

Reports in Electron Physics 2000/24
Espoo, September 2000

Ultrahigh Vacuum Plasma Oxidation in the Fabrication of Ultrathin Silicon Dioxide Films

Tero Majamaa

Dissertation for the degree of Doctor of Science in Technology to be presented with due permission for public examination and debate in Auditorium S4 at Helsinki University of Technology (Espoo, Finland) on the 20th of October, 2000, at 12 o'clock noon.

Helsinki University of Technology
Department of Electrical and Communications Engineering
Electron Physics Laboratory

Teknillinen korkeakoulu
Sähkö- ja tietoliikennetekniikan osasto
Elektronifysiikan laboratorio

Distribution:
Helsinki University of Technology
Electron Physics Laboratory
P.O. Box 3000
FIN-02015 HUT
Finland
Tel. +358-9-451 2322
Fax. +358-9-451 5008

ISBN 951-22-5146-9
ISSN 0781-4984

Otamedia Oy
Espoo 2000

Abstract

This thesis discusses the fabrication and characterization of ultrathin insulator films. These are essential for nanoscale semiconductor device fabrication. When the insulator layer thickness is only a few times the molecule diameter, it is crucial that both film homogeneity and the insulator/semiconductor interface quality are as high as possible. The small dimensions must also be taken into account in characterization, because the established measurement techniques and analyses used in more traditional MOS characterization are not necessarily valid any more.

In this work, the metal/silicon dioxide/silicon structure (especially oxide and oxide/semiconductor interface quality) and various silicon dioxide fabrication methods are discussed. The focus in the experimental work is on the studying the plasma assisted oxidation of silicon in an ultra high vacuum chamber and on characterizing the fabricated films. Some of the film properties are found to be excellent: interface smoothness is of a very high quality and interface state densities are low (10^{11} eV⁻¹cm⁻² or lower in the mid-gap) even without any annealing. Process control also seems to be good, as is the breakdown field. The oxide charge, however, is quite high. This may cause considerable harm. One of the consequences is an increased leakage current. This also significantly decreases the device life time by increasing current generated defects. In the annealing experiments carried out, the oxide charge was seen to decrease, indicating that the quality of the silicon dioxide films can be significantly improved by optimization of the thermal treatments.

The molecular beam epitaxy system used in processing is designed mainly for research purposes, offering possibility to gain much information about the oxidation process itself. Other, and cheaper, thermal oxidation procedures have in recent years already been developed to a very high level, which means that the process developed is not necessarily the best choice for conventional IC manufacturing purposes. It offers, however, useful applications for research into silicon-based nanostructures, such as silicon/silicon dioxide heterostructures.

Keywords: Ultrathin silicon dioxide, ultrahigh vacuum, plasma oxidation

Preface

This thesis for the degree of Doctor of Science in Technology was mainly researched at the Helsinki University of Technology Electron Physics Laboratory during the years 1995 - 1999. I would specially like to thank Professor Juha Sinkkonen who has supervised my work. I also thank the other personnel of the Electron Physics Laboratory, where I had the great privilege of working from 1990 to 1998. I owe a special debt of gratitude to the personnel of the Microelectronics Centre at the Helsinki University of Technology and VTT Electronics for the use of their clean room facilities and equipment as well as for many helpful conversations and invaluable advise.

I am also grateful to the Academy of Finland for the generous financial support they granted to the research project during my years at the Electron Physics Laboratory as well as for the encouragement of my co-workers and superiors since I started at the Academy.

Tero Majamaa

List of Symbols

$a, A, a_0, B, G_0, k, l_0, r, \tau$

constants

A	area
Ai	Airy function
B	susceptance
C	capacitance
C_0	oxygen concentration on sample surface
C_1	concentration of oxidizing molecules
C_{gr}	growth site concentration
C_{HF}	high frequency capacitance
C_{it}	interface trap state capacitance
C_{LF}	low frequency capacitance
C_{OX}	oxygen concentration on silicon dioxide surface
C_S	oxygen concentration at silicon/silicon dioxide interface
$C_{ox,s}$	saturated oxide capacitance
C_{mos}	MOS capacitance
C_{SI}	oxygen concentration on silicon surface
D	diffusion coefficient
D_{it}	surface state density
d_0	initial thickness of the silicon dioxide layer
d_{cr}	critical thickness of a silicon dioxide layer
d_{ox}	oxide thickness
d_{sc}	depletion region length
E	energy, electric field
E_A	activation energy
E_c	conduction band energy
E_f	Fermi energy
E_g	energy bandgap
E_{n0}	electron ground state energy in potential well
E_{ox}	electric field in the silicon dioxide
E_{p0}	hole ground state energy in potential well
E_s	electric field at the silicon/silicon dioxide interface

