

MASTER THESIS

Ultrasound Transducer Design for Continuous Fetal Heartbeat Monitoring

by Assel Rakhmetova

This work has been carried out at HSN, Department of Micro- and Nanosystem Technology, under the supervision of

Associate Professor Kristin Imenes

Vestfold, 4 July 2016

Submitted to

Faculty of Technology and Maritime Sciences, University of Southeast Norway, in partial fulfilment of the requirements for the degree Joint International Master in Smart Systems Integration (SSI)

Abstract

Stillbirth prevention requires high quality healthcare and early detection. Continuous monitoring of fetal heartbeat can be one of the ways to reduce pregnancy complications and even stillbirths. A Doppler ultrasound transducer is found to be one of the possible devices that can be adopted for home long-term monitoring of the fetal heartbeat.

Computer simulations and laboratory work were conducted to research a question of continuous monitoring of the fetal heart with the use of the Doppler transducer. Simulations were performed in MATLAB and Field II software environments. B-mode and M-mode images of the heart phantom were created with changed positions and amplitudes of the scatterers to imitate movements of the tissue structure. The received signals from the moving scatterers are then analyzed and time shifts are extracted. Transducer aperture was varied in order to increase transducer robustness for the application. Phantom position was changed in *x*- and *z*-directions. This allowed simulating the cases when either position of the transducer is displaced or fetus moves. Laboratory work was performed to create a test environment with a transmit/receive hardware system and a single-element transducer. LabVIEW graphical software was used to drive the hardware electronics. This test environment was performed to understand functioning of the transducer and the system in a real-time performance.

The velocity of the heart phantom was obtained from the time shifts between the consecutive received signals. It was possible to measure the heart velocity if the heart position is changed from the central lobe of the transducer beam in *x*-direction. However, the measurements are not accurate if the radius of the transducer aperture is decreased to 2 mm. The measured velocity of the heart phantom is in a good agreement with the actual velocity if heart is moved in *z*-direction. However, the received RF signals from the back heart wall are much weaker as compared to the received signals for the front heart wall.

The proposed idea to have a continuous fetal heartbeat monitoring is found to be one of the solutions to reduce number of stillbirths. Doppler transducer shows improved robustness with decreased size of the aperture which lead to a wider beam profile and better detectability of the heart movements.



Dedicated to my stillborn daughter named Aisha, who is always in my heart

Acknowledgements

I am very thankful to my supervisor Dr. Kristin Imenes, who supported my idea to propose a topic for a master thesis and who agreed to lead me all those months of the hard work. I am also very thankful to Prof. Lars Hoff and his ultrasound group who helped me to understand a new field for me –Acoustics.

I appreciate all my teachers in the Department of Micro- and Nanosystem Technology, who are the greatest and the kindest lecturers I had in my life.

I would like to express special thank you to Merete Hovet. She became my second mother during the whole SSI program. I express my sincere gratitude to Prof. Knut Aasmundtveit for his great support throughout all two years of the master program.

I am very thankful to midwives and doctors in Horten Kommune Helsestasjon, especially Hilde Haug for her time and guidance.

My sincere gratitude goes to my parents, sisters, friends and groupmates for their endless love and support.

Acronyms

2D	Two Dimensional
AFI	Amniotic Fluid Index
Bpm	Beats per minute
D	Diastole
DRM	Detection and Recording Module
FDA	Food and Drug Administration
LV	Left Ventricle
MI	Mechanical Index
Ν	Number of scatterers
PDM	Processing and Display Module
RF	Radio Frequency
RV	Right Ventricle
S	Systole
SFH	Symphysis-Fundal Height

Table of Contents

1	Int	rodu	iction	1
	1.1	Pro	oblem Statement	1
	1.2	Ob	jective of the Study	2
	1.3	Re	search Questions	2
	1.4	Th	esis Organization	3
2	Lit	erati	ure Background	4
	2.1	Pre	egnancy Evaluation Tests	4
	2.2	Pre	egnancy Evaluation Tests for Home Monitoring	9
	2.3	Fet	tal Heartbeat Monitoring Systems	12
3	Th	eore	tical Background	13
	3.1	Ph	ysics of Ultrasound	13
	3.1	.1	Reflection and Transmission	14
	3.1	.2	Scattering	15
	3.1	3	Attenuation	16
	3.2	Ult	rasound Transducers	17
	3.3	Ult	rasound Imaging	19
	3.4	Pri	nciples of Doppler Ultrasound	20
	3.4	.1	The Doppler Effect	20
	3.4	.2	Continuous-wave and Pulse-wave Doppler	21
	3.5	Ult	rasound Safety	22
	3.5	5.1	Thermal Effects	22
	3.5	5.2	Non-thermal effects	23
4	Me	ethod	lology	24
	4.1	MA	ATLAB and FIELD II Summary	24
	4.2	Fet	tus and Pregnant Woman Design Parameters	26
	4.3	Sin	nulation Design	29
	4.3	8.1	Ultrasound Transducer Design Model	29
	4.3	8.2	Heart Model	30
	4.3	3.3	Simulation model	33
	4.4	Lal	boratory work	

5	Res	ults	40
	5.1	B-mode Heart Image	40
	5.2	Case studies	41
	5.2	1 Case study A	
	5.2	2 Case study B	
	5.2	3 Case Study C	54
	5.2	4 Case study D	61
	5.3	Laboratory work	64
6	Dis	cussion	67
	6.1	Simulations	
	6.1	1 Case study A and B	
	6.1	2 Case study C and D	70
	6.2	Laboratory work	72
	6.3	Ultrasound safety	72
7	Cor	clusion And Future Work	
G	lossar	7	74
A	ppend	ix A: MATLAB code for a Heart Phantom Image	75
A	ppend	ix B: MATLAB codes used for Case Study A, B, C & D	80
A	ppend	ix C: MATLAB code for M-Mode Image	
A	ppend	ix D: Piston Transducer Beam Profile	
Δ.	nnend	iv F: Flat I ARVIFW 'Process' State	83
<u>л</u>	ppenu	IX E. Flat LAD VIE VV TIOCESS State	
R	eferen	Ces	
Li	st of F	igures	92
Li	st of T	able	

1 INTRODUCTION

1.1 Problem Statement

Many women and children still die during the pregnancy. It is estimated that there are 303 000 maternal deaths and 2.6 millions of stillbirths in the world registered in 2015. Every day 7 300 women suffer the loss of their babies in the last three months of gestation [1]. Stillbirth is a baby born with no signs of life after 28 weeks of pregnancy[2]. There are more than 3600 cases of stillbirths every year in the UK. Eleven babies die every day and one in every 200 labours ends up in a stillbirth [3].

Stillbirth prevention requires high quality healthcare, early detection and diagnosis. It is not affordable in many countries and especially not in developing low-income countries. The majority of stillbirths are preventable and it mostly depends on the access to maternal healthcare and immediate diagnosis of possible complications.

Various pregnancy-monitoring tests are usually performed to control normal and high-risk pregnancy in women. Sometimes, additional tests of fetal well-being may be needed to predict and to prevent serious pregnancy complications and even stillbirth. However, none of the traditional tests have a definitely proven effect when it comes to decreasing intrauterine stillbirths [4].

Moreover, for women with no signs of problems during the pregnancy, such tests are not usually performed. Not detecting pregnancy complications can lead to critical incidents and, for women in their first pregnancy, even sudden fetus death can occur. Also, these tests require the woman to come to hospital twice a week (or more) which can be inconvenient for some women who have health problems. Perhaps most importantly, the fetus is not observed in between the tests which brings difficulties to catch any fetal disorders.

There is no sole strategy to predict a stillbirth [5]. For example, if a pregnant woman feels less baby movements inside the uterus, the non-stress test and ultrasound related tests will be performed to assess fetus health at the moment [6], [7]. The fetal can behave healthy at the moment of the screening. However, the well-being of the fetus can change later once woman has left the hospital. One of the reasons for a stillbirth can

be an infection. This includes bacterial infections (Escherichia coli, streptococcus), viruses (parvovirus B19) or parasites (toxoplasmosis, malaria) [5]. However, if a woman complains about the decrease of the baby movements, blood, urine and vaginal tests are not performed to diagnose for the presence of any infections. Existing Doppler instruments for heartbeat listening that can be used by woman at home to listen to a fetal heartbeat cannot be used as medical devices to evaluate any fetal abnormalities. Moreover, these instruments can be used improperly leading to false measurements [8]. They are rather used by midwives as a psychological tool to instill future mothers with an idea of having a baby in the near future.

1.2 Objective of the Study

The objective of the study is to investigate the possibility of having a continuous monitoring system to record a fetal heartbeat during the third gestation trimester to prevent stillbirth. The objective may be fulfilled by designing an ultrasound transducer with a suitable analysis system that can be attached to a woman abdomen to continuously monitor a fetal heartbeat.

1.3 Research Questions

The diagnostic ultrasound is a noninvasive and safe technique used in clinical practice. It has also advantages of low cost and portability. The future of ultrasound systems to be miniature, mobile and portable is improving and promising.

A fetal heartbeat continuous monitoring system can be implemented by a wellestablished technique called Doppler ultrasound. The question arises whether traditional Doppler ultrasound techniques can be adopted for continuous evaluation of the fetal heartbeat and be used at home for long term monitoring. Will it always be possible to catch the fetal heartbeat during the long term screening if a transducer is attached on the female abdomen?

Another issue is whether continuous heartbeat monitoring can help to reduce stillbirth accidents. Can we rely on the heartbeat as a reflection of the fetus well-being status?

It is also important to investigate whether there are any bio-effects of the ultrasound if a fetus is exposed to it throughout a day. Will it be safe for a baby and the mother?

1.4 Thesis Organization

Thesis consists of literature background, theoretical overview, methodology, results, discussion, conclusion and future work chapters. Literature background introduces current pregnancy evaluation tests that available in hospitals and on the market. It also discusses their limitations and drawbacks. Theoretical background gives an overview of the ultrasound physics that is used to build a single-element transducer and to analyze Doppler ultrasound theory. Methodology describes case studies design parameters that were created to investigate problem stepwise. Results and discussion chapters introduce main theoretical and experimental results and discuss their outcomes. Conclusion and future work chapter summarizes the work performed and proposes the future improvements and development of the topic.

2 LITERATURE BACKGROUND

2.1 Pregnancy Evaluation Tests

Usually, pregnant woman does not need to have any extra check-ups during her pregnancy unless there are pre-existing health conditions which may cause miscarriage and even stillbirth.

The possible reasons causing stillbirth include[2],[3], [5], [9]:

- childbirth complications
- infections during the pregnancy
- maternal disorders (hypertension, diabetes)
- fetal growth restrictions
- congenital abnormalities

It is important to perform the examination of fetus well-being in high risk pregnancies for women who have [10],[11]:

- complications in the previous pregnancies or stillbirth
- existing health conditions such as diabetes, heart disease
- complications during pregnancy such as fetal growth problems, placenta abnormalities
- decreased fetal movement, fetal hypoxia
- prolonged pregnancy (beyond 40 weeks)

Certain evaluation tests may be used to assess fetal well-being which are performed in hospital if woman experiences less fetal movements or she has above mentioned health conditions [4], [12]:

- fetal movement assessment
- non-stress test including uterine contraction test
- amniotic fluid index
- contraction stress test
- biophysical profile
- umbilical artery Doppler

Fetal movement assessment [13] is a routine method used to monitor normal pregnancy. It is also a method that can be used in high risk pregnancy. Fetal movement assessment is a method when woman counts the number of kicks or movements of a baby during the day. Usually woman can feel the baby's movement from 20-24 weeks of gestation. According to several studies, there should be at least 10 distinct baby movements in 11-34 minutes. If a woman feels or counts a decreased number of movements then she must take more tests to evaluate fetus well-being. However, sometimes woman may feel anxious while counting the movements, which may be destructive to accomplish the test. Also, some woman can be busy at work or with other children during the day and they are not able to monitor baby's activity [12], [14]. Pregnant women have to select two hours during a day when the baby is the most active to count the fetal movements [13].

Non-stress test (cardiotocography) [12][15] – is a non-invasive test widely used in obstetrics. The principle of the test is the association of the fetal heart rate with fetal movements. Healthy babies will have an increase in the heart rate while moving, and a reduction in the heart rate while at rest.

Figure 2-1 illustrates the basic set-up of the monitoring system usually installed for non-stress test. The set-up consists of the ultrasound system, tokodynamometer and auxiliary electronics. The ultrasound system utilizes Doppler ultrasound (Doppler effect) and records the heartbeat of the fetus. Tokodynamometer [16] is a pressure transducer placed around a woman's abdomen with an elastic belt. When the uterine muscles contract, they raise the abdominal wall depressing the plunger and intrauterine pressure is measured.



Figure 2-1: *Fetal monitoring system. An ultrasound transducer measures fetal heartbeat while a tokodynamometer measures uterine activity* [17].

It requires 20-40 minutes to complete the test. The test is performed as frequent as required for high-risk pregnancies. Usually, pregnant women are asked to eat before the test as it can make the baby move more actively. During the test, the woman lies on her side while the ultrasound system and the tokodynamometer are attached around woman's abdomen by elastic belts. The woman is asked to press a button when she feels a movement. There are two results of the test: reactive and non-reactive. Reactive means that the baby's heart rate increases as the baby moves. There should be an increase of the fetal heart rate of 15 bpm from the baseline for 15 seconds occurring two or more times during a 20 or 30 minutes period in conjunction with fetal movement. Non-reactive means the baby's heart rate does not increase as the baby moves. *Figure 2-2* shows an example of a non-stress test result provided by the "Sykehuset i Vestfold" hospital.



Figure 2-2: Non-stress test shows normal fetal heart rate accelerations.

Amniotic fluid index (AFI) [12], [18] - is an ultrasound measurement technique of amniotic fluid volume. The uterus is divided into four quadrants and the largest vertical amniotic fluid pocket in each quadrant is measured. AFI is the sum of these four measurements. In the third trimester AFI is between 8 and 25 cm.

Olygohydraminos is defined for AFI less than 5 cm and associated with increase of caesarean section for fetal distress.

Polyhydramios is defined for AFI which is greater than 25 cm and associated with an increase of perinatal mortality rate, fetal abnormalities, increased caesarean ection rate.

Contraction stress test [19]- is an invasive procedure which is performed to evaluate the fetus response to the stimulated uterine contraction. As the uterine contracts, the changes in the baby's heart rate are recorded.

In a healthy baby, cardiorespiratory reserves are adequate to tolerate the decreased or interrupted intravillous blood flow of the placenta as the uterine contracts. In the fetus with inadequate fetal cardiorespiratory reserves, uterine contractions may not be tolerated and fetus heart rate has late deceleration.

There are some contraindications to perform this test. Since it is an invasive techniques possibility of a preterm labour and gestational age should be considered before taking the test.

Biophysical profile [20] - is a test which consists of five components: fetal movement, fetal breathing, fetal tone, amniotic fluid volume and non-stress test. The presence of each components is scored with a value of 2 if present and zero if response is found not adequate. Biophysical profile method is the overall method to identify fetus condition.

Umbilical artery Doppler evaluation [12], [21], [22]- is a widely used test to analyse uteroplacental blood flow. This test is used to evaluate various abnormalities such as placenta abnormalities, intrauterine growth retardation for prolonged pregnancies, fetal growth restriction. Doppler systems produce flow velocity waveforms that reflect the distribution and intensity of the Doppler frequency shifts over time. The frequency shifts are proportional to changes in the flow velocity within the umbilical vessels.

The result of the Doppler evaluation is given by the S/D ratio which is a ratio of peak velocity of systolic velocity waveform to the nadir to a diastole. By 30 weeks of pregnancy, the S/D ratio should be less than 3.0. Other methods of reporting the Doppler evaluation results are pulsatility index and resistance index.

Pulsatility index is given by the systolic minus the diastolic values divided by the mean of the velocity waveform profile (S-D/mean).

The resistance index is given by S-D/S. *Figure 2-3* illustrates the ultrasound results obtained after the test.



