

# Impaired Fasting Glucose, Diabetes Mellitus and Coronary Risk Factors

KIERTIJAI BHURIPANYO, M.D.\*,  
NITHI MAHANONDA, M.D.\*,

CHARN SRIRATANASATHAAVORN, M.D.\*,  
CHARUWAN KANGKAGATE, M.S. (Biostat)\*,

ONGKARN RUANGRATANAAMPORN, M.D.\*,  
WATTANA LEOWATTANA, M.D.\*\*,

CHUNHAKASEM CHOTINAIWATTAKUL, M.D.\*,  
SUPHACHAI CHAITHIRAPHAN, M.D.\*

## Abstract

The authors conducted a prevalence survey of impaired fasting glucose and diabetes mellitus in 3,615 Shinawatra employees, and we also determined various risk factors of coronary artery disease such as blood pressure level, body mass index and serum lipids. The prevalence of impaired fasting glucose and diabetes mellitus were 1.7 per cent and 0.8 per cent respectively. The prevalences were more common in males and increased with increasing age. Coronary risk factors were higher in impaired fasting glucose (IFG) and diabetes mellitus (DM) when compared with normal glucose levels. There were also significant differences between impaired fasting glucose and diabetes mellitus, except for pulse pressure, serum cholesterol level, LDL-cholesterol level and HDL-cholesterol level.

**Key word :** Impaired Fasting Glucose, Diabetes Mellitus, Coronary Risk Factors

**BHURIPANYO K, RUANGRATANAAMPORN O, MAHANONDA N, et al**  
**J Med Assoc Thai 2000; 83 (Suppl. 2): S146-S152**

## Background and Rationale

In 1997, the American Diabetes Association (ADA) published criteria for the diagnosis and classification of diabetes mellitus and related glucose disturbances(1). In the absence of weight loss, polyuria, polydipsia, and a blood glucose concentra-

tion 2 hours after load of at least 11.1 mmol/L, these new criteria rely primarily on fasting glucose values. The advantage of the ADA criteria is that they do not require oral glucose tolerance testing with 2 hour glucose measurements, as set forth by the WHO criteria(2). By simplifying the diagnosis

\* Her Majesty Cardiac Center,

\*\* Department of Clinical Pathology, Faculty of Medicine Siriraj Hospital, Mahidol University, Bangkok 10700, Thailand.

of diabetes mellitus, ADA hoped that the diagnosis of glucose metabolic disturbances would be simpler and less of a burden to patients, more readily accepted by the medical community, and would lead to earlier detection and treatment of individuals with diabetes mellitus.

The fasting ADA criteria establish a fasting glucose concentration of at least 7.0 mmol/L for the diagnosis of diabetes mellitus, which is lower than the corresponding WHO criteria of at least 7.8 mmol/L. This lower cutoff point was chosen because the microvascular eye and kidney diseases that are unique to diabetes mellitus begin to become more prevalent at this concentration. The fasting ADA criteria also define a new class of glucose disturbance called impaired fasting glucose (fasting glucose 6.1-6.9 mmol/L), which differs from the WHO category of impaired glucose tolerance (fasting glucose <7.8 mmol/L and 2 hour glucose >7.7 mmol/L and less than 11.1 mmol/L). Several studies had also shown that impaired glucose tolerance was associated with increased prevalence of coronary risk factors and cardiovascular mortality(3-26), but little is known concerning impaired fasting glucose. Our study examined the prevalence of coronary risk factors in impaired fasting glucose and diabetes mellitus in a population survey of 3,615 Shinawatra employees.

## MATERIAL AND METHOD

A cluster sampling survey was performed amongst Shinawatra employees, who were relatively young, highly educated and have high socioeconomic status. A self-administered questionnaires concerning demographics, education, family income, presence of heart disease and risk factors, physical activity, stress, alcohol consumption and angina history using the Rose and Blackburn questionnaire was conducted.

