methods to facilitate comparisons and build on existing learnings, but is novel in the use of manual screening of all hospital emergency department presentations to check for cases, the inclusion of cases with suicidal ideation only and the first international use of a Paediatric Surveillance system to monitor SH as a condition of interest.

Several small, short-term studies of emergency department (ED) presentations of SH in NZ have shown that a surveillance approach can be achieved.²⁶²⁷ However, these studies are relatively old, and focused on a single hospital, a specific sub-group of patients or a single method of presentation. Using the WHO practice manual for establishing and maintaining surveillance systems for suicide attempts and SH,²⁸ this study aims to achieve the following primary objectives:

1. To establish sentinel surveillance for SH at several large hospitals and a monthly survey of all practicing paediatricians in Aotearoa/NZ in conjunction with the New Zealand Paediatric Surveillance Unit (NZPSU) (The NZPSU, based at the University of Otago, facilitates national surveillance to improve the knowledge of a number of uncommon high-impact childhood conditions in NZ, www.otago.ac.nz/nzpsu).

2. To establish and test robust data collection methods.

3. Identify the epidemiology of current presentations for SH in terms of age, gender, ethnicity, methods of SH, alcohol/substance misuse, prior history of SH, degree of intention to die, exposure to SH/suicide, mental health assessments and discharge outcomes.

The secondary objectives of this study are to: (1) identify potential gaps in SH services; (2) identify systemic levers for suicide prevention, and (3) identify predictors of repetition of SH.

The study objectives and methods will be comparable with international SH surveillance programmes in...
METHOD AND ANALYSIS

Study design and setting

This observational study will adopt the WHO practice guidelines for sentinel surveillance of suicide attempts and SH.22 We have also taken advice from academic and advocate with lived experience of SH regarding the design of this study. Sentinel surveillance sites will be set up at three NZ hospital emergency departments (EDs) and nationally in partnership with NZPSU.

Hospital sites: SH surveillance sites will be established in the EDs of Middlemore Hospital in Auckland (North Island), Dunedin Hospital in Dunedin, and Southland Hospital in Invercargill (South Island). Surveillance data will be collected for hospital presenting cases of SH and suicidal ideation among all age groups at Dunedin and Southland Hospitals and for paediatric age group (<15 years) at Middlemore Hospital. Study sites have been selected as they serve some of the most socioeconomically deprived and ethnically diverse communities who also experience relatively high burden of suicide deaths.20 Middlemore Hospital has approximately 110000 ED presentations per annum, of which there are around 800 episodes of SH in the paediatric population (up to 18 years). Middlemore is the largest of three hospitals with a 24/7 emergency department in the Auckland region which a total population of 1.7 million people. Middlemore serves large Māori and Pacific communities,11 and was selected to better understand the urgent suicide prevention opportunities for young people in these groups. Oversampling Māori and Pacific presentations is important, as they are typically undercounted in most health datasets in NZ. Findings are likely generalisable to the wider Auckland region which is very multicultural. Dunedin Hospital has 36000 ED presentations and around 1800 SH episodes per annum. Southland Hospital has around 30000 ED presentations annually.30 The Southern DHB, that operates these hospitals has experienced suicide rates significantly higher than the national average in most years since 2009 (https://www.health.govt.nz/publication/suicide-web-tool). In combination these two hospitals cover around one third of the population who live in the South Island making the findings generalisable while also allowing greater understanding of rural populations.

NZPSU (https://www.otago.ac.nz/nzpsu/index.html): The NZPSU is a surveillance system to detect rare diseases and meets NZ reporting obligations to the WHO. This has been in operation for many years31 and has previously been used to collect information on sensitive topics such as fetal alcohol syndrome and perinatal HIV exposure. SH will be added to the current list of diseases that are monitored by the NZPSU surveillance system that paediatricians are asked to complete a surveillance ‘card’ for. Study data will be collected via registered Paediatricians (approximately 200 throughout NZ) on a monthly basis, using the same data collection variables using a secure electronic survey link. This will ensure national coverage and a good sampling of smaller and regional hospitals where younger patients tend to be admitted to paediatric units due to difficulty in accessing mental health services. The number of SH cases seen by paediatricians is currently unknown as there has been no previous study of this type in NZ. The weakness of this approach is the reliance on busy paediatricians to report on cases and the inability to know (in most hospitals) what proportion of cases presenting are reported on. However, we will be able to estimate this ratio in the hospitals covered by both systems without double counting cases.

