

The State of Lipid Control in Patients with Diabetes in a Public Health Care Centre

JS Wong, F. Tan and PY Lee
Asia Pac J Public Health 2007 19: 16
DOI: 10.1177/101053950701900304

The online version of this article can be found at:
<http://aph.sagepub.com/content/19/3/16>

Published by:



<http://www.sagepublications.com>

On behalf of:



[Asia-Pacific Academic Consortium for Public Health](http://www.sagepublications.com)

Additional services and information for *Asia-Pacific Journal of Public Health* can be found at:

Email Alerts: <http://aph.sagepub.com/cgi/alerts>

Subscriptions: <http://aph.sagepub.com/subscriptions>

Reprints: <http://www.sagepub.com/journalsReprints.nav>

Permissions: <http://www.sagepub.com/journalsPermissions.nav>

Citations: <http://aph.sagepub.com/content/19/3/16.refs.html>

>> [Version of Record](#) - Jul 1, 2007

[What is This?](#)

Abstract

Achieving treatment targets has been difficult in treating diabetic patients. This cross-sectional study describes the lipid profiles of patients with diabetes mellitus at a public primary health care centre in Sarawak, Malaysia. The targets for lipid control were based on the International Diabetes Federation recommendation (2002). 1031 patients (98% Type 2 Diabetes) were studied. Fasting lipid profiles were available in 990 (96%) patients. The mean total cholesterol was 5.3 ± 1.0 mmol/L, Triglycerides 1.90 ± 1.26 mmol/L, HDL-C 1.28 ± 0.33 mmol/L and LDL-C 3.2 ± 0.9 mmol/L. Overall, 22% of patients achieved the treatment target for LDL-C level <2.6 mmol/L. 67% of patients had HDL-C >1.1 mmol/L and 42% of patients had a target TG level below 1.5 mmol/L. Of the 40% of patients who received lipid-lowering drug, 17% achieved LDL-C target, 50% had LDL-C 2.6-4.4 mmol/L and 33% have LDL-C >4.0 mmol/L. For the remaining 60% not receiving any lipid lowering therapy, 68% had LDL-C between 2.6-4.0 mmol/L and 7% had LDL-C level >4 mmol/L. Dyslipidemia is still under-treated despite the availability of effective pharmacological agents and the greatly increased risk of cardiovascular diseases in diabetic patients. *Asia Pac J Public Health* 2007; 19(3): 16–21.

Keywords: Cholesterol, dyslipidemia, diabetes mellitus, primary health care.

Address for correspondence:

Dr Wong Jin Shyan
Department of Medicine,
Sarawak General Hospital,
Jalan Tun Ahmad Zaidi Aduce,
93586 Kuching,
Sarawak, Malaysia.
E-mail:
wongjinshyan@health.gov.my
Mobile: +6016-5208295
Fax: +6082-240767

The State of Lipid Control in Patients with Diabetes in a Public Health Care Centre

JS Wong¹, MD
F Tan¹, MRCP
PY Lee², MMed

¹ Department of Medicine, Sarawak General Hospital, Kuching, Sarawak, Malaysia

² Department of Medicine, Faculty of Medicine and Health Sciences, University Malaysia Sarawak, Sarawak, Malaysia

Introduction

Diabetes mellitus (DM) is one of the most common disorders seen in primary health care clinic. The prevalence of DM in Malaysian adults was estimated to be 8.2% in 1996¹ but the number is escalating rapidly. People with diabetes have increased morbidity and mortality particularly from cardiovascular diseases. In fact, the absolute risk of cardiovascular death was three times higher for diabetic than non-diabetic men² and people with diabetes have a similar risk of cardiovascular events as non-diabetic patients with established cardiovascular diseases³. DM is thus considered to be coronary heart disease risk equivalents and diabetic patients should have their cardiovascular risk factors treated as those with established coronary heart disease⁴.

