







Improvements in fetal heart rate analysis by the removal of maternal-fetal heart rate ambiguities

Paula Pinto^{1,2}, Cristina Costa-Santos³, Hernâni Gonçalves³, Diogo Ayres-De-Campos^{2,4,5} and João Bernardes^{2,3,4,6*}

Abstract

Background: Misinterpretation of the maternal heart rate (MHR) as fetal may lead to significant errors in fetal heart rate (FHR) interpretation. In this study we hypothesized that the removal of these MHR-FHR ambiguities would improve FHR analysis during the final hour of labor.

Methods: Sixty-one MHR and FHR recordings were simultaneously acquired in the final hour of labor. Removal of MHR-FHR ambiguities was performed by subtracting MHR signals from their FHR counterparts when the absolute difference between the two was less or equal to 5 beats per minute. Major MHR-FHR ambiguities were defined when they exceeded 1 % of the tracing. Maternal, fetal and neonatal characteristics were evaluated in cases where major MHR-FHR ambiguities occurred and computer analysis of FHR recordings was compared, before and after removal of the ambiguities.

Results: Seventy-two percent of tracings (44/61) exhibited episodes of major MHR-FHR ambiguities, which were not significantly associated with any maternal, fetal or neonatal characteristics, but were associated with MHR accelerations, FHR signal loss and decelerations. Removal of MHR-FHR ambiguities resulted in a significant decrease in FHR decelerations, and improvement in FHR tracing classification.

Conclusions: FHR interpretation during the final hour of labor can be significantly improved by the removal of MHR-FHR ambiguities.

Keywords: Heart rate, Fetal, Maternal heart rate, Cardiotocography, Misinterpretation, Computer analysis, FIGO guidelines

Background

Fetal heart rate (FHR) monitoring may be affected by the temporary acquisition of the maternal heart rate (MHR), both when using external monitoring with Doppler ultrasound [1], and when using internal monitoring with electrocardiography (ECG) [2–6]. Inadvertent MHR acquisition with external monitoring has been reported in up to 90 % of intrapartum recordings, for an average of 6.2 % of tracing length [1], while with internal monitoring it is usually due to the detection of maternal signals transmitted via the fetal electrode in cases of fetal death [2, 5]. Overall, significant errors in FHR interpretation may occur, including missing the diagnoses of newborn acidemia [7, 8] and fetal death [2–6].

Some methods reduce the occurrence of these MHR-FHR ambiguities. FHR signal acquisition with internal [9, 10] or transabdominal [1] ECG, rather than with Doppler ultrasound, has been shown to improve signal quality, making it less prone to MHR-FHR ambiguities [1, 9, 10]. Simultaneous registration of the MHR using ECG or oximetry [7, 11] allows a comparison of MHR and FHR recordings and facilitates the detection of overlapping segments. Detection of P waves in the fetal ECG may be also helpful as a marker of fetal signals [12]. Visualization of FHR movements on ultrasound is recommended in doubtful cases, namely when fetal death is suspected [11].



© 2015 Pinto et al. **Open Access** This article is distributed under the terms of the Creative Commons Attribution 4.0 International License (http://creativecommons.org/licenses/by/4.0/), which permits unrestricted use, distribution, and reproduction in any medium, provided you give appropriate credit to the original author(s) and the source, provide a link to the Creative Commons license, and indicate if changes were made. The Creative Commons Public Domain Dedication waiver (http://creativecommons.org/publicdomain/zero/1.0/) applies to the data made available in this article, unless otherwise stated.

^{*} Correspondence: jbernardes59@gmail.com

²Departamento de Obstetrícia e Ginecologia, Faculdade de Medicina da Universidade do Porto, Porto, Portugal

³Center for Health Technology and Services Research (CINTESIS), Porto, Portugal

Full list of author information is available at the end of the article

While there is a reasonable amount of evidence to document cases of missed newborn acidemia and fetal death associated with MHR-FHR ambiguities, little research has been published on more subtle signal contaminations [1]. In particular, the effect of systematic cleaning of the FHR signal when MHR is suspected has, to our knowledge, not been previously evaluated.

