

Blood pressure centiles for Great Britain

Lisa V Jackson, Nandu K S Thalange, Tim J Cole

Arch Dis Child 2007;**92**:298–303. doi: 10.1136/adc.2005.081216

See end of article for authors' affiliations

Correspondence to:
Dr L V Jackson, Willow
Wood Surgery, Aslake
Close, Sprowston, Norwich
NR7 8TT, UK; lisa.jackson@
nhs.net

Accepted 10 July 2006
Published Online First
11 August 2006

Objective: To produce representative cross-sectional blood pressure reference centiles for children and young people living in Great Britain.

Design: Analysis of blood pressure data from seven nationally representative surveys: Health Surveys for England 1995–8, Scottish Health Surveys 1995 and 1998, and National Diet & Nutrition Survey 1997.

Methods: Blood pressure was measured using the Dinamap 8100 with the same protocol throughout. Weight and height were also measured. Data for 11 364 males and 11 537 females aged 4–23 years were included in the analysis, after excluding 0.3% missing or outlying data. Centiles were derived for systolic, diastolic, mean arterial and pulse pressure using the latent moderated structural (LMS) equations method.

Results: Blood pressure in the two sexes was similar in childhood, rising progressively with age and more rapidly during puberty. Systolic pressure rose faster and was appreciably higher in adult men than in adult women. After adjustment for age, blood pressure was related more to weight than height, the effect being stronger for systolic blood pressure. Pulse pressure peaked at 18 years in males and 16 years in females.

Conclusions: These centiles increase our knowledge of blood pressure norms in contemporary British children and young people. High blood pressure for age should be defined as blood pressure above the 98th centile, and high-normal blood pressure for age as blood pressure between the 91st and 98th centiles. The centiles identify children and young people with increased blood pressure, and will be of benefit to both clinical practice and research.

There is no satisfactory definition of hypertension in children.¹ As a result, blood pressure is often not measured in paediatric clinical practice, and understanding the clinical significance of blood pressure readings in children is hampered by the lack of satisfactory reference data with which to interpret them.

Reference blood pressure centiles should therefore improve the understanding of blood pressure variation in childhood. In Britain and worldwide, there have been many studies of childhood blood pressure, but all are of limited use in Great Britain owing to the use of non-representative populations, limited age ranges and mixed methodologies for blood pressure measurement. Accordingly, we have developed representative cross-sectional blood pressure references for children and young people living in Great Britain.

METHODS

Blood pressure data from seven national health and social surveys carried out between 1995 and 1998 were obtained from the UK Data Archive (<http://www.data-archive.ac.uk/>) (table 1). The data were originally collected on behalf of the Departments of Health and the Ministry of Agriculture Fisheries and Food, by the Joint Health Surveys Unit of Social and Community Planning Research and University College London, London, UK and the Social Survey Division of the Office for National Statistics and Medical Research Council, Human Nutrition Research, Cambridge, UK.

The survey samples were obtained by stratified multistage sampling techniques to ensure that there was a proportional representation of the population at large by sex, age, geographical region and social class.² In brief, the demographic characteristics of a geographical area are known from census and other data. Using this information, a representative sample of individuals from the target age groups for each survey was obtained. Households in geographical areas selected by post-code were contacted and asked to fill in a questionnaire to identify eligible young people. A subset of this initial sample

was then contacted by trained interviewers. The demographic characteristics of those agreeing to take part were determined and further targeted sampling undertaken to ensure the study sample remained representative. More information may be found in the published surveys.

Ethical approval was obtained from all areas in which the surveys were carried out. Participation was subject to informed consent. Data for the present analysis were excluded for participants who had eaten, consumed alcohol or smoked in the 30 min before being measured, and for those on anti-hypertensive drugs.

All seven surveys used the Dinamap 8100 (Critikon, Tampa, Florida, USA) with the same protocol to measure blood pressure. The use of an automatic oscillometric method was necessary for practicality, accuracy and reproducibility.^{3,4} Briefly, the blood pressure cuff was applied to the right arm. The lower margin of the cuff was placed about 2 cm above the elbow crease, with the arrow marked on the cuff placed over the brachial artery. The cuff was wrapped to a tightness allowing two fingers to be inserted under the top and bottom of the cuff. Four cuff sizes were available, the appropriate cuff size being determined by measurement of the mid-upper arm circumference (child cuff 10–19 cm, small adult cuff 17–25 cm, adult cuff 23–33 cm, large adult cuff 31–40 cm). The participants were comfortably seated, with their feet flat to the floor. Measurements of systolic, mean arterial and diastolic pressure were obtained after a 10–15 min rest period in triplicate, at minute intervals. The first reading was discarded and the mean of the second and third readings was used for analysis, as the first reading of a series of blood pressure measurements is typically higher with oscillometric devices.^{4,5} Pulse pressure was calculated by subtracting diastolic from systolic pressure.

For 73 (0.3%) participants, the blood pressure data were found to be either outliers or inconsistent with age, lying more than five SD from the median for age and sex. Hence blood

Abbreviation: SDS, standard deviation score

Table 1 Demographic characteristics of 22 974 participants aged 4–23.9 years from seven national health and social surveys

Survey	Year	England	Scotland	Wales	Age range (years)	Sample size
Health Survey for England	1995	✓			5–23	3485
Health Survey for England	1996	✓			5–23	4198
Health Survey for England	1997	✓			5–23	5520
Health Survey for England	1998	✓			4–23	3756
Scottish Health Survey	1995		✓		16–23	707
Scottish Health Survey	1998		✓		5–23	3043
National Diet & Nutrition Survey	1997	✓	✓	✓	4–19	1905
Overall						22 974

pressure data for 22 901 participants, 11 364 male and 11 537 female, aged 4–23.9 years were analysed.

Sex-specific smoothed centiles were derived using the latent moderated structural equations (LMS) method⁶ for age and sex. The LMS method summarises the age-changing frequency distribution of blood pressure in terms of three curves: the L curve defines the skewness, the M curve the median and the S curve the coefficient of variation as functions of age. Centile charts were drawn with centiles spaced two-thirds of an SD score (SDS) apart, ranging from the 0.4th centile (−2.67 SDS) through to the 99.6th centile (+2.67 SDS), consistent with other anthropometric charts in current use in the UK.⁷

The relationship of systolic and diastolic blood pressure, weight and height was investigated through the multiple regression of blood pressure on weight and height, after adjusting the three variables for age and sex by converting them to SDS. The British 1990 reference⁸ was used for height and weight, and the internal reference for blood pressure. For measuring weight and height in subjects age ≥23 years was taken as 22.99 (the upper limit of the British reference). Sex effects were tested for in the regression by including sex and its interactions with height and weight.

RESULTS

Table 2 summarises the data for 22 901 participants with both systolic and diastolic blood pressure. Mean arterial pressure, height and weight were missing for 8%, 1% and 2% of participants, respectively. By year of age the sample consisted of 114 participants aged 4 years, 1181–1581 per year between 5 and 16 years, and 715–950 per year between 17 and 23 years. Height was very similar to the British 1990 reference (mean SDS 0.0), while weight and body mass index (weight (kg)/height² (m²)) were slightly increased (mean SDS 0.3–0.4).

The data were used to construct blood pressure centile charts for systolic, diastolic, mean arterial and pulse pressure (figs 1–4). Blood pressure in the two sexes was similar before puberty,

but the pubertal rise was more marked in boys. Pulse pressure peaked at 18 years in male participants and at 16 years in female participants, corresponding to the end of puberty.

Table 3 summarises the multiple regression of blood pressure on weight and height, each adjusted for age and sex by

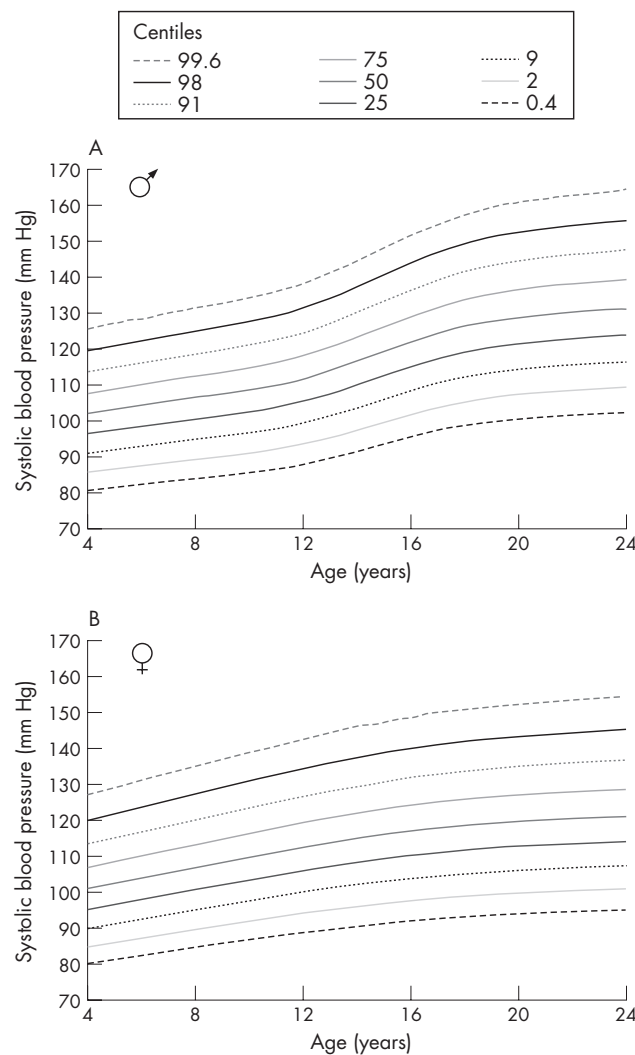


Figure 1 Systolic blood pressure centiles in male (A) and female participants (B). The centiles are spaced two-thirds of a standard deviation score apart. Systolic pressure rises progressively with age, but rises more steeply in puberty, particularly in boys.

Table 2 Summary statistics for 22 901 participants with valid data

Variable	n	Mean	SD
Men (%)	22 901	49.6	—
Age (years)	22 901	13.1	5.2
Height (cm)	22 676	148.9	21.3
Weight (kg)	22 485	46.4	19.9
Height SDS (British 1990)	22 676	−0.03	1.08
Weight SDS (British 1990)	22 485	0.27	1.14
Body mass index SDS (British 1990)	22 425	0.36	1.11

SDS, standard deviation score.

