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**ADVANCED MATERIALS AND STRUCTURES  
FOR NANOSCALE CMOS DEVICES**

by

Dae-Won Ha

Memorandum No. UCB/ERL M04/46

6 December 2004

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**ELECTRONICS RESEARCH LABORATORY**

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University of California, Berkeley  
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**Advanced Materials and Structures for Nanoscale CMOS Devices**

by

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**B.S. (Yonsei University, Korea) 1993**

**M.S. (Yonsei University, Korea) 1995**

**A dissertation submitted in partial satisfaction of the  
requirements for the degree of**

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**in**

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**in the**

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**of the**

**UNIVERSITY OF CALIFORNIA, BERKELEY**

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**Fall 2004**

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University of California, Berkeley

Fall 2004

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Abstract

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**Doctor of Philosophy in Engineering –**

**Electrical Engineering and Computer Sciences**

**University of California, Berkeley**

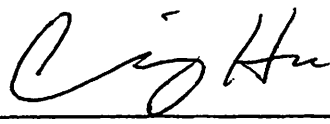
**Professor Chenming Calvin Hu, Chair**

Technological innovations have accomplished the continued scaling of CMOS devices well into nanometer regime. Short channel effects have been suppressed by the use of thinner gate oxide, shallower junction depth, and sophisticated channel doping profile without changing its basic structure. However, further CMOS scaling will be much more intricate due to fundamental materials and process limitations. The advanced thin-body transistor structures can effectively suppress the short channel effects, separating the need of heavy channel doping and the state-of-the-art thin gate oxide. This dissertation investigates the integration of advanced materials in front-end processes with advanced thin-body transistor structures.

Two essential processes in molybdenum (Mo) metal gate technology are developed: damage-free sputtering and high-selectivity dry etching. The physical origin of Mo gate work function engineering is simultaneous modification of the microstructure and chemistry at the gate dielectric interface, as discussed in chapter 2.

The impact of gate process technology on hafnium oxide ( $\text{HfO}_2$ ) gate dielectric is explored. The increase in equivalent oxide thickness and leakage current is resulted from the generation of oxygen vacancy, which causes silicon interfacial layer formation and gate electrode Fermi-level pinning in MOS devices, as discussed in chapter 3.

Tunable work function Mo gate is demonstrated for adjusting threshold voltages of ultra-thin body MOSFETs for the first time. Integration of metal gate and high- $k$  gate dielectric employing FinFETs is demonstrated for the first time; Mo-gated  $\text{HfO}_2$  CMOS FinFETs reduce gate leakage current over 3 orders-of-magnitude for inversion equivalent oxide thickness (1.72 nm) with comparable carrier mobility, as discussed in chapter 4.

 Nov. 16, 2004

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Professor Chenming Calvin Hu  
Dissertation Committee Chair



**To my parents,  
for their unforgettable love and sacrifices**

**To my wife,  
for her everlasting love, support and encouragements**

**To my son,  
for his brilliant and ambitious future**

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# Chapter 1

## Introduction

### 1.1 Future of CMOSFETs

Remarkable progress in semiconductor industry has led to the proliferation of microelectronic devices and information technology. The predominant technology is based on silicon CMOSFETs, the basic building blocks of integrated circuits. Over the past three decades, rapid and steady improvements in circuit performance and packing density have been achieved through the scaling of CMOS devices. The CMOS scaling has been accomplished with technological innovations, and leads the device dimensions well into the nanometer regime, as illustrated in **Figure 1.1** [1.1, 1.2].

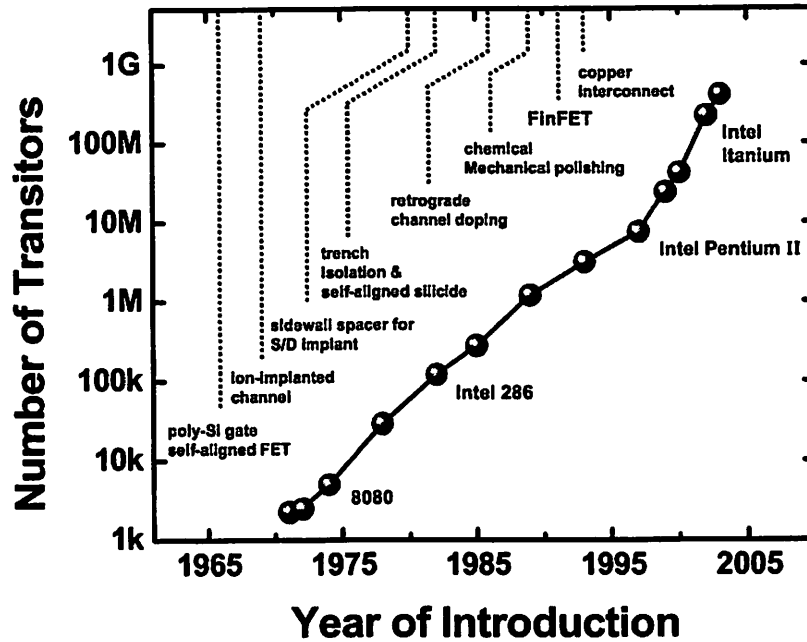


Figure 1.1 Remarkable progress in integration of transistors into a central processing unit (CPU). Technological innovations have accomplished the continued scaling of silicon-based CMOS devices, and shrunk them into the nanometer regime [1.1, 1.2].

However, new technological barriers should be overcome for future nanoscale CMOS scaling, because some fundamental materials and process limits are rapidly approached [1.3]; emphasis will be on the reduction of standby power consumption without aggravating the device performance.

To achieve low off-state leakage current of a bulk-Si MOSFET, the channel potential should be controlled by the gate electrode rather than the drain electrode; thus, the gate-to-channel capacitance should be larger than the drain-to-channel capacitance.



Historically, this has been achieved through the use of thinner gate oxide, shallower source/drain junction depth, and complex channel doping engineering [1.4, 1.5].

Thinner gate oxide, *silicon dioxide*, increases gate capacitance. However, for a silicon dioxide thickness of 2.0 nm or below, direct quantum mechanical tunneling through the dielectric leads to an exponential increase in leakage current with decreasing oxide thickness [1.6]. Therefore, thicker and higher permittivity dielectrics need to be investigated as an alternative gate dielectric to replace the traditional silicon dioxide.

Shallow junction depth decreases the capacitive coupling of the drain to the channel. However, the formation of ultra-shallow junction depth is limited due to the finite energy of ion implantation and transient enhanced dopant diffusion during the thermal budget for dopant activation [1.7]. Parasitic series resistance in the junction regions could degrade the transistor on-state current, demanding technological innovations such as raised source/drain structures or activation of dopants at concentrations higher than the solid solubility.

Higher channel doping concentration with a localized halo in a bulk-Si transistor shields the channel from the influence of the drain potential [1.8]. However, too heavy channel doping decreases carrier mobility, increases parasitic junction capacitance and

leakage current, and worsens the subthreshold slope. These factors combined can significantly deteriorate either performance or standby power consumption. Variations in device properties due to random dopant fluctuation effects may limit chip yield due to the intolerable clock skews or malfunction of the circuits [1.2].

Recently, advanced thin-body transistor structures such as ultra-thin body (UTB) single gate and double gate MOSFETs have been proposed to suppress the off-state leakage current without degrading the on-state current [1.9, 1.10]. The use of ultra-thin undoped channel can effectively suppress the off-state leakage current, overcoming the aforementioned challenges in nanoscale bulk-Si MOSFETs. Reduced vertical electric field improves the gate dielectric reliability and reduces gate tunneling current. Circuit performance can be further increased due to the higher channel carrier mobility and reduced parasitic junction capacitance. However, implementation of such device structures has difficulties in process integration, except for the FinFET which provides self-alignment of top and bottom gates, highest packing density and easy access to all four transistor electrodes [1.11]. Remaining challenges for FinFETs to maximize the performance and minimize the standby power consumption are illustrated in **Figure 1.2**, and addressed in the subsequent chapters.

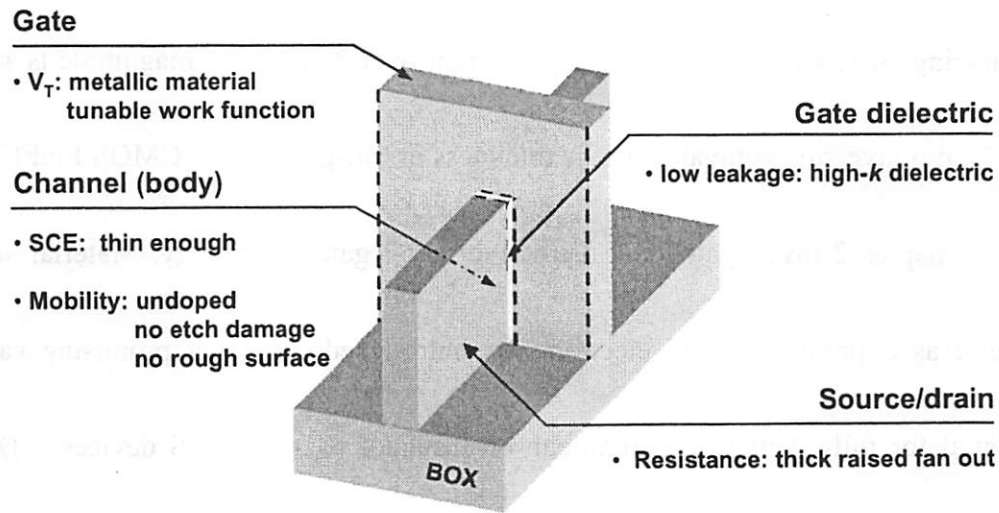


Figure 1.2 The FinFET provides relatively simple process integration, thus is one of the most promising double gate structures manufacturable below sub-25 nm gate length. However, technological innovations illustrated are required to maximize the performance benefits.

## 1.2 Objectives

In this dissertation, the key technological innovations illustrated in Figure 1.2 are addressed for the future nanoscale CMOS technologies. The process integration challenges and device performance of the advanced gate stack materials employing the advanced thin-body transistor structures are experimentally explored for the first time. A method of adjusting threshold voltages of ultra-thin body MOSFET and FinFET is

demonstrated for the first time with tunable molybdenum (Mo) gate work function engineering. A reduction of gate leakage current over 3 orders-of-magnitude is achieved for 1.72 nm inversion equivalent oxide thickness in Mo-gated HfO<sub>2</sub> CMOS FinFETs.

Chapter 2 investigates the alternative metal gate technology. Material selection criteria as a promising gate electrode are introduced; Mo is a promising candidate material for fully depleted (FD) silicon-on-insulator (SOI) CMOS devices. Damage-free sputtering with *plasma charge trap* and highly-selective dry etching processes are discussed and the analysis results of the work function shift for nitrogen implanted Mo gate are presented.

Chapter 3 explores the alternative high-*k* gate dielectric technology. The key basic material properties as a promising gate dielectric are summarized; HfO<sub>2</sub> is one of the most promising materials to replace SiO<sub>2</sub>. The impact of gate process technology is experimentally investigated. Successful integration of HfO<sub>2</sub> depends on minimizing the oxygen vacancy formation during CMOS fabrication through a reduction of post gate deposition thermal budget, and careful selection of the gate electrode material.

Chapter 4 investigates design methodology and process integration challenges for the advanced thin-body transistor structures. Adjustment of the desired threshold

voltages without degrading the performance benefits becomes a serious challenge. Mo gate work function engineering is experimentally demonstrated to be an attractive approach for adjusting the threshold voltages of advanced thin-body transistor structures. Integration of Mo gate and HfO<sub>2</sub> gate dielectric employing FinFET is experimentally demonstrated to reduce the gate leakage current over 3 orders-of-magnitude for 1.72 nm inversion equivalent oxide thickness.

Chapter 5 provides an overall summary of this dissertation. Key research contributions and suggestions for future research directions and possibilities are highlighted.

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# Chapter 2

## Metal Gate Technology

### 2.1 Introduction

Polycrystalline silicon (poly-Si) gate technology has been used in CMOS devices for several decades. One of the primary reasons is its excellent thermal stability on SiO<sub>2</sub> during the high temperature source/drain annealing steps. This allows the source/drain regions to be self-aligned to the gate, thus eliminating parasitic capacitance from overlay misalignment during the lithography process [2.1]. Another reason is the easy controllability of its work function at the SiO<sub>2</sub> interface. Its work function can be modified from a low value (~4.17 eV) to a high value (~5.1 eV) by the ion implantation of appropriate dopants; dual-doped poly-Si gates allow the threshold voltages of NMOS

and PMOS to be adequate for CMOS applications, thus dramatically reducing the circuit power consumption [2.2].

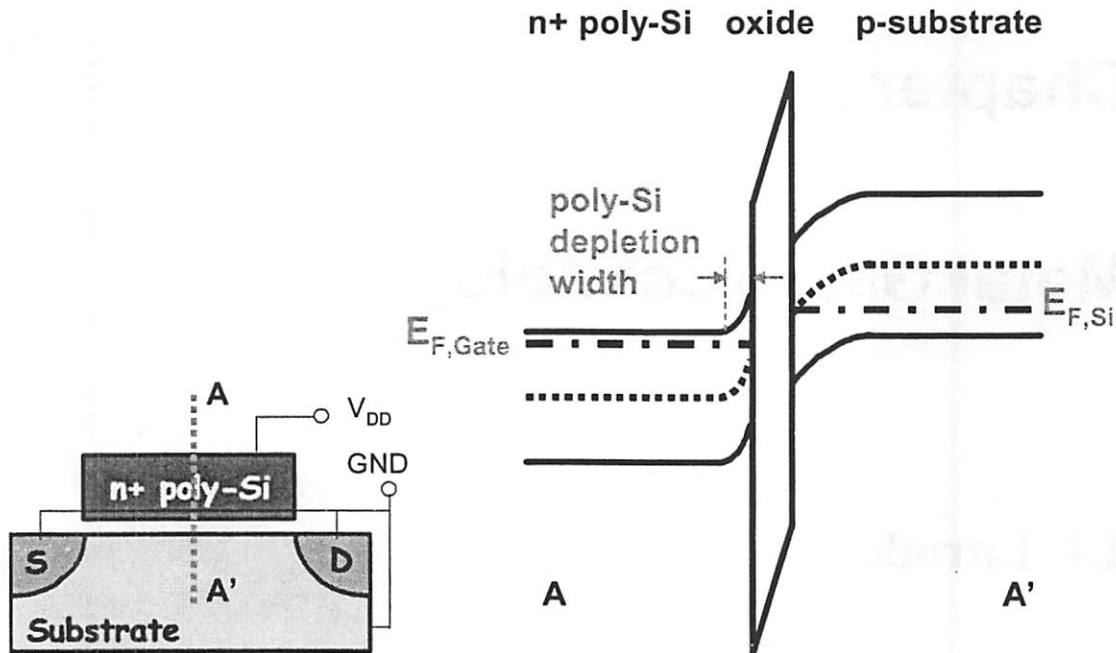


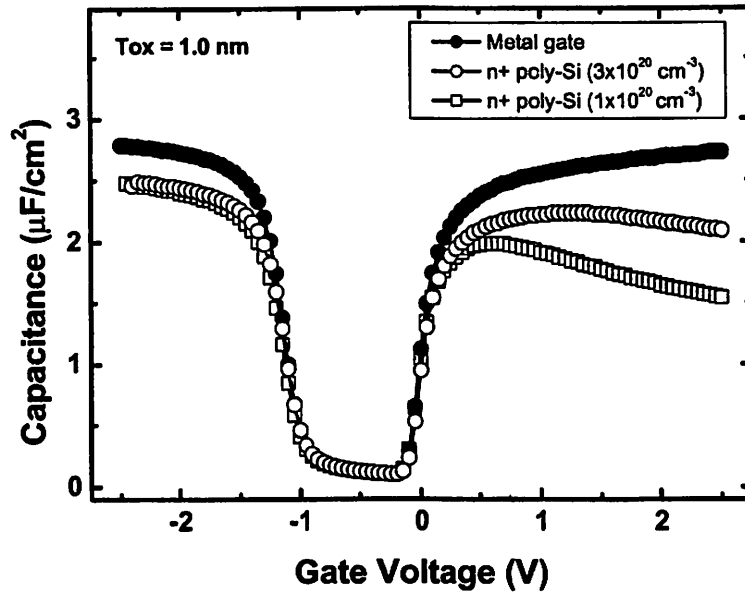
Figure 2.1 Energy band diagram of a n-channel MOS capacitor showing the depletion layer in the poly-Si gate, which typically adds several angstroms to the capacitance equivalent gate dielectric thickness (CET) when the channel is in strong inversion.

However, dual-doped poly-Si gates may not be promising any longer for nanoscale CMOS transistors due to poly-Si gate depletion effect (PDE), high sheet resistance, and boron dopant penetration through the gate oxide, etc. Among these concerns, the poly-Si gate depletion effect (PDE) is the most compelling reason; that is, a heavily doped poly-Si gate becomes slightly depleted at the gate oxide interface when the

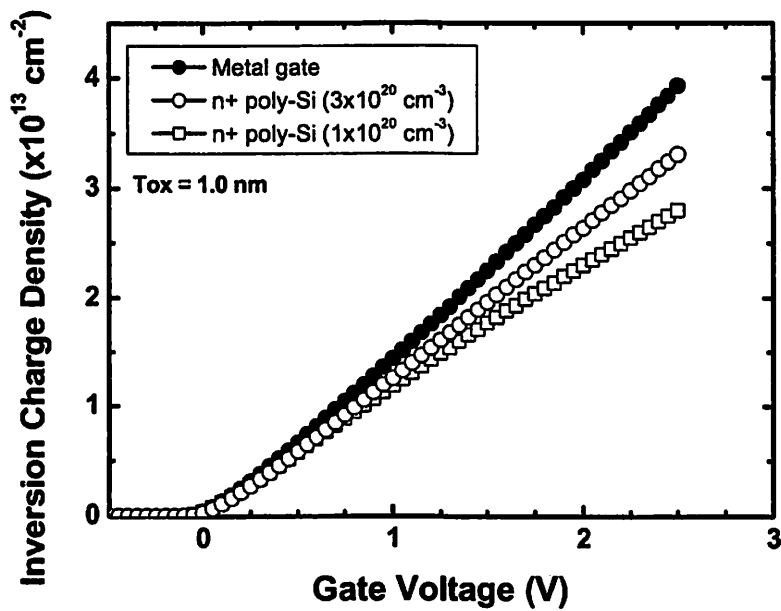


channel is in strong inversion [2.3]. **Figure 2.1** illustrates a schematic band diagram across a MOS transistor in strong inversion. When the MOS transistor is biased into the strong inversion, an electric field is developed across the gate oxide. Since the oxide field points in the direction of accelerating a negative charge toward the gate, the bands in the poly-Si gate bend slightly upward toward the oxide interface, which depletes the surface of electrons and forms a thin depletion region ( $W_{d,poly}$ ) in the poly-Si gate. This effectively increases the capacitance-equivalent thickness (CET) of the gate dielectric and decreases the inversion charge density in the channel, thus lowering the total gate capacitance and driving current of transistors. **Figure 2.2** shows the quantum-mechanical simulation results of (a) C-V characteristics, and (b) inversion charge densities of poly-Si and metal gate NMOS transistors [2.4]. Since the oxide field becomes stronger with a thinner oxide ( $< 1.5$  nm), the poly-Si gate depletion effect has more detrimental effects on nanoscale CMOS transistor performance.

Another concern with the poly-Si gate electrode is the high sheet resistance. Since the sheet resistance ( $R_S$ ) is a function of thickness ( $t$ ) and resistivity ( $\rho$ ) of the gate electrode material, a thicker poly-Si gate and/or a lower resistivity is helpful to decrease the sheet resistance. However, a thicker gate stack proves to a poor choice due to the



(a)



(b)

Figure 2.2 Quantum-mechanical (QM) simulation results of (a) C-V characteristics, and (b) inversion charge density of poly-Si (open symbols) and metal gate (solid circle) n-channel MOS transistors.

increased parasitic gate-to-source/drain capacitance, thus compromising the circuit performance [2.5]. Also, in the poly-Si gate, the resistivity is determined by the active dopant concentration, which is limited by the solid solubility and thermal annealing of dopants in silicon. While the solid solubility of dopants is a material property, the use of high temperature annealing is limited due to the need for an abrupt retrograded channel doping and shallow source/drain junction doping profiles in nanoscale CMOS transistors.

Other concerns with dual-doped poly-Si gates include dopant (boron) penetration through thin gate oxide due to the large variation in threshold voltage and degradation in gate oxide reliability [2.6], thermodynamic incompatibility with future high-k gate dielectrics such as  $ZrO_2$  [2.7],  $HfO_2$  [2.8], and so on.

Metal gate electrodes can eliminate all of the aforementioned concerns, thus are indispensable for sub-65 nm CMOS technology node [2.9]. However, there are challenges in process integration that should be overcome to apply the metal gate technology to nanoscale CMOS devices.

## 2.2 Material Selection for Gate Electrode

In selecting alternative CMOS gate electrode materials, several factors must be considered [2.10]; these include the thermal stability on gate dielectric during high-temperature annealing step(s), the compatibility of the material growth and patterning with the conventional CMOS processes, the work function on a gate dielectric, the impact of gate processes on device performance and reliability, and the extendibility to future high- $k$  gate dielectrics.

The highest-temperature annealing step(s) used in today's CMOS fabrication occurs during the source/drain dopant activation. Typically, the dopant activation is accomplished through rapid thermal annealing (RTA) at temperatures around or slightly above 1000°C for a few seconds in an inert ambient. Therefore, prospective gate materials should be thermodynamically stable on the gate dielectric, and have a higher melting point than 1000°C. It is worthwhile to note that choosing a material with a thermal expansion coefficient close to that of silicon is very desirable because the thermally induced stress may result in cracking or peeling of the gate stack materials.

The processes for material deposition and patterning should be compatible with the conventional CMOS processes. The deposition process plays an important role on

the interface quality between gate electrode and dielectric. Most thin metal films are deposited using either chemical vapor deposition (CVD) or physical vapor deposition (PVD) method. While CVD method leads to minimal damage and a smooth interface with the underneath gate dielectric, the precursors for only a limited number of metal films like tungsten (W), aluminum (Al), titanium nitride (TiN) have been well identified. On the other hand, PVD method can be used for most of the metal films with minimal incorporation of impurities, although new techniques need to be developed to prevent physical damage to the gate dielectric from the energetic particles (metal ions and/or electrons). Also, the dry etching process with high selectivity and vertical sidewall profile is indispensable for nanoscale CMOS device fabrication. While wet chemical etching results in high selectivity to the gate dielectric, the inherent isotropic etching characteristic prevents it from patterning the gate electrode in nanoscale CMOS device fabrication. Therefore, the dry etching process condition should be optimized (or developed) for the gate material.

The effect of the work function of gate electrode on the device operation is through its control of the threshold voltage ( $V_T$ ), which determines how much a gate voltage should be applied to invert the channel. For conventional bulk transistors, the

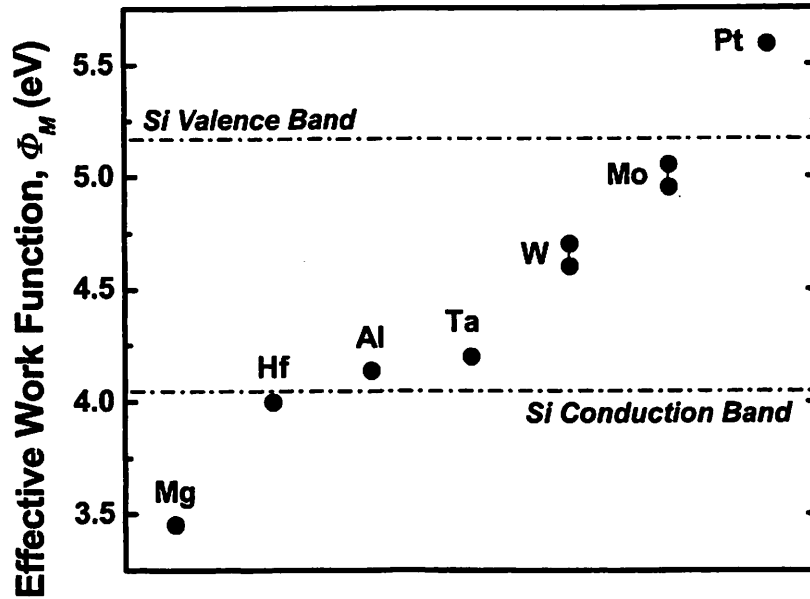


Figure 2.3 Experimental work function data for metals on silicon oxide ( $\text{SiO}_2$ ) [2.12].

silicon channel has a heavily ( $10^{16} - 10^{18} \text{ cm}^{-3}$ ) retrograded-doping profile to suppress the short channel effects. Thus, the work function difference between gate electrode and silicon channel affects the threshold voltage. A comprehensive quantum-mechanical (QM) simulation study shows that the device performance and short channel effects can be optimized simultaneously with metal gate electrode whose work function is near the conduction band ( $E_C$ ) for NMOS device and the valence band ( $E_V$ ) for PMOS device [2.11]. **Figure 2.3** shows the experimental work function data for metals on silicon dioxide ( $\text{SiO}_2$ ) [2.12]. On the other hand, for advanced thin-body transistors, the silicon

channel has an undoped or lightly ( $< 10^{16} \text{ cm}^{-3}$ ) uniform doping profile to minimize the dopant fluctuation effect and maximize the carrier mobility. Thus, threshold voltage adjustment should be achieved by the work function of gate electrode due to the negligible depletion charge [2.12]. It is worth to note that a lightly doped thin-body can effectively suppress the short channel effects through the proper design of the device parameters such as the ratio of gate length to body thickness,  $\zeta = L_G/T_{Si}$ . The optimized gate electrode work function is 0.35 eV below (above) the conduction (valence) band for NMOS (PMOS) devices [2.10]. **Figure 2.4** shows the theoretical data for threshold voltage of the thin-body MOSFET as a function of the gate work function [2.13].

Several material systems and process integration approaches which satisfy the aforementioned criteria have been investigated to date for CMOS applications: dual metal (Ti/Mo) [2.14], metal interdiffusion (Ti/Ni) [2.15], metal alloying (Ru/Ta) [2.16], fully silicided ( $\text{NiSi}_x$ ) doped polysilicon [2.17], and tunable-work-function metal (Mo) gate [2.18, 2.19, 2.20].

The tunable-work-function Mo gate technology offers the simplest CMOS process integration and better compatibility with high- $k$  gate dielectric materials [2.19].

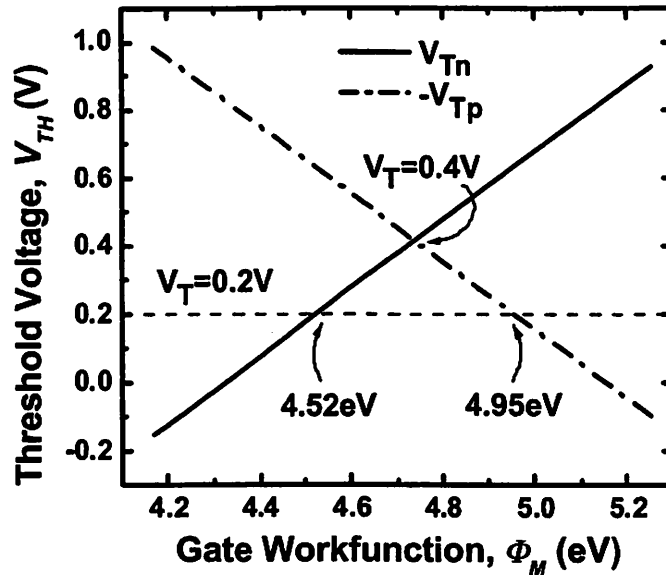


Figure 2.3 Theoretical data for threshold voltage of the thin-body MOSFET as a function of the gate work function [2.13].

Gate work function engineering via selective nitrogen implantation into the Mo gate electrode will provide multiple values of threshold voltage ( $V_T$ ) for NMOS and PMOS transistors on a single substrate. A higher melting point (2617°C) [2.21] and thermodynamic stability on SiO<sub>2</sub> and high- $k$  dielectrics such as ZrO<sub>2</sub>, HfO<sub>2</sub>, etc [2.22, 2.23] allow for the self-aligned (gate-first) CMOS transistor fabrication. It should be noted that the adjustable-work-function fully silicided (NiSi<sub>x</sub>) method may not be applied on the future high- $k$  dielectrics due to the thermodynamic instability [2.24]. The following sections will discuss the process details of the tunable-work-function Mo gate technology.



## 2.3 Damage-free Molybdenum Sputtering

Physical vapor deposition (PVD), also known as sputter deposition, and chemical vapor deposition (CVD) are the most widely used techniques for the fabrication of thin-film structures on semiconductor wafers. The physical vapor deposition as compared with chemical vapor deposition provides an important advantage of minimal incorporation of impurities. However, the physical vapor deposition is a relatively violent, atomic-scale process in which an energetic particle strikes a solid, resulting in the emission of one or more atoms from the solid sputtering target; thus, sputtering process may introduce damage to the gate dielectric.

### 2.3.1 *Sputtering System for Molybdenum*

A DC-magnetron sputtering system (*Novellus m2i with Quantum source*) was used to deposit the Mo gate films. **Figure 2.5** illustrates the schematics of the sputtering system; it has two load-lock stations, a wafer degassing station, sputtering chambers, and a cooling station. The base pressure of the sputtering chamber is maintained below  $5 \times 10^{-8}$  Torr, and argon (Ar) gas was used for Mo target sputtering.

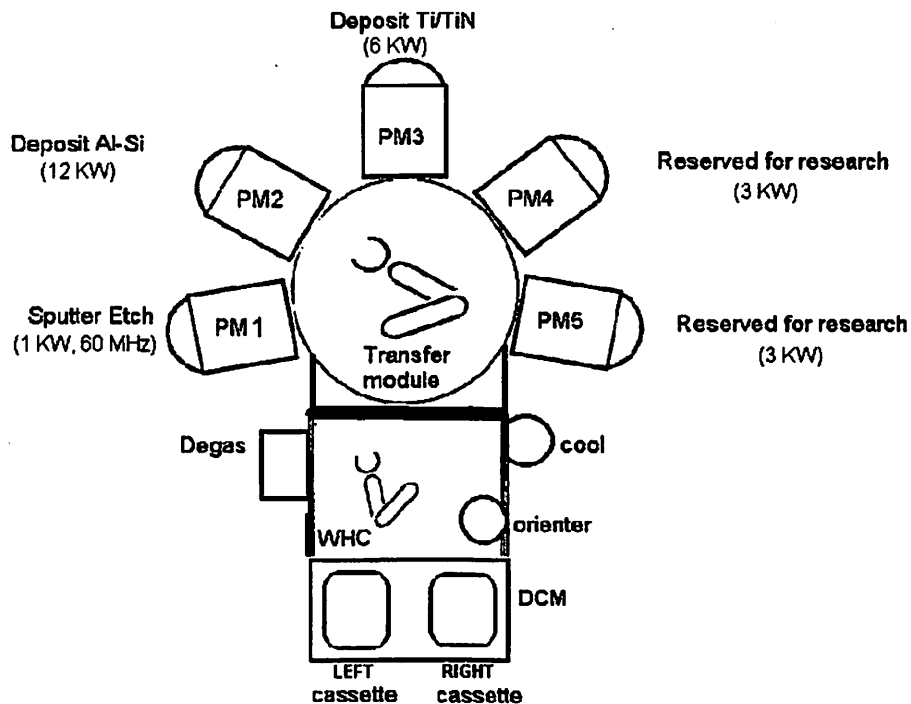


Figure 2.5 Schematic of the Mo sputtering system; loaded wafers in load-lock station are transported to process module 5 for Mo sputtering.