The objective of this study was to assess the effect of removal of MHR-FHR ambiguities on the analysis of FHR recordings, during the last hour of labor. The hypothesis was that this would alter the identification of some FHR features and thus improve overall tracing classification.

Methods

The study followed the Helsinki Declaration, was approved by the local Ethics Committee ("Comissão de Ética do SESARAM") and all women gave their informed written consent to participate. Sixty-two consecutively acquired simultaneous recordings of MHR and external FHR, with good signal quality were selected, from the same number of labouring women, with uneventful singleton pregnancies, with fetuses in cephalic presentations, in the last recorded hour before birth (with a maximum 10minute interval before vaginal delivery or 30 min before caesarean delivery). All but two women were under epidural analgesia on request. Labor protraction or arrest, secondary to poor uterine activity, was treated with oxytocin, according to the local protocol. No other drugs with a potential to effect MHR or FHR were administered, namely salbutamol or parasympathetic agonists.

Gestational ages were confirmed by first trimester ultrasound dating, Apgar scores were estimated by the attending healthcare professional, and umbilical artery blood pH was obtained by paired sampling immediately after birth.

For acquisition of MHR signals, an ECG sensor was connected to three electrodes positioned on the maternal thorax, and for FHR signals a Doppler ultrasound probe was placed on the maternal abdomen. One case was excluded because of poor signal quality that required conversion to internal FHR monitoring. Acquisition of signals at 1600 Hz and heart rate (HR) extraction was performed using a STAN[®] 31 fetal monitor (Neoventa Medical, Gothemburg, Sweden).

HR, in beats per minute (bpm), was then conveyed at 4-Hz via the digital port of the fetal monitor to the Omniview-SisPorto[®] system (Speculum, Lisbon, Portugal) for signal recording and analysis. The system closely follows the International Federation of Gynaecology and Obstetrics (FIGO) guidelines for fetal monitoring [13–16] and has been extensively validated both in the ante [16, 17] and intrapartum periods [15, 17–19]. Analysis is carried out without signal reduction or averaging (Fig. 1). The baseline is estimated using a complex algorithm based on histogram and STV analysis, aimed at finding the HR level of stable segments with normal variability, within the physiological limits of 110–150 bpm [14–16]. STV is determined as the difference between two adjacent HR beats, and considered abnormal when lower than one beat per minute. Accelerations and decelerations are detected as HR deviations, above or below baselines, with at least 15 bpm of amplitude and 15 s duration. LTV is estimated in HR segments that do not display accelerations or decelerations, as the difference between the highest and lowest values, in a sliding window of 1 min, and is classified as abnormal when <5 bpm [13-16]. All of these basic FHR features are then used by the system to elicit an overall tracing classification, as green or blue (normal tracing), yellow (suspicious tracing) or red (pathologic tracing)) [13, 14]. The system was developed essentially for FHR analysis, but the same principles were applied in this study for analysis of the MRH, after scale conversion [13].

To remove MHR-FHR ambiguities, FHR signals were subtracted of their MHR counterparts when the absolute difference between them was equal or less than 5 bpm. This threshold was based on a report by Reinhard et al. [1] and on a pilot test performed in five cases with typical MHR-FHR ambiguities detected by visual analysis. Major MHR-FHR ambiguities were defined when more than 1 % of the tracing was affected and minor ambiguities when this value was between 0 and 1 % (Table 1).

Maternal, fetal and neonatal characteristics associated with major and minor MHR-FHR ambiguities were evaluated (Table 2). In tracings with and without major ambiguities, computer analysis of FHR recordings before and after the deletion of ambiguities was compared, (Table 3), and the association between FHR classification and newborn umbilical artery blood pH < or \geq 7.15 was assessed.