Figure 2-3: *Umbilical Doppler velocimetry. Normal umbilical artery blood flow as seen with a forward flow in diastole and normal S/D ratio* [12].

However, these tests are only performed in hospitals. Woman with no signs of problems and pre-existing chronic conditions will usually not be examined with the mentioned tests.

Home monitoring system can be one of the aids that woman may have at home and which can help her to monitor fetal well-being if she feels worried. It also could help to collect more data to evaluate fetal well-being and to seek immediate medical help if something goes wrong to avoid complications and stillbirth.

Table 2-1 summarizes drawbacks of the pregnancy evaluation tests used in hospitals in assessment of the fetal well-being.

Examination Tests	Drawbacks
Fetal movement assessment	Not reliable, difficult to judge whether the changes in the observed fetal activity signify good or adverse outcomes of pregnancy [23].
Non-stress test	The position of the ultrasound transducer should be always adjusted [24]. Interpretation of the test can be misleading: high rate of false-positive results [25], [26]. Results are interpreted manually, possibility for a human error [27].

 Table 2-1: Examinations tests and their drawbacks for fetal well-being assessment.

AFI test	Cannot be used as a stand-alone test to assess fetal status. Amniotic fluid volume measurements are still not precise [28].
Contraction stress test	High rate of ambiguous results requiring repeat testing, expensive, undesirable for labour/uterine contraction [29].
Umbilical artery Doppler evaluation	There is a potential of variability and inaccuracy, measurements should be performed by experts, who are able to determine significance of the Doppler changes [30].

According to the studies, number of fetal movements and fetal heart rate can be considered as the first indicators of the fetus well-being [31], [32]. *Table 2-2* summarizes the indicators of the fetal status that can be gained from fetal movements and fetal heart rate assessments.

Assessment	Healthy	Non-healthy	
Number of fetal movements	>10 movements within 1 hour [12]	Less than 10 movements within 2 hours [13]	
Fetal heart rate	Reactive [12], with average 140 bpm [33]	Non-reactive [12], fetal bradycardia - < 120 bpm [33], fetal tachycardia - > 200 bpm [34]	

 Table 2-2: Fetal well-being assessment parameters.

2.2 Pregnancy Evaluation Tests for Home Monitoring

Movement count is one of the first tests that can be the sign weather a baby is healthy or not. The woman is required to count the number of kicks to evaluate fetus well-being. However, sometimes it can be difficult to catch the movements. Also, many women may have insufficient time to count movements due to work or being busy, with other children. Kick counter wristband are now available on the market to help women to count kicks. The basic principle of the wristband is when woman feels a baby movement she should move the counter of the wristband in order to track and count the kicks [35], [36]. *Figure 2-4* shows the wristband that may be used daily by pregnant woman.



Figure 2-4: Kick counter wristband [37].

Kickme-Baby Kicks Counter [38] - is an android mobile application which is used as a dairy to keep track of the baby movements. The appearance of the application is shown in *Figure 2-5*. The woman should press "KICKED NOW" or "KICKES EARLIER" button if she feels the movement. This application creates a statistics of the most active hour, most active day and kicks per day.

HOME	LOGS	STATS	MORE	HOME	LOGS	STATS	MORE	HOME	LOGS	STATS
	6 KICKS IN	05:21:59			SESSIO	N HISTORY		2011	5-08-13 17:33 TO	2015-08-13 21:
KICK	DATE		TIME	STAF	π	DURATION	KICKS	киск	DATE	тімі
1	2015 AUGUST	09	16:09:00	2015-08-0	4 12:25	09:01:03	10	1	2015 AUGUST 1	13 17:33
2	2015 AUGUST	09	17:05:07	2015-08-0	5 14:35	06:52:01	9	2	2015 AUGUST	3 17:48:
3	2015 AUGUST	09	18:11:12	2015-08-0	6 13:49	07:38:04	10	3	2015 AUGUST 1	3 18:24:
4	2015 AUGUST	09	19:36:18	2015-08-0	7 15:43	05:45:22	8	4	2015 AUGUST 1	19:07:
5	2015 AUGUST	09	20.03.24	2015-08-0	8 14:49	05:40:02	9	5	2015 AUGUST 1	13 19:39.
6	2015 AUGUST	09	20.45-31	2015-08-0	9 16:09	05:22:20	7	6	2015 AUGUST 1	3 20.47
RICKED	KICKED	ADD	STOP	SESSION	DAILY	MONTHLY	EMAIL	ADJUST	ADJUST	EDIT D

Figure 2-5: *Kickme- baby kicks counter Android mobile application* [38].

There are many various Doppler heart listening devices available on the market. *Figure 2-6* shows an example of the device. These devices allow listening and recording the sound of the fetus heartbeat. They are not for medical purpose but for entertainment mostly.



Figure 2-6: Fetal Doppler heart sound monitor device displaying fetal heart rate [39].

This fetal Doppler monitor should point directly at the fetal heart location in order to be able to listen to a heartbeat.

Baby's Heartbeat Listener [40] is a mobile application used for entertainment purpose to listen to the heart beat of the fetus. The phone can be applied onto the woman's tummy and the heartbeat can be listened through the headphones and recorded. *Figure 2-7* shows the appearance of the application for a mobile phone.



Figure 2-7: Baby's heartbeat listener mobile application [40].

These applications and devices are mostly aimed for entertainment purposes. They cannot be used to adequately assess the fetal well-being. Improper use of domestic fetal monitors may mislead a woman. There were cases when women did not seek medical help when they noticed less movements of their babies after listening to the heartbeats using home Doppler devices [8], [41]. Therefore, it is important to have more reliable devices for fetus well-being assessment to be used for medical purposes with remote hospital control.

2.3 Fetal Heartbeat Monitoring Systems

There are published papers available that described possible solutions for fetal heartbeat monitoring and sending the results to mobile phone.

S. Bhong and S. D. Lokhande [42] proposed a wireless fetal monitoring system for home use. In the proposed system, a mobile software application transforms existing fetal monitoring devices (Doppler ultrasound transducer and tokodynamometer) into one system that evaluates the fetal heart rate and uterine contractions, while saving and converting the data to a hospital standard.

A. K. Mittra and N. K. Choudhari [43] developed a low cost fetal heart sound monitoring system for home care application. The system consists of two parts which are Detection and Recording Module (DRM) and Processing and Display Module (PDM). DRM is a hardware placed on a woman abdomen used to detect and record fetal heartbeat. PDM is software which is aimed to record, save and generate results. DRM consists of acoustic cone, microphone, piezoelectric sensor, power amplifiers, and filters.

Nowadays, wireless fetal monitoring is approved by Food and Drug Administration (FDA) [44], [45]. Doppler ultrasound transducer and tokodynamometer can now be fixed around the women's abdomen using belts. The data are transmitted wirelessly from the device to a recorder. However, this still requires the woman to be present at hospital. It was also researched that the data can be sent via Bluetooth for the hospital evaluation [46].

3 THEORETICAL BACKGROUND

3.1 Physics of Ultrasound

Ultrasound [47], [48] is a mechanical vibration of matter with a frequency greater than 20 kHz. The acoustic particle is introduced to understand the concept of a wave propagating through the tissue. This particle is assumed to be a small volume element. The wave is propagating through the tissue as a disturbance of the particles in the medium. Initially, particles are at rest and spaced uniformly. With the presence of the ultrasound wave, the particles start to oscillate. The important types of waves are plane, spherical and cylindrical waves. In the current work, plane longitudinal wave propagation is assumed. A plane wave travels in one direction. In longitudinal wave propagation, the displacement of the particle is parallel to the propagation to the wave. The propagation speed, c depends on the medium and is given as [49]:

$$c = \sqrt{\frac{1}{\rho_0 \kappa}},\tag{3.1}$$

where ho_0 is the equilibrium density and κ is the adiabatic compressibility.

During the plane wave propagation, the acoustic particles which lie on the plane normal to the direction of propagation will undergo to the same incremental pressure change. Assuming that the wave propagation is linear, the acoustic pressure, p of the plane harmonic wave propagating in z-direction is given as:

$$p(t,z) = p_0 exp(j(\omega t - kz)), \qquad (3.2)$$

where ω is the angular frequency of the wave, $k = \omega/c$ and is the wave number and p_0 is the acoustic pressure amplitude, and *j* is the imaginary unit.

The assumption that waves obey the linearity principle means that they keep the same shape as they change amplitude and scaled versions of waves at the same location can be combined to form more complicated waves [50].

For a plane propagating wave, the particle speed, u is related to the pressure p by the acoustic impedance Z as [49]:

$$u = \frac{p}{Z} \quad , \tag{3.3}$$

where *Z* is the characteristic acoustic impedance, a material constant, equal to the product of density, ρ and the speed of sound, *c* [49]:

$$Z = \rho c. \tag{3.4}$$

The unit for characteristic impedance, *Z* is Rayls, 1 Rayl equals kg/m^2s .

The characteristic acoustic impedance is an important term for characterizing the plane wave propagation. *Table 3-1* shows the values of the sound speeds, density and characteristic impedance in different mediums and in human tissue.

Table 3-1: Speeds of sound, densities and characteristic impedances data for differentmediums and human tissues [48].

Medium	Medium Density [kg/m ³]		Characteristic acoustic impedance [kg/m²s]
Air	1.2	333	0.4×10^{3}
Blood	1.06×10 ³	1566	1.66×10 ⁶
Bone	1.38-1.81×10 ³	2070-5350	3.75-7.38×10 ⁶
Fat	0.92×10 ³	1446	1.33×10 ⁶
Muscle	1.07×10 ³	1542-1626	$1.65 - 11.74 \times 10^{6}$
Distilled water	1.00×10 ³	1480	1.48×10 ⁶

3.1.1 Reflection and Transmission

An incident wave propagating in a medium with the wavenumber, k_1 and the characteristic impedance, Z_1 will be partially transmitted and reflected at the boundary

of a different medium with the wavenumber, k_2 and the characteristic impedance, Z_2 . Tissues are usually modelled as liquids [47], [51].



Figure 3-1: The model of the one-dimensional wave propagation hitting the boundary.

For the normal incidence, the reflection coefficient, *RF* is derived as [49]:

$$RF = \frac{Z_2 - Z_1}{Z_1 + Z_2} \tag{3.5}$$

In case of oblique incidence as depicted in *Figure 3-1* where longitudinal plane wave is incident at an angle on a boundary, the reflection coefficient is given as [49]:

$$RF = \frac{Z_2 cos\theta_i - Z_1 cos\theta_t}{Z_2 cos\theta_i + Z_1 cos\theta_t}$$
(3.6)

The transmission wave is a wave that continues to propagate further through the medium. The transmission coefficient, *T* for this wave is:

$$T = \frac{2Z_2 \cos\theta_i}{Z_2 \cos\theta_i + Z_1 \cos\theta_t} \tag{3.7}$$

3.1.2 Scattering

The ability of the ultrasonic wave to penetrate the matter is due to two acoustic parameters: absorption and scattering. Absorption is the quantity of the ultrasound energy that is transformed into heat, chemical energy and light. Scattering is the radiation of all, or part of the energy in an ultrasonic wave when incident on an obstacle. The scattering can be in any direction. Reflection and refraction can be considered as special cases of the scattering [51], [52].

Backscattering [52] is a reflection of the waves back to the direction from which they were originated. Backscattering is useful for ultrasound imaging. Pulse echo technique is used to detect the backscattered signal. Ultrasound transducers transmit a pulse into a specimen to investigate, for example a heart. First, the transducer receives the echo from the front face of the specimen and later from the back face. Other echoes that are produced in between the two surfaces depend on the structural composition of the tissue there.

The signal power, P_s generated by one scatterer is characterized by the scattering cross section which is a measure of the scattering magnitude and is derived as [48]:

$$P_s = I_i \sigma_{sc} , \qquad (3.8)$$

where σ_{sc} is a scattering cross section and I_i is a uniform intensity. The backscattering cross section depends on the material type and denotes the strength of the material scattering [48].

Provided that energy is scattered uniformly in all directions, the scattered intensity, I_s is then given as [48]:

$$I_{s} = \frac{P_{s}}{4\pi R^{2}} = \frac{\sigma_{sc}}{4\pi R^{2}} I_{i},$$
(3.9)

where *R* is the distance to the scattering region. The received power, P_r with a transducer radius, *r* is then [48]:

$$P_r = I_s \pi r^2 = \sigma_{sc} \frac{r^2}{4R^2} I_i, \tag{3.10}$$

Therefore, the received power depends on the scattering cross section, emitted intensity, distance to the transducer and the size of the transducer aperture.

3.1.3 Attenuation

The ultrasound wave propagating in the tissue will experience energy loss or attenuation due to absorption and scattering (reflection and refraction). Attenuation can be expressed with exponential law as functions of distance. The amplitude loss term, A(z, t) can be added for single frequency plane wave propagation [53]:

$$A(z,t) = A_0 \exp(i(\omega_c t - kz)) \exp(-\alpha z), \qquad (3.11)$$

where ω_c is the angular centre frequency and α is the attenuation factor.

The amplitude attenuation coefficient, α is the sum of the scattering and absorption amplitude attenuation coefficients. The attenuation coefficient is expressed in nepers per meters, but in medical ultrasound commonly given in decibel per centimetre (dB/cm).

The attenuation that happens in tissue depends on the frequency. Attenuation increases as frequency increases. The centre frequency, f_0 of a transmitted signal will be shifted down in frequency as the emitted pulse propagates through the tissue. The shift in frequency of a Gaussian pulse is given as [48]:

$$p(t) = \exp(-2(B_r f_0 \pi)^2 t^2) \cos(2\pi f_0 t)$$
(3.12)

where B_r is a relative bandwidth and f_0 is a centre frequency.

3.2 Ultrasound Transducers

Ultrasound transducers are widely used in diagnostic imaging systems. Transducers come in a wide range of sizes and shapes and are often designed to operate around specific centre frequencies. The piezoelectric transducer, depicted in *Figure 3-2* is a single element transducer that consists of a piezoelectric plate, a mechanical damping material, one or several matching layers and a casing. The ultrasound transducer is connected to the medium by a physical contact. To increase the contact between the transducer and the medium an ultrasound couplant can be used, e.g. a gel. The piezoelectric plate is an important element in the transducer as this is where the acoustical waves are generated and received. The ultrasound waves are generated by the inverse piezoelectric effect, i.e. when the electric field is applied to the plate, the plate responds to the corresponding stress field by contracting or expanding, depending on the polarisation of the field [54]. In reception, when the ultrasound waves are incident on the plate, the plate generates an electric potential in response to the applied stress field. This is referred to as the piezoelectric effect. The choice of the dimensions of the piezoelectric plate depends on the transducer's application. The beam shape is to

large extent given by the ratio between the transducer diameter, *D* relative to ultrasonic wavelength, λ . To achieve a directive beam, *D* should be much larger than the wavelength. This can be estimated from the opening angle in the far field, $\vartheta_0 = \lambda/D$, for example $D = 10\lambda$ will give an opening angle of approximately 5⁰. Aperture diameters close to λ , for example 0.7 λ will lead to a much wider transmitted beam [54].



Figure 3-2: The schematic of the single element transducer [54].

The mechanical damping material is located behind the piezoelectric element and is used to prevent excessive vibrations and to generate signals with shorter pulse length that provides better spatial resolution in imaging. For a good damping, the backing material should be matched with the piezoelectric acoustic impedance and have high attenuation to eliminate reverberation within itself [54].

Often a large difference in the acoustical impedances of the media and the piezoelectric material exist. For example, a common piezoceramic from Meggit Ferroperm, Pz27 has an acoustical impedance of approximately 33 MRayl, whereas the human body has an acoustical impedance of approximately 1.5 MRayl; a ratio of 22 [55]. A consequence of this is that most of the acoustical energy will be reflected at the boundary between the piezoceramic and the human body. To reduce the reflection and increase the transduction at the boundary between the transducer and the human body, one or several matching layers are used. The matching layers form part of the transducer construction and the impedance of the matching layer (for a transducer with one matching layer) are chosen as the geometric mean between the impedance of the

piezoceramic and the media, and the thickness of the matching layer is chosen as onequarter of the wavelength in the matching layer. Matching a transducer to the medium like this helps the ultrasonic waves to propagate efficiently into the object.

