The correlation between full biophysical profile and rapid biophysical profile in antepartum fetal surveillance

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ABSTRACT

Background: The aim of this study was to assess the efficiency of rapid biophysical profile in antepartum fetal surveillance and ability to predict adverse perinatal outcome and to compare it with the gold standard full biophysical profile. The objective was to determine the correlation between the rapid biophysical profile (RBP) and the full biophysical profile (FBP) and to compare the individual scores with pregnancy outcomes.

Methods: A prospective study was performed in 153 singleton pregnancies with no fetal anomalies between 34-42 weeks of gestation. All participants received both the standard (FBP) and the new RBP. Abnormal fetal test was defined as having a score of ≤ 6 for FBP or ≤ 2 for RBP. The main outcome measured was Spearman’s correlation coefficient (rs) between both examinations and also between each examination and pregnancy outcomes measured in terms of Apgar scores and NICU (Neonatal Intensive Care Unit) admissions.

Results: The data showed a positive correlation between the two tests (rs = 0.62; p < 0.0001). Out of the individual biophysical variables, only NST (Non Stress Test) had a positive correlation with RBP. The sensitivity, specificity, positive predictive value and negative predictive value of RBP in predicting adverse outcomes was found to be 71.4%, 87.1%, 35.7%, 96.8% respectively.

Conclusions: The statistically significant positive correlation between RBP and FBP has been revealed. Due to its simplicity, rapidity, and no need for experienced interpreter, the RBP may be alternatively used as a primary screening antepartum fetal test in the overcrowded obstetric center.

Keywords: Rapid biophysical profile, Sound provoked fetal movements, Antepartum surveillance
sound-provoked fetal movement (SPFM) test has been proposed by many researchers as a promising technique for fetal surveillance.\textsuperscript{2,3} Despite the extensive use of RBP, the study of the correlation between RBP and FBP is very limited. Hence, the present study analysed the correlation between the two tests in terms of abnormal and normal test detection and also with adverse perinatal outcomes.

**METHODS**

After the approval of the Institute’s Ethics Committee, 153 pregnant women who met the inclusion criteria of singleton high risk pregnancies between 34 – 42 weeks gestational age were invited to join the study. Multiple pregnancies and anomalous fetuses were excluded. The indications for fetal surveillance were hypertensive disorders of pregnancy, prolonged pregnancy, gestational diabetes mellitus, decreased fetal growth restriction. Informed consent was taken from all the participants. The study was designed in such a way that all participants received both FBP and RBP. NST was performed in all patients and then the remaining fetal ultrasound parameters (AFI, fetal breathing, fetal tone, and fetal movement) were examined to complete the FBP test. After a 10 minute break, SPFM was carried out by the same examiner to finish the RBP test. The obstetric care absolutely relied on the result obtained by the gold standard FBP technique.

The patient was made to lie down in the supine position and Doppler transducer (Edan fetal monitor model F3, Germany) applied to the maternal abdomen to record the FHR. In case no FHR accelerations occurred in 20 minutes, the recording was extended up to 40 minutes. The FHR tracing was considered reactive if 2 or more accelerations occurred in 20 minutes. If the criterion was not met even in 40 minutes or significant decelerations occur during this period, the test was interpreted as non-reactive.

The AFI and fetal biophysical variables were evaluated using real time scanner (Philips HD7XC) with a 5 MHz abdominal transducer. To obtain AFI, the uterus was divided into 4 equal quadrants, then the transducer was placed along the maternal longitudinal axis and held perpendicular to the floor. AFI was calculated by adding the vertical, cord free depth of the largest amniotic fluid pocket in each quadrant. The other fetal biophysical variables (fetal breathing, gross body movements and tone) was observed subsequently during the examination.

The SPFM was performed 10 minutes after completion of FBP by the same examiner. With the help of a fetal acoustic stimulator (Maestros, Fetal stimulator HX1, 80 MHz) a sound stimulus of 110 db was applied to the abdomen near the position of fetal head for a maximum of 3 seconds and the appearance of fetal movement were looked for on the screen of ultrasound machine. If movement occurred within 15 seconds of the application of stimulus the result was said to be response or normal. If movement does not occur within 15seconds the stimulus can be repeated up to 3 times before terming the test abnormal or no response.

According to Manning et al.,\textsuperscript{1} each of five biophysical variables has a possible score of 2, for a total of 10. The FBP scoring system is shown in Table 1. The score of ≤ 6 is said to be abnormal for FBP and indicates fetal hypoxia. For RBP, the scoring system used is shown in Table 2. The RBP score of 4, characterizes the reassuring fetal circumstance while the score of ≤ 2 represents the non-reassuring fetal status.

