

## PP\_P002

### CLINICAL CHARACTERISTICS AND OUTCOMES OF PEDIATRIC TYPE 2 DIABETES IN HOSPITAL PUTRAJAYA

<https://doi.org/10.15605/jafes.038.S2.126>

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#### INTRODUCTION

In Malaysia, the estimated prevalence of children and adolescents with type 2 Diabetes mellitus (T2DM) was 17.7% of all paediatric patients with diabetes (DiCARE Annual Report 2006-2007). We report the characteristics and outcomes of patients with type 2 DM in Hospital Putrajaya.

#### METHODOLOGY

Data retrieved from the electronic database were reviewed. All patients diagnosed with type 2 diabetes attending the Paediatric Clinic Hospital Putrajaya from January 2012 until December 2022 were included. Data were presented as mean, median and percentages.

#### RESULT

There was a total of 41 patients reviewed. The mean age of presentation was  $11 \pm 1.9$  years old with a median diabetes duration of  $4.3 \pm 2.1$  years. A majority (68.3%) presented between 10 to < 15 years old, predominantly females (63.4%) and of Malay ethnicity (65.8%). All patients had negative diabetes autoantibodies, 82.9% were obese and 92.6% had positive family history of diabetes. At presentation, 14.6% of patients had diabetic ketoacidosis (DKA). The mean fasting blood sugar was  $11.3 \pm 3.8$  mmol/L and the mean HbA1c was  $11.4 \pm 3.2\%$ . Diabetes-related complications namely hypertension, dyslipidaemia, microalbuminuria and retinopathy were observed in 24.4%, 17.5%, 17.1% and 4.9% of patients, respectively. Dyslipidaemia was detected at  $2.4 \pm 2.6$  years, retinopathy at  $2.5 \pm 0.7$ , hypertension at  $3 \pm 2.3$  years and microalbuminuria was detected at  $3.5 \pm 1.3$  years from onset of DM.

There were 80.6% of patients who required insulin at initial presentations and 74.3% still needed combination therapy with metformin later on. At the end of the follow-up, only 12.1% of patients achieved HbA1c < 6.5%, while 51.2% had HbA1c > 10%.

#### CONCLUSION

Paediatric Type 2 diabetes patients had poor glycaemic control with early development of complications. A multi-disciplinary approach with an individualised management plan is needed to prevent the progression of the disease.