

Diabetes case finding in the emergency department, using HbA1c: an opportunity to improve diabetes detection, prevention, and care

Tien-Ming Hng,^{1,2} Amanda Hor,^{1,3} Sumathy Ravi,¹ Xiaoqi Feng,^{1,3,4} Jaime Lin,¹ Thomas Astell-Burt,^{5,6} David Chipps,¹ Mark McLean,^{1,2} Glen Maberly^{1,2}

To cite: Hng T-M, Hor A, Ravi S, *et al.* Diabetes case finding in the emergency department, using HbA1c: an opportunity to improve diabetes detection, prevention, and care. *BMJ Open Diabetes Research and Care* 2016;**4**:e000191. doi:10.1136/bmjdr-2015-000191

Received 31 December 2015
Revised 19 April 2016
Accepted 21 April 2016

ABSTRACT

Objective: We assessed the efficacy of routine glycosylated hemoglobin (HbA1c) testing to detect undiagnosed diabetes and prediabetes in an urban Australian public hospital emergency department (ED) located in an area of high diabetes prevalence. **Methods:** Over 6 weeks, all patients undergoing blood sampling in the ED had their random blood glucose measured. If ≥ 5.5 mmol/L (99 mg/dL), HbA1c was measured on the same sample. HbA1c levels $\geq 6.5\%$ (48 mmol/mol) and 5.7–6.4% (39–46 mmol/mol) were diagnostic of diabetes and prediabetes, respectively. Hospital records were reviewed to identify patients with previously diagnosed diabetes.

Results: Among 4580 presentations, 2652 had blood sampled of which 1267 samples had HbA1c measured. Of these, 487 (38.4%) had diabetes (either HbA1c $\geq 6.5\%$ or a prior diagnosis), and a further 347 (27.4%) had prediabetes. Among those with diabetes, 32.2% were previously undiagnosed.

Conclusions: Routine HbA1c testing in the ED identifies a large number of people with undiagnosed diabetes and prediabetes, and provides an opportunity to improve their care.

INTRODUCTION

Diabetes has traditionally been diagnosed by the 75 g oral glucose tolerance test (75 g OGTT), but recent adoption of glycosylated hemoglobin (HbA1c) as a diagnostic tool for high-risk individuals has simplified the diagnostic process.^{1–3} Underdetection remains a major barrier to prevention of diabetes and associated complications. Targeted HbA1c testing of high-risk populations could enhance the detection of diabetes and prediabetes. The purpose of this study was to test this hypothesis in patients presenting to a hospital emergency department (ED).

METHODS

The study was conducted over 6 weeks in the ED of Blacktown-Mt Druitt Hospital, located in the western suburbs of Sydney, Australia. This is

Key messages

- We demonstrate that glycosylated hemoglobin measurement undertaken in the setting of an emergency department is an effective and feasible means of finding cases of diabetes and prediabetes particularly in an area known to have a high prevalence of diabetes.
- Approximately a third of patients tested were previously unaware that they had diabetes.
- Earlier detection of prediabetes provides an opportunity to introduce measures that may prevent progression to diabetes.

an area of high diabetes prevalence.⁴ Opportunistic blood glucose measurements were undertaken in all non-pregnant individuals ≥ 16 years of age who had blood collected after presenting to the ED, irrespective of the presenting problem. HbA1c was automatically measured if the random glucose was ≥ 5.5 mmol/L (99 mg/dL). This cut-off level was chosen based on another Australian ED study⁵ and a substudy that we undertook (unpublished). In this substudy, HbA1c measurements were undertaken in 50 patients presenting through ED with random glucose levels < 5.5 mmol/L (< 99 mg/dL). Only three individuals were noted to have HbA1c levels $\geq 6.5\%$ and the glucose cut-off of ≥ 5.5 mmol/L (99 mg/dL) was deemed reasonable.

In patients who re-presented within the study period, HbA1c testing was undertaken only on the first presentation. As per American Diabetes Association recommendations, diabetes was diagnosed if HbA1c $\geq 6.5\%$ (48 mmol/mol) and prediabetes was diagnosed if HbA1c was between 5.7% and 6.4% (39–46 mmol/mol).¹ HbA1c was measured using a turbidimetric inhibition immunoassay on a Siemens Dimension Vista 1500 platform. Coefficient of variation was 2.9% and 2.4%, at a HbA1c level of 5.7% and



CrossMark

For numbered affiliations see end of article.

Correspondence to

Dr Tien-Ming Hng; tien-ming.hng@health.nsw.gov.au

10.1%, respectively. Demographic data, previous diagnosis of diabetes, and coding information were obtained from the hospital's database. This study was discussed with and approved by the Human Research Ethics Committee of the Western Sydney Local Health District.

