

NONINVASIVE FETAL HEART RATE MONITORING: VALIDATION OF PHONOCARDIOGRAPHY-BASED FIBER-OPTIC SENSING AND ADAPTIVE FILTERING USING THE NLMS ALGORITHM

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Abstract. Here we present the evaluation results of our novel noninvasive phonocardiographic-based fiber-optic sensor for fetal Heart Rate (fHR) detection using adaptive filtering and the NLMS Algorithm. The sensor uses two interferometric probes encapsulated inside a PolyDiMethylSiloxane (PDMS) polymer. Based on real data acquired from pregnant women in a suitable research laboratory environment, once they had given their written informed consents, we created a simplified dynamic signal model of the distribution of maternal and fetal heart sounds inside the maternal body. Building upon this signal model, we verified the functionality of our novel fiber-optic sensor and its associated adaptive filtering system using the NLMS Algorithm. The main reason why we chose this technology to develop our system was that it allows monitoring the fHR without exposing the fetus to any external energies or radiation (in contrast to the ultrasound-based Cardiotocography Method). We used objective criteria such as: Signal to Noise Ratios: SNR_{in} , SNR_{out} and Percentage Root-mean-square Difference (PRD) for our evaluations.

Keywords

ElectroMagnetic Interference (EMI), fetal Heart Rate (fHR), fetal PhonoCardioGraphy (fPCG), Fiber-optic sensor, maternal PhonoCardioGraphy (mPCG), Normalized Least Mean Square (NLMS) algorithm, Poly-DiMethylSiloxane (PDMS).

1. Introduction

In this article, we report on the evaluation of a noninvasive method for fetal Heart Rate (fHR) detection and monitoring during gestation, labor, and delivery based on fetal PhonoCardioGraphy (fPCG). Our proposed method relies on the combined capabilities of fiber-optic sensing and adaptive filtering (implementing the Normalized Least Mean Square - NLMS - Algorithm). In our recent work reported elsewhere [1] and [2], we developed an adaptive system, which enabled

us to measure the fetal Heart Rate (fHR) by means of fPCG signal peak detection using the maternal abdominal PhonoCardioGrams (aPCGs). We observed that diagnostic-quality fPCG signals required for accurate fHR detection are contaminated by an unwanted maternal component (the mPCG signals) in addition to other technical and biological interferences. We showed that as the spectral contents of the fPCG and mPCG signals overlap in the frequency domain, common filtering methods such as signal subtraction, linear filtering, and others are ineffective in extracting reliable fHR information and therefore cannot be used.

Our recent research as well as others have also indicated that Fiber-optic technologies such as Fiber Bragg Gratings (FBGs) or interferometers are used increasingly in many biomedical applications; see articles [3], [4], [5], [6], [7], [8], [9], [10] and [11]. Building upon these advancements, we developed our novel sensor that uses two non-invasive interferometric probes encapsulated in a PolyDiMethylSiloxane (PDMS) polymer with the designation Sylgard 184.

The well-established conventional Phonocardiography is based on the scanning of acoustic signals by means of a microphone placed on the thorax. As for fetal Phonocardiography, the microphone is placed on the maternal abdomen [12], [13] and [14].

Our solution described here is based on the scanning of acoustic signals by means of two Mach-Zehnder interferometric fiber-optic probes. The advantages of these interferometers are their immunity to Electro-Magnetic Interferences (EMI), and their ability to measure any changes in the optical path length (such as the core refraction index, fiber length and the wavelength used). Therefore, the smallest measurable frequency due to any phenomena resulting in the change of the above-mentioned physical properties is theoretically unlimited [15] and [16].

To perform our system evaluations, we needed to use synthetic data. For generating suitable synthetic signals, we conducted a set of measurements on pregnant women in a suitable research laboratory environment after obtaining their written consents. We then created a simplified dynamic signal model for the distribution of maternal and fetal heart sounds inside the maternal body. Based upon this signal model, we generated synthetic data with properties as close as possible to the real data. The necessity to use synthetic data at this stage of our research was further justified by considering the fact that our patent-pending interferometric sensors have yet to be legislatively approved for clinical testing on pregnant women. It is important to emphasize that legislative regulations for use of new technology on pregnant women are extremely strict (as an unborn fetus is critically sensitive to external energies

such as mechanical pressure, electromagnetic radiation, change in temperature, and so on).