Study status: The data collection both, at the hospital sites and via the NZPSU, commenced on 1 July 2020 and will continue until 2024.

Eligible participants: The study population is outlined below:
1. Dunedin and Southland Hospitals: individuals of all ages who present with either SH or suicidal ideation.22
2. Middlemore Hospital: individuals under the age of 15 years presenting with either SH or suicidal ideation.11
3. NZPSU: individuals under the age of 15 years presenting with SH and coming under the inpatient care of a Registered Paediatrician at any public hospital in NZ.

For all sites, SH is defined as intentional self-injury or self-poisoning, irrespective of motivation or degree of suicidal intent.32 This definition incorporates non-suicidal self-injury and is used by the study team to reflect the complexity of SH in clinical settings. It recognises that reported motivation/intent may vary episode to episode and there also may be disagreement between clinical staff and service users themselves about the meaning of an act of SH.33 34 Self-poisoning is defined as intentional ingestion of more than the prescribed or recommended dose of any drug (eg, analgesics, antidepressants), and includes poisoning with non-ingestible substances (eg, household bleach), overdoses of ‘recreational drugs’, and severe alcohol intoxication where clinical staff consider such cases to be acts of SH. Self-injury is defined as any injury that has been deliberately self-inflicted (eg, self-cutting, jumping from a height, hanging, strangulation, self-immolation, gunshot). Suicidal ideation is defined as
having a suicide plan or having suicidal thoughts but not having acted on these thoughts or plans at the time of hospital presentation. Details of the method of SH will be recorded verbatim to facilitate a more nuanced understanding in addition to allowing coding to International Classification of Disease (ICD-10) codes.

Inclusion criteria
Individuals of any age group presenting to the ED at Dunedin or Southland Hospital or individuals less than 15 years of age presenting to the ED at Middlemore Hospital or to paediatricians reporting via NZPSU. The individual has engaged in either of the following:
- Self-injury or self-poisoning that is intentional and not accidental.
- Suicidal ideation only (at EDs but not via NZPSU).

Exclusion criteria
- Accidental overdose of alcohol in an individual who drinks alcohol to excess requiring hospital treatment, but without any intention to SH, and who does not combine alcohol with other methods of SH.
- Accidental overdose of illicit drugs in an individual who takes illicit drugs on a regular basis but without any intention to SH.
- Accidental overdose of prescription or over-the-counter medications by incorrectly following a prescribed dosage.
- SH in the context of significant intellectual disability or developmental disorder.

Sampling
All consecutive cases of SH or suicidal ideation presenting to EDs at Dunedin, Southland and Middlemore Hospital (<15 years) will be identified by the study staff as per the case-ascertainment definition. For NZPSU, all consecutive cases of SH (<15 years) under the inpatient care of a registered paediatricians are identified and reported by the paediatricians as per the case-ascertainment definition.

Participant recruitment
This is a surveillance study using routinely collected clinical data. There is no direct contact with patients/cases/participants.

Hospital sites
All patients who present to ED receive usual care including triage and a routine clinical assessment by ED clinicians. The study team will manually review the triage descriptions described on electronic ED lists. Cases which include terms such as suicide ideation, SH, suicide attempt, overdose, hanging, jumping, cutting will be reviewed in addition to broader terms such as abdominal pain, collapse due to unknown cause will be reviewed for inclusion. Once an episode of SH or suicidal ideation is identified, further investigation of the patient records via relevant electronic record system will be undertaken. Additionally, in Middlemore Hospital, if the electronic medical record is unclear, the study team will review relevant paper records. For all eligible participants in the hospital settings, routine clinical data will be extracted from the clinical record by a member of the research team (see Table 1) and entered directly into the Research Electronic Data Capture tool (REDCap, see further description in Data Management section of this protocol).