The critical requirements for a gate deposition process are: zero damage to the gate dielectric and good step coverage for non-planar transistor structures. Sputtering damage can result in degraded gate-oxide integrity (GOI) and transistor drive current due to degraded field-effect carrier mobility. The sputtering damage can be minimized by eliminating the high-energy particle bombardment by interposing an electrically grounded *Plasma Charge Trap* (PCT) between the Mo target and substrate, separating the plasma excitation and Mo film deposition [2.25]. Figure 2.6 illustrates a schematic

of the sputtering chamber with the PCT, which is made of 1.66 mm thick stainless steel and has arrays of closely packed 3.32 mm diameter holes (40.3% aperture area). Since the PCT is electrically grounded, it collects high energy ions as well as a fraction of the neutral Mo atoms which are traveling toward the cathode/sample plane. These particles are deposited onto the PCT, and hence are not deposited onto the wafer surface. Therefore, the sputtering damage can be minimized. It is worth noting that the Mo deposition rate with the PCT is reduced by about 70% and is proportional to the PCT aperture area.

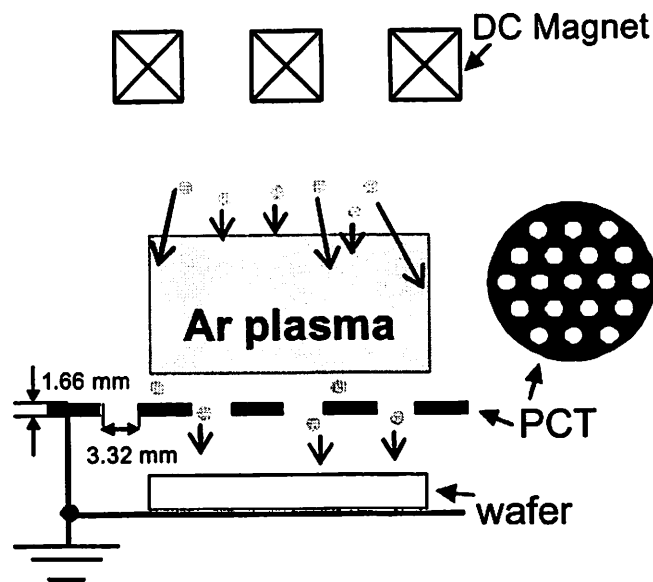


Figure 2.6 Schematic of the sputtering chamber with the plasma charge trap (PCT), which is inserted between the sputtering target (Mo) and wafer.

### **2.3.2 Experimental Results and Discussion**

To investigate the efficacy of the plasma charge trap (PCT) for minimizing the sputtering damage, conventional bulk silicon p-channel MOSFETs were fabricated with Mo gates sputtered either with or without the PCT. Mo-gate films with 50 – 150 nm thicknesses were deposited at 200°C with 10 mTorr process pressure and 300 W sputtering power.

Damage to the gate dielectric can be assessed by measuring the gate leakage current density and gate-dielectric lifetime. Gate leakage current densities for Mo-gated PMOS capacitors are compared against those of p+ poly-Si gated PMOS capacitors (control samples) as shown in **Figure 2.7**. The p+ poly-Si gated devices show a large variation in gate leakage current due to boron dopant penetration through the gate oxide. The Mo-gated devices have lower gate leakage with much tighter distributions; the PCT provides ~30% lower gate leakage due to the lower (more negative) threshold voltage of the Mo-gated devices, even though the electrical stressing condition is more severe compared to the control samples.

**Figure 2.8** shows the Charge-to-Breakdown ( $Q_{BD}$ ) and measured Time Dependent Dielectric Breakdown (TDDB) of gate oxide. The measured transistor gate

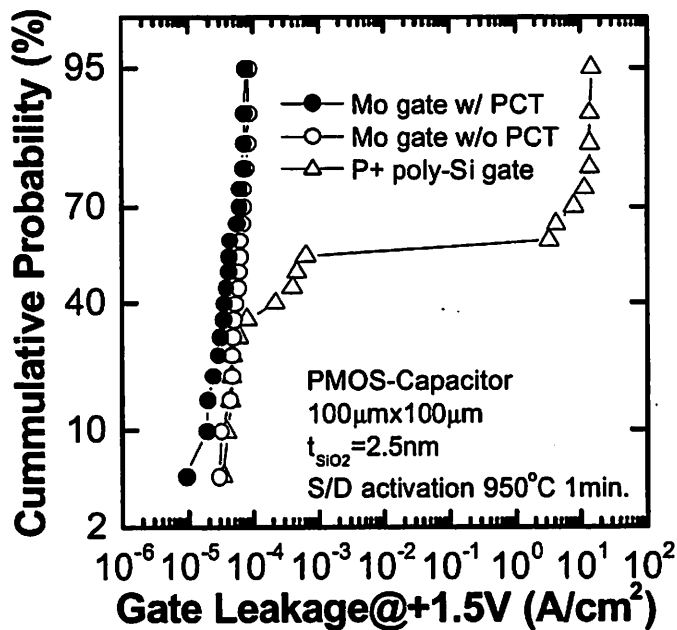


Figure 2.7 Gate leakage current densities of Mo-gated PMOS capacitors are compared against those of p+ poly-Si gated PMOS capacitors.

area is  $150\mu\text{m}^2$ . For the  $Q_{BD}$  measurements, the devices were biased at a constant gate voltage  $+4.4\text{V}$ . Since the devices have the same threshold voltage, the effective stressing condition is the same for both groups of samples. It is worth to note that the breakdown of a thinner gate oxide occurs by bulk trap generation, not by interface trap generation, and is therefore driven by gate voltage [2.26]. With PCT Mo sputtering, the gate oxide integrity is significantly improved; indicating that low sputtering damage to gate dielectric is achieved.

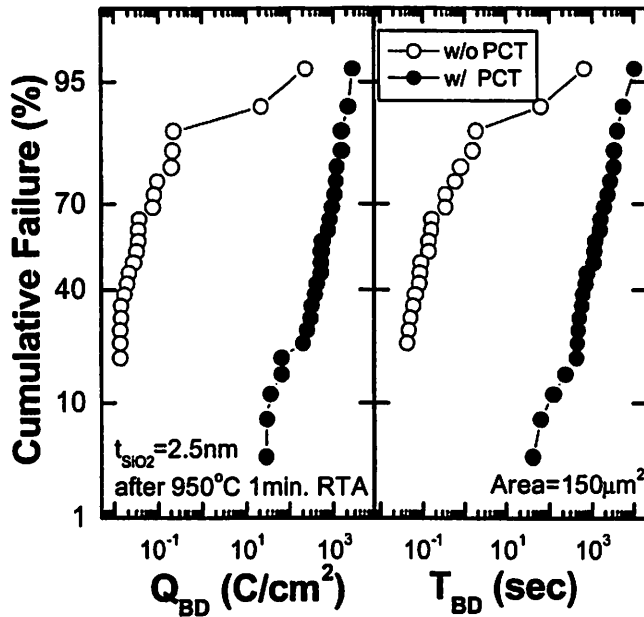
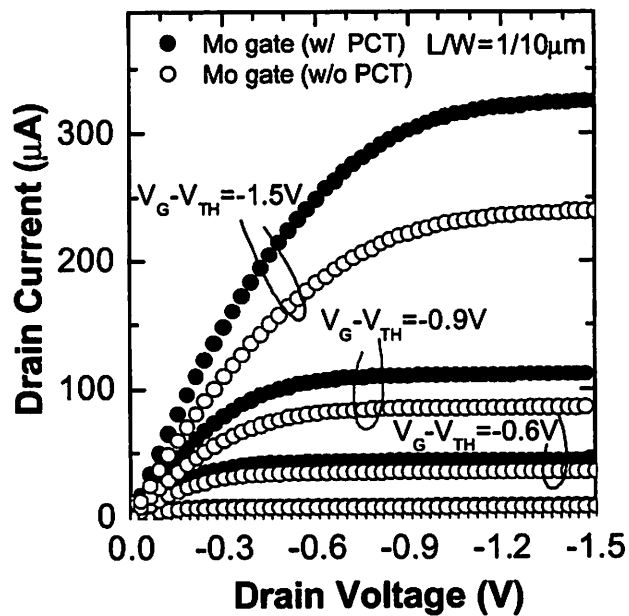
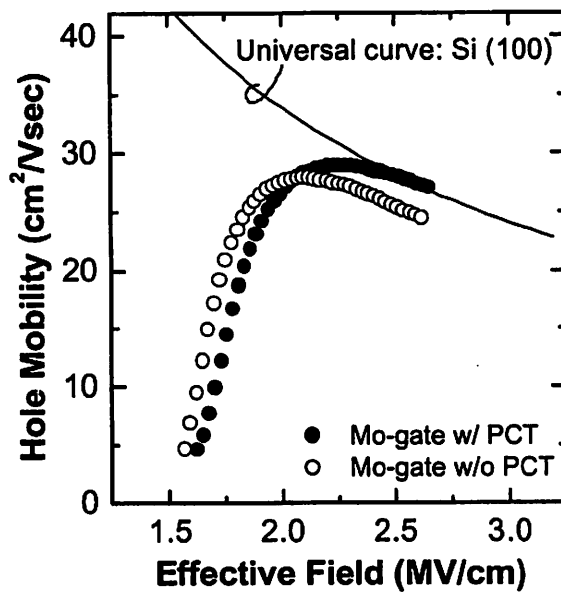


Figure 2.8 Measured Charge-to-Breakdown ( $Q_{BD}$ ) and Time Dependent Dielectric Breakdown (TDDB) of Mo-gated p-channel MOSFETs. With PCT Mo sputtering, the gate oxide integrity is significantly improved, indicating that low sputtering damage to gate dielectric is achieved.

Figure 2.9 (a) shows the measured  $I_D - V_D$  characteristics of Mo-gated bulk-Si p-channel MOSFETs. The drive current achieved with a PCT sputtered Mo gate is higher due to improved hole mobility as shown in Figure 2.9 (b), which matches the universal mobility curve. The improved hole mobility is attributed to reduced interface traps ( $Q_{it}$ ) and/or fixed charge ( $Q_f$ ) due to the damageless Mo gate sputtering process [2.27].



(a)



(b)

Figure 2.9 (a) Measured  $I_D - V_D$  characteristics, and (b) measured effective hole mobility of Mo-gated p-channel bulk-Si MOSFETs.

## 2.4 Highly Selective Molybdenum Dry Etching

It is known that fluorine (F) and chlorine (Cl) radicals can etch Mo films at an appreciable rate [2.28]. Therefore, Mo-gate films can be etched using either F-based or Cl-based etching gases such as  $\text{CF}_4$ ,  $\text{CF}_3\text{Cl}$ ,  $\text{Cl}_2$ , etc. However, fluorinated carbon compounds like  $\text{CF}_4$  and/or  $\text{CF}_3\text{Cl}$ , which may attack the underneath gate dielectric ( $\text{SiO}_2$ ) [2.29], are precluded due to the selectivity requirement. Thus, focusing on a  $\text{Cl}_2/\text{O}_2$  gas mixture, the dependences of etch rate and selectivity to  $\text{SiO}_2$  on  $\text{Cl}_2/\text{O}_2$  gas flow rates, source and bias RF powers were investigated.

### 2.4.1 Dry Etching System for Molybdenum

A Transformer Coupled Plasma etcher (*LAM Research TCP 9400*) was used to etch the Mo gate films. **Figure 2.10** illustrates the schematic of the dry etching system; it has load (unload)-lock stations and a process chamber. The process chamber has two RF power (source, bias) supplies running at the same time during the dry etching process, which results in a high-density plasma with adjustable substrate bias.

The critical requirements for a gate etching process are high selectivity to the gate dielectric ( $\text{SiO}_2$ ) and vertically etched profile. **Figure 2.11** shows the minimum



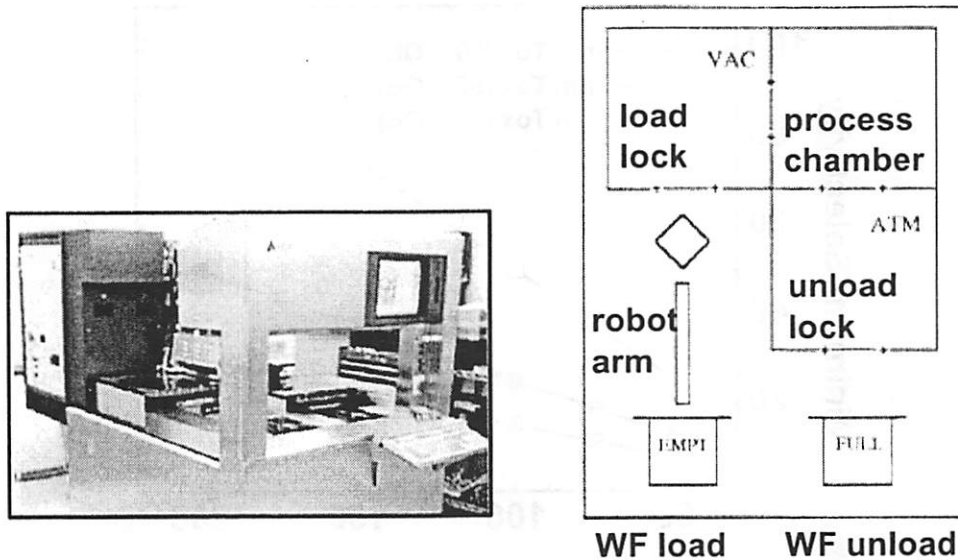


Figure 2.10 Schematics of the Mo dry etching system (LAM Research TCP9400).

etch selectivity required, as a function of gate stack thickness. For example, the etching selectivity must be higher than 25:1, assuming that the thicknesses of the Mo gate and SiO<sub>2</sub> gate oxide are 100 nm and 2 nm, respectively, with a 50% over-etch.

#### 2.4.2 Experiment Results and Discussion

Conventional bulk-Si sample wafers with 100 nm thick thermally grown SiO<sub>2</sub> and 300 nm thick Mo film (or 1 μm thick photo resist) on the SiO<sub>2</sub> were prepared to investigate the etching characteristics of Cl<sub>2</sub>/O<sub>2</sub> gas mixture. To calculate the etch rate of the thin films, physical step-height and optical thickness measurements were adopted

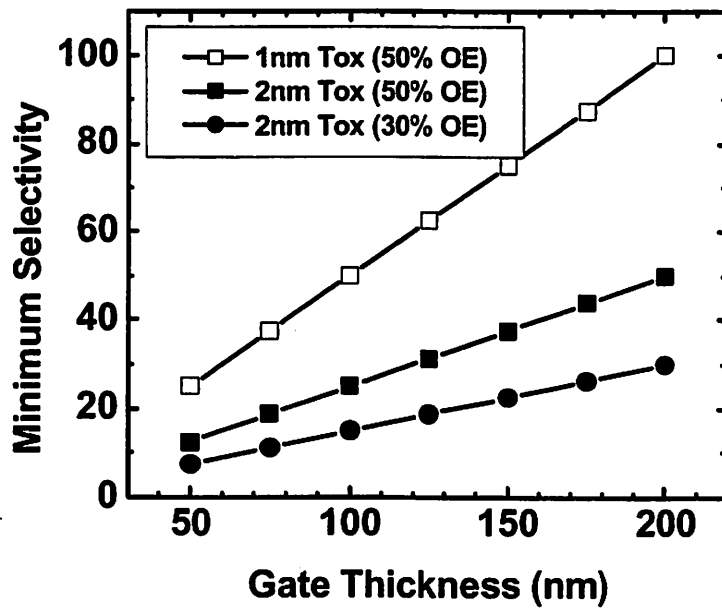


Figure 2.11 Minimum etch selectivity required as a function of gate stack thickness.

before and after the dry etching process of each sample wafer. The Mo film was sputtered in the Novellus system, SiO<sub>2</sub> was thermally grown in a wet oxidation furnace at 900°C, and *i*-line (OCG OiR-10i) photo resist was hard baked in 10 minutes at 120°C.

Figure 2.12 shows the dependence of the etch rates of Mo, SiO<sub>2</sub> and photo resist (PR) on the oxygen partial pressure in a Cl<sub>2</sub>/O<sub>2</sub> gas mixture. The total gas flow rate was held constant at 130 sccm, and the process pressure was 13 mTorr with 150W/100W source/bias RF powers. With sufficient chlorine and oxygen radicals, the addition of oxygen gas increases the Mo-gate etching rate and decreases the SiO<sub>2</sub> etch rate, and hence improves the selectivity. This implies that the surface of the Mo film can be

easily oxidized and reacted with chlorinated gas to sublime at  $\sim 100^\circ\text{C}$  in the form of molybdenum oxychlorine ( $\text{MoO}_x\text{Cl}_y$ ) [2.30]. It should be noted that more than 75% oxygen results in significant etching of the photo resist (PR), which may prevent it from being used as a useful masking material.

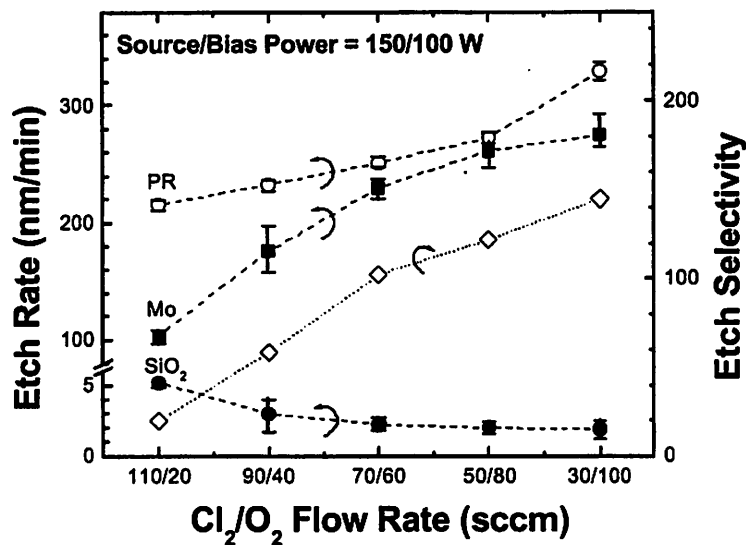


Figure 2.12 Etch-rate dependences of Mo, SiO<sub>2</sub>, and photoresist (PR) on the oxygen partial pressure in the Cl<sub>2</sub>/O<sub>2</sub> gas mixture. The total gas flow rate was held constant at 130 sccm, and the process pressure was 13 mTorr with 150 W source RF power and 100 W bias RF power.

Figure 2.13 shows the dependence of Mo and SiO<sub>2</sub> etch-rate on source power with a constant gas flow rate (Cl<sub>2</sub>/O<sub>2</sub> = 70/60 sccm), process pressure (13 mTorr), and bias power (100 W). The etch rates of Mo and SiO<sub>2</sub> increase linearly with source power

because more chlorine and oxygen radicals are available, but the selectivity decreases due to the faster increase in SiO<sub>2</sub> etch rate. For source powers below 150 W, the etch rate of SiO<sub>2</sub> is constant due to the DC self-bias voltage, so that the etch selectivity is degraded.

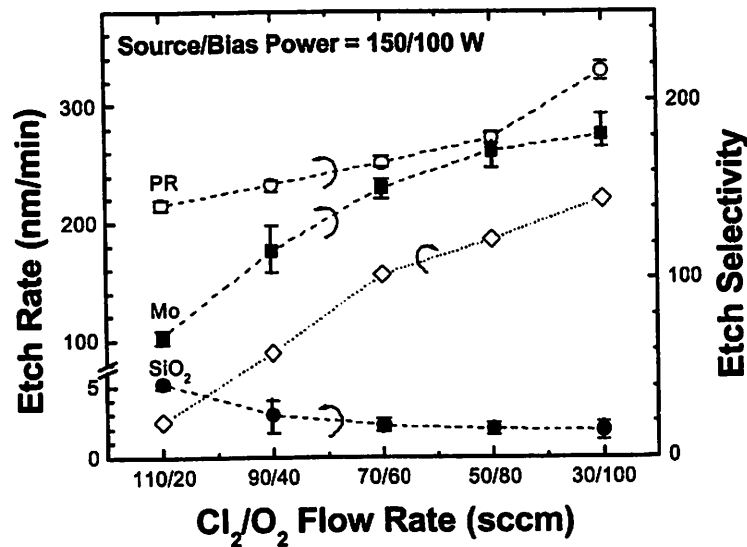


Figure 2.13 Etch-rate dependences of Mo and SiO<sub>2</sub> on source power with constant gas flow rate (Cl<sub>2</sub>/O<sub>2</sub> = 70/60 sccm), process pressure (13 mTorr), and bias power (100 W).

Figure 2.14 shows the dependence of Mo and SiO<sub>2</sub> etch-rate on bias power with constant gas flow rate (Cl<sub>2</sub>/O<sub>2</sub> = 70/60 sccm), process pressure (13 mTorr), and source power (150 W). The Mo etch rate saturates above 100 W bias power, whereas SiO<sub>2</sub> etching rate increases monotonically. Therefore, the etch selectivity decreases with increasing bias power. This indicates that energetic ion bombardment does not always

increase the Mo etch rate, and that chemical etching plays an important role. It is worth to note that the bond dissociation energy of SiO<sub>2</sub> is larger than that of Mo [2.31].

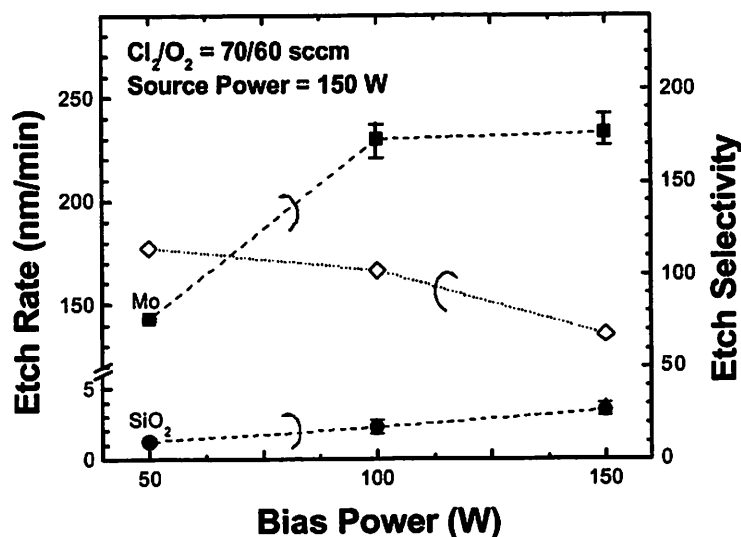


Figure 2.14 Etch-rate dependences of Mo and SiO<sub>2</sub> on bias power with constant gas flow rate (Cl<sub>2</sub>/O<sub>2</sub> = 70/60 sccm), process pressure (13 mTorr), and source power (150 W).

Table 2.1 summarizes the optimized Mo gate etch process condition which yields 230 nm/min etch rate and ~100:1 (Mo:SiO<sub>2</sub>) etch selectivity: Cl<sub>2</sub>/O<sub>2</sub> = 70/60 sccm, process pressure = 13 mTorr, source/bias RF power = 150/100 W.

Figure 2.15 shows the plane- and tilted-view scanning electron microscopy (SEM) picture of the etched Mo gate profiles using the optimized etch recipe. Neither micro-trenching nor Mo gate residue was observed due to the high etching selectivity;

thus, this process condition can be directly applicable to the thin-body transistor structures including FinFETs.

Table 2.1 Optimized Mo gate dry-etching process condition.

	Step 1	Step 2	Step 3	Step 4	Step 5
	Stabilize	Stabilize	Time	Stabilize	EPD
Gap (cm)	5.8	5.8	5.8	5.8	5.8
Pressure (mTorr)	-	30	30	13	13
Source Power (W)	0	0	300	0	150
Bias Power (W)	0	0	120	0	100
CF <sub>4</sub> (sccm)	0	100	100	0	0
Cl <sub>2</sub> (sccm)	0	0	0	70	70
O <sub>2</sub> (sccm)	0	0	0	60	60
Time (sec)	30	30	7	30	30
End-Point-Detect	-	-	-	-	Ch A

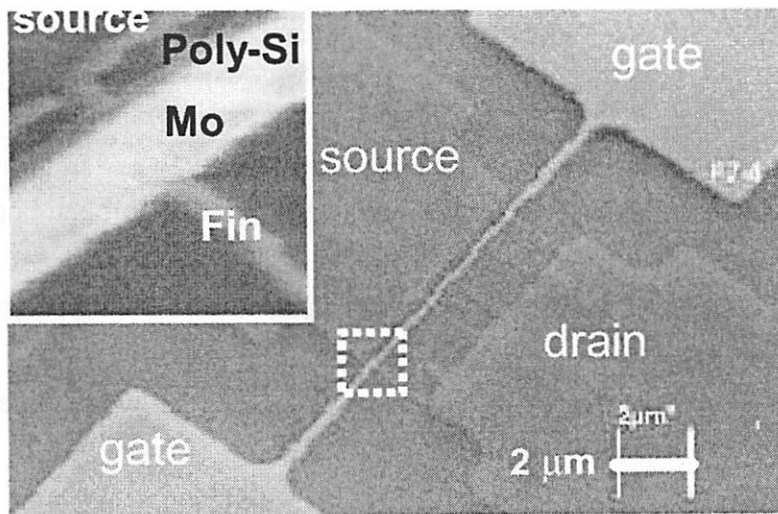
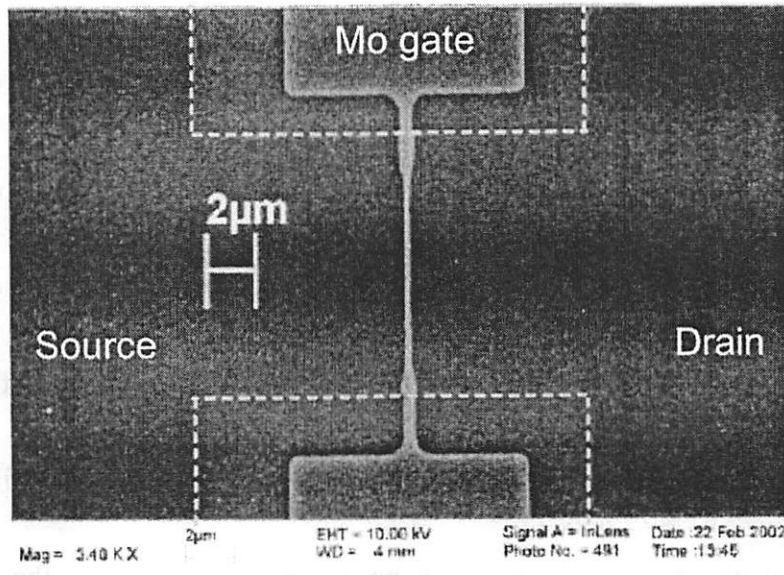


Figure 2.15 Plane- and tilted view scanning electron microscopy (SEM) picture of the etched Mo gate profiles using the optimized process condition.

## 2.5 Tunable-Work-Function Molybdenum Process

A *work function*, defined as a minimum energy necessary to extract an electron from a metal surface, is a fundamental property of metals that depends on both the electronic and ionic structure at that metal surface. Thus, the work function can be engineered by modifying the microstructure and/or chemistry of a metal film.

Microstructural modification can be achieved by changing the film texture or preferred orientation, crystalline phase, and crystal lattice, etc. For example, the work function of a metal film is, in general, dependent on the surface orientation (also known as the *work function anisotropy*); closely packed crystallographic surfaces show high work functions due to the smooth surface and relatively fewer broken bonds, while open crystallographic surfaces show low work functions due to a great number of broken bonds [2.32]. Experimentally it has been observed the work function in metals decreases (110) (100) (111) trend [2.33]. In addition, this was experimentally demonstrated for molybdenum (Mo) and tungsten (W) with modification of the film microstructures (surface orientation, crystalline phase) by implanting inert Ar<sup>+</sup> ions and/or controlling the deposition parameters [2.17, 2.34].



Chemical modification can be achieved by nitridizing, oxidizing, and alloying the metal film. It should be noted that the metal film after chemical modification should maintain high metallic conductivity to be used as a gate electrode. Modifying the film chemistry through nitridizing Mo film [2.35, 2.36], and alloying Ru-Ta [2.15] was experimentally demonstrated.

Both microstructural and chemical modifications can be simultaneously achieved by introducing foreign elements such as oxygen and nitrogen into a metal. Ion implantation is the most widely used technique to introduce dopant ions into films due to its precise control of implant dose ( $\#/cm^2$ ), energy (eV) and tilt angle. For example, the implantation of nitrogen ions ( $N^+$ ) into Mo film with subsequent annealing can modify the microstructure and chemical properties with precise control through inducing a nitridation reaction of Mo films [2.17].

### ***2.5.1 Experimental Results and Discussion***

Mo-gated capacitors were fabricated on lightly doped ( $\sim 10^{15} \text{ cm}^{-3}$ ) p-type Si substrates with thermally grown  $\text{SiO}_2$  as a gate dielectric. In order to account for the influence of oxide fixed charge ( $Q_F$ ), multiple  $\text{SiO}_2$  thicknesses were obtained on a single

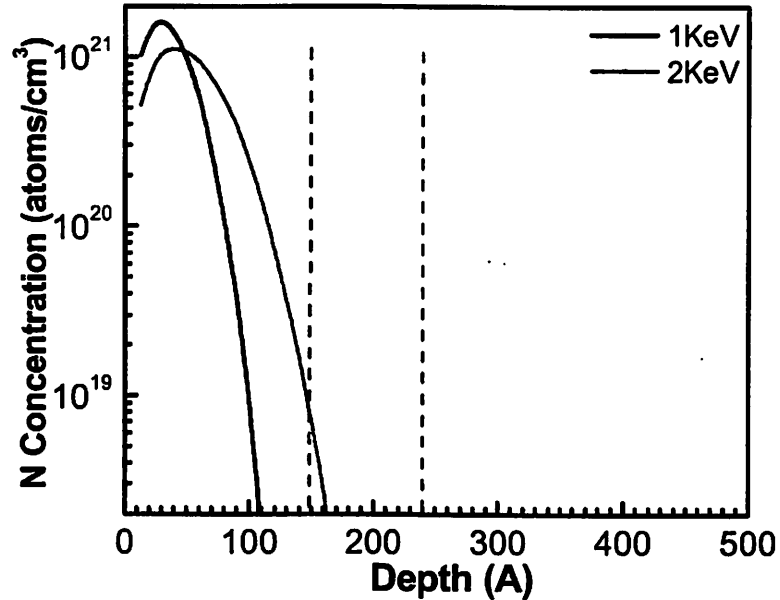


Figure 2.16 Simulated nitrogen concentration-depth profiles in the Mo/SiO<sub>2</sub>/Si stack (Dose =  $5 \times 10^{15}/\text{cm}^2$ ).

substrate by selective etchback using dilute HF solution. 15 nm thick Mo films were sputtered at 200°C in Novellus sputtering system, as described in Chapter 2.3, with a base pressure of  $1 \times 10^{-8}$  Torr and a process pressure of 10 mTorr. For some wafers, the Mo gate was implanted with nitrogen ions. The implant energies were 1 keV and 2 keV with implant dose of  $5 \times 10^{15}$  and  $1 \times 10^{16} \text{ cm}^{-2}$ . Ultra-shallow nitrogen ion implantation to very thin Mo gate films was adopted to minimize the ion penetration into the gate dielectric due to the shallower projected range ( $R_p$ ) and smaller straggle ( $\Delta R_p$ ) while allowing for high atomic nitrogen content inside the Mo films. **Figure 2.16** shows

SRIM simulation results of the nitrogen concentration-depth profiles in the Mo/SiO<sub>2</sub>/Si stacks with nitrogen implant dose of  $5 \times 10^{15} \text{ cm}^{-2}$ , which is estimated to introduce about 5 % nitrogen inside the 15 nm thick Mo gate [2.37]. After gate patterning, wafers were annealed from 500°C to 900°C in N<sub>2</sub> ambient, followed by forming gas anneal (400°C, 15 min).