Numerous clinical studies have shown that reducing lipid level effectively decreases the risk of major cardiovascular events⁵⁻⁷. The low-density lipoprotein cholesterol (LDL-C) was targeted as the primary goal in cholesterol reduction⁸ and current guidelines recommend an LDL-C level of less than 2.6 mmol/L for people with diabetes. However, few studies have explored to what extent

the guideline is being followed and the treatment goal is being achieved in diabetic patients, especially in a community care setting where the majority of diabetic patients are being managed. We therefore examine the rate of testing, treatment and goal attainment for lipid control in people with diabetes managed in a busy public health care centre.

Methods

This is a cross-sectional study conducted as an audit of a diabetes clinic in a public primary health care clinic - Tanah Puteh Health Centre (KKTP) in Sarawak, East Malaysia. Sarawak is one of the thirteen states in Malaysia with a population of 2.1 million where 460,000 live in Kuching, the state capital⁹. There are three public primary health care clinics in Kuching and KKTP is one of them, registering 77,301 consultation visits in 2002 with 54,429 consultations by adult patients. Twelve percent (6,460) of the consultations were held at the diabetes clinic¹⁰. The diabetes clinic in KKTP was established in 2002 and managed by three doctors, three medical assistants and one trained diabetic nurse educator. The clinic registered 1,337 patients by 31st of December

2002. A cross-sectional study was conducted on these patients from 1 January 2003 to 31 March 2003 to assess the effectiveness of the diabetes clinic. This report describes the state of lipid control in the clinic with regards to the rate of testing, treatment and goal attainment.

All data were captured from the clinic based diabetes cards and entered manually into an SPSS version 10 (SPSS Inc., Chicago, Illinois, USA). Information collected included patient demography, type of diabetes, cardiovascular risk factors (blood pressure, lipids, BMI and smoking history), glycemic control (HbA_{1c} and FBS), renal function (serum creatinine, micro-albuminuria and proteinuria) and diabetes treatment (pharmacological and non-pharmacological). The patients fasted overnight for 8 to 10 hours for the purpose of testing for fasting lipids and blood sugar profile. Laboratory data represent the most recent results available within the preceding twelve months. Data fields were left blank if no data were available. All data were tabulated and descriptive statistical analysis performed.

The diagnosis of diabetes mellitus was made according to the WHO criteria¹¹. The target for control of diabetes mellitus was based on the recommendation by the IDF¹² (Table 1). The target of cholesterol treatment was Total cholesterol <4.5 mmol/L, Triglycerides <1.5 mmol/L, HDL-C >1.1 mmol/L, and LDL-C <2.6 mmol/L. Patients were managed according to the national guidelines and consensus for diabetes, hypertension and hyperlipidemia¹³⁻¹⁵.

Lipid profiles were measured after an overnight fast (clinical chemistry workhorse system Roche/Hitachi 912®, Roche Diagnostics). HbA_{1c} was measured by a high performance liquid chromatography method (the Bio-Rad Variant® Hemoglobin testing system).

Results

The diabetes clinic registry recorded

a total of 1,337 patients by 31 December 2002. The study was conducted over a period of three months. The final analysis had excluded 23% of the patients as shown in Figure 1. The 1,031 remaining patients on active follow-up constituted the final study population.

Patient demography

The majority of the patients were Chinese (69.0%). Other ethnic groups were Malays (21.2%), Sarawak Natives (9.1%), Indians (0.5%), and others (0.2%). Women (63%) outnumbered men. Majority (98%) of the patients had type 2 diabetes.

The mean (\pm SD) duration of diabetes was 6.4 \pm 6 years. The mean age was 59 \pm 12 years. The mean BMI was 26.8 \pm 4.5 kg/m² with 42% of patients overweight (BMI 25-30 kg/m²) and

21% were obese (BMI > 30 kg/m²). 18% of men and 22% of women were obese. The mean waist circumference (WC) was 90 \pm 10 cm. More women (54%) had central obesity (WC >88 cm) when compared to men (WC > 102 cm) (21%)⁸.

