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Data in Brief

journal homepage: www.elsevier.com/locate/dib



Annotation dataset of the cardiotocographic recordings constituting the "CTU-CHB intra-partum CTG database"



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ARTICLE INFO

Article history: Received 27 February 2020 Revised 9 April 2020 Accepted 4 May 2020 Available online 19 May 2020

Keywords: Cardiotocography Electronic fetal monitoring Fetal distress Fetal heart rate Uterine contractions Labor Childbirth

ABSTRACT

The proposed dataset provides annotations for the 552 cardiotocographic (CTG) recordings included in the publicly available "CTU-CHB intra-partum CTG database" from Physionet (https://physionet.org/content/ctu-uhb-ctgdb/1.0.0/). Each CTG recording is composed by two simultaneously acquired signals: i) the fetal heart rate (FHR) and ii) the maternal tocogram (representing uterine activity). Annotations consist in the detection of starting and ending points of specific CTG events on both FHR signal and maternal tocogram. Annotated events for the FHR signal are the bradycardia, tachycardia, acceleration and deceleration episodes. Annotated events for the maternal tocogram are the uterine contractions. The dataset also reports classification of each deceleration as early, late, variable or prolonged, in relation to the presence of a uterine contraction. Annotations were obtained by an expert gynecologist with the support of CTG Analyzer, a dedicated software application for automatic analysis of digital CTG recordings. These annotations can be useful in the development, testing and comparison of algorithms for the automatic analysis of digital CTG

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https://doi.org/10.1016/j.dib.2020.105690

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recordings, which can make CTG interpretation more objective and independent from clinician's experience.

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Specifications Table

Subject	Biomedical Engineering
Specific subject area	Biomedical signal processing, specifically digital cardiotocographic (CTG) recordings
Type of data	MATLAB® data file (.mat)
How data were acquired	Visual inspection by an expert gynecologist, supported by a dedicated software application.
	Instruments: CTG Analyzer, dedicated software application for the automatic analysis of digital CTG recordings
	(https://dx.doi.org/10.1109/EMBC.2017.8037391).
Data format	Raw and derived
Parameters for data collection	Annotations refer to a total of 552 CTG recordings, each composed by two simultaneously acquired signals: the fetal heart rate and the maternal tocogram (uterine activity). Annotations are based on the analysis of both
	signals and indicate starting and ending points of specific events such as bradycardia, tachycardia, accelerations, decelerations and uterine contractions.
Description of data collection	This dataset contains the annotations of the digital CTG recordings constituting the "CTU-CHB intra-partum CTG database" available at Physionet
	(http://physionet.org). Annotations were visually identified by an expert
	gynecologist supported by CTG Analyzer, a dedicated software application (https://dx.doi.org/10.1109/EMBC.2017.8037391).
Data source location	Raw CTG data (digital CTG recordings) were obtained from Physionet
	(https://physionet.org/content/ctu-uhb-ctgdb/1.0.0/).
	Derived CTG annotation data were obtained and are stored at the
	Cardiovascular Bioengineering Lab, Department of Information Engineering,
	Università Politecnica delle Marche, Ancona, Italy.
Data accessibility	Raw CTG data are publicly available at the following Physionet link:
	https://physionet.org/content/ctu-uhb-ctgdb/1.0.0/.
	Derived CTG annotations are available with the article as supplementary material.
Related research article	S. Romagnoli, A. Sbrollini, L. Burattini, I. Marcantoni, M. Morettini, L. Burattini,
	Digital cardiotocography: What is the optimal sampling frequency?, Biomed.
	Signal Process. Control. 51 (2019) 210–215.
	https://doi.org/10.1016/j.bspc.2019.02.016 [1]

Value of the Data

- This annotation dataset, together with the associated raw data ("CTU-CHB intra-partum CTG database") may represent a gold standard useful for studies on digital CTG.
- Beneficiaries of this dataset include biomedical engineers, who will have data to develop algorithms; clinicians, who will have computerized tools to support their diagnosis; pregnant women, who will receive objective diagnosis.
- This annotation dataset is useful to support development, testing and comparison of algorithms for the automatic analysis of digital CTG recordings, possibly also obtained by scanning paper CTG reports.
- The "CTU-CHB intra-partum CTG database" includes only tracings; by adding annotations, this dataset supports technical developments and thus passage from usual subjective CTG interpretation, mainly based on clinician's experience, to objective CTG interpretation.