In current clinical practice, clinicians use either ultrasound-based methods such as CardioTocoGraphy (CTG), which measures the fetal heart rate along with maternal uterine contractions, or fetal Echocardiography (fECHO) to diagnose fetal congenital heart defects from the 20th to the 23rd week of pregnancy [17] and [18]. These sophisticated technologies are now integral parts of routine modern obstetrics. It is important to emphasize that the CTG technology has helped clinicians reduce the mortality rate of newborn babies during delivery. In spite of this considerable impact, it is generally recognized that this technology has some disadvantages such as high sensitivity to noise caused by maternal movements and the need to frequently reposition the ultrasound transducers. Also, this method is not suitable for long-term continuous fetal heart rate monitoring due to the potentially harmful influence of ultrasonic radiation on the fetus.

Our method and system, once statistically and clinically proven and validated, offer a number of advantages (in contrast to the currently used ultrasound-based CTG and other conventional methods), including their applicability to continuous long-term fHR monitoring without exposing the fetus to any radiation as well as their compatibility with Magnetic Resonance Imaging (MRI) environments. The continuous long-term monitoring capability of our system is highly desirable, especially in those cases in which the pregnant woman faces a dangerous situation (such as after an accident), and it becomes absolutely essential to perform a time consuming MRI examination to ensure that the unborn child is intact and safe. The other specific advantage of our technology is that it can be used in water deliveries.

2. Methods

2.1. Fetal Phonocardiography

Fetal PhonoCardioGraphy (fPCG) was discovered during the 17th century by Kergardec, Marsac, and Kennedy [19]. Although fPCG was discovered a very long time ago, interest in this research area has only grown over the past few years. This figure shows the number of peer-reviewed articles that appear in the Science Direct, the Institute of Electrical and Electronics Engineers (IEEE) and the National Institute of Health (PubMed) databases.

The PCG signal is composed of two main acoustic components (the first heart sound S1 and second heart sound S2), see Fig. 1, and two additional heart sounds (S3 and S4). S1 is systolic and is connected with the

closure of bicuspid and tricuspid valves at the beginning of ventricular contraction. S2 is diastolic and is produced by the closure of semilunar valves. The third heart sound (S3) is pro-diastolic and appears when a valve muscle quivers during the fast phase of blood flow into the valve. The fourth heart sound (S4) is pre-systolic and is a sign of the quivering of valve muscle during systole in the atrium. The last two mentioned heart sounds (S3 and S4) are not common for adults, and their presence is a sign of cardiac insufficiency [20].

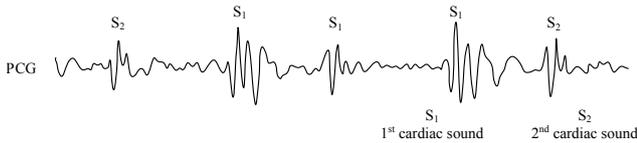


Fig. 1: Basic components of PCG signals.

2.2. PCG-Based Fibre-Optic Sensor

Our fiber-optic sensor is encapsulated inside polydimethylsiloxane [21], [22] and [23] and is comprised of two Mach-Zehnder interferometric components formed by 1×2 and 3×3 power couplers with an even split ratio; see Fig. 2.



Fig. 2: Our noninvasive fiber-optic measurement probe.

The reference fiber is stored in a stable environment. The output beams are recombined at a second 3×3 coupler. The output signal is detected by photodetectors. The resultant optical intensity after 3×3 coupler can be described by the following Eq. (1).

$$I_n = A_n + B_n \cos \left[\phi(t) + \phi_{drift}(t) + (n-1) \frac{2\pi}{3} \right], \quad (1)$$

where n represents the coupler output index with a value of 1, 2 or 3. The symbol A_n represents the mean value of optical intensity (DC component). Symbol B_n represents the optical intensity variation amplitude depending on fringe visibility, $\phi(t)$ represents the signal of interest, and $\phi_{drift}(t)$ is a quasi-static phase shift due to coupler properties. For the extraction of the

proper signal, it is necessary to use a demodulation algorithm [24].