Standing height was measured with the subject in bare feet, back square against the wall and eyes looking straight ahead. Weight was measured in undergarments, using balanced scales to the nearest 200 grams. The scale was standardized to 0 before each use. Waist and hip circumference was performed to the nearest 0.1 cm using a nonstretchable standard tape measure attached to a spring balance exerting a force of 750 grams (Ohaus tape). The waist circumference was taken over the unclothed abdomen at the smallest diameter, between the costal margin and the iliac crest. The tape mea-

sure was kept horizontal and just tight enough to allow the little finger to be inserted between the tape and the subject's skin. The hip circumference was taken at the level of greater trochanters (usually the widest diameter around the buttocks).

The blood pressure was done using standard mercury sphygmomanometer, twice on the right arm and the exact values recorded to the nearest 2 mmHg. A third measurement would be performed if there was 10 mmHg or more difference between the first two readings, and the average of the two closest values were used for the analysis. The correlation coefficients between the two measurements of systolic and diastolic blood pressure were 0.94 and 0.89 respectively.

Blood samples were taken after 10-12 hours fasting and were processed within 4 hours and serum stored in -70° C for further analysis. The DNA material was extracted from the WBC and stored for further analysis. The laboratory performance included complete blood count, fasting blood sugar, serum lipids (cholesterol, triglycerides and HDL-cholesterol), serum creatinine and uric acid using Hitachi 717 and 917 automation systems. The coefficients of variation between run and within run were performed every day and were less than 5 per cent. The external quality control was performed every 4 week by joining the QAP (quality assurance program) from Roche Diagnostics. The coefficients of variation for fasting blood glucose, serum cholesterol, triglycerides and HDL-cholesterol were 1.36 per cent, 2.29 per cent, 3.09 per cent and 3.45 per cent respectively. Twelve lead electrocardiography were performed on those subjects aged more than 30 years old, using the HP- Playwriter XII with the autoanalyzer and was confirmed by one of the investigators.

The data was recorded twice in the Dbase Foxpro II by two separate research assistants. If there was any discrepancy between the two values, the data would be checked and corrected. A telephone call would be made directly to the subject to obtain any missing data from the questionnaires. The completeness of our data ranged from 99.0-99.9 per cent. The statistical analysis was performed by a biostatistician using the SPSS for Window. Chi-square and analysis of variance were applied where appropriate. Multivariate analysis of covariance was used to adjust the age, sex and body mass index difference which was found amongst impaired fasting glucose, diabetes mellitus and normal.

## RESULTS

A total of 3,615 subjects were studied, 1,250 (34.3%) were male and 2,365 (65.7%) were female. The mean age was  $30.0 \pm 5.6$  years (range 18-58). Most of the population had a family income of more than 10,000 baht/month and graduated from a university or higher. A history of smoking was more common in males. The prevalence of current frequent smoking and social smoking in males was 18.5 per cent and 13.1 per cent respectively while the prevalence in females was 1.6 per cent and 6.5 per cent. The mean body mass index was  $21.5 \pm 3.4$  kg/m<sup>2</sup> (range 13.7-45.1). The mean systolic and diastolic blood pressures were  $113.4 \pm 12.5$  (range 78-188) and  $75.6 \pm 9.4$  (range 35-139) mmHg respectively. The mean cholesterol was  $200.5 \pm 36.6$  mg/dL (range 98-377) while the means of triglycerides and HDL-cholesterol were  $91.5 \pm 60.0$  (range 21-817) and  $58.1 \pm 14.7$  mg/dL (range 13-127) respectively.

The prevalence of impaired fasting glucose (IFG) and diabetes mellitus (by blood sugar level

only) were 1.7 per cent and 0.8 per cent respectively. The prevalence of impaired fasting glucose and diabetes mellitus was more common in males (3.4% vs 0.8% and 2.2% vs 0.1% respectively). The prevalence increased with increasing age as shown in Table 1.

The prevalence of hypertension, diabetes mellitus, hyperlipidemia, obesity, physical inactivity and smoking (blood pressure or blood levels and history) were 7.4 per cent, 1.4 per cent, 21.1 per cent, 13.9 per cent, 76.3 per cent and 16.3 per cent respectively. The awareness of hypertension, diabetes mellitus and hyperlipidemia was 42.2 per cent, 78 per cent and 32.9 per cent respectively.