E_v	valence band energy
F	flux, modulation factor
f	frequency
(g)	gas
G	conductance
G_{mos}	leakage conductance through MOS structure
h	Planck constant
i	imaginary unit
J	current density
J_{DT}	current density in direct tunneling
J_{FN}	current density in Fowler-Nordheim tunneling
k	wave number
k_B	Boltzmann coefficient
m	mass
m^*	effective mass
n_i	intrinsic carrier concentration
n_s	electron density at the silicon/silicon dioxide interface
N_A	acceptor density
N_A^-	density of ionized acceptors
N_D	donor density
N_D^+	density of ionized donors
N_{ox}	oxide charge density
N_s	silicon/silicon dioxide interface charge density
p	hole density
p_s	hole density at the silicon/silicon dioxide interface
P	pressure
P_0	reference pressure
q	unit charge
Q	charge
r	relative sticking coefficient
R	resistance
R_{it}	interface trap state resistance
R_s	series resistance
(s)	solid

t	time
T	temperature
V	potential, voltage
V_{bi}	built in voltage
V_{fb}	flat band voltage
ΔV_{fb}	flat band voltage shift
V_g	gate voltage
V_{ox}	voltage across the oxide
V_s	potential at the silicon/silicon dioxide interface
V_{trap}	potential of a interface trap level
X	reactance
Y	admittance
Y_m	measured admittance
Z	impedance
Z_m	measured impedance
δ	Dirac delta function
ϵ	dielectric constant
ϵ_{Si}	dielectric constant of silicon
ϵ_{Ox}	dielectric constant of silicon dioxide
Φ_M	barrier height at metal/oxide interface
Φ_{Si}	barrier height at silicon/oxide interface
ρ	charge density
ω	angular frequency

Abbreviations

ALCVD	Atomic Layer CVD
ALD	Atomic Layer Deposition
ALE	Atomic Layer Epitaxy
CMOS	Complementary MOS
CV	Capacitance - Voltage
CVD	Chemical Vapor Deposition

FET	Field Effect Transistor
FN	Fowler-Nordheim (tunneling)
IC	Integrated Circuit
Imag	imaginary part
LF	Low Frequency
LPCVD	Low Pressure CVD
MBD	Molecular Beam Deposition
MBE	Molecular Beam Epitaxy
MOS	Metal Oxide Semiconductor
MOSFET	Metal Oxide Semiconductor Field Effect Transistor
NMOS	n-channel MOS
NTP	Normal Temperature and Pressure
PECVD	Plasma Enhanced CVD
PEE	Plasma Enhanced Evaporation
PMOS	p-channel MOS
Real	real part
RF	Radio Frequency
RHEED	Reflection High Energy Electron Diffraction
RPECVD	Remote Plasma Enhanced CVD
RTD	Resonance Tunneling Diode
RTA	Rapid Thermal Annealing
RTO	Rapid Thermal Oxidation
SILC	Stress Induced Leakage Current
SOI	Silicon on Insulator
TEM	Transmission Electron Microscopy
UHV	Ultra High Vacuum
UPO	Ultrahigh vacuum Plasma Oxidation
XPS	X-ray Photo emission Spectroscopy

Contents

1	Introduction	1
2	MOS Structure	5
2.1	Silicon/Silicon Dioxide Interface	6
2.2	Tunneling, Leakage, Breakdown and Defect Generation	10
3	Fabrication of Thin Silicon Dioxide Films	16
3.1	Cleaning of the Substrate Surface	16
3.2	Thermal Oxidation.....	19
3.2.1	Deal-Grove Model	24
3.3	Other Fabrication Methods.....	26
3.4	Annealing	29
4	Electrical Characterization of Ultrathin Insulators	34
4.1	Oxide Charge.....	36
4.2	Impedance of Thin Oxide MOS Structure.....	37
4.3	Oxide Thickness	42
4.4	Surface State Density	45
5	Ultrahigh Vacuum Plasma Oxidation (UPO).....	49
5.1	Equipment.....	50
5.2	Electrical Characterization	51
5.3	Oxidation Rate.....	51
5.3.1	Experiment.....	52
5.4	Room Temperature Oxidation.....	58
5.4.1	Experiment.....	59
5.5	Effect of Oxidation Temperature on the Silicon Dioxide Quality	63
5.5.1	Experiment.....	63
5.6	The Effect of Post Oxidation Annealing	68
5.6.1	Experiment.....	69
5.7	Evaporation Technique.....	72
5.7.1	Experiment.....	73
6	Conclusions	78

1 Introduction

During 1994 research interests at the Helsinki University of Technology Electron Physics Laboratory started to shift increasingly from III-V compounds towards silicon based devices. With the growing interest to use our molecular beam epitaxy (MBE) system to fabricate silicon devices, it became obvious that, in addition to epitaxial semiconductor layers, it was also necessary to have some thin insulator films. This paved the way to our research on plasma oxidation in ultra high vacuum (UHV)^{1,2,3,4}.