Single element transducer is a transducer which is widely used for underwater sonar systems, A single element transducer offers no control over the transmitted ultrasonic beam. However, controlling the beam of a transducer is invaluable in e.g. medical imaging and underwater sonar systems where the beam can be focused at a finite spatial region. For this kind of applications transducer arrays are used with various beamforming techniques. Generally, the array elements are activated at different times, resulting in the beams from the individual elements reaching a specific point at the same time. A piezoelectric array transducer, consisting of a number of piezoelectric elements, is shown in *Figure 3-3*.



Figure 3-3: The schematic of a piezoelectric array transducer [54].

3.3 Ultrasound Imaging

Ultrasound imaging is based on the pulse-echo principle. The ultrasound pulse is emitted from the transducer and directed into the tissue. The echoes are generated due to the outgoing pulses interacting with tissue. The three major modes used in imaging system are A-mode, B-mode and M-mode [51], [56].

A-mode is an amplitude mode. Once the echoes are received from the object, they are amplified and displayed as amplitude versus time record, similar to an oscilloscope screen [50].

B-mode is known as a brightness mode. Brightness is proportional to the echo amplitude. In a B-mode, the image is composed of many beams aimed in different directions, creating a 2D image, typically in the zx-plane, where z refers to depth and x to lateral direction. The brightness is related to the echo amplitude [51].

M-mode is a motion mode used to visualize time variation. The vertical *z* axis is depth downwards and horizontal *t* axis is time. The images look similar to B-mode, but the lateral dimension is time, creating 2D *zt*-image. This mode is particular useful to monitor heart motion and to receive an "image" of heart valves by observing distinct patterns of heart along the time [56], [50].

3.4 Principles of Doppler Ultrasound

Doppler ultrasound is a non-invasive technique which is widely used to monitor and measure the blood flow in the human body. It is sometimes difficult to visualize blood circulation and Doppler ultrasound is a good solution to detect its movement in the vessels. Doppler effect is also applied for heart valves imaging to track heart contractions. It provides comprehensive information about fluid flow and heart dynamics and abnormalities [50].

3.4.1 The Doppler Effect

The Doppler effect is a change in frequency as a sound source moves toward or away from an observer. The Doppler frequency, f_D or observed frequency for one-way moving source is given as [57]:

$$f_D = \frac{f_0}{1 - (c_s/c_0)\cos\theta'}$$
(3.13)

where f_0 is a transmitted frequency, c_s is the velocity of the source, c_0 is the speed of sound and θ is an angle between the observer location and the source vector. From the equation 3.13, Doppler-shifted frequency is then found as [57]:

$$\Delta f = f_D - f_0 = f_0 \left(\frac{c_s}{c_0}\right) \cos\theta.$$
(3.14)

The observed frequency depends on θ which means it is important to take care of the direction of the sound relative to the observer in terms of vectors [57]. *Figure 3-4* shows Doppler-shifted wave frequencies as seen by the observers at four locations and

the angles relative to the direction of the source, where Δf is a Doppler frequency and f_s is a source frequency. Observers at B and D do not hear any Doppler shift.



Figure 3-4: Doppler frequencies seen by observers at different location and at the angles relative to the direction of the source: (A) 0°, (B) 90°, (C) 270°, (D) 45° [57].

If the observer is moving and the source is stationary, then the formula for Doppler-shifted frequency is [57]:

$$\Delta f = [1 + (v_{obs}/c_0)\cos\theta]f_0, \qquad (3.15)$$

where v_{obs} is the velocity of the observer. The Doppler-shifted frequency for the case when both the source and the observer are moving turns to [57]:

$$\Delta f = f_0 [c_0 + c_{obs} \cos\theta] / [c_0 - c_s \cos\theta], \qquad (3.16)$$

where c_s is a velocity of the source.

3.4.2 Continuous-wave and Pulse-wave Doppler

In continuous-wave Doppler mode, the transducer is divided into halves where one part continuously transmits and another part continuously receives signals [57]. This mode records the velocities of all moving targets in the ultrasound beam [58]. It allows measurement of high velocities. In pulse-wave mode, transducer transmits and then receives a signal after a pre-set time delay. One sample is acquired for each transmitted pulse. The sampling area can be moved along the path of the ultrasound beam for examination [58]. If the tissue is stationary then constant sampled values will result. If the scatterer is moving then the pulse will move past the depth of sampling and will be sampled due to its motion [49]. *Figure 3-5* shows an example of the simulated received signals from blood vessel acquired for each transmitted pulse. It is seen how the scatterers move away from the transducer and the received signals are shifted in relation to each other. The dotted line indicates the time instance when the sampling takes place, and the sampled signal is then plotted.



Figure 3-5: *Sampled signal from blood vessel. The left graph shows the received signals for each transmitted pulse, and the right graph is the sampled signal* [49].

3.5 Ultrasound Safety

There are thermal and mechanical interactions of ultrasound with tissue during ultrasound exposure. Most of the ultrasound side effects are reversible. Thermal and non-thermal bio-effects are defined to occur during clinical usage of ultrasound.

3.5.1 Thermal Effects

During the ultrasound exposure, human tissue may experience a temperature rise. Based on the statistical data, empirical relationship of the temperature rise and exposure time, *t* below which there are no side effects of ultrasound was derived [59]:

$$t = 4^{43-T}$$
 (minutes), (3.18)

where *T* is temperature in Celsius. *Table 3-2* shows the effects of the temperature elevated due to ultrasound on a human body [59].

Temperature range [ºC]	Effect		
37-39	No harm for extended exposure period		
39-43	Adverse effect after long exposure time		
>41	Threshold for fetal problems for long period		
>41.8	Damage threshold		
44-46	Protein coagulation		

 Table 3-2: Temperature effects induced by ultrasound on a human body [59].

The fetus is considered as a sensitive biological site for long term ultrasound exposure especially during the first trimester. If the transducer is placed directly on the fetus skull, the elevated temperature of the bone may damage brain tissue. The exposure of the ultrasound onto a fetus above 41 °C for 5 minutes or more should be considered unsafe. According to a equation 3.18, the rise of temperature to 1.5°C corresponds to 158 minutes and a rise to 4°C corresponds to 5 minutes provided that transducer is located unmoved during the specified time [59].

3.5.2 Non-thermal Effects

Cavitation is a major type of the non-thermal effect of ultrasound [59]. Cavitation is a collapsing of gas bubbles due to ultrasound exposure. This non-thermal effect of ultrasound hardly occurs in human body naturally. For example, during imaging of the fetus it does not show signs of any damage cases due to cavitation. Microbubbles that are located in the intestine may cause small local damage. However, damages due to cavitation are reversible and heal completely [59]. MI is a mechanical index used to estimate a degree of the bio-effects due to cavitation that ultrasound may induce. MI is defined as the maximum value of the peak negative pressure over the square root of the centre frequency [60]. According to FDA, the ultrasound system is considered safe if MI index does not exceed a maximum of 1.9 [61].

4 METHODOLOGY

Simulation is a method that extensively used in biomedical ultrasound. This thesis uses computational model or computer based simulation to investigate the stated problem. The simulation is based on the quantitative calculations and mathematical model to determine numeric behaviour of the acoustic field in a test environment. MATLAB R2016a version and Field II software programs were chosen to perform simulations. Laboratory work was the second part of the research. Transmit/receive hardware system and test environment were set up to couple a single-element piezoelectric transducer. LABVIEW 2015 software was used to acquire and process the signals from the hardware system that used to analyse the received signals from the test environment.

There also were few discussions with the midwives in Horten Kommune Helsestasjon who helped to understand the check-ups procedures for pregnant women and available techniques. The continuous fetal heartbeat monitoring system as a solution to a stillbirth reduction and fetal well-being monitoring was discussed with them.

4.1 MATLAB and FIELD II Summary

Field II [62], [63], [64] is a simulation program for ultrasound systems which can be interfaced with MATLAB. It uses the spatial impulse concept based on Tupholme-Stepanishen method. The field of any kind of excitation is found by the convolution of the spatial impulse response and excitation function. The excitation pulse is a voltage applied to the transducer terminals. The impulse response is called spatial impulse response since it varies as a function of the position relative to the transducer.

Field II allows the simulations of any kind of transducer geometries and excitation. Pulsed-wave and continuous-wave fields can be calculated for transmit and pulse-echo. Apodization and various focusing schemes can be utilized though introduced time lines. The main application of Field II is to simulate an image. It is also applied for the simulation of the velocity measurements with ultrasound using Doppler effect. It is mostly applied for the measurement of the blood flow in the vessels.

There are some approximations made in Field II:

- Spatial impulse response assumes linear propagation;
- Transducer surface is divided into smaller squares, called 'mathematical elements' to imitate piston vibrations. The edges in piston may vibrate less than the centre. The responses from each of the squares are then summed up to produce a response;
- The sound field from each mathematical element is calculated using the far field approximation for the element to make calculation fast. The sound field from the whole aperture is found by summing the contributions from the mathematical elements. These results will be valid also in the near field of the whole aperture.

Point scatterer approach is used to construct a phantom. The ultrasound image consists of a clutter and larger tissue structures. Clutter is a noise artifact in ultrasound images that obscures the image target and complicates anatomical measurements [65]. The image is then built from the collection of the randomly placed point scatterers with Gaussian amplitude. The relative amplitude between the different types of scatterers is scaled to generate clutter and larger structures [66]. The fully developed realistic image should have at least 10 randomly generated scatterers per resolution cell. Usually, 200 000 to 1 million scatterers are generated to yield a normal ultrasound image.

The received signals from the point scatterers are then calculated for each line in the image defined by focusing scheme. The resulted RF signal is then found by the summing the received signals from the scatterers. RF signal is an adopted term from the communication engineering field and it states for "radio frequency". This term is accepted and used in ultrasound field. The received RF signal is a voltage output signal of the beamformer [67].

It is possible to simulate a Doppler ultrasound scanning in Field II. The Doppler frequency shift can be determined through the estimation of the time difference between the successive snap shots of images. Different images are created with changed positions or amplitudes of the scatterers to imitate movements of the tissue structure. The received signals from moving scatterers are then analyzed and frequency shifts are extracted.

4.2 Fetus and Pregnant Woman Design Parameters

It is important to investigate the anatomy of the abdomen of the pregnant woman for the transducer to be attached. The third trimester of gestation is chosen for the study with a woman having one fetus. During the third trimester, the baby does not make big movements and eventually stays at one position with a head down [68]. The abdomen of a pregnant woman consists of the abdominal wall, uterine wall and amniotic fluid surrounding a fetus [68], as depicted in *Figure 4-1*. Since baby is big enough to occupy uterus during the most of the third trimester period, the thickness of the amniotic fluid is neglected for the simulations.



Figure 4-1: Anatomy of the woman in the third trimester of gestation.

Uterine wall consists of three layers such as endometrium, myometrium and perimetrium. Myometrium is the major constituent of the uterine wall [69]. The abdominal wall consists of a subcutaneous tissue, muscle and skin [70], [71].

It is important to define woman weight which will be considered for the study. The weight of the woman affects the thickness of the abdominal and uterine walls. These thicknesses will define the distance of the transducer to a fetal heart. *Table 4-1* shows the parameters used to perform simulation which were taken from the studies [69], [72].

Variable	Range
Maternal age [years]	21-44
Maternal weight [kg]	42-110
	Mean, (range)
Abdominal wall depth [mm]	38, (9-92)

 Table 4-1: Woman size parameters.

Abdominal wall depth is measured from the abdominal wall surface to the anterior wall of the amniotic sac. Posterior uterine wall depth is defined as the distance from the abdominal wall surface to the posterior uterine wall surface. *Figure 4-2* shows the schematics of the woman abdomen.



Figure 4-2: Schematic drawing of the woman abdomen, 1 – abdominal wall depth, 2 uterine wall thickness, 3 – posterior uterine wall depth.

Posterior uterine wall depth was calculated from the symphysis-fundal height (SFH) measurements. For simplicity of calculations, it was assumed that uterine has a circular shape. SFH is a height of the uterus which changes according to a gestation age [73], depicted in *Figure 4-3*. It is measured from the top of the uterus to the pubic bone as shown in *Figure 4-4*.



Figure 4-3: *SFH measurements versus gestational age, provided by midwives from Horten kommune helsetjenesten for barn og unge.*

SFH is approximated to behave as a circumference of a circle, *C* and it equals to:

$$C = 2SFH, \tag{4.1}$$

and posterior uterine wall depth, *D* is then derived as:

$$D = \frac{2SFH}{\pi},\tag{4.2}$$

Table 4-2 shows SFH measurements extracted from the *Figure 4-3* and calculated posterior uterine all depth for the third gestation trimester.

 Table 4-2: Posterior uterine wall depth and SFH change according to a gestation age.

Gestation age [weeks]	SFH [cm]	D [cm]
28-31	26.0-28.5	16.6-18.1
32-35	29.5-32.0	18.9-20.4
36-40	32.5-34.5	20.7-22.0



Figure 4-4: Schematic of SFH measurement procedure.

The heart wall consists of the three major layers such as endocardium, myocardium and epicardium surrounded by pericardium sac. The total thickness of these layers is around 1-2 mm for a fetal heart [74], [75]. The heart sizes change during the gestational age progression. *Table 4-3* shows the cardiac sizes for the third trimester which are assumed for the simulation.

Gestation age [weeks]	Cardiac circumference [mm]	Cardiac radius [mm]
28-31	114-129	18-20
32-35	134-148	21-23
36-40	152-169	24-26

 Table 4-3: Cardiac sizes for the third trimester [76], [77].

4.3 Simulation Design

4.3.1 Ultrasound Transducer Design Model

Single-element piston transducer is used to perform simulations in Field II for Doppler ultrasound. The circular plate disk is divided into mathematical elements, as
shown in *Figure 4-5*. The size of each element is 1 mm. The transducer is always positioned at (0, 0, 0) coordinates.



Figure 4-5: Piston transducer divided into 1mm mathematical elements.

Transducer aperture radius is the design parameter for the transducer that will be varied for simulations to investigate the problem. *Table 4-4* summarizes the design parameters used during the simulations.

Parameters	Value
Central frequency	2.0 MHz
Transdu aan na diwa	5 mm
Transducer radius	2 mm

Table 4-4: Transducer design parameters.

4.3.2 Heart Model

Ultrasound image and schematic of the fetal heart are shown in *Figure 4-6*. Based on these images, the fetal heart is assumed to have a circular shape for 2-D B-mode image construction in Field II.



Figure 4-6: Ultrasound image of the fetal heart (right) and its schematics with identified heart constituents: LV (left ventricle), RV (right ventricle), ventricular septum, moderator band, pulmonary veins, atrial septum and crux [78].

The heart model consists of the heart with a heart wall filled with blood and a surrounding tissue. Two radii, r_1 and r_2 are introduced to define the thickness of the heart wall. *Figure 4-7* illustrates a proposed heart model. Heart phantom is then constructed by the generation of random point scatterers and deterministic scaled amplitudes. The point scatterers are given the amplitude properties of tissue or blood. The blood cells are mainly responsible for the scattering when ultrasound interacts with blood. Scattering is very weak from the blood cells since blood cells have very small micrometre sizes as compared to a heart wall muscle tissue. Therefore, it is assumed that the amplitude of point scatterers inside the heart is zero. Heart tissue is a highly scattered region and its amplitude is set to 10. The background tissue is responsible to simulate the realistic surrounding environment around the fetal heart. The number of background point scatterers is a varied parameter to simulate different background conditions. Increasing the amplitude and number of scatterers will increase their scattering properties which will make difficult to distinguish the heart wall boundaries in a "noisy" environment.



Figure 4-7: Heart model for Field II simulation, r1 and r2 are heart wall radii.

Table 4-5 summarizes the amplitude properties of the point scatterers for blood, heart muscle and background tissue used for simulation.