For statistical analysis, the results were expressed as medians (with inter-quartile ranges), for skewed continuous variables. The Mann-Whitney test was used to compare the groups with minor or major ambiguities (Table 1) and the Wilcoxon sign rank was used to compare the classification results before and after removal of ambiguities (Table 2). For other continuous variables, results were expressed as means and standard deviations. The independent sample *t* test was used to compare the groups with minor or major ambiguities (Table 1) and the paired t test was used to evaluate results before and after removal of ambiguities (Table 2). To compare categorical variables between minor or major ambiguity groups (Table 1) and before and after ambiguity removal (Tables 2 and 3), the Chi-square (or Fisher exact test) and McNemar tests were used, respectively. Ninety-five percent confidence intervals were calculated as appropriate and p-values considered significant at less than 0.05.



Fig. 1 Maternal heart rate (MHR) baseline misinterpreted as a prolonged fetal heart rate (FHR) deceleration. *Top*: simultaneous recording of the last 45 min of the MHR (*black*), FHR (*pink*) and uterine contractions (UC) signals; *Middle and Bottom*: computer analysis before and after removal of MHR-FHR ambiguities. For computerized analysis of MHR tracings, a scale change (MHR + 50 beats per minute) was performed (*last hour*). After the removal of ambiguities, the FHR alarm changed from red to blue

Results

The general maternal, fetal, delivery and newborn characteristics, in relation with the presence of minor and major MHR-FHR ambiguities, are presented in Table 1. Overall, average maternal body mass index and age were, respectively, 27 Kg/m² (SD:3) and 28 years (SD:5). Average gestational age was 39 weeks (SD:1). Twenty-six percent of women were multiparous and all fetuses were in cephalic presentation. The average systolic and diastolic blood pressures were, respectively, 122 (SD:11) and

	MHR-FHR ambiguity			
	$\overline{\text{Minor } (n=17)}$	Major ($n = 44$)	р	
Maternal characteristics				
Age (years) mean (SD)	26 (6)	28 (5)	0.134	
BMI mean (SD)	27 (3)	28 (3)	0.186	
Multiparity n (%)	5 (29)	11 (25)	0.752	
Labour				
Epidural <i>n (%)</i>	17 (100)	42 (95)	1.000	
Mode of delivery			0.105	
Cesarean n (%)	4 (24)	2 (4)		
Operative vaginal n (%)	5 (29)	18 (41)		
Normal vaginal <i>n (%)</i>	8 (47)	24 (55)		
MHR				
Signal loss (%) median (IQR)	9 (1 to 11)	6 (1 to 16)	0.834	
Basal MHR (bpm) <i>mean (SD)</i>	80 (12)	77 (12)	0.292	
Accelerations (n) mean (SD)	12 (6)	18 (10)	0.013	
Decelerations (n) median (IQR)	0 (0 to 0.5)	0 (0 to 0)	0.913	
Abnormal LTV (%) median (IQR)	0 (0 to 0)	0 (0 to 0)	0.705	
FHR				
Signal loss (%) mean (SD)	13 (7)	22 (11)	0.002	
Baseline (bpm) <i>mean (SD)</i>	140 (10)	136 (12)	0.180	
Accelerations (n) median (IQR)	9 (3 to 15)	7 (4 to10)	0.415	
Decelerations (n) median (IQR)	3 (1 to 7)	8 (6 to 10)	0.002	
Prolonged decelerations (n) median (IQR)	0 (0 to 0) 0 (0 to 1)		0.073	
Abnormal LTV (%) median (IQR)	3 (1 to 9)	1 (0 to 3)	0.055	
Alarm <i>n (%)</i>			0.001	
Green-blue	14 (82)	15 (34)		
Yellow-red	3 (18)	29 (66)		
Newborn				
Gestational age (weeks) mean (SD)	39 (1)	39 (1)	0.672	
Male <i>n</i> (%)	8 (47)	26 (59)	0.396	
Birthweight (grams) mean (SD)	3171 (366)	3254 (318)	0.384	
Apgar 1 <i>median (IQR)</i>	9 (9 to 10)	9 (9 to 10)	0.530	
Apgar 5 <i>median (IQR)</i>	10 (10 to 10)	10 (10 to 10)	0.298	
UAB pH <i>mean (SD)</i>	7.25 (0.08)	7.24 (0.07)	0.586	

