Quantitative Analysis of Fetal Actocardiogram: Update

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Abstract

Aims: The actocardiogram (ACG), which recorded fetal heart rate (FHR) and movements, was quantitatively analyzed, due to its advantage to the FHR patterns diagnosis, to recognize the effect of fetal movement on FHR, to solve controversial FHR problems, to correctly evaluate fetal disorders, and to evaluate the loss of variability and acceleration.

Methods: FHR changes were diagnosed by the FHR score, fetal movements were evaluated by 4 ACG parameters, fetal behavior and abnormal FHR were quantitatively determined, physiologic sinusoidal FHR was diagnosed by the ACG and FHR frequency analysis, the developing mechanism of FHR acceleration and variability were studied to diagnose the brain damage in the loss of FHR variability. 1.3 Results: The FHR score, neural network analysis and A/B ratio predicted short and long term outcomes in the 1st stage of labor or even in pregnancy. The ACG and frequency analysis differentiated physiologic sinusoidal from the true one, Controversial problems in FHR were solved by the quantitative ACG analysis. The loss of FHR variability was the sign of fetal brain damage even in fetal non-hypoxic insults.

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Conclusion: Quantitative analyses of FHR and fetal movements in ACG were indispensable in the fetal diagnosis even in general insults. Since cerebral palsy (CP) could develop in the loss of FHR variability, C-section is recommended to perform before the loss of FHR variability.

Keywords: FHR; Fetal movement; Actocardiogram; FHR score; A/B ratios; Outcome prediction; FHR variability; Acceleration; Fetal disorders

Definition

Actocardiogram (ACG): The ACG simultaneously traced FHR curve and fetal movement signals. The MHz level continuous wave (CW) ultrasound was used, of which SPTA intensity was as weak as 1 mW/cm². The fetal movement Doppler signals obtained by 2 MHz source ultrasound was 20-50 Hz. The ultrasound detected Doppler fetal heart beat signals at the same time, of which Doppler frequency was 100 or more Hz, i.e. single ultrasonic probe detected two phenomena, FHR and fetal movements. Fetal movement Doppler signal was separated by a band-pass filter, and formed low frequency spikes, of which amplitude was parallel to that of fetal movement (Figures 1 and 2) [1]. Uterine contraction was also simultaneously recorded to prepare the CTG function.

Cardiotocogram (CTG): is simultaneous record of FHR and uterine contraction curves.

Fetal movements: The ACG motion signal developed by fetal trunk movements, because the ultrasound targeted fetal heart. The motion was fetal respiratory movements (continuous and periodic respiratory changes), fetal hiccupping, and the trunk motion caused by conducted extremity movements, and mouthing movements [2].

Fetal heart rate (FHR): More than 100 Hz Doppler signals developed by fetal heart beats were introduced into an autocorrelation FHR meter, which detected the FHR up to 210 beats per minutes (bpm) [1].

Uterine contraction: It is simultaneously recorded on the ACG chart usually by external tocodynamometry, by which no intrauterine pressure is evaluated, but used for the diagnosis of late FHR deceleration.

FHR baseline: Continuous FHR tracing without acceleration or deceleration. Normal baseline is 110 to 159 bpm.

Fetal bradycardia: The baseline FHR less than 110 bpm.

Fetal tachycardia: The baseline FHR of 160 bpm or more.

FHR variability: Minor variation of FHR baseline (long term variability, LTV) in the FHR recorded by ultrasonic Doppler autocorrelation FHR meter. The LTV amplitude is normal when it is 5 to 24 bpm.

Acceleration: Transient FHR increase for 10 or more bpm and 10 or longer sec duration is normal before 30 weeks of pregnancy, and 15 bpm and 15 sec or more is normal in 30 or more weeks of pregnancy.