For glycemic control, the mean HbA_{1c} was 7.4 \pm 1.6%. The optimal target of HbA_{1c} <6.5% was achieved by 28% of the patients. Pharmacological treatment for hyperglycaemia was prescribed to 96% of the patients while the remaining 4% were on lifestyle modification alone.

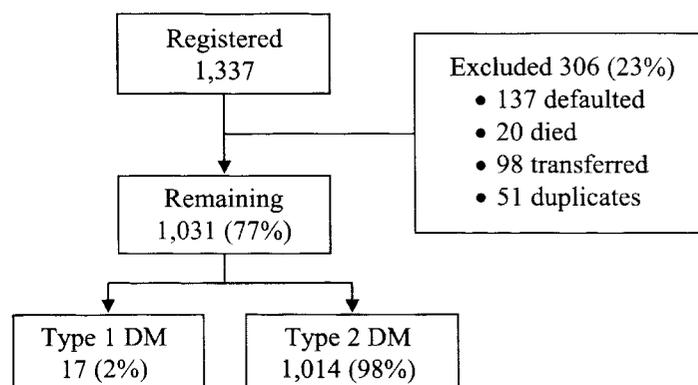
Some 75% of patients had a diagnosis of hypertension and were on treatment. The mean systolic and diastolic blood pressures were 137 \pm 17 mmHg and 85 \pm 10 mmHg respectively. However, only 6% of the patients had an optimal blood pressure

Table 1. Targets for glycemic, blood pressure and lipid control for diabetic patients

	optimal	Fair	poor
Fasting blood glucose (mmol/l)	4.4–6.0	6.1–7.0	>7.0
HbA _{1c} (%)	<6.5	6.5–7.5	>7.5
Blood pressure (mmHg)	<130/80	130/80–140/90	>140/90
Total cholesterol (mmol/l)	<4.5	4.5–6.0	>6.0
Triglycerides (mmol/l)	<1.5	1.5–2.2	>2.2
HDL-cholesterol (mmol/l)	>1.1	0.9–1.1	<0.9
LDL-cholesterol (mmol/l)	<2.6	2.6–4.0	>4.0

Modified from The Asian-Pacific Type 2 Diabetes Policy Group. Type 2 Diabetes, Practical targets and treatments, 3rd Ed. 2002: WHO. IDF.

Figure 1. Patients constituting the final analyzed population and reason for exclusion



<130/80 mmHg.

Among the 970 patients tested for proteinuria, 48% had evidence of diabetic nephropathy (with 67% overt proteinuria and 33% micro-albuminuria). 78% of them received treatment with either an Angiotensin Converting Enzyme Inhibitors or an Angiotensin Receptor Blockers. Table 2 shows the various cardiovascular risk factors of the study population. Detailed results of these clinical parameters were described elsewhere^{16,17}.

Lipid control

Figure 2 shows the fasting lipid profile of the patients. 990 patients (96%) had a fasting lipid profile tested. The mean total cholesterol was 5.3 ± 1.0 mmol/L. There were 3% (31 patients) with the serum triglycerides exceeding 4.5 mmol/L. The LDL-C results were therefore not valid. The LDL-C was available in 959 patients. The mean LDL-C was 3.2 ± 0.9 mmol/L and the median was also 3.2 mmol/L. The majority of patients (78%) did not achieve the LDL-C treatment goal. Only 22% (207 patients) achieved the optimal LDL-C level of <2.6 mmol/L. Another 61% (589 patients) had an LDL-C between 2.6-4.0 mmol/L and 17% (163 patients) had LDL-C >4.0 mmol/L.

