A Consensus of Key Opinion Leaders on the Management of Pre-diabetes in the Asia-Pacific Region

Roberto Mirasol,1 Ah Chuan Thai,2 Aftab Ahmad Salahuddin,3 Kathryn Tan,4 Chaicharn Deerohonawanong,5 Mafauzy Mohamed,6 Made Ratna Saraswati,7 Bipin Kumar Sethi,8 Sanjiv Shah,9 Nanny Natalia Soetedjo,10 Swangjit Suraamornkul,11 Rima Tan,12 Farid Uddin13

1St. Luke’s Medical Center, Philippines
2Department of Medicine, Yong Loo Lin School of Medicine, National University of Singapore, Singapore
3Jinnah Teaching Hospital, Pakistan
4Department of Medicine, University of Hong Kong, Hong Kong
5Rajavithi Hospital, Ramgosit University, Thailand
6Universiti Sains Malaysia, Malaysia
7Division of Endocrinology and Metabolism, Sanglah General Hospital, Department of Internal Medicine, Faculty of Medicine, Udayana University, Denpasar, Bali, Indonesia
8CARE Hospital, Hyderabad, India
9Navasari Superspeciality Hospital, Mumbai, India
10Division of Endocrinology and Metabolism, Hasan Sadikin General Hospital, Department of Internal Medicine, Faculty of Medicine, University of Pajajaran, Bandung, Indonesia
11Faculty of Medicine, Vajira Hospital, Navamindradhipit University, Bangkok, Thailand
12Institute for Studies on Diabetes Foundation, Inc., Philippines
13National Institute of Diabetes and Endocrinology, Dow University of Health Sciences, Pakistan

Abstract

The Asia-Pacific region carries a high disease burden, with over half of the global diabetic population residing in this region. Increasing evidence shows that without targeted intervention, the progression from impaired glucose tolerance (IGT) to type 2 diabetes occurs more frequently in Asians compared with Caucasians. Furthermore, IGT is independently associated with an increased risk of cardiovascular disease, and should be managed as early as possible. Because diabetes is now a major public health issue, strategies aimed at prevention and treatment are urgently required. Lifestyle modification, including weight loss, dietary changes and increased physical activity, play a major role in controlling the disease. Significant evidence also supports the effectiveness of a combination of lifestyle modification and pharmacologic therapy, such as metformin, in delaying the onset of diabetes. Although the importance of lifestyle interventions is well recognized throughout Asia, many countries do not have formal recommendations to guide the diagnosis and management of individuals at risk of progression to diabetes. At a recent regional meeting, experts from the Asian region convened to develop consensus recommendations to guide clinicians in the management of Asian patients with pre-diabetes. These consensus recommendations provide a clear and concise approach to the management of individuals with IGT based on the available evidence and current best clinical practice.

Key words: impaired glucose tolerance, pre-diabetes, Asia

INTRODUCTION

Globally, there are 387 million individuals living with type 2 diabetes, 46.3% of whom are undiagnosed.1 A disproportionate diabetes burden is carried by the South East Asian and Western Pacific nations, as it affects 75 million people in South East Asia (8.3% of the adult population) and 138 million in the Western Pacific (8.5% of the adult population).1 With this rising trend, it is estimated that Asia will contribute to more than 60% of the world’s diabetic population.2 Furthermore, Asians have a strong ethnic and genetic predisposition for diabetes and lower thresholds for environmental risk factors.2 Within the Asian region, diabetes and pre-diabetes are more prevalent among Indians (37.9 and 18.9%, respectively) and Malays (23.8 and 22.6%, respectively). Pre-diabetes is more prevalent in women (21.9%) and urban dwellers (21.5%).3

Of particular concern is the evidence indicating that a significant proportion of individuals with diabetes or pre-diabetes are unaware of their condition. Consequently, adoption of risk reduction behavior is suboptimal. An evaluation of the United States National Health and...
Nutrition Examination Survey (NHANES 2005-2006) data revealed that almost 30% of the adult population had pre-diabetes but only 7.3% were aware of their condition. Further, estimates for the prevalence of undiagnosed diabetes in South East Asia and the Western Pacific region were high (52.8 and 53.6%, respectively). A cross-sectional survey of Malaysians found that the prevalence of newly diagnosed type 2 diabetes reached 12.6% adults, with an increasing prevalence of undiagnosed diabetes observed with age. This high prevalence of undiagnosed diabetes poses a serious problem for public health, warranting an escalated effort to address and improve the burden of diabetes.

**METHODOLOGY**

The consensus working group consisted of an assembly of 13 regional experts working as academics, researchers, clinicians, and policy makers from national and international organizations. The experts have substantial knowledge in relevant disciplines, including endocrinology and metabolism.

