

Dr Paul Rheeder, (MMed) (Int), FCP (SA), MSc (Clinical Epidemiology) Department of Internal Medicine, University of Pretoria

r Paul Rheeder (MMed) (Int), FCP (SA), MSc (Clinical Epidemiology) is a consultant in the Department of Internal Medicine at the University of Pretoria.

He has a specific interest in diabetes and is involved in research in diabetes at Mamelodi Hospital in Pretoria.

This community hospi-

tal reflects accurately the real-world situation of diabetes care for most of South Africa's citizens. The support systems are deficient (lost file, lost results etc.) and the workload excessive.

Dr Rheeder's interest is in providing optimum care in sub-optimum conditions. He is a "hands-on" practical physician working daily with the real problem in the real world.

He understands the circumstances that many GP's work in and offers his practical solutions to practical everyday care.

South Africa's diabetics need his approach and this series of articles will help many clinicians provide an improved service to their diabetics.

### Summary

The prevalence of diabetes in South Africa is approximately 5% amongst Blacks and Caucasians but much higher in the Indian population.

Obesity and physical inactivity are the most amenable risk factors for NIDDM.

Cardiovascular disease is the most impor-

tant coexisting problem and thus both microvascular and macrovascular complications contribute to the increased morbidity and mortality. NIDDM and IDDM are beterogenous groups of diseases with latent onset auto immune diabetes (LADA) added into its classification.

## **EPIDEMIOLOGY OF DIABETES**

The tip of an iceberg...

The incidence of diabetes among school children in the USA is similar to the incidence of all childhood cancers combined.

In this series
of articles various
experts share their
understanding and
management of
medical problems
with us. The
emphasis is on
practical approaches
to the problem
concerned and

**Ivory Tower and** 

the Coalface.

n the USA diabetes has been diagnosed in 8-million adults, approximately 90-95% of whom have non-insulin dependent diabetes mellitus (NIDDM). Studies indicate that at least an additional 8-million people have undiagnosed NIDDM.

In the USA the prevalence of known cases of diabetes is 6-7% for people aged 45-64 years and 10-12% for those aged 65 years and older. Higher rates are found in US minority populations<sup>1</sup>.

On average NIDDM is present for 10-12 years before clinical diagnosis in the United States<sup>2</sup>.

The incidence of diabetes among school children in the USA is similar to the incidence of all childhood cancers combined<sup>3</sup>.

A comprehensive epidemiological survey done in Cape Town in the 1970s showed that diabetes (mainly NIDDM) had an age-adjusted prevalence of 4,2 % amongst urban Blacks, 3,6% amongst Caucasians and 19,7% amongst Asians<sup>4</sup>. *(Table I)* 

The chief risk factors are obesity and

Table I: Modifiable risk factors for NIDDM.

Magnitude	Risk factor
strong (RR $> 4$ )	obesity
moderate (RR 2-4)	none
weak (RR < 2)	physical inactivity

possible smoking,

high fat/low fibre diet

(RR = relative risk)

advancing age. Approximately 80% of NIDDM patients are obese at the time of diagnosis. An increased waist-to-hip ratio appears to predict NIDDM. A lack of physical activity and smoking have also been implicated.

The modified WHO classification gives a clear picture of the heterogeneity of diabetes mellitus. (*Table II*)

It is now clear that IDDM is more common in adults than formerly believed and in fact close to 60% of cases develop after the age of 20 years<sup>12</sup>. This often appears as subjects in the NIDDM group who actually have IDDM but in a slowly evolving form, masquerading as NIDDM at their first presentation<sup>15</sup>

This syndrome was initially called Type 1 1/2 diabetes<sup>14</sup>. Most cases can now be recognised as latent auto immune diabetes in adults (LADA). LADA has been reported from various countries such as the USA<sup>15</sup>, New Zealand<sup>16</sup> and Scandinavia<sup>17</sup>. No data exists concerning Africa, however.