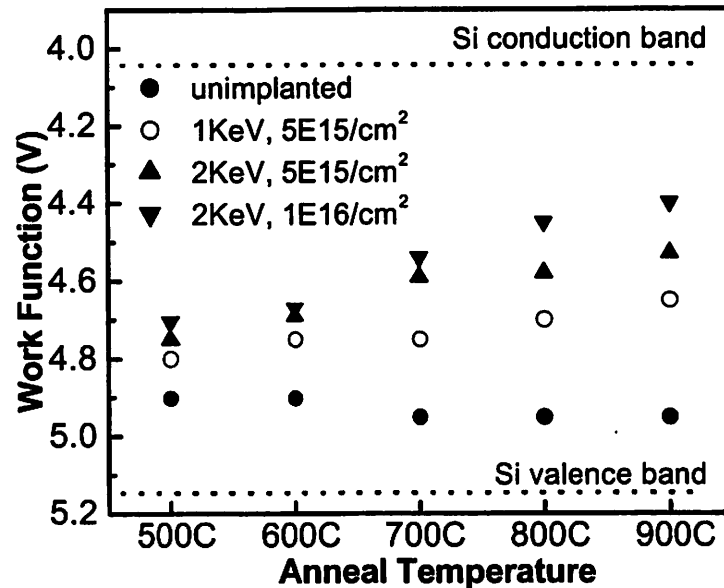


Figure 2.17 The extracted Mo work function with subsequent thermal annealing temperature. All anneals were 15 min long except for 900°C anneal (15s).

Figure 2.17 shows the extracted work function of Mo gate as a function of subsequent annealing temperature. The flatband voltages ( $V_{FB}$ ) of the Mo-gated

capacitors was determined from capacitance–voltage ( $C-V$ ) measurements, and used to extract the work function ( $\Phi_M$ ) [2.38]. The work function of un-implanted Mo gates increased slightly upon annealing, but is fairly stable at 4.95 eV. This increase can be ascribed to an improvement in the crystalline quality of the Mo film (increase in the columnar grain size). It is worth to note that this work function value is in good agreement with the published values for the (110) Mo work function. X-Ray Diffraction (XRD) analysis, as shown in Chapter 2.5.1, indicates that the deposited Mo film has a (110) preferred orientation.

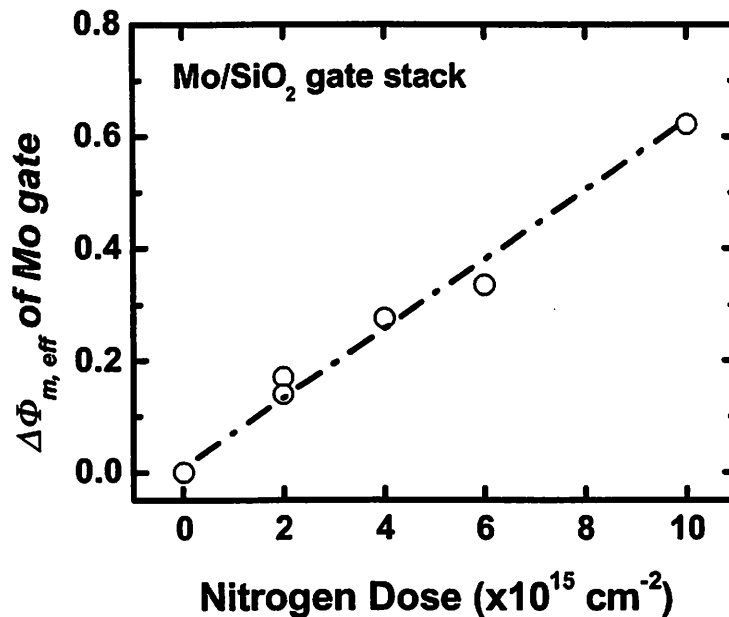


Figure 2.18 The effective work function shift of Mo gate as a function of nitrogen implant dose.

**Figure 2.18** shows the effective work function shift ( $\Delta\Phi_{m,eff}$ ) of Mo gate as a function of nitrogen implant dose. All samples were annealed at 900°C in N<sub>2</sub> ambient. It can be seen that the Mo gate work function decreases with a decrease in nitrogen implant doses. The amount of reduction in gate work function is proportional to the nitrogen implant dose.

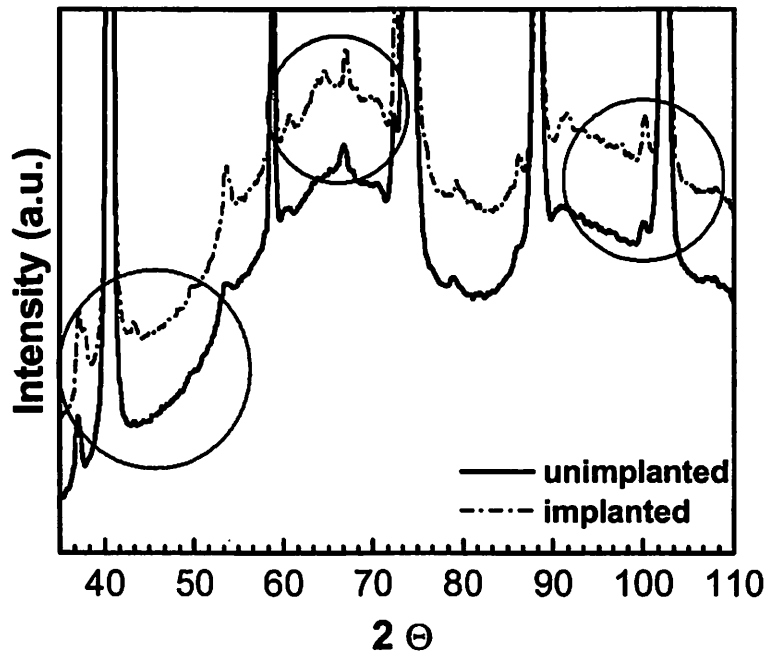


Figure 2.19 XRD analysis of nitrogen implanted and un-implanted Mo films.

In order to investigate the origin of the work function shift, x-ray diffraction (XRD) analysis of nitrogen implanted (dose =  $1 \times 10^{16} \text{ cm}^{-2}$ ) and un-implanted wafers were performed (**Figure 2.19**). Several diffraction peaks corresponding to Mo<sub>2</sub>N can be

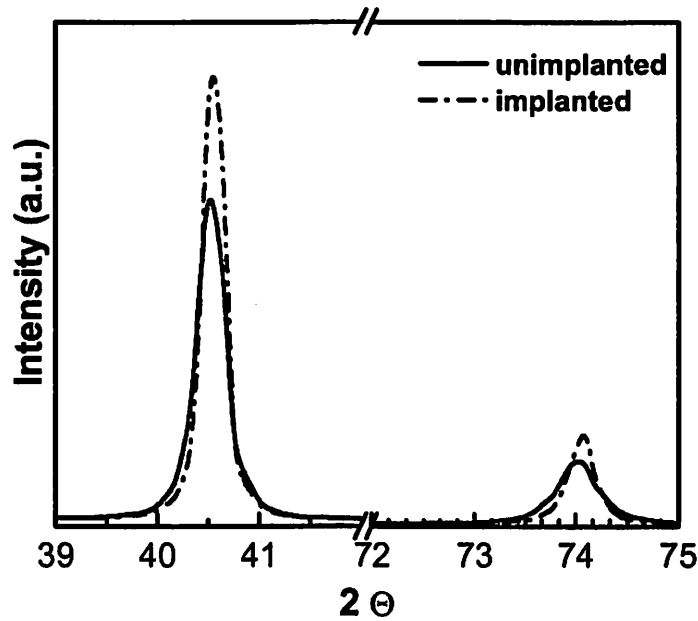


Figure 2.20 Comparison of the XRD intensities for the 110 and 112 peaks.

observed in nitrogen implanted wafers. **Figure 2.20** compares the intensities for the (110) and (112) peaks, indicating that both Mo gates have predominantly a (110) texture. **Figures 2.21 (a–d)** shows the details of diffraction peaks, indicating that peaks for  $\text{Mo}_2\text{N}$  are observed in the nitrogen implanted sample while no such peaks are observed in the unimplanted sample. **Figure 2.22 (a–b)** compares the dependence of  $\text{Mo}_2\text{N}$  (112) peak intensity on nitrogen implant dose ( $5 \times 10^{15}$ ,  $1 \times 10^{16} \text{ cm}^{-2}$ ) and subsequent annealing temperature (700, 900°C). It can be seen that increasing the dose or annealing temperature leads to an increase in the  $\text{Mo}_2\text{N}$  peak intensity. It is believed that an increase

in annealing temperature enhances segregation of nitrogen at the Mo/SiO<sub>2</sub> interface, resulting in a lowering of the interfacial work function.

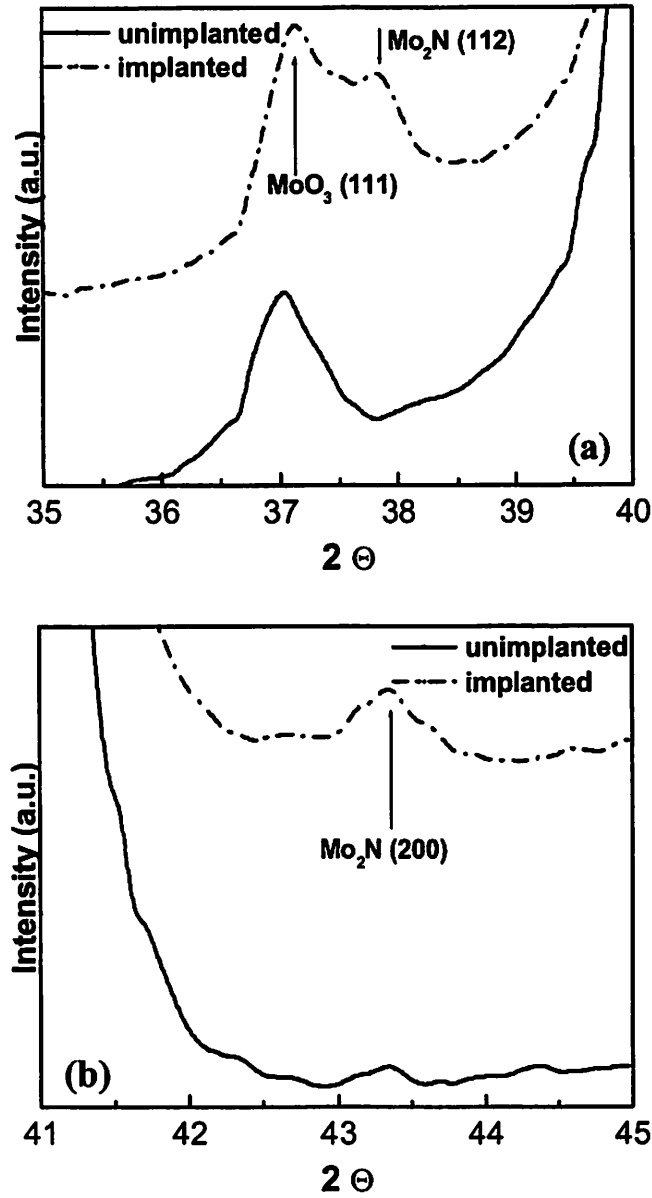


Figure 2.21 (a-d) Details of XRD peaks highlighted in Figure 2.19. Peaks indicative of Mo<sub>2</sub>N formation are observed in the nitrogen implanted sample while no such peaks are observed in the control (un-implanted) sample.

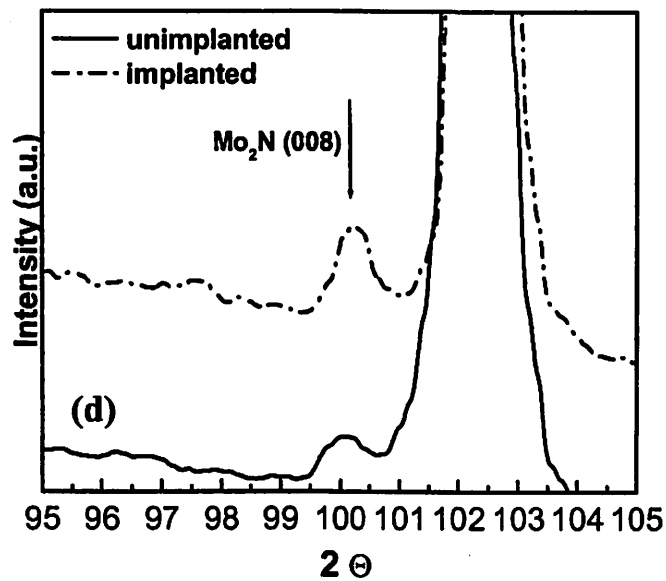
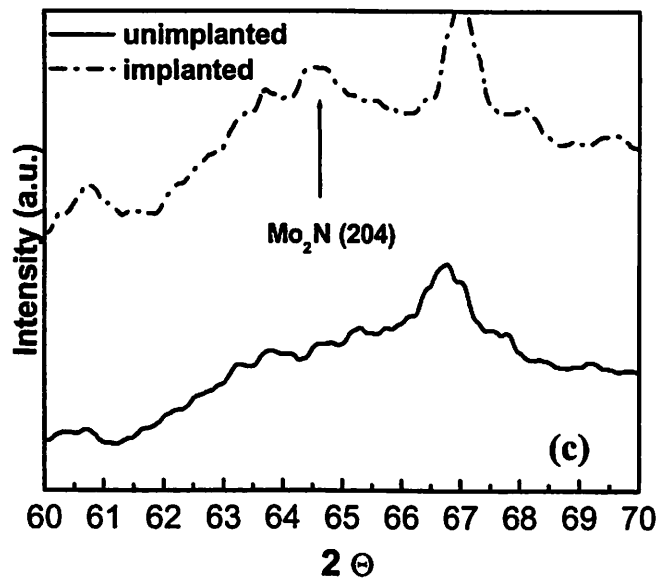


Figure 2.21 (a-d) Details of XRD peaks highlighted in Figure 2.19. Peaks indicative of  $\text{Mo}_2\text{N}$  formation are observed in the nitrogen implanted sample while no such peaks are observed in the control (un-implanted) sample.



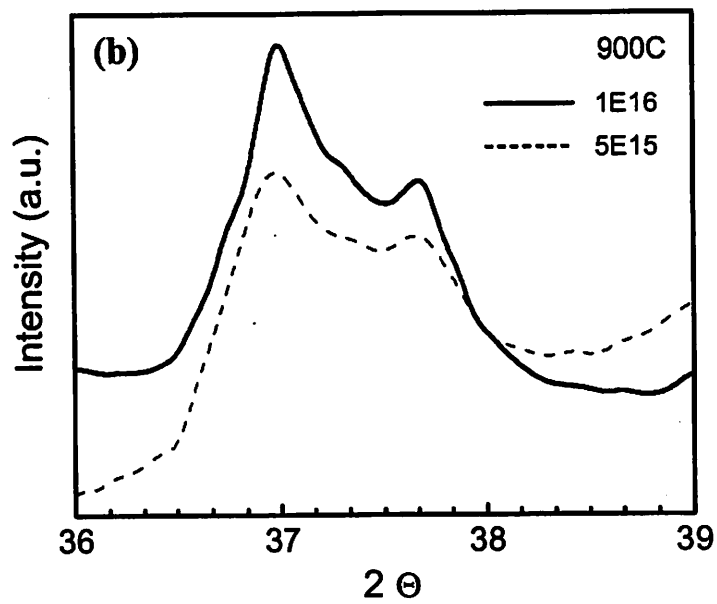
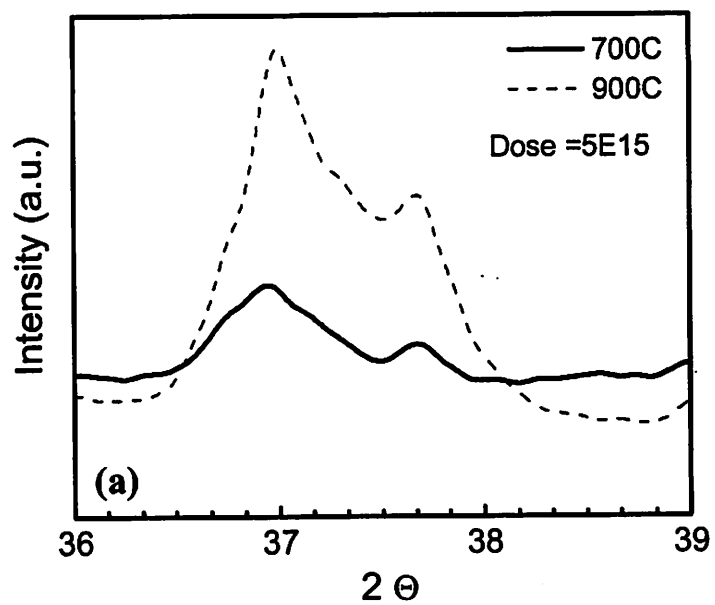
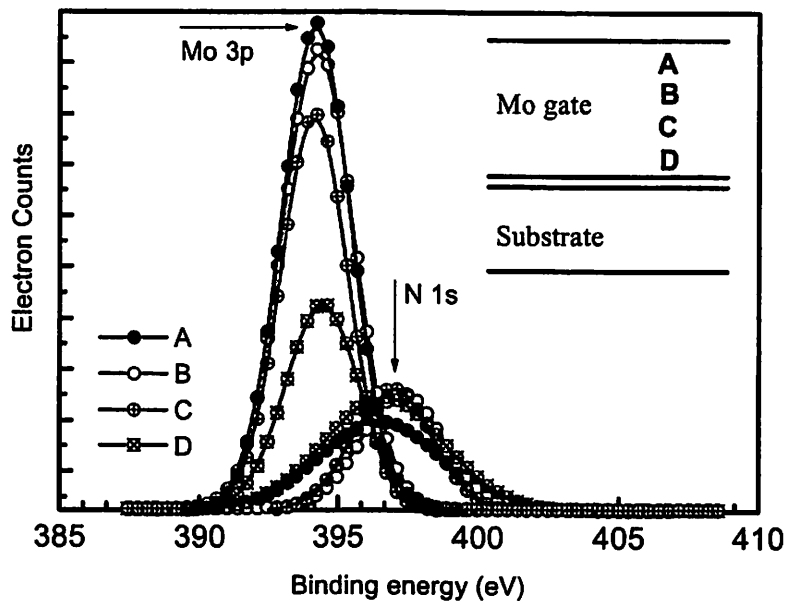


Figure 2.22 (a-b) Details of XRD peaks highlighted in Figure 2.19. Peaks indicative of  $Mo_2N$  formation are observed in the nitrogen implanted sample while no such peaks are observed in the control (un-implanted) sample.

To further analyze the chemical changes in the nitrogen implanted Mo films, x-ray photoelectron spectroscopy (XPS) analysis was performed. **Figure 2.23** shows the results of XPS analysis illustrating electron counts vs. binding energy for Mo and N signals at several points through the depth of Mo film. There is a significant overlap between the Mo 3p peak and the N 1s peak since they are separated by ~4 eV. Hence, the overall peak was deconvoluted to isolate the Mo and N contributions. A stronger N signal accompanies the Mo signal when approaching the Mo/SiO<sub>2</sub> interface (point D). It is worth to note that the XPS analysis does not show the formation of MoO<sub>2</sub>, MoO<sub>3</sub> or Mo-silicide at the SiO<sub>2</sub>



**Figure 2.23** Mo 3p and N 1s binding energy at different positions across the Mo film thickness. A significant N signal is observed throughout the Mo film, indicating a diffusion of the implanted N across the film thickness.

interface, indicating the thermodynamic stability of Mo on SiO<sub>2</sub> at 900°C.

The diffusivity and solubility of N inside Mo at high temperatures (> 1000°C) have been evaluated [2.39,2.40]. While the relevant data for lower temperatures is not available, the extrapolated N diffusion coefficient at 900°C is very high ( $\sim 2 \times 10^{-8}$  cm<sup>2</sup>/s). Low temperature diffusivities in poly-crystalline thin films are expected to be much higher, since the grain boundary diffusion would tend to dominate over bulk diffusion in the low temperature regime [2.41]. The low solid solubility of N in Mo (<1% at 1000°C) supports the observed formation of Mo<sub>2</sub>N after N implantation. The implanted N atomic content for a dose of  $1 \times 10^{16}$  cm<sup>-2</sup> into a 15 nm Mo film is  $\sim 10\%$ . Thus, a combination of enhanced diffusivity and low solubility leads to the segregation of N and formation of Mo<sub>2</sub>N in N implanted Mo films. Therefore, the work function shift of nitrogen implanted Mo gate results from a simultaneous evolution of the Mo film morphology and chemistry at the gate dielectric interface.

**Figure 2.24** shows the sheet resistance ( $R_S$ ) of 100-nm-thick Mo films for various nitrogen ion implantation doses after 900°C, 1-minute rapid thermal annealing (RTA) in nitrogen ambient. The sheet resistance increases slightly with implantation dose because of the increased impurity (nitrogen in Mo) scattering. The average grain

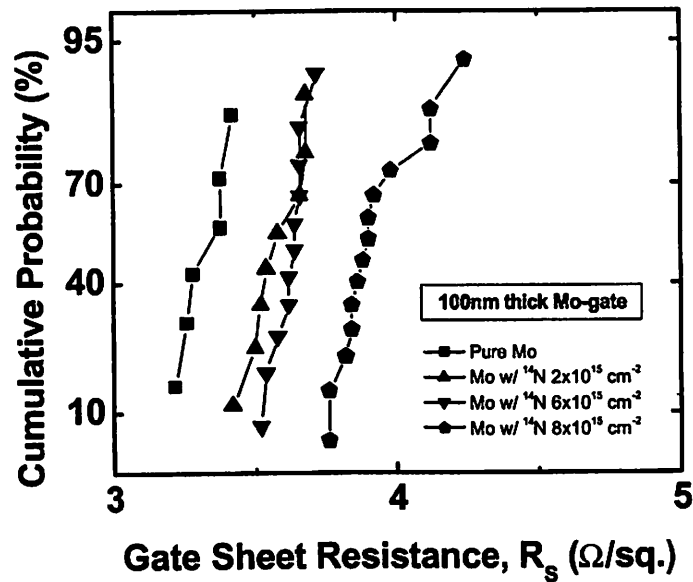


Figure 2.24 Sheet resistance ( $R_s$ ) of 100 nm-thick Mo films for various  $^{14}\text{N}^+$  implantation doses.

size was estimated from the full-width at half maximum (FWHM) of X-ray diffraction peaks using the Scherrer formula, as shown in Figure 2.20 [2.42]. The estimated grain sizes for pure and nitrogen implanted (dose of  $1 \times 10^{16} \text{ cm}^{-2}$ ) Mo films are 46.3 nm and 63.8 nm, respectively. It is worth noting that the grain size increases as more nitrogen is incorporated [2.43].

## 2.6 Summary

This chapter summarized the requirements of metal gate technology for the future nanoscale CMOS technology generations. Molybdenum (Mo) is a promising

candidate gate material for fully depleted (FD) silicon-on-insulator (SOI) CMOS devices. Damage-free sputtering and high-selectivity dry etching processes for Mo gate films have been developed. With a plasma charge trap (PCT), sputtering damage to the gate dielectric can be minimized for improved drive current and gate oxide integrity (GOI). Mo gate electrodes can be patterned without leaving any residue or stringers due to the highly selective etch process. The work function of a Mo gate can be tuned by nitrogen implantation followed by a thermal anneal, to provide a means for adjusting the threshold voltage of thin-body MOSFETs. The work function shift of nitrogen implanted Mo gate is resulted from the simultaneous modification of the microstructure and chemistry of Mo film at the gate dielectric interface. The nitrogen implantation induces a slight increase in sheet resistance due to impurity scattering, although it results in a slightly larger average grain size.

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## 2.7 References

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# Chapter 3

## High- $k$ Gate Dielectric Technology

### 3.1 Introduction

Over the past several decades, the successful improvement in CMOS transistor performance and integration of ultra-large scale (ULSI) semiconductor devices has relied on scaling down the silicon dioxide ( $\text{SiO}_2$ ) or its derivatives ( $\text{SiO}_x\text{N}_y$ ) gate dielectric. The advantage of primary reasons are that thermally grown amorphous  $\text{SiO}_2$  dielectric are low fixed charge densities ( $Q_f$ ) on the order of  $q10^{10} \text{ cm}^{-2}$ , mid-gap interface state densities ( $Q_{it}$ ) of  $\sim q10^{10} \text{ cm}^{-2}$  and a large breakdown electric field over 10 MV/cm [3.1]. Thinner gate oxide increases the drive current of transistors by increasing the areal gate

capacitance and suppresses short channel effects by providing strong gate control of the channel potential.

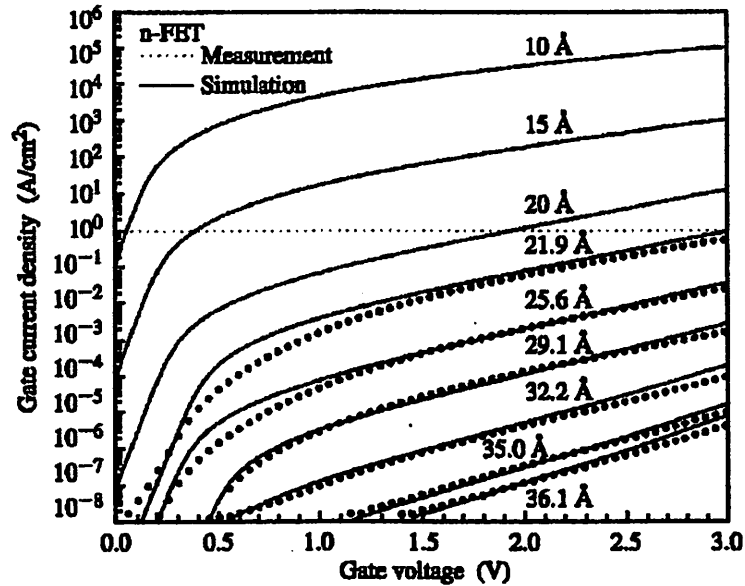


Figure 3.1 Measured and simulated  $I_G$ - $V_G$  characteristics under inversion conditions of n-channel MOSFETs. The dotted line is indicates the 1 A/cm<sup>2</sup> limit for gate leakage current [3.2]

However, silicon dioxide is not attractive as a gate dielectric when its thickness approaches 1 nm; one of the fundamental issues for scaling of the gate dielectric thickness is the exponential increase in direct tunneling current with decreasing SiO<sub>2</sub> thickness. For example, gate leakage current density can rise to 0.1 – 10 A/cm<sup>2</sup> at 1.0 V gate voltage for 1.5 – 2 nm thick SiO<sub>2</sub>, as shown in **Figure 3.1** [3.2] This high current can affect the circuit operations and dissipate power. Even though high power

dissipation may be tolerable for high-performance micro-processor applications, this will lead to serious problems for low-power portable applications. Theoretical modeling studies of the SiO<sub>2</sub>/Si interface predicted that the physical thickness of SiO<sub>2</sub> will be fundamentally limited to 0.7 nm due to the overlap of Si-rich interface regions from the silicon channel and the poly-silicon gate, causing an effective electrical short through the dielectric [3.3]. However, the practical scaling limit of SiO<sub>2</sub> as the gate dielectric will be 1.2 nm mainly due to interface roughness, degraded transistor performance, dielectric reliability, etc. [3.4, 3.5, 3.6].

The reduction of the gate leakage current density is, therefore, the most compelling reason for replacing SiO<sub>2</sub>-based dielectrics. For a given CMOS technology generation, devices are designed with a specific areal gate capacitance, which is a function of the dielectric constant ( $k$ ) and physical thickness ( $T_{ox}$ ). In order to decrease the leakage current while maintaining the same gate capacitance (or equivalent oxide thickness,  $EOT$ ), a thicker film with a higher dielectric constant should be adopted. Many materials appear favorable as an alternative gate dielectric, but very few materials

are promising with respect to the key basic material properties, compatibility with CMOS processes and integration, as will be discussed in the following section.

## 3.2 Material Selection for Gate Dielectric

Several materials have been investigated for replacing silicon dioxide or silicon nitride as the gate dielectric for sub-65nm CMOS technology generations [3.7]; these include the simple metal oxides such as tantalum oxide ( $\text{Ta}_2\text{O}_5$ ) [3.8], titanium oxide ( $\text{TiO}_2$ ) [3.9], lanthanum oxide ( $\text{La}_2\text{O}_3$ ) [3.10], yttrium oxide ( $\text{Y}_2\text{O}_3$ ) [3.11], cerium oxide ( $\text{CeO}_2$ ) [3.12], aluminum oxide ( $\text{Al}_2\text{O}_3$ ) [3.13], zirconium oxide ( $\text{ZrO}_2$ ) [3.14], and hafnium oxide ( $\text{HfO}_2$ ) [3.15, 3.16], and the ferroelectric materials such as barium strontium titanate (BST) [3.17]. As a promising candidate gate dielectric, however, the new materials should demonstrate the key basic material properties including high permittivity (or dielectric constant), large potential barrier heights to prevent tunneling current, thermodynamic stability in contact with silicon, good interface quality and film morphology [3.18]. In addition, the high-k dielectric should be compatible with the gate electrode material(s) and CMOS process integration without compromising device reliability and circuit performance [3.19].

To minimize the gate tunneling current, it seems reasonable to adopt a material whose dielectric constant ( $k$ ) is as high as possible, allowing an increase in the high- $k$  film thickness for the same SiO<sub>2</sub> equivalent oxide thickness. However, if the high- $k$  dielectric thickness approaches the gate length, fringing fields from source/drain regions to the channel becomes significant, resulting in weak control over the channel potential, hence compromising short channel behavior; this is the so called fringing-induced-barrier-lowering (FIBL) effect [3.20]. Intensive simulation study [3.21] predicted that

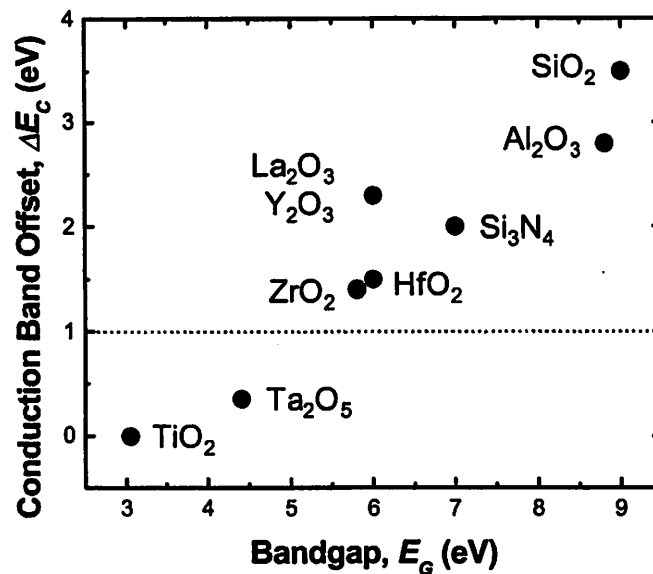


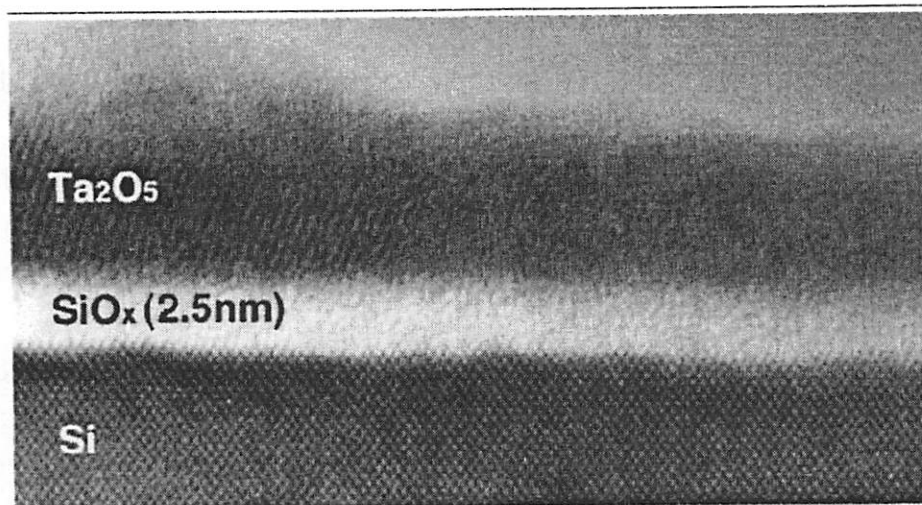
Figure 3.2 Theoretically calculated conduction band barrier height ( $\Delta E_C$ ) vs. energy bandgap ( $E_G$ ). In general, a larger bandgap leads to a larger conduction band barrier height [3.3].

when the ratio of dielectric thickness to gate length ( $T_{ox}/L_G$ ) is great than 1/5, more than 10% degradation in subthreshold swing occurs, ruling out the use of very high- $k$  ( $> 40$ ) materials such as  $TiO_2$  and BST.

