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Research Article

Importance of Investigating Mild Hyperglycaemia Using Oral Glucose Tolerance Test in the Setting of Acute Coronary Syndrome: Prospective Investigational Study

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Abstract

The aim of this article is to estimate the prevalence of true dysglycaemic syndromes (DS) (includes presence of diabetes mellitus [DM], impaired fasting glucose [IFG] and impaired glucose tolerance [IGT]) in patients with acute coronary syndromes (ACS) and mild fasting hyperglycaemia from an area in New Zealand with the country's highest prevalence of dysglycaemia. Cardiac risk factor data and laboratory findings for 565 consecutive ACS patients were obtained from the hospital database between January 2007 and August 2008. On admission, 175 patients had DS and 168 were normoglycaemic (FBG <5.6 mmol/L). The remaining 222 patients (with previously unknown glycaemic status) had mild fasting hyperglycaemia (FBG 5.6-6.9 mmol/L) on admission. This group with mild fasting

hyperglycaemia were offered oral glucose tolerance testing (OGTT) >3 months post-discharge to eliminate cases of stress hyperglycaemia. 121 patients underwent OGTT and 41 patients (a third) were found to have true DS (DM in 13, IGT or IFG in 28). Patients who did not undergo OGTT due to a variety of reasons (n=101) had significantly higher body mass index, waist circumference, and higher concentrations of triglycerides and low-density lipoprotein cholesterol, and therefore might be at a higher risk for true DS than those tested. The conclusion is that a third of the OGTT cohort had true DS and therefore even mild hyperglycaemia in this setting should be carefully investigated.

Keywords: Acute coronary syndrome; diabetes mellitus; dysglycaemic syndromes; oral glucose tolerance test.

Introduction

There is increasing recognition of the significant relationship between diabetes and cardiovascular disease, and the contribution that each of these conditions makes to the risk of morbidity and mortality from both diseases (Rydén et al 2013; Gholap et al 2012; De Caterina et al 2010). Coronary artery disease is common in patients with true dysglycaemic syndromes (DS) which includes diabetes mellitus (DM), impaired fasting glucose (IFG), and impaired glucose tolerance (IGT) (Gerstein et al 1996). However, DS are often not diagnosed in patients presenting with acute coronary syndromes (ACS) (Conaway et al 2005). Patients with undiagnosed DS admitted with ACS have a 1-year mortality rate which is similar to that in patients with proven DS (Schiele et al 2006). Establishing a diagnosis of DS in

patients admitted to hospital with ACS is therefore important because appropriate management has the potential to improve long-term prognosis (Gerstein et al 1996, Conaway et al 2005, Schiele et al 2006, Tenerz et al 2003).

Caution is required when diagnosing DS during ACS for a number of reasons. A significant proportion of patients with no previous DS may have stress hyperglycaemia that normalises once the initial stress associated with ACS has lessened (Goran Koracevic et al 2006). Previous studies have shown that stress hyperglycaemia is common in acutely ill patients (such as those with ACS) who did not have a previous diagnosis of diabetes (McCowan et al 2001, Capes et al 2000). This results from stress-related increases in catecholamine and cortisol concentrations, stimulating glucose production by gluconeogenesis and

glycogenolysis (Goran Koracevic et al 2006). Therefore, to identify patients with true DS, it is important to test patients once the initial stress period is over.

Current screening criteria for DS include two consecutive fasting blood glucose (FBG) measurements, random blood sugar samples, an oral glucose tolerance test (OGTT) and, more recently, glycosylated haemoglobin (HbA1c). Whilst HbA1c is a measurement of chronic hyperglycaemia it may not identify IGT and IFG (American Diabetes Association 2013). Research has emphasised the benefits of OGTT over FBG in terms of both reliability and sensitivity (Wallender et al 2008). In fact, FBG may miss DS in up to 70% of patients (Jessani et al 2007, Henareh et al 2004). Data suggest that IGT, but not IFG, is a risk factor for cardiovascular diseases, although IFG is associated with an

increased risk of progression to IGT (Tominaga et al 1999). Therefore, the OGTT is an important tool for the diagnosis of patients with DS.

