

## ORIGINAL ARTICLE

## A comparison of transcutaneous bilirubinometers: SpectRx BiliCheck versus Minolta AirShields

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Neonatal hyperbilirubinaemia is a common phenomenon but relatively few affected infants require intervention. Whatever the cause, total serum bilirubin (SBR) levels above defined thresholds warrant treatment to prevent the development of kernicterus.

The clinical evaluation of hyperbilirubinaemia involves visual estimation of the yellowness of skin, known as jaundice. However, quantification of SBR based on visual assessment of the depth of jaundice is subjective and inaccurate, and confounded by skin colour and haemoglobin.<sup>1</sup> Various methods have therefore been developed to aid non-invasive diagnosis of hyperbilirubinaemia and approximation of SBR levels. They include cephalocaudal staging of jaundice<sup>2,3</sup> and transcutaneous reference devices. However, because of limited accuracy, these transcutaneous devices have only been useful as screening rather than diagnostic tools for determining hyperbilirubinaemia. They include the Ingram Ictermeter<sup>2,4</sup> Chromatics Colormate III,<sup>5</sup> and Minolta AirShields Jaundice Meter.<sup>4,6</sup> A new device, the SpectRx BiliCheck, is claimed to have high accuracy and reproducibility across different ethnic groups and gestational ages, although the authors do state that further assessment of efficacy is needed for SBR  $\geq 256 \mu\text{mol/l}$ .<sup>7</sup> It has also been suggested that the SpectRx device is accurate enough to be a substitute for SBR sampling.<sup>8</sup>

Two devices are available for making transcutaneous estimates of SBR, the Minolta AirShields JM102 and the new SpectRx BiliCheck. The aims of our study were to (a) measure how well the readings produced by these devices agree with SBR measured in the laboratory, and (b) estimate for each device the proportion of infants with clinical jaundice who would require blood sampling if the device was used as a screening tool to detect infants with SBR  $\geq 250 \mu\text{mol/l}$ . This threshold was used to define significant hyperbilirubinaemia, as most infants affected would require either treatment or further investigation of their hyperbilirubinaemia, and hence

**Background:** Two devices are available for making transcutaneous estimates of serum bilirubin (SBR): the Minolta AirShields JM102 and the new SpectRx BiliCheck.

**Objectives:** (a) To measure how well the readings produced by these devices agree with SBR measured in the laboratory; (b) to estimate for each device, the proportion of infants with clinical jaundice who would require blood sampling if the device was used as a screening tool to detect infants with SBR  $\geq 250 \mu\text{mol/l}$ .

**Design:** Prospective cohort study of jaundiced infants who required SBR at  $\leq 20$  days of postnatal age. Those who had received phototherapy or exchange transfusion were excluded.

**Setting:** Tertiary neonatal service in South-East Scotland.

**Interventions:** Within 30 minutes of SBR sampling, transcutaneous bilirubinometry was performed using one Minolta and two SpectRx devices (designated A and B).

**Results:** Sixty four neonates were enrolled, 19 of which were preterm (31–35 weeks). The 95% confidence intervals of a device reading corresponding to SBR were  $\pm 66.7$ ,  $\pm 67.9$ , and  $\pm 66.4 \mu\text{mol/l}$  respectively. Using the devices to identify all SBR  $\geq 250 \mu\text{mol/l}$  would reduce SBR sampling by 23%, 16%, and 20% respectively.

**Conclusions:** Given that SBR levels range from 50 to 400  $\mu\text{mol/l}$  in jaundiced infants, the 95% confidence intervals of the devices are wide at  $\pm 67 \mu\text{mol/l}$ . The SpectRx can be used as a screening tool for hyperbilirubinaemia but there is no advantage in using it over the Minolta.

would be need to be identified.<sup>9</sup> This is with the exception of clinical jaundice in the first 24 hours, which warrants investigation whatever the SBR level.<sup>9</sup> We also assessed the intra-device precision of the Minolta and SpectRx devices. Inter-device precision between two SpectRx devices was evaluated.

## METHODS

Neonates at the Simpson Memorial Maternity Pavilion, Edinburgh, who required blood sampling for SBR were enrolled in the study with parental consent. All neonates were clinically jaundiced but otherwise well. Three bilirubinometers (see below) were applied to the infant's forehead within 30 minutes of SBR sampling by venepuncture or heel lance. All transcutaneous bilirubinometry (TcB) measurements were performed by one author (PvD) with the infant lying supine. The forehead was not exposed to direct sunlight and care was taken to avoid skin areas that were bruised, excessively hairy, or hypermelanotic. Gestation, birth weight, postnatal age, and ethnicity were recorded. Infants who had received phototherapy or exchange transfusion were excluded.