The mean HDL-C level was 1.28 ± 0.33 mmol/L. The HDL-C fared much better with 67% of the patients achieving the desired target of > 1.1 mmol/L. Another 24% had a HDL-C level between 0.9-1.1 mmol/L and 8% had HDL-C <0.9 mmol/L. The mean HDL-C was higher in women (1.34 ± 0.34 mmol/L) than in men (1.16 ± 0.29 mmol/L). 76% of the women and 54% of the men had a HDL-C of >1.1 mmol/L. The mean triglyceride level was 1.90 ± 1.26 mmol/L. 42% of the patients had an optimal level of serum triglycerides (<1.5 mmol/L). 32% and 24% of the patients had a serum triglyceride level of 1.5-2.2 mmol/L and >2.2 mmol/L respectively.

Forty percent (392 patients) of the 990 patients received cholesterol-lowering drugs. 85% (333 patients)

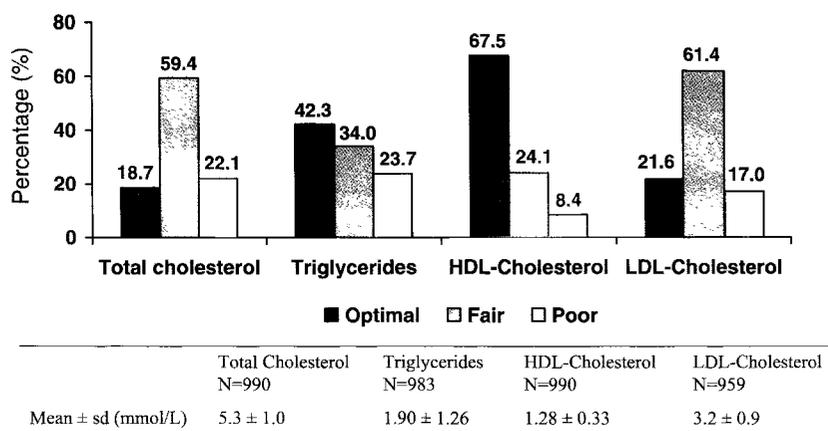
Table 2. Cardiovascular risk profiles

	Type 1 (n=17)	Type 2 (n=1014)
Age		
Male >45	78% (7)	95% (353)
Female >55	62% (5)	88% (565)
On treatment for hypertension	59% (10)	75% (762)
SBP/DBP >130/80	94% (16)	94% (951)
SBP>130	65% (11)	77% (775)
DBP>80	71% (12)	88% (893)
Obesity ^a		
BMI 25-30 (kg/m ²)	53% (9)	42% (422)
BMI >30 (kg/m ²)	18% (3)	21% (204)
Smoking	24% (4)	6% (66)
Proteinuria ^b	65% (11)	48% (458)
LDL-cholesterol ^c >2.6 (mmol/L)	82% (14)	78% (739)
Glycaemia control		
FBS>8 (mmol/L)	53% (9)	37% (375)
HbA _{1c} ^d >6.5 (%)	76% (13)	72% (687)

Some patients have missing data and are not included in the calculation of percentages.

Total number of patients with type 2 DM included in analysis -- a: 999 subjects, b: 954 subjects, c: 943 subjects, d: 959 subjects.

Figure 2. Fasting lipid profile and level of lipid control according to IDF¹²

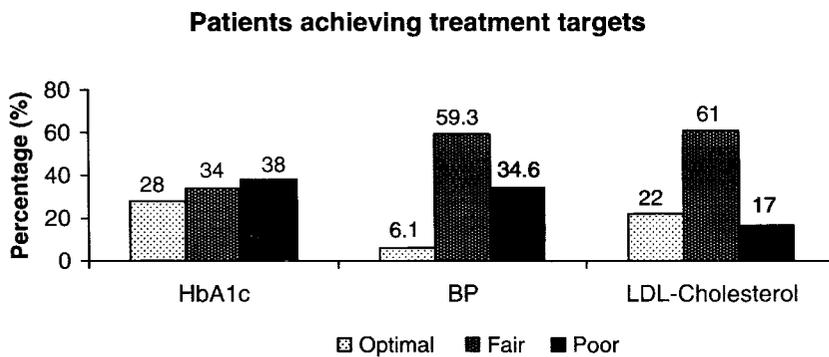


were treated with statins where lovastatin was prescribed in 99%. Another 14% (56 patients) were prescribed gemfibrozil alone. Combination therapy of gemfibrozil and lovastatin was given in 1% (three patients). For those who received lipid-lowering therapy, 17% (65/392) reached the LDL-C treatment goal. For the 60% (598 of 990) who did