Table 2 showed that the unadjusted coronary risk factors including systolic blood pressure, diastolic blood pressure, pulse pressure, body mass index, waist and hip circumference, waist/hip ratio, serum cholesterol, triglycerides and LDL-cholesterol were higher in IFG than in normal and were higher in DM than in IFG. While the HDL-cholesterol level was lower in IFG than in normal. There

**Table 1. Age and sex stratified prevalence of impaired fasting glucose (IFG) and diabetes mellitus (DM).**

		Age group (years)				
		<30 (%)	30-39 (%)	40-49 (%)	50-59 (%)	Total (%)
Male	IFG	2.3	2.3	13.2	4.5	3.4
	DM	0.5	2.7	6.6	9.1	2.2
Female	IFG	0.6	0.9	3.7	-	0.8
	DM	0.1	0.1	-	-	0.1

**Table 2. Unadjusted means standard error of coronary risk factors in impaired fasting glucose, diabetes mellitus and normal population.**

	Normal (n = 3,486)		Impaired fasting glucose (n = 61)		Diabetes mellitus (n = 30)	
	mean	SE	mean	SE	mean	SE
Systolic blood pressure (mmHg)	113.1	0.21	122.6	2.04	128.7	2.69
Diastolic blood pressure (mmHg)	75.37	0.16	81.3	1.47	88.6	2.20
Pulse Pressure (mmHg)	37.8	0.14	41.1	1.19	40.2	1.48
Body mass index (kg/m <sup>2</sup> )	21.4	0.06	23.6	0.51	26.0	0.90
Waist circumference (cm)	69.1	0.17	78.4	1.51	86.8	2.52
Hip circumference (cm)	90.1	0.12	94.2	1.0	97.8	1.7
Waist / Hip ratio	0.77	0.001	0.83	0.009	0.88	0.02
Cholesterol (mg/dL)	200	0.61	220.5	5.29	222.8	9.99
Triglyceride (mg/dL)	89.3	0.95	156.2	15.64	208.3	22.45
HDL-cholesterol (mg/dL)	58.4	0.25	50.5	1.66	46.4	2.71
LDL-cholesterol (mg/dL)	123.7	0.55	138.8	4.56	134.8	8.22

SE = standard error

Remark: Normal - fasting plasma glucose < 110 mg/dL, Impaired Fasting Glucose - fasting plasma glucose 111-125 mg/dL, Diabetes Mellitus - fasting plasma glucose > 125 mg/dL.

were statistical differences between normal, impaired fasting glucose and diabetes mellitus in all parameters, except there was no significant difference between impaired fasting glucose and diabetes mellitus in pulse pressure, serum cholesterol, LDL-cholesterol and HDL-cholesterol.

Age and sex adjusted values of the risk factors were shown in Table 3. The age, sex and body mass index adjusted values of these risk factors were shown in Table 4. Coronary risk factors even after age, sex and body mass index adjusted were also higher in impaired fasting glucose than in normal.

## DISCUSSIONS

The purpose of the diagnostic criteria that are based on blood glucose concentration is to iden-

tify individuals who have no symptoms of diabetes but who have hyperglycemia(27-32) and are, therefore, at increased risk of subsequent complications and mortality. Their study showed that those people who had impaired fasting glucose had a higher level of risk factors, i.e. systolic blood pressure, diastolic blood pressure, pulse pressure, body mass index, abdominal obesity and serum lipids level although they did not have any symptoms. If the purpose of screening for abnormal glucose disorder is to identify the maximum number of people at risk of cardiovascular disease, and there is an effective treatment to prevent cardiovascular disease events or death when glucose is slightly raised, then it would seem that the fasting ADA criteria of impaired fasting glucose are superior to the fasting ADA criteria for diabetes mellitus.