Table 1	Maternal, fetal,	delivery and	newborn	characteristics	associated with	minor and	l major MHR-FH	R ambiguities
---------	------------------	--------------	---------	-----------------	-----------------	-----------	----------------	---------------

SD standard deviation, *IQR* inter-quartile range, *BMI* body mass index *p* values with statistical significance are displayed in bold

74 mmHg (SD:8) and the average axillary temperature was 36.2 $^{\circ}$ C (SD:0).

C-section was performed in 10 % of the group and instrumental vaginal delivery in 38 %. All C-sections were performed during the first stage of labor because of protracted or arrested labour. Forty-four percent of newborns were males, with an average one and five-minute Apgar score of 9 (range: 9-10) and 10 (range:10-10), respectively. The average umbilical artery blood pH was 7.24 (SD:0.07). The average

length of the active phase of labour was 297 min (SD:170).

Five tracings did not exhibit any MHR-FHR ambiguity, 12 exhibited 1 % of ambiguities and 44 (72 %) exhibited major MHR-FHR ambiguities (11 tracings with 2 %, 7 with 3 %, 5 with 4 % and 21 with 6-33 %).

In the group with minor MHR-FHR ambiguities, maternal, fetal, delivery and newborn characteristics were similar in the five cases that did not exhibit any ambiguity compared with the 12 cases that exhibited 1 % of

	Ν	Before	After	р
No major MHR-FHR ambiguity				
Signal loss (%) <i>mean (SD)</i>	17	13 (7)	14 (7)	<0.001
Basal FHR (bpm) <i>mean (SD)</i>	17	140 (10)	140 (10)	1.000
Accelerations (n) median (IQR)	17	9 (3 to 15)	9 (3 to 15)	0.157
Sporadic decelerations (n) median (IQR)	17	3 (2 to 11)	3 (1 to 8)	0.317
Prolonged decelerations (n) median (IQR)	17	0 (0 to 0)	0 (0 to 0)	1.000
Abnormal LTV (%) median (IQR)	17	3 (0 to 7)	3 (1 to 11)	0.317
Alarm <i>n (%)</i>	17			1.000
Green-Blue		14 (82)	14 (82)	
Yellow-red		3 (18)	3 (18)	
Major MHR-FHR ambiguity				
Signal loss (%) <i>mean (SD)</i>	44	22 (11)	28 (13)	<0.001
Basal FHR (bpm) <i>mean (SD)</i>	44	136 (13)	136 (12)	0.129
Accelerations (n) median (IQR)	44	7 (4 to 10)	6 (3 to 10)	0.285
Sporadic decelerations (n) median (IQR)	44	8 (6 to 10)	6 (4 to 10)	0.001
Prolonged decelerations (n) median (IQR)	44	0 (0 to 1)	0 (0 to 0)	0.046
Abnormal LTV (%) median (IQR)	44	1 (0 to 3)	1 (0 to 3)	0.414
Alarm <i>n (%)</i>	44			0.021
Green-Blue		15 (34)	23 (52)	
Yellow-red		29 (66)	21 (48)	
Overall				
Signal loss (%) <i>mean (SD)</i>	61	20 (11)	24 (13)	<0.001
Basal FHR (bpm) <i>mean (SD)</i>	61	137 (12)	137 (12)	0.129
Accelerations (n) median (IQR)	61	7 (4 to 10)	7 (3 to 10)	0.170
Sporadic decelerations (n) median (IQR)	61	7 (4 to 10)	6 (3 to 9)	0.001
Prolonged decelerations (n) median (IQR)	61	0 (0 to 1)	0 (0 to 0)	0.046
Abnormal LTV (%) median (IQR)	61	1 (0 to 4)	2 (0 to 5)	0.197
Alarm n (%)				0.021
Green-Blue		29 (48)	37 (61)	
Yellow-red		32 (52)	24 (39)	