not receive the lipid lowering drug, 31 patients did not have LDL-C reading, 24% did not need drugs as the LDL-C were <2.6 mmol/L, 68% and 7% had LDL-C level between 2.6-4.0 mmol/L and >4mmol/L respectively.

There were more patients achieving the target of optimal glycaemic control than those achieving the target for

Figure 3. Percentage of patients achieving treatment targets in terms of HbA_{1c}, overall blood pressure and LDL-C level



Optimal: HbA_{1c} <6.5%; BP <130/80mmHg; LDL-Cholesterol <2.6mmol/L

Fair: HbA_{1c} 6.5-7.5%; BP 130/80-140/90 mmHg; LDL-Cholesterol 2.6-4.0mmol/L

Poor: HbA_{1c} >7.5%; BP >140/90mmHg; LDL-Cholesterol >4.0mmol/L

LDL-C or blood pressure. The HbA_{1c} treatment target of <6.5% was achieved in 28% of patients while 22% achieved target LDL-C <2.6 mmol/L. Only 6% achieved the target blood pressure <130/80 mmHg (Figure 3).

Discussion

Diabetes mellitus independently predicts mortality as well as cardiovascular complications. The successful treatment of DM requires more than the control of hyperglycemia. A multifaceted approach paying particular attention to the multiple cardiovascular risk factors is crucial in reducing excess morbidity and mortality associated with diabetes¹⁸⁻²⁰. Dyslipidemia outweighs other risk factors as a risk factor for coronary artery disease²¹. Large clinical trials have established the effectiveness and benefits of lipid lowering drugs, especially statin, in reducing cardiovascular events in diabetic patients. The importance is further highlighted in two recent large clinical trials which demonstrate similar beneficial effect in diabetic patients even in the absence of cardiovascular disease or elevated LDL-C^{22,23}. This has led to increasingly stringent lipid treatment goals being put forward in recent guidelines^{24,25}.

This study is an audit of the healthcare centre showing that the management of hyperlipidemia in diabetic patients is suboptimal and illustrating that doctors still target glucose alone in the management of diabetic patients. Although testing for dyslipidemia was performed in 96% of the diabetic patients, only 40% received a lipid lowering drugs and even less patients, i.e 22% achieved the recommended treatment target with an LDL-C >2.6 mmol/L. Even for patients receiving the lipid-lowering drugs, the LDL-C goal attainment rate was only 17%. This treatment gap between recommended standard of care and what the patients actually received had been reported in other studies²⁶⁻²⁸. Our results are better than the findings reported in a large managed care organisation where only 54% of their diabetic patients were tested for an LDL-C and only 28% received lipid lowering drugs²⁹. Similar to our study, they also reported a low goal attainment rate of 23% for LDL-C for those tested. The AusDiab study, a large population based survey conducted in Australia, showed a similar disappointing result with 80% of their diabetic patients not achieving the target LDL-C of < 2.6 mmol/L and only 36% of the patients receiving the lipid lowering treatment³⁰. Comparatively better results with 47%

goal attainment rate for LDL-C < 2.6 mmol/L had been reported for diabetic patients managed in a University primary care clinic³¹.