Objects	Amplitude scaling factor
Blood	0
Heart wall	10
Surrounding tissue	1

 Table 4-5: Amplitude scaling factors for blood, heart wall and surrounding tissue.

The simulated B-mode images of the heart were received in Field II. B-mode images were simulated with linear array transducer with 192 elements. These images are used to view the heart phantom and its position relative to the transducer surface. Number of scatterers was varied to receive different granular textures. *Table 4-6* summarizes the number of simulations performed for this part.

Simulation No.	Number of point scatterers
1	1 000
2	10 000
3	200 000
4	1 000 000

 Table 4-6: Simulation of the heart phantom with varied number of scatterers.

4.3.3 Simulation Model

The heart movements or beatings were imitated by the changing the heart radius. The radius change was assumed to behave as a sinusoid, as depicted in *Figure 4-8*.



Figure 4-8: Heart movement assumption used for simulation.

The radii of the heart wall were changed imitating the heart beating. *Figure 4-9* shows the flow chart of the simulation design.



Figure 4-9: *The flow chart of the simulation model, where r1 and r2 are the heart wall radii, R is a radius of the transducer aperture, f0 is the central frequency.*

The received signal is recorded once the heart wall radii change. One cardiac cycle was simulated. RF signals are analytically analysed after they were recorded. The simulation is divided into four case studies which are case study A, B, C and D. Case study A is a case where the heart phantom is a highly scattered region and the background tissue is a weakly scattered region. In case study B, the surrounding tissue is now highly scattered region and heart is a weak scattered region. Case studies C and D analyse the situation when the heart position due to baby movements or transducer location can be displaced. The heart position is slightly shifted in *x*-direction in Case study C. The phantom is then shifted in *z*-direction in case study D. The design parameters are further specified in below subsections.

4.3.3.1 Case Study A

The first simulation named as Case study A is performed in a least harsh environment. The heart phantom is generated by points positioned in a box of scatterers within 100 mm× 30 mm× 140 mm region starting from the 35 mm from the transducer surface. In this case heart wall is a highly scattered region as compared to a surrounding tissue region. It is achieved by assigning scaled amplitudes. The gestational age of 28-31 weeks is chosen with a posterior uterine wall depth, *D* of 16.6-18.1 cm or 17.4 cm in average. This posterior uterine wall depth includes abdominal wall thickness. The radius of the outer heart wall is 18 mm and inner radius is 16 mm which creates 2 mm heart wall thickness. The position of the heart is then assumed to be located approximately in the middle of the posterior uterine wall depth (*Figure 4-10*). The design parameters are summarized in *Table 4-7*.



Figure 4-10: Transducer placement assumption for simulation of Case study A.

Fetus and	Gestational age Outer heart radiu [weeks] [mm]		Transducer-heart distance [mm]
woman	28-31	18	87
Transducer	Centre frequency, f_0 Aperture radius [MHz] [mm]		-
	2	5	-
Phantom	Heart amplitude	Blood amplitude	Background amplitude
	10	0	1

 Table 4-7: Simulation design parameters for Case study A.

One cardiac cycle which includes systole and diastole is simulated with a heart rate of 140 bpm or 2.33 Hz of the heart frequency.

4.3.3.2 Case Study B

The design parameters used in the case study B are shown in *Table 4-8*. In this study, heart muscle is a weakly scattered region and background tissue is a highly scattered region. The amplitude of the heart muscle is now scaled to 1, and the amplitude of the background is scaled to 10. Other design parameters remain unchanged from the case study A.

Fetus and	Gestational age (weeks)	Heart radius [mm]	Transducer-heart distance [mm]
woman	28-31	18	87
Transducer	Centre frequency, f_0 [MHz]	ntre frequency, <i>f</i> ₀ Aperture radius [MHz] [mm]	
	2	5	-
Phantom Heart amplitude		Blood amplitude	Background amplitude
	1	0	10

 Table 4-8: Design parameters.

4.3.3.3 Case Study C

The heart centre position is slightly moved in *x*-direction. The previous centre of the heart (x_c , z_c) was (0, 70). The changed position in this case is (10, 70). The scatterer box is widened and it has dimensions of 140 mm× 30 mm× 140 mm. The design parameters are summarized in *Table 4-9*.

Fetus and	Gestational age (weeks)	Heart radius [mm]	Transducer-heart distance [mm]
woman	28-31	18	87
Transducer	Centre frequency, <i>f</i> ₀ [MHz]	Aperture radius [mm]	
	2	5	2
Phantom	Heart amplitude	Blood amplitude	Background amplitude
	1	0	10

 Table 4-9: Design parameters for case study C.



Figure 4-11: *Possible movements of the transducer and a fetus assumed in simulation for case study C and D.*

4.3.3.4 Case Study D

For this simulation, the heart position is then placed further away from the transducer surface. This simulation is also used to analyse the situation if the fetus is moved in *z*-direction away from the transducer or posterior uterine wall depth is thicker. The heart centre coordinates are changed from (0, 70) to (0, 90). Phantom is then placed 107 mm away from the transducer surface. Dimension of the scatterer box is changed to 140 mm× 30 mm× 160 mm. Design parameters are summarized in *Table 4-10*.

Fetus and	Gestational age (weeks)	Heart radius [mm]	Transducer-heart distance [mm]
woman	28-31	18	107
Transducer	Centre frequency, f_0 [MHz]	Aperture radius [mm]	
	2	5	-
Phantom	Heart amplitude	Blood amplitude	Background amplitude
	1	0	10

 Table 4-10: Design parameters for case study D.

4.4 Laboratory Work

Laboratory work was conducted to better analyse the stated problem. Transmit/receive hardware system was used to perform tests. The schematic of the system was designed by the ultrasound group of the Department of Micro- and Nanosystem Technology. The electronic hardware then was soldered and constructed according to the given schematic. This hardware system was designed to:

- generate high electrical pulses to be supplied to the piezoelectric transducer
- receive electric signals from the piezoelectric transducer and then amplify them
- filter the received signals
- convert from analog to digital representation
- send data to a computer for signal processing

More detailed explanation of the system performance can be obtained from the ultrasound group in the Department of Micro- and Nanosystem Technology.

LabVIEW 2015 is a system-design platform utilizes the graphical language. It was used to acquire and process the signals received from the transmit/receive hardware system. The LabVIEW template to drive the hardware system was supplied by the ultrasound group. The code for the "Process" state was only implemented for required application which is to compute the phase shifts between the received signals.

5 Results

5.1 B-mode Heart Image

B-mode images were simulated with varied number of randomly generated point scatterers *(Figure 5-1).* This allows modelling a speckle. Speckle is a grainy texture arises from the constructive and destructive interference of these scatterers [79]. The simulated phantom can be used for Doppler shift simulations.



Figure 5-1: *B-mode images of the phantom with varied number of scatterers (N): (a) N*=1000, (b) *N*=10 000, (c) 200 000 and (d) *N*=1 000 000.

B-mode image with number of scatterers $N=200\ 000$ and $1\ 000\ 000$ are statistically equal. It was required 18 hours to build the heart phantom with one million scatterers as compared to the image with 200 000 scatterers which took 4 hours. B-mode image with N equal to 200 000 is fully developed speckle and these number of scatterers will be used to construct phantom for Doppler ultrasound.

5.2 Case Studies

Theoretical analysis is needed in order to analyse the received RF signals. Doppler frequency shift is generated by cardiac motion. The velocity of the heart depends on a time instance when it is recorded since heart experiences accelerations and decelerations *(Figure 5-2)*. The displacement of the heart during the cardiac cycle is considered to be a sinusoid:

$$r(t) = 18 + \sin(2\pi f_h t), \tag{5.1}$$

where f_h is a frequency of the heartbeat. The velocity is found as a derivative of the radius change over time the heart beats:

$$v = \frac{dr(t)}{dt} = \frac{d}{dt}(18 + \sin(2\pi f_h t)) = 2\pi f_h \cos(2\pi f_h t)$$
(5.2)



Figure 5-2: The velocity of the heart movement for one cardiac cycle.

The maximum velocity, v_{max} of the heart motion is achieved when $\cos(2\pi f_h t) = 1$ and it equals to:

$$v_{max} = 2\pi f_h \tag{5.3}$$

The frequency of the heartbeat is calculated from the heart rate data. *Table 5-1* summarizes the heart rate and corresponding frequency of the heartbeat and maximum velocity data.

Heart rate [bpm]	<i>f</i> _h [Hz]	$v_{max}[m mm/s]$
80	1.33	8.36
100	1.67	10.49
120	2.00	12.57
140	2.33	14.64
160	2.67	16.78
180	3.00	18.85

Table 5-1: Heart rate, frequency of the heartbeat, fh and corresponding maximum heartvelocity, vmax data.

The emitted pulse consists of the sinusoidal oscillations [48]:

$$e(t) = g(t)\sin(2\pi f_0 t),$$
(5.4)

where g(t) is the envelope of the pulse and is given as:

$$g(t) = \begin{cases} 1, 0 < t < \frac{M}{f_0}, \\ 0, & else \end{cases}$$
(5.5)

M stands for number of emitted cycles and f_0 is a centre frequency.

The received signal is then found as [48]:

$$r_{s}(t) = a \cdot g(\alpha t - t_{0}) \sin(2\pi f_{0}\alpha(t - t_{0})), \qquad (5.6)$$

where α is a compression factor, t_0 is the time between pulse emission to reception and αf_0 is a frequency of the received signal. The compression factor, α is approximated to be [48]:

$$\alpha = 1 - \frac{2v_{heart}}{c},\tag{5.7}$$

where v_{heart} is a velocity of the heart motion and c is a speed of sound in tissue and equals to 1540 m/s.

The Doppler frequency, f_d is the difference of the transmitted and received frequency and found as [57]:

$$f_d = \alpha f_0 - f_0 = (\alpha - 1)f_0 = -\frac{2|v_{heart}|cos\theta}{c}f_0,$$
(5.8)

where θ is an angle between the ultrasound beam and the velocity vector.

Assuming that the velocity vector and ultrasound beam are at 0^{0} lead to $cos\theta$ = 1. The Doppler frequencies for heartbeat frequencies of 1.67, 2.33 and 3 Hz are calculated for the centre frequency of 2 MHz, as shown in *Table 5-2*.

Centre frequency, f_0 [MHz]	<i>f</i> _{<i>h</i>} [Hz]	v _{max} [mm/s]	<i>f</i> _{<i>d</i>} [Hz]
2.00	1.67	10.49	-27.24
	2.33	14.64	-38.03
	3.00	18.85	-48.96

 Table 5-2: Calculated Doppler frequencies for 2 MHz centre frequency.

The task of true and simulated Doppler instrument is to be able to detect such small frequency shifts. It is mostly impossible to detect small shifts with pulsed Doppler since the downshift in frequency due to attenuation will dominate over the Doppler frequency shift. Another method to analyse the Doppler system is a computation of the time shifts between the consecutive received pulses. Two consecutive received signals are compared. The time between the transmit pulses is T_{PRF} . The movement of the heart scatterers will yield a small displacement in their positions which can be countered as a shift in time relative to the pulse shift. The second received signal (r_{s2}) will be shifted in time as compare to the first received signals (r_{s1}) as [48]:

$$r_{s2} = r_{s1}(t - t_s) , (5.9)$$

where t_s is a time shift. The velocity can be estimated by measuring the distance travelled during a certain time interval. The mean velocity is then the distance travelled divided by the time. The time displacement or time shift, t_s between the successive received signals is [57]:

$$t_s = \frac{2\Delta z}{c} = \frac{2T_{PRF}v_{heart}cos\theta}{c},$$
(5.10)

where Δz is an imaging depth. Therefore, the time shift is proportional to the velocity of the heart movement. The equation 5.10 can be rewritten as:

$$t_s = A v_{heart}, \tag{5.11}$$

where A is then a constant and equals to:

$$A = \frac{2T_{PRF}cos\theta}{c}.$$
(5.12)

Therefore instead of measurement of the frequency shifts, phase shift measurement can be employed by estimating the time delay of the received signals due to the displacement of the scatterers.

5.2.1 Case Study A

Figure 5-3 shows B-mode image of the heart located at 8.7 cm from the transducer surface. The image shows that the heart wall is highly scattered and a surrounding tissue is a weakly scattered region.



Figure 5-3: *B-mode image of the heart phantom located at 87mm from the transducer surface.*

The beam profile and heart position in the pressure field can be seen in the *Figure 5-4* with a piston transducer having 5 mm aperture.



Figure 5-4: Beam profile for a round flat aperture with 5 mm radius with a heart position in the pressure field for case study A.

The time, t_d for the first transmitted and received signal by a transducer should be:

$$t_d = \frac{2\Delta z}{c} = \frac{2(87mm)}{1540m/s} = 1.13 \times 10^{-4}s.$$

First, signals were received from the stationary structure when the radius of the heart does not change. *Figure 5-5* shows the summed RF signals and *Figure 5-6* shows individual RF lines for each transmitted pulses. Delay time, t_d for the first received signal is found from the *Figure 5-5* and it is equal to $1.217 \times 10^{-4}s$ which is slightly different from the computed value. Transmitted pulse is reflected from the front and back wall of the heart, as seen in *Figure 5-5*. The amplitudes of the received signals are much lower for the back heart wall as compared to the amplitudes of the received signals from the front heart wall. The reason for it may be the position of the back heart wall which is located in a pressure field with a lower intensity. The blood region is observed where no RF lines are received. There are no time shifts observed between the received signals since the phantom is stationary (*Figure 5-6*).



Figure 5-5: *RF data received by a transducer.*



Figure 5-6: Individual RF lines for non-moving heart wall.

RF data were received when radii of the heart changed imitating a heartbeat for one cardiac cycle. *Figure 5-7* shows the heart wall radii change. The phase shifts between consecutive received signals were calculated and analysed *(Table 5-3). Figure 5-9* illustrates the received signals due to a number of pulse emissions. The received signals are shown on top of each other. The phase shifts with respect to the red line in the figure are extracted. Velocity of the heart scatterers were calculated using equation 5.10 and assuming that the velocity vector and ultrasound beam are at 0⁰.



Figure 5-7: The change of the heart radii over one cardiac cycle.



Figure 5-8: Summed received signals from the heart phantom for case study A.



Figure 5-9: Received signals for each RF line.

Table 5-3: Time shifts between consecutive signals and measured velocity of the heartscatterers for case study A.

Time shift, t _s	Time shift, $t_s \ [imes 10^{-6}s]$	Measured velocity of heart scatterers, $v_{heart,m}$ [mm/s]
$t_0 - t_1$	0.8	14.54
$t_1 - t_2$	0.5	8.95
$t_2 - t_3$	0.0	0.00
$t_{3} - t_{4}$	-0.2	-3.58
$t_4 - t_5$	-0.8	-14.32
$t_{5} - t_{6}$	-0.4	-7.16
$t_{6} - t_{7}$	-0.8	-14.32
$t_{7} - t_{8}$	0.0	0.00
$t_{8} - t_{9}$	0.8	14.32
$t_9 - t_{10}$	0.4	7.16

Figure 5-10 illustrates the resulting sampled signal which is the time shifts between the received signals for 11 transmitted pulses. The time shift is proportional to the velocity of the heart movement. The constant sampled values will result if the heart is stationary. The changing values are measured when heart beats as shown in the *Figure 5-10*.

The measured velocities from the time shifts are compared with actual velocities of the heart scatterers, as shown in *Figure 5-11*.



Figure 5-10: Sampled signal. Time shifts between the received signals for 11 transmitted pulses. Case study A.



Figure 5-11: *Graph of the measured and actual velocities of the heart scatterers for case study A.*

Time shifts from t_2 to t_3 and from t_7 to t_8 are expected to be 0 s since there are no changes of the heart radii as seen in *Figure 5-7*.

Figure 5-12 shows the M-mode image of the heartbeat over one cardiac cycle. This image shows the movement of the heart over time. It can be seen that the movement has a sinusoidal shape with increased and decreased heart wall radii.