**Table 3. Adjusted means and standard error by sex and age of coronary risk factors in normal, impaired fasting glucose and diabetes mellitus using analysis of covariance.**

	Normal (n = 3,486)		Impaired fasting glucose (n = 61)		Diabetes mellitus (n = 30)	
	mean	SE	mean	SE	mean	SE
Systolic blood pressure (mmHg)	114.5	0.2	117.5	1.6	124.6	3.4
Diastolic blood pressure(mmHg)	76.4	0.2	78.2	1.2	87.0	2.6
Pulse Pressure (mmHg)	38.1	0.1	39.4	1.2	37.6	2.5
Body mass index (kg/m <sup>2</sup> )	21.8	0.06	22.7	0.4	22.2	0.9
Waist circumference (cm)	71.2	0.1	74.2	1.1	73.9	2.3
Hip circumference (cm)	90.9	0.1	92.6	0.9	91.8	2.0
Waist / Hip ratio	0.78	0.001	0.80	0.007	0.80	0.016
Cholesterol (mg/dL)	201.2	0.6	208.4	4.9	193.99	10.8
Triglyceride (mg/dL)	95.2	1.0	130.5	7.6	133.9	16.6
HDL-cholesterol (mg/dL)	56.7	0.2	52.7	1.9	55.8	4.2
LDL-cholesterol (mg/dL)	125.3	0.6	129.6	4.4	111.4	9.7

SE = standard error

Remark: Normal - fasting plasma glucose < 110 mg/dl, Impaired Fasting Glucose - fasting plasma glucose 111-125 mg/dl, Diabetes Mellitus - fasting plasma glucose > 125 mg/dl.

**Table 4. Adjusted means and standard error by sex, age and body mass index of coronary risk factors in normal, impaired fasting glucose and diabetes mellitus.**

	Normal (n = 3,486)		Impaired fasting glucose (n = 61)		Diabetes mellitus (n = 30)	
	mean	SE	mean	SE	mean	SE
Systolic blood pressure (mmHg)	114.3	0.2	116.5	1.5	124.0	3.3
Diastolic blood pressure(mmHg)	76.3	0.2	77.5	1.2	86.6	2.5
Pulse Pressure (mmHg)	38.0	0.2	39.0	1.2	37.5	2.5
Waist circumference (cm)	70.6	0.08	71.8	0.6	72.6	1.4
Hip circumference (cm)	90.4	0.07	90.6	0.6	90.8	1.2
Waist / Hip ratio	0.78	0.001	0.79	0.006	0.79	0.01
Cholesterol (mg/dL)	200.7	0.6	206.3	4.9	192.9	10.7
Triglyceride (mg/dL)	93.8	0.9	124.6	7.3	130.6	15.9
HDL-cholesterol (mg/dL)	57.1	0.2	54.1	1.8	56.6	4.0
LDL-cholesterol (mg/dL)	124.8	0.6	127.3	4.4	110.2	9.5

SE = standard error

Remark: Normal - fasting plasma glucose < 110 mg/dl, Impaired Fasting Glucose - fasting plasma glucose 111-125 mg/dl, Diabetes Mellitus - fasting plasma glucose > 125 mg/dl.

The UK Prospective Diabetes Study of newly diagnosed individuals with diabetes<sup>(33)</sup> suggests that keeping glucose concentrations low at an early stage of hyperglycaemia is important in the prevention of cardiovascular disease. Likewise, early recognition of abnormal glucose concentrations may alert the physician to more aggressively managed comorbidities such as hyperlipidemia, obesity, and hypertension. Treatment strategies for these disorders vary and depend on whether glucose abnormalities are present<sup>(34)</sup>. From our study, we cannot conclude impaired fasting glucose will progress to diabetes mellitus or lead to cardiovascular disease, although the Decode study and the study by Barzilay JI *et al* showed there was an increase in cardiovascular disease and death in impaired fasting glucose<sup>(35,36)</sup>. From these two studies, the impaired fasting glucose (IFG) by the new ADA criteria is less sensitive than the impaired glucose tolerance (IGT). At baseline, cardiovascular events attributable to abnormal glucose states were clearly fewer with the ADA criteria than with the WHO criteria (53 vs 159 cases per 10,000), largely because many more people were classified as having an abnormal glucose state by the WHO criteria (46.8%) than by the ADA criteria (22.3%).