Several factors may explain the under-treatment of dyslipidemia observed in our study. The high testing rate for lipid profile in our centre contrasted with the low rate of drugs prescription for dyslipidemia. In our centre, the nurses or medical assistants had been instructed to test for fasting lipid profile in diabetic patients annually. This may account for the high lipid-testing rate. However, initiation and titration of lipid-lowering drugs was dependant entirely on the practising doctors. The under treatment reported in the study may be due to lack of awareness for both patients and physicians on the importance of treating dyslipidemia in diabetic patients. Only 40% of patients received lipid-lowering drugs whereas 96% received the pharmacological treatment for hyperglycaemia, 78% for proteinuria and 75% for hypertension. Even in those treated, only 17% achieved the treatment goal. An alarming 50% and 33% of those treated had LDL-C of 2.6-4.0 mmol/L and > 4.0 mmol/L respectively (Table 1). If the NCEP ATP III treatment goal was applied, more patients would need pharmacological agents and intensification of therapy. The healthcare providers need to lower the threshold for initiating and titrating pharmacological agents in order to maximise the potential benefits of treatment.

Increasing the awareness of the healthcare providers may increase the number of patients receiving the lipid-lowering therapy. However, this may not necessarily translate to better control. A study in Europe showed that although the proportion of patients treated with statin increased from 21% to 49%, 83% of patients on lipid lowering drugs still did not achieve the goal targeted for LDL-C³². Poor adherence to prescribed medication by patients, lack of follow up, inappropriate starting doses or failure to titrate lipid lowering therapy are all

potential barriers to goal achievement. A study in a managed care diabetes programme demonstrated that adherence to statin therapy by patients is an important determinant of the LDL-C goal attainment with a marked rise in the probability of successful treatments when the medication possession rate is above 80%³³. As medical therapy for patients with diabetes becomes more complex, reinforcement of practice guidelines should go hand in hand with education and empowerment of patients in order to improve compliance and achieve mutual goals.

Another potential contributing factor to the under-treatment of dyslipidemia in our study may be due to the limited access to lipid lowering drugs, despite the availability of generic preparations. In the Malaysian government health service formulary, the use of these drugs is a privilege restricted to family medicine specialist and physicians. This can be a hindrance to treatment of dyslipidemia in diabetes. A change of policy is very much desired. This may encourage the use of effective and proven therapy like statin in the treatment of dyslipidemia in high-risk population like people with diabetes.

The strength of our study is that this large study population consists of unselected people with diabetes receiving primary care in the community, and is thus likely to be representative of the status of care in similar setting in our country where the majority of diabetic patients are being managed. There are limitations to our study. As the data were collected from the clinic-based card, we did not assess patient's awareness of the importance of dyslipidemia, their compliance to treatment or any potential adverse effect with lipid lowering drugs. We also did not have the data on doctor's knowledge on guidelines and attitude towards management of dyslipidemia in diabetics. It would also be interesting to examine demographic data of goal achievers against the non-achievers to determine the reason for poor goal attainment. Similarly, although 99% of treated patients

received lovastatin, we did not examine the dosage used. However, given the low treatment rate of 40% in the study, failures to initiate and to titrate the dose were both likely to be important contributing factors towards poor goal attainment.

Conclusion

This audit study shows that hyperlipidemia in diabetic patients is still untreated or under-treated despite their greatly increase risk of cardiovascular disease. This may be due to several contributory factors as highlighted. Given the high lipid-testing rate in our clinic, more efforts should now focus on the implementation of clinical practice guidelines in terms of initiation, monitoring and titration of lipid lowering therapy, as well as on creating greater awareness among patients and health care providers on the importance of controlling dyslipidemia in people with diabetes.

Acknowledgements

The authors would like to thank the diabetes educator and staff of KKTP for their assistance and support in conducting the study. We would also like to acknowledge the Central Medical Laboratory, Sarawak General Hospital for performing the biochemical tests and the Department of Health, Ministry of Health, Sarawak, Malaysia.

