國立交通大學

電子工程學系 電子研究所碩士班

碩士論文

生醫元件可程式前端放大器

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AN 8-CHANNEL LOW NOISE FRONT-END CIRCUIT FOR NEURAL RECODING SYSTEMS

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生醫元件低雜訊前端放大器

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Abstract (Chinese)

1887年,英國科學家 Augustus Waller 第一次向世人發表了心電圖。1912年, 俄國生理學者 Vladimir Vladimirovich Pravdich-Neminsky 紀錄下腦部的電波訊號。 而十九世紀末科學家也紀錄到了肌肉裡的電訊號。自從這些生理訊號一一被發現 後,人類發現這些訊號和許多神經上的疾病或身體上的活動息息相關。所以深入 的研究這些生理訊號變成了重要的課題。

由於生理訊號有低頻、低振幅且獨特等特性,所以訊號的放大、低雜訊且有可調性成為訊號處理的一大挑戰。本論文即為實現一個可以用於放大Electrocochleography(ECoG)、Electromyography(EMG)和 Electrocardiography(ECG)等多種生理訊號的類比前端放大器,且為了能夠同時觀察多個通道,本設計使用八通道架構,此類比前端放大器包含了前級放大器,八通道多工器和可程式增益放大器。其中新架構的前級放大器可防止電極造成的直流電壓偏移飽和電路,高頻截止頻率可調於 1Hz-7KHz,放大增益 40 dB,輸入參考雜訊電壓在 0.5 Hz 到7 KHz 積分範圍為 0.9 μV rms,雜訊效率因子(NEF)為 1.3。

整體的類比前端放大器的放大增益為 40-74.1dB,輸入雜訊在 0.5 Hz-7 KHz 頻段積分為 5.1 μV rms,並且有很高的共模拒斥比和電源拒斥比,功率消耗 446μW,在台灣積體電路製造股份有限公司與國家晶片系統中心的幫助下,此類 比前端電路晶片將以 0.18 微米製程實現。

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An 8-Channel Low Noise Front-End Circuit for

Neural Recoding Systems

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Abstract (English)

This paper presents an 8-channel analog front-end (AFEA) circuit for neural

recording systems. An eight-channel chopper stabilized pre-amplifier, an 8-to-1

analog multiplexer, and a programmable gain amplifier are designed. It is designed

for amplification of different kinds of biopotential signals, such as electrocorticogram

(ECoG), electrocardiogram (ECG) and electromyogram (EMG). These biopotential

signals have the characteristics of small amplitudes, low frequency and variability.

The maximum gain of AFEA is 74.1dB. The high-pass corner can achieve as low as

0.8Hz and low-pass corner can be adjusted to several kilohertz to suit for different

kinds of biosignals. It is fabricated in TSMC 0.18 µm CMOS process and achieves an

input-referred noise of 5.1 µVrms (input referred noise of pre-amplifier is only

0.9μVrms), bandwidth up to 7 KHz while only consumes 446 μW at 1.8V power

supply. And the noise efficiency factor (NEF) is only 1.3 for pre-amplifier.

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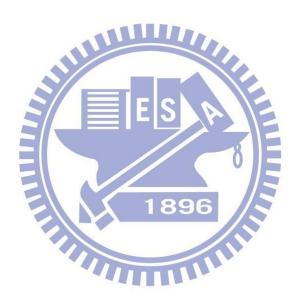
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Chapter1 Introduction

1.1 Background

In 1887, the British physiologist Augustus Waller published the first human electrocardiogram. In 1912, Russian physiologist Vladimir Vladimirovich Pravdich-Neminsky recorded the potential of the mammalian brain and evoked potentials on photographs and coins the term 'electrocerebrogram'. The scientist also recorded electrical activity during a voluntary muscle contraction in late 19th century. And then scientists discover these biopotential signals and some neurological disorders or physiological activities are closely related. It's become an important issue to explore the meaning behind the biopotential signals. The study is about the amplifier for neural recording applications.

The circuits of recording biopotential signals have to overcome various problems, while extracting the biopotential signals from the human body. These problems are not only due to the exceedingly weak characteristics of the biopotential signals but also due to the environment and the instruments that are being used during the signal acquisition. Therefore, the design of readout circuits for the biopotential acquisition systems requires a comprehensive understanding of not only the analog circuit design techniques but also the origin and the characteristics of the biopotential signals.

This chapter gives a brief introduction to the challenges of biopotential readout circuit. Section 1.2 introduces about the characteristics of the biopotential signals, which including Electroencephalogram (EEG), electrocorticogram (ECoG), electrocardiogram (ECG) and electromyogram (EMG). Section 1.3 describes review

on fronted-end amplifiers for neural recording systems. Section 1.4 explains the motivation and specifications of this work. Section 1.5 briefly introduces about main results and thesis organization.

1.2 Overview of Biopotential signals

Biopotential signals are electric potential signals that is measured between points in living cells, tissues, and organisms, and which accompanies all biochemical processes. These cells generate an potential when they are stimulated and they present a resting potential electrically. The membrane potential of an inactive cell is called the resting potential. The potentials are generated by the exchange of ions between inside and outside cell. Intracellular concentration of potassium (K⁺) ions is 30-50 times higher than extracellular it. Sodium ion (Na⁺) concentration is 10 times higher outside the membrane than inside. In resting state, the membrane is permeable only for potassium ions. K+ concentration of the interior of the cell is much higher than the exterior. Therefore, a diffusion gradient of K+ occurs towards the exterior of the cell making the interior more negative relative to the exterior, which results in an electrical field direct inward cell being built up. In steady state, the diffusion gradient of the K+ ions is balanced by the electrical field and the equilibrium is reached with a polarization voltage of nearly -70 mV. When membrane stimulation exceeds a threshold level of about 20 mV, so called action potential occurs. Sodium (Na⁺) and potassium (K⁺) ionic permeability of the membrane is changed. Sodium ion permeability increases very rapidly, allowing sodium ions to flow from outside to inside, result in the polarity of cell being more positive. Oppositely, the more slowly increasing K⁺ ion permeability permits K+ ions to flow from inside to outside, thus

returning membrane potential to its resting value. As the potential reaches to 40 mV, the permeability of the membrane to Na+ ions decreases and to K+ increases, resulting in the membrane potential swiftly decrease towards its rest state. The biopotential signals, such as EEG, ECG, EMG, are the result of several action potentials produced by a combination of different cells [1].

Many organs in the human body, such as the heart, brain, and muscles, display their function through electric activity [2]. The heart, for example, produces a signal called the electrocardiogram (ECG). The brain produces a signal called an electroencephalogram (EEG) or electrocorticogram (ECoG). The activity of muscles, such as contraction and relaxation, produces an electromyogram (EMG). Measurement of these biopotential signals can help doctor to diagnose what happened with these organs. Therefore, more and more studies are about the biopotential signals readout circuit. Biopotential signals have the characteristics of small amplitude, low frequency and variability. Table I shows the frequency and amplitude characteristics of EEG, ECoG, ECG and EMG. The following articles briefly introduce about these biopotential signals individually:

Table I The electrical characteristics and applications of these biopotential signals [3].

		0 1 3	
Source	Amplitude (mV)	Bandwidth (Hz)	Clinical and Research Use
EEG	0.001-0.01	0.5-40	Seizure detection, the diagnosis
			of encephalopathy
ECoG	0.1-1	0.5-40*	Sleep studies, seizure detection,
			cortical mapping
ECG	1-5	0.05-100	Diagnosis of ischemia,
			arrhythmia, conduction defects
EMG	1-10	20-2K	Muscle function, neuromuscular
			disease, prosthesis

*Note: In order to observe the relations between KHz signals with seizure detection, the bandwidth is extended to 7KHz in this work.

EEG & ECoG:

The brain transmit electrical signals when subject's intention to do some action. These signals are generated by neurons in the cortical layers. There are several modalities to capture the electrical activities of brain in clinical practice. The methods are classified by the locations the electrodes are placed on, and by the spatial and spectral frequency of their captured signals. Electroencephalogram (EEG) is measured by electrodes are placed on scalp which is 2-3 cm away from the surface of the cortex. This is safest way to record brain activities. In contrast, electrocorticogram (ECoG) which need to use invasive electrodes is more unsafe. The recording electrodes are approximated on the cortical surface, and therefore ECoG has larger amplitude than EEG [4].

The EEG is typically described in terms of rhythmic activity and transients. By means of Fourier transform power spectrum from the raw EEG signal is derived. In power spectrum contribution of sine waves with different frequencies are visible. Although the spectrum is continuous, ranging from 0 Hz up to one half of sampling

frequency, the brain state of the individual may make certain frequencies more dominant [5].

The normal EEG wave is classified five wave groups. Delta waves arise in the frequency below 3.5 Hz. They happen in deep sleep, in children, and in serious organic brain disease. Theta waves include all the waves between 4 and 7 Hz. These occur mainly in parietal and temporal regions in children, but they also occur during mental stress in some adult, particular in feeling of disappointment and frustration. The best-known and most extensively studied rhythm of the human brain is the normal alpha rhythm. The frequency range is between 8 and 13 Hz. They are found in normal persons when they are awake in a quiet, resting state of cerebration. When the awake subject is asleep, the alpha waves disappear completely. Beta wave normally occur in the frequency range of 14 to 30 Hz, and sometime as high as 50 Hz. They can be divided into Beta I and Beta II. The frequency of about 14 to 26 Hz of they is called Beta I, affected by mental activity as alpha waves. The rest waves of Beta is Beta II, on the other hand, appear during intense activation of the central nervous system and during tension [6]. Gamma waves are patterns of brain waves in humans with a frequency >30 Hz. Recent research have pointed out GAMMA waves observation can help to comprehend epilepsy [7][8]

The intensities of brain waves on the surface of the brain may be as large as 10mV [6]. The amplitude of EEG measured on the scalp is only few micro-volts. The electrodes of ECoG are placed more close to cortical layer. Therefore, the amplitude is larger 100 times than EEG. It's important about noise issue at the level of amplitude. The noises come from physiological, environmental, and electronic sources. Physiological sources of interference are motion artifact, muscle noise, eye motion or blink artifact, and sometimes even heartbeat signals. Electrical interference arises from the usual sources: 60 Hz power lines, radio frequencies (RF), and electrically or

magnetically induced interference. Moreover, the electronic components in the amplifier also contribute noise. Good design and measuring techniques can mitigate the effects of such noise and interference. Electrode locations and names are specified by the International 10-20 system [9] for most clinical and research applications. This system ensures that the naming of electrodes is consistent across laboratories. In most clinical applications, 19 recording electrodes (plus ground and system reference) are used. A smaller number of electrodes are typically used when recording EEG from neonates. Additional electrodes can be added to the standard set-up when a clinical or research application demands increased spatial resolution for a particular area of the brain. High-density arrays (typically via cap or net) can contain up to 256 electrodes more-or-less evenly spaced around the scalp.