Deceleration: Fifteen or more bpm deep and 15 or more sec long transient FHR decrease. V-shaped slow deceleration is a periodic deceleration including early (ED) and late (LD) decelerations comparing to uterine contraction. U-shaped irregular deceleration with sudden decrease and recovery is variable deceleration, divided into mild, moderate and severe variable deceleration (MVD and SVD). The LD and SVD are pathological in the pattern classification.

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A/B ratio: Total sum of acceleration duration ratio to the total sum of fetal movement duration in a study period.

Introduction

Although fetal movements were evaluated by maternal perception or abdominal motion detected by mechanical actograph at abdominal surface in old time, the results were insufficient to scientific fetal movement study [3]. A direct detection of fetal movement at fetal surface was required, and the device was invented by Maeda in 1984 [1]. Although the CTG was diagnosed mainly by FHR pattern classification [4], several vague problems and diagnostic difficulty were found in the past. It was, therefore, necessary to analyze the CTG with quantified techniques and numeric evaluation, and the result was the creation of FHR score [3-6]. Still there was the difficulty of differentiation of resting fetal state from non-reactive FHR, the differentiation of physiological sinusoidal FHR from true ominous sinusoidal FHR, and so on, which were related to the lack of fetal movement study. Although some problems were solve by the real-time B-mode ultrasound [7], most diagnostic difficulties were solved by the handling of ultrasonic Doppler fetal movement signals [1]. The prototype actocardiograph was hand-made by Maeda (Figures 1 and 2) and reported after the confirmation of its basic properties [1].

Methods

The FHR was quantified by the FHR score, and fetal movements by the ACG. There were various commercial ACG models after the prototype ACG (Figure 1), including MT-320, MT-325, MT-332, MT-333U, MT-340, MT-516, MT-522 and MT-540 (TOITU, Tokyo). Recent ACG works were mainly studied by the ACG records offered by friends of the author, therefore, deep appreciation is expressed to the kind offering. Four parameters were quantitatively analyzed in the quantified studies on fetal behavior [2]. Physiologic sinusoidal FHR was differentiated from true ominous one by fetal periodic respiratory and swallowing movements [8]. Developmental mechanism of acceleration and LTV was studied. Short and long fetal outcomes were studied by the A/B ratio [9]. New prognostic value of the loss of LTV was clarified in this report.

Quantitative diagnosis of CTG by FHR score

It was the first trial in 1960s to quantitatively analyze intrapartum CTG instead of visual pattern classification (Figure 3). The percentage of low Apgar score gave evaluation scores from 1 to 4 in abnormal FHR changes (Table 1). The method was statistically a goodness measure. The evaluation scores were summarized in 5 min to obtain the FHR score [2], which was compared to Apgar scores and significant correlation was found (Figure 4) [5]. The CTG analysis with FHR score were programmed in the computer for automated CTG diagnosis [5,6,10].

Fetal state is abnormal, if FHR score is 10-19, and highly abnormal if the score is 20 or more. Thus the FHR score comprehensively evaluates fetal status.
Quantitative analysis of fetal actocardiogram: The fetal ACG was analyzed by four parameters as follows:

1. Burst duration (sec): Mean duration of fetal movement burst
2. Occupancy (%): Percentage of the sum of movement burst durations against the whole analysis time.
3. Frequency (cycles per minute, cpm): The incidence of movement bursts in one min
4. A/B ratio (%): The ratio of the sum of acceleration durations to the sum of movement burst durations.

Four parameters' values were determined in the fetal ACGs of normal fetuses in the late stage of pregnancy (Table 2). Fetal behavior will be determined by 4 parameters in new cases. The importance of A/B ratio was confirmed, because it is not influenced by fetal behavior in the evaluation of fetal statuses.

Difference of ACG parameters in normal fetus and fetal hypoxia:
There was significant difference of quantified parameters between normal and hypoxic fetuses, i.e. occupancy was 32.67% in normal fetus and 10.00% in hypoxia, frequency was 0.65 and 0.24 cpm, A/B ratio was 1.03 and 0, respectively [13].