Figure 5-12: M-mode image of one cardiac cycle.

5.2.2 Case Study B

The amplitude of the heart wall is scaled to 10 and the amplitude of the surrounding tissue is scaled to 1. Heart phantom B-mode image is shown in *Figure 5-13*. It is seen that the heart wall is hardly visible due to increased background amplitude.



Figure 5-13: *B-mode image of the heart phantom for case study B. The heart wall boundaries are hardly distinguishable.*

Figure 5-14 shows the summed received RF signals for 11 transmit pulses. There are additional signals present as compared with the summed received signals in case study A. These extra signals should come from the highly scattered surrounding background tissue. *Figure 5-15* illustrates individual RF lines for each transmit pulse. Received signals from background tissue, front and back heart walls and blood can be visualized in the figure.



Figure 5-14: Summed received RF lines for case study B.



Figure 5-15: *Received signals for 11 transmit pulses for case study B. The point indicates the maximum amplitude found in overall received signals.*

Table 5-4 shows the time shifts between the received RF lines. *Figure 5-15* is used to calculate time shifts between the consecutive received RF lines.

Time shift, <i>t_s</i>	Time shift, $t_s[imes 10^{-6} s]$	Measured velocity of heart scatterers, $v_{heart,m} [mm/s]$
$t_0 - t_1$	0.5	8.95
$t_1 - t_2$	0.5	8.95
$t_2 - t_3$	0.0	0.00
$t_3 - t_4$	-1.0	-17.91
$t_4 - t_5$	-0.4	-7.16
$t_{5} - t_{6}$	-0.6	-10.74
$t_{6} - t_{7}$	-0.7	-12.53
$t_7 - t_8$	0.0	0.00
$t_8 - t_9$	1.1	19.70
$t_9 - t_{10}$	0.7	12.53

Table 5-4: Time shift calculation between consecutive received RF lines and measuredvelocities of the heart scatterers. Case study B.

Time shifts from t_2 to t_3 and from t_7 to t_8 equal to 0 s since there is no change of radius of the heart wall. The velocity was found using the equation 5.10 with $\theta = 0^0$ which is an angle between the transducer probe and velocity vector. *Figure 5-16* illustrates the sampled signal plot. The plot shows times shifts between the successive received signals for 11 transmit pulses.



Figure 5-16: Sampled signal. Phase shift between the consecutive received signals for case study B.

It is possible to find velocities of the heart scatterers from the time shifts. *Figure 5-17* shows measured and actual velocities of the phantom scatterers.



Figure 5-17: Measured and actual velocities of the heart scatterers. Case study B.

Figure 5-18 illustrates M-mode image of the heartbeat. This image shows the movement of the heart over time. It can be seen that the movement has a sinusoidal shape with increased and decreased heart wall radii. There are additional movements present in the image which can be due to scattering from the surrounding tissue.



Figure 5-18: M-mode image of one cardiac cycle.

5.2.3 Case Study C

The position of the heart phantom was moved to 2 cm in x-direction, as depicted in *Figure 5-19*.

Figure 5-20 shows the beam profile for 5 mm transducer aperture and the position of the heart in the pressure field. This case study is aimed to analyze the situation if the transducer is slightly shifted during the scan. It can also be used to investigate a situation when the fetus is also slightly moved from the direct beam of the transducer.



Figure 5-19: *B-mode image of the heart phantom for case study C.*



Figure 5-20: *Beam profile of the 5 mm transducer aperture and position of the heart in the pressure field. Case study C.*

5.2.3.1 Aperture Radius 5 mm

The heart position is moved to 2 cm away from the transducer central beam. The radius of the transducer aperture is 5 mm. *Figure 5-21* and *Figure 5-22* depict received RF lines. The received signals from the back heart wall are weak. The time shifts from the front heart wall received signals are calculated using *Figure 5-22*.



Figure 5-21: *Received RF lines for 11 transmit pulses for case study C with a transducer aperture of 5 mm.*



Figure 5-22: Individual RF lines for 11 transmit pulses. Case study C.

Table 5-5 shows calculated time shifts between the successive received signals and measured velocity. The velocity was found using the equation 5.10 with $\theta = 0^0$ which is an angle between the transducer probe and velocity vector. The blood region is reduced as seen in *Figure 5-21*. The reason may be due to the position of the heart which is not directly located in the central lobe of the pressure field.

Time shift, t _s	Time shift, $t_s[imes 10^{-6}s]$	Velocity of heart scatterers, v_{heart} [mm/s]
$t_0 - t_1$	1.0	17.91
$t_1 - t_2$	0.6	10.74
$t_2 - t_3$	0.0	0.00
$t_3 - t_4$	-0.6	-10.74
$t_4 - t_5$	-0.9	-16.12
$t_{5} - t_{6}$	-1.2	-21.49
$t_{6} - t_{7}$	-1.0	-17.91
$t_7 - t_8$	0.0	0.00
$t_8 - t_9$	1.0	17.91
$t_9 - t_{10}$	0.6	10.74

 Table 5-5: Time shifts calculation between consecutive received RF lines. Case study C.

Figure 5-23 shows the sampled signal resulted from the sampling of the received signals at one time instance. The time shifts are proportional to velocity of the heart movement. Velocity of the heart phantom is then plotted and shown in *Figure 5-24*.



Figure 5-23: Sampled signal. Phase shifts between the successive signals for 11 transmit pulses.



Figure 5-24: Measured and actual velocities of the heart scatterers. Case study C.

5.2.3.2 Aperture Radius is 2 mm

The transducer aperture radius was changed to 2 mm. *Figure 5-25* depicts the beam profile from the transducer and the position of the heart in the pressure field.



Figure 5-25: *The beam profile for a round flat transducer with 2 mm aperture. The position of the heart is shown in the pressure field. Case study C.*

It is seen in *Figure 5-26* that the received signals from the back heart wall are stronger as compared to the 5 mm radius aperture.



Figure 5-26: Summed received RF lines for 11 transmit pulses. Case study C.



Figure 5-27: Individual RF lines for 11 transmit pulses. Transmit and received aperture radius is 2 mm. Case study C.

Time shifts between the successive received signals were extracted from *Figure 5-27. Table 5-6* shows calculated time shifts and measured velocity of the heart phantom. Velocity of the heart was calculated with equation 5.10. The angle between the velocity vector and a transducer beam is assumed to be 0^{0} .

Time shift, t _s	Time shift, $t_s[imes 10^{-6}s]$	Velocity of heart scatterers, v_{heart} [mm/s]
$t_0 - t_1$	0.9	16.12
$t_1 - t_2$	0.7	12.53
$t_2 - t_3$	0.0	0.00
$t_{3} - t_{4}$	-0.6	-10.74
$t_4 - t_5$	-1.5	-26.86
$t_{5} - t_{6}$	-0.7	-12.53
$t_{6} - t_{7}$	-0.9	-16.12
$t_7 - t_8$	0.0	0.00
$t_8 - t_9$	0.9	16.12
$t_9 - t_{10}$	0.7	12.53

Table 5-6: Time shifts calculation between consecutive received RF lines.

Figure 5-28 shows the sampled signal which is the time shifts between the consecutive received signals for 11 transmit pulses. Velocity of the heart phantom is then found from the time shifts. *Figure 5-29* shows actual and measured velocity of the heart phantom.



Figure 5-28: Sampled signal. Phase shift between the successive received signals.



Figure 5-29: Actual and measured velocity of the heart phantom. Case study C.

5.2.4 Case Study D

Heart phantom is placed further away from the transducer surface. Phantom is moved to 2 cm away from the transducer surface. The distance between transducer and the phantom is 10.7 cm. *Figure 5-30* depicts B-mode image of the heart phantom position. *Figure 5-31* shows the beam profile of the transducer and the heart position in the pressure field.



Figure 5-30: *B-mode image of the phantom placed at 107 mm away from the transducer surface.*



Figure 5-31: Beam profile for a transducer of 5 mm aperture. The heart is positioned in the pressure field.

Figure 5-32 shows the summed received RF signals. Amplitude of the received signals from the back heart wall is much lower as for the amplitudes of the front heart wall. It can be due to the position of the heart since the intensity of the pressure field is lower for the back side of the cardiac muscle. *Figure 5-33* illustrates individual traces of the received RF lines for 11 transmit pulses. Signals from the back heart wall are hardly visible. Time shifts at one time instance are extracted from the figure.



Figure 5-32: Summed RF lines for received signals for case study D.



Figure 5-33: RF lines for 11 transmit pulses. Case study D.

Table 5-7 shows the calculated time shifts between the successive received RF lines and measured velocity of the heart phantom. Time shifts are proportional to velocities of the heart scatterers. Velocity of the phantom is calculated with the equation 5.10. The angle between the velocity vector and transducer beam is assumed to be 0^o.

Time shift, <i>t_s</i>	Time shift, $t_s[imes 10^{-6}s]$	Velocity of heart scatterers, $v_{heart} \text{ [mm/s]}$
$t_0 - t_1$	0.4	7.16
$t_1 - t_2$	0.3	5.37
$t_2 - t_3$	0.0	0.00
$t_3 - t_4$	-0.3	-5.37
$t_4 - t_5$	-0.9	-16.11
$t_{5} - t_{6}$	-0.5	-8.95
$t_{6} - t_{7}$	-0.2	-3.58
$t_7 - t_8$	0.0	0.00
$t_{8} - t_{9}$	0.7	12.53
$t_9 - t_{10}$	0.5	8.95

 Table 5-7: Time shifts calculation between consecutive received RF line and measured velocity of the heart phantom.

Figure 5-34 shows the sampled signal which is time shifts obtained at one time instance for successive RF lines. Measured and actual velocities of the heart phantom are shown in *Figure 5-35*.



Figure 5-34: Sampled signal. Phase shift between the successive received signals.



Figure 5-35: Measured and actual velocity of the heart phantom. Case study D.

5.3 Laboratory Work

Figure 5-36 shows the schematic representations of the transmit/receive system. The hardware consists of pulse generator, receive/transmit, transducer input, voltage gain amplifier, analog/digital converter and signal processing units. The set up used to perform experiment is shown in *Figure 5-37*. The set up consists of the transmit/receive hardware system, piezoelectric transducer and a computer. KB Aerotech 2.25 MHz/13 mm non-focused transducer was used for measurements.

Resulting received digital signal is complex and can be written as [48]:

$$r_s(t) = x(t) - jy(t).$$
 (5.13)

The phase is then found as [48]:

$$\varphi(t) = \arctan\left(\frac{y(t)}{x(t)}\right),\tag{5.14}$$

The phase difference between two consecutive signals is [48]:

$$\Delta \varphi(i) = \varphi(i) - \varphi(i-1). \tag{5.15}$$

There is another way to find a phase difference. The received signal can be multiplied with a conjugate of the previous received signal [48]:

$$\Delta\varphi(i) = \arctan\left(\frac{y(i)}{x(i)}\right) \cdot \overline{\arctan\left(\frac{y(i-1)}{x(i-1)}\right)}.$$
(5.16)



This strategy was used to find the phase difference between the received signals.

Figure 5-36: Schematic representation of the transmit/receive system for ultrasound transducer.



Figure 5-37: Hardware set up for transmit and receive system for a transducer.

Round small and soft ball was taken for the experiment. The ball was placed into the container with water. First, the transducer was moved up and down vertically and phase shifts were displayed. *Figure 5-38* shows two windows with resulted M-mode image and phase shifts.


Figure 5-38: (a) *M*-mode image and (b) phase difference between the successive received signals.

Second, the transducer was moved across the ball which allowed visualizing a ball and its depth. *Figure 5-39* shows M-mode image and obtained phase difference between the consecutive received signals. The shape of the ball can be imaged when transducer is moved across it.



Figure 5-39: (*a*) *M*-mode image and (b) phase difference between the successive received signals.

6 DISCUSSION

Most of the stillbirths are preventable and depend on the access to a healthcare and early detection. However, access to a healthcare is not always possible, especially in low-income countries. Even in high-income countries, it is not always possible to provide an immediate medical help to women who may complain about reduced fetal movements [1], [80]. It is economically expensive to keep pregnant women in the hospital if she feels worried about her baby. It is a common practice that pregnant woman should return home if her worries were not justified. Therefore, it is important to have a portable cheap device that can track fetal well-being independently of the medical staff availability.

Fetus well-being can be evaluated by fetal movement observations and fetal heart rate measurements. These are the first indicators of the fetus status that can be used daily to judge fetus well-being. The counting of the fetal movements is a good approach to observe the fetus. However, it is not a reliable technique. Fetal movement assessment is affected by maternal perception of the fetus's movements [23]. Continuous monitoring of the fetal heart rate is another way to monitor fetal health. Fetal heart reactivity has a high association with fetal well-being and, therefore, fetal heartbeat is a reliable way to judge fetus's health status [17]. Doppler single-element transducer can be implemented to track fetal heartbeat over time at home. They are portable, easy in exploitation, reasonably cheap and accurate in heart rate measurements. These transducers have good sensitivity and accuracy for heart rate detection [81]. However, traditional Doppler transducers are only used by future mothers or midwives to listen to a heartbeat for short period of time. In this thesis, application of Doppler transducer was reconsidered. Traditional Doppler ultrasound transducers due to their design are able to detect a heartbeat only if they are pointing directly at the fetus heart. This research studied whether traditional Doppler transducer will be able to track continuously a fetal heartbeat if it is attached to a woman's abdomen. Since the transducer will be attached for long time, the fetus may move and the device itself may be displaced. Therefore, transducer should tolerate it and be able to provide accurate results. Therefore, it was considered to design a transducer with a smaller aperture in order to increase its robustness.

6.1 Simulations

This simulated Doppler ultrasound transducer operates by transmitting and receiving acoustic echoes from the moving heart phantom. RF lines were recorded for different positions of the scatterers over time instants. The scatterer motion was tracked based on sampling over a defined time period. The background image was static and the radii of the heart changed imitating a heartbeat. Received RF lines for front heart wall have higher amplitudes as compared to the amplitudes of the received signals from the back heart wall for all case studies. Since no attenuation was set in Field II, the possible reason is that less energy propagates through the heart to the back heart wall. The back heart wall is positioned in the pressure field with a lower intensity. The time shifts between the successive RF line signals were used to analyse Doppler shifts and to find velocity of the heart movement. The velocity of the heart phantom movement was found from the sampled signal when one sample value is taken from each RF line.

6.1.1 Case Study A and B

In case study A, heart is a highly scattered region and background is a weakly scattered region. It was expected that simulated velocity of the heart phantom was close to the theoretical values. However, the actual and measured velocities have slight differences, especially for the measured velocity recorded with sixth transmitted pulse. In case study B, background is now a highly scattered region and heart muscle is hardly distinguishable in surrounding tissue. The measured and actual velocities have differences. The possible reason is that the scattering may happen most likely from the surrounding tissue rather than from the heart wall leading to non-accurate results for received signals. Also, there are computational artifacts present during the simulations that may lead to errors in calculating velocities. The simulation may not always give the same results as theoretical calculations. The artifacts may come from the interaction of the simulation tool with operational parameters and input data. Some of the computational artifacts can be deduced from transmit and receive signals or spatial impulse responses applied in Field II. The aperture of the transducer is divided into mathematical elements and the responses from these elements are superimposed due to linearity. The spatial impulse responses are calculated at the discrete points by taking

the difference from sample point to another sample point. The discrete time can be a factor for the numerical artifacts. Quantitative analysis of the simulation errors was performed by Jensen [82]. The accuracy of the pulse-echo response was found to be a function of the sampling frequency.



Figure 6-1: *The accuracy of the pulse-echo impulse response for rectangles and bounding lines transducer elements as a function of the sampling frequency* [82].