2.3. Implementation of the Adaptive NLMS Algorithm

The measurands sensed by our interferometric sensors generated the fetal heart rate information, which was then fed into an adaptive stochastic system using the Root Mean Square Error (RMSE) criterion. This stochastic approach required a large number of measurements to produce powerful statistics. This consideration led to the utilization of the Normalized Least Mean Square (NLMS) Algorithm, which is a representative of basic stochastic gradient-based adaptation methods; see articles [25] and [26].

The Normalized Least Mean Square (NLMS) Algorithm is a variant of the Least Mean Square Algorithm. The former is able to accelerate the convergence speed with a reasonable computational cost and selects a normalized step-size μ_n , which results in both a stable and fast converging adaptation algorithm, see [27] and [28]. Implementation of the NLMS Algorithm can be summarized as follows:

BEGIN $\vec{w}(n=0) = \vec{0}$
FOR ($n=1, 2, \dots, N$):
 $y(n) = \vec{w}^T(n) \cdot \vec{x}(n)$
 $e(n) = d(n) - y(n)$
 $\vec{w}(n+1) = \vec{w}(n) + \mu(n) \cdot e(n) \cdot \vec{x}(n)$.

The step-size μ_n can be described as follows Eq. (2).

$$\mu(n) = \frac{\mu}{\delta + \vec{x}^T(n) \cdot \vec{x}(n)}. \quad (2)$$

Finally, we obtain the following Eq. (3).

$$\vec{w}(n) = \vec{w}(n-1) + \mu \frac{e(n) \cdot \vec{x}(n)}{\delta + \vec{x}^T(n) \cdot \vec{x}(n)}, \quad (3)$$

where $\mu \in (0, 2]$ and $\delta > 0$. Parameter δ represents the regularization parameter (prevents the denominator of Eq. (4) becoming zero).

3. Results

Our measurement system comprised of a novel fiber-optic sensor and its associated adaptive filtering system for fetal Heart Rate (fHR) monitoring is shown in Fig. 3. The adaptive system consists of two measurement sensors (FC/APC type) which were placed on the chest and abdomen, optical interrogator and DSP (Digital Signal Processing) unit for the recording,

amplification, digitalization, demodulation and filtering the measured signals. Optical interrogator consists of DFB (Distributed Feedback Laser) laser with wavelength 1549.5 nm and output power of 3 mW and three InGaAs Amplified Photodetectors (Indium Gallium Arsenide). Signal was digitalized by National Instruments card NI-USB 6210 with the sampling frequency of 250 kHz and analyzed by software application written in the LabView (2015, National Instruments, Austin, Texas, USA) [29] and [30].

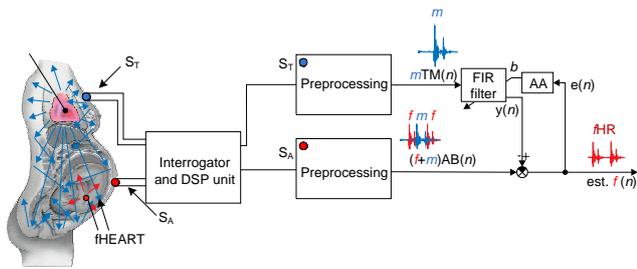


Fig. 3: Basic scheme of our sensor and its associated NLMS adaptive system for fHR monitoring.

Measurements (Fig. 4) were performed in a suitable research laboratory environment on 8 pregnant women (GA = 36–42 weeks) after obtaining their written informed consents. The test subjects were between the age of 21 and 37, their weight was between 57 kg and 103 kg, and their height was between 156 cm and 196 cm. Based on the obtained results we can state that no significant differences were found in the quality of the collected data based on the subjects’ age, weight, and height.