Differentiation of physiologic sinusoidal FHR from true ominous one by the ACG:
The CTG hardly differentiated physiologic sinusoidal FHR from the truly ominous one. A physiologic one was easily diagnosed to be harmless in fetal ACG when the periodic fetal movements synchronized to the sine wave-like FHR, e.g. periodic fetal respiratory or mouth movements provoke physiologic sinusoidal FHR (Figure 5) [8].

Developmental mechanism of FHR acceleration and LTV studied by the ACG:

### Table 2: Quantitative values in four fetal behavioral states detected by fetal ACGs.

<table>
<thead>
<tr>
<th>Behavioral States</th>
<th>Duration (sec)</th>
<th>Occupancy</th>
<th>% Frequency (cpm)</th>
<th>A/B ratio (%)</th>
<th>N</th>
</tr>
</thead>
<tbody>
<tr>
<td>Resting</td>
<td>29.7 ± 10.3</td>
<td>32.7 ± 14.8</td>
<td>0.7 ± 0.2</td>
<td>1.4 ± 0.4</td>
<td>14</td>
</tr>
<tr>
<td>Active</td>
<td>17.7 ± 3.7</td>
<td>6.4 ± 1.8</td>
<td>0.2 ± 0.03</td>
<td>1.2 ± 0.3</td>
<td>5</td>
</tr>
<tr>
<td>Highly active</td>
<td>86.6 ± 14.6</td>
<td>44.1 ± 5.3</td>
<td>0.5 ± 0.3</td>
<td>1.2 ± 0.1</td>
<td>3</td>
</tr>
</tbody>
</table>

It was clear also in the fetus, i.e. a fetal movement burst accompanies triangular FHR acceleration (Figure 6), where the movement preceds acceleration for approx. 7 sec. The acceleration was lost in the non-
reactive FHR, which follows severely hypoxic FHR changes some days later, including the bradycardia, late deceleration and the loss of variability [14].

The mechanism to produce triangular shape of acceleration

A triangular curve developed in electric wave bursts after passing through an integral circuit with 7 sec delay time constant. Also triangular heart rate curves developed in the 1 min lasting continuous leg motions in an adult [15]. The electric and physiologic simulations explained the triangular FHR developing mechanism, which was the brain excitation in the mid brain by the movements, i.e., Terao et al. [16] reported that the acceleration developed in the mid brain.

The mechanism to develop FHR variability in the brain

A repeated moderate fetal movements provoked synchronized FHR accelerations (Figure 6), and minor fetal movements also provoked FHR baseline changes (Figure 7). These facts indicate that the long term variability (LTV) develops as the response of fetal brain to minor fetal movements. Since the FHR delay was 7 sec, the location to respond the movement is the same as FHR acceleration, i.e. it is the mid brain [16].

Prenatal quantitative diagnosis of outcome with the A/B ratios of ACG

The A/B ratio was standardized dividing the total duration of acceleration by the total duration of movement bursts (Figure 8). ACGs of 15 common fetal disorders were quantitatively analysed with A/B ratios, and compared to the Apgar scores and the numeric long term outcome after births [9].

The short term outcome analysed by 1 and 5 min Apgar score closely correlated A/B ratios, i.e. Y (1 min Apgar)=7.68X (A/B ratio)-1.75, R²=0.85, p<0.001, Y (5 min Apgar)=6.44X+0.58, R²=0.68, p<0.001, and Y (numeric long term outcome)=6.42X+0.05, R²=0.71, p<0.001. Short and long term outcomes were abnormal, when the A/B ratio was less than 1. The fact was remarkable that a spastic quadriplegia case was found among cases of lower A/B ratio than 1. Therefore, A/B ratio of ACG is a useful parameter to expect fetal short and long term outcomes. Regression equations are listed [9].
COMMENT

Comprehensive evaluation of the fetus

Fetal diagnosis with the pattern classification tends to the single item diagnosis but not to cover the total condition of the fetus. In quantitative diagnosis, the FHR score predicts Apgar score and umbilical cord arterial blood pH even in the first stage of labor. Neural network analysis reports the pathological outcome probability. In the ACG, the A/B ratio predicts pathological Apgar score and long term outcome after the birth. Automated comprehensive diagnosis and outcome prediction are big advantage of quantitative analysis of FHR and ACG.

