

## Quality control of ultrasound for fetal biometry: results from the INTERGROWTH-21<sup>st</sup> Project

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## ABSTRACT

**Objectives:** To assess a comprehensive package of ultrasound quality control in a large multicentre study of fetal growth – the Fetal Growth Longitudinal Study of the INTERGROWTH-21<sup>st</sup> Project.

**Methods:** We performed quality control (QC) measures on 20,313 ultrasound scan images taken prospectively from 4,321 fetuses at 14-41 weeks' gestation in eight geographical locations. At the time of each ultrasound examination, three fetal biometric variables were measured in triplicate on separately generated images: head circumference (HC), abdominal circumference (AC) and femur length (FL). All measurements were taken in a blinded fashion. QC had two elements: 1) qualitative QC: visual assessment by sonographers at each study site of their images based on specific criteria with 10% of images being re-assessed at the Oxford-based Ultrasound Quality Unit (compared using an adjusted kappa statistic), and 2) quantitative QC: measurement data were assessed by (a) comparing the first, second and third measurement (intraobserver variability); (b) re-measurement of caliper replacement in 10% (interobserver variability), both by Bland-Altman plots, and (c) plotting frequency histograms of the SDs of triplicate measurements and assessing how many were above or below 2SDs of the expected distribution. The system allowed the sonographers' performance to be regularly monitored.

**Results:** A high level of agreement between the self- and external scoring was demonstrated for all measurements (kappa = 0.99 [95% confidence interval: 0.98, 0.99] for HC, 0.98 [0.97, 0.99] for AC, and 0.96 [0.95, 0.98] for FL. Intraobserver variability (95% limits of agreement (LoA)) of ultrasound measures for HC, AC and FL were  $\pm 3\%$ ,  $\pm 6\%$  and  $\pm 6\%$ , respectively; the corresponding values for interobserver variability were  $\pm 4\%$ ,  $\pm 6\%$  and  $\pm 6\%$ . The SD distribution of triplicate measurements for all biometric

variables showed excessive variability for three of 31 sonographers, allowing prompt identification and retraining.

**Conclusions:** Qualitative and quantitative QC monitoring was feasible and highly reproducible in a large multicentre research study, which facilitated the production of high-quality ultrasound images. We recommend that the QC system we developed is implemented in future research studies and clinical practice.

## INTRODUCTION

Standardisation and quality control (QC) of fetal ultrasound biometry are essential to ensure high levels of reproducibility among operators and ultrasound facilities. This particularly applies to multicentre studies because reproducibility and measurement consistency – among even well-trained sonographers – improve as a result of introducing QC systems<sup>1</sup>. Unfortunately, however, a common failing in this field is a complete absence of QC systems: for example, in studies designed to create charts for pregnancy dating, fetal and neonatal growth<sup>2-5</sup>.

Although the effects of QC on measurement reproducibility have been demonstrated in research settings, their relevance may be even greater in routine clinical practice because measurement accuracy is critical for detecting abnormal fetal growth patterns, especially in the absence of blinding of measurements to the sonographer. In fact, avoiding false positive findings, with their attendant anxiety and risks of unnecessary interventions<sup>6</sup>, is almost as important in antenatal care as diagnostic failures.

For a QC system in fetal biometry to be useful clinically, multiple strategies need to be employed<sup>7</sup>, such as:

- (i) Qualitative scoring of ultrasound images against predefined criteria<sup>8</sup>;
- (ii) Quantitative assessment of measurements and comparison with their expected distributions as, for example, occurs in fetal nuchal translucency QC<sup>9-13</sup> although, until now, these approaches have largely only been utilised in small studies<sup>8-10, 13, 14</sup>.

Here we describe and assess the value of the comprehensive QC package used in the Fetal Growth Longitudinal Study (FGLS) of the INTERGROWTH-21<sup>st</sup> Project.

## METHODS

Women at low risk of adverse pregnancy outcomes were recruited into FGLS, one of the three main components of INTERGROWTH-21<sup>st</sup> ([www.intergrowth21.org.uk](http://www.intergrowth21.org.uk)), a multicentre, multi-country, population-based project, conducted between 2008 and 2014 in eight countries <sup>5, 15, 16</sup>, which aimed to construct international fetal growth standards. Serial ultrasound scans were performed every 5±1 weeks from 14+0 to 41+6 weeks' gestation. Gestational age was calculated on the basis of the last menstrual period (LMP) provided that: a) it was known and certain; b) the menstrual cycles were regular; c) there was no hormonal contraceptive use or breastfeeding during the 2 months prior to natural conception, and d) standardised <sup>17</sup> ultrasound measurement of the fetal crown-rump length between 9+0 and 13+6 weeks' gestation agreed with the LMP-based estimate of gestational age within 7 days <sup>18</sup>.