New Zealand Paediatric Surveillance Unit (NZPSU)
The NZPSU team will inform the study team about any reported cases of paediatric SH as notified by Registered Paediatricians on their monthly report card. The reporting paediatrician is contacted by the study team and provided a unique URL which will link them directly to the secure REDCap data capture tool. Paper-based data collection forms will be available to any paediatrician who requests this format.

Determination of SH/suicidal ideation relatedness, data quality and reliability
Initial training in case ascertainment will involve 2–3 members of the research team who will review all ED

| Table 1 | Data collection fields |
|---------|------------------------|
| 1. Sociodemographic information: NHI, gender, age, ethnicity, usual residential address, occupation |
| 2. Description of presenting complaint (narrative) |
| 3. Date and time of SH |
| 4. Date and time of hospital presentation and discharge |
| 5. How the patient arrived at the hospital (ambulance/police/own transport) |
| 6. Did the patient present with self-injury, self-poisoning, suicidal ideation |
| 7. Location of SH (home/public place) |
| 8. How did the patient harm themselves (record verbatim) |
| 9. Was alcohol and/or an illegal substance involved |
| 10. Did the patient make a statement of intention to die |
| 11. Problems/difficulties associated with the current episode of SH or suicidal ideation |
| 12. History of exposure to SH and suicide among peers/family |
| 13. Prior history of SH |
| 14. Referrals to specialist services such as social worker, ICU |
| 15. Referral for mental health assessment during this episode |
| 16. Did this episode lead to a hospital admission |
| 17. Discharge location from ED |
| 18. For cases of self-poisoning, name, amount, strength and source of substance used (Codes/categories available on request) |
presentations to each hospital study site. Each case will be reviewed by the team against the inclusion/exclusion criteria, robust discussion will ensure capture of all cases. Once accurate case ascertainment has been achieved, an ongoing review process to determine inter-rater reliability of case determinant and accurate extraction of key variables will include:

1. An independent review by a second team member of the routinely collected hospital presentation data to check accurate identification of cases for inclusion at each hospital site on a regular basis.

2. An independent review by a second team member, cross-checking the routinely collected clinical data for 10% participants registered every month at each hospital site. Cases will be selected using a random number generator. Any discrepancies in case ascertainment will be documented and resolved by the study Principal Investigator (PI).

3. An independent review, by a second researcher, of extracted key variables for a random sample of 20 identified cases over a period of 6 months, against routinely collected clinical records to check for consistency in data extraction. Cases will be selected using the random number generator in statistical software.

Intrarater (within reviewer) reliability will also be determined by comparing the determination of SH/suicidal ideation, and consistency in extraction of key variables, between the beginning and end of the data collection period. The reasons behind any inconsistencies or discrepancies between reviewers will be examined further and if necessary, cases will be re-coded.

Consent
Participant consent is not required; the study does not require patients to take part in any additional procedures nor does it affect care or treatment they receive. The study research team will not have any direct patient contact as the study is based only on routinely collected information.

Stakeholder engagement
A project advisory committee, comprising of national and international experts on suicide prevention including clinicians, academics, and those with lived experience, will be established in line with international best practice. The committee will assist in identifying issues that need to be addressed and opportunities for the surveillance system. The committee will also provide advice to the data management plan and delivering culturally relevant research outcomes.

Data management
A customised data collection mechanism, specific to this study, will be developed using REDCap. REDCap is a secure web-based application designed to support data capture for research studies. It is suitable for collecting personal information which is covered by Health Information Privacy Principles, The Privacy Act and Ethics Committee specifications that require the use of a secure survey tool. All survey data collected via REDCap will be securely stored at the University of Otago.

Any missing values or variables will be checked against records to ensure that all possible information is included. This may require the study team to revisit some case files to collect any missing information. A preliminary statistical analysis will be conducted after every 100 cases (for the initial period of 6 months) to ensure that necessary data has been captured correctly.