ECG:

Electrocardiogram (ECG) records the activities of heart by placing the electrodes on the skin. It is a noninvasive recording produced by an electrocardiographic device. In clinical applications, a lead system called 12-lead system has been used usually. Ten electrodes placed on the torso, arms, and legs are used for a 12-lead ECG. And these electrodes can be combined into a number of pairs. (For example: Left arm (LA), right arm (RA) and left leg (LL) electrodes form the pairs: LA+RA, LA+LL, RA+LL) The output from each pair is known as a lead. ECGs from these different leads help define the nature of the activity on a specific part of the heart muscle. Different types of ECGs can be referred to by the number of leads that are recorded, for example 3-lead, 5-lead or 12-lead ECGs. A 12-lead ECG is one in which 12 different electrical signals are recorded at approximately the same time and are common in research and clinical application.

The ECG signals at the surface of the body are small in amplitude, which make

the measurements be interfered with noise. An important consideration in good ECG signal acquisition is the use of high-quality electrodes. Electrodes made out of silver coated with silver chloride or of sintered Ag-AgCl material, are recommended. An electrolytic gel is used to enhance conduction between the skin and the electrode metal. Artifacts at the electrode-skin contact as well as electromagnetic interference from all sources must be minimized. Since ECG instruments are often used in critical-care environments, they must be electrically isolated for safety and protected from the high voltages generated by defibrillators. [3]

EMG

Muscle fibers generate electric activity whenever muscles are active. EMG signals are recorded by placing electrodes close to the muscle group. For example, a pair of electrodes placed on the biceps and another pair placed on the triceps can capture the EMG signals generated when these muscles contract. EMG signals recorded in this manner have been shown to give a rough indication of the force generated by the muscle group. Electrodes used for such applications should be small, securely attached and should provide recordings free of artifacts. Either silver-silver chloride or gold plated electrodes perform quite well, although inexpensive stainless steel electrodes may also suffice.

Since the frequency range of EMG signals is higher than that of ECG and EEG signals, and since the signals are of comparable or larger amplitudes, the problem of motion artifact and other interference is relatively less severe. Filtering can reduce the artifact and interference: for example, setting the bandwidth to above 20 Hz can greatly reduce the skin potentials and motion artifacts.

Recording activity directly from the muscle fibers themselves can be clinically valuable in identifying neuromuscular disorders. Therefore, invasive electrodes are needed to access the muscle fibers or the neuromuscular junction. Fine-needle

electrodes or thin stainless steel wires are inserted or implanted to obtain local recording from the fibers or neuromuscular junctions. [3]

Other biopotential acquisition systems follow similar principles of measurement. In conclusion, the most of the biopotential signals have small amplitude, low frequency, and narrow bandwidth. Therefore, a circuit compatible all classifications of biopotential signals is difficult to design. The effort is required to minimize the noise and interference by improving electrode design and placement and optimizing the amplifier circuit.

1.3 Review on Biopotential Amplifiers

It's necessary about a front-end biopotential amplifier for the readout system due to the slight amplitude biopotential signal. The amplifier must have two characteristics: to amplify the signal and property of low noise. Considering the portable systems and implantation of human security, the power dissipation of the circuit must be minimized for long-term biopotential signal monitoring. High common-mode rejection ratio (CMRR) and high input impedance design to reduce the effect of interference from human body at 50/60Hz. Another important issue is the differential DC offset, which is generated by electrodes, results in the saturation of the amplifier. In conclusion, design the biopotential amplifier has a lot of challenges to overcome. Recently, there are four structures of circuit apply to biopotential amplifier, and are described below.

A type of biopotential amplifiers is chopper stabilized amplifier. It uses chopper technique to cancel the dynamic offset and low frequency noise of the amplifier. The low frequency noise especially means flicker noise (i.e. 1/f noise), power spectral density of which is inversely proportional to the frequency. The operation principle of

the chopper technique is that the input signal is modulated to the chopping frequency, amplified and demodulated to the baseband. The offset and noise is modulated only once, and its frequency is shifted to the chopping frequency f_{chop} and its odd harmonics. These modulated offset and noise can be filtered by low pass filter. f_{chop} must be larger than two time than the bandwidth of the input signal to prevent aliasing. And that is limited by noise issue, too. To remove effectively the 1/f noise, it should be higher than the 1/f noise corner frequency (at which means the frequency thermal noise and flicker noise of a circuit contributes the same spectral density).

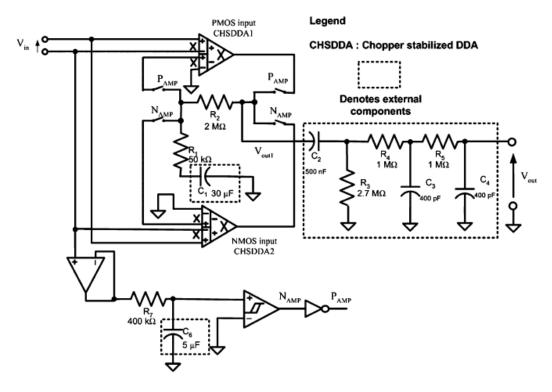
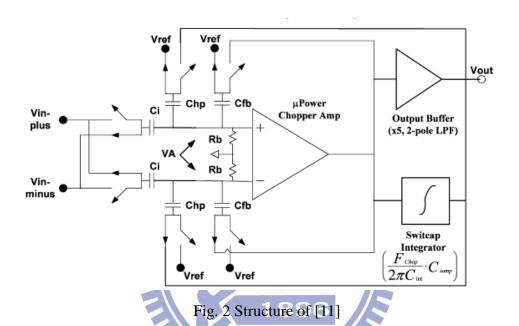


Fig. 1 The architecture of rail to rail instrumentation amplifier [10]

Fig. 1 shows a rail to rail instrumentation amplifier for portable EEG/ECG monitoring applications. The parallel input structure utilized PMOS and NMOS replica structures provides large input common-mode range. Furthermore, it use chopper stabilized technique to reduce noise and low frequency disturbance. To reject DC offset occurred at electrode-tissue interface, R1, R2, and C1 are used to generate a

very large time constants. For achieving the goal, the capacitors are made of off-chip device. The size of off-chip devices are usually large than integrated chip. Therefore, external component is NOT suitable for implantation. Architecture of [10] consumes large power due to replica input stage. Meanwhile, this ICs is designed for conventional EEG/ECG monitoring, them possess a narrow bandwidth (150 Hz).



Another chopper stabilized technique biopotential amplifier is shown in Fig. 2. The architecture consists of a micro-power chopper amplifier and multi-loop feedbacks. Gain of whole circuit is determined by the ratio of Ci and Cfb and output buffer, gain of which is generated by ratio of resistor. It improves the issue of using external component. The high pass corner is generated by Chp and switch integrator. The overall gain is associated with low cut-off frequency. It results in the complication of the design. Although it has very low power consumption, the bandwidth of [11] is merely 180 Hz. It possesses moderated noise performance and not well enough noise efficiency factor (NEF).

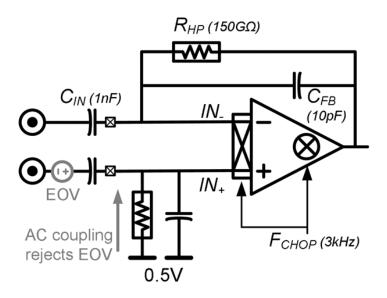


Fig. 3 Architecture of [12]

Fig. 3 shows the structure of [12]. That architectures are composed of a chopper stabilized amplifier and negative feedback loop is very similar to this thesis. That case has low input referred noise and very excellent noise efficiency factor performance. Switch capacitor resistors are used to generate a $150G\Omega$ resistance. Therefore, to let high pass corner closed to DC and provide appropriate gain, the external components are still used. Although it has very low power consumption, the bandwidth of [12] is only 100 Hz and the CMRR performance is NOT good enough.

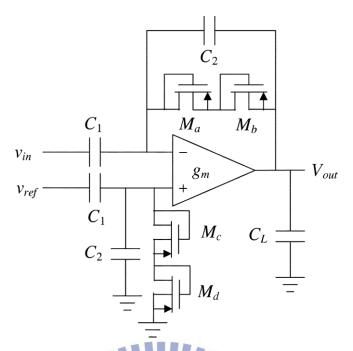


Fig. 4 The schematic of biopotential amplifier rejects DC offset [13].

The bandwidths of biopotential signals are from the millihertz range to few kilohertz range. Therefore, to rejects DC electrode offset without influence biopotential signal is difficult challenge. Fig. 4 shows the schematic of one of biopotential amplifier design [13], it claim that can generate a tens millihertz low frequency cutoff given by $1/(R_P \cdot C_2)$. C_2 is set to 200fP, and R_2 is a MOS-bipolar pseudoresistor consist of transistor M_a - M_d . The resistance of the pseudoresistor can reach more than 10^{12} ohm when the voltage across this device is between ± 0.2 V. the mid-band gain A_M of this architecture is set by C_1/C_2 . And -3dB corner is approximately $g_m/(A_M C_L)$ as C_1 , $C_L \gg C_2$. Where g_m is the transconductance of the operational transconductance amplifier.

There are a lot of interferences when measure the biopotential signals. Environmental noise (introduce in chapter 3) is one of them. Device electronic noise is the other. It is the important issue in the topic of biopotential amplifier design, since it limits the minimum signal level that the amplifier can process. Thermal and flicker noise is common type of device electronic noise. Since electrons in a conductor move

randomly by heat, the voltage across the conductor is fluctuant. The noise is called as thermal noise. The power spectral density is

$$S_{\nu}(f) = 4kTR, f \ge 0. \tag{1}$$

Where $K = 1.38 \times 10^{-23}$ J/K is the Boltzmann constant. T is absolute temperature. R is the resistance of the conductor. According to eq. (1) the power spectral density of thermal noise does not change by frequency. It's a typical white noise. The flicker noise spectral density, however, depends on the frequency. Due to many dangling bonds are formed at the interface between the gate oxide and the silicon substrate in a MOSFET, the interface rises to extra energy states. The movements of the change carriers are influenced by such energy states. The flicker noise can be modeled as a voltage source with the gate terminal, and the noise spectral density is expressed as

$$\overline{V_n^2} = \frac{\kappa}{c_{ox}wL} \cdot \frac{1}{f},$$
(2)

Where K is a process-dependent constant on the order of 10^{-25} V²F. C_{ox} is the gate oxide capacitance per unit area. Therefore, flicker noise is also called 1/f noise [14].

The noise level of amplifier depends on the power dissipation of circuit. To consider the tradeoff between power and noise and therefore a factor called noise efficiency factor (NEF) is first described by [15]. NEF normalize the power-noise performance and bandwidth of different circuits and can be expressed as:

$$NEF = V_{in,rms} \sqrt{\frac{2I_{tot}}{\pi \cdot V_t \cdot 4KT \cdot BW}}$$
 (2)

Where $V_{in,rms}$ is the total input referred voltage noise of the amplifier, I_{tot} is the total amplifier supply current, and BW is the -3dB bandwidth of the amplifier in Hertz. The NEF of a single bipolar transistor without 1/f noise is 1. Therefore, NEF is

a factor which implies the performance of the biopotential amplifier in comparison with a single bipolar transistor. In [13], the amplifier reduces the noise by using large MOS transistor size and the NEF of the circuit is down to 4.