References

1. Rugayah B. Diabetes Mellitus among Adults Aged 30 years and Above. National Health Morbidity Survey-2. Public Health Institute, Ministry of Health, Malaysia.
2. Stamler J, Vaccaro O, Neaton JD, Wenworth D. Diabetes, other risk factors, and 12-yr cardiovascular mortality for men screened in the Multiple Risk Factor Intervention Trial. *Diabetes Care*. 1993; 16(2): 434-44.

3. Haffner SM, Lehto S, Rönnemaa T, Pyörälä 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 Eng J Med* 1998; 339(4): 229-34.
4. Malmberg K, Yusuf S, Gerstein HC, *et al*. Impact of diabetes on long-term prognosis in patients with unstable angina and non-Q-wave myocardial infarction: results of the OASIS (Organization to Assess Strategies for Ischemic Syndromes) Registry. *Circulation* 2000; 102(9): 1014-9.
5. Goldberg RB, Mellies MJ, Sacks FM, *et al*. Cardiovascular events and their reduction with pravastatin in diabetic and glucose-intolerant myocardial infarction survivors with average cholesterol levels: subgroup analysis in the Cholesterol and Recurrent Events (CARE) trial. *Circulation* 1998; 98(23): 2513-9.
6. Pyörälä K, Pedersen TR, Kjekshus J, Faergeman O, Olsson AG, Thorgeirsson G. Cholesterol lowering with simvastatin improves prognosis of diabetic patients with coronary heart disease: a subgroup analysis of the Scandinavian Simvastatin Survival Study (4S). *Diabetes Care* 1997; 20(14): 614-20.
7. Rubins HB, Robins SJ, Collins D, Fye CL, Anderson JW, Elam MB, *et al*. Gemfibrozil for the secondary prevention of coronary heart disease in men with low levels of high-density lipoprotein cholesterol. *N Eng J Med* 1999; 341(6): 410-8.
8. Expert panel on detection, evaluation, and treatment of high blood cholesterol in adults. Executive summary of the third report of the National Cholesterol Education Program (NCEP) expert panel on detection, evaluation, and treatment of high blood cholesterol in adults (Adult Treatment Panel III). *JAMA* 2001; 285(19): 2486-97.