#### Limitations

Several potential limitations of this study should be noted. First, our population was a selected

group which represented a relatively young age, high socioeconomic status, high education and had a high level of stress. Second, we used only one baseline glucose measurement to classify individuals into glucose categories. Over time, many individuals who were normal, developed abnormal glucose concentration and vice versa.

#### SUMMARY

This study showed that the prevalence of impaired fasting glucose and diabetes mellitus in our population were 1.7 per cent and 0.8 per cent respectively. The prevalences were more common in males and with increasing age. Coronary risk factors such as high blood pressure, excessive weight and obesity, and abnormal lipid levels were more common in impaired fasting glucose even adjusted for age and sex. Progression to diabetes mellitus and whether or not impaired fasting glucose was associated with cardiovascular death need further study in a prospective fashion. Lifestyle modification in those with impaired fasting glucose may be needed for the prevention of coronary disease.

#### ACKNOWLEDGEMENT

The authors would like to thank the Shina-watra Corporation for support of the study. We would also like to thank all the staff of Her Majesty Cardiac Center for their help throughout the study.

(Received for publication on October 2, 2000)

#### REFERENCES

1. American Diabetes Association. Report of the expert committees on the diagnosis and classification of diabetes mellitus. *Diabetes Care* 1997;20: 1183-97.
2. WHO. WHO expert committee on diabetes mellitus: second report-WHO Technical Report Series 646. Geneva: WHO, 1980.
3. Fuller JH, Shipley MJ, Rose G, *et al*. Mortality from coronary heart disease and stroke in relation to degree of glycaemia: the Whitehall Study. *BMJ* 1983;287:867-70.
4. Jarrett RJ, Keen H, McCartney M. Worsening to diabetes in men with impaired glucose tolerance. *Diabetologia* 1979;16:25-30.
5. Bhuripanyo K, Bhuripanyo P, Muktaphan B, Kusalertjariya S, Khumsuk K, Saensongsri V. Cardiovascular risk factors in impaired glucose tolerance (IGT) compared with normal and diabetic patients. *Thai Heart J* 1990;3:83-9.
6. Welborn TA, Wearne K. Coronary heart disease incidence and cardiovascular mortality in Buselton with reference to glucose and insulin concentration. *Diabetes Care* 1979;2:154-60.
7. Stamler R, Stamler J, Lindberg HA, *et al*. Asymptomatic hyperglycaemia and coronary heart disease in middle aged men in two employed populations in Chicago. *J Chronic Dis* 1979;32: 805-15.
8. Pyorala K, Savolainen E, Lehtovirta E, Punar S.

Siltanen P. Glucose tolerance and coronary heart disease: Helsinki Policemen Study. *J Chronic Dis* 1979;32:729-45.

9. Schroll M, Hagerup L. Relationship of fasting blood glucose to prevalence of ECG abnormalities and 10 year risk of mortality from cardiovascular diseases in men born in 1914: from the Glostrup Population Studies. *J Chronic Dis* 1979; 32:699-707.

10. Reunanan A, Pyorala K, Aromaa A, Maatela J, Knek P. Glucose tolerance and coronary heart disease in middle aged Finnish men : Social Insurance Institution's Coronary Hearat Disease Study. *J Chronic Dis* 1979;32:747-58.

11. Fuller JH, Shipley MJ, Rose G, Jarrett RJ, Keen H. Mortality from coronary heart disease and stroke in relation to degree of glycaemia: the Whitehall study. *BMJ* 1983;287:867-70.

12. Barrett-Connor E, Wingard DL. Sex differential in ischemic heart disease mortality in diabetics: a prospective population based study. *Am J Epidemiol* 1983;118:489-96.

13. Cruz-Vidal M, Garcia-Palmieri MR, Costas R Jr, Sorlie PD, Havlik RJ. Abnormal blood glucose and coronary heart disease: the Puerto Rico Heart Health Program. *Diabetes Care* 1983;6:556-61.