The expert group used a qualitative approach involving a question-and-answer format to shape and direct the flow of the discussions. This was followed by a comprehensive literature review of published academic articles for identifying the research evidence to guide recommendations. The research findings were subsequently triangulated and circulated electronically among all consensus group members.

The recommendations were formulated by the chairperson and members of the advisory board committee, after the initial group discussion and multiple e-mail communications. The diagnostic cut-off fasting plasma glucose (FPG), glycated haemoglobin (HbA1c), and oral glucose tolerance test (OGTT) values for pre-diabetes are based on the consensus-based outcomes of that meeting and the subsequent literature review.

Consensus was a priori defined as agreement of a large majority of advisory group members, without strong disagreements. If consensus was not reached, the working group would take a vote, where at least a simple majority vote would be required for the recommendation to pass. Any dissenting opinion would be captured and presented in the report. Every effort was made to achieve consensus among the committee members, and consensus was reached on every recommendation.

A preliminary review of the literature showed that several consensus statements exist, including the official position of the American Association of Clinical Endocrinologists and American College of Endocrinology, and another by the working group of the Spanish Diabetes Society. However, this consensus statement was written to reflect the various aspects of pre-diabetes management from an Asian perspective. Majority of the consensus statements were crafted based on studies conducted in Western populations without consideration of the cultural nuances that are sensitive and specific to Asian populations.

**Recommendations for Pre-diabetes Management in Asian Patients**

At a regional consensus meeting held in Manila, Philippines on June 6, 2014 sponsored by Merck, experts from the Asia-Pacific region convened to develop consensus recommendations to guide clinicians in the management of Asian patients with pre-diabetes. The following are their recommendations:

**Recommendation 1: Screening and Diagnosis of Pre-diabetes**

Screening and treatment of IGT can delay or prevent the development of type 2 diabetes, providing a window of opportunity for primary prevention of diabetes and CVD. The authors recommend targeted screening for patients aged ≥35 years and/or high risk individuals, followed by laboratory tests (i.e., FPG, HbA1c and/or 75-gram OGTT). High risk individuals include overweight or obese patients (country-specific), family history of diabetes, high blood pressure, dyslipidemia, history of large babies or gestational diabetes.

Pre-diabetes is diagnosed if the FPG ≥100 mg/dL (5.5 mmol/L), 2-hour OGTT is 140-199 mg/dL (7.8-11.0 mmol/L), and HbA1c of >5.7%. If initial screening of pre-diabetes is negative, patients should be rescreened every 1 to 3 years, depending on risk factors (based on clinical judgment) and local resource availability. The optimal cut-offs for diagnosing pre-diabetes in Chinese patients are HbA1c of 5.6% (38 mmol/mol) in the young and middle-aged and 5.7% (39 mmol/mol) in the elderly.

The bases for the cut-off of 35 years and older for routine screening were reports from Bangladesh and Eastern Uganda that these individuals could have abnormal glucose regulation with a normal body mass index (BMI). Together with majority consensus voting, these formed the rationale for periodic screening of all persons older than 35 years for pre-diabetes. The HbA1c threshold set at 5.7% when screening for pre-diabetes was based on a previously published cost-effectiveness strategy. Therefore, the authors recommend age-specific cut-offs for detecting pre-diabetes or diabetes in populations where such differences have been observed.

**Recommendation 2: Treatment of Pre-diabetes**

Current guidelines on diabetes prevention recommend intensive lifestyle intervention as the cornerstone of pre-diabetes management. The authors strongly recommend lifestyle intervention, preferably with a dietitian referral specifically encompassing the following:

- Reduced intake of simple sugars
- Reduced fat intake, specifically saturated fats and oils
- Reduced trans fatty acid intake
5 to 10% weight loss from baseline
Total caloric intake deficit based on target weight loss
30 minutes of exercise 5 to 7 times per week
Patients should ideally undergo a review after a period of 3 to 6 months.

**Recommendation 3: Pharmacologic Therapy**
Pharmacologic intervention is recommended if there is inadequate response to lifestyle intervention after 3 to 6 months. Metformin should be initiated at a starting dose of 500 mg/day titrated up to a maximum of 2,000 mg/day as required. Alternative treatment should be considered if the patient is nonresponsive or intolerant to metformin (e.g., acarbose), or when it is contraindicated. Follow up is recommended at 3 to 6 months. These recommendations are in line with international and local guidelines and reflect current practice within the region.12-15,17 However, important considerations concerning pharmacotherapy with metformin for high-risk individuals should be emphasized, because the impact of duration of therapy with metformin, and long term cost-effectiveness of such early intervention, remain unclear.19

**Burden Attributable to Pre-diabetes in Asia**
Individuals with impaired fasting glucose (IFG) and/or IGT are considered to have pre-diabetes, indicating a relatively high risk for future development of diabetes.19 IfG and IGT should not be viewed as clinical entities in their own right but rather as risk factors for diabetes as well as cardiovascular disease (CVD).19 Both IFG and IGT are associated with obesity or adiposity (especially abdominal or visceral obesity), dyslipidemia with high triglycerides and/or low high density lipoprotein (HDL) cholesterol, and hypertension.