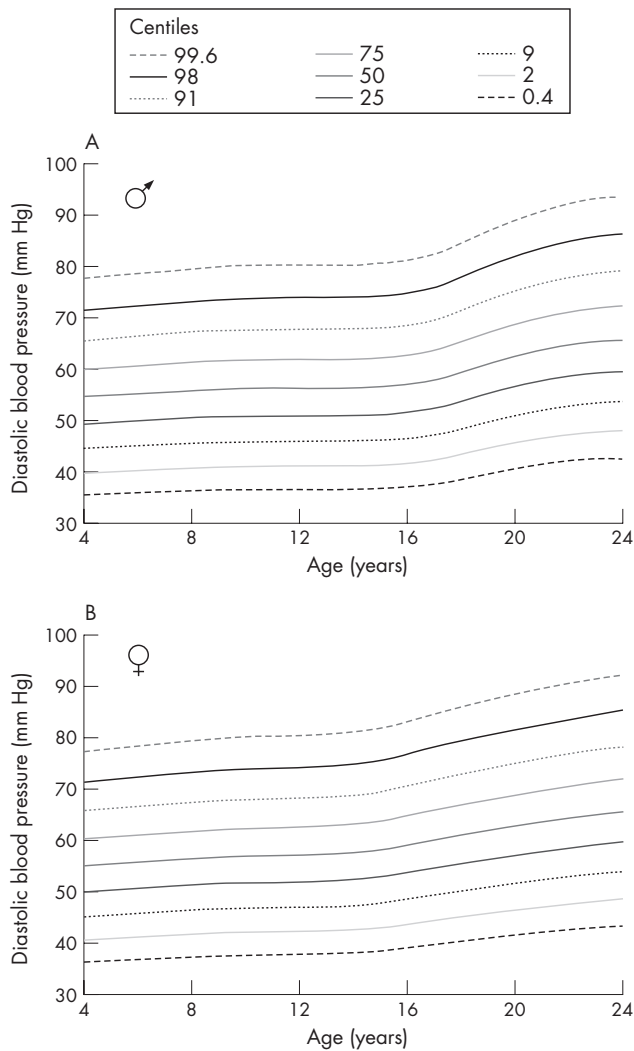


Figure 2 Diastolic blood pressure centiles in male (A) and female participants (B). The centiles are spaced two-thirds of a standard deviation score apart. Diastolic pressure rises slowly in childhood, but as with systolic pressure, rises more steeply in puberty.

converting to SDS. This adjustment allowed the data for both sexes and all ages to be combined. Results are also given by sex, although they do not differ significantly; hence the combined results are valid. Weight had a large and positive effect on blood pressure ($p < 0.001$), whereas height had a smaller negative effect ($0.005 < p < 0.001$). A 1 SD increase in weight was associated with a 0.3 SD increase in systolic pressure and a 0.08 SD increase in diastolic pressure, whereas a 1 SD increase in height was associated with a 0.03 SD reduction in both systolic and diastolic pressure. Thus, on average, for any given weight, a taller (and hence thinner) individual had lower blood pressure. Analysing the data in separate age groups showed the associations in late puberty to be stronger than before or after.

These results suggest that body size (ie, weight) and obesity (weight adjusted for height) both play a role in raising blood pressure, particularly systolic blood pressure, 8% of the variation of which was explained by weight and height. The effect on diastolic blood pressure (0.5% of variance explained) was much smaller.

Using the British Hypertension Society cut-offs for hypertension,⁹ 23% of men and 6% of women exceeded the systolic

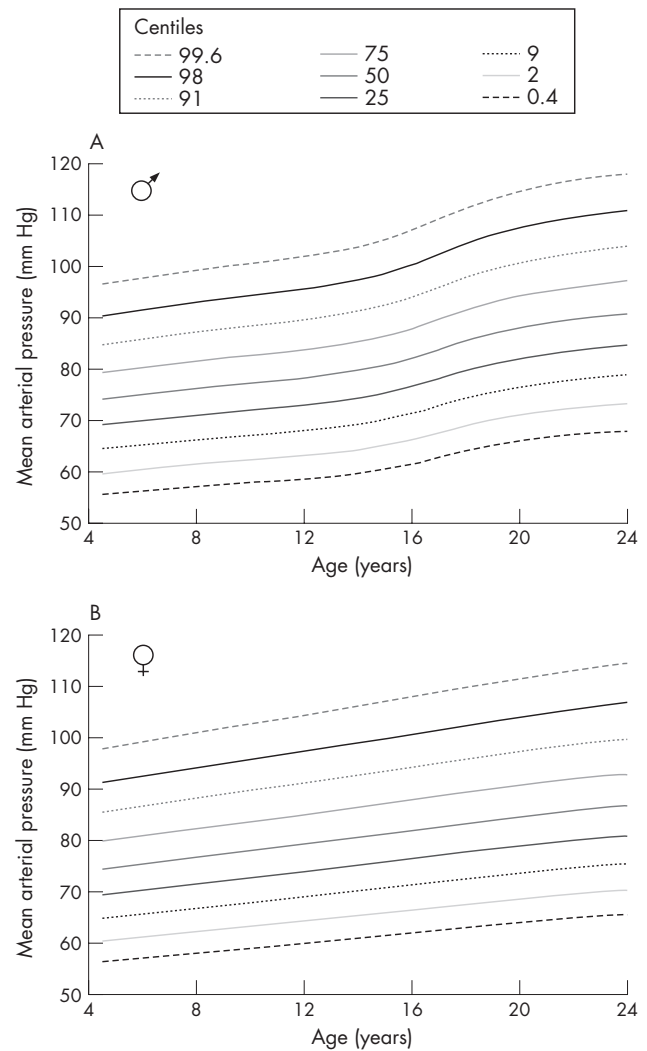


Figure 3 Mean arterial pressure centiles in male (A) and female participants (B). The centiles are spaced two-thirds of a standard deviation score apart. Mean arterial pressure rises progressively with age.

cut-off, and 1.0% of men and 0.8% of women exceeded the diastolic cut-off by age 24 years.

DISCUSSION

The blood pressure centiles presented here are based on data collected using a consistent and rigorous method in representative samples of nearly 23 000 children and young people living in Great Britain. As such, we believe they are the most accurate characterisation of normal blood pressure in any country to date.

It is well recognised that children's blood pressure tends to "track" over time.¹⁰⁻¹⁴ Moreover, high blood pressure in children is associated with the development of atherosclerosis,¹⁵⁻¹⁹ especially in those with additional risk factors, notably obesity.¹⁶⁻²⁰ The charts will aid the timely recognition and monitoring of individuals with high blood pressure and hypertension, and facilitate the detection of children with secondary hypertension, consequent on renal, endocrine or other disease.¹ Blood pressure monitoring is also important in children at risk of hypertension and/or vascular disease, such as those with obesity, diabetes, renal disease, or those receiving

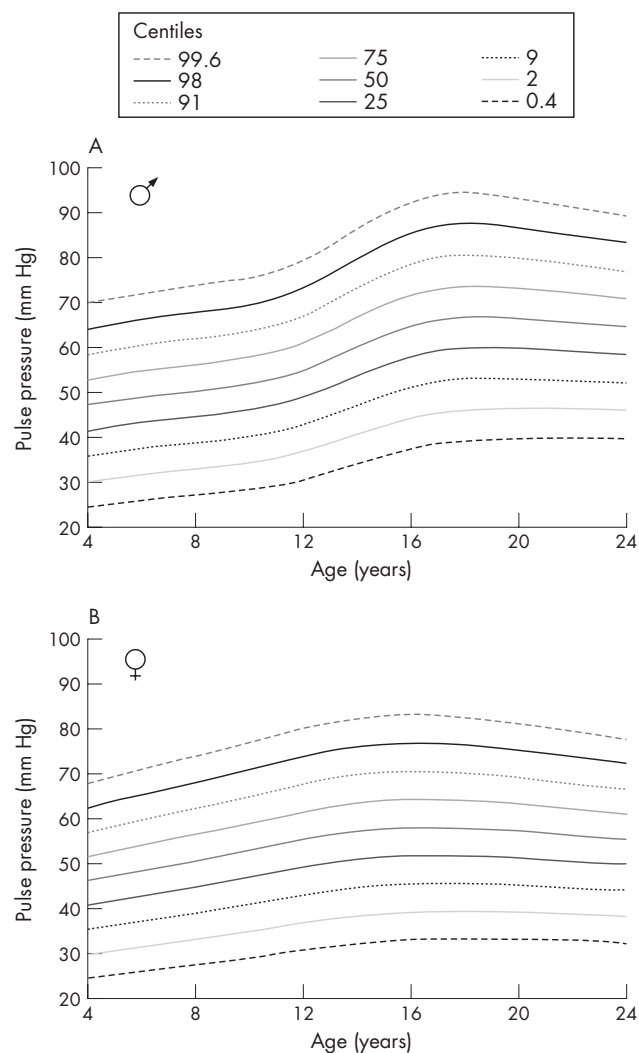


Figure 4 Pulse pressure centiles in male (A) and female participants (B). The centiles are spaced two-thirds of a standard deviation score apart. Pulse pressure rises progressively until the end of puberty and then falls again.

steroids or stimulant drugs and where a family history of hypertension is present.

We have used a standard nine-centile format consistent with other charts in use in the UK.⁷ The charts show a progressive rise in systolic and diastolic pressure with increasing age, the rise being more marked in males during puberty. This is consistent with an effect of body size (indicated by weight) and obesity (weight adjusted for height) on blood pressure, the

effect being stronger for systolic blood pressure. Thus males, who gain more weight at puberty than females, have significantly higher blood pressures, with almost a quarter satisfying the British Hypertension Society definition of hypertension,⁹ defined as systolic pressure >140 mm Hg and/or diastolic pressure >90 mm Hg, by the age of 24 years. The high systolic pressures in older teenagers and young adults, particularly men, are of special concern. However, blood pressure measurements on a single occasion are insufficient for the determination of high or high-normal blood pressure for age,^{21, 22} in the absence of evidence of a pathological cause or end organ damage, especially in children, who are more prone to “white coat” hypertension.²³ Repeated measurements typically show that the majority of children with increased blood pressure on a single occasion subsequently have normal blood pressure.²⁴

Pulse pressure^{25, 26} and mean arterial pressure²⁷ have been found to be significant determinants of morbidity and mortality in adults. The significance of these measures in children is unknown, but pulse pressure may be an indicator of early arterial disease, as has been found in young adults with type 1 diabetes.²⁷ Of note, we found that pulse pressure peaks at the end of puberty in both sexes, before falling in young adult life (fig 4) in contrast with systolic, diastolic and mean arterial pressures, which rise progressively with age (figs 1–3). A knowledge of normal ranges for pulse pressure and mean arterial pressure should aid research in this area.