SBR samples were sent immediately for analysis by an automated Hitachi 911 multichannel analyser. The method used involves oxidation of bilirubin to biliverdin by potassium ferricyanide in the presence of caffeine.<sup>10</sup> This technique is reliable even in the presence of haemolysis up to 5.0 g/l and is therefore particularly suitable for neonatal samples. Inter-batch quality control coefficients of variation are 3.0% at 53  $\mu\text{mol/l}$ , 1.4% at 124  $\mu\text{mol/l}$ , and 1.6% at 231  $\mu\text{mol/l}$ . The laboratory participates in the United Kingdom External Quality Assessment Scheme, and shows a mean + 2.4% bias for SBR analysis.

**Abbreviations:** SBR, serum bilirubin; TcB, transcutaneous bilirubinometry; HPLC, high performance liquid chromatography

**Table 1** Details of patients enrolled in the study

	Mean	SD	Range
All infants (n=64)			
Gestation (weeks)	37.4	3.0	31–42
Birth weight (g)	2920.8	755.5	3000–4610
SBR ( $\mu\text{mol/l}$ )	207.3	68.8	50–349
Postnatal age (days)	4.6	3.4	1–17
Term infants (n=45)			
Gestation (weeks)	39.1	1.4	37–42
Birth weight (g)	3258.9	605.4	1750–4610
SBR ( $\mu\text{mol/l}$ )	212.1	72.5	50–349
Postnatal age (days)	3.6	2.5	1–14
Preterm infants (n=19)			
Gestation (weeks)	33.4	1.2	31–35
Birth weight (g)	2120.0	373.1	1610–3120
SBR ( $\mu\text{mol/l}$ )	195.9	59.4	85–307
Postnatal age (days)	7.0	4.0	2–17

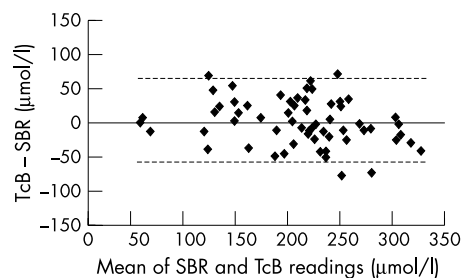
SBR, Serum bilirubin.

### Bilirubinometer specifications and method of use

The devices compared were two SpectRx BiliCheck (SpectRx, USA), designated A and B, and one Minolta AirShields JM102 (now Hill-Rom AirShields International). Both systems rely on spectrophotometric methods.

Detailed information on the SpectRx device has been given elsewhere.<sup>7</sup> The SpectRx instruction manual and website claim the device to be accurate at gestational age 27–42 weeks, postnatal age 0–20 days, and weight 950–5200 g. It is also claimed to be accurate regardless of skin colour by compensating for the confounding effects of dermal maturity, melanin, and haemoglobin on spectral reflectance. The SpectRx uses disposable probe tips which require calibration before each measurement. The tip is then applied with light pressure to the skin. The SpectRx device is designed to be triggered for five consecutive spectral analyses before it provides one displayed averaged reading in either  $\mu\text{mol/l}$  or  $\text{mg/dl}$ . In other words, it has an internal averaging system in an attempt to improve accuracy. For the study, three of these readings were obtained per device to determine intradevice precision. The average of these readings was used as the operative result in all comparisons with the Minolta, and for calculating interdevice precision.

The Minolta AirShields also measures the spectral reflectance of bilirubin but does not claim to correct for confounding effects. Detail on operation principles is described elsewhere.<sup>6</sup> It has been extensively studied and validated for use as a screening tool for hyperbilirubinaemia in term infants.<sup>4 11–13</sup> There have also been reports in the literature on its use in ethnic minorities with various skin colouring.<sup>11 12 14 15</sup> The Minolta device does not use disposable parts or require repeated calibration. The probe tip is applied to the skin with increasing pressure until a flash is activated and a reading obtained. Readings are produced as a numeric index. The Minolta was used according to the protocol of the study centre, recording three consecutive readings, and choosing the maximum value for clinical decision making.