Rectangular transducer elements were used for the simulation and the sampling frequency was set to 100 MHz. From the above *Figure 6-1*, it is seen that 100 MHz sampling frequency will result in 20% relative error for accurate computations of the received RF signals. Therefore, the accuracy of the simulation depends on the choice of the sampling frequency. It is considered that increasing the sampling frequency provides fine transmitted pulse which will results in accurate results for further summations and convolutions operations. However, limitations should be applied for the choice of the sampling frequency. First, high sampling frequency above 100 MHz will result in increased number of samples to compute which may be a non-realistic during the real-time performance. Second, Field II functions reliably with 100 MHz sampling frequency and does better approximate calculations [62]. Third, the process of finding sampling frequency arbitrary for each case can tedious and non-reliable.

6.1.2 Case Study C and D

Case study C and D are the most realistic situations if Doppler ultrasound technique is used for continuous monitoring. The challenge of this continuous monitoring is that transducer central beam may not be directed always at the fetal heart due to baby movements or if transducer is displaced during exploitation. Therefore, it is important to investigate whether it is still possible to detect heartbeat if the central beam of the transducer is not at the heart position. It was considered to use a transducer with a smaller aperture as in case study C. The beam profile can be analyzed from the beam divergence half-angle, θ_R . The angle is given by [51]:

$$\theta_R = \sin^{-1}\left(\frac{0.61\lambda}{a}\right),\tag{6.1}$$

where *a* is a aperture radius. The beam divergence half-angles were found for 5 mm and 2 mm apertures and equal to 5.39^o and 13.58^o respectively. *Figure 6-2* shows the dependence of the beam divergence half-angle on the aperture radius. As the radius increases the central lobe narrows and the number of side lobes increases. This tendency is also observed from the beam profile figures for 5 and 2 mm aperture radius *(Figure 5-20* and *Figure 5-25)*.



Figure 6-2: The far-field pressure directivity function $D(\theta)$ for $a=2\lambda$ (top polar graph) and for $a=4\lambda$ (bottom polar graph) with given beam divergence half-angle, θ_R .

The measured and actual velocities are in a good agreement with each other for 5 mm transducer aperture. However, when the transducer aperture was changed to 2 mm, the measured velocity of the heart phantom has more errors. There are more

received signals from the surrounding tissue with 2 mm transducer aperture leading to non-accurate result for measured velocity.

If the heart position is moved further away in depth (z-direction) from the transducer surface as in case study D, it is still possible to analyze velocity of the heartbeat. The back heart wall is hardly visible. However, the movement of the front heart wall should be enough to extract velocity movement data. Therefore, the detectability of the heart movement is mostly affected when transducer is shifted in x-direction. It was noticed that when the aperture is decreased, there were more scattered signals from the surrounding tissue. However, with 2 mm aperture it was still possible to detect heart movements. If the baby is moved further away from the central lobe then it will get more difficult to detect a heartbeat. However, when the position of the heart is moved further in *x*- and *z*-directions, no significant changes in the received RF signals were observed. This might be a limitation of the Field II that it cannot incorporate these changes.

Convex shape of the transducer can be considered for this application. Convex transducer has wider field of view radiating energy broadly. This shape of the transducer can be more suitable to detect heart movements and can solve problems that flat piston cannot. *Figure 6-3* shows the beam profile for convex shape transducer with 5 mm aperture simulated by Field II. The beam width is much wider as compared with the flat transducer of 5 mm aperture used for case studies.



Figure 6-3: Beam profile for a one element convex transducer with 5 mm aperture.

Therefore, convex shape provides a wide beam profile with a large aperture as compared to a flat piston transducer.

6.2 Laboratory work

Laboratory work was performed to analyse how single-element transducer functions. Phase shifts between the received signals can be extracted from the moving object. The test environment consists of the ball placed into container filled with water. It was not possible to have any real-life environment as it would require a pregnant woman to participate. Ultrasound is considered to be safe for a mother and a fetus. However, at this stage of the research, it is not ethical to perform any real-life experiments. This set up can be improved and used for further studies. Other types of the transducers can be connected to a transmit/receive system. Based on the phase shifts, this system is also able to produce M-mode image which also can be used to track heart movements. M-mode imaging can be one of the options that can be included into continuous monitoring system to track fetal well-being.

6.3 Ultrasound safety

Diagnostic ultrasound is considered to be a safe technique used in medicine. Medical ultrasound offers many opportunities for diagnostics due to its impressive safety [59]. The safety of ultrasound mainly depends on the magnitude of the temperature elevation due to ultrasound exposure and the duration of the acoustic exposure [83]. When it comes to the bio-effects induced by a transducer, I_{SPTA} is introduced and stands for spatial peak temporal averaged intensity. I_{SPTA} is a power divided by effective radiating area [59]. FDA approves I_{SPTA} to be 720 mW/cm² [84]. Intensities less than 100 mW/cm² do not generate heat regardless of the length of the exposure [17]. Pulsed-wave Doppler transducer generate 1.5 to 5 mW/cm² [17]. Also, this Doppler ultrasound is a single-element unfocused piston-like transducer operating at 2 MHz which is unlikely to excite any cavitation in fetal tissue [59]. However, more research is required to study bio-effects for long term Doppler transducers monitoring.

7 CONCLUSION AND FUTURE WORK

The proposed idea to have a continuous fetal heartbeat monitoring is found to be one of the solutions to reduce number of stillbirths. Discussions with midwives and doctor from Horten Kommune Helsestasjon and also from literature [27], [5], [85], [86] led to a conclusion that continuous monitoring can help to reduce pregnancy complications and even stillbirth. Continuous monitoring will provide better tracking of the fetus health and small changes of the fetus well-being will be recorded. Therefore, pregnant women can get immediate help if it is required.

It is hard to adopt traditional Doppler ultrasound transducer for home use. The robustness of the system can be improved if transducer aperture decreases. Wider transducer beam will be able to catch the fetal heartbeat. However, transducer aperture of 2 mm results only in 13.580 of beam divergence half-angle. Therefore, it may not be always possible to detect fetal heartbeat due to the transducer displacement or fetus's movements. The system then becomes unreliable and problematic for long-term monitoring.

Convex-shaped transducer is a promising device for a continuous fetal heartbeat monitoring and also for future research topic. Both fetal heart rate and fetal movements tracking can be a better solution to observe fetus well-being [87]. For example, fetal movement assessment can be advanced by adopting an accelerometer. The system then can be further advanced with the ability to analyse, record and transmit results for a hospital review and evaluation. It also should be light and small and easy to be used for a pregnant woman. Transmit/receive hardware system can then be used to perform experimental work in a test environment and later in a real-life environment.

GLOSSARY

Amniotic fluid	A fluid surrounding a fetus
Amniotic sac	A fluid-filled sac that contains and protects a fetus in a womb
Diastole	Part of the cardiac cycle when the heart refills with blood
Endocardium	Inner layer of the heart wall that lines the chambers of the heart
Endometrium	Inner membrane of the uterus
Epicardium	Layer immediately outside of the myocardium heart muscle
Fundus	Top portion of the uterus, located opposite to the cervix
Myocardium	Middle layer of the heart wall
Pericardium	Doubled-wall sac containing the heart and the roots of the great vessels.
Perimetrium	Outer layer of the uterus
Placenta	Organ that connects the fetus to the uterine wall to allow nutrient uptake
Pubic bone	One of the bones that make up the pelvis
Systole	Contraction phase of the cardiac cycle
Umbilical artery	Paired artery that is found in the abdominal and pelvic regions. It extends into the umbilical cord.

APPENDIX A: MATLAB CODE FOR A HEART PHANTOM IMAGE

```
% Calling: [positions, amp] = heart phantom (N);
2
8
  Parameters: N - Number of scatterers in the phantom
2
                positions - Positions of the scatterers.
8
  Output:
00
                            - amplitude of the scatterers.
                 amp
function [positions, amp] = heart phantom (N)
x size = 140/1000;
                     % Width of phantom [mm]
y_size = 30/1000; % Transverse width of phantom [mm]
z_size = 160/1000; % Height of phantom [mm]
z_start = 35/1000; % Start of phantom surface [mm];
% Create the general scatterers
x = (rand (N, 1) - 0.5) * x size;
y = (rand (N, 1) - 0.5) * y size;
z = rand (N, 1) * z size + z start;
% Generate the amplitudes with a Gaussian distribution
amp=10*randn(N,1);
% Make the heart and set the amplitudes to zero inside
% heart radii
r1=18/1000;
                 % Radii [mm]
r2=16/1000;
               % Place of heart [mm]
xc=0/1000;
zc=90/1000+z start;
inside=((x-xc).^2+(z-zc).^2)<r1^2;
inside=inside.*((x-xc).^2+(z-zc).^2)>r2^2;
A heart=1;
ampHEART=amp.*inside*A heart;
ampMEDIUM=amp.*(inside-1);
amp=ampHEART+ampMEDIUM;
inside = ((x-xc).^2 + (z-zc).^2) < r2^2);
amp = amp .* (1-inside);
positions=[x y z];
```

Transducer Simulation Script

```
% Generate the transducer apertures for send and receive
f0=2e6;
                       % Transducer center frequency [Hz]
fs=100e6;
                        % Sampling frequency [Hz]
c=1540;
                        % Speed of sound [m/s]
lambda=c/f0;
                        % Wavelength [m]
width=lambda;
                       % Width of element
element height=5/1000; % Height of element [m]
kerf=0.05/1000;
                       % Kerf [m]
                      % Fixed focal point [m]
focus=[0 0 70]/1000;
                       % Number of physical elements
N elements=192;
                       % Number of active elements
N active=64;
% Set the sampling frequency
set sampling(fs);
% Generate aperture for emission
xmit aperture = xdc linear array (N elements, width, element height, kerf,
1, 10, focus);
% Set the impulse response and excitation of the xmit aperture
impulse response=sin(2*pi*f0*(0:1/fs:2/f0));
impulse response=impulse response.*hanning(max(size(impulse response)))';
xdc impulse (xmit aperture, impulse response);
excitation=sin(2*pi*f0*(0:1/fs:2/f0));
xdc excitation (xmit aperture, excitation);
% Generate aperture for reception
receive aperture = xdc linear array (N elements, width, element height,
kerf, 1, 10, focus);
% Set the impulse response for the receive aperture
xdc impulse (receive aperture, impulse response);
  Load the computer phantom
if ~exist('pht data.mat')
 disp('Scatterer positions should be made by the script mk pht')
 disp('before this script can be run')
 return
else
 load pht data
 end
% Set the different focal zones for reception
focal zones=[30:20:200]'/1000;
Nf=max(size(focal zones));
focus times=(focal zones-10/1000)/1540;
```

z focus=60/1000; % Transmit focus % Set the apodization apo=hanning(N active)'; % Do linear array imaging no lines=120; % Number of lines in image image width=80/1000; % Size of image sector % Increment for image d x=image width/no lines; % Do imaging line by line for i=[1:no lines] % Test if the file for the line exist. % Skip the simulation, if the line exits and % go the next line. Else make the simulation file name=['rf data/rf ln',num2str(i),'.mat']; if ~exist(file name) % Save a file to reserve the calculation cmd=['save rf data/rf ln',num2str(i),'.mat i']; eval(cmd); disp(['Now making line ',num2str(i)]) % The the imaging direction x = -image width/2 + (i-1) * d x;% Set the focus for this direction with the proper reference point xdc center focus (xmit aperture, [x 0 0]); xdc_focus (xmit_aperture, 0, [x 0 z_focus]); xdc_center_focus (receive_aperture, [x 0 0]); xdc focus (receive aperture, focus times, [x*ones(Nf,1), zeros(Nf,1), focal zones]); % Calculate the apodization N pre = round(x/(width+kerf) + N elements/2 - N active/2); N post = N elements - N pre - N active; apo_vector=[zeros(1,N_pre) apo zeros(1,N_post)]; xdc_apodization (xmit_aperture, 0, apo_vector);

% Calculate the received response

[rf_data, tstart]=calc_scat(xmit_aperture, receive_aperture, phantom positions, phantom amplitudes);

xdc apodization (receive aperture, 0, apo vector);

% Store the result

```
cmd=['save rf_data/rf_ln',num2str(i),'.mat rf_data tstart'];
disp(cmd)
eval(cmd);
else
disp(['Line ',num2str(i),' is being made by another machine.'])
end
end
% Free space for apertures
xdc_free (xmit_aperture)
xdc_free (receive_aperture)
disp('You should now run make image to display the image')
```

Make Image Script

```
f0=2e6;
                        % Transducer center frequency [Hz]
fs=100e6;
                          % Sampling frequency [Hz]
                          % Speed of sound [m/s]
c = 1540;
no lines=120;
                          % Number of lines in image
                          % Size of image sector
image width=140/1000;
d_x=image_width/no_lines; % Increment for image
% Read the data and adjust it in time
min sample=0;
for i=1:no lines
  % Load the result
  cmd=['load rf data/rf ln',num2str(i),'.mat'];
  disp(cmd)
  eval(cmd)
  % Find the envelope
  rf env=abs(hilbert([zeros(round(tstart*fs-min sample),1); rf data]));
  env(1:max(size(rf env)),i)=rf env;
  end
% Do logarithmic compression
              % Sampling frequency decimation factor
D=10:
dB range=50; % Dynamic range for display in dB
disp('Finding the envelope')
log env=env(1:D:max(size(env)),:)/max(max(env));
log env=20*log10(log env);
log env=127/dB range*(log env+dB range);
```

```
\% Make an interpolated image
```

```
disp('Doing interpolation')
ID=20;
[n,m]=size(log_env);
new_env=zeros(n,m*ID);
for i=1:n
 new_env(i,:)=interp(log_env(i,:),ID);
 end
[n,m]=size(new_env);
fn=fs/D;
clf
image(((1:(ID*no lines-1))*d x/ID-
no lines*d x/2)*1000,((1:n)/fn+min sample/fs)*1540/2*1000,new env)
xlabel('Lateral distance [mm]')
ylabel('Axial distance [mm]')
colormap(gray(128))
axis('image')
axis([-70 70 35 160])
```

APPENDIX B: MATLAB CODES USED FOR CASE STUDY A, B, C & D

```
%% Main script to simulate heart wall movements and to receive RF lines
clc; clear; close all;
%% variables
N = 200000;
f heart = 2.33;
Thm = 1/f heart;
Nhm = 10; %Number of transmit pulses=Nhm+1
t 2bps=[0:Thm/Nhm:Thm];
%t 2bps=[0 0 0 0 0 0 0 0 0 0 0];
 r1=(18 + 1*sin(2*pi*f_heart*(t_2bps)))./1000;
 r2=(16 + 1*sin(2*pi*f heart*(t 2bps)))./1000;
% create struct
results = struct(); % empty struct
88
for ii = 1:length(r1)
    % call function heart pht
    [phantom positions, phantom amplitudes] = heart pht(N, r1(ii), r2(ii));
    save pht data.mat phantom positions phantom amplitudes
    % call script sim img
   %sim img;
    % call script make image
    %make image;
    % call script single el tr pulse
    signle el tr pulse;
    % allocate data to struct
  results(ii).v = v;
     results(ii).t = t;
     results(ii).t2 = t2;
end
응응
figure(91); hold on
for ii =0:size(results,2)-1
   %plot(results(ii+1).t2, ii+abs(hilbert(results(ii+1).v./2)), 'k')
   plot(results(ii+1).t2, ii+((results(ii+1).v./2)), 'k')
   [tmax, indtmax] = max(results(ii+1).v);
    plot(results(ii+1).t2(indtmax), ii+results(ii+1).v(indtmax)./2, 'ob')
    xlabel('Time [s]', 'FontSize', 12)
ylabel('RF lines', 'Fontsize', 12)
end
[tmax, indtmax] = max(results(1).v);
plot([results(1).t2(indtmax), results(1).t2(indtmax)], [0, ii+1], 'r')
xlim([1.1e-4, 1.8e-4])
save('results', 'results')
```