The use of oscillometric blood pressure measurements was dictated by the nature of the health and social surveys, which required a reliable, reproducible and accurate method for determining blood pressure, using multiple observers.^{3, 4} The Dinamap 8100 was subject to a rigorous calibration study³ to ensure its validity (although the calibration study did not include participants aged <16 years). However, the Dinamap monitor has been compared with direct radial artery pressure and central aortic pressure measurements in infants and children and was found to be superior to the auscultatory method.^{28, 29} Moreover, particularly in young children, the conventional mercury sphygmomanometer can be difficult to use,^{4, 20} with the Korotkoff sounds hard to distinguish, so, increasingly, automated oscillometric devices are being used in clinical practice.³⁰

O’Brien *et al.*,³¹ using the British Hypertension Society protocol, graded the Dinamap 8100 B for systolic blood pressure and D for diastolic blood pressure compared with the conventional mercury sphygmomanometer in adults.³² Paediatric studies have generally found significant differences, particularly between diastolic pressure assessed by fourth-phase Korotkoff sounds.^{33–35} However, an Australian study of prepubertal children with type 1 diabetes using the British Hypertension Society protocol graded the Dinamap B for both systolic and diastolic pressure.³⁶

The Dinamap 8100 and other oscillometric devices produce results that differ in comparison with the mercury sphygmomanometer. These differences have been attributed to

Table 3 Relationship of systolic and diastolic blood pressure with weight and height by sex (all variables expressed as standard deviation score)

Outcome measure	Sex	n	Weight (SDS)	Height (SDS)	R ² (%)
Systolic blood pressure (SDS)	Men	11 153	0.279 (0.011)	−0.033 (0.011)	8.6
	Women	11 272	0.258 (0.010)	−0.039 (0.010)	7.6
	Combined	22 425	0.268 (0.007)	−0.035 (0.008)	8.1
Diastolic blood pressure (SDS)	Men	11 153	0.070 (0.011)	−0.033 (0.012)	0.4
	Women	11 272	0.081 (0.010)	−0.032 (0.011)	0.6
	Combined	22 425	0.076 (0.008)	−0.033 (0.008)	0.5

SDS, standard deviation score. The table gives regression coefficients (SEs).

What is already known on this topic

- Blood pressure rises through childhood and childhood blood pressure strongly predicts adult blood pressure.
- This rise in blood pressure is substantially determined by weight.
- As with growth, blood pressure is an important parameter of child health.
- Furthermore, atherosclerosis and hypertension may have their origins in childhood, particularly in those with additional risk factors—for example, obesity, renal disease or diabetes.

What this study adds

- These blood pressure centiles compiled from nationally representative data are the most comprehensive attempt to characterise normal blood pressure in childhood in Great Britain.
- The centiles complement existing charts for height, weight and body mass index and other parameters in evaluating the health of children.
- This information will contribute to a better understanding of blood pressure in childhood and aid further research.

inaccuracies³⁷ but simply reflect the fact that different methods yield different results.^{1 38} However, in view of these differences, blood pressure results recorded with the mercury sphygmomanometer should be referenced to these centiles with caution.

The definition of hypertension in children is problematic. Use of the British Hypertension Society cut-offs in adults is justified by adverse health outcomes in association with hypertension.⁹ However, no single cut-off can define hypertension in children owing to the normal rise in blood pressure with age, and the paucity of evidence about what constitutes hypertension in children.¹ Consequently, we suggest that, in children, those above the 98th centile on repeated occasions are stated to have high blood pressure for age, whereas those lying between the 91st and 98th centiles are stated to have high-normal blood pressure for age. These cut-offs are similar to recommendations made in the Taskforce Report on High Blood Pressure in Children and Adolescents in the USA.³⁹ Our centile-based definitions predict a prevalence of 2.3% for high blood pressure (>2 SDS) and 6.9% for high-normal blood pressure (>1.33 SDS). These centiles should facilitate ongoing research into the importance of high or high-normal blood pressure in children, and serve as a basis for defining hypertension in childhood.

The strong association between high blood pressure and weight/obesity that we and others have found^{40 41} is of particular concern given the well documented rise in childhood obesity.⁴² Childhood obesity, and its health consequences—including hypertension, metabolic syndrome and type 2 diabetes—present a major challenge for the coming years and demand vigilance and concerted action from all healthcare professionals to mitigate the adverse health consequences for children and young people.

ACKNOWLEDGEMENTS

We thank Dr Graham Derrick, Consultant Paediatric Cardiologist, Great Ormond Street Hospital for Children NHS Trust, and Dr Carlo Acerini, University Lecturer in Paediatrics, Cambridge University, for their helpful comments.

Authors' affiliations

Lisa V Jackson, Willow Wood Medical Practice and University of East Anglia School of Medicine, Health Policy and Practice, Norwich, UK
Nandu K S Thalange, Norfolk and Norwich University Hospital, Norwich, UK

Tim J Cole, Centre for Paediatric Epidemiology and Biostatistics, Institute of Child Health, University College London, London, UK

Funding: TJC is funded by the Medical Research Council.

Research at the UCL Institute of Child Health and Great Ormond Street Hospital for Children NHS Trust benefits from R&D funding received from the NHS Executive.

Competing interests: Subsequent to the presentation of data at the Spring Meeting in April 2003, blood pressure centile charts using these data were published commercially.

Previous publication: Some of this work was presented as an oral presentation at the Royal College of Paediatrics & Child Health Spring Meeting, April 2003: G203. Blood pressure centiles for children and young people aged 4–24 years in Great Britain. *Arch Dis Child* 2003; **88**:A66.

Contributors: all authors contributed to the design, analysis and writing up of the paper. LVJ is guarantor.

REFERENCES

- 1 **Goonasekera CDA**, Dillon MJ. Measurement and interpretation of blood pressure. *Arch Dis Child* 2000;**82**:261–5.
- 2 **Elliot D**. Optimising sample design for surveys of health and related behaviour and attitudes. *Survey Methodol Bull* 1995;**36**:8–17.
- 3 **Bolling K**. *The Dinamap 8100 calibration study*. London: HMSO, 1994.
- 4 **Gillman MW**, Cook NR. Blood pressure measurement in childhood epidemiological studies. *Circulation* 1995;**92**:1049–57.
- 5 **Park MK**, Menard SM. Normative oscillometric blood pressure values in the first 5 years in an office setting. *Am J Dis Child* 1989;**143**:860–4.
- 6 **Cole TJ**, Green PJ. Smoothing reference centile curves: the LMS method and penalized likelihood. *Stat Med* 1992;**11**:1305–19.
- 7 **Cole TJ**. Do growth chart centiles need a face lift? *BMJ* 1994;**308**:641–2.
- 8 **Freeman JV**, Cole TJ, Chinn S, *et al*. Cross sectional stature and weight reference curves for the UK, 1990. *Arch Dis Child* 1995;**73**:17–24.
- 9 **Williams B**, Poulter NR, Brown MJ, *et al*. British Hypertension Society guidelines for hypertension management 2004 (BHS-IV): summary. *BMJ* 2004;**328**:634–40.
- 10 **Bao W**, Threefoot SA, Srinivasan SR, *et al*. Essential hypertension predicted by tracking of elevated blood pressure from childhood to adulthood: the Bogalusa Heart Study. *Am J Hypertens* 1995;**8**:657–65.
- 11 **Klumbiene J**, Sileikiene L, Milasauskiene Z, *et al*. The relationship of childhood to adult blood pressure: longitudinal study of juvenile hypertension in Lithuania. *J Hypertens* 2000;**18**:531–8.
- 12 **O'Sullivan JJ**, Derrick G, Foxall RJ. Tracking of 24-hour and casual blood pressure: a 1-year follow-up study in adolescents. *J Hypertens* 2000;**18**:1193–6.
- 13 **Cook NR**, Gillman MW, Rosner BA, *et al*. Prediction of young adult blood pressure from childhood blood pressure, height, and weight. *J Clin Epidemiol* 1997;**50**:571–9.
- 14 **Lambrechtsen J**, Rasmussen F, Hansen HS, *et al*. Tracking and factors predicting rising in 'tracking quartile' in blood pressure from childhood to adulthood: Odense Schoolchild Study. *J Hum Hypertens* 1999;**13**:385–91.
- 15 **Newman WP III**, Freedman DS, Voors AW, *et al*. Relation of serum lipoprotein levels and systolic blood pressure to early atherosclerosis: the Bogalusa Heart Study. *N Engl J Med* 1986;**314**:138–144.
- 16 **Berenson GS**, Srinivasan SR, Bao W, *et al*. Association between multiple cardiovascular risk factors and atherosclerosis in children and young adults. The Bogalusa Heart Study. *N Engl J Med* 1998;**338**:1650–6.
- 17 **Mahoney LT**, Burns TL, Stanford W, *et al*. Coronary risk factors measured in childhood and young adult life are associated with coronary artery calcification in young adults: the Muscatine Study. *J Am Coll Cardiol* 1996;**27**:277–84.
- 18 **Li S**, Chen W, Srinivasan SR, *et al*. Childhood cardiovascular risk factors and carotid vascular changes in adulthood: the Bogalusa Heart Study. *JAMA* 2003;**290**:2271–6.
- 19 **Raitakari OT**, Juonala M, Kahonen M, *et al*. Cardiovascular risk factors in childhood and carotid artery intima-media thickness in adulthood: the Cardiovascular Risk in Young Finns Study. *JAMA* 2003;**290**:2277–83.
- 20 **Reinehr T**, Andler W. Changes in the atherogenic risk factor profile according to degree of weight loss. *Arch Dis Child* 2004;**89**:419–22.
- 21 **de Swiet M**. The epidemiology of hypertension in children. *Br Med Bull* 1986;**42**:172–5.
- 22 **Adroque HE**, Sinaiko AR. Prevalence of hypertension in junior high school-aged children: effect of new recommendations in the 1996 Updated Task Force Report. *Am J Hypertens* 2001;**14**:412–14.
- 23 **Hornsby JL**, Mongan PF, Taylor AT, *et al*. 'White coat' hypertension in children. *J Fam Pract* 1991;**33**:617–23.
- 24 **Madhavan S**, Ooi WL, Cohen H, *et al*. Relation of pulse pressure and blood pressure reduction to the incidence of myocardial infarction. *Hypertension* 1994;**23**:395–401.