SD standard deviation, IQR inter-quartile range

p values with statistical significance are displayed in bold

Table 3 Changes in overall FHR classification before and after
the removal of MHR-FHR ambiguities, in acidemic and non-
acidemic cases (umbilical artery blood pH <7.15 or ≥7.15)

	Before n (%)	After n (%)	Р
Acidemic newborns ($n = 7$)			1.000
Green-Blue	3 (57)	3 (57)	
Yellow-red	4 (43)	4 (43)	
Non-acidemic newborns ($n = 54$)			0.021
Green-Blue	26 (48)	34 (63)	
Yellow-red	28 (52)	20 (37)	

p values with statistical significance are displayed in bold

ambiguities. Major ambiguities were significantly associated with MHR accelerations, FHR signal loss, decelerations and yellow or red alarms, when compared with minor ambiguities (Table 1).

Removal of ambiguities resulted in a significant increases in FHR signal loss, both in cases with minor and major ambiguities. When major ambiguities occurred, a decrease in the number of FHR decelerations, and a decrease in the false positive rate of yellow-red alarms for the detection of acidemic fetuses, were also observed, but this was not seen with minor ambiguities (Tables 2 and 3).

Discussion

In this study, 72 % of tracings acquired in the last recorded hour of labour exhibited episodes of major MHR-FHR ambiguity. This concurs with the report by Reinhard et al. [1] showing that MHR-FHR ambiguities with clinical significance may be much more frequent and subtle than suggested by anecdotal reports published in the literature, most of them confined to extreme cases of fetal acidemia or death [2–8].

Another finding in the present study is that MHR-FHR ambiguities should not only be suspected when accelerations coincide with uterine contractions [2–8, 10], but also when FHR decelerations occur in association with signal loss (Table 1 and Fig. 1). Most MHR tracings are characterized by repetitive accelerations, which may overlap FHR signals and mimic FHR decelerations or baseline segments during episodes of FHR signal loss [15, 20, 21] (Fig. 1). With the high epidural rate observed in this population, it is possible that there was a lower increase in MHR during active pushing, and therefore only a few cases of suspected FHR accelerations due to inadvertent MHR recording were detected. Only 7 % cases had a MHR baseline above 100 bpm, and in no cases it exceeded 110 bpm. MHR accelerations rarely reached the FHR baseline and this resulted in frequent MHR-FHR ambiguities being caused by MHR baseline segments being confused with FHR decelerations (Fig. 1). It may be almost impossible to detect subtle misinterpretations without computer analysis of simultaneous MHR and FHR recordings [20], with automated detection of ambiguities.

C-sections were not evaluated separately as risk factor of MHR-FHR ambiguities because of the small number of cases. However, it was observed that C-sections were associated with fewer MHR-FHR ambiguities, although without assessing statistical significance. This probably occurred because C-sections were performed predominantly during the first stage of labour. The second stage is characterized by higher signal loss, more frequent MHR accelerations and FHR decelerations. Further studies to assess if C-section is an independent risk factor for major MHR-FHR ambiguities are warranted.

To our knowledge, the effect of systematic removal of MHR-FHR ambiguities has not been previously assessed. In this study, removal of ambiguities resulted in a significant increase in FHR signal loss, along with a decrease in the number of FHR decelerations and an improvement in tracing classification (Tables 2 and 3). Again, the improvements were more often related with the misinterpretation of FHR decelerations than accelerations. If our results are confirmed in larger studies, it may be possible to decrease the unnecessary intervention associated with false positive yellow-red alarm in non-acidemic fetuses, without compromising sensitivity.