Together with the high dielectric constant, the conduction barrier height ( $\Delta E_C$ ) should be high enough to prevent a significant tunneling current because both the thickness and height of the barrier have an exponential influence upon the tunneling leakage current. Barrier heights have been theoretically calculated for several potential high- $k$  materials [3.22]. In general, a larger bandgap ( $E_G$ ) leads to a larger conduction band offset ( $\Delta E_C$ ), as shown in **Figure 3.2**. However, for most high- $k$  dielectrics, the bandgap is inversely proportional to the dielectric constant [3.23]. Hence, even if there is a significant reduction of tunneling leakage current due to the increased thickness, this may be compensated by the reduction of the bandgap, or barrier height. Therefore, a high- $k$  dielectric, whose conduction band offset ( $\Delta E_C$ ) is smaller than 1.0 eV, may not replace the silicon dioxide [3.22]. Identifying a dielectric that provides even a moderate increase in dielectric constant, but that also produces a sufficiently large tunneling barrier height is indispensable.  $La_2O_3$ ,  $Y_2O_3$ ,  $ZrO_2$ , and  $HfO_2$  offer relatively high dielectric constant and conduction band-offset.



For very thin gate dielectrics, the interface with silicon determines the overall electrical properties. Thermodynamic stability in direct contact with silicon during CMOS fabrication processes is therefore indispensable [3.24]. Most high- $k$  metal oxides including  $Ta_2O_5$  and  $TiO_2$  are not thermally stable on silicon; reacting with silicon to form an undesirable  $SiO_2$  or medium- $k$  interfacial layer, or metal silicide [3.25, 3.26], as shown in **Figure 3.3**. The underlying  $SiO_2$  or interfacial layer limits the scaling of the equivalent oxide thickness (EOT). For example, if 1.5 nm EOT is required and 1.0 nm  $SiO_2$  is formed at the interface, only 0.5 nm remains for the high- $k$  material, making the use of high- $k$  material ineffective due to the significant increase in tunneling leakage



**Figure 3.3** An example of thermodynamic instability of  $Ta_2O_5$  in direct contact with silicon. A reaction at the interface results in the formation of an undesirable thin  $SiO_2$  layer [3.26]

current, as illustrated in **Figure 3.4**.

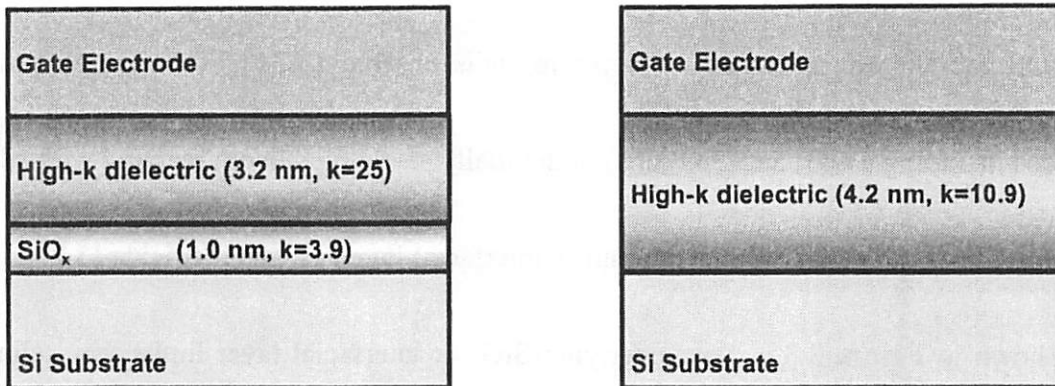


Figure 3.4 Comparison of stacked gate dielectric and single layer gate dielectrics in transistor. Both gate dielectrics result in the same equivalent oxide thickness,  $EOT = 1.5$  nm. Ultimate scaling of  $EOT$  can be limited due to the interface layer formation.

Glassy phase (amorphous) gate dielectrics during CMOS fabrication are very desirable, because they will have isotropic electrical properties and not have grain boundaries, which typically serve as leakage current paths [3.18]. In addition, grain size and orientation changes lead to significant variations in dielectric constant. However, nearly all metal oxides under investigation form polycrystalline films upon source/drain annealing, which may necessitate an intentionally grown amorphous interfacial layer to reduce the leakage current. It is worthwhile to note that despite relatively low dielectric

constant (9~15),  $\text{Al}_2\text{O}_3$  [3.27],  $\text{ZrSi}_x\text{O}_y$  [3.28] and  $\text{HfSi}_x\text{O}_y$  [3.29] appears promising because their morphology remains amorphous during high temperature annealing.

Low midgap interface state density ( $Q_{it}$ ) and fixed charge density ( $Q_f$ ) are crucial to meet the performance requirements of nanoscale CMOS transistors; channel carrier mobilities, threshold voltage shift, and subthreshold swing can be greatly affected. This will lead to the need for a sufficiently high quality interface with the silicon channel, challenging for high- $k$  dielectrics as compared to  $\text{SiO}_2$  [3.19].

Thus, it may be appropriate to replace  $\text{SiO}_2$  gate dielectric with a material which provides only moderate increase (15 – 30) in dielectric constant, but also provides a large tunneling barrier height ( $> 1.0$  eV) and high-quality ( $Q_{it}, Q_f < 10^{11} \text{ cm}^{-2}$ ) interface to the silicon channel.

### **3.3 $\text{HfO}_2$ Gate Dielectric**

Hafnium dioxide ( $\text{HfO}_2$ ) has shown much promise as a candidate to replace  $\text{SiO}_2$ , as discussed in Chapter 3.2; it has a high dielectric constant ( $k \sim 25$ ) with the bandgap of 5.7 eV, large energy-band offsets ( $\Delta E_C \sim 1.5$  eV), thermodynamic stability in direct contact with silicon substrate, and negligible frequency dispersion [3.22, 3.30, 3.31,

3.32]. For the gate electrode, polycrystalline silicon (poly-Si) has been the preferred material because it offers simplicity of process integration. However, when poly-Si is used as the gate material together with HfO<sub>2</sub> as the gate dielectric, increases in equivalent oxide thickness (EOT) and gate leakage current can result upon high-temperature annealing (*e.g.* during source/drain dopant activation) due to interfacial layer formation at the HfO<sub>2</sub>/Si interface [3.33]. Recently, polycrystalline silicon-germanium (poly-SiGe) has received much attention as an alternative gate-electrode material, because it alleviates gate depletion and boron penetration issues, and yields thinner EOT for HfO<sub>2</sub> gate dielectric [3.34, 3.35, 3.36].

### ***3.3.1 Fabrication of HfO<sub>2</sub> MOS Capacitors***

MOS capacitors were fabricated on epitaxially grown silicon (p-type dopant concentration ( $N_A$ ) below  $10^{15}$  cm<sup>-3</sup>) on heavily doped p+ silicon wafers. **Figure 3.5** summarizes the fabrication sequences for HfO<sub>2</sub> MOS capacitors. 100 nm thick SiO<sub>2</sub> was thermally grown in wet oxidation furnace at 950°C and patterned to define the device active regions. Pure Hf was then deposited by DC magnetron sputtering at 300°C and 210 W power (base pressure of  $5 \times 10^{-8}$  Torr), and rapid thermal annealing at 700°C, 30s in

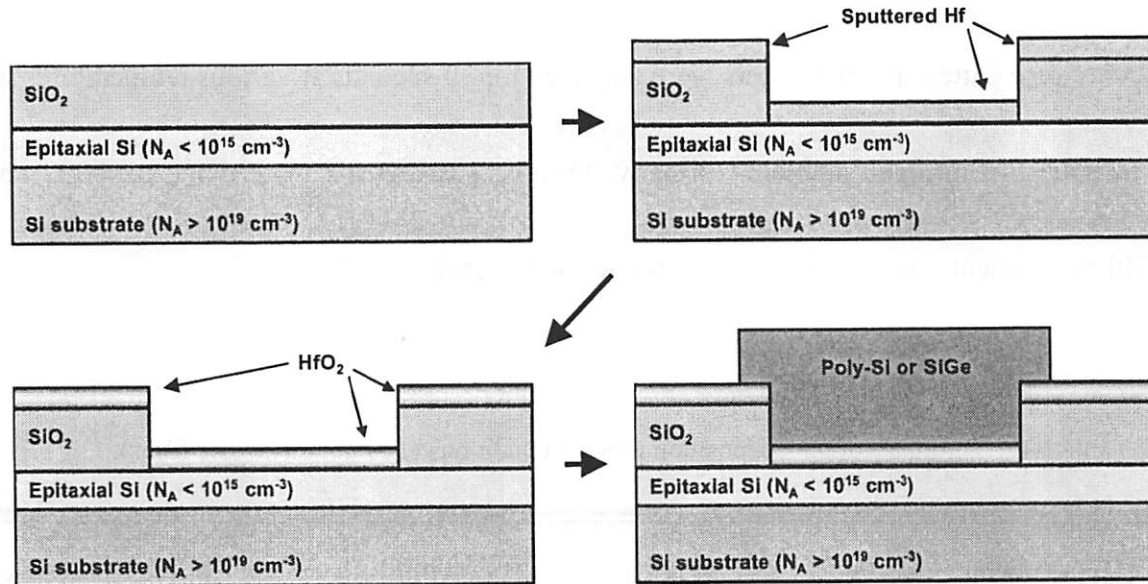


Figure 3.5 Fabrication sequences for HfO<sub>2</sub> gate dielectric MOS capacitors.

oxygen ambient was used to form HfO<sub>2</sub>. It is worth noting that the wafers were precleaned in a diluted HF solution to remove the chemical oxide and no intentional interfacial layer was grown before HfO<sub>2</sub> formation. Afterwards, the gate electrode materials (either poly-Si or Si<sub>0.75</sub>Ge<sub>0.25</sub>) were deposited in a conventional low-pressure chemical vapor deposition (LPCVD) furnace at 550°C, using either SiH<sub>4</sub> or Si<sub>2</sub>H<sub>6</sub> as the gaseous Si source, and GeH<sub>4</sub> as the gaseous Ge source. In order to minimize gate depletion, all the gate films were *in-situ* n<sup>+</sup> doped using phosphine (PH<sub>3</sub>). The gate deposition conditions are summarized in **Table 3.1**. Shorter deposition times were used

for poly-Si<sub>0.75</sub>Ge<sub>0.25</sub> and Si<sub>2</sub>H<sub>6</sub>-based poly-Si in order to achieve the same film thickness. After gate patterning, the wafers were annealed for 30 seconds at various temperatures up to 950°C in nitrogen ambient. Finally, forming gas anneals were done at 600°C and 400°C sequentially to improve the interface state density [3.37].

Table 3.1 Summary of gate deposition process conditions (100 nm thick gate films).

Gate material	Poly-Si	Poly-Si	Poly-Si <sub>0.75</sub> Ge <sub>0.25</sub>
Precursor gas	SiH <sub>4</sub>	Si <sub>2</sub> H <sub>6</sub>	SiH <sub>4</sub> + GeH <sub>4</sub>
Deposition temperature	550°C	550°C	550°C
Deposition time	270 min	7.5 min	54 min

### 3.3.2 *Experimental Results and Discussion*

Figure 3.6 shows the measured high-frequency  $C-V$  characteristics of the HfO<sub>2</sub> capacitors. For the poly-Si gated capacitors, the capacitance decreases with increasing the subsequent annealing temperature. This is more pronounced for SiH<sub>4</sub>-based poly-Si

gated capacitors ( $\Delta EOT = 1.3\text{nm}$ ) compared to  $\text{Si}_2\text{H}_6$ -based poly-Si gated capacitors ( $\Delta EOT = 0.65\text{nm}$ ).

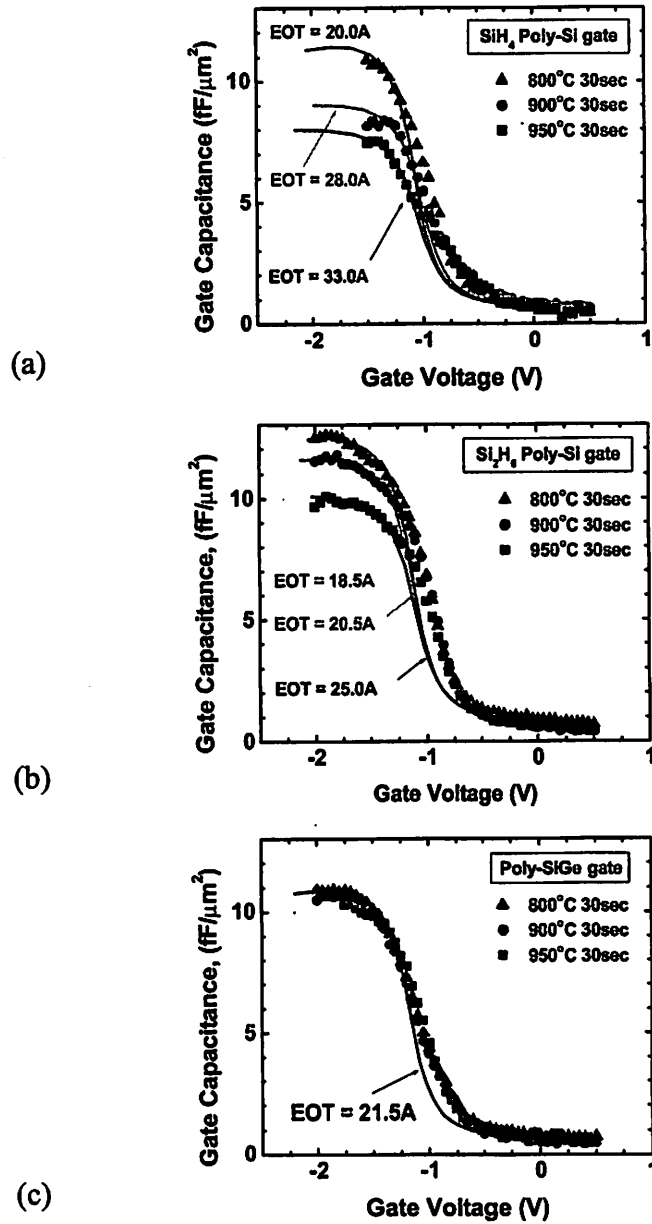


Figure 3.6 High-frequency (100 KHz) C-V curves for  $\text{HfO}_2$  MOS capacitors with (a) poly-Si gate deposited using  $\text{SiH}_4$ , (b) poly-Si gate deposited using  $\text{Si}_2\text{H}_6$ , and (c) poly- $\text{Si}_{0.75}\text{Ge}_{0.25}$  gate. The solid lines shown were obtained from quantum-mechanical simulation and used to extract the EOT values as indicated.

These results indicate that a reduction in the gate deposition time is helpful to prevent increases in EOT with high-temperature annealing. However, for poly-SiGe gated capacitors, negligible change in capacitance ( $\Delta EOT \cong 0$ ) was observed for annealing temperatures up to 950°C. This result can only be partially attributed to the reduced gate deposition time (refer to Table 3.1).

Figure 3.7 shows the measured leakage current characteristics of the HfO<sub>2</sub> capacitors. The leakage current for SiH<sub>4</sub>-based poly-Si capacitors is much larger than that of Si<sub>2</sub>H<sub>6</sub>-based poly-Si capacitors. This indicates that reduction of thermal budget in gate deposition is very effective for reducing leakage current through the HfO<sub>2</sub> gate dielectric. It is worthwhile to note that the leakage current increases with annealing temperature, *i.e.*, thicker EOT. Both *C-V* and *I-V* measurements show that poly-Si as a gate material yields a thicker EOT with higher leakage current for HfO<sub>2</sub> gate dielectric, as compared with poly-Si<sub>0.75</sub>Ge<sub>0.25</sub>. These results cannot be explained simply by the formation of an interfacial layer.

In order to elucidate the leakage mechanism, the dependence of gate leakage current on temperature was investigated for HfO<sub>2</sub> capacitors with either SiH<sub>4</sub>-based poly-Si or poly-SiGe gate material, as shown in Figure 3.8. The leakage current shows an



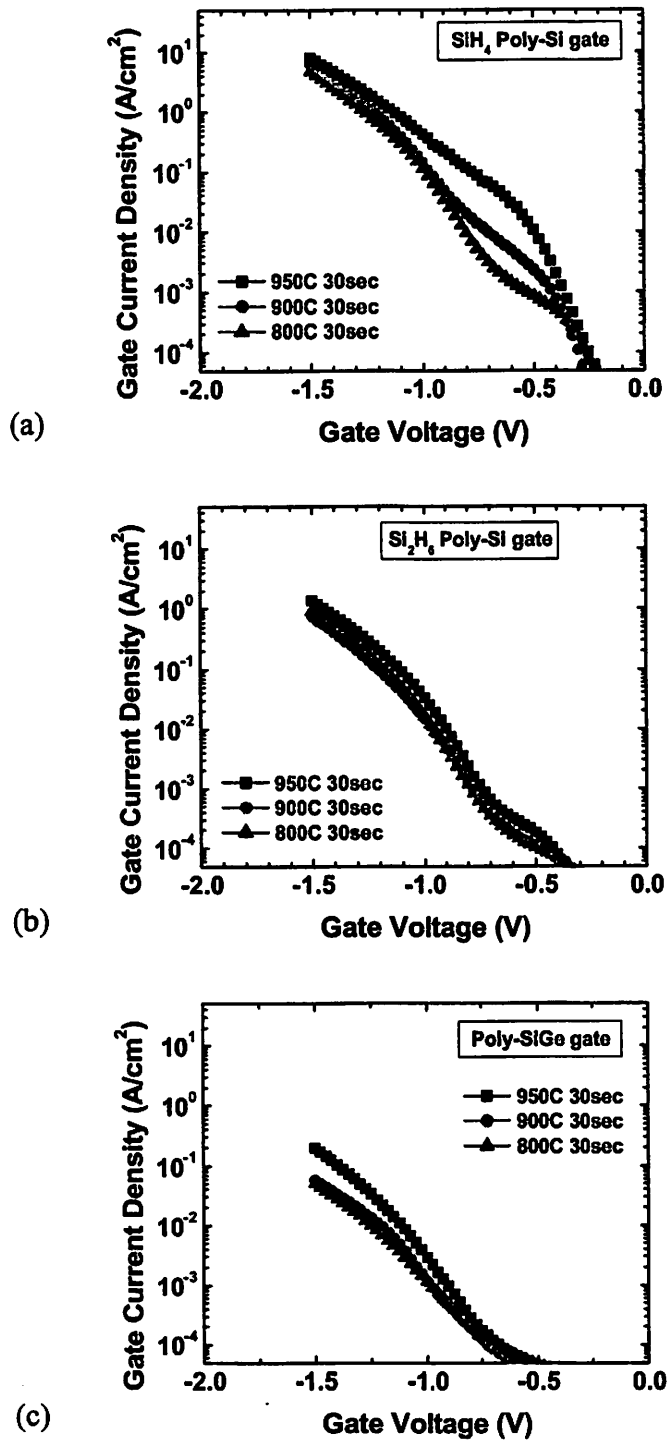


Figure 3.7 Gate leakage current characteristics of HfO<sub>2</sub> MOS capacitors with (a) poly-Si gate deposited using SiH<sub>4</sub>, (b) poly-Si gate deposited using Si<sub>2</sub>H<sub>6</sub>, and (c) poly-SiGe gate.

exponential dependence on voltage and temperature, which corresponds to Frenkel-Poole emission [3.38]. From the characteristic slope for different gate voltages from  $-1.0\text{V}$  to  $-1.5\text{V}$ , the trap energy level in  $\text{HfO}_2$  can be extracted. Since the same trap energy level can be seen for each gate material, and the poly-SiGe gated capacitors exhibit  $\sim 100$  times lower leakage current, it is concluded that the  $\text{HfO}_2$  defect density is lower for poly-SiGe gate material.

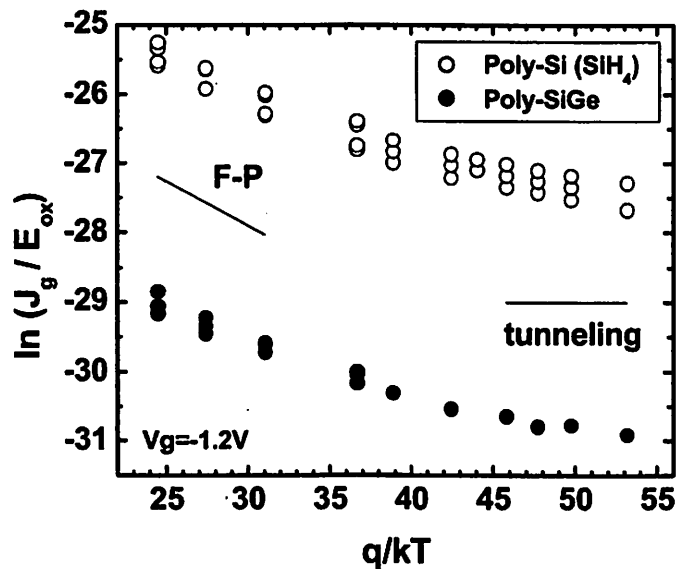


Figure 3.8 Leakage current vs. temperature for  $\text{HfO}_2$  MOS capacitors with either poly-Si gate deposited using  $\text{SiH}_4$ , or poly-SiGe. The devices were annealed at  $900^\circ\text{C}$ .

Figure 3.9 shows the cross-sectional transmission electron microscopy (XTEM), which reveals that a bottom interfacial layer exists at the  $\text{HfO}_2/\text{Si}$  interface before and

after high-temperature annealing for both poly-Si and poly-SiGe gate materials. This interfacial layer was likely formed during the Hf oxidation process due to excessive oxidation. Thus, the increase in leakage current upon high-temperature annealing cannot be attributed to the bottom interfacial layer.

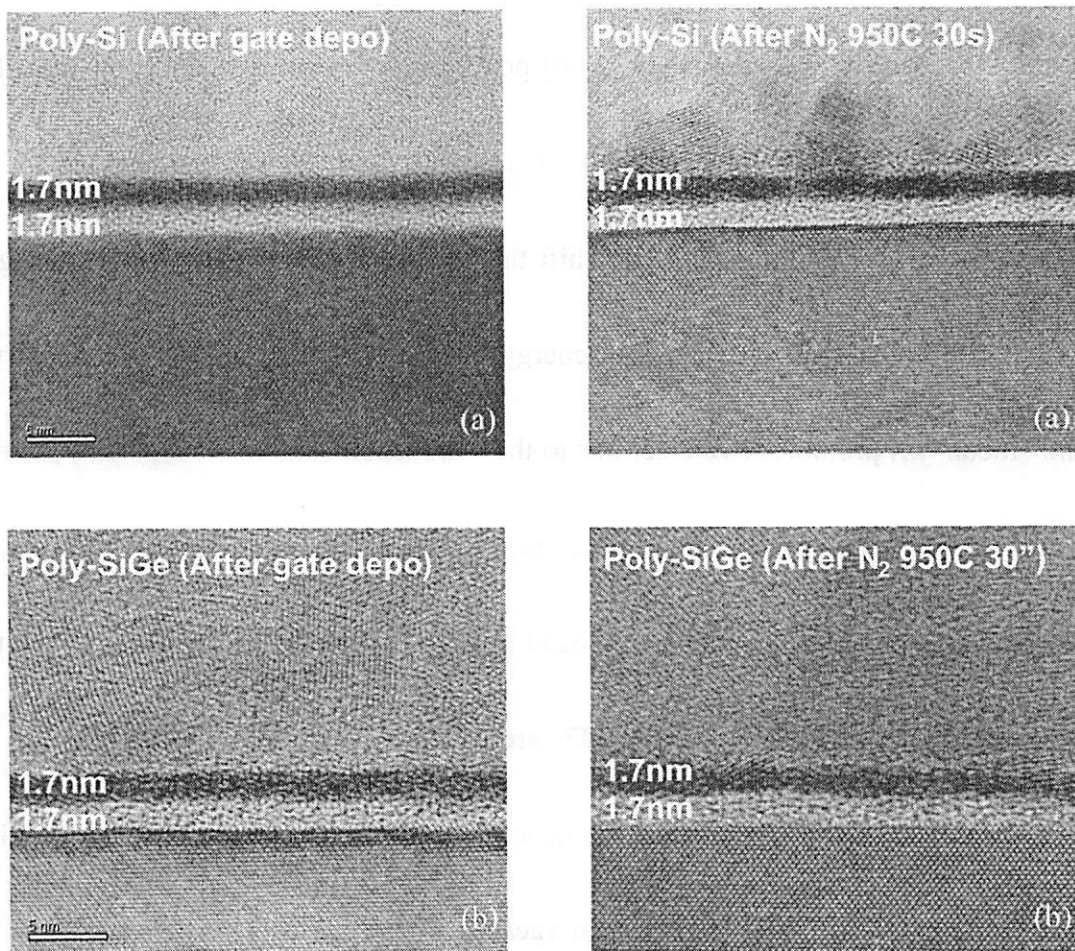


Figure 3.9 Cross-sectional transmission electron microscopy (X-TEM) for HfO<sub>2</sub> gate stacks annealed at 950°C with (a) poly-Si and (b) poly-SiGe gates. Bottom interface layer exists at the HfO<sub>2</sub>/Si interface before and after high-temperature annealing.

On the other hand, at the top interface between the  $\text{HfO}_2$  and the gate electrode, the formation of an interfacial film upon annealing was observed. Although it has been predicted to be thermodynamically stable in direct contact with silicon [3.39],  $\text{HfO}_2$  readily forms an interfacial silicate layer; the reaction  $\text{HfO}_2 + \text{Si} \rightarrow \text{Hf} + \text{SiO}_2$  does not seem likely to occur, since the Gibbs free energy change is positive ( $\Delta G = + 1.14 \text{ eV}$  at 1000K). However, a recent study [3.40] pointed out that oxygen transport from  $\text{HfO}_2$  into Si generates conduction electrons in the  $\text{HfO}_2$ , and the energy supplied by their transition to lower energy states can shift the overall  $\Delta G$  to negative values. Figures 3.10 illustrate the sequence using an energy band diagram. Transport of oxygen (O) into silicon (Si) promotes two electrons to the conduction band in  $\text{HfO}_2$ . The promoted electrons then fall into available lower-energy states: the unoccupied oxygen vacancy ( $V_o$ ) level in  $\text{HfO}_2$  and the conduction band in Si. The energy difference between these states and the conduction band of  $\text{HfO}_2$  are  $-1.2 \text{ eV}$  and  $-1.5 \text{ eV}$ , respectively. The electron transfer results in a charge dipole at the oxide/Si interface as depicted in Figure 3.10 (b). Transport of O (i.e., oxygen vacancy formation inside  $\text{HfO}_2$ ) continues until the electrostatic dipole energy balances the effective reaction energy. Hence, interfacial silicate layer formation is thermodynamically driven and imposes thermal process

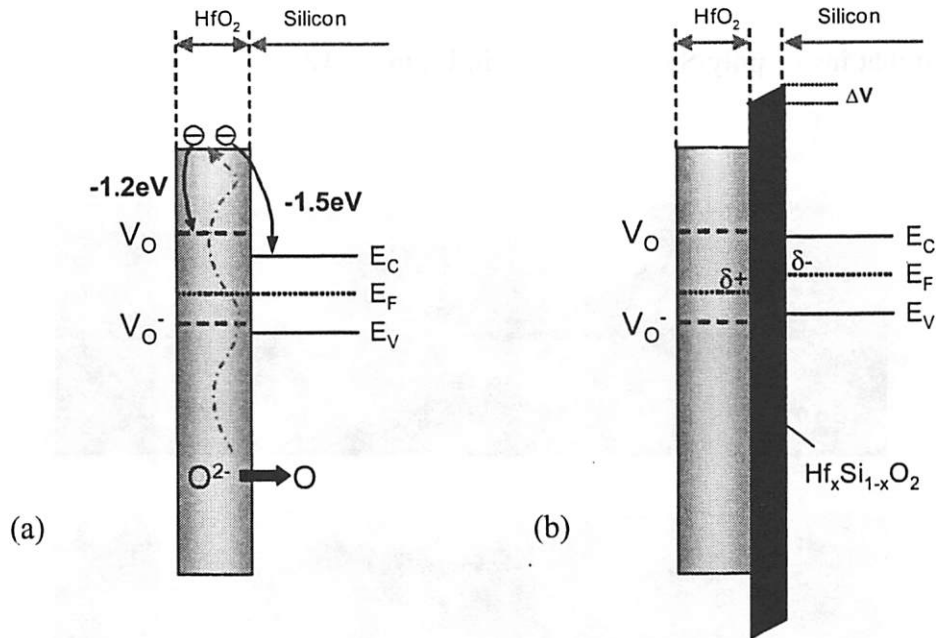


Figure 3.10 (a) Electron transfer resulting from oxygen transport across the interface. Two electrons promoted to  $E_C$  in  $\text{HfO}_2$  fall to unoccupied states,  $V_O$  in  $\text{HfO}_2$  and  $E_C$  in the silicon substrate. (b) an interfacial silicate layer and potential drop is formed.

constraints for  $\text{HfO}_2$  on Si. It is interesting to note that 0.2 V potential drop has been observed by x-ray photoelectron spectroscopy (XPS) [3.41]. In contrast, high-k dielectrics in contact with Ge do not form interfacial layers because of the larger  $\Delta G$  for  $\text{GeO}_2$  formation [3.42].

**Figure 3.11** shows that the density of this interfacial layer is much higher for poly-Si gate capacitors after 950°C 30s RTA in nitrogen ambient. From compositional analyses using energy dispersive x-ray spectrometry (EDX), it was found that both

oxygen (O) and hafnium (Hf) concentration inside poly-Si near the top interface is much higher than that inside poly-SiGe, as shown in **Figure 3.12**.

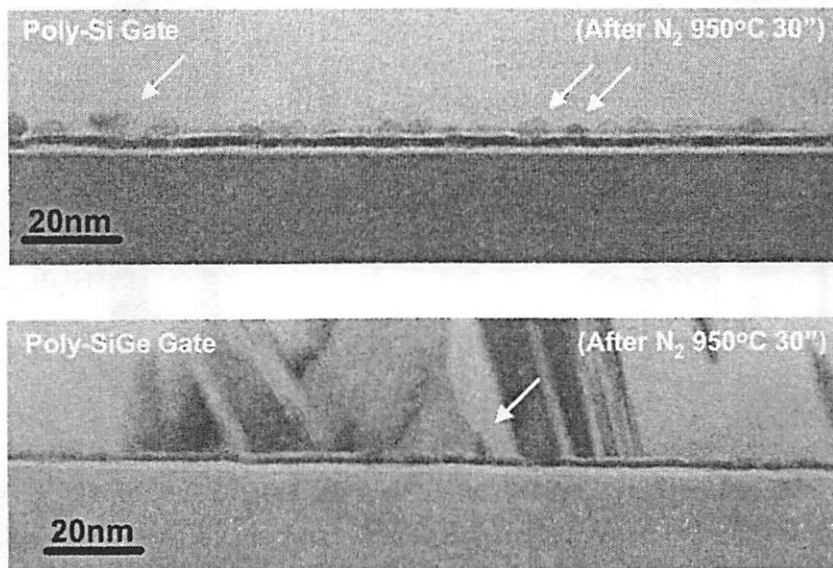


Figure 3.11 Cross-sectional transmission electron microscopy (X-TEM) for HfO<sub>2</sub> gate stacks annealed at 950°C with (a) poly-Si and (b) poly-SiGe gates. Defect density at top interface layer for poly-Si gate is much higher than that for poly-SiGe gate.

