Identification of factors affecting infant growth in developing countries

M P ECCLES,* T J COLE,† AND R G WHITEHEAD†


SUMMARY The anthropometric progress of seven infants was followed throughout their first year of life. Weight, length, mid upper arm circumference, triceps and subscapular skinfold thicknesses were measured on a mean (SD) of 30 (6) occasions with weight alone being measured on a further 6 (2) occasions. The effects of infection and energy intake were identified and illustrated using both standard deviation (SD) score graphs for individual subjects, and more traditionally, regression analysis for the group. Diarrhoea and vomiting, pneumonia, and diarrhoea alone each resulted in significant reductions in growth velocity of 30 g/day. Multi-measurement SD score graphs showed the effects of all illnesses, and permitted simultaneous comparison of anthropometric measurements.

The effect of acute infection on growth has been widely studied, predominantly in third world countries. In Mexico1 and Guatemala2 the infective causes of weight loss were diarrhoea, dysentery, and bronchopneumonia. In rural Gambia they were gastroenteritis and malaria,3 4 and in Uganda they were gastroenteritis and measles.4 A recent study from urban Gambia again identified diarrhoea and also lower respiratory tract infection.5

Such studies have traditionally relied upon mean weight deficit calculated by regression analysis to quantify the effects of infection, showing reductions in weight gain of as much as 746 g/month6 or 22 g/day for the various infections.5

An alternative approach is to express weight, and any other available anthropometric measurements, as standard deviation (SD) scores adjusted for age. The graphical presentation of such information has, however, always been limited by an inability to provide a comprehensive anthropometric picture of an individual subject without recourse to five or six separate centile charts.

There is also the question of the appropriateness of the standards used. In adopting the National Center for Health Statistics (NCHS) growth curves6 as their international standard, the World Health Organisation (WHO) defined their standards for body weight, length, and head circumference.7 The values that they chose for mid upper arm circumference were based on the data of Wolanski.8 There were no values adopted for skinfold thickness. In the United Kingdom the most accessible skinfold values are those of Tanner and Whitehouse.9 The data for all these standards were collected between 15 and (in the case of the NCHS values for infants) 59 years ago. These standards are taken from different populations and their use to provide a composite set of multiple anthropometric measurements for an individual subject is of doubtful value.

A set of centiles that allows a comparison of several measurements is that derived from the data of an infant growth study in Cambridge, United Kingdom.10–12 The centiles cover weight, length, mid upper arm and head circumference, and triceps and subscapular skinfold thicknesses. They are constructed using the LMS method, which gives smooth centile curves while adjusting for skewness thus permitting skewed data (such as weight or skinfold thickness) to be presented in SD form.13 14 The method also allows the conversion of SD scores directly into centile values using normal distribution tables. An SD (or Z) score of, for example, an infant’s weight at a given age, expresses weight relative to the population mean in the form of a normally distributed variable that has a mean of zero and an SD of one. An SD score can be positive or negative, depending on whether the weight is above or below the mean, and an infant growing along the 50th centile will, on an SD score graph, appear on the horizontal line corresponding to zero. The range of values can be set as required, but the range of normal is conventionally from −2 SD to +2 SD. The use of SD scores is in line with the WHO recommendations7 15 and using these centiles.
it is possible to construct, on one graph, a set of SD scores for multiple measurements, which permit all a child's anthropometric measurements and the factors affecting them to be viewed at once.

In the present study we prospectively followed seven infants through their first year of life to identify those factors affecting their growth such as frequent acute infection—and to assess SD score graphs of multiple measurements as a means of presenting a comprehensive anthropometric picture of individual infants.

Subjects and methods

The study took place in the village of Keneba, which is in the Kiang West district of The Gambia, West Africa. It is a rural subsistence farming community and has been the site of a nutritional field station for 15 years. It has a population of 1250 and the community has previously been described in detail.16 17

Seven sequential births from the village were recruited to a study18 of which this formed one part. The study lasted for the first year of the infants' lives.

Measurements were made of body weight, length, mid upper arm circumference, and triceps and subscapular skinfold thickness. These were measured every 14 days when the infants were well and up to six times in the 21 days after the onset of an episode of acute illness. All the measurements were made by one person (MPE). The values were converted into SD scores using the methods described by Cole,13 and were then plotted, with all of an individual infants' five sets of values on one graph.

To establish a coefficient of variation for each measurement, repeated short term measurements were made on a group of 10 children. Coefficients of variation were low, being <0-1% for weight and head circumference, 2-8% for mid upper arm circumference, and 6-6% for skinfold thicknesses.

I llnesses were recorded and treated at the daily, open access, medical clinic held in the village. Diagnoses were made by either of the resident clinicians according to predetermined criteria.18 The infants did not attend any other medical service. The illnesses fell into 11 diagnostic groups and for the purposes of the regression analysis each was represented by a two letter code, for instance pneumonia was coded as RP and diarrhoea as DU.