Increasing evidence shows that without targeted intervention, the progression from IGT to type 2 diabetes occurs more frequently in Asians compared with Caucasians.20-23 The population-based controlled Da Qing study (n=110,660, aged 25 to 74 years), which explored the incidence of progression to diabetes among 577 IGT subjects, reported an incidence rate of 15.7% per 100 person-years in Chinese individuals with IGT.20 In Indians, the progression rate of pre-diabetes to diabetes was found to be as high as 18.3% per year.21 These data are in stark contrast to Finnish (with an average progression rate of 6% per year) and American (11% per 100 person-years) individuals with pre-diabetes.22,23

The prevalence of overweight/obesity (BMI ≥25 kg/m²) is increasing rapidly in the Asia Pacific region. It is important to note that a significant proportion of adults have undiagnosed diabetes, suggesting that the prevalence within the region is likely to be higher than is currently reported (Table 1).12,24 An IGT comparative prevalence in excess of 10% has been reported in Malaysia (15.19%), Hong Kong (13.30%), Japan (12.64%), Singapore (12.40%) and Taiwan (11.19%).24

| Table 1. Estimated prevalence of DM and IGT in Asia compared with the US and the UK^
|---|---|---|
| Country | DM comparative prevalence, 2014 (%) | Adults with undiagnosed DM, 2014 (in 10,000s) | IGT comparative prevalence, 2013 (%) |
|---|---|---|---|
| Hong Kong | 7.66 | 307 | 13.30 |
| Indonesia | 6.03 | 4,854 | 9.38 |
| Japan | 5.12 | 3,891 | 12.64 |
| Malaysia | 17.64 | 1,717 | 15.19 |
| Mainland China | 8.64 | 51,273 | 5.16 |
| Philippines | 6.71 | 1,743 | 6.61 |
| Republic of Korea | 5.97 | 1,493 | 8.75 to 8.45 |
| Singapore | 10.79 | 288 | 12.40 |
| Taiwan | 8.3 | 948 | 11.19 |
| Thailand | 7.36 | 2,223 | 7.88 |
| Vietnam | 5.71 | 1,776 | 1.0 |
| United Kingdom | 3.9 | 834 | 6.63 |
| United States | 9.39 | 7,143 | 12.37 |

1. DM, diabetes mellitus
2. IGT, impaired glucose tolerance
3. US, United States
4. UK, United Kingdom
5. North Korea
6. South Korea

In Thailand, it is estimated that 4.1 million adults have IGT.24 India is estimated to have 21.5 million adults with IGT, making it the country with the highest number of persons with diabetes in the South East Asian region.24,25 Almost one-third of adults with IGT are younger than 40 years of age. It is projected that by 2035, one in 8 adults will have IGT, while one in 10 will have diabetes. Therefore, prompt and effective IGT management is essential.24 A similar pattern is emerging in the Philippines where the prevalence of pre-diabetes (i.e., combined IFG and IGT) was 31.3%, while it was 17.5% for IFG and 23.9% for IGT.26

Of note, the high prevalence of IGT throughout the region indicates that the incidence of diabetes is likely to increase over the coming decade. The alarming prevalence of pre-diabetes (IGF and IGT) and diabetes in the Asian region warrants urgent strategies aimed at preventing the conversion from pre-diabetes to diabetes.27

**Current Management of Patients with Pre-diabetes**
Not everyone with IGT will subsequently develop type 2 diabetes. A large body of evidence supports the effectiveness of lifestyle interventions, specifically healthy diet and physical exercise, to prevent progression to diabetes.20,22,23 The primary aim of lifestyle interventions is to prevent or delay the development of type 2 diabetes and its complications by targeting obesity and physical inactivity, which are the two most important modifiable risk factors of diabetes development.

The beneficial effect of lifestyle interventions has been confirmed in Asian populations. Among 577 Chinese individuals with IGT, long-term lifestyle interventions involving diet and/or exercise have been shown to significantly decrease the incidence of diabetes.20 Over a 6-year period, there was a reduction in the incidence of
diabetes in individuals practicing lifestyle interventions. Compared with the control group, both diet and exercise resulted in reductions in the risk of developing diabetes (31% and 46%, respectively).\(^\text{20}\) These findings were replicated in the Finnish Diabetes Prevention Study (DPS) in 522 overweight (mean BMI $\geq$31 kg/m\(^2\)) adults aged 40 to 65 years with IGT.\(^\text{22,28}\) At 4 years, the cumulative incidence of diabetes was 11% in the intervention group compared with 23% in the control group.\(^\text{21}\) From these results the authors concluded that one case of diabetes could be prevented by treating 22 IGT patients with lifestyle intervention for one year, or 5 patients for a period of 5 years.\(^\text{22}\) The Diabetes Prevention Program (DPP) in the US, one of the largest randomized controlled clinical trials to date, found similar results. In a population of 3234 adults with IGT and/or IFG (with a mean BMI of 34.0 kg/m\(^2\) and a mean age of 51 years), the incidence of diabetes for the lifestyle intervention group was lower than in the placebo group (4.8 and 11.0 cases per 100 person-years, respectively).\(^\text{23}\) The authors concluded that one case of diabetes could be prevented by treating 7 patients with intensive lifestyle modification for 3 years.\(^\text{23}\) These studies found that type 2 diabetes can be prevented by lifestyle changes in those deemed to be at high-risk, such as those with IGT (Table 2).\(^\text{20,22,23}\)

