Blood pressure centiles for Great Britain

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


**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 height than to weight, 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. 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. 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 postcode 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. Briefly, the blood pressure cuff was applied to the right arm. The lower margin of the cuff was placed about 2 cm above the brachial artery, with the arrow marked on the cuff placed over 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. 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
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) method 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.

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 reference 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 converting them to SDS.

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

<table>
<thead>
<tr>
<th>Survey</th>
<th>Year</th>
<th>England</th>
<th>Scotland</th>
<th>Wales</th>
<th>Age range (years)</th>
<th>Sample size</th>
</tr>
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<tbody>
<tr>
<td>Health Survey for England</td>
<td>1995</td>
<td></td>
<td></td>
<td></td>
<td>5-23</td>
<td>3485</td>
</tr>
<tr>
<td>Health Survey for England</td>
<td>1996</td>
<td></td>
<td></td>
<td></td>
<td>5-23</td>
<td>4198</td>
</tr>
<tr>
<td>Health Survey for England</td>
<td>1997</td>
<td></td>
<td></td>
<td></td>
<td>5-23</td>
<td>5520</td>
</tr>
<tr>
<td>Health Survey for Scotland</td>
<td>1998</td>
<td></td>
<td></td>
<td></td>
<td>4-23</td>
<td>3756</td>
</tr>
<tr>
<td>Scottish Health Survey</td>
<td>1995</td>
<td></td>
<td></td>
<td></td>
<td>16-23</td>
<td>707</td>
</tr>
<tr>
<td>Scottish Health Survey</td>
<td>1998</td>
<td></td>
<td></td>
<td></td>
<td>5-23</td>
<td>3043</td>
</tr>
<tr>
<td>National Diet &amp; Nutrition Survey</td>
<td>1997</td>
<td></td>
<td></td>
<td></td>
<td>4-19</td>
<td>1905</td>
</tr>
<tr>
<td>Overall</td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
<td>22,974</td>
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## Table 2
Summary statistics for 22,901 participants with valid data

<table>
<thead>
<tr>
<th>Variable</th>
<th>n</th>
<th>Mean</th>
<th>SD</th>
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<tbody>
<tr>
<td>Men (%)</td>
<td>22,901</td>
<td>49.6</td>
<td></td>
</tr>
<tr>
<td>Age (years)</td>
<td>22,901</td>
<td>13.1</td>
<td>5.2</td>
</tr>
<tr>
<td>Height (cm)</td>
<td>22,676</td>
<td>148.9</td>
<td>21.2</td>
</tr>
<tr>
<td>Weight (kg)</td>
<td>22,485</td>
<td>46.4</td>
<td>19.9</td>
</tr>
<tr>
<td>Height SDS (British 1990)</td>
<td>22,676</td>
<td>−0.03</td>
<td>1.08</td>
</tr>
<tr>
<td>Weight SDS (British 1990)</td>
<td>22,485</td>
<td>0.27</td>
<td>1.14</td>
</tr>
<tr>
<td>Body mass index SDS (British 1990)</td>
<td>22,425</td>
<td>0.36</td>
<td>1.11</td>
</tr>
</tbody>
</table>

SDS, standard deviation score.

![Figure 1](http://www.archdischild.com) Systolic blood pressure centiles in male (A) and female (B) participants. 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.
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 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.

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.
stereoskopic 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. 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 (indexed 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, 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, in the absence of evidence of a pathological cause or end organ damage, especially in children, who are more prone to "white coat" hypertension. Repeated measurements typically show that the majority of children with increased blood pressure on a single occasion subsequently have normal blood pressure.

Pulse pressure rise being more marked in males during puberty. This is consistent with the relationship of systolic and diastolic blood pressure with weight and height by sex (all variables expressed as standard deviation score).

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

<table>
<thead>
<tr>
<th>Outcome measure</th>
<th>Sex</th>
<th>n</th>
<th>Weight (SDS)</th>
<th>Height (SDS)</th>
<th>R² (%)</th>
</tr>
</thead>
<tbody>
<tr>
<td>Systolic blood pressure (SDS)</td>
<td>Men</td>
<td>11 153</td>
<td>0.279 (0.011)</td>
<td>-0.033 (0.011)</td>
<td>8.6</td>
</tr>
<tr>
<td></td>
<td>Women</td>
<td>11 272</td>
<td>0.258 (0.010)</td>
<td>-0.039 (0.010)</td>
<td>7.6</td>
</tr>
<tr>
<td></td>
<td>Combined</td>
<td>22 425</td>
<td>0.268 (0.007)</td>
<td>-0.035 (0.008)</td>
<td>8.1</td>
</tr>
<tr>
<td>Diastolic blood pressure (SDS)</td>
<td>Men</td>
<td>11 113</td>
<td>0.070 (0.011)</td>
<td>-0.033 (0.012)</td>
<td>0.4</td>
</tr>
<tr>
<td></td>
<td>Women</td>
<td>11 272</td>
<td>0.081 (0.010)</td>
<td>-0.032 (0.011)</td>
<td>0.6</td>
</tr>
<tr>
<td></td>
<td>Combined</td>
<td>22 425</td>
<td>0.076 (0.008)</td>
<td>-0.033 (0.008)</td>
<td>0.5</td>
</tr>
</tbody>
</table>

SDS, standard deviation score.
The table gives regression coefficients (SEs).
importance of high or high-normal blood pressure in children, definitions predict a prevalence of 2.3% for high blood pressure. Consequently, we suggest that, in children, those with blood pressure in excess of 95th centile for age may be considered hypertensive. Paucity of evidence about what constitutes hypertension in childhood, owing to the normal rise in blood pressure with age, and the absence of a single cut-off can define hypertension in children mandates 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.

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.

REFERENCES

An interesting facial rash

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-inflamatory 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. It is the most commonly misdiagnosed dermatophytosis. Other differential diagnoses include eczema, seborrhoeic dermatitis 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.

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Competing interests: None declared.

**REFERENCES**

13. Blood pressure centiles

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


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


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 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 AkB. 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 LV, Thalange NK, Cole TJ. 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-mu-sigma.” We apologise for this error.