Several limitations need to be considered in this study and should be taken in consideration of future research. Firstly, a higher-risk population needs to be evaluated, as the number of cases with umbilical artery pH under 7.15 was low, and this 10th percentile cut-off value has limited clinical significance [22, 23]. Secondly, other methods for simultaneous MHR signal acquisition need to be evaluated, namely those using oximetry. Thirdly, the criterion used for removal of MHR-FHR ambiguities [1] may benefit from refinement after analysis of cases with adverse perinatal outcome, ideally using simultaneous internal FHR monitoring as the gold standard for fetal signals.

In conclusion, removal of MHR-FHR ambiguities resulted in a significant decrease in FHR decelerations and an improvement in tracing classification. Larger studies are needed to confirm the impact of this methodology on the diagnostic accuracy of intrapartum cardiotocography.

Competing interests

Diogo Ayres de Campos and João Bernardes have been involved in the development of Omniview-SisPorto system. Royalties are fully converted to institutional research funds.

Authors' contributions

All authors significantly contributed for the study conception, results interpretation, discussion and writing. PP was the main researcher and data collector, CS the main statistical analyser, HG the main signal pre-processor and processor, DAC the main reviewer of the text and JB the main coordinator, concept developer and final text co-reviewer. All authors read and approved the final manuscript.

Acknowledgments

Hernâni Gonçalves is financed by a post-doctoral grant (SFRH/BPD/69671/ 2010) from the Fundação para a Ciência e a Tecnologia (FCT), Portugal.

Author details

¹Hospital Nélio Mendonça, Funchal, Portugal. ²Departamento de Obstetrícia e Ginecologia, Faculdade de Medicina da Universidade do Porto, Porto, Portugal. ³Center for Health Technology and Services Research (CINTESIS), Porto, Portugal. ⁴Centro Hospitalar São João, Porto, Portugal. ⁵INEB – Institute of Biomedical Engineering; I3S – Institute for Research and Innovation in Health, University of Porto, Portugal. ⁶Unidade Local de Saúde de Matosinhos, Hospital Pedro Hispano, Matosinhos, Portugal.

Received: 20 November 2014 Accepted: 10 November 2015 Published online: 19 November 2015

References

- Reinhard J, Hayes-Gill BR, Schiermeier S, Hatzmann H, Heinrich TM, Louwen F. Intrapartum heart rate ambiguity: a comparison of cardiotocogram and abdominal fetal electrocardiogram with maternal electrocardiogram. Gynecol Obstet Invest. 2013;75:101–228 108.
- Odendaal HJ. False interpretation of fetal heart role monitoring in cases of intra uterine death. S Afr Med J. 1976;50:1963–5.
- Barrett JM, Boehm FH. Documentation of recent fetal demise with simultaneous maternal and fetal heart rate monitoring. Obstet Gynecol. 1980;55 Suppl 3:28S–30S.
- Achiron R, Zakut H. Misinterpretation of fetal heart rate monitoring in case of intrauterine death. Clin Exp Obstet Gynecol. 1984;11:126–9.
- Herman A, Ron-El R, Arieli S, Schreyer P, Caspi E. Maternal ECG recorded by internal monitoring closely mimicking fetal heart rate in a recent fetal death. Int J Gynaecol Obstet. 1990;33:269–71.
- Ramsey PS, Johnston BW, Welter VE, Ogburn Jr PL. Artifactual fetal electrocardiographic detection using internal monitoring following intrapartum fetal demise during VBAC trial. J Matern Fetal Med. 2000;9:360–1.
- Neilson Jr DR, Freeman RK, Mangan S. Signal ambiguity resulting in unexpected outcome with external fetal heart rate monitoring. Am J Obstet Gynecol. 2008;198:717–24.