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1. Data Description

1.1. Raw CTG Data

The raw data consist of 552 digital CTG recordings constituting the "CTU-CHB intra-partum CTG database" [2,3] and are publicly available in Physionet at the link: https://physionet.org/ content/ctu-uhb-ctgdb/1.0.0/. Each CTG recording is composed by two signals: i) the fetal heart rate (FHR, sometimes called tachogram); and ii) the maternal tocogram (representing the uterine activity).

1.2. Derived CTG Annotations

This dataset contains the annotations of the digital CTG recordings constituting the "CTU-CHB intra-partum CTG database" available at Physionet (http://physionet.org). Annotations were visually identified by an expert gynecologist supported by CTG Analyzer [4], a dedicated software application. They consist in the detection of starting and ending sample points of specific events on both FHR signal and maternal tocogram. Annotated events for the FHR signal are the bradycardia, tachycardia, acceleration and deceleration episodes. Annotated events for the maternal tocogram are the uterine contractions. Table 1 reports guidelines for identification of events on the CTG recording. The dataset also includes classification of each deceleration as early, late, variable or prolonged, in relation to the presence of a uterine contraction. Table 2 reports classification of the decelerations in relation to uterine contractions.

The dataset includes a total of 552 MATLAB® data file; each data file is called *annota-tion_i.mat* where *i* corresponds to the patient identifier in the "CTU-CHB intra-partum CTG database" [2,3]. Each *annotation_i.mat* includes two variables called *dataloss* and *ann*.

The *dataloss* variable $(1 \times 3$ vector) contains information about the quality of the FHR signal divided in 3 consecutive segments of equal length (see Experimental Design, Materials, and Methods section, Signal-Quality Assessment paragraph). Percentage of data loss, contained in the *dataloss* variable and automatically determined by CTG Analyzer [4] (Fig. 1 to Fig. 4), ranges from 0% to 100%. FHR-signal quality decreases with increasing percentage [1].

The *ann* variable ($5 \times L$ cell array, where L is the length of the CTG recording in samples) contains the annotations (i.e. strings indicating start and ending points; see Experimental Design, Materials, and Methods section, Annotations of CTG events and Decelerations Classification paragraph) for the 5 considered events: 1-bradycardia, 2-tachycardia, 3-acceleration, 4-deceleration

Table 1

|--|

Bradycardia	FHR baseline below 110 bpm for longer than 10 minutes
Tachycardia	FHR baseline over 160 bpm for longer than 10 minutes
Acceleration	FHR increments above baseline of more than 15 bpm lasting longer than 15 s
Deceleration	FHR decrements below baseline of more than 15 bpm lasting longer than 15 s
Uterine contraction	Bell-shaped temporary increment of the amplitude of the uterine activity signal lasting 45–120 s

Table 2

Classification of the decelerations in relation to uterine contractions.

Early	Deceleration onset synchronized with uterine-contraction onset; shallow short-lasting deceleration with normal variability within the deceleration
Late	Deceleration onset during the second half of uterine contraction; gradual onset and/or gradual return to the baseline and/or reduced variability within the deceleration
Variable	Deceleration uncorrelated to any uterine contraction; a rapid drop and good variability within the deceleration; variable in size and shape
Prolonged	Deceleration lasting more than 3 minutes



Fig. 1. Signal-quality assessment of a case (initial segment of signal '1016') with data loss equal to 38.5% and for which automatic annotation by CTG Analyzer was not performed.

Table 3

Symbols used for the annotations of the CTG events.

Bradycardia	Starting point of the k th bradycardia event	(BCk
	Ending point of the k th bradycardia event)BCk
Tachycardia	Starting point of the k th tachycardia event	(TCk
	Ending point of the k th tachycardia event)TCk
Acceleration	Starting point of the k th acceleration event	(ACCk
	Ending point of the k th acceleration event)ACCk
Deceleration	Starting point of the k th deceleration event, classification in	(DECkE
	relation to uterine contraction	(DECkL
		(DECkV
		(DECkP
	Ending point of the k th deceleration event, classification in)DECkE
	relation to uterine contraction)DECkL
)DECkV
)DECkP
Uterine contraction	Starting point of the k th uterine contraction	(UCk
	Ending point of the k th uterine contraction)UCk

and 5-uterine contraction. CTG annotations, contained in *ann* variable, were visually identified by an expert gynecologist supported by CTG Analyzer [4] (Fig. 2 to Fig. 4). Table 3 reports the symbols used in the proposed dataset for the annotations of the CTG events.