1.4 Motivation

Some published amplifier circuit for biopotential signal applications have been reviewed and surveyed. According to the research mentioned before, an attempt to make further improvements about the noise, high PSRR and high CMRR of chip, based on proper performance and power dissipation. Furthermore, small chip area and no external component are suitable for implantation. Meanwhile, in recent study of the epilepsy monitoring, the wider bandwidth detection system is necessary for observing the relation between high frequency component of ECoG and seizure of epilepsy. Therefore, we would like to design an multi-channel analog front-end amplifier (AFEA) to achieve low noise, excellent NEF performance, wide tunable bandwidth, programmable gain, and rejecting DC offset without external components for neural recording system applications.

1.5 Main Results and Thesis Organization

An 8-channel analog front-end amplifier (AFEA) for neural recording systems is designed and fabricated. The AFEA consists of a new chopper stabilized pre-amplifier, 8-to-1 analog multiplexer (MUX), and programmable gain amplifier (PGA). The new pre-amplifier is designed with combining chopper stabilized and negative feedback with pseudoresistor to perform low noise and low frequency high

pass corner. The AFEA possess 40.3/50/60.8/74.1 (dB) four steps programmable gains. The high-pass corner can achieve as low as 0.8Hz and low-pass corner can be adjusted to several kilohertz to suit for different kinds of biosignals. AFEA achieves an input-referred noise of $5.1~\mu V$ rms, and the noise of pre-amplifier is only $0.9~\mu V$ rms, tunable bandwidth up to 10~KHz while only consumes $446~\mu W$ at 1.8V power supply. And the noise efficiency factor (NEF) is 1.3 for pre-amplifier. Meanwhile, the AFEA has more than 91dB~PSRR and 105dB~CMRR

The first chapter, chapter 1, introduces about the characteristics of the biopotential signals, the recent development of biopotential amplifier, the motivation and target specifications of this thesis, main results and thesis organization.

The second chapter, chapter 2, presents the flow of the circuit design includes three blocks of circuit: The new pre-amplifier combines a MOS-bipolar pseudoresistor and chopper technique. An 8-to-1 analog multiplexer and a programmable gain amplifier consists of capacitive-reset gain circuit are introduced after pre-amplifier. This chapter shows how to design them and theirs post-simulation results.

The chapter 3 of this thesis includes several parts. First, chapter 3.1 describes the consideration of the chip layout and shows the diagram of the layout. Second part introduces the measurement setup. Then, third part shows the results of measurement. The final part of chapter 3 describes the discussion of these circuits.

The last chapter, chapter 4, summarizes the major consideration of this thesis and concludes with suggestions for future investigation.

Chapter2 Circuit Design and Simulation Results

The block diagram of the implantable epilepsy detection and stimulation system can be described as shown in Fig. 5. The system includes signal process loop, monitoring and controller, and power delivery. Neural recording and stimulation loop is composed of electrode array, multi-channel pre-amplifiers, analog multiplexer (MUX), second stage amplifier (or programmable gain amplifier), analog to digital converter, DSP, Digital to analog converter, and driver for stimulation which are all implanted under skull. Monitoring and controller includes two transceivers, one is for implanted, and the other is placed outside. Power delivery supplies power to implanted device by coils. Some segments of circuits of power delivery and monitoring and controller are implanted with the loop. This thesis introduces about implementing a front-end circuit of the epilepsy detection and stimulation system. The AFEA consists of pre-amplifier, analog MUX, and programmable gain amplifier. And this chapter describes the design consideration, design flows, and post simulation results of the AFEA.

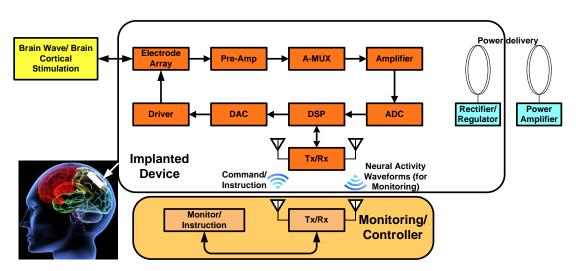


Fig. 5 An implantable epilepsy detection and stimulation system includes signal process loop, monitoring and controller, and power delivery.

2.1 Target Specifications

Therefore, an 8-channel biopotential amplifier should be designed. This work would try to realize an 8-channel front-end circuit in TSMC 018-µm technology under 1.8V supply voltage. Its specification is listed in Table III. The constructing circuit components of the AFEA would consist a pre-amplifier, aimed at possessing high common mode rejection ratio (CMRR), low noise, DC blocking, and around 40dB mid-band gain with proper power consumption; a 8-to-1 analog multiplexer is designed to provide the 8-channel signals could be selected by next stage circuit; a programmable gain amplifier provide various gains to select.

Table II Target specification of this work.

Parameter	Value (Unit)
Technology	TSMC 018μm
Power Supply	1.8 (V)
Input Voltage Range	100μ-1m (V)
Channel No.	1896
Gain	Programmable Gain in the range 100-5000
	(V/V)
High Pass Corner	<5 (Hz)
Low Pass Corner	adjustable in the range 1K-7K (Hz)
CMRR	≥66(dB)
PSRR	≥66(dB)

2.2 Design Consideration

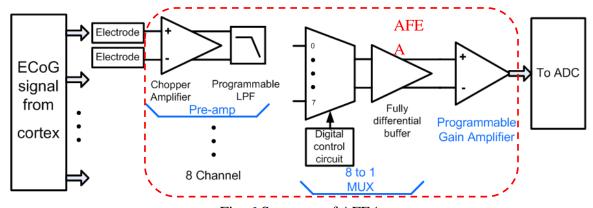


Fig. 6 Structure of AFEA.

Fig. 1 shows the main block of the circuits of this thesis. Since the system is designed for recording ECoG, the specifications of this work are made by the characteristics of ECoG. However, we make the specification of the front-end circuit flexible that let it could be appropriate for the other applications (see Table III).

Brain wave signal pass by electrode array into the pre-amplifier first stage of the AFEA. Like other biopotential amplifier as mentioned chapter 1, the brain wave like ECoG is a small amplitude signal (see Table II) and therefore the pre-amplifier must possess the characteristic of low noise, high common mode rejection ratio (CMRR), and high power supply rejection ratio (PSRR). Another reason of having high CMRR is for rejecting the DC offset voltage by differential electrodes. Hence, the pre-amplifier needs a high pass filter, too. And it also provides an adjustable high frequency cutoff. A new architecture of circuit is proposed for the pre-amplifier of the front-end circuit described in chapter 2.2.1. For practical application, the circuit needs to deal with multi-channel inputs. In this case, we choose eight-channel input to implement circuit. To reduce area and power of chip, an 8-to-1 channel analog multiplexer (MUX) is connected after pre-amplifier. The 2nd stage amplifier is composed of a Capacitive-reset gain circuit. It provides a programmable gain and stable voltage of amplified signals for ADC which follow the AFEA. These are described explicitly below.

2.3 Circuit Design

Circuit design is divided into three parts, and a detailed description of the following three sections.

2.3.1 Low Noise Pre-Amplifier Circuit

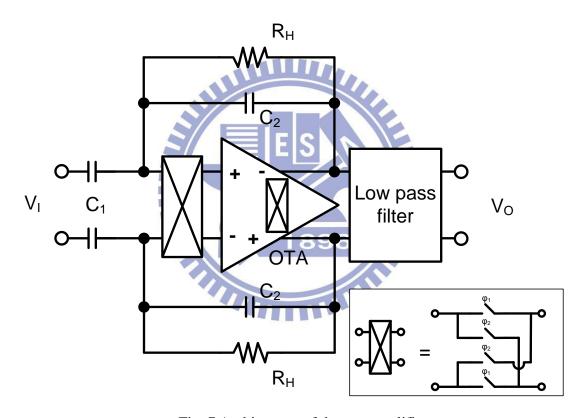


Fig. 7 Architecture of the pre-amplifier.

The structure of the first stage of AFEA consists of a negative feedback loop chopper stabilized amplifier with MOS-bipolar pseudoresistors and a passive low pass filter is shown in Fig. 7. As mentioned in chapter 1, the chopper stabilized technique could reduce the effect of the low frequency noise. To effectively achieve the purpose,

the chopping frequency f_{chop} must be larger than the 1/f noise corner frequency. The 1/f noise corner frequency of operational transconductance amplifier is about 60 KHz in this case (see Fig. 8). Therefore, the f_{chop} is chosen at 125 KHz.

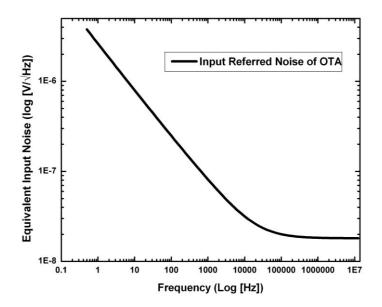


Fig. 8 Simulated noise power spectrum of the operational transconductance amplifier of pre-amplifier circuit.

And negative feedback and high impedance MOS-bipolar pseudoresistors R_H generate a low frequency cutoff like [16] S. Farshchi, A. Pesterev, E. Guenterberg, I. Mody, J.W. Judy; "An Embedded System Architecture for Wireless Neural Recording,", 3^{rd} International IEEE/EMBS Conf. on Neural Engineering, pp. 327-332, May 2007.] to block DC offset voltage. The R_H are implemented using PMOS with source-and-body connected shown in Fig. 9 (a). Its impedance can reach several hundred giga-ohm in simulation (see Fig. 9 (b)). The structure of the OTA is a telescopic op-amp with embedded chopper-modulators as illustrated in Fig. 10 and similar to the design in [17]. The chopper switches are placed at low impedance nodes

to avoid affecting the bandwidth of the amplifier.

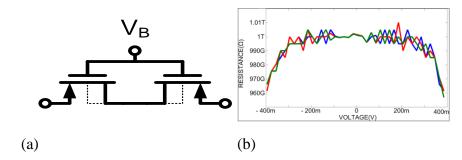
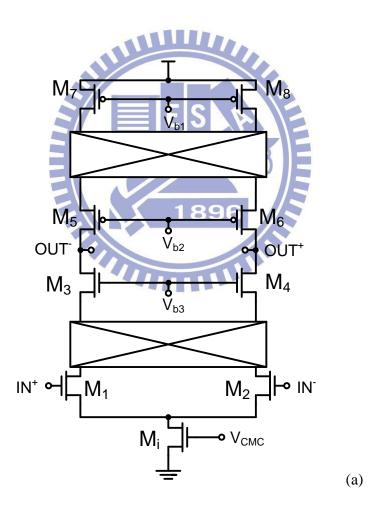


Fig. 9 (a) schematic of pseudoresistor R_H . (b) The diagram is the simulation results of the impedance of the pseudoresistor, when V_B is connected to VDD. Blue line, red line, and green line respectively represent results in the condition of TT corner, FF corner, and SS corner.