Advantage of fetal movement record

Since the change of movement signal of ACG is the same as those of fetal movement (Figure 2), the influence of fetal movement on FHR is precisely evaluated, i.e. fetal movement precedes FHR change, e.g. periodic fetal movements provoked sine wave-like FHR changes (Figure 5), where the physiologic sinusoidal FHR is differentiated from the truly ominous sinusoidal heart rate by the ACG. Resting state of the fetus is differentiated by the absence of fetal movement bursts from non-reactive FHR. The loss of acceleration against fetal movement burst shows hypoxic suppression of fetal brain predicting the severe hypoxia.

Frequency analysis of FHR traces

The technique was introduced with the purpose to automatically diagnose truly ominous sinusoidal FHR, which appeared without fetal movement record, and quantitatively confirmed by the FFT frequency analysis [11]. It was a surprise to diagnose severe loss of FHR variability less than the resting fetal state also by a FFT frequency analysis [12].

The fetal state in severe loss of variability (LTV)

Since anencephalic fetal ACG recorded neither acceleration nor LTV, normal fetal brain is responsible to the development of acceleration and LTV (Figures 9-11), but not in the brain cortex because no heart rate change was recognized by the person who developed the triangular acceleration in adult exercise [15], and the fetal acceleration was reported to be produced in the midbrain in the studies on anencephalic fetus [16]. The FHR acceleration disappeared in the non-reactive FHR, while the LTV was preserved, and severe hypoxic FHR changes appear some days later in non-reactive FHR cases, including bradycardia, late deceleration and the loss of LTV. The outcome of C-section in this state was ominous, if compared to reactive FHR [14], i.e. fetal damage was heavy in cases of the loss of LTV, which will be severe brain damage. In an intrapartum fetus of heavy loss of LTV less than 1 bpm similar to anencephalic fetus (Figure 12) whose mother refused C-section and a severe loss of LTV means such brain damage as the loss of brain in anencephaly [15].
Abnormal FHR in general insults without hypoxia

Abnormal FHR changes were reported in cytomegalovirus (CMV) infected fetus [17] and in congenital syphilis [18]. Therapeutic decision still seems controversial in the occasion because of no hypoxia. However, as discussed above, such abnormal FHR as severe loss of LTV is caused by fetal brain damage, the FHR changes can develop by viral or bacterial toxin or other general insults. As the loss of LTV means severe damage of fetal brain, FHR changes in various general insults should be treated by the same strategy as hypoxic FHR changes, i.e. C-section before the loss of variability. The A/B ratio of ACG will be helpful for the outcome prediction.

Computerization of fetal monitoring

The FHR score calculation, neural network analysis of FHR, frequency analysis of FHR, discussed in this report, were programmed and working in the centralized fetal monitoring, improving perinatal status [5,6,10]. Quantified ACG and hypoxic index will be added to the system in the near future. An attending obstetrician will input continuous FHR into the computer system, and receive analyzed results rapidly and directly from the system. The time consuming visual CTG analysis will be changed to the computerization to receive objective, comprehensive and precise diagnostic data [6].

Conclusion

Since the ACG and its quantitative analysis proposed new field of fetal evaluation, which is totally objective in quantitative as well as visual analysis of ACG as discussed in this report, the subjective visual FHR pattern diagnosis, which was vague with big interobserver difference, is definitely improved and the fetal management is greatly progressed by the introduction of quantified FHR analysis and fetal ACG, where C-section was recommended to be performed before the loss of FHR variability. In addition, computerized automatic fetal diagnosis, which utilizes various quantified analyses, extensively improves obstetric statuses.