The aim of this study was to estimate the prevalence of true DS in patients with ACS and mild fasting hyperglycaemia.

Patients and Methods

Data for 565 consecutive patients with confirmed ACS who presented to Middlemore Hospital, Auckland, New Zealand, between 1 August 2007 and 31 August 2008 were identified from the Acute PREDICT database. Middlemore Hospital is the base hospital for the South Auckland region and serves approximately half a million people. Acute PREDICT is a prospectively collected

electronic registry of all patients admitted to the coronary care unit. Myocardial infarction was diagnosed according to established criteria, including changes in cardiac biomarkers, ischaemic symptoms, ECG changes and/or cardiac imaging. Diagnosis of unstable angina was based on acute worsening of angina pectoris but with negative or insignificant increases in troponin concentrations. The study protocol was approved by the Northern X Regional Ethics Committee (NTX/08/101/EXP).

Clinical, biochemical and demographic data were obtained from the Acute PREDICT database, and additional information (past medical history, type of ACS, admission glucose and FBG results during the most recent ACS, and prior OGTT history) were extracted from the hospital online patient information system.

Three hundred and forty-three patients were excluded from OGTT testing (group 1); 175 had a prior diagnosis of DS and 168 had a normal fasting blood sugar (FBS), at the time of the coronary event. Two hundred and twenty-two patients were eligible for an OGTT test (group 2) as per criteria (FBS 5.6-6.9 mmol/L). All of these patients were >3 months after their acute cardiac event and were well in the community and therefore unlikely to have persistent stress induced hyperglycaemia. These 222 patients were contacted via their general practitioner (GP) and 121 agreed to undergo an OGTT (Figure 1). OGTT was not performed in 101 patients; reasons are detailed in Figure 1. The results of testing were forwarded to GPs to provide ongoing care. The standardised OGTT was performed (75 g glucose load after a fasting period of 9-12 hours). FBG and 2-hour post-load blood glucose (2h-BG) tests were also undertaken. Analysis of all

samples was performed at a central laboratory (Diagnostic Medlab) using the Hexokinase UV method with Roche Diagnostics reagents via the Roche Diagnostic Hitachi Modular method. The calibrator was analyser specific and standardised against the isotope dilution mass spectrometry (ID MS) method.

OGTT data were interpreted using the World Health Organisation definition of IFG and the American Diabetes Association definition of IGT. Normal ranges were defined as follows: normal glucose tolerance (NGT)/normoglycaemia = FBG <5.6 mmol/L and 2h-BG <7.8 mmol/L; IFG = FBG 5.6-6.9 mmol/L; IGT = 2h-BG 7.8-10.9 mmol/L; DM = FBG \geq 7.0 mmol/L or 2h-BG \geq 11.0 mmol/L. Mild fasting hyperglycaemia was defined as FBG 5.6-6.9 mmol/L.

Statistical Analysis

Continuous variables were presented as mean values with 95% confidence intervals; categorical variables were summarised as frequencies and percentages. Two-sample student t-tests were used to assess the mean differences between groups when data were normally distributed and Mann Whitney U tests were used for data with skewed distributions. Chi square test was used to assess if there were differences in the frequency distributions of categorical variables between groups.

Multiple logistic regression analysis was used to examine the differentiating factors for patients who undertook OGTT and did versus did not have dysglycaemia, in the presence of confounding variables. Variables were included in the multiple regression

models if the p-value was ≤ 0.25 in the univariate logistic regression analysis. The final model was chosen based on the Akaike information criteria (AIC) and the estimates were represented as odds ratios (OR) with 95% confidence intervals. The software SAS version 9.3 was used to carry out the statistical analyses. All tests were two tailed and a p-value of < 0.05 was considered statistically significant.

