Change of HbA_{1c} reporting to the new SI units

TO THE EDITOR: The position statement by Jones and colleagues regarding the change of HbA_{1c} reporting to the new Système International (SI) units — which has been recommended by the Australasian Association of Clinical Biochemists, the Australian Diabetes Educators Association, the Australian Diabetes Society and the Royal College of Pathologists of Australasia — provides a comprehensive summary of the rationale behind the proposed change and suggests a 2-year period of dual reporting.¹

However, Jones et al did not specify targets for children and adolescents, and we believe that it is important to do so. The incidence of type 1 diabetes in Australian children and adolescents is among the highest in the world² and, in New South Wales, type 2 diabetes represents at least 10% of cases of newonset diabetes in adolescents.³ National evidence-based clinical care guidelines for type 1 diabetes in children, adolescents and adults⁴ include agespecific targets for HbA_{1c}, while recognising that such targets are predominantly consensus based.

 HbA_{1c} targets for young people with type 1 diabetes are higher, with a level of <7.5% recommended for children and adolescents in the Australian guidelines⁴ and in those produced by the International Society for Pediatric and Adolescent Diabetes (ISPAD).⁵ Jones et al note that "Achievement of HbA_{1c} targets must be balanced against risk of severe hypoglycaemia, especially among older people";¹ this is also the case for young people. For children and adolescents with type 2 diabetes, the ISPAD guidelines recommend an HbA_{1c} target of <7%.⁵

The move to SI units represents a major change in the established, widely recognised outcome measure of glycaemia; during the transition period, the specific needs of young people with diabetes must not be forgotten.

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Letters

IN REPLY: We appreciate Craig and colleagues' comments regarding the importance of reporting general HbA_{1c} targets for children and adolescents with type 1 and type 2 diabetes. In our position statement, the headings of Box 2 and Box 3 indicated that the targets listed were for adults with type 1 diabetes and adults with type 2 diabetes, respectively.¹ While it is not possible to highlight every clinical situation, we agree that providing general HbA_{1c} targets for children and adolescents will add value to our article, and we have updated it accordingly,¹ recognising the differences in these targets for type 1 diabetes ($\leq 58 \text{ mmol/mol}, \leq 7.5\%$) and type 2 diabetes ($\leq 53 \text{ mmol/mol}$, \leq 7.0%).²⁻⁴ In the interests of uniformity and simplicity, the paediatric targets expressed as "<" $^{\rm \prime\prime 2-4}$ have been adjusted to " \leq ", which represents differences of less than 1.5% of the target values.

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