Fig. 4: An example of real data acquisition from a volunteer subject.

Using real data, we created a simplified dynamic model of sound distribution in the human body to generate suitable synthetic signals such as: ST (signals from sensors placed on the chest) and SA signals (from sensors placed on the abdomen). Our PCG signal model was inspired by contributions made by ALMASI et al. [31] and [32], who devoted considerable

efforts to generating synthetic PCG signals. In addition, we greatly benefited from our own research in generating realistic synthetic physiological and pathological fECG signals [33] and [34] in order to evaluate the performance of our system.

Figure 5 shows an ideal mPCG signal after removing the mother’s breathing artifacts (using a Butterworth second-order band-pass filter with corner frequencies: $f_L = 10$ Hz, and $f_H = 400$ Hz, respectively). This signal served as a reference input for our adaptive system running the NLMS Algorithm. The filtered results enabled us to determine the mHR (by performing mPCG signal peak detection). Maternal first and second heart sounds are denoted as mS_1 and mS_2 , respectively, in Fig. 5.

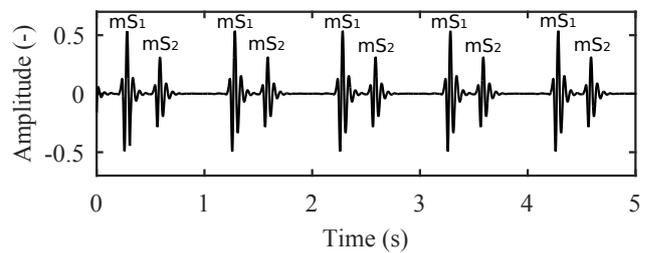


Fig. 5: The reference synthetic mPCG signal based on real measurements made from thoracic (S_T) sensors.

Figure 6 shows an ideal fPCG waveform after preprocessing the maternal signal. We need to emphasize here that the first fetal heart sounds (fS_1) result from the closing of the fetal tricuspid and mitral valves and the second fetal heart sounds (fS_2) are produced by the closure of the fetal pulmonic and aortic valves.

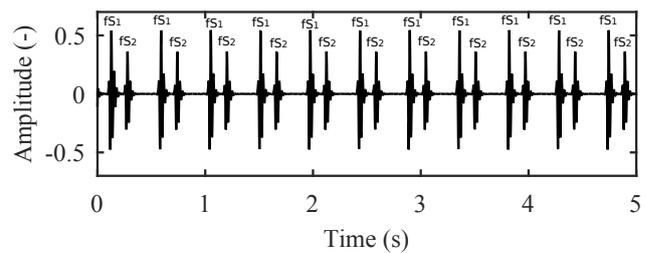


Fig. 6: The reference ideal synthetic fPCG signals based on real measurements from abdominal (S_A) sensors.

Figure 7 shows an example of the primary abdominal PCG (aPCG) synthetic input signal measured by the abdominal sensor. The aPCG signal (made up of the fPCG and mPCG components) is applied to the adaptive NLMS Algorithm. For determination of the fetal Heart Rate (fHR), it is necessary to detect fS_1 components in the composite aPCG signals, which is a difficult task without advanced signal processing.

Figure 8 shows an example of the output from our adaptive system using the NLMS Algorithm. Based on these results we can observe that: the mPCG compo-

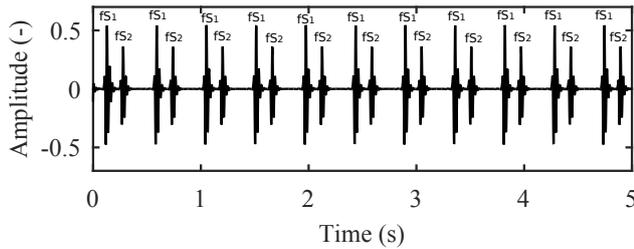


Fig. 7: The reference ideal synthetic fPCG signals based on real measurements from abdominal (S_A) sensors.