The DPP study also showed that treatment with metformin could also delay or prevent type 2 diabetes. Compared with placebo, metformin reduced the incidence of type 2 diabetes by 31% (Table 2).\(^\text{23}\) A subsequent washout study showed that approximately one-quarter of this effect could be accounted for by the pharmacologic effect of metformin that disappeared following discontinuation of the therapy.\(^\text{29}\) Notably, even after the washout period, a significant 25% reduction in the incidence of diabetes persisted.\(^\text{29}\) Although the DPP study was not powered to demonstrate statistical significance for between subgroup effects, analysis revealed that metformin was more effective in patients with higher fasting plasma glucose (FPG) levels ($\geq$110 mg/dL), in those younger than 60 years of age, and in individuals with a BMI $\geq$35 kg/m\(^2\).\(^\text{23}\)

Despite the benefits of delaying or preventing the onset of diabetes, it has been shown that metformin is rarely prescribed as preventive therapy in working-age adults with pre-diabetes. Over a period of 3 years, only a minority of US adults with pre-diabetes (3.7%, or one in 27) were prescribed metformin, and only 7.8% (fewer than one in 12) of high-risk patients as identified by the national guidelines received metformin in a retrospective cohort study.\(^\text{30}\) These findings highlight the need for intensive lifestyle modification programs. Patients should, at a minimum, be educated on the benefits of metformin and should ideally also be offered this option as preventive treatment for diabetes.

Acarbose has also been shown to effectively reduce the risk of progression to diabetes in individuals with IGT (Table 2).\(^\text{31}\) The STOP-NIDDM trial reported a decrease in progression to diabetes by 25% of patients with IGT.\(^\text{31}\) Patients on acarbose additionally improved in their tolerance to glucose as the probability of reversing to normal glucose tolerance was significantly higher in these patients than in those on placebo ($p<0.0001$).\(^\text{31}\)

### Positions on Specific Questions Addressed to Guide Recommendations

1. **Do prevention interventions have sustained effects?**

A 20-year follow-up of the Da Qing study determined that combined lifestyle intervention resulted in a 51% reduction in the incidence of diabetes during active intervention and a 43% reduction over 20 years (Table 2).\(^\text{32}\) The benefits of the active lifestyle intervention translated to an average delay of diabetes onset of 3.6 years.\(^\text{32}\) The Da Qing study found that the 20-year cumulative diabetes incidence was 93% in the controls versus 80% among those who received the combined lifestyle intervention.\(^\text{32}\) The 20-year follow-up did not detect significant differences in the incidence of first CVD events, CVD mortality, or all-cause mortality between the combined lifestyle intervention and control group, as it was not powered to detect statistical differences in these outcomes.\(^\text{32}\) Nevertheless, the follow-up study showed that lifestyle interventions over 6 years

### Table 2. Overview of lifestyle and pharmacologic interventions in individuals with IGT\(^a\) or IFG\(^b\)

| Intervention | Study | Patient Characteristics | N | Duration Main Study and Follow-up (years) | Risk Reduction Main Study and Follow-up (%) |
|--------------|-------|-------------------------|---|------------------------------------------|------------------------------------------|
| **Lifestyle** | Da Qing\(^\text{20}\) | IGT | 577 | 6 | 31 - 46 |
| | Finnish DPS\(^\text{22,28}\) | IGT | 522 | 3.2 | 43 |
| | DPP/DPPOS\(^\text{23,34}\) | IGT and/or IFG | 3324 | 2.8 | 58 |
| **Pharmacologic** | DPP/DPPOS\(^\text{23,34}\) | IGT and/or IFG | 3234 | 2.8 | 31 |
| | Acarbose | STOP-NIDDM\(^\text{31}\) | 14 | 3.3 | 25 |

\(^a\)IGT, impaired glucose tolerance  
\(^b\)IFG, impaired fasting glucose  
\(^c\)DPS, Diabetes Prevention Study  
\(^d\)DPP, Diabetes Prevention Program  
\(^e\)DPPOS, Diabetes Prevention Program Outcomes Study  
\(^f\)STOP-NIDDM, The Study to Prevent Non-insulin Dependent Diabetes Mellitus