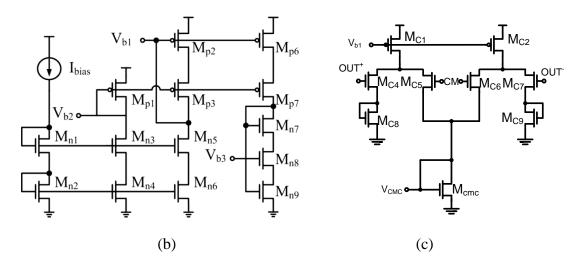


Fig. 10 (a) Schematic of OTA used in the pre-amplifier. (b) Bias circuit. (c) common-mode feedback circuit.

The telescopic op-amp has properties of low power, good stability, and low noise contribution [18]. Therefore, that is suitable for pre-amplifier circuit. A common mode feedback (CMFB) consists of two differential pair. The continue CMFB is more linear and could avoid the effect of chopper switch. The input referred noise of op-amp (0.1Hz to 10 KHz) is $14\mu V$ rms, the overall dc voltage gain is larger than 61dB. It has the properties of GBW more than 55MHz, phase margin > 74° , slew rate > $25V/\mu s$, common mode rejection ratio > 85dB, power supply rejection ratio (PSRR+) > 79dB, and PSRR- > 78dB. The op-amp of PGA dissipate $18\mu W$.

The block next the chopper stabilized amplifier is a low pass filter. The filter is implemented by passive device shown in Fig. 11. Its function is used to filter the modulated noise and offset voltage, and therefore its cutoff frequency must be much smaller than f_{chop} . For the specification the system of an adjustable high low corner, a gate-voltage control variable MOS resistors R_{f1} and R_{f2} are used. The signal transfer function of the low pass is given by

$$\frac{V_{out}}{V_{in}} = \frac{1}{1 + s \cdot \left(C_{f1}R_{f2} + C_L'R_{f1} + C_L'R_{f2}\right) + s^2 \cdot \left(C_{f1}C_L'R_{f1}R_{f2}\right)}.$$
 (4)

Where C'_L includes C_{f2} and output loading of filter C_L . Therefor, C_{f1} and C_{f2}

are selected 2pF and 1pF respectively.

as

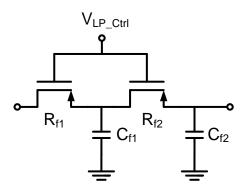


Fig. 11 Schematic of adjustable passive low pass filter.

To simplify the analysis of the pre-amplifier, a single-ended block diagram which noise source from each device have been include in is represented as shown in Fig. 12. Where $V_{n,OTA}$ represents the noise of the OTA, and $V_{n,RH}$ represents the noise comes from R_H . A(s) is the open loop gain of the OTA can be written as

$$A(s) = \frac{A_0}{s + \omega_0},\tag{5}$$

Where A_0 is DC gain of OTA, ω_0 is the dominant pole of the OTA. The transfer function of pre-amp is then given by

$$H(s) = \frac{-s \frac{C_1}{C_1 + C_2} \omega_o A_o}{s^2 + s \left(\omega_o + \frac{G_H}{C_1 + C_2} + \frac{C_2}{C_1 + C_2} \omega_o A_o\right) + \frac{(A_o + 1)\omega_o G_H}{C_1 + C_2}}.$$
 (6)

Assuming gain of OTA is approximated to A_0 in baseband. (6) can be rewritten

$$H(s) \approx \frac{-s \frac{C_1 A_0}{C_2 A_0 + C_1 + C_2}}{s + \frac{G_H (A_0 + 1)}{C_2 A_0 + C_1 + C_2}}.$$
 (7)

The expression shows low cutoff frequency is determined by G_H , C_1 , C_2 , and A_0 . And mid-band gain is $C_1A_0/(C_2A_0+C_1+C_2)$ which can be approximated to C_1/C_2 . This stage is designed to provide a gain of 40dB, settling C_1 to 10pF and C_2

to 100fF.

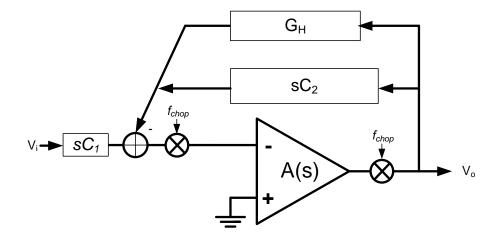


Fig. 12 Equivalent circuit model of pre-amplifier

2.3.2 Analog Multiplexer

The analog multiplexer includes a single stage CMOS switch array, switch control circuit, and a fully differential buffer as shown in Fig. 13. There are two kind of conventional switch array, one is multi-stage switch array (i.e., switch-tree array), and the other is what we adopt. The advantage of the former is the switch control circuit is simpler than the later. However, it generates some glitch in some case. For example, when control circuit trips switches from 011 to 100, owing to the LSB change faster than the other bits, switches might change from 011 to 010, than to 000, finally to 100. Nodes of switch-tree array pre-charge to some wrong voltage. Than charge injection is produced and leads to appear the glitch at output node. This affects the accuracy of the output of the multiplexer. Single stage switch array improves the issue and applies dummy switch (see Fig. 14) to reduce the effect of charge injection.

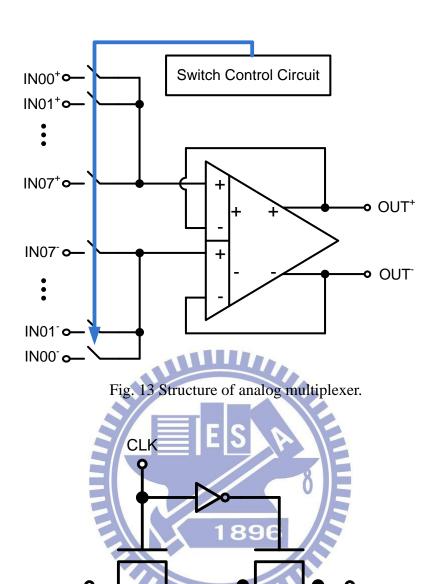


Fig. 14 Switch of analog multiplexer consists of NMOS switch, inverter, and dummy switch. The dummy switch size is half of NMOS switch.

Comparing with switch-tree array, the control circuit of the single stage switch array is more complicated. The switch control circuit includes 3-digital counter and 3-to 2³ decoder as shown in Fig. 15. 3-digital counter is composed of three resettable D-flip flops and generate three synchronous pulses and individual frequency are 250 kHz, 125 kHz, and 62.5 kHz. The three pulses are decoded to eight 500K channel/sec switch control signals (ch00-ch07) by 3-to 2³ decoder. In order to synchronize the eight switch signals, D-flip flops are place at output of decoder. It might reduce glitch

made by Asynchronous control signals.

The fully differential buffer is composed of a differential difference amplifier [19] (DDA) as shown in Fig. 16. The DDA circuit results in fully balanced outputs expressed as follow

$$V_{out}^{+} = -V_{out}^{-} = A_O \cdot [(V_{pp} - V_{pn}) - (V_{np} - V_{nn})].$$
 (8)

Where A_0 is the gain of DDA. The negative feedback topology with connection between outputs and both input differential pair is utilized to produce fully differential buffer.

An important issue of this stage is offset voltage. Since the next stage, programmable gain amplifier, provide up to 50 of gain and tolerate at least 1-V output swing, the offset voltage of this stage must be much less than 8mV. Fig. 17 shows the simulation result of the offset of the fully differential buffer by using Monte Carlo analysis.

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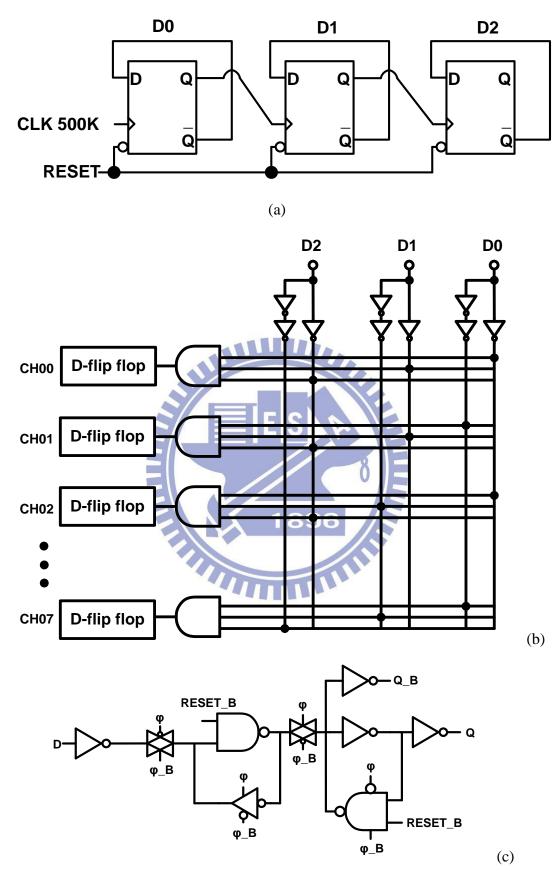


Fig. 15 Architecture of switch control circuit (a) 3-digital counter (b) 3-to 2³ decoder (c) resettable D-flip flop.

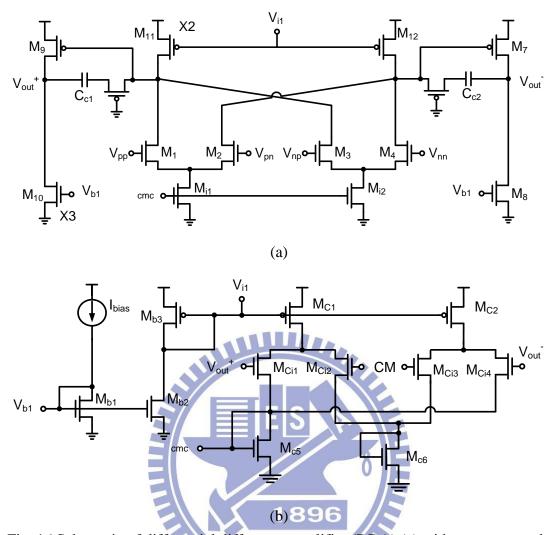


Fig. 16 Schematic of differential difference amplifier (DDA) (a) with common mode feedback and bias circuit (b).

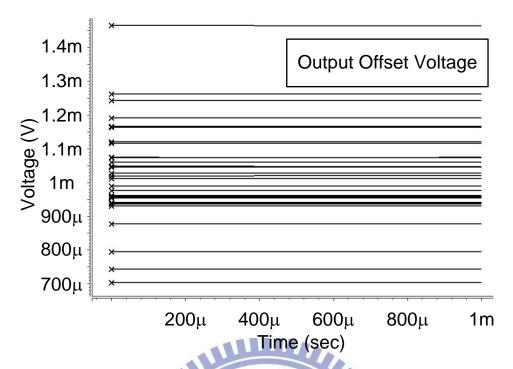
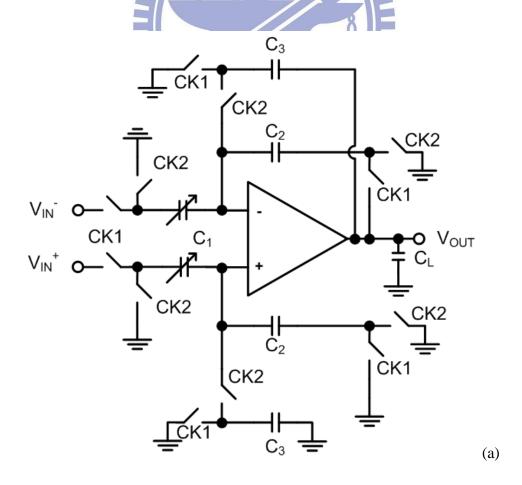


Fig. 17 According to the simulation result of Monte Carlo analysis, output offset of analog multiplexer is less than 1.5mV.