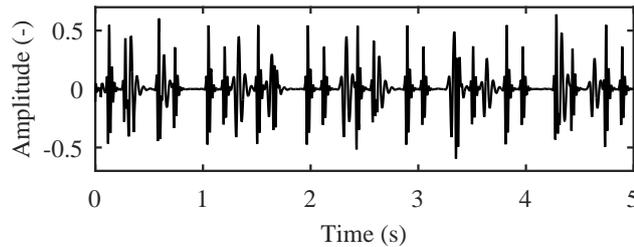


Fig. 8: Output of the adaptive NLMS system.

ment has been significantly reduced. This figure clearly shows that the elimination of the maternal component is not ideal; nevertheless, this component is reduced well under the level of fPCG signals. Using the filtered signal, we can use conventional techniques [35], [36] and [37] to determine the fHR information from the fPCG signals.

Table 1 summarizes our experimental results. The performance of our adaptive system using the NLMS Algorithm was evaluated by finding the differences between input (SNR_{in}) and output (SNR_{out}) values as well as the objective measure known as the Percentage Root-mean-square Difference (PRD) [38].

Tab. 1: Statistical results of the tested NLMS Algorithm.

SNR_{in} (dB)	SNR_{out} (dB)	PRD (%)
-7	0.98	14.61
-6	1.12	13.14
-5	1.33	10.74
-4	1.43	9.98
-3	1.48	9.35
-2	1.57	7.9
-1	1.65	6.74
0	1.71	5.69
1	1.70	5.74

The SNR_{in} value can be calculated by using the following Eq. (4):

$$SNR_{in} = 10\log \left(\frac{\sum_{n=1}^{N-1} [sig_{usef}(n)]^2}{\sum_{n=1}^{N-1} [sig_{noise}(n) - sig_{usef}(n)]^2} \right), \quad (4)$$

where $sig_{usef}(n)$ is a desired signal (modelled reference course of S_T) and $sig_{noise}(n)$ is a noise signal (mPCG is measured up in the abdominal part - S_A).

The SNR_{OUT} value can be calculated by using the following equation Eq. (5):

$$SNR_{out} = 10\log \left(\frac{\sum_{n=1}^{N-1} [sig_{des}(n)]^2}{\sum_{n=1}^{N-1} [sig_{pre}(n) - sig_{usef}(n)]^2} \right), \quad (5)$$

where $sig_{pre}(n)$ represents a predicted (estimated) signal, or more precisely, the output from the proposed NLMS adaptive system and $sig_{des}(n)$ represents the desired signal.

$$PRD (\%) = \left(\frac{\sum_{n=1}^N [sig_{usef}(n) - sig_{pre}(n)]^2}{\sum_{n=1}^N sig_{usef}(n)} \right) \cdot 100. \quad (6)$$

One way to quantify the difference between the reference and the output signal: $sig_{pre}(n)$ is by using the PRD as given by equation Eq. (6) below:

4. Conclusion

In this article we focused on the validation of our novel patent-pending interferometric PPG-based sensor and its associated adaptive filtering system using the NLMS Algorithm for effective processing of aPCG signals to extract fPCG signals and fHR information. In the evaluations of the signal filtering quality of our system, we used objective parameters such as SNR and PRD.

The main reason why we chose the fiber-optic technology to develop our system was that it enables fHR monitoring without exposing the fetus to any radiation (in contrast to the ultrasound-based CTG method). Our innovative system offers a number of advantages including applicability to continuous long-term fHR monitoring without exposing the fetus to any radiation as well as compatibility with Magnetic Resonance Imaging (MRI) environments. The long-term monitoring capacity of our system is highly desirable, especially in those cases when the pregnant woman faces a dangerous situation (such as after an accident), and it becomes absolutely necessary to perform a time consuming MRI examination to ensure that the unborn fetus is intact and safe. The other specific advantage of our technology is that it can be used in water deliveries.

In our future research, we intend to use data from clinical practice to investigate a variety of challenging

research topics such as the influence of sensor placement, fetal position and gestational age on aPCG signal filtering, fPCG signal extraction, and fHR monitoring.

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