---

**Vol. 32 No. 1 May 2017**

---

www.asean-endocrinejournal.org
were able to delay or prevent diabetes onset for up to 14 years after active intervention ceased.\textsuperscript{32}

The most recent 23-year follow-up of the Da Qing study confirmed that active lifestyle intervention significantly reduced the risk of CVD and all-cause mortality. The cumulative incidence of CVD mortality was 11.9\% for patients in the lifestyle intervention group and 19.6\% in the control group. In terms of all-cause mortality, the cumulative incidence was 28.1\% in the lifestyle intervention and 38.4\% in the control group.\textsuperscript{33} The significant differences in the incidence of diabetes between the two groups persisted during the 23-year follow-up: the cumulative incidence was 72.6\% in the intervention group and 89.9\% in the control group.\textsuperscript{33} These findings justify the adoption of lifestyle interventions in patients with IGT.

2. Are we preventing type 2 diabetes or delaying it?
Interventions that may prevent or delay IGT, which is associated with cardiovascular disease and conversion to type 2 diabetes, are clinically important. The 10-year follow-up of the DPP, the Diabetes Prevention Program Outcomes Study (DPPOS), found that patients who were on intensive lifestyle had a 34\% reduction in the incidence rate of diabetes, with an average delay of diabetes progression by about 4 years versus placebo (Table 2).\textsuperscript{34} Those treated with metformin had an 18\% reduction in diabetes incidence rate, with the onset of diabetes delayed by an average of 2 years.\textsuperscript{34}

It is clear that high risk patients must be identified and lifestyle changes should be implemented and sustained over the long term. Delaying or preventing type 2 diabetes is cost-effective and will help turn the tide in the diabetes epidemic.\textsuperscript{35}

3. What is the current management of pre-diabetes in countries of the Asia-Pacific region?
Most countries within the region do not have country-specific guidelines and therefore follow the American Diabetes Association (ADA) or the International Diabetes Federation (IDF) recommendations.\textsuperscript{12,13} The IDF consensus guidelines on the prevention of type 2 diabetes recommend the following three steps for the prevention of diabetes development: (1) identification of individuals at high risk of developing diabetes, (2) assessment of risk levels by measuring plasma glucose levels, and (3) initiation of lifestyle interventions with or without pharmacologic therapy.\textsuperscript{36} Once individuals with prediabetes have been identified, they are advised to undergo structured lifestyle modifications, with the aim of achieving gradual and sustained weight loss and maintaining a healthy body composition through physical activity and change of dietary habits.\textsuperscript{36} The World Health Organization (WHO) and IDF also recommend to address other risk factors, including smoking.\textsuperscript{13,37} In addition, the WHO highlights the need for a global approach to reduce the growing global burden of diabetes.\textsuperscript{37}

The ADA recommends the referral of patients with IGT, IFG or HbA1c of 5.7 to 6.4\% to an ongoing support program targeting weight loss of 7\% of body weight and moderate exercise of ≥150 minutes per week.\textsuperscript{12} The ADA states that metformin therapy for prevention of type 2 diabetes may be considered in individuals with IGT, IFG or an HbA1c of 5.7 to 6.4\%, especially for those with BMI >35 kg/m\(^2\), individuals aged <60 years, and women with prior gestational diabetes.\textsuperscript{12} In addition, the ADA guidelines encompass recommendations for follow-up counselling for successful lifestyle interventions, annual monitoring of individuals with pre-diabetes for development of diabetes, as well as screening for and treating of modifiable risk factors for CVD.\textsuperscript{12}

In Asia-Pacific countries, lifestyle modifications remain the mainstay of recommended first-line interventions for patients with pre-diabetes. In Malaysia, lifestyle interventions, such as diet and physical therapy, are the pillars for pre-diabetes therapy.\textsuperscript{14} In addition, the 2015 Ministry of Health (MOH) guidelines state that metformin (as the preferred first-line oral anti-diabetic agent) should be considered for patients at very high risk of progressing to diabetes (combined IFG and IGT, IGT plus other risk factors, or failed lifestyle intervention after 6 months).\textsuperscript{14} Off-label metformin may be initiated at the discretion of the prescribing physician.

In Thailand, there are no available guidelines for the management of pre-diabetes. Majority of physicians follow the IDF recommendations, ADA guidelines or findings from randomized controlled trials on pre-diabetes prevention.\textsuperscript{12,13,20,22,34} Likewise, given a lack of guidelines in the Philippines, physicians generally follow the ADA recommendations. However, in the Philippines, metformin is approved for the treatment of pre-diabetes after failed lifestyle intervention.