- 25 **Benetos A**, Rudnichi A, Safar M, *et al*. Pulse pressure and cardiovascular mortality in normotensive and hypertensive subjects. *Hypertension* 1998;**32**:560–4.
- 26 **van Trijp MJ**, Grobbee DE, Peeters PH, *et al*. Average blood pressure and cardiovascular disease-related mortality in middle-aged women. *Am J Hypertens* 2005;**18**:197–201.
- 27 **Schram MT**, Chaturvedi N, Fuller JH, *et al*. Pulse pressure is associated with age and cardiovascular disease in type 1 diabetes: the Eurodiab Prospective Complications Study. *J Hypertens* 2003;**21**:2035–44.
- 28 **Park MK**, Menard SM. Accuracy of blood pressure measurement by the Dinamap monitor in infants and children. *Pediatrics* 1987;**79**:907–14.
- 29 **Colan SD**, Fujii A, Borow KM, *et al*. Noninvasive determination of systolic, diastolic and end-systolic blood pressure in neonates, infants and young children: comparison with central aortic pressure measurements. *Am J Cardiol* 1983;**52**:867–70.
- 30 **O'Brien E**. Demise of the mercury sphygmomanometer and the dawning of a new era in blood pressure measurement. *Blood Press Monit* 2003;**8**:19–21.
- 31 **O'Brien E**, Petrie J, Littler W, *et al*. The British Hypertension Society protocol for the evaluation of automated and semi-automated blood pressure measuring devices with special reference to ambulatory systems. *J Hypertens* 1990;**8**:607–19.
- 32 **O'Brien E**, Mee F, Atkins N, *et al*. Short report: accuracy of the Dinamap portable monitor, model 8100 determined by the British Hypertension Society protocol. *J Hypertens* 1993;**11**:761–3.
- 33 **Wattigney WA**, Webber LS, Lawrence MD, *et al*. Utility of an automatic instrument for blood pressure measurement in children. The Bogalusa Heart Study. *Am J Hypertens* 1996;**9**:256–62.
- 34 **Barker ME**, Shiell AW, Law CM. Evaluation of the Dinamap 8100 and Omron M1 blood pressure monitors for use in children. *Paediatr Perinat Epidemiol* 2000;**14**:179–86.
- 35 **Park MK**, Menard SW, Yuan C. Comparison of auscultatory and oscillometric blood pressures. *Arch Pediatr Adolesc Med* 2001;**155**:50–3.
- 36 **Jin RZ**, Donaghue KC, Fairchild J, *et al*. Comparison of Dinamap 8100 with sphygmomanometer blood pressure measurement in a prepubertal diabetes cohort. *J Paediatr Child Health* 2001;**37**:545–9.
- 37 **O'Brien E**, Atkins N. Inaccuracy of the Dinamap 8100 portable monitor. *Lancet* 1997;**349**:1026.
- 38 **Friedman B**. Accuracy of Dinamap monitors. *Lancet* 1997;**350**:217–18.
- 39 **National Blood Pressure Education Working Group on High Blood Pressure in Children and Adolescents**. Fourth report on the diagnosis, evaluation and treatment of high blood pressure in children and adolescents: a working group report from the National High Blood Pressure Education Program. *Pediatrics* 2004;**114**:555–76.
- 40 **St George IM**, Williams SM, Silva PA. The stability of high blood pressure in Dunedin children: an eight year longitudinal study. *NZ Med J* 1990;**103**:115–17.
- 41 **de Swiet M**, Fayers P, Shinebourne EA. Blood pressure in first 10 years of life: the Brompton study. *BMJ* 1992;**304**:23–6.
- 42 **Chinn S**, Rona RJ. Prevalence and trends in overweight and obesity in three cross sectional studies of British children 1974–94. *BMJ* 2001;**322**:24–6.

IMAGES IN PAEDIATRICS

doi: 10.1136/adc.2006.112771

An interesting facial rash

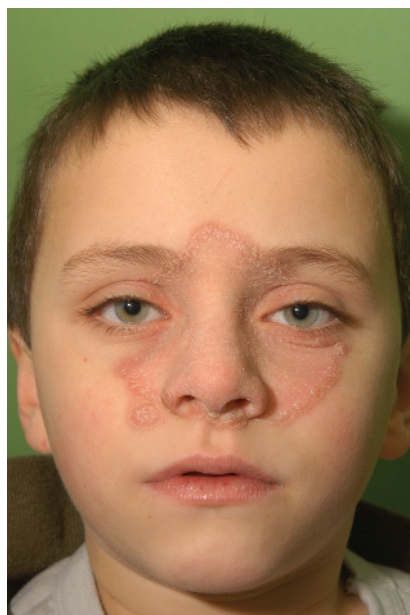


Figure 1 Informed consent was obtained for publication of this figure.

A 7-year-old boy presented to paediatrics with a 3-week history of a worsening facial rash, which was intermittently itchy. He was otherwise well. He had two guinea pigs as pets. On examination, an extensive scaling erythema was noticed with a definite edge involving the upper eyelids, the bridge of the nose and extending onto both cheeks (see fig 1).

A provisional diagnosis of tinea faciei was made; however, cutaneous lupus was also considered. While mycology results were awaited, topical terbinafine was given, with little effect. Microscopy revealed a dermatophyte infection with *Trichophyton mentagrophytes*, and a 3-week course of oral terbinafine (125 mg oral dosage once daily) was given. The rash resolved completely, leaving post-inflammatory hyperpigmentation only.

Tinea facialis/faciei is a dermatophytosis of the glabrous facial skin, characterised by a well-circumscribed, often asymmetric, erythematous patch with an elevated border and central regression. It may be asymptomatic or present with pruritus, or, occasionally, photosensitivity that may lead to diagnostic confusion with cutaneous lupus.¹ It is the most commonly misdiagnosed dermatophytosis. Other differential diagnoses include eczema, seborrhoeic dermatitis² and rosacea.

It is most common in children, with predisposing factors including exposure to animals, chronic topical steroid use and spread from tinea capitis. The most frequent organisms involved are *T mentagrophytes*, *T rubrum* and *T tonsurans*. However, cases caused by *Microsporum audouinii* and *M canis* occur worldwide. Most cases are given short-term oral antifungal treatment, but milder cases may respond to topical imidazoles. Affected animals and family members should also be treated.

Caroline A Love

Department of Dermatology, Rowan House, Whiston Hospital, Merseyside, UK

John A Sills, Judith M Ellison

St Helens and Knowsley NHS Trust, Merseyside, UK

Correspondence to: Dr C A Love, Department of Dermatology, Rowan House, Whiston Hospital, Warrington Road, Prescot, Merseyside L35 5DR, UK; drcalove@hotmail.com

Competing interests: None declared.

REFERENCES

- 1 **Cirillo-Hyland V**, Humphreys T, Elenitsas R. Tinea faciei. *J Am Acad Dermatol* 1993;**29**:119–20.
- 2 **Gorani A**, Oriani A, Cambiaghi S. Seborrhoeic dermatitis-like tinea faciei. *Pediatr Dermatol* 2005;**22**:243–4.

to date. An infant had presented with increased work of breathing from birth, and tracheomalacia had been found on flexible bronchoscopy. The section on tracheomalacia was again succinct and pragmatic. The chapter on bronchoscopy is by the undisputed king of this investigation, and there was a nice section on interpretation of bronchoalveolar lavage. We had a teenager with a pleural effusion from likely auto-immune disease; there was a solid 23 page chapter on pulmonary involvement in rheumatic disorders. I was therefore very quickly sold on this book – I had confidence that it would deliver what I needed to know and point to further reading.

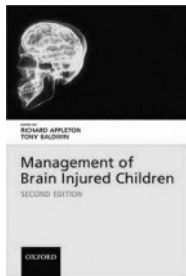
Sometimes a large specialist textbook like this is not so good on more common conditions, especially where a clear and pragmatic guide is needed. There are five chapters in the section on asthma, and in fact the advice on the management of chronic and acute asthma was still pretty good. Although more accessible guidance on acute severe asthma might be found elsewhere, it did give brief up-to-date reviews of the use of intravenous bronchodilators and magnesium sulphate. However, this is not what this book is primarily for; its real strength is the ability to provide highly readable but comprehensive information on the whole range of paediatric respiratory problems.

I am going to keep this book right on my desk. I would consider it the best textbook in paediatric respiratory medicine and therefore a vital resource for specialist paediatric pulmonologists, trainees and paediatricians with an interest in respiratory medicine. While I don't think I can recommend to our trainees that they should keep it by their bed and read a daily passage, this is as close to a bible as it gets....

Tom Hilliard

Management of brain injured children, 2nd edition

Edited by Richard Appleton, Tony Baldwin. Published by Oxford University Press, Oxford, 2006, pp 398, £32.95 (paperback). ISBN 0-198-56724-3



The concept of multi-disciplinary working in child health is frequently paid lip-service by professionals but is less frequently achieved in practice. This important book on the management of brain injured children is a truly multi-disciplinary production from the head injury rehabilitation team led by Richard

Appleton at Alder Hey, Liverpool. The book is now in its second edition; the first edition, published in 1998, has been revised to provide a

comprehensive guide for professionals managing brain injured children. New information on long-term effects of acquired brain injury (ABI) and resuscitation advances has been included.

The 15 contributors cover acute treatment of brain injury, through nursing and therapy needs to the assessment of cognitive problems and re-integration into the home and educational environments. There is an excellent personal contribution by a survivor of ABI and her mother, which gives some insight into the effect on individuals and their families.

Advances in the management of children with ABI have meant improved survival rates but consequently higher morbidity in survivors, ranging from transient memory deficits to complex, multiple difficulties.

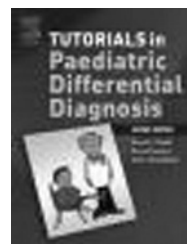
The book discusses the issues around giving long-term prognostic information to families following ABI and highlights problems such as the "sleeper effect", where an individual who has apparently made a good recovery presents years later with cognitive difficulties or school failure. The book is well-referenced with good quality neuro-imaging examples, but it could have benefited from more diagrams, particularly to help explain the chapter on cognitive assessment, and the images in the feeding assessment chapter are of disappointing quality.