### **Epidemiology of complications**

Again, most of the experience comes from the USA. Their experience shows the following 18.

**Heart disease:** Cardiovascular disease is 2-4 times more common among people with diabetes and it is present in 75% of diabetes-related deaths.

**Stroke:** The risk of stroke is 2-4 times higher among persons with diabetes.

**Hypertension:** An estimated 60-65% of persons with diabetes have high blood pressure.

**Blindness:** In the USA diabetes is the leading cause of new cases of blindness among adults 20-74 years of age.

**Kidney disease:** Diabetes is the leading cause of end stage renal disease, accounting for 30% of new cases.

**Neuropathy:** Approximately 60-70% of people with diabetes have mild to severe forms of neuropathy.

**Amputations:** More than half of lower limb amputations occur among people with diabetes.

**Dental disease:** Periodontal disease, which can lead to tooth loss, occurs with greater frequency and severity among persons with diabetes. In one study, 30% of IDDM patients aged 19 years and over had periodontal disease.

### **Complications in South Africa**

Omar and Asmal<sup>®</sup> studied the prevalence of acute and chronic complications in Blacks and Indians.

Of the 92 Blacks, almost 70% developed ketoacidosis on one or more occasions, whereas 50% of the Indians manifested this complication. Most of the chronic complications were related to the duration of IDDM. Retinopathy was found in 14% of Blacks and 22% of Indian patients, nephropathy in 3% and 7% and neuropathy in 22% and 32% respectively.

In a retrospective study by Motala<sup>20</sup> *et al* to assess the development of microvascular risk factors in African and Indian patients, they found that of the 219 patients (172 NIDDM and 47 IDDM), persistent proteinuria was present in 24,4% of NIDDM and 25,5% of IDDM patients. Hypertension developed in 64,5% of NIDDM and 34% of IDDM patients. Abnormal serum creatinine was observed in 25,1 % of NIDDM and 17,8 % of IDDM patients. ●

# Table II: Modified WHO classification of Diabetes Mellitus and allied categories of glucose intolerance

### A. Clinical classes

Insulin dependent diabetes mellitus (IDDM) Non-insulin dependent diabetes mellitus (NIDDM)

- a) non-obese
- b) obese

Malnutrition-related diabetes mellitus (MRDM)

### Impaired glucose tolerance (IGT)

- a) non-obese
- b) obese
- c) associated with certain syndromes

### Gestational diabetes mellitus (GDM)

Latent autoimmune diabetes in adults (LADA)

### B. Statistical risk classes

Previous abnormality of glucose tolerance.
Potential abnormality of glucose tolerance.
(Normal glucose tolerance but at an increased risk for developing DM)

# C. Other types of diabetes associated with certain conditions and syndromes

- 1. Pancreatic disease
- 2. Diseases of hormonal aetiology
- 3.Drug induced or chemical induced conditions
- 4. Abnormality of insulin or its receptors
- 5. Certain genetic syndromes
- 6. Miscellaneous

# Table III: Features of latent autoimmune diabetes in adults (LADA)

- Age usually > 35 years
- Clinical presentation as NIDDM
- Initial control with diet/oral agents
- Insulin dependency within 1-3 years
- Other features of IDDM:

Low serum C-peptide Island cell antibody positive or anti GAD antibody positive Obesity and physical inactivity are the most amenable risk factors for NIDDM.