Guidelines in Singapore indicate that lifestyle modification should be the first-line treatment of choice.\textsuperscript{15} Metformin may be considered for individuals with a very high risk of progressing to diabetes, particularly patients with IFG, IGT, less than 60 years of age, or BMI ≥35 kg/m\(^2\).\textsuperscript{15} Hong Kong has pre-diabetes management guidelines aimed at the primary care sector.\textsuperscript{16} The emphasis is mainly on lifestyle modifications using dietary or behavioral interventions to reduce and maintain body weight and practice healthy lifestyle. Pharmacologic therapy is not routinely recommended at present.\textsuperscript{16}

The Indonesian guidebook on the management of prediabetes and prevention of type 2 diabetes 2009 [Buku Panduan - Pengurus Besar Persatuan Diabetes Indonesia (PB Persadia)] states that the diabetes prevention strategy should encompass a three-step process that includes identification of high-risk individuals (step 1), risk calculation, (step 2) and intervention (step 3).\textsuperscript{17} Step 3 involves lifestyle changes, body weight management
(reduction by 5 to 7% of baseline body weight, 0.5 to 1 kg/week), physical activity and pharmacologic intervention. The latter involves either metformin given 250 to 850 mg twice daily in individuals 60 years or younger, with BMI >25 kg/m² and FBS >110 mg/dL (6.1 mmol/L) if no contraindications are present; or acarbose 50 to 100 mg thrice daily.17

Within the region, despite the lack of formal recommendations, metformin is often used off-label for certain patient populations. For example, there are no pre-diabetes guidelines in India, but metformin is used off-label by physicians if pre-diabetes patients require pharmacotherapy. Pakistan also has no formal guidelines for pre-diabetes treatment. In view of long-standing safety information about metformin, this drug is prescribed to individuals who are noncompliant with lifestyle interventions. For other potential drugs, further long-term studies are needed on safety and vascular outcomes before lifelong treatment can be safely recommended.

CONCLUSIONS

Pre-diabetes represents a window of opportunity to prevent or delay the progression to diabetes and its associated complications, underscoring the critical need for screening at the primary care level. Lifestyle modification including weight loss, dietary changes and increased physical activity play a major role in controlling the disease. Furthermore, significant evidence support the effectiveness of combining lifestyle modification and pharmacologic therapy on certain patient populations in delaying the onset of diabetes. A cost-effectiveness analysis of lifestyle intervention and metformin therapy for the prevention of diabetes in Singapore concluded that both lifestyle modification and metformin are likely to be cost-effective and worth implementing in Singapore to prevent or delay the onset of type 2 diabetes.35 Although the importance of lifestyle interventions is well recognized throughout Asia, many countries do not have formal recommendations to guide the diagnosis and management of individuals at risk of progression to diabetes.

Overall, these consensus recommendations provide a clear and concise approach to the management of individuals with IGT based on the available evidence and current best clinical practice. Furthermore, local applicability of these recommendations will be far-reaching, particularly in guiding action and policy for pre-diabetes and other related endocrine and metabolic disorders at the regional, national and local levels.

Acknowledgements

The authors would like to thank Merck Pte Ltd (Singapore) for providing an educational grant in support of the consensus meeting. Meeting and editorial support was provided by MIMS Pte Ltd through this grant.

Statement of Authorship

All authors certified fulfillment of ICMJE authorship criteria.

Author Disclosure

The authors have received honoraria for their participation in the APAC Regional Pre-Diabetes Advisory Board Meeting 2014. Professor Mirasol is an advisory board and speaker for AstraZeneca, Merck, Novo, Genzyme, Lilly, MSD.

Funding Source

Merck Pte Ltd (Singapore) provided logistic support and funding for the consensus meeting. The authors received fair market honorarium for their time spent attending the consensus meeting and developing the manuscript. Merck Pte Ltd (Singapore) also paid for MIMS Pte Ltd to provide editorial support. RM, ACT, AA, KT, CD, MM, MRS, BKS, SS, NNS, SS, RTI and FU reported personal fees from Merck Pte Ltd (Singapore) during the preparation of this manuscript.