9. Sarawak Online. Available at: <http://www.sarawak.gov.my/contents/population/population.shtml>. Accessed 28 June 2005.
10. *Laporan Penilaian Prestasi Program Kesihatan Tahunan (January-December) 2001/2002*. Klinik Kesihatan Tanah Puteh, Kuching.
11. Alberti KCM, Zimmet PZ. For the WHO Consultation. Definition, diagnosis and classification of diabetes mellitus and its complications. Part 1: Diagnosis and classification of diabetes mellitus. Provisional report of a WHO Consultation. *Diabet Med* 1998; 15(7): 539-53.
12. The Asian-Pacific Type 2 Diabetes Policy Group. Type 2 Diabetes, Practical Targets and Treatments, 3rd ed. 2002: WHO. IDF.
13. *Practise Guidelines for Diabetes Mellitus Type 2 (NIDDM)*. The Malaysian Consensus 2000. Kuala Lumpur: Ministry of Health Malaysia.
14. *Consensus Statement on Management of Hyperlipidaemia*, 2nd Edition. 2000. Kuala Lumpur: Ministry of Health Malaysia.
15. *Clinical Practice Guidelines on the Management of Hypertension 2002*. Kuala Lumpur: Ministry of Health Malaysia.
16. Wong JS, Rahimah N. Glycaemic control of diabetic patients in an urban primary health care setting in Sarawak: The Tanah Puteh Health Centre experience. *Med J Malaysia* 2004; 59(3): 411-17.
17. Wong JS. Proteinuria in diabetic patients in a primary health care setting in Sarawak. *Med J Malaysia* 2005; 60(2): 146-150.
18. Gæde P, Vedel P, Larsen N, Jensen GVH, Parving HH, Pedersen O. Multifactorial intervention and cardiovascular disease in patients with Type 2 Diabetes. *N Eng J Med* 2003; 348(5): 383-89.
19. The Diabetes Control and Complications Trial Research Group. The effect of intensive treatment of diabetes on the development and progression of long-term complications in insulin-dependent diabetes mellitus. *N Eng J Med* 1993; 329(14): 977-86.
20. UK Prospective Diabetes Study Group. Tight blood pressure control and risk of macrovascular and microvascular complications in type 2 diabetes, UKPDS 38 UK. *Prospective diabetes study group*. *BMJ* 1998; 317(7,160): 703-13.
21. Assmann G, Schulte H. The prospective cardiovascular Munster (PROCAM) Study: Prevalence of hyperlipidaemia in persons with hypertension and/or diabetes mellitus and the relationship to coronary heart disease. *Am Heart J* 1988; 116(2 Part2): 1,713-24.
22. Heart Protection Study Collaborative Group. MRC/BHF Heart Protection study of cholesterol lowering with simvastatin in 20 536 high-risk individuals: a randomised placebo controlled trial. *Lancet* 2002; 360(9,326): 7-22.
23. Colhoun HM, Betteridge DJ, Durrington PN, *et al*. Primary prevention of cardiovascular disease with atorvastatin in type 2 diabetes in the Collaborative Atorvastatin Diabetes Study (CARDS): multicentre randomised placebo-controlled trial. *Lancet* 2004; 364: 685-696.
24. American Diabetes Association. Position Statement: Standards of medical care in diabetes. *Diabetes Care* 2005; 28(Suppl. 1): S4-36.
25. Grundy SM, Cleeman JI, Merz CN, Brewer HB Jr, Clark LT, Hunninghake DB. Implications of recent clinical trials for the National Cholesterol Education Program Adult Treatment Panel III Guidelines. *Circulation* 2004; 110(2): 227-239.
26. Saaddine JB, Engelgau MM, Beckles GL, Gregg EW, Thompson TJ, Narayan KM. A diabetes report card for the United States: quality of care in the 1990s. *Ann Intern Med* 2002; 136(8): 565-74.
27. Massing MW, Foley KA, Sueta CA, *et al*. Trends in lipid management among patients with coronary artery disease: has diabetes received the attention it deserves? *Diabetes Care* 2003; 26(4): 991-7.
28. Pearson TA, Laurora I, Chu H, Kafonek S. The lipid treatment assessment project (L-TAP): a multicenter survey to evaluate the percentages of dyslipidemic patients receiving lipid-lowering therapy and achieving low-density lipoprotein cholesterol goals. *Arch Intern Med* 2000; 160(4): 459-67.
29. Beaton SJ, Nag SS, Gunter MJ, Gleeson JM, Sajjan SS, Alexander CM. Adequacy of glycemic, lipid, and blood pressure management for patients with diabetes in a managed care setting. *Diabetes Care* 2004; 27(3): 694-8.
30. Kemp TM, Barr EL, Zimmet PZ, *et al*. Glucose, lipid, and blood pressure control in Australian adults with type 2 diabetes: the 1999-2000 AusDiab. *Diabetes Care* 2005; 28(6): 1,490-2.
31. Putzer G, Roetzheim R, Ramirez AM, Sneed K, Brownlee HJ Jr, Campbell RJ. Compliance with recommendations for lipid management among patients with type 2 diabetes in an academic family practice. *J Am Board Fam Pract* 2004; 17(2): 101-7.
32. EUROASPIRE I and II Group; European action on secondary prevention by intervention to reduce events. Clinical reality of coronary prevention guidelines: a comparison of EUROASPIRE I and II in nine countries. European action on secondary prevention by intervention to reduce events. *Lancet* 2001; 357(9,261): 995-1001.
33. Parris ES, Lawrence DB, Mohn LA, Long LB. Adherence to statin therapy and LDL cholesterol goal attainment by patients with diabetes and dyslipidemia. *Diabetes Care* 2005; 28(3): 595-9.