2. Experimental Design, Materials, and Methods

2.1. Data Acquisition

The raw CTG recordings constituting the "CTU-CHB intra-partum CTG database" were acquired from pregnant women during labor at the Czech Technical University in Prague (Czech





Fig. 2. Example (initial segment of signal '1317') of automatic bradycardia identification by CTG Analyzer.



Fig. 3. Example (initial segment of signal '1036') of automatic tachycardia identification by CTG Analyzer.

Republic) and at the University Hospital in Brno (Czech Republic) using, as electronic fetal monitors [3], the STAN S21 and S31 (Neoventa Medical, Mölndal, Sweden) and the Avalon FM40 and FM50 (Philips Healthcare, Andover, MA). CTG recordings have a duration ranging from 30 minutes to 90 minutes. Both FHR signal and maternal tocogram were sampled at 4 Hz [1,3]. These



Fig. 4. Example (initial segment of signal '1176') of automatic identification of accelerations, decelerations and uterine contractions and classification of decelerations by CTG Analyzer.

data, as all Physionet data, were fully anonymized and may be used without further approval of the Institutional Review Board.

2.2. Data Annotation

2.2.1. CTG Analyzer

CTG Analyzer [4] is a dedicated software application developed under MATLAB® GUI (Graphical User Interface) for the automatic analysis of digital CTG recordings obtained by digital recorders or by scanning paper CTG reports [5,6]. Details of CTG Analyzer can be found in [4]. Briefly, it provides a measure of the FHR signal quality and quantitative characterization of FHR signal and maternal tocogram according to FIGO (International Federation of Gynecology and Obstetrics) guidelines [7].

2.2.2. Signal-Quality Assessment

FHR signals are often affected by artifacts due to data loss represented by samples the value of which is set to zero. Since data loss may occur locally, FHR signals were divided into 3 consecutive segments of equal length (initial, middle and last segment) and data loss correction was performed in each segment independently by using the CTG Analyzer [1,4].

Quality of each FHR segment was assessed by computing the percentage of data loss, computed as 100 by the number of corrected samples over the total number of samples in the FHR segment. If FHR-segment quality was greater than 10%, the FHR segment was considered of bad quality [4,8] and no further automatic analysis by CTG Analyzer was performed; however, where possible, annotations were still obtained by visual inspection. Fig. 1 reports signal-quality assessment of a case with data loss equal to 38.5% and for which automatic annotation by CTG Analyzer was not performed. If FHR-segment quality was no more than 10%, linear interpolation was performed to compensate the data loss before automatic CTG events identification [4].

2.2.3. Annotations of CTG events and Decelerations Classification

Each CTG recording was annotated by an expert gynecologist by visual inspection and with the support of CTG Analyzer. All CTG events, that are bradycardia, tachycardia, acceleration, deceleration and uterine contraction, were identified according to their definitions provided by FIGO guidelines as reported in Table 1 [7,8] and indicated as BC, TC, ACC, DEC and UC, respectively. Classification of decelerations in early, late, variable and prolonged, was also performed according to the definitions of FIGO guidelines as reported in Table 2 [7] and deceleration class was indicated as E, L, V and P, respectively. Fig. 2 to Fig. 4 report examples of automatic CTG events identification and classification by CTG Analyzer. The starting sample point of an event is indicated as a string composed by an opening bracket '(', the acronym of the event, a progressive number and, if a deceleration, the deceleration class as reported in Table 3. Analogously, the ending sample point of an event is indicated as a string composed by a nopening as a string composed by a closing bracket ')', the acronym of the event, a progressive number and, if a deceleration class as reported in Table 3. When a sample does not represent a starting/ending point of an event, the cell has been left empty.

Declaration of Competing Interest

The authors declare that they have no known competing financial interests or personal relationships which have, or could be perceived to have, influenced the work reported in this article.

Supplementary materials

Supplementary material associated with this article can be found, in the online version, at doi:10.1016/j.dib.2020.105690.

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