14. Barrett-Connor E, Wingard DL, Criqui MH, Suarex L. Is borderline hyperglycaemia a risk factor for cardiovascular death? *J Chronic Dis* 1984;37: 773-9.

15. Eschwege E, Richard JL, Thibault N, et al. Coronary heart disease mortality in relation with diabetes, blood glucose and plasma insulin levels: the Paris Prospective Study, ten years later. *Horm Metab Res* 1985;15(suppl):41-6.

16. Butler MU, jOstrander LD Jr, Carman WJ, Lamphier DE. Mortality from coronary heart disease in the Tecumseh Study: long term effect of diabetes mellitus, glucose tolerance and other risk factors. *Am J Epidemiol* 1985;121:541-7.

17. Pan WH, Cedres LB, Liu K, et al. Relationship of clinical diabetes and asymptomatic hyperglycaemia to risk fo coronary heart disease mortality in men and women. *Am J Epidemiol* 1986;123: 504-16.

18. Scheidt-Nave C, Barrett-Connor E, Wingard DL, Cohn BA, Edelstein SL. Sex differences in fasting glycemia as a risk factor for ischemic heart disease death. *Am J Epidemiol* 1991;133:565-76.

19. Wilson PWF, Cupples LA, Kannel WB. Is hyperglycaemia associated with cardiovascular disease? The Framingham Study. *Am Heart J* 1991;121: 586-90.

20. Vaccaro O, Ruth KJ, Stamler J. Relationship of postload plasma glucose to mortality with 19 year follow-up: comparison of one versus two plasma glucose measurements in the Chicago Peoples Gas Company Study. *Diabetes Care* 1992;15:1328-34.

21. Feskens EJM, Kromhout D. Glucose tolerance and the risk of cardiovascular diseases: the Zutphen Study. *J Clin Epidemiol* 1992;45:1327-34.

22. Yarnell JWG, Pickering JE, Elwood PC, et al. Does nondiabetic hyperglycaemia predict future IHD? Evidence from the Caerphilly and Speedwell studies. *J Clin Epidemiol* 1994;47:383-8.

23. Yano K, Kagan A, McGee D, Rhoads G. Glucose intolerance and nine year mortality in Japanese men in Hawaii. *Am J Med* 1982;72:71-80.

24. Ohlson LO, Svardsudd K, Welin L, et al. Fasting blood glucose and risk of coronary heart disease, stroke and all cause mortality: a 17 year follow-up study of men born in 1913. *Diabet Med* 1986;3: 33-7.

25. Wei M, Gibbons L, Mitchell T, Kampert J, Blair S. Undiagnosed diabetes and impaired fasting glucose as predictors of cardiovascular disease and all cause mortality. *Cardiovasc Dis Prev* 1998;1: 123-8.

26. Balkau B, Shipley M, Jarrett RJ, et al. High blood glucose concentration is a risk factor for mortality in middle-aged non-diabetic men. *Diabetes Care* 1998;21:360-74.

27. The expert committee on the diagnosis and classification of diabetes mellitus. Report of the expert committee on the diagnosis and classification of diabetes mellitus. *Diabetes Care* 1997;20:1183-97.

28. Alberti KGMM, Zimmet PZ, for the WHO consultation. Definition, diagnosis and classification of diabetes mellitus and its complications. Part 1: dianosis and classification of diagnosis and classification of diabetes mellitus. Provisional report of a WHO Consultaiton. *Diabet Med* 1998;15: 539-53.

29. DECODE Study Group. Will new diagnostic criteria for diabetes mellitus change phenotype of patients with diabetics? Reanalysis of European epidemiological data. *BMJ* 1998 ;317:371-5.