The effect of infection on weight gain was calculated by regression analysis using the Genstat 5 statistical language. Weight gain was calculated, in g/day, as the difference between successive weights divided by the time interval in days. Individual weights were coded in two ways. Absence of illness was coded as 'well' and a weight measured during an episode of illness was coded by a single diagnosis with time from onset. Thus weights measured on the third, seventh, 10th, and 13th days of an episode of pneumonia were coded as RP03, RP07, RP10, and RP13. Well measurements were coded as WE00. Using five days as the duration of an illness, binary independent variables were constructed for each illness category to indicate whether or not they were present during a time interval. Measurements were also coded by stage of illness as acutely unwell (defined clinically), convalescent (14 days after being acutely unwell), and well.

The study was approved by the joint Medical Research Council and Gambian government ethics committee.

Table  Number of episodes of illness for each infant. Total number of days that the infant was ill is shown in parentheses

<table>
<thead>
<tr>
<th>Illness</th>
<th>Case No</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>1</td>
</tr>
<tr>
<td>Upper respiratory tract infection</td>
<td>6 (31)</td>
</tr>
<tr>
<td>Tonsillitis</td>
<td>1 (5)</td>
</tr>
<tr>
<td>Otitis media</td>
<td>2 (11)</td>
</tr>
<tr>
<td>Pneumonia</td>
<td>2 (9)</td>
</tr>
<tr>
<td>Diarrhoea alone</td>
<td>3 (17)</td>
</tr>
<tr>
<td>Diarrhoea and vomiting</td>
<td>0</td>
</tr>
<tr>
<td>Vomiting alone</td>
<td>0</td>
</tr>
<tr>
<td>Malaria</td>
<td>1 (3)</td>
</tr>
<tr>
<td>Abscess</td>
<td>1 (6)</td>
</tr>
<tr>
<td>Urinary tract infection</td>
<td>0</td>
</tr>
<tr>
<td>Fever</td>
<td>2 (9)</td>
</tr>
<tr>
<td>All illnesses</td>
<td>18 (91)</td>
</tr>
</tbody>
</table>
Identification of factors affecting infant growth in developing countries 1561

Results

The illnesses from which the infants suffered were divided into 11 diagnostic groups (table). These patterns of morbidity, which are extreme, produced a mean (SD) period of acute illness of 75 (29) days/infant. The mean duration of an episode of illness was 5-7 (2-4) days.

The infants had all five anthropometric measurements made on a mean of 30 (6) occasions, with weight alone being measured on a further 6 (2) occasions. The mean interval between weights used for the calculation of weight gain was 10 (7) days.

Falls in weight gain were respectively: 33 g/day (p<0.01), 32 g/day (p<0.01), and 34 g/day (p<0.05). Despite the significance of the difference from when they were well, they were not significantly different from one another. The growth rate increments for convalescence and 'well', expressed as positive increments compared with when ill, were 20 g/day (p<0.001) and 16 g/day (p<0.01), respectively. Again, despite their significant differences from when ill, they were not significantly different from each other.

The SD score graphs for the individual infants are shown in figs 1–7. The effect of illness on growth is clearly seen in the graphs, as with case 1 at 30 weeks or case 6 at 39 weeks. The effects of infection are reflected in all measurements, even length. With

Fig 1  SD scores for case 1. OM=otitis media, URTI=upper respiratory tract infection; 1=length, 2=subscapular skinfold thickness, 3=weight, 4=triceps skinfold thickness, and 5=mid upper arm circumference.

Fig 2  SD scores for case 2. OM=otitis media, URTI=upper respiratory tract infection; 1=length, 2=subscapular skinfold thickness, 3=weight, 4=triceps skinfold thickness, and 5=mid upper arm circumference.
Eccles, Cole, and Whitehead

the exception of length and mid upper arm circumference all the measurements are significantly correlated with each other (p<0.001). It is noticeable that the illnesses producing the largest deviations are those identified by regression analysis as causing significant falls in weight gain. With the exception of case 5, growth in length is maintained while the other measurements, particularly weight and mid upper arm circumference, fall consistently, showing that the infants are wasting. Case 5 shows a degree of stunting. With the exception of cases 2 and 3, the graphs also show that infection produces modulations on an underlying trend. There is an initial variable rise (pronounced in cases 5 and 7) with a universal subsequent fall with most of the values finishing up at -1.5 to -2 SD.

Discussion

The results of this study confirm the growth faltering effects of gastroenteritic and lower respiratory tract disease in infants in third world countries. This is shown qualitatively for individual infants by the multiple measurement standard deviation score graphs, and quantitatively for all infants by the regression analysis.

The multiple measurement SD score graphs allow the whole anthropometric progress of an infant to be viewed at once. The significant effects of gastroenteritis and pneumonia are easily seen on the graphs, as are the growth faltering effects of other episodes of illness.

It is important to understand the justification for using the Cambridge growth curves in a West African population. The Cambridge data set has all its measurements drawn from the same infants, giving it a cohesion not present in other data sets and allowing 'across measurement' comparisons to be soundly based. The LMS method provides (uniquely) SD scores for skewed variables like

Figure 3 SD scores for case 3. OM=otitis media, URTI=upper respiratory tract infection; 1=length, 2=subscapular skinfold thickness, 3=weight, 4=triceps skinfold thickness, and 5=mid upper arm circumference.