**Figure 1** Bland-Altman plots for the Minolta device. The dotted line represents 95% confidence intervals of the difference,  $\pm 66.7 \mu\text{mol/l}$ . SBR, Serum bilirubin; TcB, transcutaneous bilirubinometry.

### Data and statistical analysis

For statistical accuracy, each patient was assessed only once. All statistical calculations were performed using Microsoft Excel 97. Pearson correlation coefficients were calculated using original Minolta index readings. For measurement of agreement by the Bland-Altman method,<sup>16</sup> Minolta index readings were transformed into  $\mu\text{mol/l}$  using linear regression of SBR on TcB readings. The transformed values were used for calculation of the differences between SBR and TcB values, the mean difference and 95% confidence intervals of the difference.

### RESULTS

From June to September 2000, 64 neonates were enrolled, 19 of whom were preterm (31–35 weeks). Table 1 gives patient details. All were less than 20 days old. Six infants were non-white. None had a haemoglobinopathy. SBR levels ranged from 50 to 349  $\mu\text{mol/l}$ .

### Agreement between SpectRx and Minolta devices and SBR

Table 2 shows agreement results for TcB readings against SBR. The 95% confidence intervals of a device reading corresponding to SBR were  $\pm 66.7$ ,  $\pm 67.9$ , and  $\pm 66.4 \mu\text{mol/l}$  for the Minolta, SpectRx A, and SpectRx B respectively. The Bland-Altman scatterplots for the devices show a tendency for both Minolta and SpectRx devices to underestimate SBR at higher levels (figs 1–3).

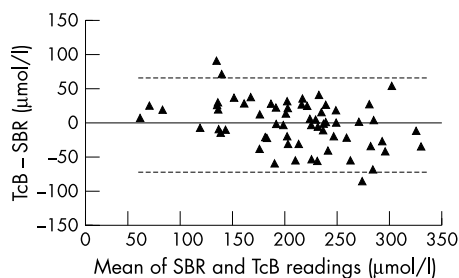
For preterm infants, the Minolta device showed a tendency to overestimate SBR but the device has not been validated for such use. A good correlation with SBR was seen in non-white infants for the Minolta, SpectRx A, and SpectRx B ( $r = 0.94$ ,  $0.95$ , and  $0.99$  respectively), but because of small numbers ( $n = 6$ ), it was deemed inappropriate to calculate statistical agreement.

Accuracy can be determined by agreement, but for practical purposes (such as defining thresholds for performing SBR), the maximum differences between device readings and SBR should be considered. Consequently 100% of infants with hyperbilirubinaemia requiring treatment would be identified. Table 2 also displays the maximum range of differences between devices and SBR. For example, to identify 100% of

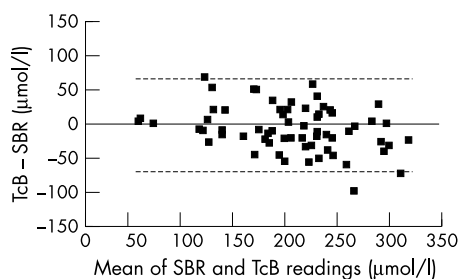
**Table 2** Mean and maximum differences, and 95% confidence intervals of the differences for respective devices

Patients	No	Minolta			SpectRx A			SpectRx B		
		m $\Delta$	95% CI	Max $\Delta$	m $\Delta$	95% CI	Max $\Delta$	m $\Delta$	95% CI	Max $\Delta$
All	64	0.0	$\pm 66.7$	–79 to +69	–4.0	$\pm 67.9$	–88 to +88	–8.6	$\pm 66.4$	–101 to +67
Term	45	–9.6	$\pm 65.1$		–5.5	$\pm 67.2$		–12.8	$\pm 62.9$	
Preterm	19	22.7	$\pm 46.0$		–0.5	$\pm 71.1$		1.3	$\pm 72.0$	

m $\Delta$  ( $\mu\text{mol/l}$ ) = mean difference (device–SBR); 95% CI ( $\mu\text{mol/l}$ ) = 95% confidence intervals of the difference; Max $\Delta$  ( $\mu\text{mol/l}$ ) = maximum range of difference (device–SBR); SBR, serum bilirubin.



**Figure 2** Bland-Altman plots for the SpectRx A device. The dotted line represents 95% confidence intervals of the difference,  $\pm 67.9$   $\mu\text{mol/l}$ . SBR, Serum bilirubin; TcB, transcutaneous bilirubinometry.