At each examination, three fetal biometric variables were measured in triplicate on separately generated two-dimensional ultrasound images: head circumference (HC), abdominal circumference (AC) and femur length (FL). Thus, each examination produced nine measurements (three per variable) in accordance with the study protocol ([www.intergrowth21.org.uk](http://www.intergrowth21.org.uk)) <sup>19</sup>.

All sonographers were recruited on the basis of being motivated, reliable and trained in ultrasound; ability to speak the local language(s) and work positively within a team structure. The goals of standardisation were, firstly to ensure that all sonographers fully understood the study protocol and take measurements in an identical fashion, and secondly that they were familiar with the equipment used. The precise details of how measurements were taken for FGLS and how data collection was standardised (through training, assessment and certification of all the sonographers) are presented in full elsewhere <sup>7, 19</sup>. Head measurements were obtained in the transthalamic plane, placing the calipers on the outer border of the skull, using both the ellipse facility and

two perpendicular diameters. Abdominal measurements were obtained in an axial plane, with the umbilical vein in the anterior third of the fetal abdomen (at the level of the portal sinus) and the stomach bubble visible. Again, both the ellipse facility and the two diameters method were used, placing the calipers on the outer border of the body outline (skin covering). In this study, we elected to analyse only the HC and AC measurements obtained using the ellipse facility, as a previous study showed that these were almost identical to those using the two diameters, but marginally more reproducible<sup>20</sup>. For FL, the femur closest to the probe was measured, with its long axis as horizontal as possible. Calipers were placed on the outer borders of the diaphysis of the femoral bone ('outer to outer').

All ultrasound scans were performed using the same commercially available ultrasound machine (Philips HD-9, Philips Ultrasound, Bothell, WA, USA) with curvilinear abdominal transducers (C5-2, C6-3 and V7-3). During the INTERGROWTH-21<sup>st</sup> Project blinding of operators to the measurement value was undertaken, thus eliminating expected value bias. For this purpose, the manufacturer programmed the machine's software so that the measurement values did not appear on screen during a scan.

The QC strategies adopted, which are described in detail below, included qualitative (i.e. image scoring) and quantitative analyses (i.e. estimating intraobserver and interobserver variability, and standard deviations (SD) of triplicate measures) for each biometric variable. Six sonographers undertook QC at the Oxford-based Ultrasound Quality Unit (USQU); any uncertainties were adjudicated by the QC Director (ATP). The analyses were performed monthly for the first 18 months of each site's participation and quarterly thereafter, or more frequently if any QC concerns were raised so as to identify sonographers performing outside accepted norms to allow corrective action (e.g. retraining) to be administered promptly<sup>7</sup>.

### **Qualitative QC: Image scoring**

Images were scored <sup>7, 8</sup> based on a set of criteria, each worth one point towards the total score, with a maximum of six points for HC and AC, and four points for FL (Table 1). All images were self-scored at the time of scanning by the sonographer taking the image. A randomly chosen sample of 10% of all these images was re-scored by a sonographer at the USQU; the highest of these three scores was used in the QC analysis.

In order to simplify the comparison between self- and USQU scoring, we divided data into low-scoring (1 to 3 for HC and AC, and 1 to 2 for FL) and high-scoring images (4 to 6 for HC and AC, and 3 to 4 for FL) <sup>7</sup>. As the quality was generally very good, higher scores were much more prevalent than lower scores; comparison between self- and USQU scoring was therefore undertaken using an adjusted kappa statistic (interobserver variability of image scoring) to account for the resulting unbalanced distributions of scores <sup>21</sup>. A kappa value of more than 0.6 was considered *a priori* an acceptable level of agreement among sonographers.

### **Quantitative QC: intraobserver and interobserver variation**

As triplicate images and measurements were taken for each fetal biometric variable (HC, AC, FL), the *intraobserver* variability of the measurements could be assessed in the full dataset using Bland-Altman plots <sup>22</sup>. Instead of simply expressing differences within observers in actual measurement units (mm), pairwise comparisons were also made in percentage terms to account for changes in fetal size with increasing gestational age. The difference between two selected measurements was calculated and expressed as a percentage of their mean, then plotted against this mean. The 95% limits of agreement (LoA) were calculated and marked on the plots, giving a quantifiable estimate for the measurement variability within the same observer associated with acquiring an image and positioning the calipers. These plots were

generated by randomly selecting two of the three triplicate measurements taken at each scan for each biometric variable.