- Muñoz Brands RM, Bakker PC, Bolte AC, van Geijn HP. Misidentification of maternal for fetal heart rate patterns after delivery of the first twin. J Perinat Med. 2009;37:177–9.
- Gonçalves H, Rocha AP, Ayres-de-Campos D, Bernardes J. Internal versus external intrapartum foetal heart rate monitoring: the effect on linear and nonlinear parameters. Physiol Meas. 2006;27:307–19.
- Nunes I, Ayres-de-Campos D, Costa-Santos C, Bernardes J. Differences between external and internal fetal heart rate monitoring during the second stage of labor: a prospective observational study. J Perinat Med. 2014;42:493–8.
- 11. RCOG. The use of electronic fetal monitoring. The use of cardiotocography in intrapartum fetal surveillance. Evidence-based clinical guideline number 8. *Clinical Effectiveness Support Unit.* London: RCOG Press; 2001.
- Nurani R, Chandraharan E, Lowe V, Ugwumadu A, Arulkumaran S. Misidentification of maternal heart rate as fetal on cardiotocography during the second stage of labor: the role of the fetal electrocardiograph. Acta Obstet Gynecol Scand. 2012;91:1428–32.
- Pinto P, Bernardes J, Costa-Santos C, Amorim-Costa C, Silva M, Ayres-de-Campos D. Development and evaluation of an algorithm for computer analysis of maternal heart rate during labor. Comput Biol Med. 2014;49:30–5.
- Ayres-de-Campos D, Sousa P, Costa A, Bernardes J. Omniview-SisPorto 3.5 a central fetal monitoring station with online alerts based on computerized cardiotocogram + ST event analysis. J Perinat Med. 2008;36:260–4.
- Ayres-de-Campos D, Bernardes J, FIGO Subcommittee. Twenty-five years after the FIGO guidelines for the use of fetal monitoring: time for a simplified approach? Int J Gynaecol Obstet. 2010;110:1–6.
- Ayres-de-Campos D, Bernardes J, Garrido A, Marques-de-Sá J, Pereira-Leite L. SisPorto 2.0 – a program for automated analysis of cardiotocograms. J Matern Fetal Med. 2000;9:311–8.
- Ayres-de-Campos D, Costa-Santos C, Bernardes J, SisPorto Multicentre Validation Study Group. Prediction of neonatal state by computer analysis of fetal heart rate tracings: the antepartum arm of the SisPorto multicentre validation study. Eur J Obstet Gynecol Reprod Biol. 2005;118:52–60.
- Costa MA, Ayres-de-Campos D, Machado AP, Santos CC, Bernardes J. Comparison of a computer system evaluation of intrapartum cardiotocographic events and a consensus of clinicians. J Perinat Med. 2010;38:191–5.
- Costa A, Ayres-de-Campos D, Costa F, Santos C, Bernardes J. Prediction of neonatal acidemia by computer analysis of fetal heart rate and ST event signals. Am J Obstet Gynecol. 2009;201:464. e1-464.e6.
- Bernardes J, Ayres-de-Campos D. Maternal heart rate analysis during labour. Has the time come for computerized analysis? Acta Obstet Gynecol Scand. 2012;91:1474.
- Sherman DJ, Frenkel E, Kurzweil Y, Padua A, Arieli S, Bahar M. Characteristics of maternal heart rate patterns during labor and delivery. Obstet Gynecol. 2002;99:542–7.
- Dudenhausen JW, Luhr C, Dimer JS. Umbilical artery blood gases in healthy term newborn infants. Int J Gynaecol Obstet. 1997;57:251–8.
- Kro GA, Yli BM, Rasmussen S, Norèn H, Amer-Wåhlin I, Rosén KG, et al. Association between umbilical cord artery pCO2 and the Apgar score; elevated levels of pCO2 may be beneficial for neonatal vitality after moderate acidemia. Acta Obstet Gynecol Scand. 2013;92:662–70.

Submit your next manuscript to BioMed Central and take full advantage of:

- Convenient online submission
- Thorough peer review
- No space constraints or color figure charges
- Immediate publication on acceptance
- Inclusion in PubMed, CAS, Scopus and Google Scholar
- Research which is freely available for redistribution

) BioMed Central

Submit your manuscript at www.biomedcentral.com/submit