2.3.3 Programmable Gain Amplifier



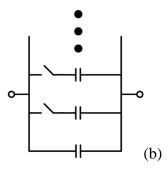


Fig. 18 (a)Schematic of PGA circuit. (b) The detail diagram of C_1 .

Analog to digital converter which AFEA connects to is designed in single input circuit. Therefore, the final stage of the front-end circuit would have a function of differential input to single-ended output and programmable gain. The programmable gain amplifier (PGA) circuit is realized by a single-ended capacitive-reset gain circuit as shown in Fig. 18 (a). It is based on the correlated double sampling (CDS) technique widely used to minimize errors due to reduction of offset voltage and 1/f noise. To see how this capacitive-reset gain circuit operates, consider it during the hold mode (CK2="1"), capacitor C_1 and C_2 both stores the input offset voltage of the op-amp. At this moment, C_3 is the feedback capacitor and was charged the voltage of the output node of previous clock phase. The principle sketch is shown as Fig. 19 (a). During the sample phase (CK1="1") is shown in Fig. 19 (b), than C_2 becomes the feedback capacitor, and the output voltage is given by

$$V_{OUT} = V_{OS} + V_{C_2}$$

$$= V_{OS} + \left(-V_{OS} - \frac{C_1}{C_2} \cdot V_{IN}\right)$$

$$= -\frac{C_1}{C_2} \cdot V_{IN}$$
(9)

Thus, the output voltage is independent of the op-amp offset voltage. And input signal import the circuit and the valid output voltage is generated.

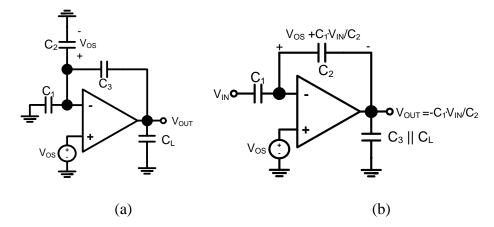


Fig. 19 Operating mode of PGA: (a) hold mode (b) sampling mode.

The transfer function of programmable gain amplifier in z domain is found to be

where
$$a = \left(\frac{1}{1 + \frac{C_1 + C_2}{C_2 \cdot A_0}}\right) \cdot \left(1 - \frac{C_1 + C_2}{A_0 \cdot \left(C_3 + \frac{C_1 + C_2 + C_3}{A_0}\right)}\right), \tag{11}$$

$$b = \frac{\frac{C_1 + C_2}{A_0} \cdot \left(C_2 + C_3 + \frac{C_1 C_2}{A_0}\right)}{C_2 \left(1 + \frac{C_1 + C_2}{C_2 A_0}\right) \left(C_3 + \frac{C_1 + C_2 + C_3}{A_0}\right)}, \tag{12}$$

and A_0 is the DC gain of op-amp. From the above equation, the gain of this stage is approximately the ratio of C_1 and C_2 at low frequency [21]. Therefore, two capacitor and switch arrays are used to be variable capacitors as shown in Fig. 18 (b). To match the speed of next stage, the analog to digital converter, sampling rate of programmable gain amplifier is 500 KHz the same as speed of analog multiplexer.

The design consideration of the PGA circuit is mainly focused on op-amp and switch design. In respect of switch, effect of charge injection and clock feedthrough, settling time, and linearity is generally considered about. To reduce effect of charge injection and clock feedthrough, the parasitic capacitor of switch should be minimized,

however, settling time limits size of it. Because transconductance is direct proportion to ratio of width and length, the minimum length is chosen. Fortunately the sampling rate we choose is not fast for this technology, we don't need to use extraordinarily large width to achieve. All switch is consist of NMOS switch except the switch connect output node to C_2 . A complementary analog switch is adopted in this structure due to 1-V swing output node.

The clock generation circuit used is shown in Fig. 20. It generates two phase and non-overlapping square wave. In this case, the delay time of the two phase clock signals is about 1ns.

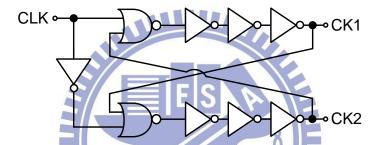


Fig. 20 Non-overlapping clock generation circuit.

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The op-amp is the primary building block of the PGA. The accuracy and speed of it are mainly determined by op-amp. During sampling mode illustrated in Fig. 19 (b), closed loop gain of PGA, A_L, is expressed as

$$A_{L} = -\frac{C_{1}}{C_{2}} \cdot \frac{1}{1 + \frac{1}{A_{0}} \cdot \left(\frac{C_{1} + C_{2} + C_{P}}{C_{2}}\right)}.$$
 (13)

Where C_P is parasitic capacitor of input of op-amp. Let feedback factor $\beta = C_2/(C_1 + C_2 + C_P)$, the error ε is represent as

$$\varepsilon = \frac{1}{A_O} \cdot \left(\frac{C_1 + C_2 + C_P}{C_2} \right) = \frac{1}{A_O \cdot \beta}.$$
 (14)

For achieving N bit resolution, error must less than $1/2^N$. Hence, the conditional expression $A_0 > 2^N/\beta$ can be got. In this case, the requirement of DC gain of the op-amp is greater than 94dB. The high gain op-amp not only reduces the gain error

but also increases the linearity.

On the other hand, the specification also limits the speed of the op-amp. By KCL equation at input node of op-amp and output node, the closed loop gain can be derived in the S domain

$$A_L(s) = -\frac{C_1}{C_2} \left[\frac{1 - s \cdot \frac{C_2}{G_m}}{1 + s \cdot \tau_a} \right]. \tag{15}$$

Where

$$\tau_a = \frac{C_1 + C_2 + C_P}{C_2} \cdot \frac{C_L' + [(C_1 + C_P)||C_2]}{G_m}$$
 (16)

, G_m is the transconductor of op-amp and $C_L{}'$ is $C_3 \mid\mid C_L$. Therefore, the unity-gain bandwidth of op-amp is

$$\omega_{u} = \frac{G_{m}}{C_{L}' + [(C_{1} + C_{P})||C_{2}]}.$$
(17)

The closed loop -3dB bandwidth is given by

$$\omega_{-3dB} = \frac{1}{\tau_a} = \beta \cdot \omega_u. \tag{18}$$

Then, the step response equation can be expressed as

$$V_{OUT}(t) = \Delta V \cdot \left(1 - e^{-\frac{t}{\tau_a}}\right) \tag{19}$$

Where ΔV is the ideal output voltage (should be equal to $V_{IN} \cdot C_1/C_2$). Let error $\varepsilon = [V_{OUT}(t) - \Delta V]/\Delta V$. To ensure $\varepsilon \leq 1/2^N$, the condition equation could be expressed as

$$\tau_a < \frac{t_{settle}}{N \cdot In2} \tag{20}$$

Where t_{settle} is settling time, it must be much less than duty period. Using the worth case (10 bits resolution, 500 KHz sampling rate, and gain of 50) to calculate, the requirement of the unity-gain bandwidth of the op-amp is more than 54 MHz.

The op-amp must achieve high gain and large output swing (comparing to 1.8V power supply). Comparing to single stage op-amp, the two stage op-amp more meet

these requirement. Therefore, a high gain two stage configuration op-amp [22] used in PGA circuit is shown in Fig. 21. It consists of a telescopic first input stage composed of M₁-M₈, and common source output stage, transistor M₉. This structure provides high open loop gain and large output swing.

The dominant pole is restricted to very low frequency due to the large telescopic structure output resistance of first stage. Therefore, if using a capacitor crossing input and output of second stage to supply Miller compensation, the bandwidth would be limited. To gain more bandwidth, the C_{C1} capacitor provides conventional current-buffering Miller compensation [23]. The C_{C2} capacitor is added to push the dominant pole to a lower frequency effectively killing peaks of frequency response. The op-amp has GBW more than 64 MHz. The overall dc voltage gain of it is larger than 99dB. It has the properties of phase margin > 63.6°, slew rate > $70V/\mu s$, common mode rejection ratio > 77dB, power supply rejection ratio (PSRR+) > 80dB, and PSRR- > 95dB. The op-amp of PGA dissipates $160\mu W$.

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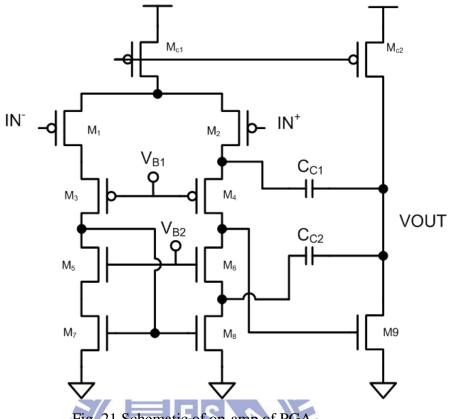


Fig. 21 Schematic of op-amp of PGA.

2.4 Post-Simulation Results

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Post-simulation is completed by SpectrRF and HSPICE simulators with process parameters of TSMC 0.18-µm mixed signal 1P6M+ salicide 1.8V/3.3V RF SPICE models. All the simulated results are considered under TT, FF, and SS three corner and process variation. Power supply for whole analog front-end circuit is 1.8Volt. The followings are the post-simulation results of the AFEA.

☆ Pre-Amplifier Circuit

The first stage of neural system front-end circuit is the pre-amplifier. It provides 40-dB gain to amplify those biopotential signals and a low cut-off frequency extremely close to DC. The Fig. 22 shows the frequency response of the first stage under three corners. The diagram reveals that pre-amplifier possesses a 38.8dB mid-band gain and a less than 2Hz high pass corner. The tunable low pass filter provides a high cut-off frequency with 1 kHz to 7 kHz range as shown in Fig. 23. The other important issue is about the noise. Fig. 24 is the simulated input noise power spectrum density of pre-amplifier under TT corner, and integration under this curve from 0.5 Hz to 7 kHz yields an rms noise voltage of 0.23 μV rms. noise efficiency factor (NEF) is only 0.34. The power consumption of pre-amplifier is 17.9μW. The simulation results of pre-amplifier circuit are summarized in Table IV.

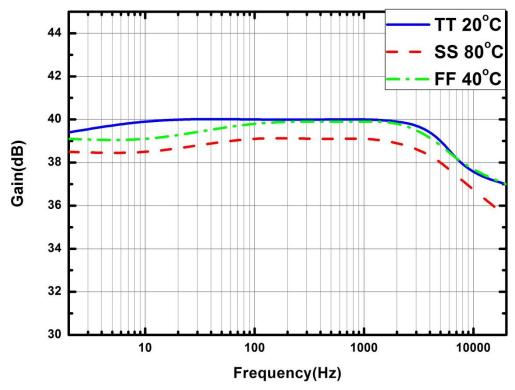


Fig. 22 The diagram shows the frequency response of pre-amplifier under in TT, SS, and FF corner and the control voltage of low pass filter (V_{LP_ctrl}) is 0.1V.