Actual (mm) and percentage difference Bland-Altman plots were also used to assess the *interobserver* variability of the measurements. As above, a sonographer at the USQU re-measured a random sample of 10% of all images from each site. The difference between the original and USQU measurements was expressed both as the actual value and as a percentage of their mean, then plotted against this mean. Again, 95% LoA were calculated and marked on the plots, giving a quantifiable estimate for the measurement reproducibility between observers associated with caliper placement.

### **Quantitative QC: data distribution**

The SD of each measurement triplet was expressed as a percentage of the mean of the three measurements, enabling each sonographer's individual variability to be compared with the expected variability<sup>14</sup> whilst accounting for changes in fetal size with increasing gestational age.

Plotting each sonographer's SDs as separate frequency histograms allowed sonographers to be identified whose SD distributions differed from those of the expected range, based on the equivalent data derived from an initial variability study<sup>14</sup>. Sonographers demonstrating disproportionately large numbers of triplets outside the expected variability distribution were identified, causes investigated and retraining undertaken if necessary. Each sonographer's SDs were also plotted sequentially with a cumulative sum control chart<sup>7, 14</sup> to identify triplets with values >2 SDs more than 10% of the time.

All QC performed by USQU sonographers was undertaken blinded to the study site, sonographer identity, original measurement and their own repeated measurements. Unblinding only occurred to provide feedback to sonographers where necessary. All

plots were generated and analyses performed using SAS software (Copyright, SAS Institute Inc. SAS and all other SAS Institute Inc. product or service names are registered trademarks or trademarks of SAS Institute Inc., Cary, NC, USA).

The INTERGROWTH-21<sup>st</sup> Project protocol was approved by the Oxfordshire Research Ethics Committee C (reference: 08/H0606/139); all the pregnant women enrolled gave informed written consent.

## RESULTS

We studied 4,321 singleton fetuses, whose intrauterine biometric measures were used to construct international fetal growth standards, all of whom were born alive without congenital malformations<sup>18</sup>. Each fetus had a median of 5 (range 1-7; mean 4.9 [SD 0.8]; total 20,313) scans. Figure 1 shows the total number of scans eligible for QC for each biometric variable (HC, AC and FL) after excluding measures >5 SD's and missing data.

Between 20,040 and 20,313 scans, depending on the biometric variable, were assessed for intraobserver variability (Figure 1, Box 1). Ten per cent (n=1,735) of the 17,350 scans performed at sites other than Oxford were randomly selected for external image re-scoring and re-measurement by a USQU sonographer. Of these, 122 scans could not be assessed (due to incomplete backup of images or data lost due to corruption of the backup file) leaving 1,613 scans from 1,322 women that underwent QC by USQU sonographers (Figure 1).

### Qualitative QC: Image scoring

Of the 1,613 scans, 1,340 (83.1%) were re-scored; the remainder were missing original image scores (n=256) or re-scoring were not logged (n=17). Overall, the quality of all measurements was high. The median self-scored image values for HC, AC and FL were 6 [interquartile range (IQR): 6-6], 6 [IQR: 6-6] and 4 [IQR: 4-4], respectively – the maximum values in the scoring system. There was a very high level of agreement between the self- and external scoring of image quality for all measurements, with adjusted kappa values of 0.99 [95% confidence interval (CI): 0.98, 0.99] for HC, 0.98 [95% CI: 0.97, 0.99] for AC, and 0.96 [95% CI: 0.95, 0.98] for FL (Table 2). In almost all cases, both the local and USQU sonographers classified the same image as high-scoring (99% scored 4 to 6 for HC and AC, 98% scored 3 to 4 for FL).

This external image assessment process resulted in six sonographers requiring retraining over the entire study period, after which improvements in performance were seen.