<table>
<thead>
<tr>
<th>Biophysical variable</th>
<th>Normal (score=2)</th>
<th>Abnormal (score=0)</th>
</tr>
</thead>
<tbody>
<tr>
<td>Fetal breathing movements (FBM)</td>
<td>One or more episodes of FBM&gt;30 sec in 30min</td>
<td>Absent or no episode of FBM&gt;30 sec in 30min</td>
</tr>
<tr>
<td>Gross body movements</td>
<td>3 or more discrete body/limb movements in 30 minutes (episodes of active continuous movement considered as single movement)</td>
<td>2 or less episodes of body/limb movements in 30 min</td>
</tr>
<tr>
<td>Fetal tone</td>
<td>1 or more episodes of extremity extension and subsequent flexion: opening and closing of hand considered normal tone</td>
<td>Either slow extension with return to partial flexion or movement of limb in full extension or absent fetal movements</td>
</tr>
<tr>
<td>NST</td>
<td>2 or more accelerations of 15 beats per minute for 15 sec within 20-40 min</td>
<td>0 or 1 acceleration within 20-40min</td>
</tr>
</tbody>
</table>

**Interpretation**

Score= 8-10 Normal fetus
Score= 6 Fetal hypoxia is suspicious
Score= 0-4 Fetal hypoxia
Table 2: Rapid BPP scoring system.4

<table>
<thead>
<tr>
<th>Biophysical variable</th>
<th>Normal (score=2)</th>
<th>Abnormal (score= 0)</th>
</tr>
</thead>
<tbody>
<tr>
<td>Sound provoked fetal movement</td>
<td>Response</td>
<td>No response</td>
</tr>
<tr>
<td>AFI</td>
<td>&gt;5cm</td>
<td>&lt;5cm</td>
</tr>
<tr>
<td>Total score</td>
<td>4</td>
<td>2</td>
</tr>
</tbody>
</table>

Interpretation
Score= 4 Normal fetus
Score=0-2 Fetal hypoxia

RESULTS

In the study population comprising 153 pregnant women, the mean maternal and gestational age of patients was found to be 26.6 ± 4.3 years and 38.5±1.7 weeks respectively. Majority (62.7%) of the cases were primigravida and more than half (55.56%) of participants had their gestational age between 37 weeks – 39 weeks + 6 days. Out of the high risk pregnancies analyzed, as shown in Figure 1, Hypertensive disorders of pregnancy accounted for 38.5% of cases, followed by post dated pregnancies (29.4%). Vaginal delivery was the result in 90 (58.8%) subjects. Among the 153 cases analyzed, 17% and 9.1% had abnormal FBP and RBP scores, respectively.

![Figure 1: Indication for antepartum fetal surveillance.](image)

Correlation among scores of RBP and FBP using Spearman’s correlation coefficient was found to be 0.62 and it was statistically significant (p <0.0001) as shown in table 3. Correlation of individual components of FBP with RBP in patients who had abnormal FBP score generated few interesting results. Among 26 patients who had an abnormal FBP score, 16 cases had non-reactive NSTs out of which 9 had abnormal RBP score. The correlation between NST and RBP in patients with an abnormal FBP proved to be r = 0.46 and statistically significant (p 0.01). In this study, only 1 case had an abnormal RBP score in spite of having a reactive NST. Baby was delivered through caesarean done in view of severe IUGR and oligohydramnios, and no adverse pregnancy outcome was noted. In the 26 cases with abnormal FBP score, 18 subjects had abnormal score for fetal movements and only 7 had an abnormal RBP score. The correlation was found to be r = 0.013 (p 0.94) and hence not statistically significant. Among the same 26 cases comparison of fetal tone, fetal breathing score with RBP score revealed a negative correlation of r=-0.118 (p 0.56) and r = -0.224 (p 0.27) respectively and hence devoid of any statistical significance.

Table 3: Correlation between FBP and RBP.

<table>
<thead>
<tr>
<th>RBP</th>
<th>FBP−Normal</th>
<th>FBP−Abnormal</th>
</tr>
</thead>
<tbody>
<tr>
<td>Normal</td>
<td>123</td>
<td>16</td>
</tr>
<tr>
<td>Abnormal</td>
<td>4</td>
<td>10</td>
</tr>
<tr>
<td></td>
<td>127</td>
<td>26</td>
</tr>
</tbody>
</table>

(FBP – full biophysical profile, RBP – rapid biophysical profile)