Results

Cohort and Glycaemic Status

In the overall cohort, the prevalence of DS was 175/565 (31%) prior to OGTT. On OGTT a dysglycaemic syndrome was identified in a further 41 of the 121 patients (DM n=13, IGT n= 24 or IFG

n=4) (Figure 1). This increased the prevalence of diagnosed DS in the overall cohort to 222/565 (39%).

OGTT Results (Table 1)

Body mass index (BMI) and waist circumference were significantly higher in the 41 patients with DS comparing to those who had normal OGTT results (BMI: 30 vs. 27 kg/m²; waist circumference: 104 vs. 98 cm). Over half the South Asian, Maori and Pacific patients who were tested had DS compared with only a quarter of European patients (Table 1).

Table 1. Demographic and Clinical Data for the Study Population

Please See Table 1 in PDF Version

In patients who did not accept the offer of an OGTT, BMI and waist circumference were significantly higher than those in patients who did (median [range] values of 31 [26-35] vs. 27 [25-30], respectively; $p=0.002$). Therefore, patients who did not have the OGTT may be at greater risk of developing DS. Patients who did not have OGTT also had a higher total cholesterol: high density lipoprotein (HDL) cholesterol ratio (4.8 [3.4-5.6] vs. 4.0 [3.2, 4.9] in those who did have OGTT; $p=0.02$). In addition, patients of Maori and Pacific Island ethnicities were least likely to undergo OGTT, with a >50% non-participation rate.

On multiple regression analysis fasting blood glucose, waist circumference, and Maori, Pacific and South Asian ethnicity were the independent factors significantly associated with DS (Table 2).

Table 2. Independent Predictors of a Diagnosis of Dysglycaemic Syndrome on Oral Glucose Tolerance Testing, in Acute Coronary Syndrome Patients with Mild Fasting Hyperglycaemia

Independent predictors	Odds ratio	95% confidence interval	p value
Fasting triglycerides	1.1	0.8, 1.4	0.59
Fasting blood glucose	1.7	1.2, 2.3	0.004
Waist circumference (per 10 cm increase)	1.5	1.1, 2.1	0.008
Maori or Pacific or South East Asian vs others	2.7	1.1, 6.7	0.03

Discussion

In a consecutive cohort of patients presenting with ACS, 39% (222/565) had mild fasting hyperglycaemia. Of those who agreed to an OGTT in the community >3 months later, one-third had a final diagnosis of DS. By combining the patients with known DS at the index admission with those diagnosed on subsequent OGTT the prevalence of DS in the whole ACS cohort was 39% (222/565). This is likely to be an underestimate because only just over a half those with mild hyperglycaemia who had an OGTT. On the basis of associated risk factors, those who were untested are likely to have at least as high a rate of DS than those tested with an OGTT. By extrapolation of the DS prevalence in those tested to those not tested the prevalence of DS in the whole ACS cohort could be as high as 45%.

DS are very common in patients presenting with ACS in South Auckland, New Zealand, and the current data are comparable to similar studies from other countries. In a prospective cohort study conducted in the US, a total of 57% of patients with ACS and no previously-recognized diabetes mellitus were shown to have abnormal glucose metabolism using a simple FPG test (Conaway et al 2005).

A significant number of patients with indeterminate FBG 5.6-6.9 mmol/L results in hospital were not followed up in our study and are therefore unlikely to have been provided with early advice and treatment. In the study by Conaway et al (2005), approximately 66% of those who met criteria for new diabetes were not diagnosed or treated as such by the physicians who managed them (Conaway et al 2005).

Patients with elevated FBG on admission may be followed up using either FBG or OGTT on discharge to identify previously undiagnosed DS (Norhammer et al 2002). The OGTT has been found to be effective and reliable in evaluating glycaemic status compared with FBG, which may miss up to 70% of cases (Jessani et al 2007). Another study found that 2-hour post prandial blood glucose during admission, as opposed to FBG, predicted DS (Srinivas-Shankar et al 2008). In our study, 41 ACS patients with DS would have remained undiagnosed if OGTT had not been performed.