#### **Quantitative QC: intraobserver and interobserver variation**

Intraobserver variability was assessed using all 20,313 scans, comparing two of the triplicate measurements for each scan, selected randomly and in random order. Overall, the reproducibility was very good (Supplementary figure 1). For HC, the mean difference was 0.0% and the 95% LoA were consistently between  $\pm 3.1$  and  $\pm 3.5\%$ . For AC, the mean difference was approximately 0.1% and the 95% LoA were consistently between  $\pm 5.4$  and  $\pm 6.0\%$ . For FL, the mean difference was approximately 0.2%; however, the 95% LoA, even when expressed as a percentage of FL, varied with gestation, showing greater variability and poorer intraobserver reproducibility at lower gestational ages. On average, the 95% LoA for FL were between  $\pm 5.8$  and  $\pm 6.4\%$  (Table 3).

Interobserver variability was assessed using 1,483 of the 1,613 scans selected in the 10% QC sample as 130 re-measured scans were erased as a result of a technical problem. Overall, the reproducibility was very good (Supplementary figure 2). For HC, AC and FL the mean difference was 1.0% or less, with 95% LoA between  $\pm 4.3$  and  $\pm 4.4\%$ , between  $\pm 5.9$  and  $\pm 6.3\%$  and between  $\pm 5.4$  and  $\pm 5.9\%$ , respectively (Table 3).

#### **Quantitative QC: data distribution**

Comparisons with the expected distribution of measures showed no cause for concern in any biometric variable for 28 out of the 31 study sonographers. In one instance, a sonographer was found to have 16.5% of HC SDs and 11.1% of AC SDs outside the expected range, whilst two other sonographers demonstrated unacceptably high FL SDs (Figure 2). In all three instances retraining was undertaken and improvements

were seen thereafter. The total number of images taken by these three sonographers made up only a very small proportion of the total dataset [HC: 188 of 20,041 (0.9%); AC: 45 of 20,135 (0.2%); FL: 267 of 20,313 (1.3%)].

## DISCUSSION

We report the implementation and results of using a comprehensive system to assess the quality of ultrasound images obtained from a large multicentre, international project. We not only demonstrate the system's feasibility, but show that it is possible to achieve a high level of reproducibility in such a study with the necessary QC measures.

Firstly, over 98% of the scored images were considered as high quality by both the local and USQU sonographers (qualitative QC); secondly, the intra- and interobserver reproducibility of measurements (quantitative QC) was high and within the limits of a previous study<sup>14</sup> (Table 3); and thirdly, we monitored images and data regularly, which enabled us to identify a few sonographers whose performance fell outside expected standards, following which corrective action was taken. It should be noted that this entire process relied upon initial training and standardisation<sup>1, 7, 19</sup> – a crucial element of the project's success.

Meticulous standardisation and ongoing monitoring of adherence to measurement protocols during data collection have been shown to ensure consistency and minimise systematic error in multicentre studies<sup>1, 7, 11, 17</sup>. In two recent systematic reviews of the literature relating to the creation of fetal crown-rump length charts and growth charts, no studies reporting a comprehensive QC process were identified, which undoubtedly contributed to the poor quality of many existing studies<sup>2, 3</sup>.

Our study has a number of strengths: a) the QC strategy was prospectively designed and implemented<sup>7, 15, 19</sup>, and based firmly on previous studies that assessed the role of feedback on image quality<sup>7, 10, 23, 24</sup>, and b) visual assessment of ultrasound images was based on an objective criterion-based scoring system, which has been shown to be significantly more reproducible than subjective methods<sup>8, 25, 26</sup>. In Salomon et al.'s<sup>8</sup> original description of this process, high reproducibility levels for the image scoring

method were demonstrated (kappa between 0.60 and 0.98); despite undertaking QC in a blinded fashion, the results from our study were even better. The high level of reproducibility of such objective methods<sup>9, 12, 25, 26</sup> are corroborated by our study, which is the largest to date. It is likely that the high level of training of the sonographers acquiring the images and those conducting the QC, and the requirement for standardisation of all staff in settings of near-optimal conditions for scanning, contributed to the overall quality.

One of the limitations of the study is that only 10% of images underwent external scoring. However, this is the largest quality control programme ever performed in the setting of a study into fetal growth. All 100% images underwent self-scoring, and those images that were externally scored were randomly selected, meaning that there was no evidence that a different proportion would have yielded different results. Of course, implementing such a QC strategy is labour-intensive. While it is relatively easy to assess data distributions routinely<sup>27</sup>, external qualitative assessment using image scoring requires additional resources.