Script for Transducer Simulation

```
% Set initial parameters
f0=2e6; % Transducer center frequency [Hz]
fs=100e6; % Sampling frequency [Hz]
% c=1540; % Speed of sound [m/s]
% lambda=c/f0; % Wavelength [m]
R=5/1000;
element size=1/1000;
M=5;
% Define the transducer
Th=xdc piston(R, element size);
% Set the impulse response and excitation of the emit aperture
impulse response=sin(2*pi*f0*(0:1/fs:3/f0));
impulse response=impulse response.*hanning(max(size(impulse response)))';
xdc impulse (Th, impulse response);
excitation=sin(2*pi*f0*(0:1/fs:M/f0));
xdc excitation (Th, excitation);
% Do the calculation
[v,t]=calc scat multi(Th, Th, phantom positions, phantom amplitudes);
%calculate the received signals from the collections of scatterers for all
the elements in the aperture
% Plot the individual responses
[N,M]=size(v)
v=v/max(max(v));
figure(1); hold on
plot((0:N-1)/fs+t,v)
% hold off
%title('Individual traces')
xlabel('Time [s]')
ylabel('Normalized response')
%% update t
t2 = (0:N-1)/fs+t;
88
L=length(v);
nfft = 2^nextpow2(10*L);
f=fs.*[-nfft/2:1:nfft/2-1]/nfft;
V=2*fft(v,nfft)/L;
% V = V(nfft/2:end);
figure(2); hold on
plot(f/le6, fftshift(abs(V)));
xlim([0,5]);
xlabel('f [MHz]')
ylabel('Amplitude')
```

APPENDIX C: MATLAB CODE FOR M-MODE IMAGE

```
f0=2e6;
                        % Transducer center frequency [Hz]
fs=100e6;
                         % Sampling frequency [Hz]
c=1540;
                         % Speed of sound [m/s]
no lines=1;
                         % Number of lines in image
image width=60/1000;
                       % Size of image sector
d x=image width/no lines; % Increment for image
  % Load the result
min sample=0;
dynamicRange=17;
%load('Result\Info.mat');
        i=1;
        for time=0:1:length(t 2bps)-1
            fileName=sprintf('rf data\\rf ln%i.mat',i);
            load(fileName,'rf data');
            echo=rf data;
            %echo=-echo(1:1:ti*fs);
            %rf env=abs(hilbert([zeros(round(time*fs-
min_sample),1);echo]));
            echo=echo(1:1:length(echo));
            rf env=abs(hilbert(echo));
            env(1:size(rf_env,1),i+1)=rf_env;
            i=i+1;
        end
  numberOfLines=i;
        c=1540;
        env_dB=20*log10(env);
        env_dB=env_dB-max(max(env_dB));
        env_gray=127*(env_dB+dynamicRange)./dynamicRange;
        %env_gray = env;
        depth=((0:size(env,1)-1)+min sample)/fs*c/2;
       x=t 2bps;
        figure
        imagesc(x,(depth*1000),env_gray,[0 dynamicRange]);
        xlabel('No. of transmit pulses');
        ylabel('Depth [mm]');
        colormap(gray(128));
        colorbar('XTick',0,'XTickLabel','Power(dB)');
        hold off
```

APPENDIX D: FLAT PISTON TRANSDUCER BEAM PROFILE

```
function Th=define_transducer(TR)
% function Th=define transducer(TR)
2
% Define transducer for use in Field II
8
% TR Struct containing transducer specification
% Th Pointer to transducer aperture, from Field II
%--- Geometry ---
% Round flat aperture following Field II specification
ele size=0.2/1000;
radius=5/1000;
Th=xdc piston(radius, ele size);
%--- Pulse ---
excitation=sin(2*pi*TR.f0*(0:1/TR.fs:5/TR.f0));
xdc excitation (Th, excitation);
%--- Transmit aperture ---
% Impulse response approximated as Hanning-windowed sine-pulse
impulse response=sin(2*pi*TR.f0*(0:1/TR.fs:3/TR.f0));
impulse response=impulse response.*hanning(max(size(impulse response)))';
xdc impulse(Th, impulse response);
%--- Receive aperture ---
xdc impulse(Th, impulse response);
return
c = 1540;
                 % m/s Speed of sound in fluid
% 'TR' is a struct containing transducer specification
TR.fs = 100e6; % Hz Sample rate
TR.f0 = 2e6;
                   % Hz Transducer center frequency
Th = define transducer(TR);
set sampling(TR.fs);
TR= calculate and plot(TR, Th, [70 160]*1e-3);
%title( sprintf('Frequency %.1f MHz. Width %.0f mm', TR.f0/1e6, TR.W*1e3) )
xdc free(Th)
```

return

Appendix E: LABVIEW 'Process' state



References

- L. De Bernis, M. V Kinney, W. Stones, P. Ten Hoope-Bender, D. Vivio, S. H. Leisher, Z. Qar,
 A. Bhutta, M. Gülmezoglu, M. Mathai, J. M. Belizán, L. Franco, L. Mcdougall, J. Zeitlin, A.
 Malata, K. E. Dickson, and J. E. Lawn, "Stillbirths: ending preventable deaths by 2030,"
 Lancet, vol. 387, pp. 703–716, 2016.
- "WHO | Stillbirths," 2015. [Online]. Available: http://www.who.int/maternal_child_adolescent/epidemiology/stillbirth/en/. [Accessed: 06-Sep-2015].
- [3] "Stillbirth Introduction NHS Choices." [Online]. Available: http://www.nhs.uk/conditions/Stillbirth/Pages/Definition.aspx. [Accessed: 06-Sep-2015].
- [4] E. O'Neill and J. Thorp, "Antepartum evaluation of the fetus and fetal well being.," *Clin. Obstet. Gynecol.*, vol. 55, no. 3, pp. 722–30, Sep. 2012.
- [5] A. Vais and L. Kean, "Stillbirth in the UK: current trends, investigation and opportunities for prevention," *Obstet. Gynaecol. Reprod. Med.*, vol. 25, no. 6, pp. 160–166, 2015.
- [6] R. A. Haws, M. Y. Yakoob, T. Soomro, E. V Menezes, G. L. Darmstadt, and Z. A. Bhutta, "Reducing stillbirths: screening and monitoring during pregnancy and labour."
- [7] "The American College of Obstetricians and Gynecologists," 2016. [Online]. Available: http://www.acog.org/Patients/FAQs/Special-Tests-for-Monitoring-Fetal-Health.
- [8] "Home foetal heart monitors 'risky," NHS Choices, 2009. [Online]. Available: http://www.nhs.uk/news/2009/11November/Pages/Home-foetal-heart-monitorsrisky.aspx. [Accessed: 09-Sep-2015].
- K. J. Gibbins, R. M. Silver, H. Pinar, U. M. Reddy, C. B. Parker, V. Thorsten, M. Willinger, D. J. Dudley, R. Bukowski, G. R. Saade, M. A. Koch, D. Conway, C. J. Hogue, B. J. Stoll, and R. L. Goldenberg, "Stillbirth, hypertensive disorders of pregnancy, and placental pathology," *Placenta*, vol. 43, pp. 61–68, 2016.
- [10] "Special Tests for Monitoring Fetal Health ACOG." [Online]. Available: http://www.acog.org/Patients/FAQs/Special-Tests-for-Monitoring-Fetal-Health.
 [Accessed: 05-Sep-2015].
- [11] R. A. Haws, M. Y. Yakoob, T. Soomro, E. V Menezes, G. L. Darmstadt, and Z. A. Bhutta, "Reducing stillbirths: screening and monitoring during pregnancy and labour.," *BMC*

Pregnancy Childbirth, vol. 9 Suppl 1, no. Suppl 1, p. S5, Jan. 2009.

- [12] A. Y. Elzouki, H. A. Harfi, F. B. Stapleton, H. Nazer, W. Oh, and R. J. Whitley, *Textbook of Clinical Pediatrics*. Springer Science & Business Media, 2012.
- [13] B. Winje, E. Saastad, N. Gunnes, J. Tveit, B. Stray-Pedersen, V. Flenady, and J. Frøen,
 "Analysis of 'count-to-ten' fetal movement charts: a prospective cohort study," *BJOG An Int. J. Obstet. Gynaecol.*, vol. 118, no. 10, pp. 1229–1238, 2011.
- [14] L. Mangesi and G. J. Hofmeyr, "Fetal movement counting for assessment of fetal wellbeing.," *Cochrane database Syst. Rev.*, no. 1, p. CD004909, Jan. 2007.
- [15] "Non-Stress Test (NST)." [Online]. Available: http://www.hopkinsmedicine.org/gynecology_obstetrics/specialty_areas/maternal_fetal _medicine/services/antepartum_testing_prenatal_diagnosis_treatment_center/non_stres s_test.html. [Accessed: 06-Sep-2015].
- [16] *Fetal Heart Monitoring: Principles and Practices.* Kendall Hunt, 1993.
- [17] R. K. Freeman, T. J. Garite, M. P. Nageotte, and L. A. Miller, *Fetal Heart Rate Monitoring*. Lippincott Williams & Wilkins, 2012.
- [18] J. P. Phelan, M. O. Ahn, C. V Smith, S. E. Rutherford, and E. Anderson, "Amniotic fluid index measurements during pregnancy.," *J. Reprod. Med.*, vol. 32, no. 8, pp. 601–4, Aug. 1987.
- [19] L. Littleton-Gibbs and J. Engebretson, *Maternity Nursing Care*. Cengage Learning, 2012.
- [20] K. Whitaker, P. Eberle, and L. Trujillo, *Comprehensive Perinatal & Pediatric Respiratory Care*. Cengage Learning, 2014.
- [21] M. I. Levene and F. A. Chervenak, *Fetal and Neonatal Neurology and Neurosurgery*. Elsevier Health Sciences, 2009.
- [22] *Doppler Ultrasound in Obstetrics and Gynecology*. Springer Science & Business Media, 2006.
- [23] J. F. Frøen, A. E. P. Heazell, J. V. H. Tveit, E. Saastad, R. C. Fretts, and V. Flenady, "Fetal Movement Assessment," *Semin. Perinatol.*, vol. 32, no. 4, pp. 243–246, 2008.
- [24] E. K. Cydney Afriat Menihan, *Electronic Fetal Monitoring: Concepts and Applications*. Lippincott Williams & Wilkins, 2007.
- [25] WebMD, "High-risk pregnancy," 2016. [Online]. Available: http://www.webmd.com/baby/checking-fetus-in-high-risk-pregnancy. [Accessed: 03-Mar-2016].

- [26] A. Biswas, S. Biswas, A. Cum Mo, M. Walliullah, and A. K. Mukhopadhyay, "Indian Medical Gazette Role of Non Stress Test in Monitoring High Risk Pregnancy," 2013.
- [27] R. Brown, J. H. B. Wijekoon, A. Fernando, E. D. Johnstone, and A. E. P. Heazell, "Continuous objective recording of fetal heart rate and fetal movements could reliably identify fetal compromise, which could reduce stillbirth rates by facilitating timely management," *Med. Hypotheses*, vol. 83, no. 3, pp. 410–417, 2014.
- [28] T. R. Moore, "The Role of Amniotic Fluid Assessment in Evaluating Fetal Well-Being."
- [29] Ronald S. Gibbs, *Danforth's Obstetrics and Gynecology*. Lippincott Williams & Wilkins, 2008.
- [30] R. Gagnon, M. Van den Hof, H. N. Stephen Bly, O. O. Duncan Farquharson, V. B. Robert
 Gagnon, L. O. Barbara Lewthwaite, W. M. Lucie Morin, M. Q. Shia Salem, and A. Skoll, "THE
 USE OF FETAL DOPPLER IN OBSTETRICS," *J Obs. Gynaecol Can*, vol. 25(7), 2003.
- [31] R. L. Goldenberg, E. M. McClure, A. H. Jobe, B. D. Kamath-Rayne, M. G. Gravette, and C. E.
 Rubens, "Stillbirths and neonatal mortality as outcomes," *Int. J. Gynecol. Obstet.*, vol. 123, no. 3, pp. 252–253, 2013.
- [32] G. Pagani, F. D'Antonio, A. Khalil, A. Papageorghiou, A. Bhide, and B. Thilaganathan,
 "Association between reduced fetal movements at term and first trimester markers of impaired placental development," *Placenta*, vol. 35, no. 8, pp. 606–610, 2014.
- [33] c and Ernst Beinder, MD,a Tomas Grancay,a Theresa Menéndez, MD,b Helmut Singer, MD and Md. Michael Hofbeck, "Fetal sinus bradycardia and the long QT syndrome," *Am J Obs. Gynecol*, vol. 185, no. 3.
- P. G. Pézard, F. Boussion, L. Sentilhes, C. Lépinard, M.-H. Couvreur, J. Victor, P. Geslin, and
 P. Descamps, "Fetal tachycardia: A role for amiodarone as first- or second-line therapy?," *Arch. Cardiovasc. Dis.*, vol. 101, no. 10, pp. 619–627, 2008.
- [35] "How can I monitor my baby's movements? | Count the Kicks." [Online]. Available: http://www.countthekicks.org.uk/mums/your-babys-movements/how-to-monitorbabys-movements/. [Accessed: 09-Sep-2015].
- [36] "Products | Count the Kicks." [Online]. Available: http://www.countthekicks.org.uk/shop/. [Accessed: 09-Sep-2015].
- [37] "Kick Counter Wristband | Count the Kicks." [Online]. Available: http://www.countthekicks.org.uk/shop/kick-counter-wristband/. [Accessed: 09-Sep-2015].

- [38] "Kickme Baby Kicks Counter Android Apps on Google Play." [Online]. Available: https://play.google.com/store/apps/details?id=com.dilmeapps.kickme&hl=en_GB. [Accessed: 09-Sep-2015].
- [39] "Popamazing LCD Prenatal Fetal Doppler Angel Heart Sound Monitor Backlight Screen Display CE & EC Medical Directive: Amazon.co.uk: Baby." [Online]. Available: http://www.amazon.co.uk/dp/B00SM7LWFK?psc=1. [Accessed: 09-Sep-2015].
- [40] "Baby Heartbeat Listener Android Apps on Google Play." [Online]. Available: https://play.google.com/store/apps/details?id=com.lema.android.babyheartbeatlistene r&hl=en_GB. [Accessed: 09-Sep-2015].
- [41] "BBC NEWS | Health | Foetal heart rate monitor warning." [Online]. Available: http://news.bbc.co.uk/2/hi/health/8343317.stm. [Accessed: 09-Sep-2015].
- [42] S. Bhong and S. D. Lokhande, "Wireless Fetal Monitoring," *Int. J. Sci. Res.*, vol. 2, no. 3, pp. 2319–7064, 2013.
- [43] I. Clausen and T. Glott, "Development of Clinically Relevant Implantable Pressure Sensors: Perspectives and Challenges.," *Sensors (Basel).*, vol. 14, no. 9, pp. 17686–17702, Jan. 2014.
- [44] A. J. Wolfberg, "The future of fetal monitoring.," *Rev. Obstet. Gynecol.*, vol. 5, no. 3–4, pp. e132–6, 2012.
- [45] Q. Plummer, "Wireless Fetal Monitoring System Gets Feds Blessings, Boasts Advanced Features," *Tech Times*, 2015. [Online]. Available: http://www.techtimes.com/articles/25689/20150113/fda-clears-marketing-ofwireless-system-for-fetal-monitoring.htm. [Accessed: 11-Sep-2015].
- [46] A. A. Boatin, B. Wylie, I. Goldfarb, R. Azevedo, E. Pittel, C. Ng, and J. Haberer, "Wireless fetal heart rate monitoring in inpatient full-term pregnant women: testing functionality and acceptability.," *PLoS One*, vol. 10, no. 1, p. e0117043, Jan. 2015.
- [47] T. L. Szabo and T. L. Szabo, "3 ACOUSTIC WAVE PROPAGATION," in *Diagnostic Ultrasound Imaging*, 2004, pp. 47–70.
- [48] Jørgen Arendt Jensen, *Estimation of Blood Velocities Using Ultrasound A Signal Processing Approach*. Technical University of Denmark: Cambridge University Press, 1996.
- [49] Jørgen Arendt Jensen, Estimation of Blood Velocities Using Ultrasound A Signal Processing Approach. Technical University of Denmark, Lyngby: Cambridge University Press, 1996.