The WHO protocol suggests it is essential to collect personal information about participants for case ascertainment, particularly where the person represents to hospital while complying with local regulations. Potential cases will be identified using clinical record identifiers. However, once cases have been selected, all identifying information such as name, home address, National Health Index (NHI) number (a unique identifier that is assigned to every person who uses health and disability support services in NZ), and date of birth variables will be partitioned off and a unique participant code will be allocated. Thus, the study team will use a de-identified dataset for aggregated statistical analysis and subsequent publications. The study data will be solely used for addressing the research objectives and will not be used for any dissemination that may potentially identify an individual. Original data will be securely stored by the study team for a minimum of 10 years. The research data will be solely owned by the University of Otago.

Data analysis
The primary outcomes of the study are: (1) annual number of episodes of hospital-presenting SH and suicidal ideation; (2) trends in frequently used methods of SH; (3) “new” or emerging methods of SH; (4) frequently used drugs in overdose related SH; (5) variations (seasonal/demographic/temporal) in presentations of SH and suicidal ideation; and (6) average time taken between time of SH and time of ED presentation.

We would also include outcomes related to incidence rates: (1) annual rates of SH per 100,000 population for the total population, for subgroups by age and sex; (2) annual rates of SH in paediatric population (<15 years); (3) annual rates of suicidal ideation; (4) variation by gender, age, ethnicity and geographical region. The secondary outcomes will focus on identifying predictors of repetition of SH, potential gaps in SH services and systemic levers for suicide prevention.

For comparison, crude and age-specific SH rates will be calculated using the NZ Census 2018 and WHO world standard population. An analysis of repeat events will be conducted using conditional risk set analysis and Cox hazard proportional models will be used to identify independent and multivariable
predictors of repetition of SH. All the study data will be analysed using SPSS V.27.0.

ETHICS AND DISSEMINATION

This study has received approval from University of Otago Human Ethics Committee (H19/165) including the Ngāi Tahu Research Consultation Committee, to review the research plan in context to the indigenous population of NZ (Māori). Institutional approvals were provided by Counties Manukau DHB which oversees Middlemore Hospital and the Southern DHB which oversees Dunedin and Southland Hospitals. Each team member has a confidentiality agreement with the DHB.

We will publish aggregated study data to both national and international audiences via publications in peer reviewed scientific journals and conference presentations. Submission of study findings will also be made to local regulatory authorities including DHB management teams, local Suicide Prevention Coordinators and the NZ Suicide Prevention Office.

The main aim of this study is to establish multi-centre sentinel surveillance of patients presenting with SH or suicidal ideation as per the recommended WHO practice guidelines on surveillance systems and to provide up to date, detailed and robust epidemiological data which will establish trends in SH in NZ. The focus on paediatric populations at Middlemore Hospital and via the NZPSU has been informed by the lack of high-quality data for this group who have historically been excluded from studies due to relatively small numbers, the prominence of SH among those who die at a very young age by suicide and the support of a specific paediatric health funder. Hospitals were selected for study inclusion because they have large numbers of patients presenting for emergency care, allow for purposeful oversampling of indigenous youth, serve large populations. The inclusion of paediatricians reporting on presentations around the country allows triangulation of data, and a greater understanding of the epidemiology of SH beyond large hospital catchment areas. In doing so, the study findings will inform suicide prevention efforts to support the development of evidence-based interventions and workforce development in a way that is most likely to reduce SH and ultimately suicide deaths.

PUBLIC AND PATIENT PARTICIPATION

The research question and priority is part of a broader programme of work in which the voice of those with living or lived experience of suicidal distress is central and based on active collaboration with lived experience academics. This study connects with systematic reviews by this group on service user experiences of SH and staff attitudes towards those seeking hospital care after SH. Contributors SF conceptualised the study and developed the study protocol with the other co-investigators—KMS, SH and GM. RTM provided inputs to the study design. LH and VS assisted in finalisation of the protocol and drafted this manuscript.

Funding This study is funded by Cure Kids (grant number—3592) for paediatric presentations of SH, and the James Hume Bequest Fund, Dunedin School of Medicine, University of Otago, for presentations at Dunedin and Southland Hospitals.

Competing interests None declared.

Patient and public involvement Patients and/or the public were involved in the design, or conduct, or reporting, or dissemination plans of this research. Refer to the Methods section for further details.

Patient consent for publication Not applicable.

Provenance and peer review Not commissioned; externally peer reviewed.

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