More cost-effective options might include:

- Voluntary submission of a small number of selected images (as, for example, in certification for nuchal translucency measurement)<sup>28</sup>; however, the small number of images and the nature of self-selection mean it is difficult to ascertain whether such images are truly representative of a sonographer's routine practice.
- Self-assessment of images which correlates well with external scoring using a 10% random sample of all images suggesting that it may be a reasonable alternative. However, there are three reasons to be cautious: firstly, we have demonstrated that self-scoring is effective *in association with* external scrutiny, and it is not known whether similarly high quality is achievable without a QC

system; secondly, such a system is feasible with a few highly trained and motivated sonographers and may not be scalable, for example, to a national screening programme; and thirdly, while we have demonstrated excellent agreement between self- and external scoring across the whole dataset, the role of QC in screening is exception reporting, i.e. to detect individual outliers, rather than to demonstrate that, *on average*, the system works. Only by integrating all the elements of our QC system were we able to identify opportunities for improvement that could not be detected by self-scoring alone.

- Automated methods for QC of routinely collected images are being studied and may, in the future, be the best option. These systems have the potential advantage of allowing all images to be assessed objectively and at low cost<sup>29</sup>,

<sup>30</sup>.

Regarding quantitative QC, a literature search was performed to identify previous publications on the evaluation of reproducibility of fetal ultrasound biometry after 14+0 weeks (17 studies identified)<sup>1, 14, 27, 31-45</sup>. Studies were selected only if reliable quantitative values were calculated as LoA or repeatability coefficients<sup>46, 47</sup>. Overall intraobserver reproducibility reported 95% LoA of less than 4% (12 mm) for HC, 6% (12 mm) for AC, and 7% (3 mm) for FL<sup>1, 14, 32-37, 40, 41, 43, 45</sup>. Similarly, for interobserver analyses 95% LoA for HC, AC and FL were within 4%, 6%, and 6%, respectively<sup>1, 14, 27, 31, 34-39, 42, 44, 45</sup>. Even though these studies were undertaken on smaller numbers of cases, mostly in single centre research settings and without blinding of the measurements, these values are not markedly different from the results of our large-scale multicentre study.

Our study has shown that, in general, both intra- and interobserver variability remain reasonably constant throughout pregnancy when reported as a percentage of fetal size. The exception is for FL, where increased variance was demonstrated at early

gestational ages, most likely due to the difficulty in accurately measuring FL when it is only 10-30 mm long.

In conclusion, both qualitative and quantitative QC monitoring were found to be feasible in a large multicentre fetal growth study. The development of a standardised fetal biometric ultrasound measurement protocol, standardisation of all sonographers (involving their training, assessment and certification), consistency and blinding of measurement are all necessary to minimise systematic error and ensure high reproducibility. Having developed a framework for ultrasound QC, we recommend that it is implemented in future similar research studies and, ideally, in clinical practice.

**Conflict of interest:** ATP is the Chief Medical Officer of Intelligent Ultrasound and receives non-financial support from Philips Ultrasound. JAN has received personal fees from Intelligent Ultrasound and grants and non-financial support from Philips Ultrasound. All other authors declare no competing interests.

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## TABLES

**Table 1** - Image scoring criteria used for standardisation and quality control.

| <b>Cephalic plane<br/>(max. 6 points)</b> | <b>Abdominal plane<br/>(max. 6 points)</b> | <b>Femoral plane<br/>(max. 4 points)</b> |
|-------------------------------------------|--------------------------------------------|------------------------------------------|
| 1 Symmetrical plane                       | 1 Symmetrical plane                        | 1 Both ends of the bone clearly visible  |
| 2 Thalami visible                         | 2 Stomach bubble visible                   | 2 Angle <45°                             |
| 3 Cavum septi pellucidi visible           | 3 Portal sinus visible                     | 3 Femur occupying at least 30% of image  |
| 4 Cerebellum not visible                  | 4 Kidneys not visible                      | 4 Callipers placed correctly             |
| 5 Head occupying at least 30% of image    | 5 Abdomen occupying at least 30% of image  | -                                        |
| 6 Callipers/ellipse placed correctly      | 6 Callipers/ellipse placed correctly       | -                                        |

**Table 2** - Matrix showing the number of scans that were self-scored (rows) versus external scoring results (columns) for head circumference, abdominal circumference and femur length, and adjusted kappa values.