ABI is an important subject – the average district general hospital can expect to see 10 children each year who will need rehabilitation – and this book is an excellent guide for the paediatrician and other professionals in the team. It deserves to be widely read.

Neil Harrower

Tutorials in paediatric differential diagnosis, 2nd edition

Edited by David J Field, David Isaacs, John Stroobant. Published by Elsevier, 2005, pp 288, £29.99 (paperback). ISBN 0-443-07100-4



As medical curricula nationally have moved towards a problem based approach, it is encouraging to find textbooks that mirror this way of learning. When a child presents in a paediatric assessment ward, they will complain of "vomiting" or "noisy breathing" not "problems with the gastrointestinal tract" or "problems with the respiratory system". This book is divided into chapters with titles that describe the child sitting in front of you, "The floppy baby" and "The crying baby". This simple labelling allows quick access to the appropriate topic to allow you to work through the problem.

This book does not cover everything you need to know in paediatrics, nor does it propose to. Its remit is described as "designed

to provide doctors involved in childcare with a logical approach to interpreting symptoms".

Divided into 40 short, accessible chapters, it covers almost all presentations you are likely to see during acute medical paediatric receiving. Within each chapter there is a brief introduction before a description of common diagnoses that should be considered in light of the particular presentation. The chapter is concluded in most instances by a clinical case which keeps the reader interested and grounded in the clinical relevance of the chapter. It is well laid out and easy to read. At times it feels a little too list based, but that is that nature of the book. It fulfils its title well by listing differential diagnoses and providing brief descriptions. This makes the book accessible for use as a quick reference during clinical work. Any further information may be sourced from weightier tomes. It never claims to replace your standard paediatric textbooks but instead directs your use of them.

An alternative use suggested by the authors is to use to topic headings to steer departmental teaching in "tutorials". I have not put this use into practice, but I think used in this way the book would help to ensure some of the most prevalent presenting complaints in paediatric practice are covered. I do not, on the other hand, feel it is a book useful for candidates preparing for MRCPCH; it is not detailed enough with its basic science information for Part 1 A&B. The information contained I would expect most doctors to have obtained through clinical practice by the time they are sitting the clinical examination.

Overall, I feel this is a good quality publication that fulfils its objectives and presents a wide variety of information in a clear and concise format. I feel it would be most useful to those just starting in acute general paediatrics, in particular, the new breed of FY2s who will need to become familiar with common presentations in a short period of time. With specialty placements changing every 4 months, books that allow easy access to core topics will become increasingly popular. I feel this book could be used as a first reference during clinical work and to assist with practice based learning.

Gemma Louise Duffy

CORRECTION

doi: 10.1136/adc.2005.081216corr1

Jackson L V, Thalange N K S, Cole T J. *Arch Dis Child* 2007;**92**:298–303. Blood pressure centiles for Great Britain. In the Abstract and in the Methods sections of this paper the expansion of the abbreviation "LMS" was published incorrectly. The correct expansion is "lambda-delta-sigma." We apologise for this error.

ORIGINAL ARTICLE

Blood pressure centiles for Great Britain

Lisa V Jackson, Nandu K S Thalange, Tim J Cole

Arch Dis Child 2007;92:298–303. doi: 10.1136/adc.2005.081216

See end of article for authors' affiliations

Correspondence to:
Dr L V Jackson, Willow
Wood Surgery, Aslake
Close, Sprowston, Norwich
NR7 8TT, UK; lisa.jackson@
nhs.net

Accepted 10 July 2006
Published Online First
11 August 2006

Objective: To produce representative cross-sectional blood pressure reference centiles for children and young people living in Great Britain.

Design: Analysis of blood pressure data from seven nationally representative surveys: Health Surveys for England 1995–8, Scottish Health Surveys 1995 and 1998, and National Diet & Nutrition Survey 1997.

Methods: Blood pressure was measured using the Dinamap 8100 with the same protocol throughout. Weight and height were also measured. Data for 11 364 males and 11 537 females aged 4–23 years were included in the analysis, after excluding 0.3% missing or outlying data. Centiles were derived for systolic, diastolic, mean arterial and pulse pressure using the lambda-mu-sigma (LMS) equations method.

Results: Blood pressure in the two sexes was similar in childhood, rising progressively with age and more rapidly during puberty. Systolic pressure rose faster and was appreciably higher in adult men than in adult women. After adjustment for age, blood pressure was related more to weight than height, the effect being stronger for systolic blood pressure. Pulse pressure peaked at 18 years in males and 16 years in females.

Conclusions: These centiles increase our knowledge of blood pressure norms in contemporary British children and young people. High blood pressure for age should be defined as blood pressure above the 98th centile, and high-normal blood pressure for age as blood pressure between the 91st and 98th centiles. The centiles identify children and young people with increased blood pressure, and will be of benefit to both clinical practice and research.

There is no satisfactory definition of hypertension in children.¹ As a result, blood pressure is often not measured in paediatric clinical practice, and understanding the clinical significance of blood pressure readings in children is hampered by the lack of satisfactory reference data with which to interpret them.

Reference blood pressure centiles should therefore improve the understanding of blood pressure variation in childhood. In Britain and worldwide, there have been many studies of childhood blood pressure, but all are of limited use in Great Britain owing to the use of non-representative populations, limited age ranges and mixed methodologies for blood pressure measurement. Accordingly, we have developed representative cross-sectional blood pressure references for children and young people living in Great Britain.

METHODS

Blood pressure data from seven national health and social surveys carried out between 1995 and 1998 were obtained from the UK Data Archive (<http://www.data-archive.ac.uk/>) (table 1). The data were originally collected on behalf of the Departments of Health and the Ministry of Agriculture Fisheries and Food, by the Joint Health Surveys Unit of Social and Community Planning Research and University College London, London, UK and the Social Survey Division of the Office for National Statistics and Medical Research Council, Human Nutrition Research, Cambridge, UK.

The survey samples were obtained by stratified multistage sampling techniques to ensure that there was a proportional representation of the population at large by sex, age, geographical region and social class.² In brief, the demographic characteristics of a geographical area are known from census and other data. Using this information, a representative sample of individuals from the target age groups for each survey was obtained. Households in geographical areas selected by post-code were contacted and asked to fill in a questionnaire to identify eligible young people. A subset of this initial sample

was then contacted by trained interviewers. The demographic characteristics of those agreeing to take part were determined and further targeted sampling undertaken to ensure the study sample remained representative. More information may be found in the published surveys.

Ethical approval was obtained from all areas in which the surveys were carried out. Participation was subject to informed consent. Data for the present analysis were excluded for participants who had eaten, consumed alcohol or smoked in the 30 min before being measured, and for those on anti-hypertensive drugs.

All seven surveys used the Dinamap 8100 (Critikon, Tampa, Florida, USA) with the same protocol to measure blood pressure. The use of an automatic oscillometric method was necessary for practicality, accuracy and reproducibility.^{3,4} Briefly, the blood pressure cuff was applied to the right arm. The lower margin of the cuff was placed about 2 cm above the elbow crease, with the arrow marked on the cuff placed over the brachial artery. The cuff was wrapped to a tightness allowing two fingers to be inserted under the top and bottom of the cuff. Four cuff sizes were available, the appropriate cuff size being determined by measurement of the mid-upper arm circumference (child cuff 10–19 cm, small adult cuff 17–25 cm, adult cuff 23–33 cm, large adult cuff 31–40 cm). The participants were comfortably seated, with their feet flat to the floor. Measurements of systolic, mean arterial and diastolic pressure were obtained after a 10–15 min rest period in triplicate, at minute intervals. The first reading was discarded and the mean of the second and third readings was used for analysis, as the first reading of a series of blood pressure measurements is typically higher with oscillometric devices.^{4,5} Pulse pressure was calculated by subtracting diastolic from systolic pressure.

For 73 (0.3%) participants, the blood pressure data were found to be either outliers or inconsistent with age, lying more than five SD from the median for age and sex. Hence blood

Abbreviation: SDS, standard deviation score

Table 1 Demographic characteristics of 22 974 participants aged 4–23.9 years from seven national health and social surveys

Survey	Year	England	Scotland	Wales	Age range (years)	Sample size
Health Survey for England	1995	✓			5–23	3485
Health Survey for England	1996	✓			5–23	4198
Health Survey for England	1997	✓			5–23	5520
Health Survey for England	1998	✓			4–23	3756
Scottish Health Survey	1995		✓		16–23	707
Scottish Health Survey	1998		✓		5–23	3043
National Diet & Nutrition Survey	1997	✓	✓	✓	4–19	1905
Overall						22 974

pressure data for 22 901 participants, 11 364 male and 11 537 female, aged 4–23.9 years were analysed.

Sex-specific smoothed centiles were derived using the lambda-mu-sigma (LMS) method⁶ for age and sex. The LMS method summarises the age-changing frequency distribution of blood pressure in terms of three curves: the L curve defines the skewness, the M curve the median and the S curve the coefficient of variation as functions of age. Centile charts were drawn with centiles spaced two-thirds of an SD score (SDS) apart, ranging from the 0.4th centile (−2.67 SDS) through to the 99.6th centile (+2.67 SDS), consistent with other anthropometric charts in current use in the UK.⁷

The relationship of systolic and diastolic blood pressure, weight and height was investigated through the multiple regression of blood pressure on weight and height, after adjusting the three variables for age and sex by converting them to SDS. The British 1990 reference⁸ was used for height and weight, and the internal reference for blood pressure. For measuring weight and height in subjects age ≥23 years was taken as 22.99 (the upper limit of the British reference). Sex effects were tested for in the regression by including sex and its interactions with height and weight.

RESULTS

Table 2 summarises the data for 22 901 participants with both systolic and diastolic blood pressure. Mean arterial pressure, height and weight were missing for 8%, 1% and 2% of participants, respectively. By year of age the sample consisted of 114 participants aged 4 years, 1181–1581 per year between 5 and 16 years, and 715–950 per year between 17 and 23 years. Height was very similar to the British 1990 reference (mean SDS 0.0), while weight and body mass index (weight (kg)/height² (m²)) were slightly increased (mean SDS 0.3–0.4).

The data were used to construct blood pressure centile charts for systolic, diastolic, mean arterial and pulse pressure (figs 1–4). Blood pressure in the two sexes was similar before puberty,

but the pubertal rise was more marked in boys. Pulse pressure peaked at 18 years in male participants and at 16 years in female participants, corresponding to the end of puberty.