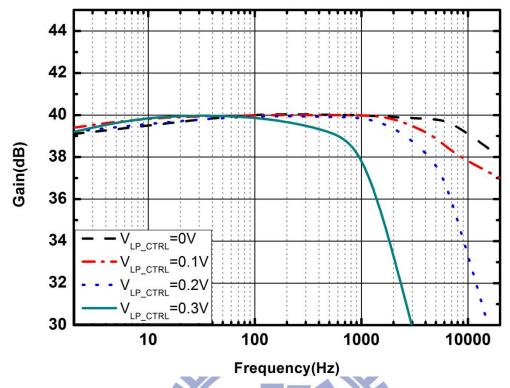


Fig. 23 Frequency response of pre-amplifier under TT corner.

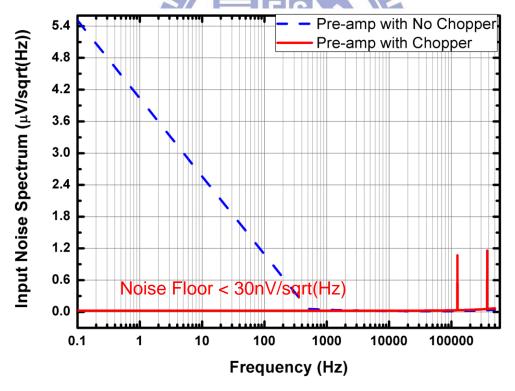


Fig. 24 Simulated pre-amplifier input-referred voltage noise spectrum.

Table III simulated characteristics of pre-amplifier circuit.

	Post-simulation
Parameter	Value (Unit)
Gain	38.5 (dB)
$\mathbf{f}_{\mathrm{CHOP}}$	125K (Hz)
Power Dissipation	17.95 (µW)
High Pass Corner Frequency	<2(Hz)
Low Pass Corner Frequency	1K~>10K (Hz)
Input referred noise (0.5 Hz-7 kHz)	0.23(µV rms)
NEF	0.34

☆ Analog Multiplexer

The analog multiplexer is composed of switch control circuit and fully differential buffer. Fig. 25 shows the eight channel switch control signal made by the digital circuit and it provides the multiplexer 500k channel/sec switch rate to work. And a FFT analysis is shown in Fig. 26, and the total harmonic distortion is 7.278e-05% (-122.7 dB) when input frequency is 10-KHz. It would tolerate maximum $200mV_{PP}$ input swing from output of pre-amplifier therefore simulated input range is about 700mV to 1.1V is far away large than target. The power dissipation of this stage is $77.83~\mu W$. Table V summarizes the simulated results of analog multiplexer.

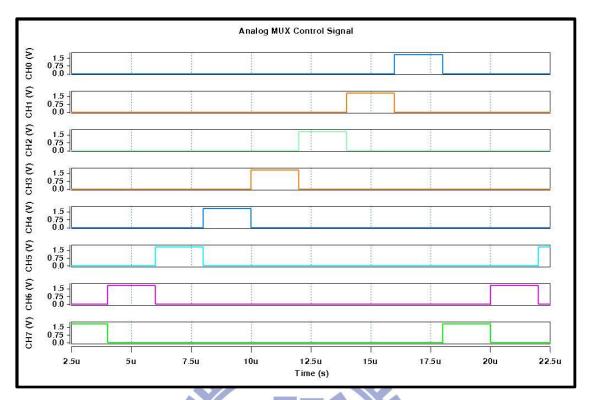


Fig. 25 A simulated controlling signals in the analog multiplexer.

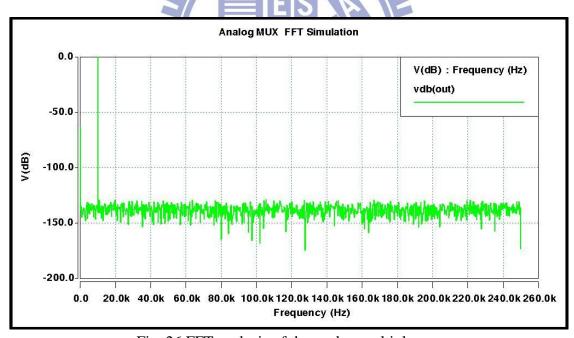


Fig. 26 FFT analysis of the analog multiplexer.

Table IV Simulated results of analog multiplexer.

	Post-simulatiom
Parameter	Value(Unit)
Channel No.	8
Switching Speed	500K(Number of Channel/sec)
Power Dissipation	77.83(µW)
Input Range	653~1142(mV)
THD	82.8dB (@Fin=10k)

☆ Programmable Gain Amplifier

The PGA circuit provides a tunable gain and 1-V output swing. In this case, Gain of 1/3/11/51 V/V are designed in PGA as shown in Fig. 27. The SNDR of PGA stage is greater than 62dB and SFDR is greater than 71 dB when full signal is used (see Fig. 28). The power consumption of PGA is 216 μ W. The summaries of simulated results of PGA are shown in Table VI.

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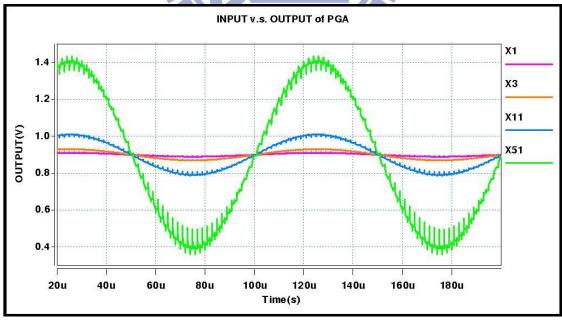


Fig. 27 The diagram shows the output waveforms under different gains as input frequency is 10-KHz and its amplitude is 20mV_{PP}. The purple line, red line, blue line, and green line represent output of gain of 1,3,11, and 51 respectively.

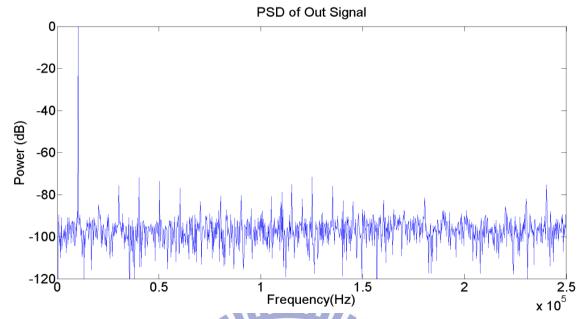


Fig. 28 Output power spectrum density of PGA.



	Post-simulation
Parameter	Value (unit)
Gain	1,3,11,51 (V/V)
SFDR (X1)	71.0 (dB)
(X51)	72.37
SNDR (X1)	62 (dB)
(X51)	63.22
THD (X1)	-59.2 (dB)
(X51)	-64.7 (dB)
$\mathbf{f}_{\mathrm{SAMPLE}}$	500 (KHz)
Bandwidth	265 (KHz)
Power Dissipation	216 (μW)
Output Range	1.6~0.2 (V)

Whole chip simulation results are summarized in Table VII. Programmable gain of front end circuit is 40/49.5/60.8/74.1 dB at mid-band and the cut-off frequency is less than 2 Hz. The tunable low pass filter provides the cut-off frequency of 1 kHz to 7 kHz achieved by a passive filter with tunable resistors controlled by external voltage. It is integrated from 0.5 Hz to 10 kHz to obtain rms noise voltage, whose value is 1.2 μ V rms. The noise efficiency factor (NEF) of whole circuit is only 4.9. The overall power consumption is 446 μ W. Whole chip simulated waveform are shown in Fig. 29 when an 10 kHz 200 μ Vpp sine waves as the input are used

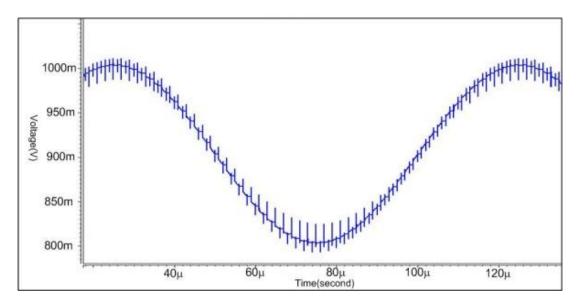


Fig. 29 Transient waveform of the AFEA when a 10 kHz 200 μ Vp-p sine wave is used.

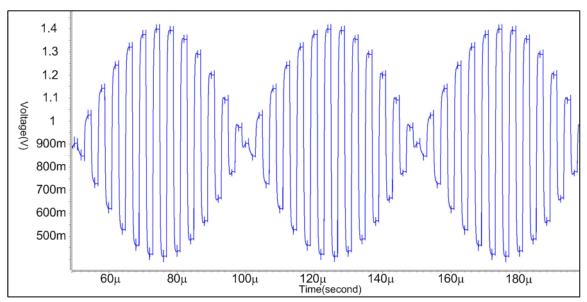
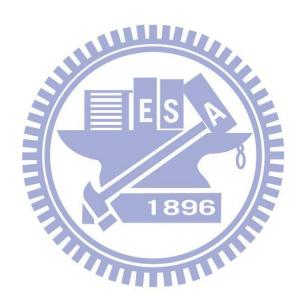


Fig. 30 The worst case simulation: The odd input channels and even input channels provide 180 different phase signals respectively. The frequency of input signals is 10-KHz and its amplitude is $200\mu V_{PP}$. The diagram shows the worst case output waveform. In this case, the AFEA is set in gain of 74.1dB.



Table VI Summary of AFEA simulated result.

	Post-simulation
Parameter	Value (Unit)
Gain (programmable)	40/49.5/60.8/74.1dB
Low Frequency Cut-off	<2 (Hz)
High Frequency Cut-off	1K~>10k (Hz)
(Tunable)	
Power Consumption	445.74 (μW)
CMRR(@f=10K)	120(dB)
PSRR(@f=10K)	95(dB)
Input referred noise	$1.11(\mu V \text{ rms})$
NEF(per channel)	4.74



Chapter3 Experimental Results

3.1 Chip Layout Descriptions

The whole chip of AFEA is fabricated in TSMC 0.18um Mixed Signal/RF process. It possesses 1 poly layer and 6 metal layers with low k inter-metal dielectric. Deep n-well topology is employed to surround the N-MOS device which allows the connection of source and body terminals to avoid body effect. MiM capacitor provides a low variation and low couple capacitor. High poly resistor, multi-Vt device and thick top metal are available for 1.8V/3.3V applications. It is suitable for logic, mixed signal, and RF designs. The layout of the front-end circuit is shown in Fig. 31. To avoid interference of digital clock to analog signal, the analog ground and power supply are separated from digital's. Furthermore, Digital signal lines closed to analog circuit are protected by shielding metal as illustrated in Fig. 32, and all NMOS of analog circuit are surrounded by deep n-well. With the aid of it, parasitic capacitance and noise coupling from substrate can be reduced.