- [50] T. L. Szabo and T. L. Szabo, "1 INTRODUCTION," in *Diagnostic Ultrasound Imaging*, 2004, pp. 1–28.
- [51] R. S. C. Cobbold, *Foundations of biomedical ultrasound*. Oxford University Press, 2007.
- [52] B. Fay, "Ultrasonic Backscattering: Fundamentals and Applications," *Phys. Acoust.*, pp. 41–53, 1991.
- [53] T. L. Szabo and T. L. Szabo, "4 ATTENUATION," in *Diagnostic Ultrasound Imaging*, 2004, pp. 71–95.
- [54] S. Cochran, "Piezoelectricity and basic configurations for piezoelectric ultrasonic transducers," in *Ultrasonic Transducers*, 2012, pp. 3–35.
- [55] FERROPERM, "Pz27 (Navy II) A soft PZT with low ageing rates and high sensitivity.".
- [56] J. E. Wilhjelm, M. Kristensson, and O. T. Andersen, "Medical diagnostic ultrasound physical principles and imaging," 2010.
- [57] T. L. Szabo and T. L. Szabo, "11 DOPPLER MODES," in *Diagnostic Ultrasound Imaging*, 2004, pp. 337–380.
- [58] G. S. Reeder, P. J. Currie, D. J. Hagler, A. J. Tajik, and J. B. Seward, "Use of Doppler techniques (continuous-wave, pulsed-wave, and color flow imaging) in the noninvasive hemodynamic assessment of congenital heart disease," *Mayo Clin Proc*, vol. 61, no. 9, pp. 725–744, Sep. 1986.
- [59] T. L. Szabo and T. L. Szabo, "15 ULTRASOUND-INDUCED BIOEFFECTS," in *Diagnostic Ultrasound Imaging*, 2004, pp. 489–516.
- [60] T. R. Nelson, J. B. Fowlkes, J. S. Abramowicz, and C. C. Church, "Ultrasound biosafety considerations for the practicing sonographer and sonologist.," *J. Ultrasound Med.*, vol. 28, no. 2, pp. 139–50, Feb. 2009.
- [61] "510(k) Summary of Safety and Effectiveness I90-IYO," 2009.
- [62] J. A. Jensen, "Users' guide for the Field II program."
- [63] J. A. Jensen, "Field: A Program for Simulating Ultrasound Systems," *Publ. Med. Biol. Eng. Comput.*, vol. 34, no. 1, pp. 351–353, 1996.
- [64] M. Schlaikjer, S. Torp-Pedersen, and J. A. Jensen, "Simulation of RF data with tissue motion for optimizing stationary echo canceling filters."
- [65] M. A. Lediju, M. J. Pihl, J. J. Dahl, and G. E. Trahey, "Quantitative assessment of the magnitude, impact and spatial extent of ultrasonic clutter.," *Ultrason. Imaging*, vol. 30, no. 3, pp. 151–68, Jul. 2008.

- [66] J. A. Jensen and S. I. Nikolov, "Fast Simulation of Ultrasound Images."
- [67] Johan Kirkhorn, "Introduction to IQ-demodulation of RF-data."
- [68] J. N. The Boston Women's Health Book Collective, *Our Bodies, Ourselves: Pregnancy and Birth.* Simon and Schuster, 2008.
- [69] D. Shimon, "Myometrial Thickness in Pregnancy :," *J. Ultrasound Med.*, pp. 661–665, 1998.
- [70] N. B. Hershfield, "The Abdominal Wall," *J. Clin. Gastroenterol.*, vol. 14, no. 3, pp. 199–202, 1992.
- [71] L. Lancerotto, C. Stecco, V. MacChi, A. Porzionato, A. Stecco, and R. De Caro, "Layers of the abdominal wall: Anatomical investigation of subcutaneous tissue and superficial fascia," *Surg. Radiol. Anat.*, vol. 33, no. 10, pp. 835–842, 2011.
- [72] E. Zohav, A. Dunsky, O. Segal, R. Peled, A. Herman, and S. Segal, "The effects of maternal and fetal parameters on the quality of nuchal translucency measurement," pp. 638–640, 2001.
- [73] "WOMEN AND NEWBORN HEALTH SERVICE MEASURING FUNDAL HEIGHT WITH A TAPE MEASURE," *King Edward Memorial Hospital*. [Online]. Available: http://www.kemh.health.wa.gov.au/development/manuals/0&G_guidelines/sectionb/1 /b1.6.3.pdf. [Accessed: 10-Mar-2016].
- S. Kunii, J. Sugawara, Y. Kimura, N. Imai, H. Chisaka, H. Hasegawa, Y. Koiwa, H. Kanai, and
 K. Okamura, "Fetal Myocardial Thickening Measured by Ultrasonic-Based Technique
 Called 'Phased-Tracking Method," *Fetal Diagn Ther*, vol. 21, pp. 458–465, 2006.
- [75] C. Firpo, J. I. E. Hoffman, and N. H. Silverman, "Evaluation of Fetal Heart Dimensions from 12 Weeks to Term," *Am J Cardiol*, vol. 87, pp. 594–600, 2001.
- [76] S. L. Kuntharee Traisrisilp, Fuanglada Tongprasert, Kasemsri Srisupundit and M. Theera Tongsong, "Reference Ranges for the Fetal Cardiac Circumference Derived by Cardio– Spatiotemporal Image Correlation From 14 to 40 Weeks' Gestation," *J Ultrasound Med*, vol. 30, pp. 1191–1196, 2011.
- [77] S. Luewan, Y. Yanase, F. Tongprasert, K. Srisupundit, and T. Tongsong, "Fetal cardiac dimensions at 14-40 weeks' gestation obtained using cardio-STIC-M," *Ultrasound Obstet. Gynecol.*, vol. 37, no. 4, pp. 416–422, 2011.
- [78] "Cardiac screening examination of the fetus: guidelines for performing the 'basic' and 'extended basic' cardiac scan," *Ultrasound Obs. Gynecol*, vol. 27, pp. 107–113, 2006.
- [79] T. L. Szabo and T. L. Szabo, "8 WAVE SCATTERING AND IMAGING," in *Diagnostic Ultrasound Imaging*, 2004, pp. 213–242.

- [80] P. Vergani, S. Cozzolino, E. Pozzi, M. S. Cuttin, M. Greco, S. Ornaghi, and V. Lucchini,
 "Identifying the causes of stillbirth: a comparison of four classification systems," *Am. J. Obstet. Gynecol.*, vol. 199, no. 3, pp. 319.e1–319.e4, 2008.
- [81] A. Mert, M. Sezd [·] I, and A. Akan, "A test and simulation device for Doppler-based fetal heart rate monitoring," vol. 23, pp. 1187–1194, 2015.
- [82] J. A. Jensen, "Speed-accuracy trade-offs in computing spatial impulse responses for simulating medical ultrasound imaging," *Publ. J. Comput. Acoust.*, vol. 9, no. 3, pp. 731– 744, 2001.
- [83] H. Azhari, "CH12: Safety and therapeutic applications," pp. 287–312.
- [84] D. L. Miller, "Safety Assurance in Obstetrical Ultrasound," *Semin. Ultrasound, CT MRI*, vol. 29, no. 2, pp. 156–164, 2008.
- [85] P. ten Hoope-Bender, K. Stenberg, and K. Sweeny, "Reductions in stillbirths a more than a triple return on investment," *Lancet*, vol. 387, pp. e14–e16, 2016.
- [86] J. Gardosi, S. Giddings, S. Buller, M. Southam, and M. Williams, "Preventing stillbirths through improved antenatal recognition of pregnancies at risk due to fetal growth restriction Stillbirth: a public health challenge," 2014.
- [87] R. Rabinowitz, E. Persitz, and E. Sadovsky, "The relation between fetal heart rate accelerations and fetal movements.," *Obstet. Gynecol.*, vol. 61, no. 1, pp. 16–8, Jan. 1983.

LIST OF FIGURES

Figure 2-1: Fetal monitoring system. An ultrasound transducer measures fetal heartbo while a tokodynamometer measures uterine activity [17]	eat 5
Figure 2-2: Non-stress test shows normal fetal heart rate accelerations	6
Figure 2-3: Umbilical Doppler velocimetry. Normal umbilical artery blood flow as seen with a forward flow in diastole and normal S/D ratio [12]	n 8
Figure 2-4: Kick counter wristband [37]	10
Figure 2-5: <i>Kickme- baby kicks counter Android mobile application</i> [38]	10
Figure 2-6: Fetal Doppler heart sound monitor device displaying fetal heart rate [39].	.11
Figure 2-7: <i>Baby`s heartbeat listener mobile application</i> [40]	11

Figure 3-1:	<i>The model of the one-dimensional wave propagation hitting the boundary.</i>
Figure 3-2:	The schematic of the single element transducer [54]18
Figure 3-3:	The schematic of a piezoelectric array transducer [54]19
Figure 3-4: relative to a	Doppler frequencies seen by observers at different location and at the angles the direction of the source: (A) 0°, (B) 90°, (C) 270°, (D) 45° [57]21

Figure 3-5: Sampled signal from blood vessel. The left graph shows the received signals for each transmitted pulse, and the right graph is the sampled signal [49].....22

Figure 4-1: Anatomy of the woman in the third trimester of gestation	5
Figure 4-2: <i>Schematic drawing of the woman abdomen, 1 – abdominal wall depth, 2 - uterine wall thickness, 3 – posterior uterine wall depth.</i> 27	7
Figure 4-3: SFH measurements versus gestational age, provided by midwives fromHorten kommune helsetjenesten for barn og unge	8
Figure 4-4: Schematic of SFH measurement procedure. 29	9
Figure 4-5: Piston transducer divided into 1mm mathematical elements	0
Figure 4-6: Ultrasound image of the fetal heart (right) and its schematics with identified heart constituents: LV (left ventricle), RV (right ventricle), ventricular septum, moderator band, pulmonary veins, atrial septum and crux [78]	ו 1
Figure 4-7: Heart model for Field II simulation	2

Figure 4-8: Heart movement assumption used for simulation
Figure 4-9: <i>The flow chart of the simulation model, where r1 and r2 are the heart wall radii, R is a radius of the transducer aperture, f0 is the central frequency</i> 34
Figure 4-10: Transducer placement assumption for simulation of Case study A35
Figure 4-11: <i>Possible movements of the transducer and a fetus assumed in simulation for case study C and D.</i>

Figure 5-1: <i>B-mode images of the phantom with varied number of scatterers (N): (a)</i> N=1000, (b) N=10 000, (c) 200 000 and (d) N=1 000 000	.40
Figure 5-2: The velocity of the heart movement for one cardiac cycle	.41
Figure 5-3: <i>B-mode image of the heart phantom located at 87mm from the transducer</i> <i>surface.</i>	r 44
Figure 5-4: <i>Beam profile for a round flat aperture with 5 mm radius with a heart position in the pressure field for case study A.</i>	.45
Figure 5-5: <i>RF data received by a transducer.</i>	.46
Figure 5-6: Individual RF lines for non-moving heart wall	.46
Figure 5-7: The change of the heart radii over one cardiac cycle	.47
Figure 5-8: Summed received signals from the heart phantom for case study A	.47
Figure 5-9: Received signals for each RF line	.48
Figure 5-10: <i>Sampled signal. Time shifts between the received signals for 11 transmit pulses. Case study A.</i>	<i>ted</i> 49
Figure 5-11: <i>Graph of the measured and actual velocities of the heart scatterers for ca</i> <i>study A</i>	<i>ase</i> 49
Figure 5-12: <i>M-mode image of one cardiac cycle.</i>	.50
Figure 5-13: <i>B-mode image of the heart phantom for case study B. The heart wall boundaries are hardly distinguishable.</i>	.50
Figure 5-14: Summed received RF lines for case study B	.51
Figure 5-15: <i>Received signals for 11 transmit pulses for case study B. The point indica</i> the maximum amplitude found in overall received signals	<i>ites</i> 51
Figure 5-16: <i>Sampled signal. Phase shift between the consecutive received signals for</i> <i>case study B</i>	53
Figure 5-17: Measured and actual velocities of the heart scatterers. Case study B	.53
Figure 5-18: <i>M-mode image of one cardiac cycle.</i>	.54

Figure 5-19: B-mode image of the heart phantom for case study C	54
Figure 5-20: <i>Beam profile of the 5 mm transducer aperture and position of the hea</i> the pressure field. Case study C	<i>rt in</i> 55
Figure 5-21: <i>Received RF lines for 11 transmit pulses for case study C with a transparent aperture of 5 mm.</i>	<i>ducer</i> 55
Figure 5-22: Individual RF lines for 11 transmit pulses. Case study C	56
Figure 5-23: <i>Sampled signal. Phase shifts between the successive signals for 11 tra</i> <i>pulses.</i>	<i>nsmit</i> 57
Figure 5-24: Measured and actual velocities of the heart scatterers. Case study C	57
Figure 5-25: The beam profile for a round flat transducer with 2 mm aperture. The position of the heart is shown in the pressure field. Case study C	, 58
Figure 5-26: Summed received RF lines for 11 transmit pulses. Case study C	58
Figure 5-27: <i>Individual RF lines for 11 transmit pulses. Transmit and received aper</i> <i>radius is 2 mm. Case study C.</i>	<i>rture</i> 59
Figure 5-28: Sampled signal. Phase shift between the successive received signals	60
Figure 5-29: Actual and measured velocity of the heart phantom. Case study C	60
Figure 5-30: <i>B-mode image of the phantom placed at 107 mm away from the trans surface.</i>	<i>ducer</i> 61
Figure 5-31: <i>Beam profile for a transducer of 5 mm aperture. The heart is position the pressure field</i>	<i>ed in</i> 61
Figure 5-32: Summed RF lines for received signals for case study D	62
Figure 5-33: RF lines for 11 transmit pulses. Case study D	62
Figure 5-34: Sampled signal. Phase shift between the successive received signals	63
Figure 5-35: Measured and actual velocity of the heart phantom. Case study D	64
Figure 5-36: <i>Schematic representation of the transmit/receive system for ultrasou transducer.</i>	<i>nd</i> 65
Figure 5-37: Hardware set up for transmit and receive system for a transducer	65
Figure 5-38: (a) <i>M-mode image and (b) phase difference between the successive received signals.</i>	66
Figure 5-39: (a) M-mode image and (b) phase difference between the successive received signals.	66

Figure 6-1: *The accuracy of the pulse-echo impulse response for rectangles and bounding lines transducer elements as a function of the sampling frequency* [82].......69

Figure 6-3: Beam profile for a one element convex transducer with 5 mm aperture......71

LIST OF TABLE

Table 2-1: Examinations tests and their drawbacks for fetal well-being assessment. 8
Table 2-2: Fetal well-being assessment parameters
Table 3-1: Speeds of sound, densities and characteristic impedances data for different mediumsand human tissues [48].14
Table 3-2: Temperature effects induced by ultrasound on a human body [59]. 23
Table 4-1: Woman size parameters27
Table 4-2: Posterior uterine wall depth and SFH change according to a gestation age
Table 4-3: Cardiac sizes for the third trimester [76], [77].
Table 4-4: Transducer design parameters
Table 4-5: Amplitude scaling factors for blood, heart wall and surrounding tissue
Table 4-6: Simulation of the heart phantom with varied number of scatterers
Table 4-7: Simulation design parameters for Case study A
Table 4-8: Design parameters
Table 4-9: Design parameters for case study C. 37
Table 4-10: Design parameters for case study D
Table 5-1: Heart rate, frequency of the heartbeat, f_h and corresponding maximum heart velocity, v max data
Table 5-2: Calculated Doppler frequencies for 2 MHz centre frequency. 43
Table 5-3: Time shifts between consecutive signals and measured velocity of the heartscatterers for case study A
Table 5-4: Time shift calculation between consecutive received RF lines and measured velocitiesof the heart scatterers. Case study B
Table 5-5: Time shifts calculation between consecutive received RF lines. Case study C. 56
Table 5-6: Time shifts calculation between consecutive received RF lines. 59
Table 5-7: Time shifts calculation between consecutive received RF line and measured velocityof the heart phantom