References

1. International Diabetes Federation. Diabetes Atlas 6th edition poster update, 2014. http://www.idf.org/sites/default/files/Atlas-poster-2014_EN.pdf. Accessed September 22, 2016.
2. Ramachandran A, Snehalatha C, Ma RC. Diabetes in South-East Asia: an update. Diabetes Res Clin Pract. 2014;103(2):231-7. PMID: 24300015. https://doi.org/10.1016/j.diabres.2013.11.011.
3. Wan Nazaimoon WM, Md Isa SH, Wan Mohamad WB, et al. Prevalence of diabetes in Malaysia and usefulness of HbA1c as a diagnostic criterion. Diabet Med. 2013;30(7):825-8. PMID: 23419441. https://doi.org/10.1111/dme.12161.
4. Geiss LS, James C, Gregg EW, Albright A, Williamson DF, Cowie CC. Diabetes risk reduction behaviors among U.S. adults with prediabetes. Am J Prev Med. 2010;38(4):403-9. PMID: 20307809. https://doi.org/10.1016/j.amepre.2009.12.001.
5. Mustafa N, Kamarudin NA, Ismail AA, et al. Prevalence of abnormal glucose tolerance and risk factors in urban and rural Malaysia. Diabetes Care. 2011;34(6):1362-4. PMID: 21496788. PMCID: PMC3114358. https://doi.org/10.23736/dci.311.005.
6. Garber AJ, Abrahamson MJ, Barzilay JL, et al. Consensus statement by the American Association of Clinical Endocrinologists and American College of Endocrinology on the comprehensive type 2 diabetes management algorithm—2016 Executive Summary. Endocr Pract. 2016;22(1):84-113. PMID: 26731084. https://doi.org/10.4158/EP151126-CS.
7. Mata-Cases M, Artola S, Escalada J, et al. Consensus on the detection and management of prediabetes. Consensus and Clinical Guidelines Working Group of the Spanish Diabetes Society. Rev Clin Esp. 2015;215(2):117-29. PMID: 25553948. https://doi.org/10.1016/j.ree.2014.10.012.
8. Yan ST, Xiao HY, Tian H, et al. The cutoffs and performance of glycated hemoglobin for diagnosing diabetes and prediabetes in a young and middle-aged population and in an elderly population. Diabetes Res Clin Pract. 2015;109(2):238-45. PMID: 26059072. https://doi.org/10.1016/j.diabres.2015.05.047.
9. Akter S, Rahman MM, Abe SK, Sultana P. Prevalence of diabetes and prediabetes and their risk factors among Bangladeshi adults: A nationwide survey. Bull World Health Organ. 2014;92(3):204-13.213E. PMID: 24700980. PMCID: PMC3949596. https://doi.org/10.2471/BLT.13.132871.
10. Mayegwa RW, Guwathule D, Makumbi F, et al. Diabetes and prediabetes among persons aged 35 to 60 years in eastern Uganda: Prevalence and associated factors. PLoS One. 2013;8(8):e72554. https://doi.org/10.1371/journal.pone.0072554.
11. Zhuo X, Zhang P, Selvin E, et al. Alternative HbA1c cutoffs to identify high-risk adults for diabetes prevention: a cost-effectiveness perspective. Am J Prev Med. 2012;42(4):374-81.PMID:22424250. https://doi.org/10.1016/j.amepre.2012.01.003.
12. American Diabetes Association. Standards of Medical Care in Diabetes—2015. Diabetes Care. 2015;38(Suppl 1):S1-89.
13. International Diabetes Federation Clinical Guidelines Task Force. Global Guideline for Type 2 Diabetes. Brussels, Belgium: International
Diabetes Federation. 2012. http://www.idf.org/library/guidelines/79-global-guideline-for-type-2-diabetes.html. Accessed September 22, 2016.

14. Ministry of Health Malaysia. Clinical practice guidelines—Management of type 2 diabetes mellitus, 5th ed. Putrajaya, Malaysia: Ministry of Health Malaysia, 2015. http://www.moh.gov.my/index.cfm?menuid=67/Endocrine_Disease.

15. Ministry of Health Singapore. Diabetes Mellitus—MOH Clinical Practice Guidelines 1/2014. Singapore: Ministry of Health, Singapore, 2014. https://www.moh.gov.sg/content/dam/moh_web/HIP/Doctors/cpg_medical/current/2014/diabetes_mellitus/cpg_Diabetes%20Mellitus%20Summary%20Card%20-%202014.pdf.

16. Task Force on Conceptual Model and Preventive Protocols, Working Group on Primary Care, Food and Health Bureau. Hong Kong Reference Framework for Diabetes Care for Adults in Primary Care Settings, 2013. http://www.pcc.gov.hk/english/resources/files/RF_DM_full.pdf.

17. Buku Panduan - Pengurus Besar Persatuan Diabetes Indonesia (PB Persada). 2009.

18. Bansal N. Prediabetes diagnosis and treatment: A review. World J Diabetes 2015;6(2):6-303. PMID: PMC4364022. https://doi.org/10.4239/wjd.v6.i2.6.

19. World Health Organization International Diabetes Federation. Definition and diagnosis of diabetes mellitus and intermediate hyperglycaemia: Report of a WHO/IDF consultation. Geneva, Switzerland: World Health Organization, 2006. http://www.who.int/diabetes/publications/Definition%20and%20diagnosis%20of%20diabetes_new.pdf.

20. Pan XR, Li GW, Hu YH, et al. Effects of diet and exercise in preventing NIDDM in people with impaired glucose tolerance. The Da Qing IGT and Diabetes Study. Diabetes Care. 1997;20(4):537-44. PMID: 9086977.