RESULTS

There were 4580 presentations to the ED. Using the methodology described, 1267 HbA1c measurements were available (figure 1). Female patients accounted for 47.3% of results obtained. Three hundred and seventy-nine HbA1c tests were not performed due to multiple presentations, insufficient blood sampling or issues related to the automation of HbA1c measurement. In this cohort, diabetes was present in 38.4% (HbA1c \geq 6.5% or a prior diagnosis if HbA1c $<$ 6.5%; table 1).

Of the patients identified with diabetes, 45.2% were female and 32.2% (157/487) were newly diagnosed. Of newly diagnosed patients, 61.8% had mild diabetes (HbA1c of 6.5–6.9% or 48–52 mmol/mol). Three hundred and forty-seven (27.4%) of HbA1c sampled patients had levels consistent with prediabetes (HbA1c 5.7%–6.4% or 39–46 mmol/mol). The sampling protocol resulted in HbA1c being measured in 59% (n=751) of admitted patients, and, of these, diabetes was present in 42.4%. Analysis of coding information revealed that 28% of admissions who were known to have diabetes (either by HbA1c assessment or previously noted in the medical record) were not coded for the diagnosis of diabetes. Of this group, 11% were previously known to have diabetes, 81% were newly diagnosed, and the remaining 8% were coded as impaired glucose regulation.

DISCUSSION

The key finding from this ED study is that almost two in every five patients tested on available HbA1c samples had diabetes. This is consistent with other studies that indicate a high risk of diabetes in Western Sydney's ethnically diverse, low socioeconomic population.^{4 6} Importantly, one in three patients found to have diabetes were previously unaware of their condition. Additionally, a further 27.4% had prediabetes, meaning that 65.7% of patients tested with HbA1c had either

diabetes or prediabetes. The use of HbA1c for diabetes case finding in the ED had a high yield in our patient population, but these findings may not necessarily be replicated in locations with a lower prevalence of diabetes. In an Australian tertiary referral hospital's ED in Melbourne, Jelinek *et al*⁷ noted similar rates of dysglycemia but, despite a high diabetes prevalence, the authors did not find screening a feasible exercise due to their reliance on a 75 g OGTT to confirm the diagnosis.

Diagnosis of diabetes was not confined to the mild end of the spectrum—with a severely elevated HbA1c level ($>$ 9% or 75 mmol/mol) seen in 7.3% of the entire cohort, and in 10.2% of newly diagnosed patients. Rapidly identifying admitted patients with poor glycemic control utilizing a test on a single blood sample that does not require any pretest preparation provides an ideal opportunity for intervention by hospital diabetes services. The majority of the newly diagnosed patients had early stage diabetes with HbA1c levels of 6.5–6.9% (48–52 mmol/mol), providing the ideal opportunity for early intervention to take place before the onset of complications. Almost a third of the individuals diagnosed as having diabetes were not aware of their diagnosis, reflecting the hidden burden of diabetes in our community. This may indicate that current screening practices in primary care are insufficient and further supports opportunistic HbA1c testing in individuals presenting to hospital, a population seemingly enriched with cases of diabetes. In Australia, universal health insurance (Medicare) covers the cost of HbA1c testing to diagnose diabetes in 'high-risk' individuals.⁸ Our data suggest that a high proportion of patients presenting to ED and requiring blood collection would meet this definition. Use of blood glucose alone for diabetes diagnosis is problematic because of the presence of 'stress hyperglycemia', a pitfall avoided by use of the HbA1c test.

The use of HbA1c to identify prediabetes is supported by the American Diabetes Association.¹ Progression to diabetes may be prevented if these individuals were provided with achievable lifestyle intervention. This has been shown to be beneficial in prediabetes in several studies such as the Diabetes Prevention Program⁹ and the Finnish Diabetes Prevention Program.¹⁰ One such lifestyle intervention programme in Australia is the government run *Get Healthy* initiative.¹¹

A novel feature of this study is the use of HbA1c to diagnose diabetes in a non-primary care setting. Previous studies have demonstrated that diabetes screening in an outpatient and a GP setting is feasible.¹² The visit to ED is an opportunity for us to detect diabetes in individuals who infrequently seek routine medical care, and who may otherwise go undetected. An earlier ED study of a tertiary referral hospital in Adelaide, Australia, noted a prevalence of undiagnosed diabetes of 11%,⁵ compared with 32.2% in this study. This may be because of the highly multicultural population in our study, compared with the overwhelmingly Caucasian population of Adelaide (table 2).

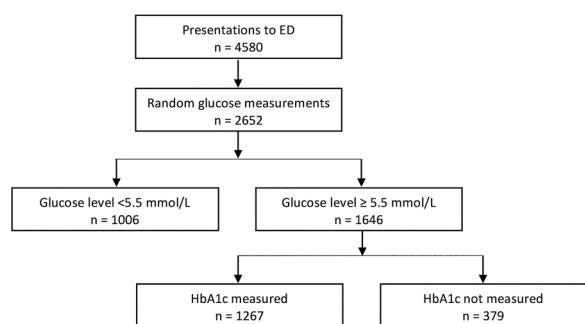


Figure 1 Study profile. ED, emergency department; HbA1c, glycated hemoglobin.