**Figure 3.13** shows the x-ray photoelectron spectroscopy (XPS) spectra of Hf 4f before and after amorphous Si deposition and after 950°C 30s RTA. The Hf 4f spectra did not exhibit an Hf-Si peak, indicating that no Hf silicide forms after 950°C 30s RTA. However, a +0.4 eV shift of the Hf-O peak was observed after silicon deposition; an additional +0.3 eV shift can be seen after 950°C 30s RTA. The positive shift of the Hf-

O peak is known to be caused by the second-neighbor effect due to Hf-O-Si formation [3.43]. These results indicate that the intermixing of Hf, Si, and O occurs during gate deposition and proceeds during subsequent thermal processing steps.

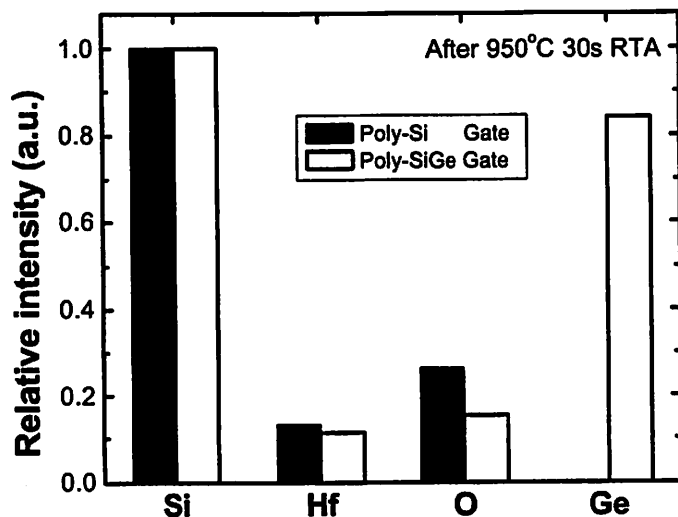


Figure 3.12 Comparison of relative intensity of energy dispersive X-ray spectrometry (EDX) analysis for gate/HfO<sub>2</sub> interface. The samples were annealed at 950°C 30s in N<sub>2</sub>.

Detail experimental investigation on the impact of the oxidation process on an HfO<sub>2</sub> film [3.44] found that insufficiently oxidized Hf films are non-stoichiometric, and the origin for the extra absorption peak (defects) detected in spectroscopic ellipsometry (SE) was attributed to the oxygen vacancies inside the HfO<sub>2</sub> films. It is worth noting that oxygen vacancies in metal oxides including ZrO<sub>2</sub> [3.45], TiO<sub>2</sub> [3.46], SnO<sub>2</sub> [3.47], and Ta<sub>2</sub>O<sub>5</sub> [3.48] are known to induce electronic states in the band gap.

Therefore, the increases in EOT and gate leakage current with an increase in thermal process budget likely result from the formation of an Hf silicate layer at the top (gate electrode) interface and oxygen-vacancy-trap assisted tunneling through the  $\text{HfO}_2$ . Also, the smaller increases in EOT and gate leakage current with poly-SiGe gate compared to those with poly-Si gate can be attributed to reduced transport of oxygen into the poly-SiGe gate electrode, hence leading to less interfacial layer formation.

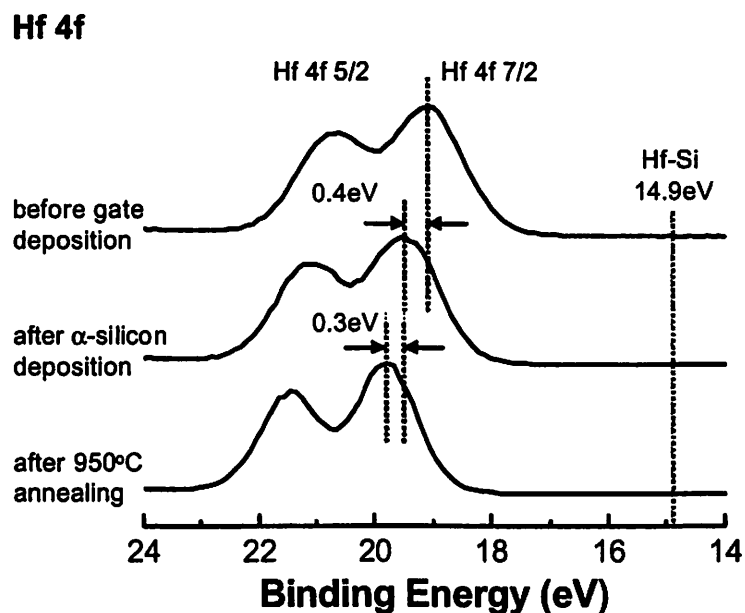


Figure 3.13 X-ray photoelectron spectroscopy (XPS) spectra of Hf 4f before and after amorphous Si deposition and after 950°C 30s RTA.



### 3.3.3 Considerations for alternative gate electrode materials

Successful integration of HfO<sub>2</sub> gate dielectric depends on the minimization of oxygen vacancy formation; a reduction in post gate deposition thermal process budget, and careful selection of the gate stack materials is therefore necessary. **Table 3.2** summarizes the heat of formation of oxide ( $E_{MOx}$ ), vacuum work function ( $\Phi_{M,vac}$ ), and effective reaction enthalpy ( $E_{eff}$ ) for candidate gate electrode materials on HfO<sub>2</sub> [3.49]. Most of these elements are found to exhibit negative  $E_{eff}$ , and thus would suffer from the Fermi-level pinning effect on HfO<sub>2</sub> [3.50]. Cobalt (Co), palladium (Pd), and ruthenium (Ru) appear to be promising candidates for a PMOS gate electrode material. However, n<sup>+</sup> doped germanium (Ge) is the only candidate with positive  $E_{eff}$  and low  $\Phi_{M,vac}$  suitable for a NMOS gate electrode material. The relatively small shift in effective work function for Mo-gate as compared to p<sup>+</sup> poly-Si gate can be expected by the difference in  $E_{eff}$  (-0.24 eV vs. -1.39eV). The addition of Al<sub>2</sub>O<sub>3</sub> into HfO<sub>2</sub> seems to be effective not only for increasing thermal stability [3.43], but also for reducing the Fermi-level pinning effect at the poly-Si gate interface [3.40] due to the lower  $E_{MOx}$  of Al<sub>2</sub>O<sub>3</sub> than HfO<sub>2</sub>.

Table 3.2 Summary of estimated effective reaction enthalpy for oxygen transfer from HfO<sub>2</sub> to gate electrode [3.49].

Gate electrode material	Vacuum work function, $\Phi_{M,vac}$ (eV)	Formation enthalpy of $E_{MOX}$ (eV) <sup>*1</sup>	Effective enthalpy, $E_{eff}$ (eV) <sup>*2</sup>
Al	4.06	-5.80 (Al <sub>2</sub> O <sub>3</sub> )	-2.57
Co	5.00	-1.65 (CoO)	0.65
Cr	4.50	-3.10 (CrO <sub>2</sub> )	-0.31
Cu	4.48	-2.47 (CuO)	0.34
Ge	4.00 <sup>*3</sup>	-3.01 (GeO <sub>2</sub> )	0.28
Hf	3.90	-5.94 (HfO <sub>2</sub> )	-2.55
Mo	4.95	-2.58 (MoO <sub>3</sub> )	-0.24
Ni	5.04	-3.39 (Ni <sub>2</sub> O <sub>3</sub> )	-1.14
Pd	5.22	-0.89 (PdO)	1.19
Pt	5.64	-4.07 (PtO <sup>*4</sup> )	-2.41
Ru	4.71	-1.58 (RuO <sub>2</sub> )	1.00
Si	4.05 <sup>*3</sup>	-4.73 (SiO <sub>2</sub> )	-1.49
Ta	4.15	-4.25 (Ta <sub>2</sub> O <sub>5</sub> )	-1.11
Ti	4.33	-4.90 (TiO <sub>2</sub> )	-1.94
V	4.30	-6.44 (V <sub>2</sub> O <sub>5</sub> )	-3.45
W	4.55	-3.06 (WO <sub>2</sub> )	-0.32
Zr	4.05	-7.62 (ZrO <sub>2</sub> )	-4.38

\*<sup>1</sup> normalized by number of oxygen atom.

\*<sup>2</sup> calculated using the following formula:  $E_{eff} = E_{MOx} - E_{HfO2} - 1.2eV - (1.5eV + \Phi_{M,vac} - E_{C,Si})$ .

\*<sup>3</sup>  $E_C$  is used for semiconductor materials.

\*<sup>4</sup> gaseous diatom bond enthalpy used.

## 3.4 Summary

This chapter reviewed the requirements for a high-k gate dielectric for future nanoscale CMOS technology generations. For hafnium oxide ( $\text{HfO}_2$ ), the impact of gate process technology on equivalent-oxide-thickness and leakage current was investigated. Even though hafnium oxide ( $\text{HfO}_2$ ) is one of the most promising candidates, successful integration of  $\text{HfO}_2$  into CMOS fabrication depends on the minimization of oxygen vacancy formation through careful selection of the gate electrode materials, and reduction of post-gate-deposition thermal budget.

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# Chapter 4

## Advanced Transistor Structures

### 4.1 Introduction

The scaling of CMOS devices has been continued over many technology generations, leading to improvement in both circuit performance and packing density; according to the Moore's law, integrated circuit complexity would increase at a rate of roughly a factor of two per year [4.1]. The semiconductor industry has kept this trend over thirty years, mainly fueled with technological innovations. The design of MOSFET at smaller dimensions has been predominantly guided by the constant electric field scaling theory [4.2] and its variants [4.3]; without changing the basic transistor

structure, lateral dimension can be scaled down together with reduction in vertical dimension and operation voltage ( $V_{DD}$ ), as illustrated in Figure 4.1.

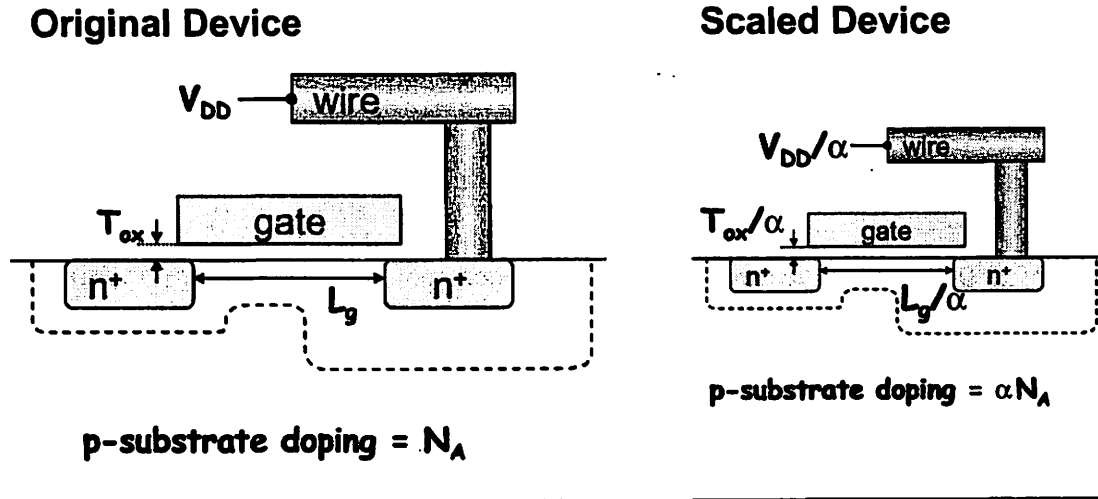


Figure 4.1 Schematic illustration of the constant electric field scaling theory of MOSFET devices [4.2].

To suppress the short channel effects, the channel potential in a MOSFET should be controlled by the gate electrode rather than the drain electrode; thus, the gate capacitance ( $C_g$ ) should be larger than the drain capacitance ( $C_d$ ). Otherwise, the characteristics of the MOSFET approach that of a resistor [4.3]. Traditionally, these issues have been successfully resolved by the reduction in gate oxide thickness ( $T_{ox}$ ), channel depletion width ( $W_D$ ) and source/drain junction depth ( $X_J$ ). However, as the device dimensions

shrink down to nanometer scale regime, the integration of ever increasing number of devices without degrading yield, and reliability faces fundamental physics barriers.

The increase in gate capacitance can be achieved through thinning down the gate oxide thickness. However, there are fundamental limitation in SiO<sub>2</sub> vertical scaling due to the direct tunneling current as discussed in Chapter 3 [4.4, 4.5], and still remaining issues in the alternative high-*k* dielectrics such as threshold voltage instability and mobility degradation [4.6, 4.7, 4.56, 4.57].

The influence of drain field on the channel potential can be shielded through increasing the channel and/or localized halo doping concentration (*i.e.*, minimizing the depletion depth) [4.8]. However, the very thin depletion depths needed for the future nanoscale CMOS require very high body doping concentration, probably above  $5 \times 10^{18}$  cm<sup>-3</sup> for sub-50 nm MOSFET. At these doping levels direct tunneling leakage current from body to drain is expected to become a serious problem [4.9]. The increased parasitic junction capacitance ( $C_J$ ) and degraded channel carrier mobility can deteriorate the circuit performance. Furthermore, the fluctuations of device properties can be induced due to the random fluctuation of the placement and number of the dopant atoms [4.10]; this may limit the number of devices which can be integrated on a single chip,

because of the intolerable clock skews or malfunction of the circuits that depends on the matched or absolute values of device properties [4.9].

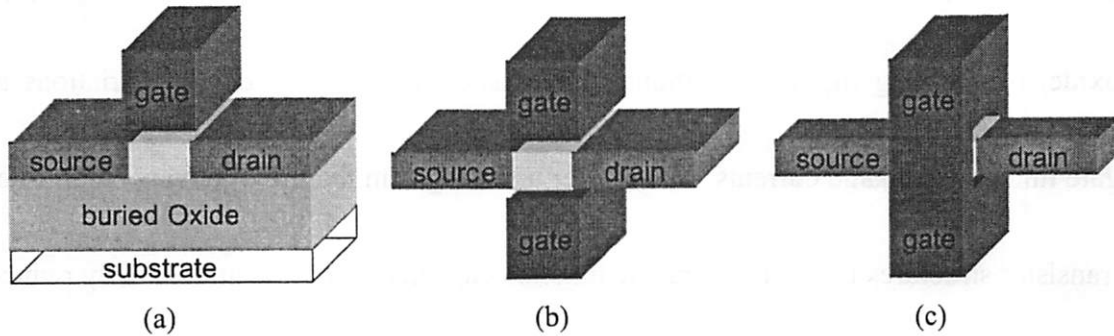


Figure 4.2 Advanced thin-body transistor structures; (a) ultra-thin body single gate (UTB) MOSFET, (b) double gate (DG) MOSFET, and (c) multiple gate (MG) MOSFET.

Alternative way to suppress the short channel effects without degrading the circuit performance is adopting the advanced thin-body transistor structures such as ultra-thin body single gate (UTB) MOSFET, double gate (DG) MOSFET, and multiple gate (MG) MOSFET, as shown in **Figure 4.2**; the use of an ultra-thin undoped silicon channel can effectively eliminate the off-state leakage current ( $I_{off}$ ) between source and drain. **Figure 4.3** illustrates the elimination of the silicon channel region whose potential is not well controlled by the gate electrode effectively suppresses the off-state leakage current; the buried oxide (BOX) prevents the channel potential from the drain field. The device parameters used for the 2-D device simulations [4.11] are 35 nm gate length ( $L_G$ ), 2.0 nm

gate oxide thickness ( $T_{OX}$ ),  $1 \times 10^{15} \text{ cm}^{-3}$  channel doping concentration ( $N_A$ ), 5 nm thick silicon body ( $T_{Si}$ ) with 1.2 V operation voltage ( $V_{DD}$ ). It should be noted that  $I_{Off}$  was quite well suppressed without using a heavy body doping and state-of-the-art thinner gate oxide, overcoming the aforementioned issues such as statistical dopant variations and gate tunneling leakage currents. The other advantages in the use of advanced thin-body transistor structures is that the parasitic junction capacitance ( $C_J$ ) is dramatically reduced, and the source/drain junction depth ( $X_J$ ) is naturally determined by the silicon body thickness ( $T_{Si}$ ).

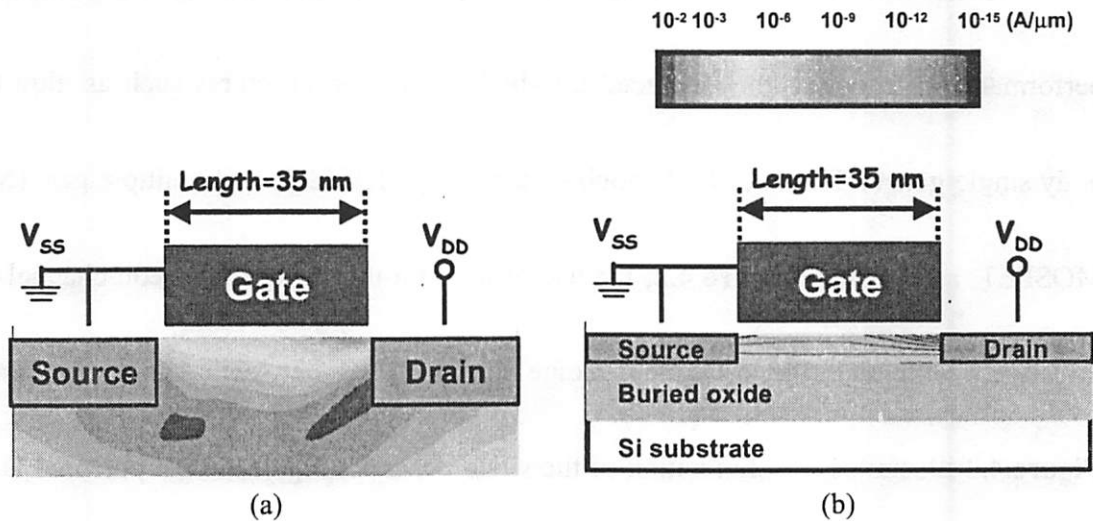


Figure 4.3 Comparison of leakage current density for (a) conventional MOSFET, and (b) UTB MOSFET ( $V_{GS}=0\text{V}$ ,  $V_{DS}=1.2\text{V}$ ,  $L_G=35 \text{ nm}$ ,  $T_{ox}=2.0 \text{ nm}$ , and  $N_A=10^{15} \text{ cm}^{-3}$ ).

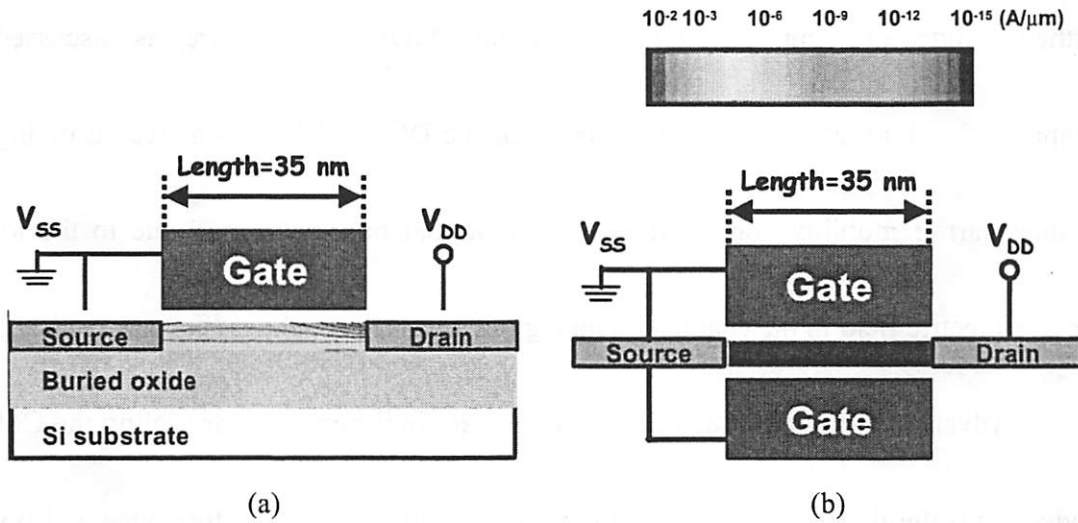


Figure 4.4 Comparison of leakage current density for (a) UTB MOSFET, and (b) double gate (DG) MOSFET ( $V_{GS}=0\text{V}$ ,  $V_{DS}=1.2\text{V}$ ,  $L_G=35\text{ nm}$ ,  $T_{Si}=5\text{ nm}$ ,  $T_{ox}=2.0\text{ nm}$ , and  $N_A=10^{15}\text{ cm}^{-3}$ ).

The double gate (DG) and multiple gate (MG) MOSFET has electrostatically much more robust control over the channel potential than the single-gated MOSFET, since two (or multiple) gates shields the channel potential from the drain field, suppressing the short channel effects, as shown in **Figure 4.4**. It is worth noting that the substrate with heavy channel doping for conventional MOSFET plays the identical role of the bottom shield, which results in a tradeoff between the degree of shielding and the reduction of the subthreshold swing [4.9]. However, such tradeoff does not exist for DG (or MG) MOSFETs, allowing the scalability of gate length down below sub-35 nm CMOS technology node [4.12]; DG MOSFET structures enable approximately 2 times

further scaling, as compared to the single gate MOSFET structure, as discussed in Chapter 4.2. Furthermore, the performance of the DG MOSFET is increased by higher channel carrier mobility compared to a conventional bulk MOSFET due to the lower average electric field in the channel, reducing the interface roughness scattering [4.13]

Advanced thin-body transistor structures are indispensable for scaling the CMOS devices into the deep nanometer regime without degrading the performance and power consumption. However, in order to apply for nanoscale CMOS devices, the unique device parameters needs to be designed adequately, and the process integration challenges should be overcome.

## **4.2 Design Methodology and Challenges for Advanced Transistor Structures.**

The most straightforward implementation of the advanced thin-body transistor structures is the UTB MOSFET, in which the device is fabricated in the silicon film of a silicon-on-insulator (SOI) wafer. The body thickness ( $T_{Si}$ ) is determined by the thickness of the SOI layer. **Figure 4.5** shows the impact of body thickness on the subthreshold characteristics of UTB MOSFET. It can be seen that the off-state leakage



current decreased by 3 orders-of-magnitude and the subthreshold swing is improved by 25 mV/dec through the reduction in body thickness from 7 nm to 3 nm for 18 nm gate length. The subthreshold swing of the UTB MOSFET with 3 nm thick body shows an almost ideal value (*i.e.*, 60 mV/dec at room temperature), despite of any heavy body doping concentration. It is worth noting that the behavior of fully depleted (FD) SOI device ( $T_{Si} = 20$  nm) is similar to that of a resistor due to the short channel effects.

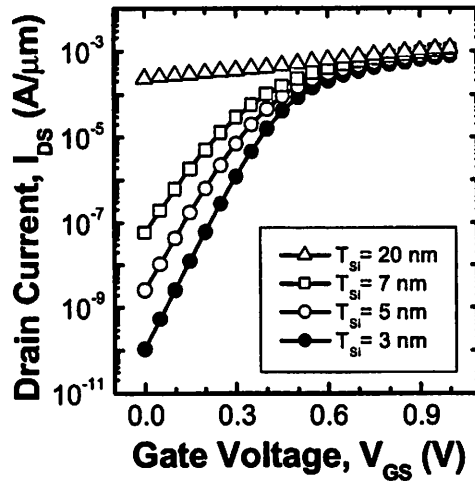


Figure 4.5 Impact of body thickness on the subthreshold characteristics of UTB MOSFET ( $L_G=18$  nm,  $T_{ox}=1.5$  nm, and  $N_A = 10^{15}$  cm<sup>-3</sup>).

Figure 4.6 shows the simulation results of drain induced barrier lowering (DIBL) as a function of the ratio of gate length to body thickness ( $\zeta = L_G/T_{Si}$ ). It can be seen that the ratio of gate length to body thickness ( $\zeta$ ) should be properly designed for a given

technology node and device structure; while  $\zeta = 4$  is necessary for UTB MOSFETs to suppress the short channel effects, the required  $\zeta$  is reduced to 2 for DG MOSFETs due to the stronger control over the channel potential. For example, for 40 nm gate length,  $T_{Si} < 10$  nm is needed for UTB MOSFETs, while  $T_{Si} < 20$  nm for DG MOSFETs.

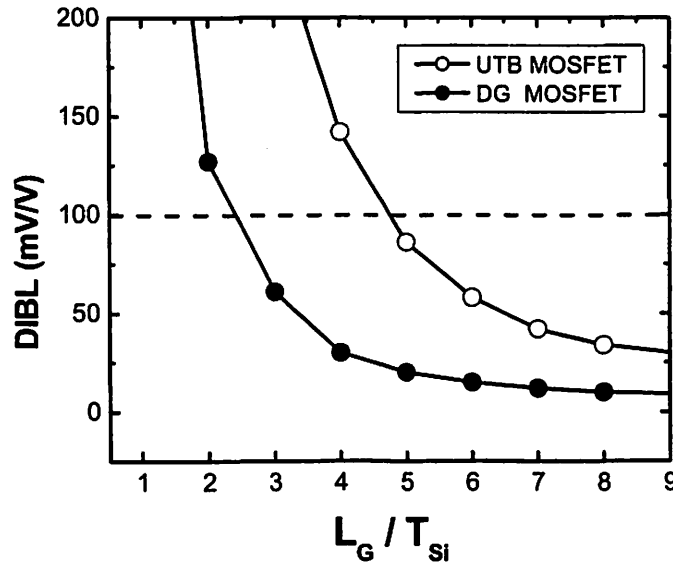
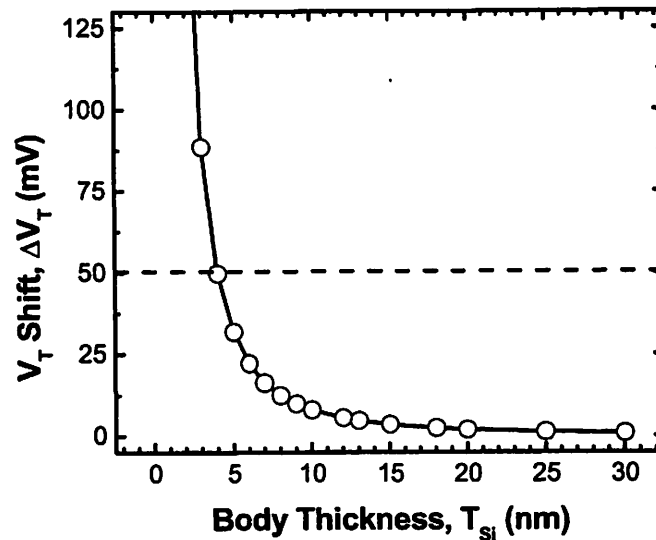


Figure 4.6 Simulation results of drain induced barrier lowering (DIBL) as a function of the ratio of gate length to body thickness ( $\zeta = L_g/T_{Si}$ ) for UTB and DG MOSFETs.

Another big benefit from DG MOSFETs is alleviating the stringent body thickness requirement, allowing approximately 2 times further scaling of the MOSFET with a given silicon body thickness. It should be noted that when the silicon body becomes thinner than 10 nm, threshold voltage ( $V_T$ ) varies because quantum mechanical (QM)

shift of the ground state energy is inversely proportional to the square of the body thickness. The uncertainty of the threshold voltage ( $\sigma V_T$ ), to the first order, can be expressed by  $\Delta V_T = (h^2\pi^2/qm^*)(1/T_{Si}^2)(\Delta T_{Si}/T_{Si})$  [4.14], as shown in **Figure 4.7**. Numerical simulation and experimental studies show that  $V_T$  shifts rapidly as the silicon body thickness scales thinner than 5 nm [4.15, 4.16, 4.17].



**Figure 4.7** Impact of the body thickness on threshold voltage shift for advanced thin-body transistor structures [4.14].

There are three basic configurations to implement DG MOSFET, as illustrated in **Figure 4.8** [4.18]. In a type I configuration, the silicon body thickness can be easily controlled uniformly compared to the other two configurations, which is the most critical

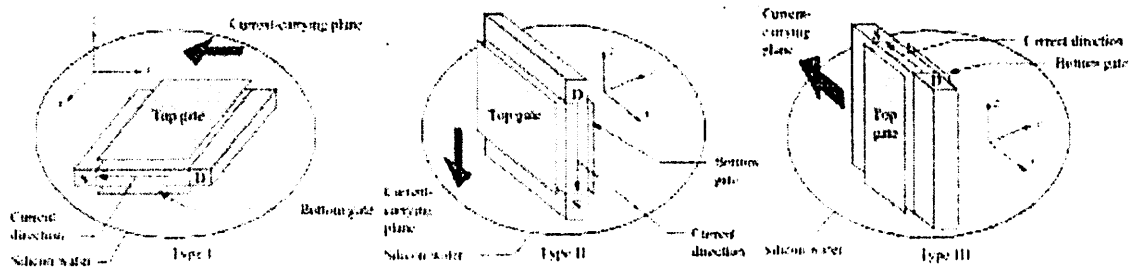


Figure 4.8 Three basic configurations of the double gate (DG) MOSFET [4.18].

dimension in DG MOSFET. However, complexity in process integration and difficulty in self-aligning the top and bottom gate without introducing substantial parasitic resistance and capacitance should be overcome; Figure 4.9 shows the scanning electron microscopy (SEM) image of a misaligned DG MOSFET [4.19]. Simulation study predicted that 25 % misalignment between top and bottom gate induced 33 % increase in delay [4.20]. Type II configuration has the better control of the gate length uniformly, and provides the most compact layout for DRAM applications. However, there are challenges in avoiding a large parasitic gate-to-source/drain overlap capacitance, obtaining more than one gate length, and connecting individual devices without sacrificing the layout area for CMOS logic applications [4.21, 4.22]. Type III configuration, the *FinFET*, has the highest packing density and allows easy access to all four electrodes with the self-alignment of top and bottom gate. However, there are

challenges in obtaining the critical dimension of body thickness and minimizing the plasma etching damage of the channel. It is worthwhile to note that *spacer lithography* process has been demonstrated to pattern the body thickness down to 30 nm with a good uniformity, which is by far below the today's lithographical limit [4.23], and *hydrogen annealing* process has shown to provide high surface quality on etched silicon surface [4.24].

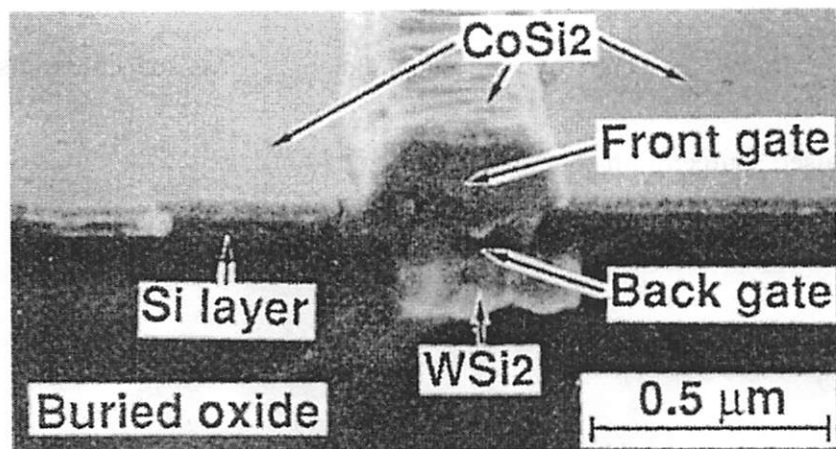


Figure 4.9 Cross-sectional scanning electron microscopy (SEM) image of a misaligned double gate (DG) MOSFET (Type I configuration) [4.19].