These findings are consistent with a previous study (Wallander et al 2008) which included 61 patients (62% male; mean age 64 ± 12.5 years), of whom 70% were White European and 30% were South Asian. Mean plasma glucose level on admission was

6.9±1.7 (range 5.8-8.1) mmol/L. Newly-diagnosed DM and IGT were detected in 31% (95% confidence interval [CI], 10%-52%) and 33% (95% CI, 12%-53%) of patients, respectively, by OGTT.

In the new diagnostic criteria of DM, HBA1c has a central role because it is convenient that the patient does not need to be in a fasting state for measurement (American Diabetes Association 2013). However, there is no perfect concordance between HBA1c and glucose based tests (American Diabetes Association 2013) and HBA1c may not detect other forms of dysglycaemia detected by OGTT. At the time the study was conducted, routine use of HBA1c for diagnosis of diabetes status was not recommended so we are unable to report the incidence of diabetes and IGT using the new definitions. HbA1c measurement is of limited value in differentiating isolated IFG, IGT and DM in subjects with IFG. It

can not be used to identify which subjects with IFG do not require an OGTT (Likhari et al 2008). It is important to detect all forms of dysglycemia in this high-risk population as appropriate management is likely to be beneficial (Gerstein et al 1996, Conaway et al 2005, Schiele et al 2006, Tenerz et al 2003; Rydén et al 2013). Newly-updated guidelines recommend that patients with cardiovascular disease are investigated for the presence of glucose metabolism disorders. It is suggested that this screening is initiated with HbA1c and FPG, and that an OGTT be added if the result of the initial testing are inconclusive (Rydén et al 2013). In patients with ACS, the recommended appropriate screening method is an OGTT, performed at least 4 to 5 days after the initial coronary event to minimize the risk of false positive results (Rydén et al 2013). Admission blood glucose was found to be strongly associated with mortality in patients with myocardial

infarction (Gholap et al 2012) and current evidence on prognostic role of hyperglycaemia in ACS allows us to suggest its direct involvement in short term complications(De Caterina et al 2010).

Our study has a number of limitations. Firstly, only 121 of the 222 patients eligible to undergo OGTT did so (reasons are listed in Figure 1). For most patients, the reason for nonparticipation was either moving out of the area or unknown. Secondly, patients with normal FBG were not requested to undergo a follow-up OGTT and some of these may have subsequently developed DS. Interestingly, patients who did not have an OGTT appeared to have a higher prevalence of features of metabolic syndrome. Therefore, the prevalence of DS in our study may have been underestimated.

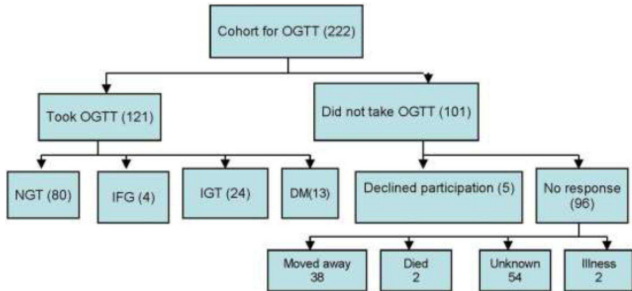


Figure 1. Flow of patients offered oral glucose tolerance testing (OGTT). NGT, normal glucose tolerance; IFG, impaired fasting glucose; IGT, impaired glucose tolerance; DM, diabetes mellitus.

A FBG test is insufficient for the screening and diagnosis of DS in patients with ACS. Routine use of OGTT in the recovery phase helps to dichotomise the large number of patients with mild hyperglycaemia at the time of admission into those with and without DS. OGTT testing in these patients with indeterminate FBG reclassifies approximately one-third as having definite DS. Further study is required to better understand how to integrate OGTT testing into the recently-adopted strategy of routine HBA1c testing.

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Conflicts of Interest

The authors have no conflicts of interest to report.

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