30. WHO Study Group. Diabetes mellitus. WHO Technical Report Series 727. Geneva: WHO, 1985.

31. Harris MI. Undiagnosed NIDDM: clinical and public health issues. *Diabetes Care* 1993;16:642-52.

32. Valle T, Tuomilehto J, Eriksson J. Epidemiology of NIDDM in Europids. In : Alberti KGMM, Zimmet P, DeFronzo RA, Keen H, eds. International textbook of diabetes mellitus, 2<sup>nd</sup> edn. London: Wiley and Sons, 1997:125-42.

33. UK Prospective Diabetes Study Group. Intensive blood-glucose control with sulphonylureas or insulin compared with conventional treatment and risk of complications in patients with type 2 diabetes (UKPDS 33). *Lancet* 1998;352:837-53.

34. Haffner SM, Lento S, Ronnemaa T, Pyorala K,

Laakso M. Mortality from coronary heart disease in subjects with type 2 diabetes and in nondiabetic subjects with and without prior myocardial infarction. *N Engl J Med* 1998;339:229-34.

35. The DECODE Study Group on behalf of the European Diabetes Epidemiology Group. Glucose tolerance and mortality: comparison of WHO and American Diabetes Association diagnostic criteria. *Lancet* 1999;354:617-21.

Barzilay JI, Spiekerman CF, Wahl PW, et al. Cardiovascular disease in older adults with glucose disorders: comparison of American Diabetes Association criteria for diabetes mellitus with WHO criteria. *Lancet* 1999;354:622-25.

## การศึกษาระดับของปัจจัยเสี่ยงของโรคหลอดเลือดหัวใจใน Impaired fasting glucose (IFG) และโรคเบาหวาน

เกียรติชัย ภูริปัญญา, พ.บ.\*, องค์การ เรืองรัตนอัมพร, พ.บ.\*; นิธิ มหานนท์, พ.บ.\*; วัฒนา เลี้ยววัฒนา, พ.บ.\*\*, ชานุ ศรีรัตน์สถาวร, พ.บ.\*; ชุณหกेषม ใจดินัยวัตรกุล, พ.บ.\*; จากรุรรณ คงคงเกตุ, วท.ม. (ชีวสัตติ)\*; ศุภชัย ไชยธีระพันธ์, พ.บ.\*

คณบัญชีได้ทำการสำรวจความซุกของ impaired fasting glucose และโรคเบาหวานในกลุ่มพนักงานบริษัท ขินวัติจำนวน 3615 คน พนักงานซุกของ impaired fasting glucose และเบาหวานเท่ากับ 1.7% และ 0.8% ความซุกของ impaired fasting glucose และเบาหวานพบมากกว่าในเพศชาย และความซุกสูงขึ้นตามอายุที่เพิ่มขึ้น การตรวจระดับของความดันโลหิต systolic, diastolic, pulse pressure, ดัชนี body mass index, รอบเอว, อัตราส่วนรอบเอว/รอบสะโพก และระดับไขมัน cholesterol, triglyceride, และ LDL-cholesterol พนักงาน impaired fasting glucose และเบาหวานสูงขึ้นกว่าในประชากรที่มีระดับน้ำตาลอ้อยในเกณฑ์ปกติ (ยกเว้น HDL-cholesterol) และระดับของปัจจัยเสี่ยงเหล่านี้ในคนที่เป็นเบาหวานสูงกว่าคนที่เป็น impaired fasting glucose (ยกเว้น pulse pressure ระดับ cholesterol, HDL-cholesterol และ LDL-cholesterol)

**คำสำคัญ** : Impaired Fasting Sugar, เบาหวาน, ปัจจัยเสี่ยงโรคหลอดเลือดหัวใจ

เกียรติชัย ภูริปัญญา, องค์การ เรืองรัตนอัมพร, นิธิ มหานนท์, และคณะ  
จตุภาคัยเทฤทธิวงศ์ฯ 2543; 83 (ฉบับพิเศษ 2): S146-S152

\* สำนักงานศูนย์โรคหัวใจสมเด็จพระบรมราชินีนาถ,

\*\* ภาควิชาพยาธิวิทยาคลินิก, คณะแพทยศาสตร์ศิริราชพยาบาล, มหาวิทยาลัยมหิดล, กรุงเทพฯ 10700