For advanced transistor structures regardless of the configurations, there are two challenges resulted from the nature of ultra-thin body thickness; parasitic resistance and threshold voltage adjustment. The parasitic resistance due to the ultra-thin body

thickness should be reduced using thick source/drain fanout structures self-aligned to gate electrode in order to fully achieve the aforementioned performance improvements [4.25, 4.26, 4.27], but this results in layout area penalty. By using selective deposition, a raised source/drain structure can be created to reduce the parasitic resistance as shown in **Figure 4.10**. Several methods have been demonstrated, including selective silicon (Si) epitaxial growth [4.26], selective silicon-germanium (SiGe) epitaxial growth [4.28], and selective germanium (Ge) deposition [4.29]. One drawback is, however, that it inevitably results in increased overlap capacitance. Thus, spacer widths between the gates and raised source/drain should be optimized [4.30].

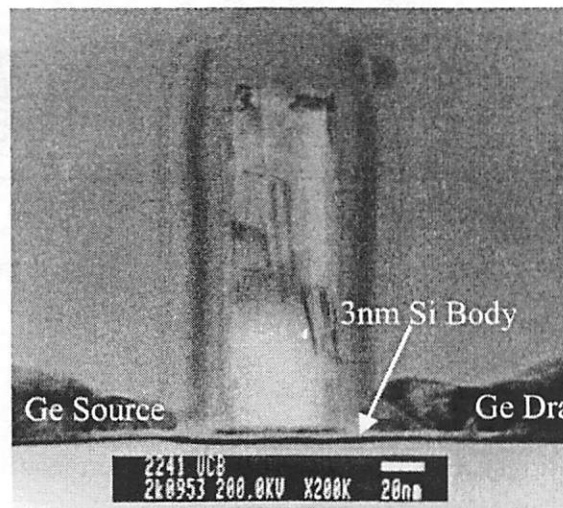


Figure 4.10 Cross-sectional transmission electron microscopy (X-TEM) image of selectively deposited raised germanium (Ge) source/drain ultra-thin body (UTB) MOSFET [4.30].

Threshold voltage adjustment should be achieved by gate work function engineering due to the lack of channel body dopants in advanced thin-body transistor structures. Even though the use of heavy channel doping ( $> 5 \times 10^{18} \text{ cm}^{-3}$ ) is an alternative way to control the threshold voltage, this reintroduces problems with reduced carrier mobility and random dopant fluctuations [4.31]. An asymmetric DG MOSFET structure can also achieve an appropriate threshold voltage by using conventional  $n^+$  and

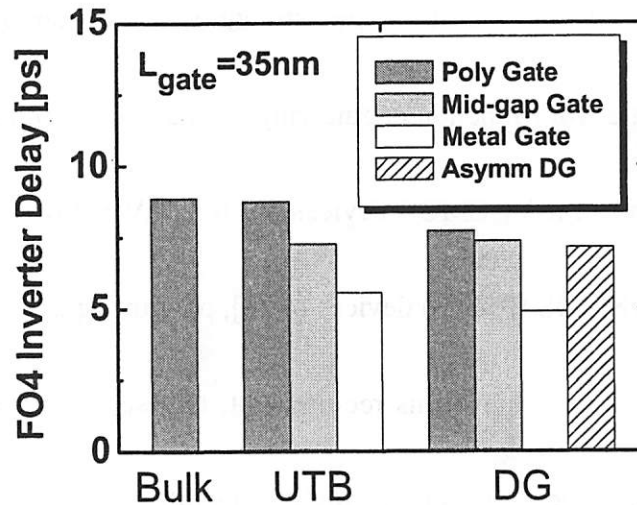


Figure 4.11 The impact of methods for controlling threshold voltage on the fanout of four (FO4) inverter delay. Note that all the devices have the identical  $V_T$  values [4.32].

$p^+$  poly-Si gates at the cost of increased transverse electric field [4.19]. As a result, both channel doping and an asymmetric DG MOSFET structure result in degraded device performance as shown in **Figure 4.11** [4.32]. Therefore, the need for gate work

function engineering is emphasized as a unique solution for the performance benefits and reliable-operations of the advanced thin-body transistor structures. The theoretical data for threshold voltage as a function of gate work function is shown in Figure 2.3

## **4.3 Tunable Work Function Mo-Gated SiO<sub>2</sub> UTB MOSFET**

For advanced thin-body transistor structures, it is desirable to adjust threshold voltage through gate work function engineering. The gate work function values needed for low and symmetrical  $V_T$  CMOS devices are 0.35 eV below (above) the conduction (valence) band for NMOS (PMOS) devices [4.33], precluding the use of doped poly-Si as the gate electrode. To achieve this requirement, two separate metals with appropriate work functions are necessary on a single substrate. However, this approach has difficulties in avoiding the process complexity and in providing the multiple  $V_T$  values for low voltage and high performance applications; that is, low  $V_T$  devices are adopted to reduce the delay in the critical paths, while high  $V_T$  devices in minimizing the standby leakage power [4.34]. Thus, it is preferable to deposit a single gate material and subsequently adjust its work function selectively. Molybdenum (Mo) is a candidate



gate material for future fully depleted (FD) SOI CMOS technology, because it is compatible with a standard CMOS process flow and its work function can be adjusted within the desired range (4.5-5.0 eV) via nitrogen implantation as discussed in Chapter 2.5 [4.24, 4.35, 4.36]. In this section, the tunability of the Mo gate work function on SiO<sub>2</sub> is demonstrated for the first time using p-channel UTB MOSFETs [4.37].

#### **4.3.1 Fabrication of Mo-gated SiO<sub>2</sub> p-channel UTB MOSFETs**

The process sequence used to fabricate p-channel UTB MOSFETs is depicted in **Figure 4.12**. The starting substrates were Unibond<sup>®</sup> SOI wafers, with 100 nm thick lightly doped ( $\rho = 10 - 15 \Omega\text{-cm}$ ) p-type (100) silicon on 400 nm thick buried oxide. The silicon film was thinned down to ~15 nm by thermal oxidation. The body thickness was chosen not to be too thin, to avoid threshold voltage ( $V_T$ ) shifts due to quantum confinement [4.15, 4.16, 4.17, 4.38]. Isolation is simply achieved by patterning the silicon into active-area islands without the need for further dielectric deposition and planarization processes because of the small step height (< 15 nm). After active-area patterning, a 3.0 nm thick SiO<sub>2</sub> gate dielectric was thermally grown, and a 40 nm thick Mo film was deposited at 200°C using a DC magnetron sputtering system (10<sup>-7</sup> Torr base

pressure). It should be noted that *plasma charge trap* (PCT), as discussed in Chapter 2.3, was not applied, thus sputtering damage may be introduced to the gate dielectric during the sputtering process. Then, nitrogen implantation was performed at various doses for  $V_T$  adjustment (see **Table 4.1**). Afterwards, *in-situ* phosphorus doped poly-Si and a hard mask low temperature oxide (LTO) were sequentially deposited. The purpose of the hard mask was to prevent shorting of the source/drain (S/D) with gate during the subsequent selective germanium (Ge) deposition to form raised S/D regions.

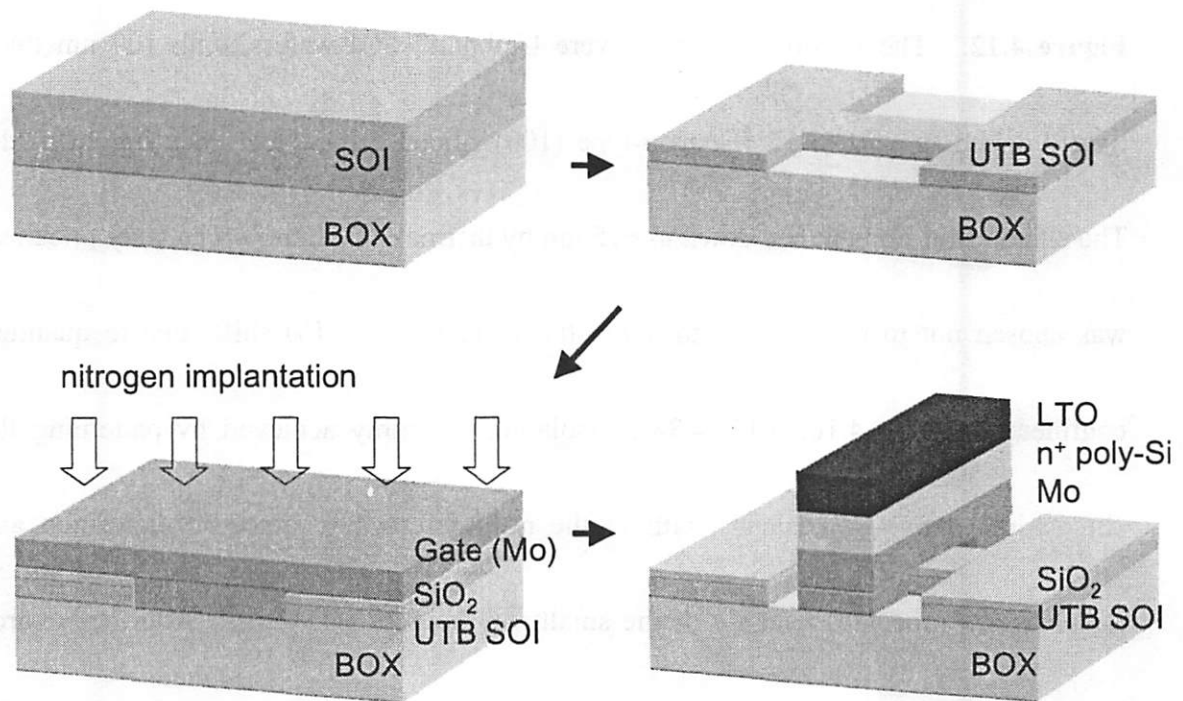


Figure 4.12 The fabrication sequences used for p-channel ultra-thin body (UTB) MOSFET.

A highly selective dry etch process, as discussed in Chapter 2.4, was adopted to etch the Mo gate on thin gate oxide. After gate-sidewall spacer formation, Ge was selectively deposited by low-pressure chemical vapor deposition (LPCVD), and S/D dopants were implanted. Device fabrication was completed with rapid thermal annealing (RTA) at 700°C, 60s in N<sub>2</sub> ambient to activate the implanted S/D dopants. This rather conservative anneal was used because Ge and Si intermixing is known to be enhanced in the presence of dopants [4.39], for the formation of abrupt p-n junctions with low thermal budget. **Figure 4.13** shows a cross-sectional transmission electron microscopy (X-TEM) of the completed p-channel UTB MOSFET.

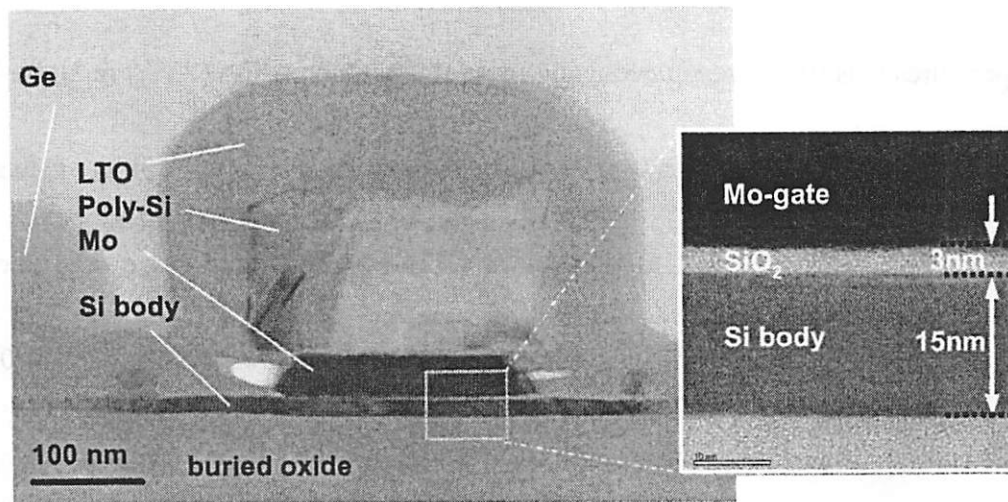


Figure 4.13 A cross-sectional transmission electron microscopy (X-TEM) of an ultra-thin (15 nm) silicon body Mo-gated MOSFET ( $T_{OX} = 3$  nm,  $L_G=290$  nm). The notched gate profile resulted from the unintended wet cleaning process.

Table 4.1 Summary of the process split conditions.

	Substrate	Channel doping ( $N_A$ )	Gate Stack	$^{14}\text{N}^+$ implant condition
WF #1	Bulk Si	$< 10^{15} \text{ cm}^{-3}$	Mo/SiO <sub>2</sub>	–
WF #2	Bulk Si	$< 10^{15} \text{ cm}^{-3}$	Mo/SiO <sub>2</sub>	20 KeV, $2 \times 10^{15} \text{ cm}^{-2}$
WF #3	Bulk Si	$< 10^{15} \text{ cm}^{-3}$	Mo/SiO <sub>2</sub>	20 KeV, $4 \times 10^{15} \text{ cm}^{-2}$
WF #4	SOI	$\sim 10^{15} \text{ cm}^{-3}$	Mo/SiO <sub>2</sub>	–
WF #5	SOI	$\sim 10^{15} \text{ cm}^{-3}$	Mo/SiO <sub>2</sub>	20 KeV, $2 \times 10^{15} \text{ cm}^{-2}$
WF #6	SOI	$\sim 10^{15} \text{ cm}^{-3}$	Mo/SiO <sub>2</sub>	20 KeV, $4 \times 10^{15} \text{ cm}^{-2}$

#### 4.3.2 Results and Discussion of Mo-gated SiO<sub>2</sub> p-channel UTB MOSFETs

The thermal stability of Mo in contact with SiO<sub>2</sub> gate dielectric at various temperatures was first investigated using rapid thermal annealing (RTA) in N<sub>2</sub> for bulk-Si MOS capacitors. It is worth noting that all the fabrication steps to form the gate stack of these wafers were processed together with UTB MOSFETs. Figure 4.14 shows the measured high-frequency C-V curves. For annealing temperatures up to 900°C, the accumulation capacitance does not change at all. The capacitance decreases after a 1000°C anneal, however, due to the reaction of Mo with SiO<sub>2</sub>. These results indicate

that Mo gate is thermodynamically stable on SiO<sub>2</sub> at temperatures up to 900°, which is sufficient for S/D dopant activation in selective Ge raised S/D UTB MOSFET structures.

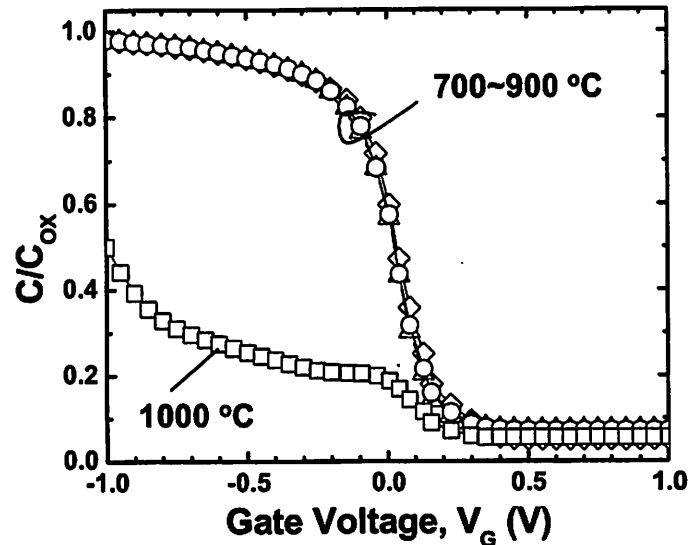


Figure 4.14 Measured high frequency (100 KHz) C-V curves of Mo-gated SiO<sub>2</sub> bulk-Si MOS capacitors annealed in N<sub>2</sub> ambient for 1 min. at various temperatures ( $T_{ox} = 3.0$  nm, Area =  $50 \times 50 \mu\text{m}^2$ ).

Figure 4.15 shows the measured high-frequency C-V curves for p-channel UTB MOSFETs implanted with various doses of nitrogen. It can be seen that the threshold voltage shift to more negative values with increasing nitrogen implant dose. The measurement set-up can be shown in the inset of Figure 4.15 [4.40]; when the channel is inverted, the measured capacitance corresponds to the gate dielectric capacitance. It is

worthwhile to note that the inversion capacitance does not decrease with increasingly negative gate voltage, *i.e.* there is no poly-Si depletion effect (PDE) due to the Mo gate.

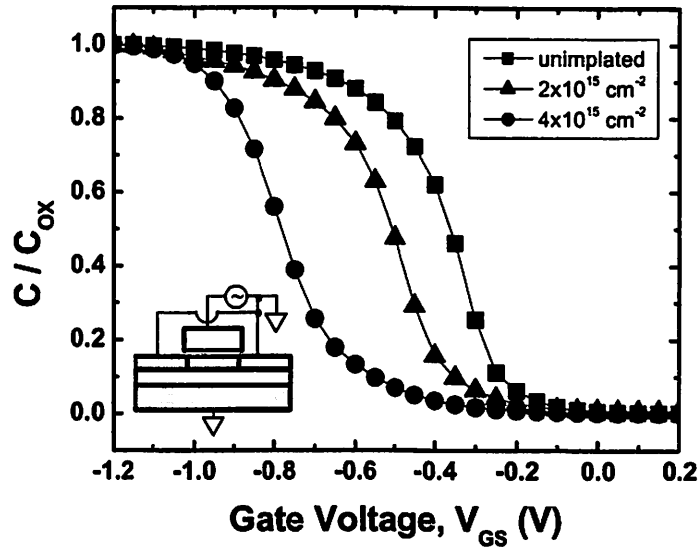


Figure 4.15 Measured high frequency (100 KHz) C-V curves of Mo-gated SiO<sub>2</sub> p-channel UTB MOSFETs implanted with various doses of nitrogen.

$V_T$  values for the Mo-gated p-channel UTB MOSFETs were extracted using the maximum transconductance ( $g_{m,max}$ ) method, as shown in Figure 4.16. The  $V_T$  for unimplanted Mo-gate device is  $-0.2$  V, and shifts by approximately  $-65\text{mV}$  for every  $1 \times 10^{15} \text{ cm}^{-2}$  increment in nitrogen implant dose. Thus, the threshold voltage for the UTB MOSFETs can be adjusted via Mo-gate work function engineering. In order to achieve

the desired  $V_T$  (0.2 ~ 0.3 V) in n-channel UTB MOSFETs, an estimated dose of  $6 - 8 \times 10^{15} \text{ cm}^{-2}$  would be needed.

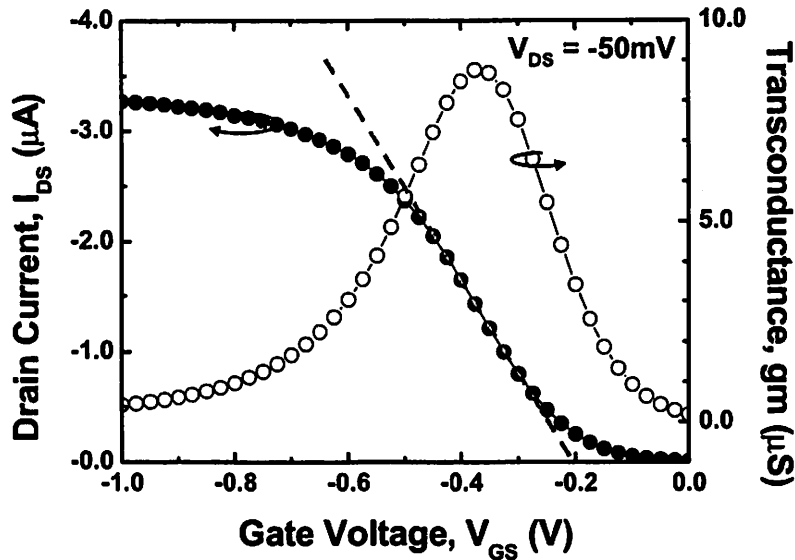


Figure 4.16 Maximum transconductance method is used to extract the  $V_T$  of Mo-gated  $\text{SiO}_2$  p-channel UTB MOSFET. Data is shown for  $L_G = 300\text{nm}$ ,  $T_{\text{Si}} = 15\text{nm}$ , and unimplanted Mo-gate.

Figure 4.17 shows the  $I_{\text{DS}}-V_{\text{GS}}$  characteristics for p-channel UTB MOSFETs.

The subthreshold slope is somewhat high (106.9 mV/dec), considering that the body thickness ( $T_{\text{Si}}$ ) is more than 4 times smaller than the gate length ( $L_G$ ), i.e.  $\zeta = 20$ , so that the subthreshold slope should be nearly ideal ( $\sim 60 \text{ mV/dec}$ ) as discussed in Chapter 4.2.

This may be due to the damage incurred during Mo film sputtering, which can be avoided

by either using a chemical vapor deposition (CVD) method or sputtering with *plasma charge trap* (PCT) as discussed in Chapter 2.3. It should be noted that since the Mo film has the columnar crystal structure, low-energy and large tilted-angle implantation can prevent  $^{14}\text{N}^+$  implant damage with reduced channeling effect [4.41].

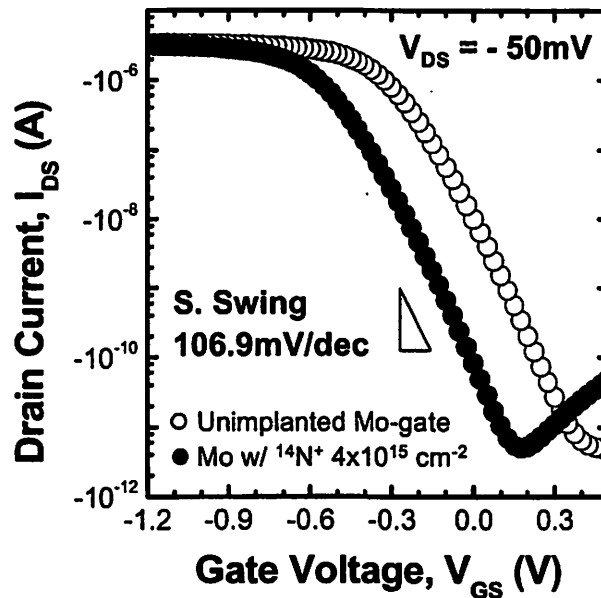


Figure 4.17 Measured  $I_{DS}$ - $V_{GS}$  characteristics for Mo-gated  $\text{SiO}_2$  p-channel UTB MOSFETs.

**Figure 4.18** shows the measured drain induced barrier lowering (DIBL) for p-channel UTB MOSFETs. Despite of the lightly doped body ( $10^{15} \text{ cm}^{-3}$ ), DIBL is greatly suppressed less than 40 mV/V for less than 300 nm gate length due to the ultra-thin body thickness (15 nm), hence  $\zeta > 4$ . It should be noted that there is no tradeoff



between the degree of drain field shielding, *i.e.*, short channel effects and the threshold voltages, *i.e.*, channel doping concentration in UTB MOSFETs.

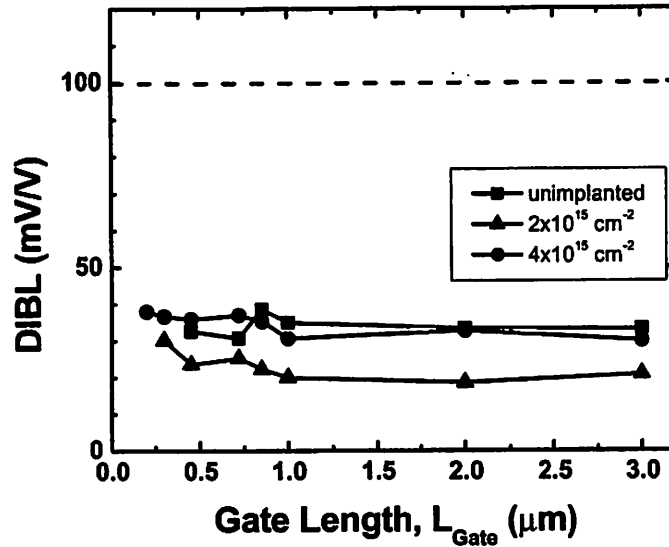


Figure 4.18 Measured drain induced barrier lowering (DIBL) as a function of the gate length of Mo-gated  $\text{SiO}_2$  p-channel UTB MOSFETs ( $T_{\text{Si}} = 15 \text{ nm}$ , and  $T_{\text{Ox}} = 3.0 \text{ nm}$ ).

Figure 4.19 shows the measured  $I_{\text{DS}}-V_{\text{DS}}$  characteristics for the p-channel UTB MOSFETs. The channel length modulation is greatly reduced because UTB MOSFETs can effectively shield the drain field from affecting the channel potential. The low drain current is mainly due to the insufficient overlap of source/drain with gate, which is caused by the unintended lateral etching of Mo film during the cleaning process (see Figure 4.13).

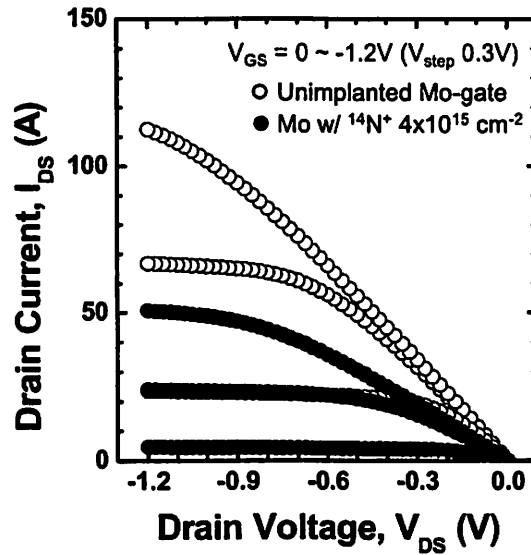


Figure 4.19 Measured  $I_{DS}$ - $V_{DS}$  characteristics of Mo-gated  $\text{SiO}_2$  p-channel UTB MOSFETs ( $L_G = 300$  nm,  $W = 10$   $\mu\text{m}$ ,  $T_{\text{Si}} = 15$  nm, and  $T_{\text{OX}} = 3.0$  nm).

## 4.4 Tunable Work Function Mo-Gated $\text{HfO}_2$ CMOS FinFETs

The double-gate (DG) FinFET is one of the most promising transistor structures for scaling CMOS technology to sub-25 nm gate lengths. Integration of metal gate and high- $k$  gate dielectric is desirable to reduce the equivalent gate-oxide thickness (EOT) while maintaining low gate leakage current, to improve transistor drive current and to relax the fin-width requirement for controlling short-channel effects [4.42, 4.43, 4.44].

In order to maximize carrier mobilities and avoid statistical dopant fluctuation effects on threshold voltage ( $V_T$ ), the body of a FinFET should be undoped;  $V_T$  adjustment must be achieved by tuning the gate work function, in this case. Molybdenum (Mo) is a candidate gate material for future FD-SOI CMOS technology, because it is compatible with a standard CMOS process flow and its work function can be adjusted within the desired range (4.5-5.0 eV) via nitrogen implantation [4.24, 4.35, 4.36]. In this section, the tunability of the Mo gate work function on  $HfO_2$  is demonstrated for the first time using FinFETs [4.45].

#### **4.4.1 Fabrication of Mo-gated $HfO_2$ CMOS FinFETs**

Figure 4.20 outlines the FinFET process flow. UNIBOND<sup>®</sup> wafers were used as the starting substrates, and oxidized to reduce the SOI thickness to 50 nm. 80-nm wide Si fins were then defined by *spacer lithography*, while the source/drain (S/D) contact regions were defined by *i-line photolithography* [4.23]. After patterning of the SOI and  $NH_3$  pretreatment of the Si fin (110) sidewall surfaces,  $HfO_2$  was deposited by CVD using Hf t-butoxide ( $Hf(OC(CH_3)_3)_4$ ) as a precursor at 500°C. A 60 nm thick

molybdenum (Mo) film was then deposited by DC magnetron sputtering with a *plasma charge trap* (PCT) to minimize sputtering damage, as described in Chapter 2.3.

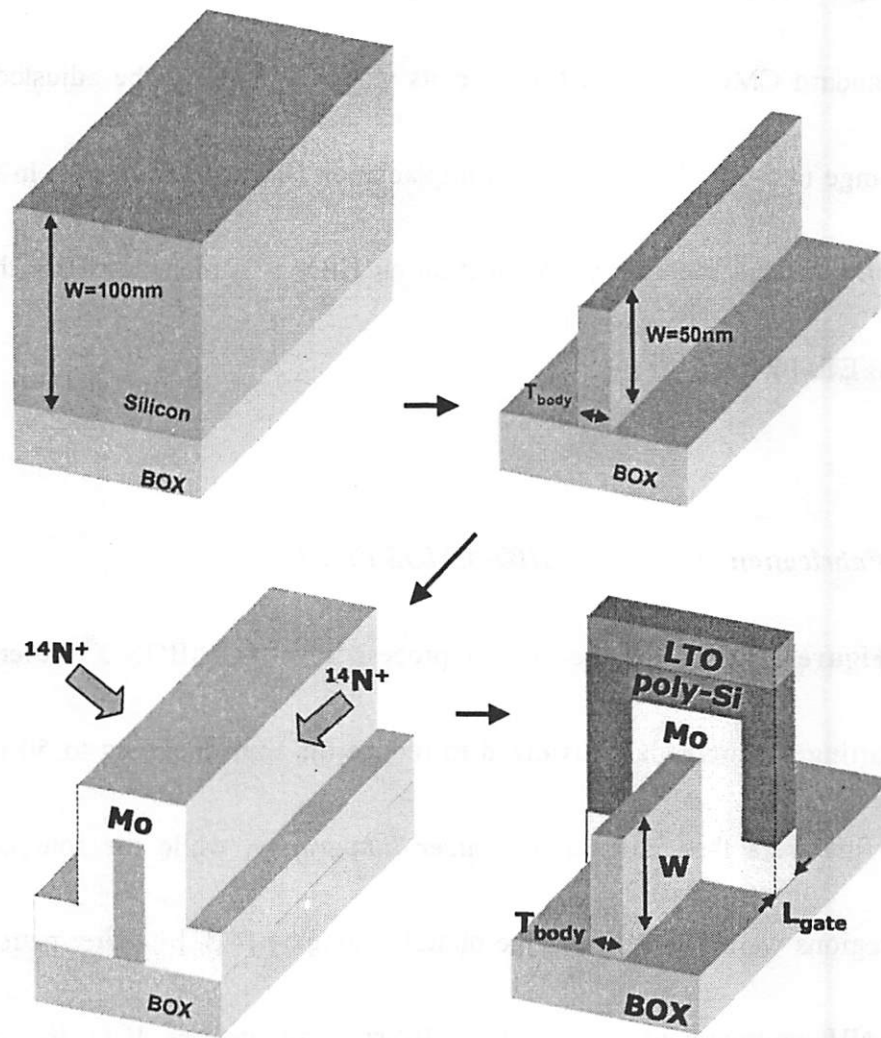


Figure 4.20 The fabrication sequences used for the Mo-gated  $\text{HfO}_2$  CMOS FinFETs

For some n-channel FinFETs, nitrogen ( $1 \times 10^{16} \text{ cm}^{-2}$ , 5 keV) was implanted into the Mo gate film at  $30^\circ$  tilt on each side of the Si fins (*i.e.*,  $60^\circ$  tilt on the normal to the Mo gate surface) in order to reduce the effective Mo work function and achieve low  $V_T$ . The use of a low-energy tilted implant prevents nitrogen penetration into the underlying gate dielectric as shown in Figure 4.21. The Mo was capped with *in-situ*  $n^+$  doped poly-Si and planarized by chemical mechanical polishing (CMP). After gate patterning and source/drain ion implantation, a  $900^\circ\text{C}$  60s RTA in  $\text{N}_2$  was used to activate the dopants. Finally, the devices were annealed at  $400^\circ\text{C}$  in forming gas.