21. Ramachandran S, Snehalatha C, Mary S, et al. The Indian Diabetes Prevention Programme shows that lifestyle modification and metformin prevent type 2 diabetes in Asian Indian subjects with impaired glucose tolerance (IDPT-1). Diabetologia 2006;49(2):289-97. PMID: 16591903. https://doi.org/10.1007/s00125-005-0097-z.

22. Tuomilehto J, Lindstrom J, Eriksson JG, et al. The Finnish Diabetes Prevention Study: A 23-year follow-up study. Lancet. 2002;359(9323):2072-7. PMID: 12086760. https://doi.org/10.1016/S0140-6736(02)08905-5.

23. Li G, Zhang P, Wang J, et al. The long-term effect of lifestyle interventions to prevent diabetes in the China Da Qing Diabetes Prevention Study: A 20-year follow-up study. Lancet. 2008;371(9626):1753-9. PMID: 18502303. https://doi.org/10.1016/S0140-6736(08)6766-7.

24. Li G, Zhang P, Wang J, et al. Cardiovascular mortality, all-cause mortality, and diabetes incidence after lifestyle intervention for people with impaired glucose tolerance in the Da Qing Diabetes Prevention Study: A 23-year follow-up study. Lancet Diabetes Endocrinol. 2014;2(6):247-54. PMID: 24731674. https://doi.org/10.1016/S2213-8587(14)70057-9.

25. Shaw JE, Sicree RA, Zimmet PZ. Global estimates of the prevalence of diabetes for 2010 and 2030. Diabetes Res Clin Pract. 2010;87(1):4-14. PMID: 19896746. https://doi.org/10.1016/j.diabres.2009.10.007.

26. Soria MB, Sy RG, Vega RS, et al. The incidence of type 2 diabetes mellitus in the Philippines: A 9-year cohort study. Diabetes Res Clin Pract. 2009;86(2):130-3. PMID: 19763434. https://doi.org/10.1016/j.diabres.2009.07.014.

27. Jayawardena R, Ranasinghe P, Byrne NM, Soares MJ, Katulanda P, Hills AP. Prevalence and trends of the diabetes epidemic in South Asia: A systematic review and meta-analysis. BMC Public Health. 2012;12:380. PMID: 22600343. PMCID: PMC344674. https://doi.org/10.1186/1471-2458-12-380.

28. Lindström J, Ilanne-Parikka P, Peltonen M, et al. Sustained reduction in the incidence of type 2 diabetes by lifestyle intervention: Follow-up of the Finnish Diabetes Prevention Study. Lancet. 2006;368(9548):1673-9. PMID: 17098085. https://doi.org/10.1016/S0140-6736(06)69701-8.

29. Diabetes Prevention Program Research Group. Effects of withdrawal from metformin on the development of diabetes in the diabetes prevention program. Diabetes Care. 2003;26(4):977-80. PMID: 12663559. PMCID: PMC166737.

30. Moin T, Li J, Duru OK, et al. Metformin prescription for insured adults with prediabetes from 2010 to 2012: A retrospective cohort study. Ann Intern Med. 2015;162(8):542-8. PMID: 25894024. PMCID: PMC4682357. https://doi.org/10.7326/M14-1773.

31. Chiasson JL, Josse RG, Gomis R, et al. Acarbose for prevention of type 2 diabetes mellitus: the STOP-NIDDM randomised trial. Lancet. 2002;359(9323):2072-7. PMID: 12086760. https://doi.org/10.1016/S0140-6736(02)08905-5.

32. Li G, Zhang P, Wang J, et al. Cardiovascular mortality, all-cause mortality, and diabetes incidence after lifestyle intervention for people with impaired glucose tolerance in the Da Qing Diabetes Prevention Study: A 23-year follow-up study. Lancet Diabetes Endocrinol. 2014;2(6):247-54. PMID: 24731674. https://doi.org/10.1016/S2213-8587(14)70057-9.

33. Diabetes Prevention Program Research Group. 10-year follow-up of diabetes incidence and weight loss in the Diabetes Prevention Program Outcomes Study. Lancet. 2009;374(9702):1677-86. PMID: 19879886. https://doi.org/10.1016/S0140-6736(09)61457-4.

34. Png ME, Yoong SYY. Evaluating the cost-effectiveness of lifestyle modification versus metformin therapy for the prevention of diabetes in Singapore. PLoS One. 2014;9(4):e107225. https://doi.org/10.1371/journal.pone.0107225.

35. Alberti KG, Zimmet P, Shaw J. International Diabetes Federation: A consensus on type 2 diabetes prevention. Diabet Med. 2007;24(5):451-63. PMID: 17470191. https://doi.org/10.1111/j.1464-5491.2007.02157.x.

36. World Health Organization. “Diabetes Fact Sheet No. 312.” Reviewed November 2016. http://www.who.int/mediacentre/factsheets/fs312/en/.