Table 3 summarises the multiple regression of blood pressure on weight and height, each adjusted for age and sex by

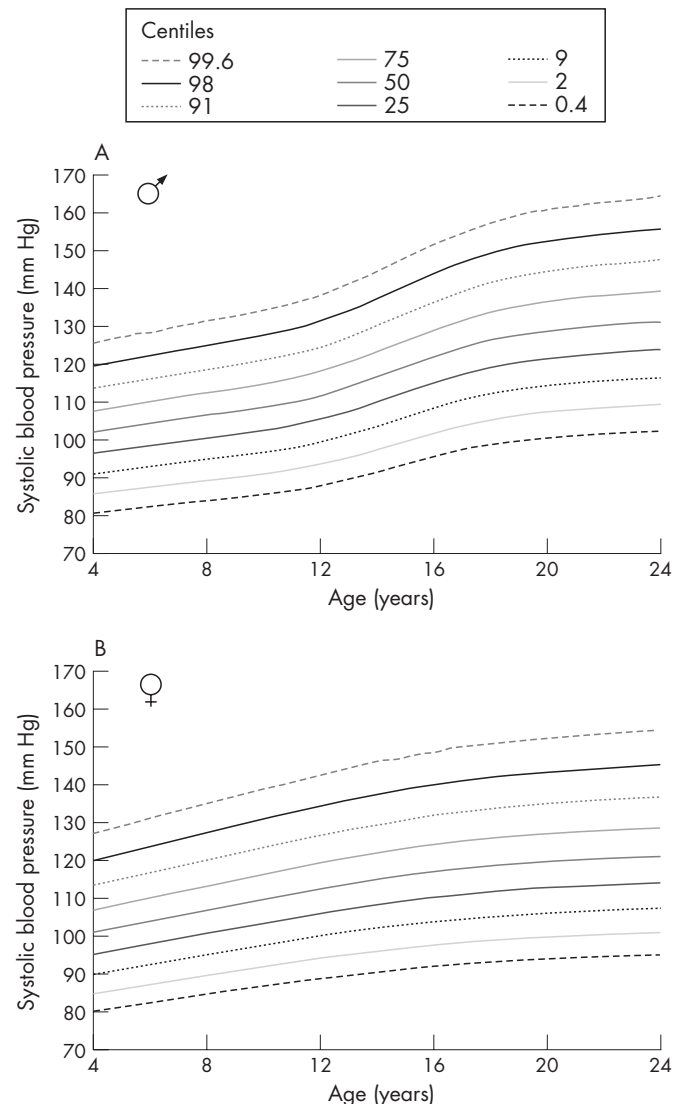


Figure 1 Systolic blood pressure centiles in male (A) and female participants (B). The centiles are spaced two-thirds of a standard deviation score apart. Systolic pressure rises progressively with age, but rises more steeply in puberty, particularly in boys.

Table 2 Summary statistics for 22 901 participants with valid data

Variable	n	Mean	SD
Men (%)	22 901	49.6	—
Age (years)	22 901	13.1	5.2
Height (cm)	22 676	148.9	21.3
Weight (kg)	22 485	46.4	19.9
Height SDS (British 1990)	22 676	−0.03	1.08
Weight SDS (British 1990)	22 485	0.27	1.14
Body mass index SDS (British 1990)	22 425	0.36	1.11

SDS, standard deviation score.

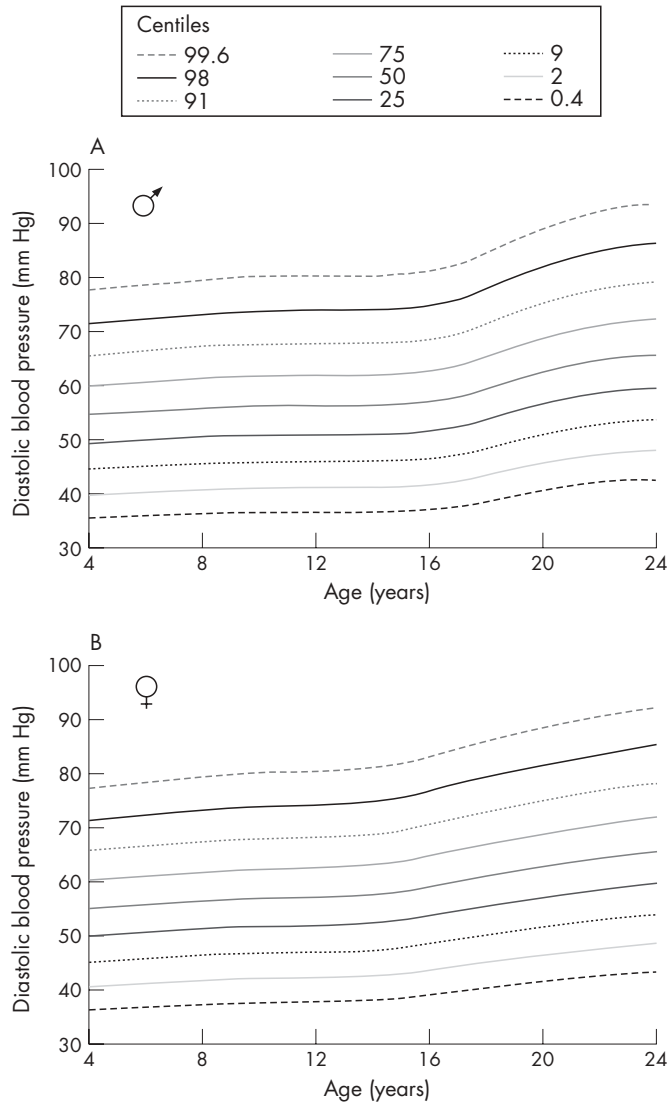


Figure 2 Diastolic blood pressure centiles in male (A) and female participants (B). The centiles are spaced two-thirds of a standard deviation apart. Diastolic pressure rises slowly in childhood, but as with systolic pressure, rises more steeply in puberty.

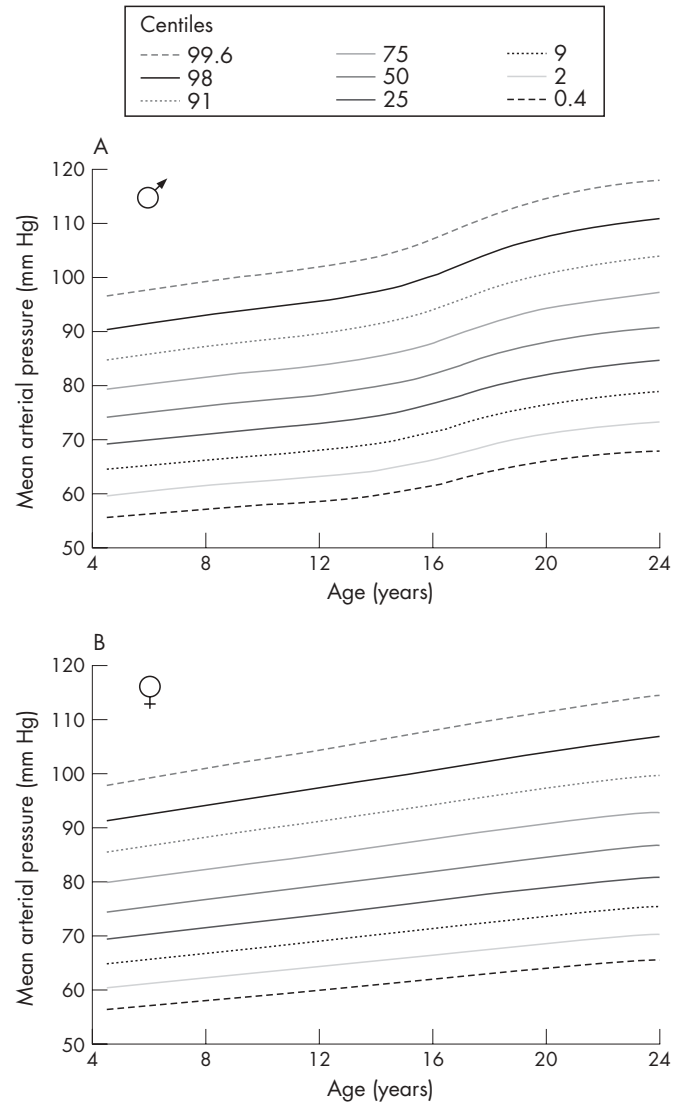


Figure 3 Mean arterial pressure centiles in male (A) and female participants (B). The centiles are spaced two-thirds of a standard deviation apart. Mean arterial pressure rises progressively with age.

converting to SDS. This adjustment allowed the data for both sexes and all ages to be combined. Results are also given by sex, although they do not differ significantly; hence the combined results are valid. Weight had a large and positive effect on blood pressure ($p < 0.001$), whereas height had a smaller negative effect ($0.005 < p < 0.001$). A 1 SD increase in weight was associated with a 0.3 SD increase in systolic pressure and a 0.08 SD increase in diastolic pressure, whereas a 1 SD increase in height was associated with a 0.03 SD reduction in both systolic and diastolic pressure. Thus, on average, for any given weight, a taller (and hence thinner) individual had lower blood pressure. Analysing the data in separate age groups showed the associations in late puberty to be stronger than before or after.

These results suggest that body size (ie, weight) and obesity (weight adjusted for height) both play a role in raising blood pressure, particularly systolic blood pressure, 8% of the variation of which was explained by weight and height. The effect on diastolic blood pressure (0.5% of variance explained) was much smaller.

Using the British Hypertension Society cut-offs for hypertension,⁹ 23% of men and 6% of women exceeded the systolic

cut-off, and 1.0% of men and 0.8% of women exceeded the diastolic cut-off by age 24 years.

DISCUSSION

The blood pressure centiles presented here are based on data collected using a consistent and rigorous method in representative samples of nearly 23 000 children and young people living in Great Britain. As such, we believe they are the most accurate characterisation of normal blood pressure in any country to date.