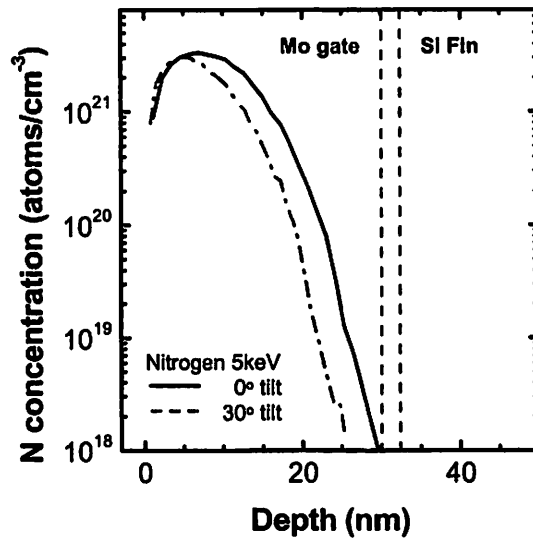


Figure 4.21 *SRIM*<sup>®</sup> simulation results for the nitrogen implantation profiles in the Mo/HfO<sub>2</sub> gate stack (dose= $5 \times 10^{15} \text{ cm}^{-2}$ ).

Figures 4.22 are tilted-view scanning electron microscopy (SEM) images of the completed Mo/HfO<sub>2</sub> FinFETs. Neither micro-trenching nor Mo gate residue was observed due to the high etching selectivity, as discussed in Chapter 2.4. It should be noted that no metallization or silicided S/D structure was used in this study. Figures 4.23 show cross-sectional transmission electron microscopy (TEM) image, and close-up view of the gate stack at the silicon fin sidewall. The undercut of buried oxide beneath the fin occurred during the sacrificial oxide removal using diluted HF. It should be noted that this undercut may introduce the yield issues, but shield the channel potential from drain field [4.46]. It is worth noting that the Mo gate layer is continuous at the bottom of the Si fin due to the improved step-coverage with PCT sputtering [4.47].

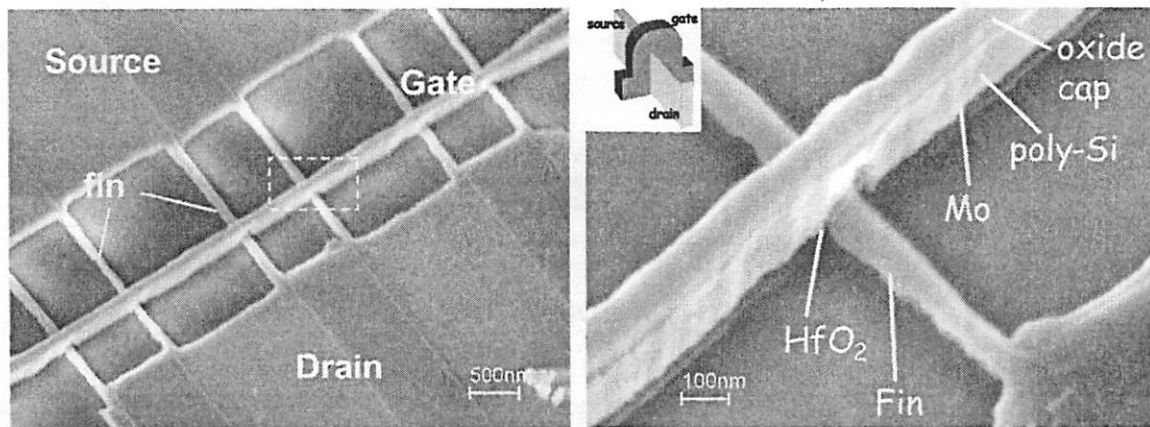


Figure 4.22 Tilted-view scanning electron microscopy (SEM) images of the Mo/HfO<sub>2</sub> FinFETs.

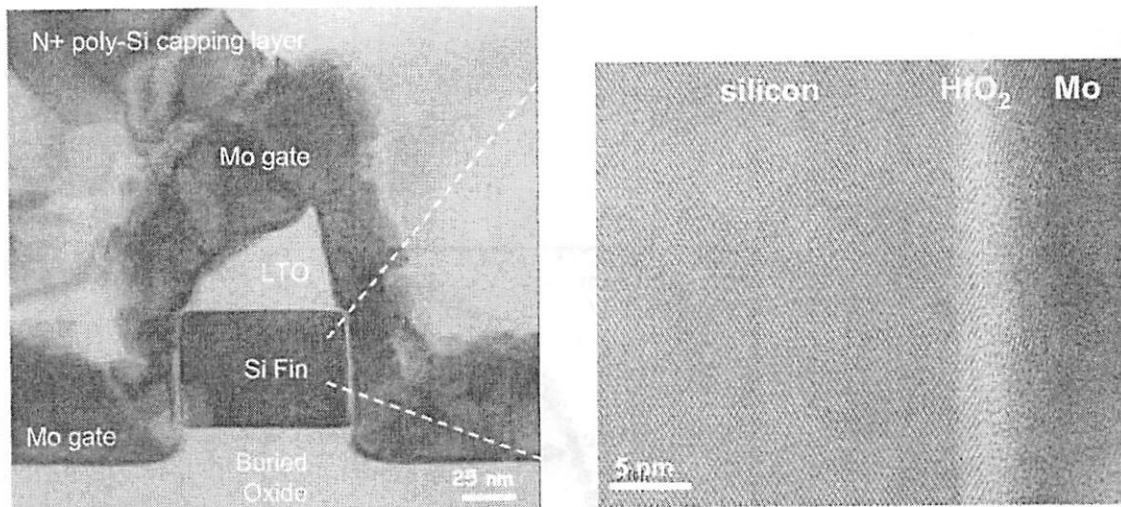


Figure 4.23 Cross-sectional transmission electron microscopy (X-TEM) images of the Mo/HfO<sub>2</sub> FinFETs.

#### 4.4.2 Results and Discussion of Mo-gated HfO<sub>2</sub> CMOS FinFETs

Figure 4.24 compares the measured  $I_{DS}$ - $V_{GS}$  characteristics for Mo-gate HfO<sub>2</sub> n-channel FinFETs with and without nitrogen gate implantation. It can be seen that  $V_T$  (defined as the gate voltage when  $I_{DS} = 100 \text{ nA}/\mu\text{m}$  for  $V_{DS} = 50 \text{ mV}$ ) is shifted from 0.73 V down to 0.28 V by the nitrogen implant. This indicates that the Mo-gate work function was effectively reduced by the nitrogen implant. Compared to the previous studies [4.24, 4.35, 4.36], the subthreshold swing is greatly improved due to the

minimization of Mo gate sputtering damage and the low-energy tilted nitrogen implantation [4.47].

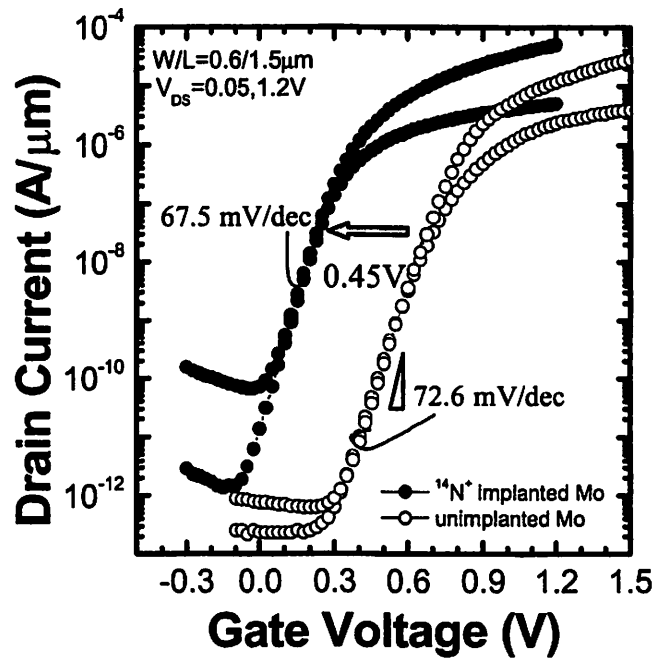


Figure 4.24 Measured  $I_{DS}$ - $V_{GS}$  characteristics of pure and nitrogen implanted Mo-gated  $HfO_2$  FinFETs. Nitrogen implantation into Mo-gate is effective for  $V_T$  control without degrading the subthreshold swing.

The amount of reduction in gate work function is proportional to the nitrogen implant dose as shown in Figure 4.25, but it is less for Mo on  $HfO_2$  than for Mo on  $SiO_2$  due to the *Fermi-level pinning effect* [4.6, 4.7, 4.48, 4.49] and nitrogen diffusion into the  $HfO_2$ . [4.45].



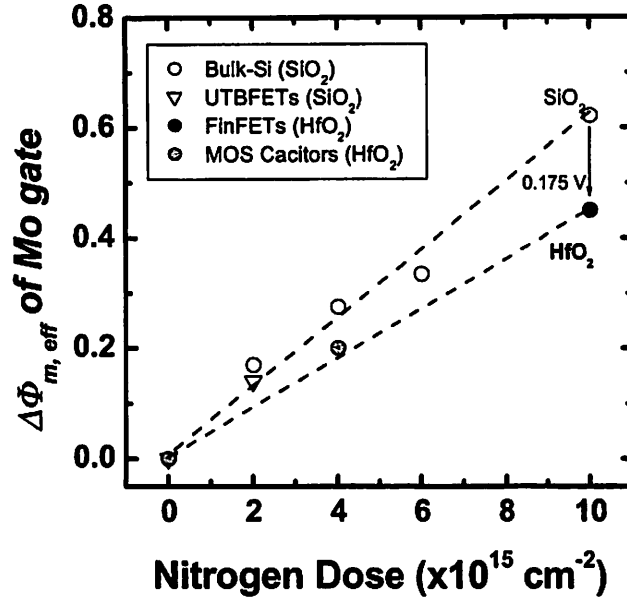
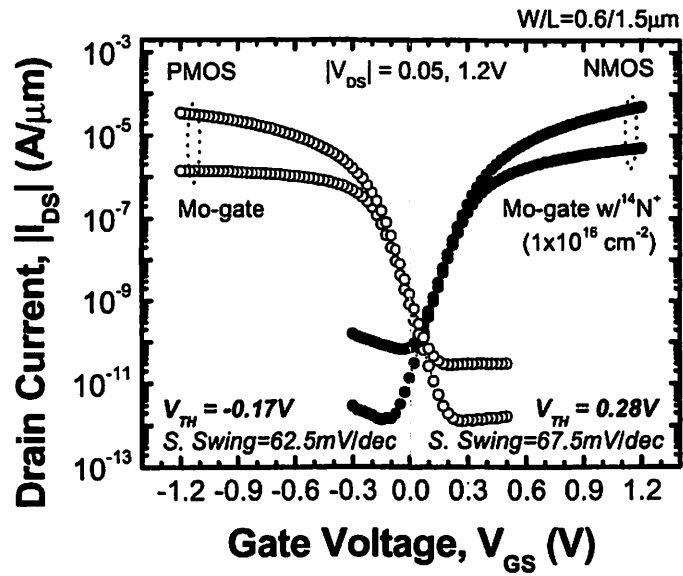
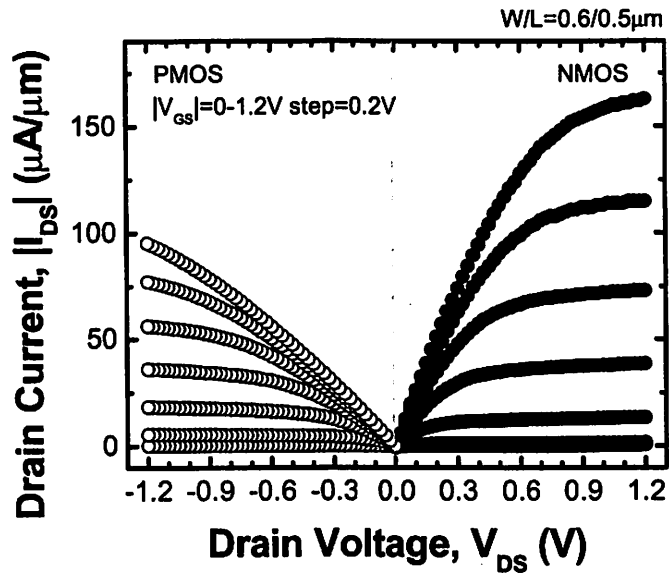


Figure 4.25 Change in effective Mo gate work function ( $\Delta\Phi_{m,eff}$ ) vs. nitrogen implantation dose.

Figure 4.26 shows measured  $I_{DS}-V_{GS}$  and  $I_{DS}-V_{DS}$  characteristics for n-channel (with nitrogen-implanted Mo gate) and p-channel (with pure Mo gate) FinFETs. The  $V_T$  and subthreshold swing values for the n-channel (p-channel) device are 0.28 (-0.17) V, and 67.5 (62.5) mV/dec, respectively. The p-channel device exhibits higher parasitic resistance than the n-channel device due to differences in p-type vs. n-type dopant redistribution in thin SOI during rapid thermal annealing [4.50].



(a)



(b)

Figure 4.26 Measured (a)  $I_{DS}$ - $V_{GS}$  and (b)  $I_{DS}$ - $V_{DS}$  characteristics of Mo-gated  $\text{HfO}_2$  CMOS FinFETs. Low energy (5 KeV) and 30-degree tilted nitrogen implantation has been applied to Mo-gate for n-channel FinFET.

**Figure 4.27** compares measured  $C_G$ - $V_G$  curves (circles) with quantum-mechanical simulations (lines) [4.51]. The measured multi-fin device structure is shown in the inset. Extracted equivalent oxide thickness (EOT) in inversion is 1.95 or 1.92 nm for n-channel or p-channel devices, respectively. Stretch-out in the measured C-V characteristic can be seen for the n-channel device, but not for the p-channel device.

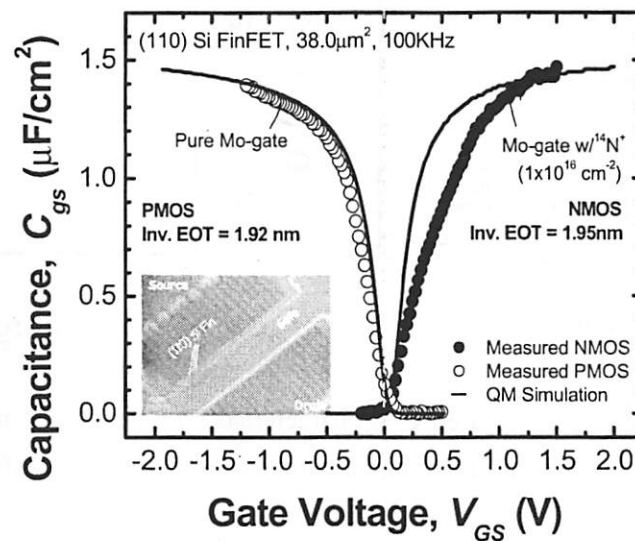


Figure 4.27 Comparison of measured C-V characteristics (circles) and quantum mechanical simulations (lines) for n-channel and p-channel Mo/HfO<sub>2</sub> FinFETs.

This result indicates an asymmetrical distribution of interface traps ( $D_{it}$ ) within the energy bandgap (*i.e.*, higher  $D_{it}$  above  $E_i$ ) [4.52], which explains the larger subthreshold swing of the n-channel FinFET as shown in Figure 4.26. As can be seen in **Figure 4.28**,

the deviation from the ideal C-V characteristic is larger for a nitrogen-implanted Mo gate, indicating a higher interfacial trap density. This is likely due to the formation of an interfacial  $\text{HfO}_x\text{N}_y$  layer at the Si surface [4.53].

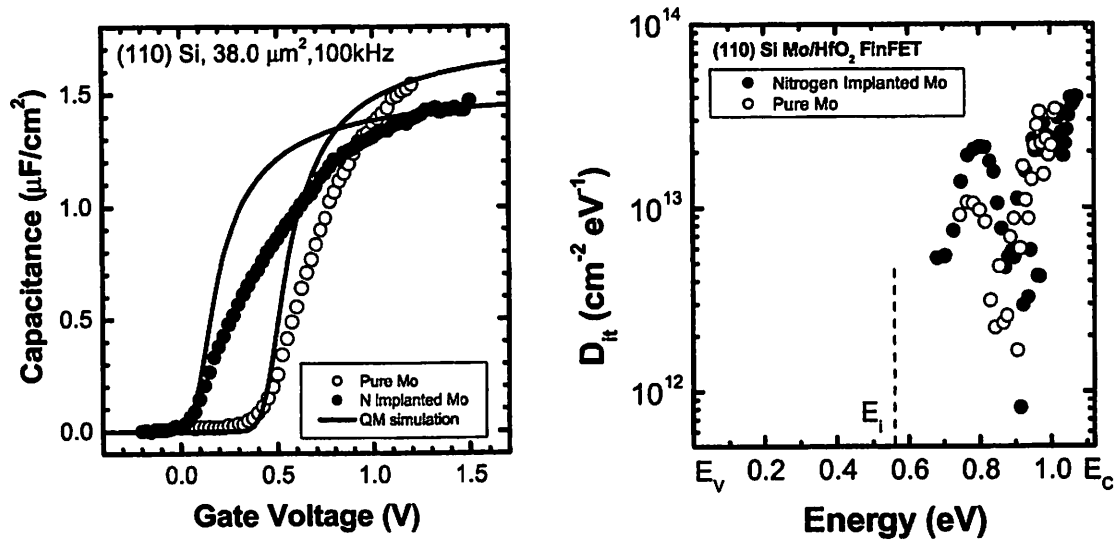


Figure 4.28 Comparison of measured C-V characteristics and extracted interface trap densities ( $D_{it}$ ) for pure and nitrogen implanted n-channel Mo/ $\text{HfO}_2$  FinFETs.

Figure 4.29 shows the field-effect electron mobility for (110) Si sidewall Mo-gated  $\text{HfO}_2$  FinFETs with (lines) and without (symbols) correcting interface trap density [4.54]. The apparent electron mobility is lower for the nitrogen-implanted device as compared to the unimplanted device, due to the higher interface trap density. Figure 4.30 shows the field-effect hole mobility for a (110) Si sidewall Mo-gated  $\text{HfO}_2$  FinFET.

Both the electron mobility and the hole mobility are significantly degraded compared to the universal mobility curves for a (110) Si surface with SiO<sub>2</sub> gate dielectric [4.55].

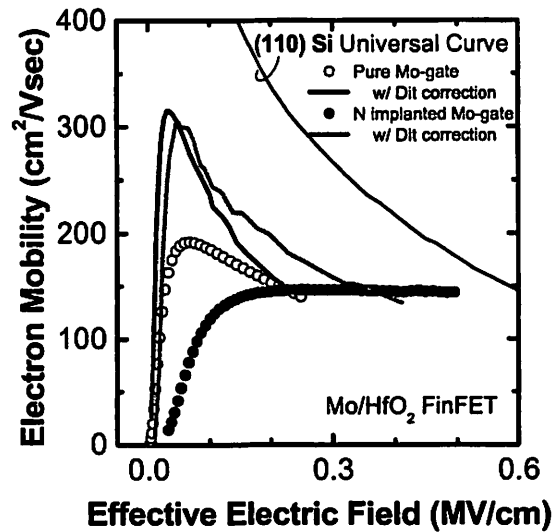


Figure 4.29 Effective electron mobility with and without interface trap density ( $D_{it}$ ) correction for pure and nitrogen implanted (110) n-channel Mo/HfO<sub>2</sub> FinFET.

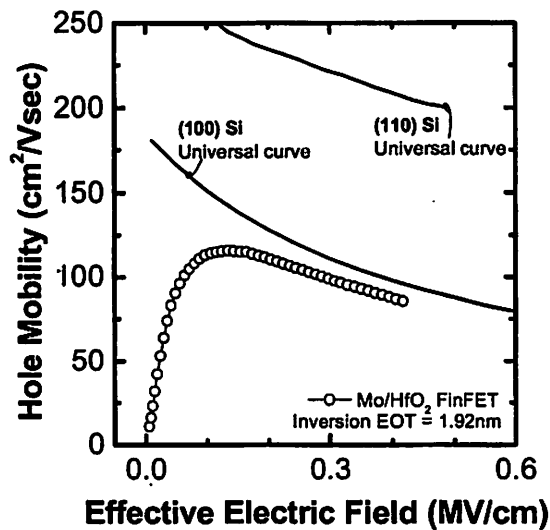


Figure 4.30 Effective hole mobility for (110) p-channel Mo/HfO<sub>2</sub> FinFET.

In order to elucidate the reason for this, the carrier mobility dependence on temperature (in the range from  $-50^{\circ}\text{C}$  to  $200^{\circ}\text{C}$ ) was investigated. Both electron and hole mobilities increase with decreasing temperature due to reduced phonon scattering as shown in Figure 4.31. However, at low field strength (0.1 MV/cm for electrons and 0.2 MV/cm for holes) they exhibit a weaker temperature dependence ( $\mu_{\text{eff}} \sim T^{-0.95}$ ) compared to mobilities for  $\text{SiO}_2$  gate dielectric ( $\mu_{\text{eff}} \sim T^{-1.5}$ ), as shown in Figure 4.32. This result indicates that the mobilities are limited by  $\text{HfO}_2$  soft phonon scattering [4.56, 4.57].

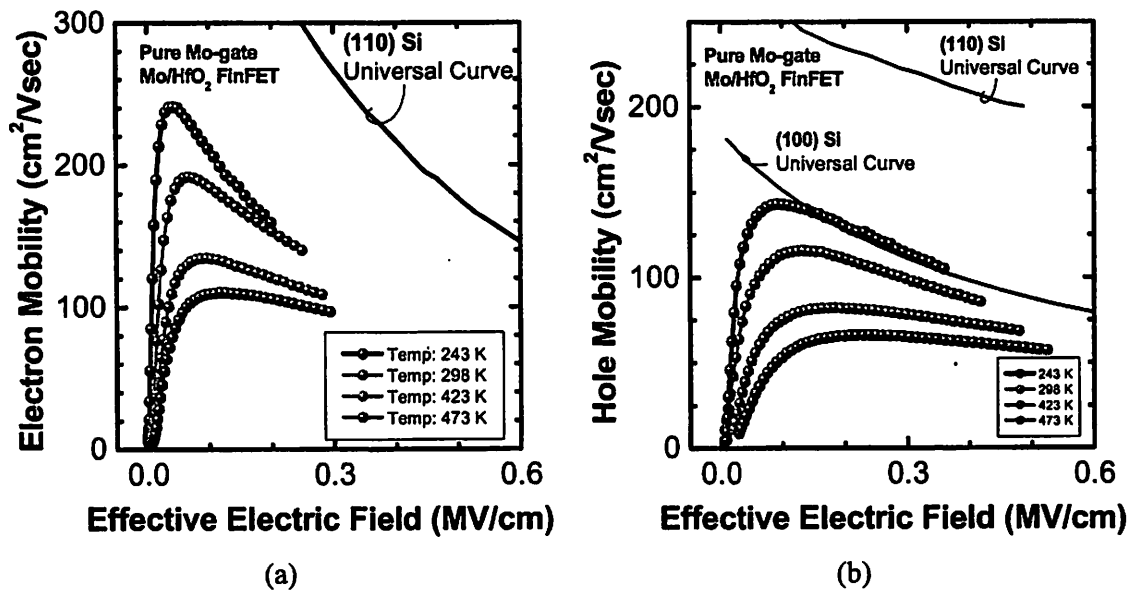


Figure 4.31 Effective (a) electron and (b) hole mobility at various measurement temperatures.

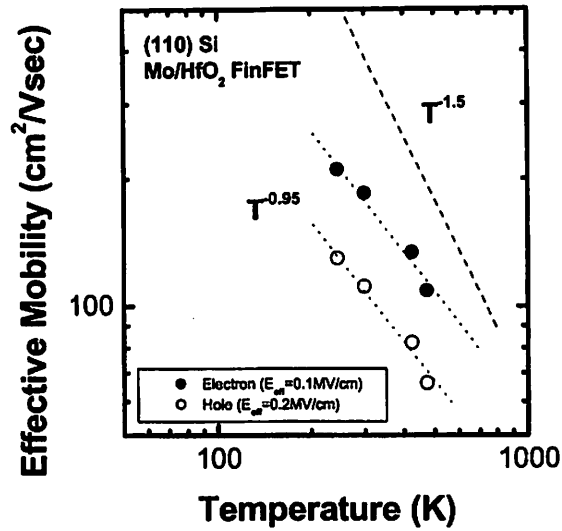


Figure 4.32 Electron and hole mobility for Mo/HfO<sub>2</sub> FinFET less depends on temperature, compared to SiO<sub>2</sub>.

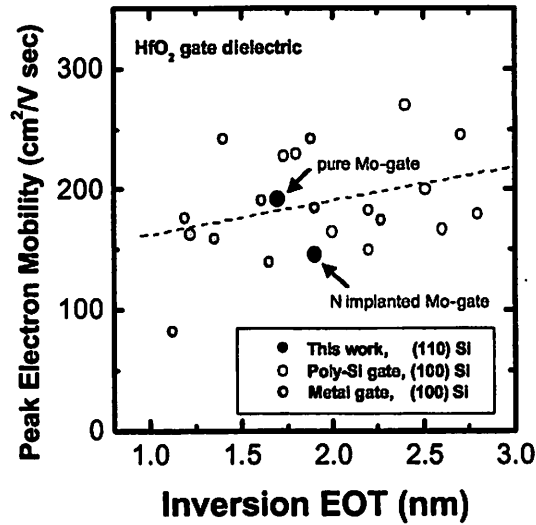


Figure 4.33 Peak electron mobility vs. inversion equivalent oxide thickness (EOT).

Figure 4.33 shows that the peak electron mobility achieved in this work is comparable to previously published data for (100) Si with HfO<sub>2</sub> gate dielectric. This is

notable because the electron mobility is lower for a (110) Si surface than for a (100) Si surface, with SiO<sub>2</sub> gate dielectric.

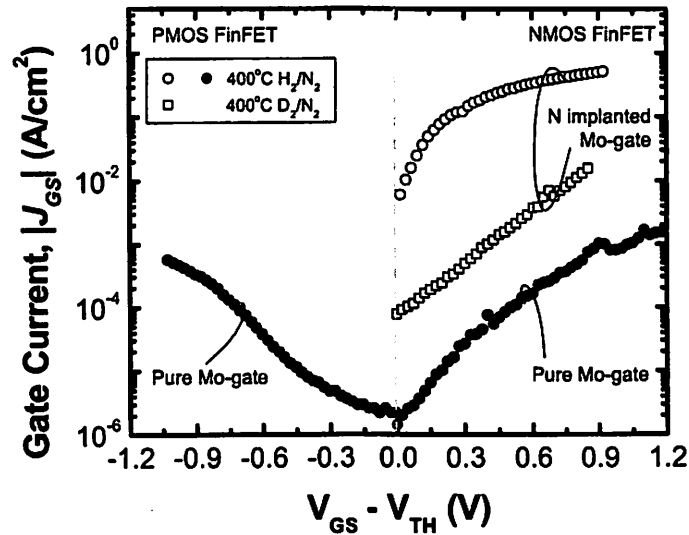


Figure 4.34 Gate leakage current characteristics for n-channel and p-channel Mo/HfO<sub>2</sub> FinFETs. Deuterium (D<sub>2</sub>) annealing is effective to reduce the gate current.

Figure 4.34 shows the measured gate leakage current density characteristics. Lower gate leakage can be seen for the p-channel FinFET as compared to the n-channel FinFET (unimplanted Mo gate), due to the larger hole barrier height for HfO<sub>2</sub> [4.58]. Compared to a poly-Si gated SiO<sub>2</sub> stack, the gate leakage current is reduced by 3-4 orders of magnitude for the same EOT as shown in Figure 4.35. The nitrogen-implanted Mo-



gate n-channel FinFET shows increased gate current, likely due to nitrogen diffusion into the  $\text{HfO}_2$  which degrades the interfacial and bulk properties of  $\text{HfO}_2$  [4.53].

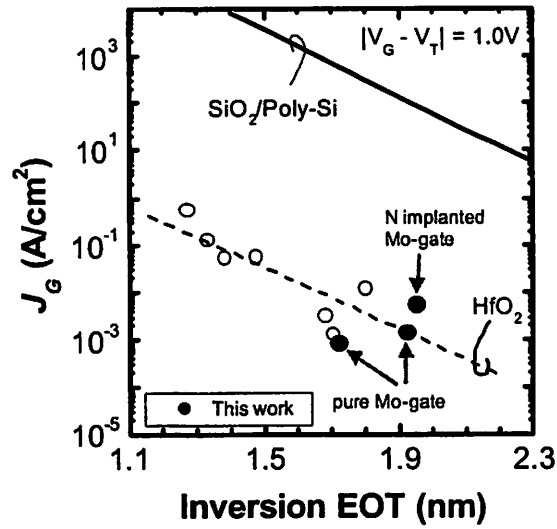


Figure 4.35 Inversion equivalent oxide thickness (EOT) vs. gate leakage current density ( $J_G$ ).

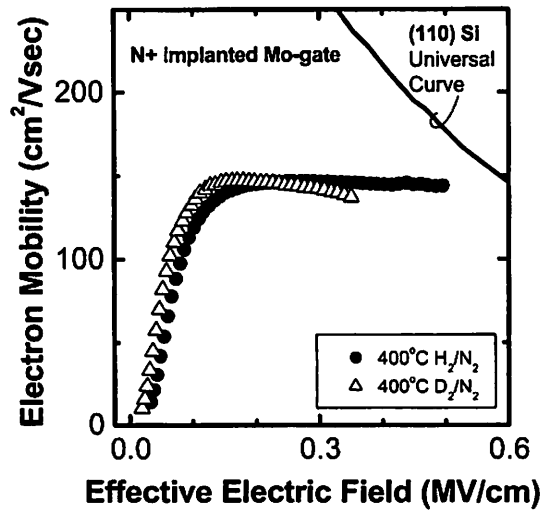


Figure 4.36 Effective electron mobility for different forming gas annealing conditions.

Deuterium annealing is effective to reduce the gate current by 1-2 orders of magnitude as shown in Figure 4.35, and to improve the effective electron mobility slightly as shown in Figure 4.36. This implies that deuterium can effectively passivate traps within  $\text{HfO}_x\text{N}_y$ .

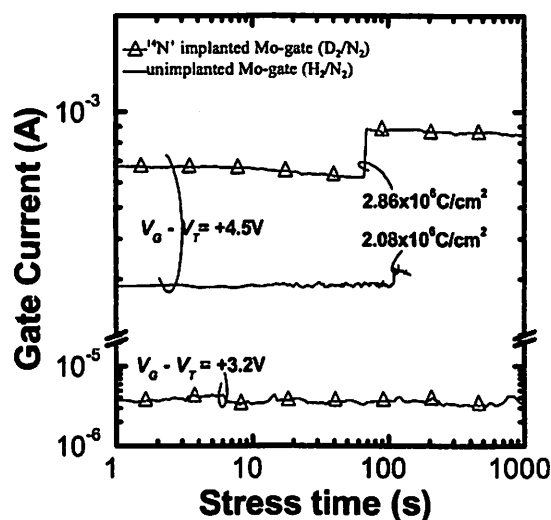


Figure 4.37 Gate dielectric breakdown characteristics for pure and nitrogen implanted Mo-gated  $\text{HfO}_2$  FinFETs.

Figure 4.37 shows the gate dielectric charge-to-breakdown ( $Q_{BD}$ ) characteristics under constant voltage stressing. These initial results indicate that the reliability of a nitrogen-implanted Mo-gate device is comparable to that of a pure Mo-gate device. Key results are summarized in Table 4.2.