Table 1 Levels of dysglycemia based on glycated hemoglobin (HbA1c) measurements with subset analysis of individuals newly diagnosed with diabetes

Status	HbA1c (%)		All patients		Admitted patients	
	Per cent	mmol/mol	N	Per cent	n	Per cent
No diabetes	<5.7	<39	433	34.2	244	32.5
Prediabetes	5.7–6.4	39–46	347	27.4	188	25.0
*Known diabetes	<6.5	<48	81	6.4	66	8.8
Diabetes	6.5–6.9	48–52	138	10.9	80	10.7
	7.0–7.9	53–63	117	9.2	76	10.1
	8.0–8.9	64–74	58	4.6	43	5.7
	≥9.0	≥75	93	7.3	54	7.2
Total			1267	100	751	100
<i>Subset analysis</i>						
New diabetes	6.5–6.9	48–52	97	61.8	52	80
	7.0–7.9	53–63	37	23.6	10	15.4
	8.0–8.9	64–74	7	4.5	2	3.1
	≥9.0	≥75	16	10.2	1	1.5
Total			157	100	65	100

*Previously diagnosed with diabetes as noted in the medical record.

For the purpose of funding and performance review, all clinical data on admitted patients are coded using the International Classification of Disease codes. In our cohort, 28% of patients diagnosed with diabetes based on an elevated HbA1c were not coded as having diabetes, leading to an underestimate in the coded data, and potential underfunding relative to true case complexity. Coding inaccuracies can also result in an overestimation of the number of newly diagnosed individuals as this calculation is dependent on whether they had previously been coded as having diabetes.

The use of HbA1c to diagnose diabetes has limitations as any factor affecting the quantity or quality of the hemoglobin molecule can result in measurement inaccuracies. This has not been accounted for in our study, but would tend to result in an underestimate of diabetes cases. Sampling was only undertaken in individuals who had blood collected, and this introduces a source of

bias. It could be argued that the pickup rate for diabetes could be further improved by advocating routine testing in all individuals presenting to the ED.

Another limitation of using HbA1c to diagnose diabetes is the existence of ethnic variability in the measurement that is not explained by glycemic state, structural abnormalities or quantity of the hemoglobin molecule.^{15–17} This is relevant in our multiethnic population where the diagnosis of diabetes can be based on a slight elevation in the HbA1c level. Our study used glucose level as the initial screening step and this would have limited the diagnostic inaccuracies that might have occurred had this not been undertaken.

Diabetes case finding in the ED is justified as there is a significant population with known and undiagnosed diabetes. Following the findings of this pilot, it is intended that routine HbA1c testing will be the norm in the ED at Blacktown Hospital.

Table 2 Comparison of the top 10 cultural groups (based on country of birth) between the study population and the postcodes within which the respective hospitals are located

Study population		Postcode 2148 (Sydney) ¹³		Postcode 5041 (Adelaide) ¹⁴	
	Population (%)		Population (%)		Population (%)
Australia	50.1	Australia	53.2	Australia	78.4
India	5.1	Born elsewhere	9.4	UK	4.6
Philippines	4.9	India	7.5	Country of birth not stated	3.7
England	2.8	Country of birth not stated	4.8	Born elsewhere	2.7
Fiji	2.3	Philippines	4.7	Greece	1.3
Sri Lanka	2.0	UK	2.3	India	1.1
Malta	2.0	New Zealand	2.2	China	1.1
New Zealand	1.7	Fiji	2.0	New Zealand	1.0
Italy	1.6	China	1.9	Italy	0.8
Egypt	1.3	Malta	1.1	Germany	0.7

Source: Australian Bureau of Statistics Census (2011) data.

Author affiliations

¹Western Sydney Local Health District, Blacktown Hospital, Blacktown, New South Wales, Australia

²School of Medicine, Western Sydney University, Parramatta, New South Wales, Australia

³School of Health and Society, University of Wollongong, Wollongong, New South Wales, Australia

⁴Early Start Research Institute, University of Wollongong, Wollongong, New South Wales, Australia

⁵School of Science and Health, Western Sydney University, Parramatta, New South Wales, Australia

⁶School of Geography and Geosciences, University of St Andrews, St Andrews, UK

Acknowledgements The authors acknowledge the assistance of Pathology West for HbA1c assessment and data retrieval, the Health Information Record Service for coding information, and the Executive of Blacktown/Mt Druitt Hospital for supporting the study, particularly for the costs of HbA1c measurements.