It is well recognised that children's blood pressure tends to "track" over time.¹⁰⁻¹⁴ Moreover, high blood pressure in children is associated with the development of atherosclerosis,¹⁵⁻¹⁹ especially in those with additional risk factors, notably obesity.¹⁶⁻²⁰ The charts will aid the timely recognition and monitoring of individuals with high blood pressure and hypertension, and facilitate the detection of children with secondary hypertension, consequent on renal, endocrine or other disease.¹ Blood pressure monitoring is also important in children at risk of hypertension and/or vascular disease, such as those with obesity, diabetes, renal disease, or those receiving

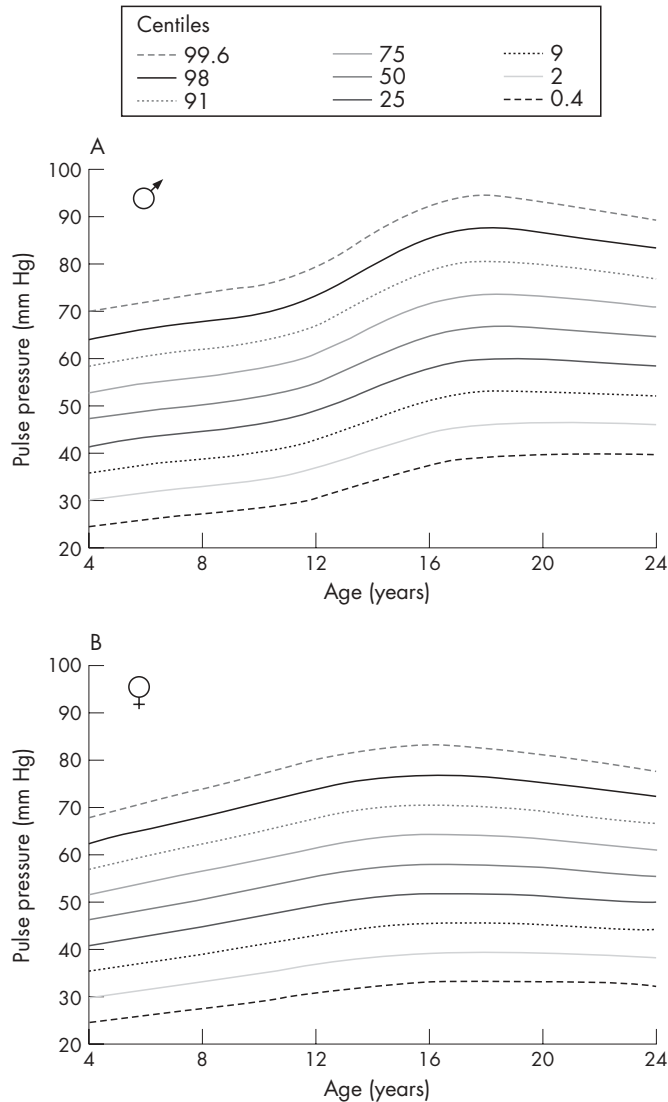


Figure 4 Pulse pressure centiles in male (A) and female participants (B). The centiles are spaced two-thirds of a standard deviation score apart. Pulse pressure rises progressively until the end of puberty and then falls again.

steroids or stimulant drugs and where a family history of hypertension is present.

We have used a standard nine-centile format consistent with other charts in use in the UK.⁷ The charts show a progressive rise in systolic and diastolic pressure with increasing age, the rise being more marked in males during puberty. This is consistent with an effect of body size (indicated by weight) and obesity (weight adjusted for height) on blood pressure, the

effect being stronger for systolic blood pressure. Thus males, who gain more weight at puberty than females, have significantly higher blood pressures, with almost a quarter satisfying the British Hypertension Society definition of hypertension,⁹ defined as systolic pressure >140 mm Hg and/or diastolic pressure >90 mm Hg, by the age of 24 years. The high systolic pressures in older teenagers and young adults, particularly men, are of special concern. However, blood pressure measurements on a single occasion are insufficient for the determination of high or high-normal blood pressure for age,^{21, 22} in the absence of evidence of a pathological cause or end organ damage, especially in children, who are more prone to “white coat” hypertension.²³ Repeated measurements typically show that the majority of children with increased blood pressure on a single occasion subsequently have normal blood pressure.²⁴

Pulse pressure^{25, 26} and mean arterial pressure²⁷ have been found to be significant determinants of morbidity and mortality in adults. The significance of these measures in children is unknown, but pulse pressure may be an indicator of early arterial disease, as has been found in young adults with type 1 diabetes.²⁷ Of note, we found that pulse pressure peaks at the end of puberty in both sexes, before falling in young adult life (fig 4) in contrast with systolic, diastolic and mean arterial pressures, which rise progressively with age (figs 1–3). A knowledge of normal ranges for pulse pressure and mean arterial pressure should aid research in this area.

The use of oscillometric blood pressure measurements was dictated by the nature of the health and social surveys, which required a reliable, reproducible and accurate method for determining blood pressure, using multiple observers.^{3, 4} The Dinamap 8100 was subject to a rigorous calibration study³ to ensure its validity (although the calibration study did not include participants aged <16 years). However, the Dinamap monitor has been compared with direct radial artery pressure and central aortic pressure measurements in infants and children and was found to be superior to the auscultatory method.^{28, 29} Moreover, particularly in young children, the conventional mercury sphygmomanometer can be difficult to use,^{4, 20} with the Korotkoff sounds hard to distinguish, so, increasingly, automated oscillometric devices are being used in clinical practice.³⁰

O'Brien *et al*,³¹ using the British Hypertension Society protocol, graded the Dinamap 8100 B for systolic blood pressure and D for diastolic blood pressure compared with the conventional mercury sphygmomanometer in adults.³² Paediatric studies have generally found significant differences, particularly between diastolic pressure assessed by fourth-phase Korotkoff sounds.^{33–35} However, an Australian study of prepubertal children with type 1 diabetes using the British Hypertension Society protocol graded the Dinamap B for both systolic and diastolic pressure.³⁶

The Dinamap 8100 and other oscillometric devices produce results that differ in comparison with the mercury sphygmomanometer. These differences have been attributed to

Table 3 Relationship of systolic and diastolic blood pressure with weight and height by sex (all variables expressed as standard deviation score)

Outcome measure	Sex	n	Weight (SDS)	Height (SDS)	R ² (%)
Systolic blood pressure (SDS)	Men	11 153	0.279 (0.011)	-0.033 (0.011)	8.6
	Women	11 272	0.258 (0.010)	-0.039 (0.010)	7.6
	Combined	22 425	0.268 (0.007)	-0.035 (0.008)	8.1
Diastolic blood pressure (SDS)	Men	11 153	0.070 (0.011)	-0.033 (0.012)	0.4
	Women	11 272	0.081 (0.010)	-0.032 (0.011)	0.6
	Combined	22 425	0.076 (0.008)	-0.033 (0.008)	0.5

SDS, standard deviation score. The table gives regression coefficients (SEs).

What is already known on this topic

- Blood pressure rises through childhood and childhood blood pressure strongly predicts adult blood pressure.
- This rise in blood pressure is substantially determined by weight.
- As with growth, blood pressure is an important parameter of child health.
- Furthermore, atherosclerosis and hypertension may have their origins in childhood, particularly in those with additional risk factors—for example, obesity, renal disease or diabetes.

What this study adds

- These blood pressure centiles compiled from nationally representative data are the most comprehensive attempt to characterise normal blood pressure in childhood in Great Britain.
- The centiles complement existing charts for height, weight and body mass index and other parameters in evaluating the health of children.
- This information will contribute to a better understanding of blood pressure in childhood and aid further research.

inaccuracies³⁷ but simply reflect the fact that different methods yield different results.^{1 38} However, in view of these differences, blood pressure results recorded with the mercury sphygmomanometer should be referenced to these centiles with caution.

The definition of hypertension in children is problematic. Use of the British Hypertension Society cut-offs in adults is justified by adverse health outcomes in association with hypertension.⁹ However, no single cut-off can define hypertension in children owing to the normal rise in blood pressure with age, and the paucity of evidence about what constitutes hypertension in children.¹ Consequently, we suggest that, in children, those above the 98th centile on repeated occasions are stated to have high blood pressure for age, whereas those lying between the 91st and 98th centiles are stated to have high-normal blood pressure for age. These cut-offs are similar to recommendations made in the Taskforce Report on High Blood Pressure in Children and Adolescents in the USA.³⁹ Our centile-based definitions predict a prevalence of 2.3% for high blood pressure (>2 SDS) and 6.9% for high-normal blood pressure (>1.33 SDS). These centiles should facilitate ongoing research into the importance of high or high-normal blood pressure in children, and serve as a basis for defining hypertension in childhood.

The strong association between high blood pressure and weight/obesity that we and others have found^{40 41} is of particular concern given the well documented rise in childhood obesity.⁴² Childhood obesity, and its health consequences—including hypertension, metabolic syndrome and type 2 diabetes—present a major challenge for the coming years and demand vigilance and concerted action from all healthcare professionals to mitigate the adverse health consequences for children and young people.

ACKNOWLEDGEMENTS

We thank Dr Graham Derrick, Consultant Paediatric Cardiologist, Great Ormond Street Hospital for Children NHS Trust, and Dr Carlo Acerini, University Lecturer in Paediatrics, Cambridge University, for their helpful comments.

Authors' affiliations

Lisa V Jackson, Willow Wood Medical Practice and University of East Anglia School of Medicine, Health Policy and Practice, Norwich, UK
Nandu K S Thalange, Norfolk and Norwich University Hospital, Norwich, UK

Tim J Cole, Centre for Paediatric Epidemiology and Biostatistics, Institute of Child Health, University College London, London, UK

Funding: TJC is funded by the Medical Research Council.

Research at the UCL Institute of Child Health and Great Ormond Street Hospital for Children NHS Trust benefits from R&D funding received from the NHS Executive.

Competing interests: Subsequent to the presentation of data at the Spring Meeting in April 2003, blood pressure centile charts using these data were published commercially.

Previous publication: Some of this work was presented as an oral presentation at the Royal College of Paediatrics & Child Health Spring Meeting, April 2003: G203. Blood pressure centiles for children and young people aged 4–24 years in Great Britain. *Arch Dis Child* 2003; **88**:A66.

Contributors: all authors contributed to the design, analysis and writing up of the paper. LVJ is guarantor.