|                             |                            |       | Quality score: External scoring |    |     |      |       | Adjusted kappa<br>(95% CI) |
|-----------------------------|----------------------------|-------|---------------------------------|----|-----|------|-------|----------------------------|
|                             |                            |       | 1 - 3                           | 4  | 5   | 6    | Total |                            |
| Quality score: Self scoring | Head<br>circumference      | 1 - 3 | 1                               | 1  | 4   | 2    | 8     | 0.99<br>(0.98 to 0.99)     |
|                             |                            | 4     | 0                               | 3  | 6   | 23   | 32    |                            |
|                             |                            | 5     | 1                               | 4  | 26  | 182  | 213   |                            |
|                             |                            | 6     | 1                               | 15 | 137 | 917  | 1070  |                            |
|                             |                            | Total | 3                               | 23 | 173 | 1124 | 1323  |                            |
|                             |                            |       |                                 |    |     |      |       |                            |
|                             | Abdominal<br>circumference | 1 - 3 | 0                               | 0  | 0   | 7    | 7     | 0.98<br>(0.97 to 0.99)     |
|                             |                            | 4     | 1                               | 3  | 6   | 36   | 46    |                            |
|                             |                            | 5     | 1                               | 2  | 20  | 234  | 257   |                            |
|                             |                            | 6     | 1                               | 10 | 90  | 906  | 1007  |                            |
|                             |                            | Total | 3                               | 15 | 116 | 1183 | 1317  |                            |
|                             |                            |       |                                 |    |     |      |       |                            |
|                             |                            |       | 1                               | 2  | 3   | 4    |       |                            |
|                             | Femur length               | 1     | 0                               | 0  | 0   | 0    | 0     | 0.96<br>(0.95 to 0.98)     |
|                             |                            | 2     | 0                               | 0  | 2   | 0    | 2     |                            |
|                             |                            | 3     | 0                               | 1  | 7   | 19   | 27    |                            |
|                             |                            | 4     | 0                               | 21 | 136 | 1145 | 1302  |                            |
|                             |                            | Total | 0                               | 22 | 145 | 1164 | 1331  |                            |

**Table 3** - Quantitative QC: intraobserver variability for image acquisition and caliper placement and interobserver variability for caliper replacement (10% of all images), expressed as percentages (%). The relevant data from a previous study<sup>14</sup> are included here for comparison. QC: quality control; HC: head circumference; AC: abdominal circumference; FL: femur length.

|                                                                               |    | Quantitative QC |                         |                 |                         |
|-------------------------------------------------------------------------------|----|-----------------|-------------------------|-----------------|-------------------------|
|                                                                               |    | Pilot study     |                         | Our study       |                         |
|                                                                               |    | Mean difference | 95% limits of agreement | Mean difference | 95% limits of agreement |
| Intraobserver reproducibility (%) for image acquisition and caliper placement | HC | 0.0             | ± 3.0                   | 0.0             | ± 3.3                   |
|                                                                               | AC | - 0.3           | ± 5.3                   | 0.0             | ± 5.6                   |
|                                                                               | FL | 0.2             | ± 6.6                   | 0.0             | ± 6.2                   |
| Interobserver reproducibility (%) for caliper replacement (10% of all images) | HC | 1.3             | ± 3.7                   | 1.0             | ± 4.4                   |
|                                                                               | AC | 1.1             | ± 5.7                   | - 0.1           | ± 6.0                   |
|                                                                               | FL | 0.8             | ± 5.8                   | - 0.8           | ± 5.6                   |

## FIGURE LEGENDS

**Figure 1** - Flow chart of patients and scans included in the analysis. QC: quality control; HC: head circumference; AC: abdominal circumference; FL: femur length.

**Figure 2** - Distribution of standard deviations (SD) (expressed as a percentage of the mean) of triplicate measurements of head circumference made by two sonographers. The vertical lines indicate the 97.5<sup>th</sup> centile value (2.42), median and 2.5<sup>th</sup> centile (0.16) taken from the reference standard study <sup>7</sup>. We illustrate two examples: in the upper panel 3% of triplicate measurements are above the 97.5<sup>th</sup> centile. In the lower panel 17% of measurements were above the accepted threshold, set at 10%; and retraining was undertaken.

**Supplementary figure 1** - Bland Altman plots: intraobserver variability in head circumference (a,b), abdominal circumference (c,d) and femur length (e,f) measurements, expressed as millimetres (mm) (a,c,e) and percentage (%) (b,d,f).

**Supplementary figure 2** - Bland Altman plots: interobserver variability in head circumference (a,b), abdominal circumference (c,d) and femur length (e,f) measurements, expressed as millimetres (mm) (a,c,e) and percentage (%) (b,d,f).