The I/O pads are selected by STC Pure 1.8V Linear I/O Library in 0.18-µm CMOS Process. These I/O pads all include electro static discharge (ESD) protection circuits. Meanwhile, digital output buffer are embed in digital output pad to improve driving ability. To reduce the effect of couple of digital signals, digital I/O pads are put far away from analog ones. In this project, we select five analog observation points (outputs of pre-amplifier, outputs of analog multiplexer, and end of whole circuit.) three digital observation points (outputs of three digital counter).

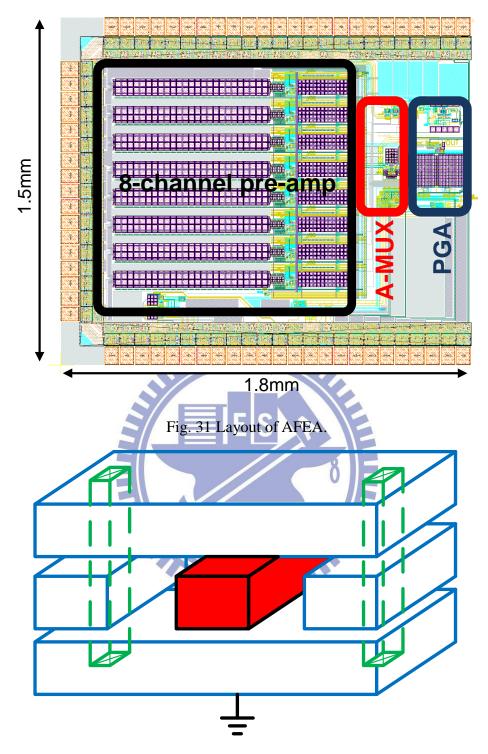


Fig. 32 Digital signal lines (Red block) closed to analog circuit are protected by shielding metal. Green block represents the VIA though all shielding metal.

3.2 Measurement Setup

This chip is directly bonded on printed circuit board (PCB). Package chip is excluded due to the reduction of more complicated parasites. And the chip microphotograph is shown in Fig. 33. On measurement consideration, we should restrict the environmental noise as possible as we can. The main environmental noise is the interference coupled from power line 50/60 Hz. Its common mode term is eliminated by front-end circuit's property of high CMRR and PSRR. However, other term of noise still affect the chip. The method is used to prevent that. We place a grounded connect metal plate under the PCB [24] and wrap up the sensitive line in aluminum foil. In case of long time observation, we take notice of some offset and disturbances at the output of circuit due to high impedance of pseudoresistors. To solve this issue, the control voltage of pseudoresistor (V_B) occasionally down to 0V to let the circuit "fast settle" [25]. The measuring environment and PCB of the chip are shown in Fig. 34. Fig. 35 to Fig. 37 present the instrument setup for different measuring requirement respectively.

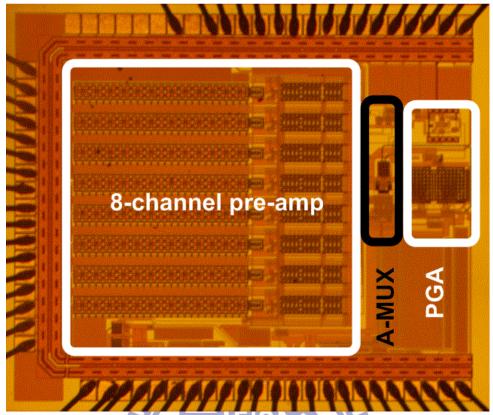


Fig. 33 Chip microphotograph.

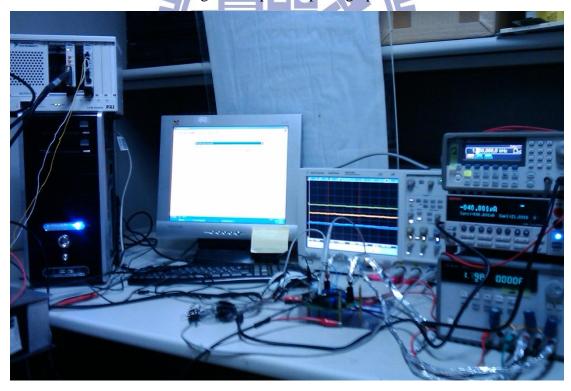


Fig. 34 Measuring environment

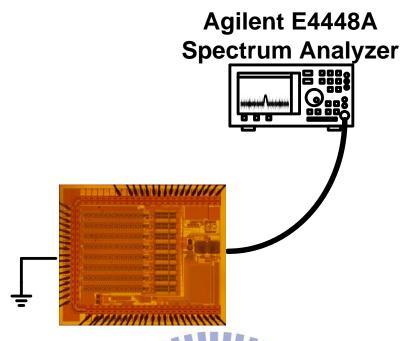


Fig. 35 Instrument setup for noise analysis.

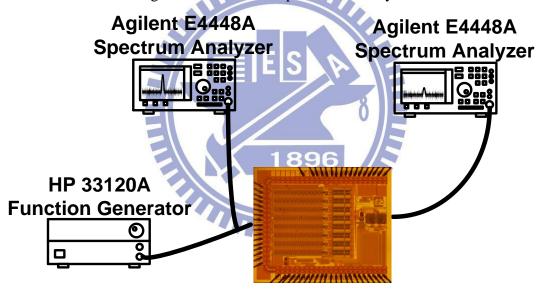


Fig. 36 Instrument setup for CMRR.

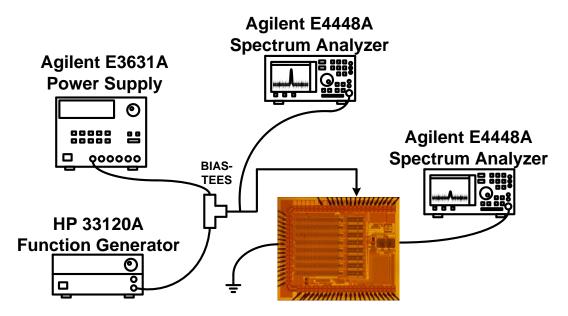


Fig. 37 Instrument setup for PSRR analysis.

3.3 Measurement Results

The measurement results are described as following. The frequency response of pre-amplifier circuit is shown in Fig. 38. We can observe the mid-band gain of pre-amplifier is about 40dB. Meanwhile, the expected tuning range of high frequency cut-off is covered in the bandwidth of tunable low pass filter. To measure the input referred noise, input ports are shorted to ground (see Fig. 35). The spectrum that we capture at output nodes is output noise spectrum. Therefore, we can divide amplifier gain into it, and get input noise spectrum as shown in Fig.36. The noise floor is less than $10\text{nV}/\sqrt{Hz}$ and the input referred noise integrated from 0.5Hz to 7 KHz is 0.9 μV rms.

To measure CMRR and PSRR of the front-end circuit, spectrum analyzer and signal generator are utilized to measure these specifications (see Fig. 36 and Fig. 37). The measured spectrum of common mode gain and gain from power supply are shown in Fig. 39 and Fig. 40 respectively. The AFEA possesses over 105dB CMRR

and 91dB PSRR. The power consumption of whole circuit is about 446.4 μ W. The voltage is enlarged 1/3.08/10.6/51 times by PGA circuit. Fig. 41 shows the frequency response of AFEA circuit, the tunable mid-band gains of whole circuit are **40.3/49.8/60.7/74.4** (**dB**) respectively. The input noise power spectrum density is shown in Fig. 42. The summary of measurement results are list in Table VIII.

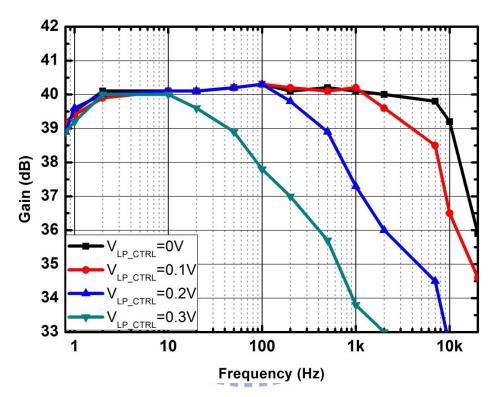
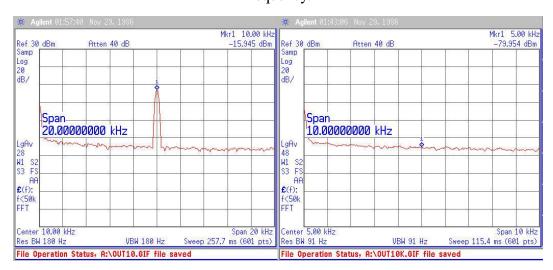


Fig. 38 The frequency response of Pre-amplifier circuit for different low pass corner frequency.



(a) (b)

Fig. 39 Spectrum results for measuring CMRR (a) input (b) output.

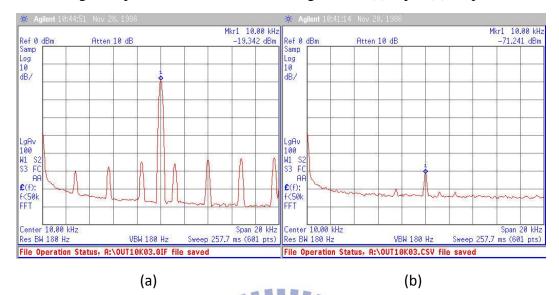


Fig. 40 Spectrum results for measuring PSRR (a) input (b) output.

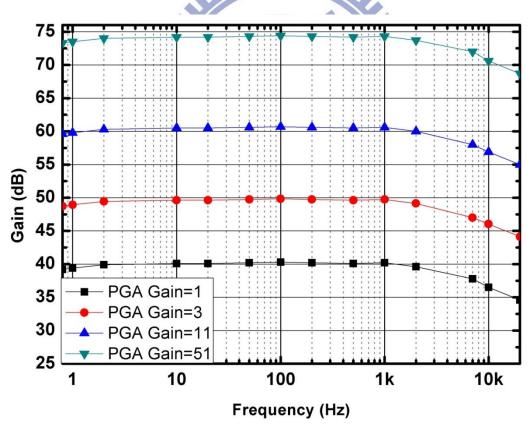


Fig. 41 Frequency response of AFEA

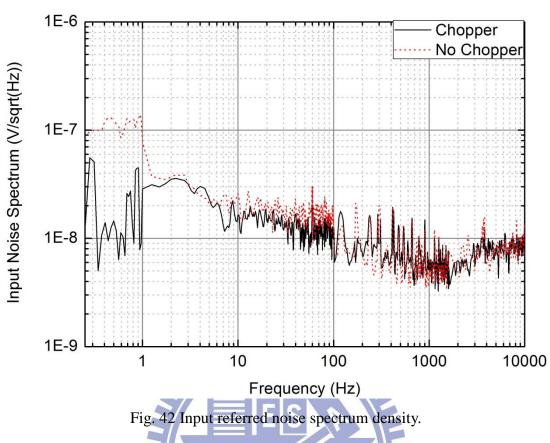


Fig. 42 Input referred noise spectrum density

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Table VII Measure Result of AFEA.