#### References

- Harris MI, Cowie CC, Reiber G, Boyko E, Stern M, Bennet P, eds. *Diabetes in America*. 2nd ed. Washington, DC: Government printing Office 1995.
- Harris MI, Klein RE, Welborn TA, Knuiman MW. Onset of NIDDM occurs at least 4-7 yr before clinical diagnosis. *Diabetes Care*. 1992; 15:815-819.
- Laporte RE, Cruikshanks KJ. *Incidence and risk factors for insulin dependent diabetes*. In: Harris MI, Hamman RF, eds. *Diabetes in America*.
  Bethesda, Md: US Dept of Health and Human Services, 1985: III: 1-12. NIH publication 85-1468.
- Jackson WPU. Epidemiology of diabetes in Southern Africa. In: Levine R, Luft R, eds. Advances in Metabolic Disorders, Vol. 9. New York: Academic Press, 1978: 111-46.
- Kaye SA, Folsom AR, Sprafka JM, Prineas RJ, Wallace RB. Increased incidence of diabetes mellitus in relation to abdominal adiposity in older women. J Clin Epidemiol. 1991;44:329 334.
- Helmrich SP, Ragland DR, Leung RW, Paffenbarger RS Jr. Physical activity and reduced occurrence of non-insulin-dependent diabetes mellitus. N Engl J Med. 1991;324: 147-152
- Manson JE, Nathan DM, Krolewski AS, Stampfer MJ, Willett WC, Hennekens CH. A prospective study of exercise and incidence of diabetes among US male physicians. JAMA. 1991;268:63-67
- Rimm EB, Manson JE, Stampfer MJ, et al. Cigarette smoking and the risk of diabetes in women. AM J Public Health. 1993;83:211-214
- Molitch ME. Diabetes mellitus: control and complications. Postgrad Med. 1989;11:258-262.
- Bishop DB, Roesler JS, Zimmerman BR, Ballerd DJ. Diabetes in: Chronic disease epidemiology and control. Brownson RC, Remington PL, Davis JR. eds. Washington DC: American Public Health Association 1993.
- World Health Organization. *Diabetes Mellitus: Second Report*. Technical Report Series No. 626, WHO Geneva, 1980;8-12.

- 12. Zimmet P. The pathogenesis and prevention of diabetes in adults: genes, autoimmunity and demography. Diabetes Care 1995;18:1050-1064
- 13. Tuomi T, Groop LC, Zimmet PZ, Rowley MJ, Knowles W, Mackay IR. Antibodies to glutamic acid decarboxylase reveal latent autoimmune diabetes mellitus in adults with a non insulin dependent onset of disease. Diabetes 1993, 42, 359-62.
- Harris MI, Zimmet P. Classification of diabetes mellitus and other categories of glucose intolerance. In The International Textbook of Diabetes Mellitus, eds, H Keen, R DeFronzo, KGMM Alberti, P Zimmet. London: John Wiley, 1992, 3-18.
- Zimmet PZ, Shaten BJ, Kuller LH, Rowley MJ, Knowles WJ, Mackay IR. Antibodies to glutamic acid decarboxylase and diabetes mellitus in the multiple risk factor intervention trial. Am J Epidemiol 1994, 140, 683-90.
- Scott R, Willis J, Brown L, Forbes L, Schmidzi R, Zimmet P, Mackay I, Rowley M. Antibodies to glutamic acid decarboxylase (GAD) predict insulin-deficiency in adult onset diabetes mellitus. Diabetes 1993, 42(Suppll):220A.
- 17. Hagopian WA, Karlsen AE, Gottsater A, Landin-Olsson M, Grubin CE, Sundkvist G, Petersen JS, Boel E, Dyrberg T, Lernmark A. Quantitative assay using recombinant buman islet glutamic acid decarboxylase (GAD65) shows that 65K autoantibody positivity at onset predicts diabetes type. J Clin Invest 1993, 91, 358-74.
- Reducing the burden of Diabetes: National Diabetes Fact Sheet. 1995. American Diabetes Association.
- Omar MA, Asmal AC. Complications of early onset insulin-dependent diabetes mellitus in Blacks and Indians. S Afr Med J. 1984; 65:75-8
- Motala AA, Omar MAK, Pirie FJ, Amod A. Microvascular Complications in African and Indian subjects with diabetes of long duration (abstract) 32nd SEMDSA. 1996