REFERENCES

- 1 **Goonasekera CDA**, Dillon MJ. Measurement and interpretation of blood pressure. *Arch Dis Child* 2000; **82**:261–5.
- 2 **Elliot D**. Optimising sample design for surveys of health and related behaviour and attitudes. *Survey Methodol Bull* 1995; **36**:8–17.
- 3 **Bolling K**. *The Dinamap 8100 calibration study*. London: HMSO, 1994.
- 4 **Gillman MW**, Cook NR. Blood pressure measurement in childhood epidemiological studies. *Circulation* 1995; **92**:1049–57.
- 5 **Park MK**, Menard SM. Normative oscillometric blood pressure values in the first 5 years in an office setting. *Am J Dis Child* 1989; **143**:860–4.
- 6 **Cole TJ**, Green PJ. Smoothing reference centile curves: the LMS method and penalized likelihood. *Stat Med* 1992; **11**:1305–19.
- 7 **Cole TJ**. Do growth chart centiles need a face lift? *BMJ* 1994; **308**:641–2.
- 8 **Freeman JV**, Cole TJ, Chinn S, *et al*. Cross sectional stature and weight reference curves for the UK, 1990. *Arch Dis Child* 1995; **73**:17–24.
- 9 **Williams B**, Poulter NR, Brown MJ, *et al*. British Hypertension Society guidelines for hypertension management 2004 (BHS-IV): summary. *BMJ* 2004; **328**:634–40.
- 10 **Bao W**, Threefoot SA, Srinivasan SR, *et al*. Essential hypertension predicted by tracking of elevated blood pressure from childhood to adulthood: the Bogalusa Heart Study. *Am J Hypertens* 1995; **8**:657–65.
- 11 **Klumbiene J**, Sileikiene L, Milasauskiene Z, *et al*. The relationship of childhood to adult blood pressure: longitudinal study of juvenile hypertension in Lithuania. *J Hypertens* 2000; **18**:531–8.
- 12 **O'Sullivan JJ**, Derrick G, Foxall RJ. Tracking of 24-hour and casual blood pressure: a 1-year follow-up study in adolescents. *J Hypertens* 2000; **18**:1193–6.
- 13 **Cook NR**, Gillman MW, Rosner BA, *et al*. Prediction of young adult blood pressure from childhood blood pressure, height, and weight. *J Clin Epidemiol* 1997; **50**:571–9.
- 14 **Lambrechtsen J**, Rasmussen F, Hansen HS, *et al*. Tracking and factors predicting rising in 'tracking quartile' in blood pressure from childhood to adulthood: Odense Schoolchild Study. *J Hum Hypertens* 1999; **13**:385–91.
- 15 **Newman WP III**, Freedman DS, Voors AW, *et al*. Relation of serum lipoprotein levels and systolic blood pressure to early atherosclerosis: the Bogalusa Heart Study. *N Engl J Med* 1986; **314**:138–144.
- 16 **Berenson GS**, Srinivasan SR, Bao W, *et al*. Association between multiple cardiovascular risk factors and atherosclerosis in children and young adults. The Bogalusa Heart Study. *N Engl J Med* 1998; **338**:1650–6.
- 17 **Mahoney LT**, Burns TL, Stanford W, *et al*. Coronary risk factors measured in childhood and young adult life are associated with coronary artery calcification in young adults: the Muscatine Study. *J Am Coll Cardiol* 1996; **27**:277–84.
- 18 **Li S**, Chen W, Srinivasan SR, *et al*. Childhood cardiovascular risk factors and carotid vascular changes in adulthood: the Bogalusa Heart Study. *JAMA* 2003; **290**:2271–6.
- 19 **Raitakari OT**, Juonala M, Kahonen M, *et al*. Cardiovascular risk factors in childhood and carotid artery intima-thickness in adulthood: the Cardiovascular Risk in Young Finns Study. *JAMA* 2003; **290**:2277–83.
- 20 **Reinehr T**, Andler W. Changes in the atherogenic risk factor profile according to degree of weight loss. *Arch Dis Child* 2004; **89**:419–22.
- 21 **de Swiet M**. The epidemiology of hypertension in children. *Br Med Bull* 1986; **42**:172–5.
- 22 **Adrogué HE**, Sinaiko AR. Prevalence of hypertension in junior high school-aged children: effect of new recommendations in the 1996 Updated Task Force Report. *Am J Hypertens* 2001; **14**:412–14.
- 23 **Hornsby JL**, Mongan PF, Taylor AT, *et al*. 'White coat' hypertension in children. *J Fam Pract* 1991; **33**:617–23.
- 24 **Madhavan S**, Ooi WL, Cohen H, *et al*. Relation of pulse pressure and blood pressure reduction to the incidence of myocardial infarction. *Hypertension* 1994; **23**:395–401.

- 25 **Benetos A**, Rudnichi A, Safar M, *et al*. Pulse pressure and cardiovascular mortality in normotensive and hypertensive subjects. *Hypertension* 1998;**32**:560–4.
- 26 **van Trijp MJ**, Grobbee DE, Peeters PH, *et al*. Average blood pressure and cardiovascular disease-related mortality in middle-aged women. *Am J Hypertens* 2005;**18**:197–201.
- 27 **Schram MT**, Chaturvedi N, Fuller JH, *et al*. Pulse pressure is associated with age and cardiovascular disease in type 1 diabetes: the Eurodiab Prospective Complications Study. *J Hypertens* 2003;**21**:2035–44.
- 28 **Park MK**, Menard SM. Accuracy of blood pressure measurement by the Dinamap monitor in infants and children. *Pediatrics* 1987;**79**:907–14.
- 29 **Colan SD**, Fujii A, Borow KM, *et al*. Noninvasive determination of systolic, diastolic and end-systolic blood pressure in neonates, infants and young children: comparison with central aortic pressure measurements. *Am J Cardiol* 1983;**52**:867–70.
- 30 **O'Brien E**. Demise of the mercury sphygmomanometer and the dawning of a new era in blood pressure measurement. *Blood Press Monit* 2003;**8**:19–21.
- 31 **O'Brien E**, Petrie J, Littler W, *et al*. The British Hypertension Society protocol for the evaluation of automated and semi-automated blood pressure measuring devices with special reference to ambulatory systems. *J Hypertens* 1990;**8**:607–19.
- 32 **O'Brien E**, Mee F, Atkins N, *et al*. Short report: accuracy of the Dinamap portable monitor, model 8100 determined by the British Hypertension Society protocol. *J Hypertens* 1993;**11**:761–3.
- 33 **Wattigney WA**, Webber LS, Lawrence MD, *et al*. Utility of an automatic instrument for blood pressure measurement in children. The Bogalusa Heart Study. *Am J Hypertens* 1996;**9**:256–62.
- 34 **Barker ME**, Shiell AW, Law CM. Evaluation of the Dinamap 8100 and Omron M1 blood pressure monitors for use in children. *Paediatr Perinat Epidemiol* 2000;**14**:179–86.
- 35 **Park MK**, Menard SW, Yuan C. Comparison of auscultatory and oscillometric blood pressures. *Arch Pediatr Adolesc Med* 2001;**155**:50–3.
- 36 **Jin RZ**, Donaghue KC, Fairchild J, *et al*. Comparison of Dinamap 8100 with sphygmomanometer blood pressure measurement in a prepubertal diabetes cohort. *J Paediatr Child Health* 2001;**37**:545–9.
- 37 **O'Brien E**, Atkins N. Inaccuracy of the Dinamap 8100 portable monitor. *Lancet* 1997;**349**:1026.
- 38 **Friedman B**. Accuracy of Dinamap monitors. *Lancet* 1997;**350**:217–18.
- 39 **National Blood Pressure Education Working Group on High Blood Pressure in Children and Adolescents**. Fourth report on the diagnosis, evaluation and treatment of high blood pressure in children and adolescents: a working group report from the National High Blood Pressure Education Program. *Pediatrics* 2004;**114**:555–76.
- 40 **St George IM**, Williams SM, Silva PA. The stability of high blood pressure in Dunedin children: an eight year longitudinal study. *NZ Med J* 1990;**103**:115–17.
- 41 **de Swiet M**, Fayers P, Shinebourne EA. Blood pressure in first 10 years of life: the Brompton study. *BMJ* 1992;**304**:23–6.
- 42 **Chinn S**, Rona RJ. Prevalence and trends in overweight and obesity in three cross sectional studies of British children 1974–94. *BMJ* 2001;**322**:24–6.

IMAGES IN PAEDIATRICS.....

doi: 10.1136/adc.2006.112771

An interesting facial rash



Figure 1 Informed consent was obtained for publication of this figure.

A 7-year-old boy presented to paediatrics with a 3-week history of a worsening facial rash, which was intermittently itchy. He was otherwise well. He had two guinea pigs as pets. On examination, an extensive scaling erythema was noticed with a definite edge involving the upper eyelids, the bridge of the nose and extending onto both cheeks (see fig 1).

A provisional diagnosis of tinea faciei was made; however, cutaneous lupus was also considered. While mycology results were awaited, topical terbinafine was given, with little effect. Microscopy revealed a dermatophyte infection with *Trichophyton mentagrophytes*, and a 3-week course of oral terbinafine (125 mg oral dosage once daily) was given. The rash resolved completely, leaving post-inflammatory hyperpigmentation only.

Tinea facialis/faciei is a dermatophytosis of the glabrous facial skin, characterised by a well-circumscribed, often asymmetric, erythematous patch with an elevated border and central regression. It may be asymptomatic or present with pruritus, or, occasionally, photosensitivity that may lead to diagnostic confusion with cutaneous lupus.¹ It is the most commonly misdiagnosed dermatophytosis. Other differential diagnoses include eczema, seborrhoeic dermatitis² and rosacea.

It is most common in children, with predisposing factors including exposure to animals, chronic topical steroid use and spread from tinea capitis. The most frequent organisms involved are *T mentagrophytes*, *T rubrum* and *T tonsurans*. However, cases caused by *Microsporum audouinii* and *M canis* occur worldwide. Most cases are given short-term oral antifungal treatment, but milder cases may respond to topical imidazoles. Affected animals and family members should also be treated.

Caroline A Love

Department of Dermatology, Rowan House, Whiston Hospital, Merseyside, UK

John A Sills, Judith M Ellison

St Helens and Knowsley NHS Trust, Merseyside, UK

Correspondence to: Dr C A Love, Department of Dermatology, Rowan House, Whiston Hospital, Warrington Road, Prescott, Merseyside L35 5DR, UK; drcalove@hotmail.com

Competing interests: None declared.

REFERENCES

- 1 **Cirillo-Hyland V**, Humphreys T, Elenitsas R. Tinea faciei. *J Am Acad Dermatol* 1993;**29**:119–20.
- 2 **Gorani A**, Oriani A, Cambiaghi S. Seborrhoeic dermatitis-like tinea faciei. *Pediatr Dermatol* 2005;**22**:243–4.