Taking Apgar score at 1 minute ≤7 as an adverse outcome, 18 (11.7%) cases had a low Apgar score. The correlation of scores of FBP, RBP with low Apgar at 1 minute was found to be r = 0.25 (p 0.001) and 0.47 (p<0.0001) respectively. Apgar score at 5 minutes ≤7 was taken as the second adverse outcome and number of cases meeting this criteria were 14 (9.1%). The correlation of scores of FBP, RBP with low Apgar at 5 minutes was worked out to be r = 0.19 (p 0.01) and 0.35 (p<0.0001) respectively and revealed the fact that correlation of abnormal RBP with Apgar at 1 minute and 5 minutes was better than that with abnormal FBP. The need for NICU admissions was taken as the last adverse outcome. In the present study 24 neonates required NICU admissions out of which prematurity being most common indication with 10 (41.7%) cases followed by meconium aspiration syndrome in 6 (25%) cases. In the present study rapid biophysical profile proved to have a sensitivity of 71.42%, specificity of 87.05%, positive predictive value of 35.71% and negative predictive value of 96.80% for predicting all adverse perinatal outcomes put together.

Table 4: Correlation between RBP and adverse perinatal outcome.

<table>
<thead>
<tr>
<th>Adverse outcome</th>
<th>RBP - Normal</th>
<th>RBP - Abnormal</th>
</tr>
</thead>
<tbody>
<tr>
<td>Present</td>
<td>121</td>
<td>10</td>
</tr>
<tr>
<td>Absent</td>
<td>18</td>
<td>4</td>
</tr>
<tr>
<td></td>
<td>139</td>
<td>14</td>
</tr>
</tbody>
</table>

(RBP – rapid biophysical profile, Adverse perinatal outcomes include low Apgar score at 1,5 minutes or NICU admission)

DISCUSSION

The FBP as a non-invasive, very accurate and applicable antenatal method to all patients is particularly attractive...
since it provides immediate individual results, does not provoke fetal distress. FBP has a low false positive rate and consists of acute markers of fetal hypoxia (fetal breathing, fetal movement, fetal tone, and FHR reactivity), and a chronic marker of fetal hypoxia that gives a better notion of uteroplacental reserve (AFI). However, there are disadvantages of this test. FBP is time-consuming as it includes at least a 30 minutes observation period of fetal biophysical activities and NST, which requires 20-40 minutes. Moreover, an expensive fetal heart rate monitor and an experienced interpreter is needed. RBP is simpler, inexpensive, and is faster. It has been developed to evaluate fetal well being when an NST machine is unavailable. The present study has demonstrated a correlation between RBP and FBP test and is very similar to results obtained by Phattanchindakun et al. Taking individual biophysical variables and comparing them with RBP in cases where the subject had an abnormal FBP score, only NST was found to have a statistically significant positive correlation with RBP. AFI, the chronic hypoxia indicator, and SPF, the acute fetal hypoxia marker, have been thoroughly examined by the RBP. Keeping low Apgar score at 1 and 5 minutes and NICU admissions as adverse perinatal outcome, the sensitivity, specificity, positive predictive value, negative predictive value of RBP in predicting adverse pregnancy outcome when all three outcomes were taken together was found to be 71.4%, 87.1%, 35.7%, 96.8% respectively. Owing to the differences in sample size and indications for fetal assessment, earlier studies by Chousawai et al. have shown wide variations in the statistical parameters of Rapid BPP. In spite of the variations the obtained data encouraged the use of RBP as an alternative antepartum test to evaluate the fetal well-being. In particular, its simplicity, shorter duration, no obligation of NST, or experienced interpreter makes the RBP a good choice for the obstetric center that is rather crowded or limited in experienced NST interpreters. In addition, the RBP does not need expensive high-resolution ultrasound equipment. If this technique is applied as a screening fetal test in rural areas, it will help in reduce the number of referral cases for FBP in tertiary care centers. However, the accuracy of RBP test (in terms of sensitivity, specificity, false positive, and false negative rates) should be extensively verified and a larger number of studied populations including more abnormal tests need to be investigated. This instrument produces the specific quality of sound with 110 dB of loudness and the frequency of 80 MHz. Despite its high efficacy, other inexpensive instrument that can generate the same quality of sound should be invented and studied to reduce the cost further.

CONCLUSION

With the positive correlation of rapid BPP with full BPP and its ability to predict adverse outcome, rapid BPP can be used as a good screening test for high risk pregnancies in busy obstetric setups where lack of experienced personnel and advanced equipment for surveillance is the main limitation. In case of an abnormal rapid BPP we suggest that the patient be subjected to a NST and followed up with FBP. As only those few patients who have an abnormal RBP score need to be subjected to further intensive surveillance, it saves a lot of time and energy.

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Ethical approval: The study was approved by the Institutional Ethics Committee

REFERENCES