**Figure 3** Bland-Altman plots for the SpectRx B device. The dotted line represents 95% confidence intervals of the difference,  $\pm 66.4$   $\mu\text{mol/l}$ . SBR, Serum bilirubin; TcB, transcutaneous bilirubinometry.

infants with  $\text{SBR} \geq 250$   $\mu\text{mol/l}$ , blood samples need to be obtained from all infants whose Minolta reading is  $\geq 170$   $\mu\text{mol/l}$  (index reading 15.5) or whose SpectRx reading is  $\geq 150$   $\mu\text{mol/l}$ .

#### Positive predictive value and number of SBR samples required to identify all $\text{SBR} \geq 250$ $\mu\text{mol/l}$

When the threshold for each device is defined to achieve 100% sensitivity for  $\text{SBR} \geq 250$   $\mu\text{mol/l}$ , the positive predictive values are similar (table 3). Using the devices to screen for this threshold would reduce SBR sampling by 23% for the Minolta device, 16% for SpectRx A, and 20% for SpectRx B. Thus the SpectRx does not result in fewer SBR samples being taken to identify all infants with significant hyperbilirubinaemia.

#### Intradvice precision

The median coefficients of variation for the Minolta, SpectRx A, and SpectRx B were 1.49%, 8.94%, and 8.51% respectively.

Maximum coefficients of variation were 6.74%, 18.67%, and 27.64% respectively.

#### Interdevice precision

The 95% confidence interval of difference in readings for the two SpectRx devices was  $\pm 37.4$   $\mu\text{mol/l}$ .

#### DISCUSSION

This study is the first to compare the performance of the new SpectRx BiliCheck with the Minolta AirShields. Given that SBR levels range from 50 to 400  $\mu\text{mol/l}$  in jaundiced infants, both instruments have wide 95% confidence intervals at  $\pm 67$   $\mu\text{mol/l}$ . A TcB reading that is different from SBR by up to 100  $\mu\text{mol/l}$  (the maximum difference in this study) can have implications for starting phototherapy or exchange transfusion.<sup>9</sup> Such a difference could possibly also have an effect on whether an infant develops kernicterus or not. These devices are useful as screening tools for hyperbilirubinaemia, but not as replacements for measuring SBR.

#### Comparison of SpectRx and Minolta devices

Our study has shown that, as a screening tool, there is no advantage in using the SpectRx device over the Minolta. Although the SpectRx provides readings in  $\mu\text{mol/l}$  for easier interpretation, and claims to be accurate regardless of skin colour, we have found that it is neither more accurate than the Minolta device nor does it reduce the number of samples needed to identify significant hyperbilirubinaemia. Agreement is a better measure of accuracy than correlation, and the agreement between the device readings and SBR is poor.

Positive predictive values are acknowledged to vary with disease prevalence, in this case the number of jaundiced infants deemed to require SBR sampling. Hence our positive predictive values should not be extrapolated to a different population of infants, but nevertheless provide useful data for cross comparisons between the devices. The positive predictive values of the devices were similar for all three devices.

Clinical usefulness of the devices as screening tools was assessed by the reduction in SBR sampling while ensuring that all samples with  $\text{SBR} \geq 250$   $\mu\text{mol/l}$  were identified. Thus for the Minolta, SpectRx A, and SpectRx B devices respectively, 15 (23%), 10 (16%), and 13 (20%) SBR samples could have been avoided. The Minolta, in fact, performed slightly better with respect to reduction of SBR sampling than the SpectRx devices. Again, these data are valuable primarily for comparing the devices with each other.

Whichever screening tool is used, the Bland-Altman scatterplots show a tendency for the devices to underestimate

**Table 3** Specificity and positive predictive values for respective devices when sensitivity is set to 100%

	SBR $\geq 250$ $\mu\text{mol/l}$	SBR $< 250$ $\mu\text{mol/l}$	Totals	Positive predictive value
<b>Minolta</b>				
TcB reading $\geq 170$ $\mu\text{mol/l}$	17	32	49	34.7
TcB reading $< 170$ $\mu\text{mol/l}$	0	15	15	
Totals	17	47	64	
	100.0 (sensitivity)	31.9 (specificity)		
<b>SpectRx A</b>				
TcB reading $\geq 150$ $\mu\text{mol/l}$	17	37	54	31.5
TcB reading $< 150$ $\mu\text{mol/l}$	0	10	10	
Totals	17	47	64	
	100.0 (sensitivity)	21.3 (specificity)		
<b>SpectRx B</b>				
TcB reading $\geq 150$ $\mu\text{mol/l}$	17	34	51	33.3
TcB reading $< 150$ $\mu\text{mol/l}$	0	13	13	
Totals	17	47	64	
	100.0 (sensitivity)	27.7 (specificity)		

SBR, Serum bilirubin; TcB, transcutaneous bilirubinometry.