Table 4.2 Summary of key device results of Mo-gated HfO<sub>2</sub> FinFETs.

	NMOS		PMOS
	Low VT	High VT	Low VT
nitrogen implant dose (cm <sup>-2</sup> )	1x10 <sup>16</sup>	—	—
measured V <sub>T</sub> (V) (@100nA/μm)	0.28	0.73	- 0.17
subthreshold swing (mV/dec)	67.5	72.6	62.5
Inversion EOT (nm)	1.95	1.75	1.92
gate current density (mA/cm <sup>2</sup> )	5.55	0.86	0.52
peak mobility (cm <sup>2</sup> /Vsec)	146.7	191.9	115.9

## 4.5 Summary

Design methodology and challenges for the advanced thin-body transistor structures have been investigated. Scalability of the gate length strongly depends on the silicon body thickness, necessitating the technological innovations in achieving critical dimensions below the limit of today's lithography process. Adjustment of the desired threshold voltages without degrading the performance benefits is a serious challenge; the ideal solution is the metal gate work function engineering.

Metal gate work function engineering is demonstrated to be a feasible approach for adjusting the threshold voltages of p-channel UTB MOSFETs for the first time. The range of work function values achieved with molybdenum (Mo) gate technology makes it an attractive candidate for the future nanoscale CMOS technologies, particularly for low power and high performance applications.

Integration of metal gate and high- $k$  dielectric employing the advanced thin-body transistor structure is demonstrated for the first time; CMOS FinFETs with Mo gate on HfO<sub>2</sub>. Low gate leakage current density was achieved for a thin inversion equivalent oxide thickness (EOT) down to 1.72 nm with carrier mobilities comparable to previously reported works, which is limited by soft phonon scattering. Threshold voltage control is shown to be feasible by tuning the effective Mo work function via nitrogen implantation. Further process optimization is needed to prevent nitrogen diffusion into the HfO<sub>2</sub> gate dielectric to make Mo-gated HfO<sub>2</sub> FinFET technology suitable for future nanoscale CMOS technology.

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# **Chapter 5**

## **Conclusion**

### **5.1 Summary of Contributions**

Technological innovation has accomplished remarkable progress in the semiconductor industry and CMOS scaling well into the nanometer regime [5.1, 5.2]. Technological challenges in the near future, however, should be overcome for the reduction of standby power consumption without aggravating the device performance. This dissertation addressed the important issues and technological barriers for the integration of advanced gate stack materials and transistor structures into nanoscale CMOS technologies.

To continue CMOS scaling, key technological innovations for advanced thin-body transistor structures are presented. The use of ultra-thin silicon body can suppress the short channel effects effectively, thus eliminating the need of heavy channel doping and state-of-the-art thin gate oxide. Undoped or lightly doped channel can improve the channel carrier mobility due to the reduced impurity scattering and vertical electric field, and eliminating the troublesome issues such as statistical random dopant fluctuation. Self-limited junction depth to ultra-thin body thickness significantly reduces parasitic junction capacitance and leakage current. These benefits combined will significantly improve the overall circuit performance without degrading the standby power consumption.

The most pressing challenge to realize these benefits is the adjustment of threshold voltage. The ideal solution is the gate work function engineering technology; other potential solutions such as heavy channel doping and asymmetrical double gate structures will lose many of the above benefits [5.3]. For the first time, tunable molybdenum (Mo) gate work function technology is demonstrated to adjust the threshold voltage of ultra-thin body (UTB) MOSFETs. Integration of Mo technology and high- $k$  dielectric ( $\text{HfO}_2$ ) employing FinFETs is demonstrated, for the first time. For these

implementations, two process technologies were developed; damage-free sputtering and high-selectivity dry etching of Mo films.

Major conclusions on the topics listed above are summarized as follows.

### ***5.1.1 Metal Gate Technology***

Metal gate electrodes in a CMOS transistor eliminate problems such as the poly-Si gate depletion (PDE), boron dopant penetration, and high sheet resistance. Material selection criteria for integration of alternative gate electrode materials were summarized.

Molybdenum (Mo) is a promising candidate gate material. Two critical process technologies were developed for Mo gate films; *damage-free sputtering and high-selectivity dry etching*. With a plasma charge trap (PCT), sputtering damage to the gate dielectric can be minimized for improved drive current and gate oxide integrity (GOI). Mo gate can be patterned without leaving any residue or stringers due to the highly selective etch process and directly applicable to advanced thin-body transistor structures.

The work function of Mo gate can be tuned by nitrogen implantation followed by a thermal anneal, to provide a means for adjusting the threshold voltage of advanced thin-body transistor structures. The work function shift of nitrogen implanted Mo gate is

resulted from the simultaneous modification of the microstructure and chemistry of Mo film at the gate dielectric interface.

### ***5.1.2 High- $k$ Gate Dielectric Technology***

High- $k$  gate dielectrics decrease the leakage current through the dielectric while maintaining the same gate capacitance, thus are indispensable for the low-power portable application of nanoscale CMOS technologies. The key basic material properties for integration of alternative gate dielectric materials were summarized.

Hafnium oxide ( $\text{HfO}_2$ ) is one of the most promising candidates to replace the silicon dioxide ( $\text{SiO}_2$ ). Through experimental device fabrication, the impact of gate process (poly-Si vs. poly-SiGe) technology on equivalent-oxide-thickness (EOT) and leakage current was investigated; thermal budget reduction and careful selection of gate electrode material leads to thinner EOT and smaller gate leakage current. The impact of oxygen vacancies in  $\text{HfO}_2$  gate dielectric was discussed. Generation of oxygen vacancies in  $\text{HfO}_2$  is thermodynamically driven and causes Si interfacial layer formation and gate Fermi-level pinned in MOS devices.

Successful integration of  $\text{HfO}_2$  into CMOS fabrication depends on the minimization of oxygen vacancy formation through careful selection of the gate electrode materials and reduction of post-gate-deposition thermal budget.

### ***5.1.3 Advanced Transistor Structures***

Advanced thin-body transistor structures suppress the short channel effects without the use of heavy channel doping and ultra thin gate oxide thickness. Design methodology and challenges were investigated. Adjustment of the desired threshold voltages to maximize the performance benefits is a serious challenge; the ideal solution is the metal gate work function engineering.

Tunable work function Mo gate is demonstrated to be a feasible approach for adjusting the threshold voltages of UTB MOSFETs for the first time. The range of work function values achieved with Mo gate technology makes it an attractive candidate for advanced thin-body transistor structures. Integration of metal gate and high- $k$  dielectric employing FinFETs is demonstrated for the first time; Mo-gated  $\text{HfO}_2$  CMOS FinFETs achieved the reduction of gate leakage current over 3 orders-of-magnitude for inversion



equivalent oxide thickness down to 1.72 nm. Measured carrier mobilities are comparable to previously reported works, which is limited by the soft phonon scattering.

Further process optimization is needed to prevent nitrogen diffusion into the HfO<sub>2</sub> gate dielectric to make Mo-gated HfO<sub>2</sub> FinFET technology suitable for future nanoscale CMOS technologies.

## **5.2 Suggestions for Future Work**

### ***5.2.1 Mobility Enhancement for Advanced Transistor Structures***

Process-induced strain in the channel of a MOSFET can enhance mobility and hence circuit performance, and is already employed in CMOS production today [5.4]. For a p-channel bulk-Si MOSFET, uniaxial compressive strain induced by epitaxially grown Si<sub>1-x</sub>Ge<sub>x</sub> source/drain regions is very effective for enhancing hole mobility [5.4]. This approach may not be directly applicable to advanced thin-body transistor structures, however, because of the lack of a crystalline Si lattice directly beneath the source/drain regions to serve as a template for epitaxial growth of Si<sub>1-x</sub>Ge<sub>x</sub>. Biaxially tensile strained (in the plane parallel to the wafer surface) silicon-on-insulator (SOI) films can be

achieved with a wafer-bonding process [5.5]; however, this approach is relatively expensive and would not provide biaxial strain in the plane of current conduction for the FinFET structure. Therefore, alternative approaches to inducing strain in the channel of advanced thin-body FET structures are necessary through a stressed capping layer, stressed gate electrode(s), or strained-Si<sub>1-x</sub>Ge<sub>x</sub> S/D regions.

### ***5.2.2 Reliability of Advanced Materials and Transistor Structures***

Investigation of reliabilities for advanced thin-body transistor structures such as hot carrier injection, gate dielectric integrity, and low-frequency noise should be carried out before putting such device structures into mass-production. Nonetheless, only a few works for the advanced transistor structures has been done to date [5.6, 5.7, 5.8].

Thinner silicon body is helpful to reduce the hot carrier injection, thus improving the hot carrier reliability due to the reduced electron temperature and generated electron-hole pair generation rate [5.6]. On the other hand, the use of silicon surface orientation other than (100) and rough surface with plasma etching damage together with nitrogen implantation into Mo-gate may cause degradation in gate oxide integrity, necessitating proper treatments such as hydrogen annealing and so on [5.9, 5.10]. While these

advanced transistor structures with advanced gate stack materials may appear immune to hot carrier reliability [5.6, 5.11], more detail and comprehensive reliability studies are in pressing need for the adaptation into IC production.

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# Appendix A

## Fabrication Processes for Mo-gated UTB MOSFETs

STEP	PROCESS	CONDITIONS	EQUIPMENT	COMMENT
	Initial wafers	4" (100) p-type 100nm thick SOI with 400 nm BOX		Unibond® SOI
<b>0.0</b>	<b>LABELING</b>			
0.1	Label		Diamond pencil	
0.2	DI Rinse	3 Cycle DI Water Rinse	Sink6	
0.3	Thickness Measure	Silicon	NanoDuv	R:4
0.4	HF Cleaning	10:1 HF 5min	Sink6	
<b>1.0</b>	<b>BODY THINNING</b>			
1.1	Piranha Cleaning	120°C 10min	Sink6	
1.2	Wet Oxidation	Wet O <sub>2</sub> 850°C 90min	Tystar2	R:SWETOXB
1.3	Wet Etching	10:1 HF 10min	Sink6	
1.4	Thickness Measure	Silicon	NanoDuv	R:4
1.5	Piranha Cleaning	120°C 10min	Sink6	
1.6	Wet Oxidation	Wet O <sub>2</sub> 850°C 60min	Tystar2	R:SWETOXB
1.7	Wet Etching	10:1 HF 5min	Sink6	
1.8	Thickness Measure	Silicon	NanoDuv	R:4
1.9	Piranha Cleaning	120°C 10min	Sink6	
1.10	Dry Oxidation	Dry O <sub>2</sub> 850°C 15min	Tylan6	R:SGATEOX
1.11	Wet Etching	10:1 HF 2min	Sink6	
1.12	Thickness Measure	Silicon	NanoDuv	R:4
<b>2.0</b>	<b>ACTIVE</b>			
2.1	HMDS	120°C 1min	Primeoven	R:0
2.2	PR Coating	1.1µm 90°C Soft Bake	Svgcoat1,2	OCG OiR 897-10i
2.3	Expose	Focus: 248 ET: 3.1sec	Gcaws1,2	Mask: Active
2.4	Post Exp. Bake	120°C 1min	Svgdev	
2.5	Develop	60sec	Svgdev	OPD4262
2.6	Hard Bake	120°C 10min	Vwroven	

STEP	PROCESS	CONDITIONS	EQUIPMENT	COMMENT
2.7	Active Dry Etching	Lam5 BT: TP/BP=200/40W 30mT CF <sub>4</sub> =100sccm 7sec ME: TP/BP=300/150W 15mT Cl <sub>2</sub> /HBr =50/150scm EPD OE: TP/BP=70/30W 15mT HBr/He/O <sub>2</sub> =50/50/1 sccm 15sec		
2.8	Thickness Measure	Buried Oxide	NanoDuv	R:1
2.9	PR Ashing	O <sub>2</sub> 300W 5min	Technics-C	
2.10	Piranha Cleaning	120°C 10min	Sink8	
2.11	CD Measure	SEM	LEO	
3.0	<b>GATE STACK</b>			
3.1	Piranha Cleaning	120°C 10min	Sink6	
3.2	HF Cleaning	25:1 HF 30sec	Sink6	
3.3	Gate Oxidation	Dry O <sub>2</sub> 750°C 13min 30sec N <sub>2</sub> 900°C 20min	Tylan5	R:Thin_Ann
3.4	Thickness Measure	Oxide (monitor wafer)	Sopra	
3.5	Mo Deposition	200°C 300W 10mT	UHV Sputtering	Without PCT
3.6	Rs Measure	Monitor wafer	4PointProbe	
4.0	<b>NITROGEN IMPLANTATION</b>			
4.1	HMDS	120°C 1min	Primeoven	R:0
4.2	PR Coating	1.1µm 90°C Soft Bake	Svgcoat1,2	OCG OiR 897-10i
4.3	Expose	Focus: 248 ET: 3.1sec	Gcaws1,2	Mask: Blank
4.5	Develop	60sec	Svgdev	OPD4262
4.6	Nitrogen Implant	<sup>14</sup> N <sup>+</sup> 20KeV (dose split)	Implant Service	Foundry Company
4.7	PR Ashing	O <sub>2</sub> 200W 7min	Technics-C	
4.8	PR Strip	Acetone	Manual Bath	
4.9	HF Cleaning	25:1 HF 20"	Sink7	
4.10	N <sup>+</sup> Poly-Si Deposition	Nucleation: 550°C 300mT SiH <sub>4</sub> =200sccm 1min Deposition: 550°C 300mT SiH <sub>4</sub> /PH <sub>3</sub> =200/5sccm		Tystar19 (dedicated boat)
4.11	Thickness Measure	Silicon (monitor wafer)	NanoDuv	R:4
4.12	LTO Deposition	450°C 6min undoped	Tystar11	R:11SULTOA
4.13	Thickness Measure	Oxide (monitor wafer)	NanoDuv	
5.0	<b>GATE</b>			
5.1	HMDS	120°C 1min	Primeoven	R:0
5.2	PR Coating	1.1µm 90°C Soft Bake	Svgcoat1,2	OCG OiR 897-10i

STEP	PROCESS	CONDITIONS	EQUIPMENT	COMMENT
5.3	Expose	Focus: 248 ET: 3.1sec	Gcaws1,2	Mask: Gate
5.4	Post Exp. Bake	120°C 1min	Svgdev	
5.5	Develop	60sec	Svgdev	OPD4262
5.6	Hard Bake	120°C 1min	Vwroven	
5.7	CD Measure	SEM	LEO	
5.8	Ashing	O <sub>2</sub> 30W 1min	Technics-C	Repeat
5.9	CD Measure	SEM	LEO	
5.10	Gate Stack Dry Etching	Equipment: Lam5 BT: TP/BP=200/40W 30mT CF <sub>4</sub> =100sccm 60sec Poly-Si ME: TP/BP=300/150W 15mT Cl <sub>2</sub> /HBr=50/150 sccm EDP + 10sec Mo ME: TP/BP=150/100W 13mT Cl <sub>2</sub> /O <sub>2</sub> =70/60 sccm EDP + 20sec		
5.11	HF Cleaning	200:1 HF 30sec	Sink7	
5.12	PR Ashing	O <sub>2</sub> 200W 7min	Technics-C	Repeat
5.13	HF Cleaning	200:1 HF 15sec	Sink7	
5.14	Polymer Removal	Acetone	Manual Bath	
5.15	LTO Deposition	450°C 1min Undoped	Tystar11	R:11SULTOA
5.16	Thickness Measure	Buried Oxide & monitor wafer	NanoDuv	
5.17	Spacer Dry Etching	Equipment: Lam5 TP/BP=200/40W 30mT CF <sub>4</sub> =100sccm 30sec		
5.18	HF Cleaning	25:1 HF 10sec	Sink7	
6.0	<b>SELECTIVELY RAISED GE SOURCE/DRAIN</b>			
6.1	HF Cleaning	200:1 HF 30sec	Sink7	
6.2	Ge Deposition	340°C 300mT GeH <sub>4</sub> =200sccm 10min	Tystar19	
6.3	LTO Deposition	450°C 90sec Undoped	Tystar11	R:11SULTOA
7.0	<b>N+ SOURCE/DRAIN IMPLANTATION</b>			
7.1	HMDS	120°C 1min	Primeoven	R:0
7.2	PR Coating	1.1µm 90°C Soft Bake	Svgcoat1,2	OCG OiR 897-10i
7.3	Expose	Focus: 248 ET: 3.1sec	Gcaws1,2	Mask: N+ Active
7.4	Post Exp. Bake	120°C 1min	Svgdev	
7.5	Develop	60sec	Svgdev	OPD4262
7.6	Hard Bake	120°C 10min	Vwroven	
7.7	N+ Implant	As 40KeV 5E15cm <sup>-2</sup> Tilt 0°	Implant Service	Foundry Company
7.8	PR Ashing	O <sub>2</sub> 300W 5min	Technics-C	

STEP	PROCESS	CONDITIONS	EQUIPMENT	COMMENT
7.9	Piranha Cleaning	120°C 5min	Sink8	
8.0	<b>P+ SOURCE/DRAIN IMPLANTATION</b>			
8.1	HMDS	120°C 1min	Primeoven	R:0
8.2	PR Coating	1.1µm 90°C Soft Bake	Svgcoat1,2	OCG OiR 897-10i
8.3	Expose	Focus: 248 ET: 3.1sec	Gcaws1,2	Mask: P+ Active
8.4	Develop	60sec	Svgdev	OPD4262
8.5	P+ Implant	B 20KeV 3E15cm <sup>-2</sup> Tilt 0°	Implant Service	Foundry Company
8.6	PR Ashing	O <sub>2</sub> 300W 5min	Technics-C	
8.7	Piranha Cleaning	120°C 5min	Sink8	
8.8	Piranha Cleaning	120°C 5min	Sink6	
8.9	RTA	N <sub>2</sub> 700°C 60sec	Heatpulse3	
9.0	<b>METAL CONTACT</b>			
9.1	LTO Deposition	450°C 20min Undoped	Tystar11	ILD1
9.2	HMDS	120°C 1min	Primeoven	R:0
9.3	PR Coating	1.1µm 90°C Soft Bake	Svgcoat1,2	OCG OiR 897-10i
9.4	Expose	Focus: 248 ET: 3.1sec	Gcaws1,2	Mask: MC
9.5	Post Exp. Bake	120°C 1min	Svgdev	
9.6	Develop	60sec	Svgdev	OPD4262
9.7	Hard Bake	120°C 20min	Vwroven	
9.8	Contact Etching	Equipment: Lam5 TP/BP=200/40W 30mT CF <sub>4</sub> =100sccm 3min		
9.9	Thickness Measure	Buried Oxide	NanoDuv	
9.10	HF Cleaning	25:1 HF 1min	Sink7	Repeat
9.11	Thickness Measure	Buried Oxide	NanoDuv	
9.12	PR Ashing	O <sub>2</sub> 300W 5min	Technics-C	
9.13	Piranha Cleaning	120°C 5min	Sink8	
9.14	HF Wet Etching	25:1 HF 30sec	Sink7	
9.15	Al Deposition	6mT 15cm/min 1pass	Cpa	
10.0	<b>METALLIZATION</b>			
10.1	HMDS	120°C 1min	Primeoven	R:0
10.2	PR Coating	1.1µm 90°C Soft Bake	Svgcoat1,2	OCG OiR 897-10i
10.3	Expose	Focus: 248 ET: 3.1sec	Gcaws1,2	Mask: Metal



STEP	PROCESS	CONDITIONS	EQUIPMENT	COMMENT
10.4	Post Exp. Bake	120°C 1min	Svgdev	
10.5	Develop	60sec	Svgdev	OPD4262
10.6	Hard Bake	120°C 20min	Vwroven	
10.7	Al Wet Etching	Al Etchant	Sink8	
10.8	Thickness Measure	Buried Oxide	NanoDuv	
10.9	PR Ashing	O <sub>2</sub> 300W 5min	Technics-C	
10.10	DI Rinse	3 Cycle DI Water Rinse	Sink8	
10.11	Sintering	400°C 10% H <sub>2</sub> 30min	Tylan13	R:VSINT400

# Appendix B

## Fabrication Processes for Mo-gated HfO<sub>2</sub> FinFETs

STEP	PROCESS	CONDITIONS	EQUIPMENT	COMMENT
	Initial wafers	4" (100) p-type 100nm thick SOI with 400 nm BOX		Unibond® SOI
<b>0.0</b>	<b>LABELING</b>			
0.1	Label		Diamond pencil	
0.2	DI Rinse	3 Cycle DI Water Rinse	Sink6	
0.3	Thickness Measure	Silicon	NanoDuv	R:4
0.4	HF Cleaning	10:1 HF 5min	Sink6	
<b>1.0</b>	<b>BODY THINNING</b>			
1.1	Piranha Cleaning	120°C 10min	Sink6	
1.2	Wet Oxidation	Wet O <sub>2</sub> 850°C 100min	Tystar2	R:SWETOXB
1.3	Wet Etching	10:1 HF 10min	Sink6	
1.4	Thickness Measure	Silicon	NanoDuv	R:4
1.5	Piranha Cleaning	120°C 10min	Sink6	
1.6	Dry Oxidation	Dry O <sub>2</sub> 900°C 4min	Tylan6	R:SGATEOX
1.7	Piranha Cleaning	120°C 10min	Sink6	
1.8	Sac. SiGe Deposition	Tystar19 Nucleation: 550°C 300mT SiH <sub>4</sub> =200sccm 30sec Deposition: 450°C 300mT SiH <sub>4</sub> /GeH <sub>4</sub> =124/80sccm 30sec 18min		
1.9	Measurement	SiGe	α-step	
<b>2.0</b>	<b>ACTIVE FIN</b>			
2.1	HMDS	120°C 1min	Primeoven	R:0
2.2	PR Coating	1.1µm 90°C Soft Bake	Svgcoat1,2	OCG OiR 897-10i
2.3	Expose	Focus: 248 ET: 3.1sec	Gcaws1,2	Mask: Fin
2.4	Post Exp. Bake	120°C 1min	Svgdev	
2.5	Develop	60sec	Svgdev	OPD4262
2.6	Hard Bake	120°C 10min	Vwroven	
2.7	Sac SiGe Etching	Lam5 BT: TP/BP=200/40W 30mT CF <sub>4</sub> =100sccm 7sec ME: TP/BP=300/150W 15mT Cl <sub>2</sub> /HBr=50/150sccm EDP+5sec		

STEP	PROCESS	CONDITIONS	EQUIPMENT	COMMENT
2.8	HF Cleaning	200:1 HF 10sec	Sink7	
2.9	PR Ashing	O <sub>2</sub> 300W 5min	Technics-C	
2.10	HF Cleaning	200:1 HF 10sec	Sink7	
2.11	Piranha Cleaning	120°C 10min	Sink8	
2.12	LTO Deposition	450°C 3min Undoped	Tystar11	Spacer Lithography
2.13	Thickness Measure	Oxide	NanoDuv	
3.0	<b>ALIGNMENT KEY PROTECTION</b>			
3.1	HMDS	120°C 1min	Primeoven	R:0
3.2	PR Coating	1.1µm 90°C Soft Bake	Svgcoat1,2	OCG OiR 897-10i
3.3	Expose	Focus: 248 ET: 3.1sec	Gcaws1,2	Mask: Alignment
3.4	Post Exp. Bake	120°C 1min	Svgdev	
3.5	Develop	60sec	Svgdev	OPD4262
3.6	Hard Bake	120°C 10min	Vwroven	
3.7	LTO Spacer Etching	Lam5 TP/BP=200/40W 30mT CF <sub>4</sub> =100sccm 30sec (EPD detection)		
3.8	SiGe Removal	Lam5 BT: TP/BP=200/40W 30mT CF <sub>4</sub> =100sccm 7sec ME:TP/BP=250/120W 35mT HBr/O <sub>2</sub> =200/5 60sec		
3.9	HF Cleaning	200:1 HF 10sec	Sink7	
3.10	PR Ashing	O <sub>2</sub> 300W 5min	Technics-C	
3.11	HF Cleaning	200:1 HF 10sec	Sink7	
3.12	Piranha Cleaning	120°C 10min	Sink8	
4.0	<b>ACTIVE PAD</b>			
4.1	HMDS	120°C 1min	Primeoven	R:0
4.2	PR Coating	1.1µm 90°C Soft Bake	Svgcoat1,2	OCG OiR 897-10i
4.3	Expose	Focus: 248 ET: 3.1sec	Gcaws1,2	Mask: Active
4.4	Post Exp. Bake	120°C 1min	Svgdev	
4.5	Develop	60sec	Svgdev	OPD4262
4.6	Hard Bake	120°C 1min	Vwroven	
4.7	Ashing	O <sub>2</sub> 30W 1min	Technics-C	Repeat
4.8	CD Measure	SEM	LEO	
4.9	Active Etching	Lam5 BT: TP/BP=200/40W 30mT CF <sub>4</sub> =100sccm 40sec ME: TP/BP=300/150W 15mT Cl <sub>2</sub> /HBr=50/150 sccm EDP + 5sec		
4.10	HF Cleaning	200:1 HF 10sec	Sink7	

STEP	PROCESS	CONDITIONS	EQUIPMENT	COMMENT
4.11	PR Ashing	O <sub>2</sub> 300W 5min	Technics-C	
4.12	HF Cleaning	200:1 HF 10sec	Sink7	
4.13	Piranha Cleaning	120°C 10min	Sink8	
4.14	CD Measure	SEM	LEO	
4.15	Piranha Cleaning	120°C 10min	Sink6	
4.16	Sac. Oxidation	Dry O <sub>2</sub> 900°C 3min	Tylan6	R:SGATEOX
4.17	Thickness Measure	Oxide	NanoDuv	
<b>5.0</b>	<b>GATE STACK</b>			
5.1	HF Cleaning	25:1 HF 60sec		Univ of Texas, Austin
5.2	Surface Treatment	NH <sub>3</sub> 700°C 500T 10sec		
5.3	HfO <sub>2</sub> Deposition	500°C 4T 3min Hf(OC(CH <sub>3</sub> ) <sub>3</sub> ) <sub>4</sub>		
5.4	Post Depo. Treatment	N <sub>2</sub> 700°C 500T 30sec		
5.5	Mo Deposition	200°C 300W 10mT	Novellus	With PCT
5.6	Rs Measure	Monitor wafer		
<b>6.0</b>	<b>NITROGEN IMPLANTATION</b>			
6.1	HMDS	120°C 1min	Primeoven	R:0
6.2	PR Coating	1.1µm 90°C Soft Bake	Svgcoat1,2	OCG OiR 897-10i
6.3	Expose	Focus: 248 ET: 5.0sec	Gcaws1,2	Mask: Blank
6.5	Develop	60sec	Svgdev	OPD4262
6.6	Nitrogen Implant	<sup>14</sup> N <sup>+</sup> 5KeV 1E16cm <sup>-2</sup> 30° tilt		Axcelis Inc.
6.7	PR Ashing	O <sub>2</sub> 200W 7min	Technics-C	
6.8	PR Strip	Acetone	Manual Bath	
6.9	HF Cleaning	25:1 HF 20sec	Sink7	
6.10	N <sup>+</sup> Poly-Si Deposition	Nucleation: 550°C 300mT SiH <sub>4</sub> =200sccm 1min Deposition: 550°C 300mT SiH <sub>4</sub> /PH <sub>3</sub> =200/5sccm		Tystar19 (dedicated boat)
6.11	Thickness Measure	Silicon (monitor wafer)	NanoDuv	
6.12	CMP	CMP DF=8psi Table/Chuck=24/6rpm BP=1psi Slurry=100ml/min		
6.13	Post Cleaning	Sink8 DI Rinse NH <sub>4</sub> OH 1min Piranha 120°C 1min 5:1 BHF 10sec		
6.14	Post Cleaning2	Manual Bath H <sub>2</sub> O/NH <sub>4</sub> OH/H <sub>2</sub> O <sub>2</sub> =5/1/1 65°C 5min		
6.15	Thickness Measure	Silicon (monitor wafer)	NanoDuv	R:4
6.16	LTO Deposition	450°C 8min30sec undoped	Tystar11	R:11SULTOA

STEP	PROCESS	CONDITIONS	EQUIPMENT	COMMENT
6.17	Thickness Measure	Oxide (monitor wafer)	NanoDuv	
<b>7.0</b>	<b>GATE</b>			
7.1	HMDS	120°C 1min	Primeoven	R:0
7.2	PR Coating	1.1µm 90°C Soft Bake	Svgcoat1,2	OCG OiR 897-10i
7.3	Expose	Focus: 248 ET: 3.1sec	Gcaws1,2	Mask: Gate
7.4	Post Exp. Bake	120°C 1min	Svgdev	
7.5	Develop	60sec	Svgdev	OPD4262
7.6	CD Measure	SEM	LEO	
7.7	Ashing	O <sub>2</sub> 30W 1min	Technics-C	Repeat
7.8	CD Measure	SEM	LEO	
7.9	LTO Dry Etching	Lam5 BT: TP/BP=200/40W 30mT CF <sub>4</sub> =100sccm 60sec		
7.10	PR Ashing	O <sub>2</sub> 200W 7min	Technics-C	
7.11	CD Measure	SEM	LEO	
7.12	LTO Trimming	200:1 HF 30sec	Sink7	Repeat
7.13	CD Measure	SEM	LEO	
7.14	Gate Stack Dry Etching	Lam5 BT: TP/BP=200/40W 30mT CF <sub>4</sub> =100sccm 60sec Poly-Si ME: TP/BP=300/150W 15mT Cl <sub>2</sub> /HBr=50/150 sccm EDP + 10sec Mo ME: TP/BP=150/100W 13mT Cl <sub>2</sub> /O <sub>2</sub> =70/60 sccm EDP + 20sec		
7.15	HF Cleaning	200:1 HF 30sec	Sink7	
7.16	Polymer Removal	Acetone	Manual Bath	
7.17	DI Rinse	3 Cycle DI Water Rinse	Sink8	
<b>8.0</b>	<b>N+ SOURCE/DRAIN IMPLANTATION</b>			
8.1	HMDS	120°C 1min	Primeoven	R:0
8.2	PR Coating	1.1µm 90°C Soft Bake	Svgcoat1,2	OCG OiR 897-10i
8.3	Expose	Focus: 248 ET: 3.1sec	Gcaws1,2	Mask: N+ Active
8.4	Develop	60sec	Svgdev	OPD4262
8.5	N+ Implant	P 40&30KeV 2E15cm <sup>-2</sup> Tilt 0°		Axcelis Inc.
8.6	PR Ashing	O <sub>2</sub> 200W 7min	Technics-C	
8.7	PR Strip	Acetone	Manual Bath	
8.8	DI Rinse	3 Cycle DI Water Rinse	Sink8	
<b>9.0</b>	<b>P+ SOURCE/DRAIN IMPLANTATION</b>			
9.1	HMDS	120°C 1min	Primeoven	R:0

STEP	PROCESS	CONDITIONS	EQUIPMENT	COMMENT
9.2	PR Coating	1.1µm 90°C Soft Bake	Svgcoat1,2	OCG OiR 897-10i
9.3	Expose	Focus: 248 ET: 3.1sec	Gcaws1,2	Mask: P+ Active
9.4	Develop	60sec	Svgdev	OPD4262
9.5	P+ Implant	BF <sub>2</sub> 25KeV 3E15cm <sup>-2</sup> Tilt 0°	Implant Service	Foundry Company
9.6	PR Ashing	O <sub>2</sub> 300W 5min	Technics-C	
9.7	PR Strip	Acetone	Manual Bath	
9.8	DI Rinse	3 Cycle DI Water Rinse	Sink8	
9.9	RTA	N <sub>2</sub> 900°C 60sec		
9.10	DI Rinse	3 Cycle DI Water Rinse	Sink8	
9.11	Sintering	600°C 10% H <sub>2</sub> 30min	Tylan18	R:H2SINT6A
9.12	Sintering	400°C 10% H <sub>2</sub> (D <sub>2</sub> ) 30min	Tylan18	R:H2SINT4A