Contributors T-MH assisted in study design and database development, contributed to the discussion, researched and analyzed data, and wrote the manuscript. AH assisted in study design, researched, collected and analyzed data, and wrote the manuscript. SR collected data and assisted with correspondence. XF analyzed data, contributed to the discussion, and reviewed/edited the manuscript. JL collected data. TA-B contributed to the discussion and reviewed/edited the manuscript. DC contributed to the discussion and concept. MM contributed to the study design and discussion, and reviewed/edited the manuscript. GM conceived the idea, secured institutional support, contributed to the discussion, and reviewed the manuscript.

Competing interests None declared.

Ethics approval Human Research and Ethics Committee of the Western Sydney Local Health District.

Provenance and peer review Not commissioned; externally peer reviewed.

Data sharing statement No additional data are available.

Open Access This is an Open Access article distributed in accordance with the Creative Commons Attribution Non Commercial (CC BY-NC 4.0) license, which permits others to distribute, remix, adapt, build upon this work non-commercially, and license their derivative works on different terms, provided the original work is properly cited and the use is non-commercial. See: <http://creativecommons.org/licenses/by-nc/4.0/>

REFERENCES

- American Diabetes Association. 2. Classification and diagnosis of diabetes. *Diabetes Care* 2015;38(Suppl 1):S8–16.
- d'Emden MC, Shaw JE, Colman PG, *et al*. The role of HbA1c in the diagnosis of diabetes mellitus in Australia. *Med J Aust* 2012;197:220–1.
- World Health Organization Consultation. Use of glycated haemoglobin (HbA1c) in the diagnosis of diabetes mellitus. *Diab Res Clin Pract* 2011;93:299–309.
- Astell-Burt T, Feng X, Kolt GS, *et al*. Understanding geographical inequities in diabetes: multilevel evidence from 114,755 adults in Sydney, Australia. *Diab Res Clin Pract* 2014;106:e68–73.
- Valentine NA, Alhawassi TM, Roberts GW, *et al*. Detecting undiagnosed diabetes using glycated haemoglobin: an automated screening test in hospitalised patients. *Med J Aust* 2011;194:160–4.
- Australian Diabetes Map Australia: Diabetes Australia; [updated 6/10/2015]. <https://http://www.diabetesaustralia.com.au/tools-e-learning>
- Jelinek GA, Weiland TJ, Moore G, *et al*. Screening for type 2 diabetes with random finger-prick glucose and bedside HbA1c in an Australian emergency department. *Emerg Med Australas* 2010;22:427–34.
- The November 2014 Medicare Benefits Schedule: Department of Health, Australian Government. <http://www.health.gov.au/internet/mbsonline/publishing.nsf/Content/Downloads-2014-11>
- Knowler WC, Barrett-Connor E, Fowler SE, *et al*. Reduction in the incidence of type 2 diabetes with lifestyle intervention or metformin. *N Engl J Med* 2002;346:393–403.
- Tuomilehto J, Lindstrom J, Eriksson JG, *et al*. Prevention of type 2 diabetes mellitus by changes in lifestyle among subjects with impaired glucose tolerance. *N Engl J Med* 2001;344:1343–50.
- NSWHealth. *Get Healthy Information and Coaching Service NSW*. Australia: NSW Health, 2015 [10 Oct 2015]. <http://www.gethealthynsw.com.au/>
- Klein Woolthuis EP, de Grauw WJ, van Gerwen WH, *et al*. Yield of opportunistic targeted screening for type 2 diabetes in primary care: the diabscreen study. *Ann Fam Med* 2009;7:422–30.
- 2011 Census of Population and Housing 2148 (POA2148) 29 sq Kms: Australian Bureau of Statistics; 2011 [updated 17 March 2016]. Census data from the Australian Bureau of Statistics]. http://www.censusdata.abs.gov.au/census_services/getproduct/census/2011/communityprofile/POA2148
- 2011 Census of Population and Housing 5041 (POA5041) 6.2 sq Kms: Australian Bureau of Statistics; 2011 [updated 17 March 2016]. Census data from the Australian Bureau of Statistics]. http://www.censusdata.abs.gov.au/census_services/getproduct/census/2011/communityprofile/POA5041
- Hare MJ, Magliano DJ, Zimmet PZ, *et al*. Glucose-independent ethnic differences in HbA1c in people without known diabetes. *Diabetes Care* 2013;36:1534–40.
- Herman WH, Cohen RM. Racial and ethnic differences in the relationship between HbA1c and blood glucose: implications for the diagnosis of diabetes. *J Clin Endocrinol Metab* 2012;97:1067–72.
- Venkataraman K, Kao SL, Thai AC, *et al*. Ethnicity modifies the relation between fasting plasma glucose and HbA1c in Indians, Malays and Chinese. *Diabet Med* 2012;29:911–17.