SBR at higher levels. This implies that the devices are less accurate when the risk for kernicterus is higher.

### Accuracy of SpectRx device

Although the aim of the study was to compare the SpectRx with the Minolta, we simultaneously evaluated accuracy for validation purposes. At that time, no published data on the SpectRx device were yet available in the medical literature. Since then, three papers have been published on the accuracy of the SpectRx device.<sup>7,8,17</sup> The correlation coefficient for the SpectRx in this study ( $r = 0.87$ ) is comparable to those obtained by Bhutani *et al*<sup>7</sup> ( $r = 0.91$ ), Rubaltelli *et al*<sup>8</sup> ( $r = 0.89$ ) and Knupfer *et al*<sup>17</sup> ( $r = 0.73$ ). The 95% confidence intervals of the difference are also similar at  $\pm 67 \mu\text{mol/l}$  in this study,  $\pm 55 \mu\text{mol/l}$  in Bhutani *et al*,<sup>7</sup> and  $\pm 76 \mu\text{mol/l}$  in Rubaltelli *et al*.<sup>8</sup> Our data are thus analogous to studies carried out elsewhere and add to the literature about the validity of the SpectRx device.

### TcB as a replacement for SBR sampling

Perhaps one limitation of this study is that SBR was analysed by a clinical laboratory method rather than the ideal method of high performance liquid chromatography (HPLC). It has been argued that, when compared with HPLC, some clinical laboratory analyses of SBR use automated methods that are subject to errors as wide as those obtained with TcB.<sup>8</sup> Our study used the Hitachi 911 multichannel analyser with the potassium ferricyanide method, which has been shown to be an accurate and precise assay for total SBR.<sup>10</sup> Furthermore, our laboratory participates in rigorous internal and external quality control procedures. Each institution should be aware of the accuracy of their local clinical laboratory method for measuring SBR before deciding whether transcutaneous devices are suitable replacements.

Rubaltelli *et al*<sup>8</sup> propose that transcutaneous measurements could be used to replace invasive SBR sampling altogether. They reported correlation coefficients for SpectRx compared with HPLC ( $r = 0.89$ ), and laboratory SBR with HPLC ( $r = 0.93$ ). The corresponding 95% confidence intervals of the difference were  $\pm 76 \mu\text{mol/l}$  and  $\pm 63 \mu\text{mol/l}$  (non-significant at  $p = 0.53$ ). Despite not reaching statistical significance, the 95% confidence intervals were superior for laboratory SBR. The authors concluded that the SpectRx performed as well as laboratory SBR. However, the overall accuracy of their laboratory SBR results was compromised by the use of nine different automated laboratory methods. It is known that certain methods perform more accurately than others,<sup>18</sup> but no information was provided about the individual performance of each of the nine methods.

In addition, no studies have addressed the question of whether skin or serum levels of bilirubin more accurately reflect its deposition in the central nervous system. Some suggest that transcutaneous readings are more representative of tissue deposition of bilirubin,<sup>19,20</sup> whereas others propose that circulating serum bilirubin is more available to move across the blood-brain barrier.<sup>8</sup> Factors that affect blood-tissue bilirubin distribution include exchange transfusion, phototherapy, albumin levels, pH, and gestational and postnatal age. This debate on the use of TcB as a replacement for SBR sampling would be ideally resolved by conducting a randomised controlled trial of the two diagnostic methods with long term outcome measurements and cost-benefit analyses. This would

require a large, multicentre setting with years of multidisciplinary follow up, and the cost would be prohibitive. The current evidence also shows that the transcutaneous methods at present available are not accurate enough to proceed with such a trial.

### Summary

The purpose of measuring bilirubin levels, by whatever method, is to identify infants who require treatment. The inaccuracy of existing transcutaneous bilirubinometers means that they should only be used as screening tools for SBR sampling. The newer SpectRx BiliCheck is no more accurate than the Minolta AirShields, and is more complicated to use.

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