

Excretion of polyamines by children with Beckwith's syndrome

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SUMMARY The urinary excretion of the polyamines—putrescine, spermidine, and spermine—was measured in 7 children with Beckwith's syndrome. Putrescine excretion was raised and spermidine excretion reduced. The raised putrescine and the low spermidine ratios were highly significant. These results are consistent with a disturbance in a metabolic pathway under growth hormone-like regulation.

The features of children with Beckwith's syndrome (sometimes known as EMG (exomphalos, macroglossia, gigantism) or Beckwith-Wiedemann syndrome) are macroglossia, somatic gigantism, visceromegaly, exomphalos, and a transient neonatal hypoglycaemia.^{1–2} The disorders in growth suggest that there is some disturbance of the control mechanism. However, in the only patient in whom growth hormone (GH) production was studied, it was normal.³

GH facilitates the formation in the liver of the

polyamines—putrescine, spermidine, and spermine—by controlling the enzyme ornithine decarboxylase (EC 4.1.1.17). Cortisol also stimulates hepatic ornithine decarboxylase activity, both by itself and synergistically with GH.⁴ If the abnormal intrauterine growth of children with Beckwith's syndrome were due to a fault in the GH-like regulating mechanism, changes might be expected in the excretion of polyamines. Although an increased excretion of total polyamines was reported in one patient with acromegaly in whom the enlargement of distal parts of the body was associated with prolonged excessive secretion of GH, no further details were given.⁵ It is also stated that 'There are no known growth processes that occur without the prior stimulation of polyamine biosynthesis'.⁶ The urinary excretion of these polyamines was measured in a group of 7 children with this syndrome.

Table 1 *Clinical features of children with Beckwith's syndrome*

Case	Age (weeks)	Weight (kg)	
1	5	4.5	Umbilical hernia, large protruding tongue, sloping forehead with prominent occiput. Blood glucose 2.4 mmol/l
2	1	3.5	Exomphalos, large protruding tongue, abnormal ear lobes, hypoglycaemic episodes
3	7	3.45	Exomphalos, enlarged tongue, perforation of colon, malrotation of gastrointestinal tract. Blood glucose 1.7 mmol/l
4	1 day	4.7	Exomphalos, enlarged protruding tongue, large slanting forehead, creases on ear lobes, large hands and feet
5	1 day	3.3	Exomphalos, glandular hypospadias, enlarged left kidney
6	13	4.95	Small exomphalos, protruberant tongue. Blood glucose 1.0 mmol/l
7	1	3.1	Exomphalos, enlarged protruding tongue. Blood glucose, 1.8 mmol/l

Conversion: SI to traditional units—glucose: 1 mmol/l \approx 18 mg/100 ml

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Methods

24-hour urine collections were made from a group of 10 children aged 2 months to 2 years who were not suffering infection or any growth disorder. These urine samples were used as controls. Similar collections were made from 7 children with Beckwith's syndrome (Table 1). Urine samples, without preservative, were stored deep-frozen (-20°C) until required for analysis. Tissue samples were obtained at necropsy (controls), or at operation (Beckwith's syndrome), and these were also stored at -20°C until required.

Aliquots of urine were hydrolysed in 6 mol/l HCl at 100°C overnight. The free polyamines were measured using a modular automatic amino-acid analyser (ChemLab Instruments) with columns and buffers similar to those described by Marton *et al.*⁷

Results

When standards were added to urine samples and taken through the whole procedure, putrescine, spermidine, and spermine were recovered in yields of $88 \pm 6.5\%$, $79 \pm 7.2\%$, and $83 \pm 4.1\%$ respectively (mean \pm coefficient of variation) (8 estimations). In the controls the 24-hourly urinary excretion of putrescine was $0.85 \pm 0.71 \mu\text{mol/kg}$ (mean \pm SD) (range 0.26–2.2) and spermidine $0.45 \pm 0.36 \mu\text{mol/kg}$ (range 0.02–1.15). Spermine was below the detection level ($\approx 0.05 \mu\text{mol/kg}$ per 24 hours) in all but one child. When the age range was extended from 2 months to 8 years there were no significant differences for the excretion of putrescine and spermidine, but the amounts of both were found to be related to body weight ($r=0.72$ with $P<0.01$ and $r=0.65$ with $P<0.05$). These control values appear to be similar to those reported for adults.⁷

The urinary excretion of spermidine in children with Beckwith's syndrome was either very low or it was undetectable (detection limit $\approx 0.02 \mu\text{mol/kg}$ per 24 hours), the difference being significant ($P<0.05$) (Table 2). The excretion of putrescine was significantly increased ($P<0.05$), while spermine could not be detected in any of the patients. The putrescine/spermidine ratio was very high in all but one patient.

In samples of tongue muscle removed at operation, the polyamine levels showed a reduced putrescine/

spermidine ratio compared with control specimens obtained at necropsy (Table 3).

Discussion

The stimulation of ornithine decarboxylase activity, on which polyamine formation depends, is controlled by GH in the kidney and liver, and by prolactin in the spleen. However, little appears to be known of the hormonal control in other tissues.⁸

The concentrations of polyamines in tissue appear to determine the rate of growth as in catch-up growth after a period of restricted food intake,⁹ and in liver regeneration after partial hepatectomy.¹⁰ The polyamines are believed to alter the rate of nucleic acid activity. Enhanced rates of ribonucleic acid and protein synthesis were observed after the rise in polyamine levels.¹¹ The growth of tissue appears to be affected not only by the absolute amounts of polyamine present but also by the ratios of the different components. Thus in various tumours changes take place in the spermidine and putrescine ratios.⁷

It is possible therefore, that the growth disorders in Beckwith's syndrome are associated with the high putrescine excretion and low spermidine and putrescine ratios in the urine. The raised output of putrescine and the reduced output of spermidine may have been due to a reduced conversion of putrescine to spermidine in the patients, or to enhanced metabolism of the spermidine. While in infancy children with Beckwith's syndrome grow faster than normal, they tend to slow down to normal rates later. During this stage it is desirable to recheck the urinary polyamine levels which may conceivably be close to the normal range. If these results can be shown to be typical in this syndrome, then they raise the possibility of a better understanding of the mechanisms of a wider series of growth disorders.

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Table 2 24-hour urinary polyamine excretion ($\mu\text{mol/kg}$) in children with Beckwith's syndrome

Case	Putrescine	Spermidine	Spermine
1	1.5*	0.1	0
	0.7*	0.1	0
	1.7*	0	0
2	1.8	0	0
3	1.0	0	0
4	0	0	0
5	2.0	0	0
6	3.4	0	0
7	0.6	0	0

*Analyses on 3 successive days.

Control levels: putrescine $0.85 \pm 0.71 \mu\text{mol/kg}$ per 24 hours (mean \pm SD), spermidine $0.45 \pm 0.36 \mu\text{mol/kg}$ per 24 hours (mean \pm SD), putrescine/spermidine ratio 3.0 ± 3.5 (mean \pm SD).

Table 3 Tissue levels of polyamines (mmol/kg wet tissue) in children with Beckwith's syndrome

Group	Tongue	Age (months)	Putrescine	Spermidine	Spermine
Control (n = 3)	Necropsy specimen	3½	190	60	160
	Necropsy specimen	4 hours	460	100	200
	Necropsy specimen	11 years	320	100	190
Patients with Beckwith's syndrome (n = 2)	At operation	14	260	120	210
	At operation	11	340	180	230

Putrescine/spermidine ratio in controls 3.7, putrescine/spermidine ratio in children with Beckwith's syndrome 2.0.

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