Figure 4 SD scores for case 4. URTI=upper respiratory tract infection; 1=length, 2=subscapular skinfold thickness, 3=weight, 4=triceps skinfold thickness, and 5=mid upper arm circumference.
Identification of factors affecting infant growth in developing countries 1563

The size of the weight deficit over infancy is calculable. Using the figures for 'convalescence/well' it is possible to calculate the mean weight deficit in the seven infants caused by acute infection, and thus the weight deficit from other causes. The infants were acutely unwell for 20% of the time, convalescent for 34% of the time, and well for 46% of the time. The deficit for acute illness is: 16 g/day × 0.2 = 3.2 g/day. Had the infants never been acutely unwell they would never have been convalescent so this figure of 3.2 g/day has to have subtracted from it the difference between being well and being convalescent: (20–16) × 0.34 = 1.4 g/day. This produces an overall figure of 1.8 g/day weight deficit as a result of infection, which produces a total of almost 700 g over the first year of life.

The mean weight deficit at 12 months, measured from the Cambridge growth curve 50th centile for weight and skinfold thickness, and it is this that allows the construction of the SD score graphs. Furthermore, the growth curves are based upon infants who were fed in the manner currently recommended by the DHSS.19 This permits prolonged breast feeding and more closely resembles the feeding patterns of the Keneba infants. Given that infant feeding patterns affect growth it is sensible to use growth curves that are as appropriate as possible in this respect.11

In all the infants the graphs identify not only the effects of illness but also a steady fall in the SD score for weight from about 15 weeks of age. As all the infants except case 5 maintained their growth in length, this represents wasting. With his steady fall off in both weight and length, case 5 shows evidence of stunting, a feature of growth that is, in Keneba, more normally seen after infancy.
When comparing Keneba and Bakau, Tomkins et al showed seasonal swings in growth rates in both communities. These swings were larger in Keneba where seasonal food shortage, as opposed to Bakau’s year round availability of food, was thought to account for much of the difference in growth performance between the two communities. Although food intake was not measured in this study, previous work in Keneba has shown infant energy intake to be low and an underlying deficiency in energy intake provides a rational explanation for the steady wasting seen in the SD score graphs.

The illnesses producing an impact on growth are similar to those identified by previous authors. Diagnostic criteria vary, but the diagnoses of diarrhoea, and diarrhoea and vomiting, in this study can be equated with diarrhoea or gastroenteritis from other studies. Previous work in Keneba identified diarrhoeal disease as an important cause of weight loss with a fall in weight gain of 25 g/day. This is similar to the figure of 34 g/day found in this study and shows that the cost in terms of weight gain associated with infant diarrhoea in this rural community has unfortunately not changed over the last 10 years. It is also worth noting that the study of Rowland et al and this study both identified lower respiratory tract infection as an important cause of failure of growth.

SD score graphs provide a powerful method of identifying and highlighting problems with growth resulting from, in this case, the effects of infection and undernutrition. This has not previously been possible because of the lack of an adequate multi-measurement standard. These effects are statistically calculable within the group, but when considering an individual subject the use of SD score graphs permits both the time scale and the accurate progress of an individual subject to be clearly seen. Moreover the curves permit simultaneous comparison across five different anthropometric measurements and the demonstration of the effects of those illnesses that cause growth faltering in an individual subject but do not achieve significance within the group. Although applied here to the acute and chronic problems of infants in the third world, these graphs could equally well be used to record the anthropometric progress of any child suffering from any acute (for example, infective) or chronic (for example, coeliac disease) condition.

We are grateful to the families and particularly the mothers of our seven subjects who were immensely helpful and understanding during the course of the study. We thank M Jarjue for technical assistance. We also acknowledge the support of the Medical Research Council Laboratories at Fajara, and The Gambian Government.

Fig 7  SD scores for case 7. OM = otitis media, URTI = upper respiratory tract infection; 1 = length, 2 = subscapular skinfold thickness, 3 = weight, 4 = triceps skinfold thickness, and 5 = mid upper arm circumference.
References


Correspondence to Dr MP Eccles, Prospect House Medical Group, Prospect House, Prospect Place, Newcastle Upon Tyne, NE4 6QD.

Accepted 1 June 1989
Identification of factors affecting infant growth in developing countries.
M P Eccles, T J Cole and R G Whitehead

Arch Dis Child 1989 64: 1559-1565
doi: 10.1136/adc.64.11.1559

Updated information and services can be found at:
http://adc.bmj.com/content/64/11/1559

These include:
Email alerting service
Receive free email alerts when new articles cite this article. Sign up in the box at the top right corner of the online article.

Notes

To request permissions go to:
http://group.bmj.com/group/rights-licensing/permissions

To order reprints go to:
http://journals.bmj.com/cgi/reprintform

To subscribe to BMJ go to:
http://group.bmj.com/subscribe/