Spec.	parameter		
Pre-amplifier			
Mid-band gain	40.3(dB)		
Low frequency cut-off	<0.8 (Hz)		
High frequency cut-off	1k~7K(Hz)(tunable)		
Input referred noise (0.5 Hz- 7 KHz)	0.9 (μV rms)		
AFEA			
Mid-band gain	40.3/50/60.8/?(dB)		
Power consumption	446.4		
PSRR	91 (dB) @ 10KHz		
CMRR	>105@ 10KHz		
Input referred noise (0.5 Hz- 7 KHz)	5.1 (μV rms)		

3.4 Discussion

The measurement results didn't completely match with simulation results due to environments, instruments, PCB route or method of simulation. Fortunately, the experiment results almost conform to the target specifications. The Fig. 43 shows the comparison of measurement and post-simulation results about frequency response of pre-amplifier. It shows the two result match within 2.5dB gain. Although CMRR and PSRR are different from simulation result, they both meet the target specification, too. However, the measured noise floor is larger than simulated. From Fig. 44 to Fig. 47 show the biosignals measurement setups and results by AFEA. The all comparisons are minutely illustrated in Table X.

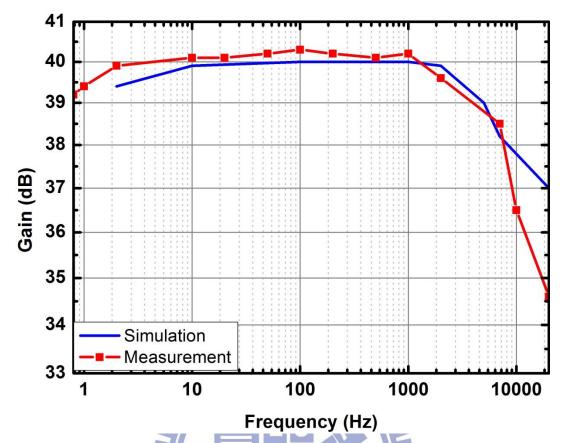


Fig. 43 Simulation and measurement results about frequency response of pre-amplifier are shown above when LP_ctrl=0.1V.

Table VIII Comparison of measurement and modified simulation results.

	Simulation	Measurement	
Pre-amplifier Pre-amplifier			
Mid-band gain	40(dB)	40.3(dB)	
Low frequency cut-off	<2(Hz)	<0.8 (Hz)	
High frequency cut-off	1k~7K(Hz)(tunable)	1k~7K(Hz)(tunable)	
Input referred noise	0.23 (µV rms)	0.9 (µV rms)	
NEF	0.34	1.3	
	AFEA		
Mid-band gain	40/49.5/60.8/74.1(dB)	40.3/50/60.8/?(dB)	
Power consumption	$445.74(\mu W)$	446.4(µW)	
PSRR	95 (dB) @ 10KHz	91 (dB) @ 10KHz	
CMRR	120 (dB) @ 10KHz	>105@ 10KHz	
Input referred noise	1.11 (µV rms)	5.1 (µV rms)	
NEF	4.74	21,8	

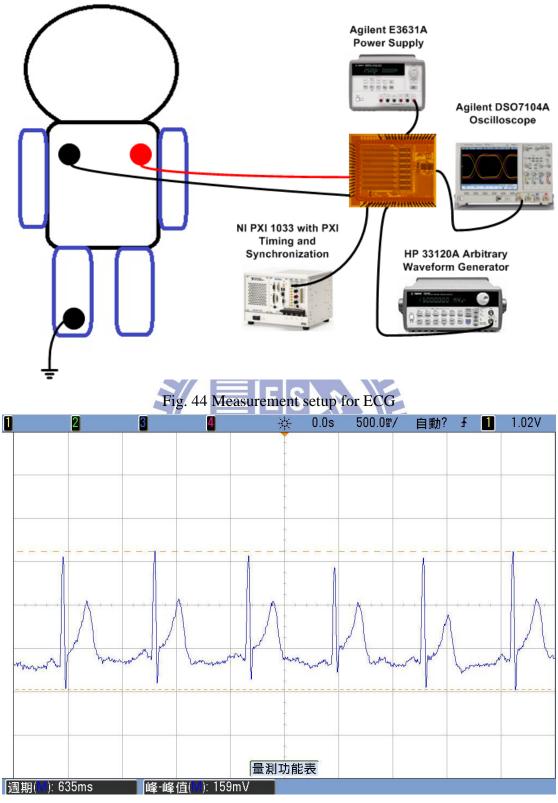


Fig. 45 ECG waveform measured at output of chopper stabilized pre-amplifier

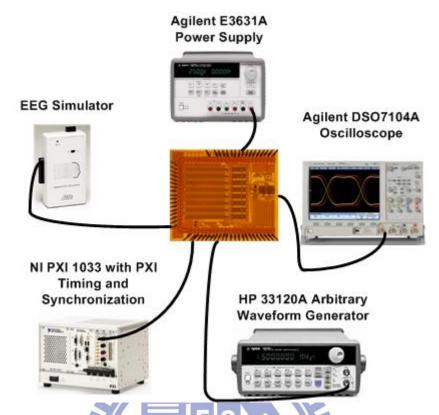


Fig. 46 Measurement setup for EEG (signal generated by EEG simulator)

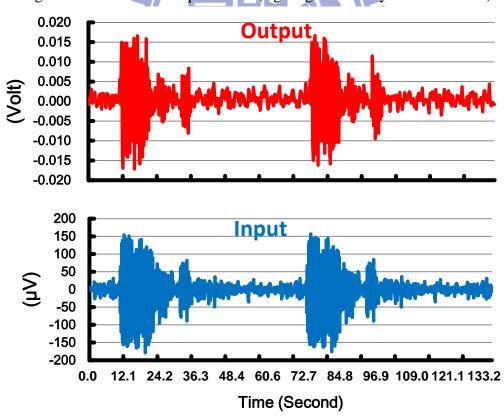


Fig. 47 EEG waveform measured at output of chopper stabilized pre-amplifier

Table IX Performance comparison with other Biopotential amplifier.

Spec.	This Work	[10]	[11]	[12]	[13]
K	· •	TCAS-I2005	JSSC2007	JSSC2010	JSSC2003
Technology	0.18μm ^a	0.5μm ^a	0.8μm ^a	0.18μm ^a	1.5μm ^a
Architecture	pre-amp+ 8	8 to 1 MUX+	Instrumentation	Instrumentation	Pre-amp
	to 1 MUX+	RRIA+ PGA	amplifier	amplifier	
	PGA				
technique	Chopper	Chopper	Chopper	-	-
	stabilized	stabilized	stabilized		
Use external	NO	YES	YES	YES	NO
component					
channel	8	1	1	1	1
Chip Area	1.8X1.5	2.3X2.1	0.7 ^b	0.3^{b}	2.2X2.2 ^c
(mm2)			N. S.		
Supply voltage	1.8	±1.5	1.8/3.3	1V	±2.5
(V)					
Power	17 ^d /446	1335 ^f	1.8/3.3	3.5	80
consumption	(pre-amp/AF		9 14	2	
(μW)	EA)		0		
DC gain (dB)	40.3/50/60.8	0-80	41/50.5	60	39.5
D 1 111 (T)	/74.1	0.5 150	808080 5 10	0.2.100	0.025.7.2
Bandwidth (Hz)	0.8~1k-7K	0.3~150	0.05/0.5/2.5~18	0.2~100	0.025~7.2
CMRR (dB)	105	117	80	60	K 83
PSRR (dB)	91	52	80	-	85
Input noise PSD	10/55	-	100	130	21
(nV/√Hz)	(pre-amp/AF		100	130	21
(1117)	EA)				
Input referred	0.9/5.1e	0.86 ^{e f}	0.95 ^e	1.3 ^e	2.2 ^e
noise (μV rms)	(pre-amp/AF				
	EA)				
NEF	1.3/21.8	56.9 ^f	5.4	0.96	4
	(pre-amp/				
	AFEA)				
_	_				

 a CMOS b w/o PAD c 6 amplifier d simulation result c integrated range of rms noise voltage is from 0.5Hz to 7 KHz for this work, from 0.3Hz to 150Hz for [10], from 0.05Hz to 100Hz for [11], from 0.5Hz to 100Hz for [12], and from 0.5Hz to 50KHz for [13] f whole circuit

Chapter4 Conclusions and Future Work

4.1 Conclusion

An 8-channel analog AFEA for neural recording system with integrated pre-amplifier, analog multiplexer, programmable gain amplifier has been designed, fabricated and tested in TSMC 0.18-µm CMOS technology. A new pre-amplifier with combining pseudoresistor and chopper stabilized technique has been proposed and protect neural signal from disturbance of internal noise. The whole AFEA circuit occupies an area of 2.7 mm², this is mainly dominated by MIM-capacitors array and a large region on the chip is filled with dummy metal, poly and oxide required to raise the yield rate.

The pre-amplifier possesses less than 0.8Hz low cut-off frequency to block DC offset, 200Hz ~10 KHz tunable low pass 3dB bandwidth, and 40.3dB mid-band gain to amplify the small amplitude signals. The input noise of power spectrum density is less than 10 nV/sqrt(Hz). It is integrated from 0.5 Hz to 7 kHz to obtain rms noise voltage, whose value is 0.9 μ V rms. The noise efficiency factor (NEF) of pre-amp is only 1.3.

The analog front-end circuit, power consumption of which is $446\mu W$ is tested under 1.8-V supply. As mentioned above chapter, AFEA has characteristics of 95dB PSRR and 120dB CMRR to prevent effect from environmental disturbance. Input referred noise integrated within baseband is 5.1 μV rms. The noise efficiency factor of the AFEA is 21.8. The comparison of specifications of others biopontial amplifier are shown in Table X.

4.2 Future Work

- 1. The whole AFEA circuit will be integrated into the implantable epilepsy detection and stimulation system and it will be test in vivo. The SOC will be packaged in a special case passing FDA inspection.
- 2. For long time monitoring biopotential signal, fast settle is necessary. Therefore, a clock generator should be designed on chip to generate the "fast settle" signal.
- 3. The proposed pre-amplifier has an advantage of their NEF performance due to decline of noise. However, the rear end circuit might dominate the noise source. For more thorough and complete front-end circuit design, we should diminish the noise and power consumption of analog multiplexer and PGA circuit.
- 4. To reduce power dissipation, we might add power gating at bias circuit of chopper stabilized pre-amplifier. Let the 8 channel pre-amplifiers will NOT consume static power at the same time.
- 5. Although the analog MUX and PGA are shared by eight channels pre-amplifier, overall power is still too large. Therefore, the power of the shared part should be more optimized.
- 6. Considering external disturbance, using pseudoresistor to generate high pass corner frequency might be NOT the best choice. However, current mode circuit may be another choice for addressing the issue.
- 5. Finally, a full understanding of neural recording system and a more careful consideration are essential to design and implement a front-end circuit for these applications. There is still large improvement available in this design.

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