Molecular beam epitaxy systems have been used in submicron MOSFET fabrication^{5,6}, but these processes usually do not include oxidation in the MBE. There has, however, been some work on cyclotron resonance plasma oxidation in UHV⁷ or some other oxidation/evaporation methods⁸, for instance. In our research the aim was to maximize the process purity and the quality of the materials produced by performing the oxidation in an ultra high vacuum MBE system. The use of an MBE system allows for highly accurate growth and there are almost no impurities at all. Our processing equipment also provides the possibility for *in situ* native oxide removal and device processing, which further improves process purity. This process not only allows us to fabricate high quality films, but also makes it possible to get a more reliable picture on the oxidation process on an atomic level.

The main impetus for our research was provided by the growing interest worldwide in silicon/silicon dioxide based nanodevices. The aim has been to fabricate ever smaller and faster devices and circuits. The decrease in the chip area, and by the same token in the processing linewidths, means that the vertical dimensions of the devices also have to decrease. This requires a fabrication method that produces very homogeneous and high quality Si/SiO₂ interfaces and SiO₂ layers with accurately defined thickness. A 50-nm channel length bulk MOSFET would have a 2-nm thick gate oxide for high performance applications. That would give a maximum electric field of 5 MV/cm in the oxide, about half of the breakdown field of good quality bulk oxide⁹. Normal operation of a MOSFET with an ultra-thin direct-tunneling gate oxide was first achieved in November 1995. In an n-MOSFET, with 1.5 nm gate oxide and 100 nm gate length, transconductance above 10 S/cm and current drive above 1 mA/μm was measured at

room temperature¹⁶. By decreasing the channel length of MOSFET to 80 nm and the oxide thickness to 3.5 nm, cut-off frequencies of about 100 GHz have been reached⁹. Other silicon based devices in which ultrathin oxides are needed include, for instance, memory structures based on silicon nanocrystals¹⁰.

The tightening of process tolerances in order to satisfy more stringent defect density and reliability requirements poses many difficult challenges. For purposes of fabricating faster sub-micrometer MOSFETs and quantum devices, the thickness of the silicon dioxide layer may be no more than a couple of nanometers. This greatly enhances the impact of surface roughness scattering on drain current and carrier mobility¹¹. The surface and the interface have to be flat on an atomic scale. Roughness may also enhance tunneling currents. Furthermore, roughness increases the amount of traps and charge at the interface, creates extra energy states, and leads to local variations in the electric field and in the channel charge. It is widely accepted that low frequency $1/f$ noise in MOS transistors is due to mobility or charge fluctuations arising from dynamic trapping and detrapping of channel carriers by interface traps.

As the dimensions are reduced to the level where quantum effects begin to appear, it also becomes possible to use silicon to fabricate versatile new devices, which could earlier only be made from compound semiconductors. This would mean increased process compatibility and reduced circuit prices. Research so far has produced such innovations as luminescent silicon/silicon dioxide heterostructures, for instance¹². Because the energy gap in silicon dioxide is above 8 eV and in silicon 1.1 eV and the affinity of silicon is over 3 eV higher than in silicon dioxide (Figure 1.1), the barriers can be much higher and quantum wells much deeper than in III-V heterostructures. This means there can exist more quantum states at the same time, and quantum phenomena can be utilized in room temperature. The state energies can be adjusted by changing the quantum well dimensions.

Research on silicon-based nanodevices has also involved some trials to fabricate resonant tunneling diodes (RTDs), but the resonant effect achieved has not been very strong¹³. The quality of the interface between these two materials is not as good as when using crystalline materials with low lattice mismatch, such as GaAs and AlGaAs. This means that there always exist interface states and consequently there are several

possible states from which and to which the carriers can tunnel. For this reason it is hard to see clear quantum effects. The strongest silicon based photon emission, also, has been reached by using porous or multicrystalline silicon: not silicon/silicon dioxide heterostructures. The origin of this luminescence is not clear, but it might be due to some three dimensional quantum structures, formed by nanocrystals.

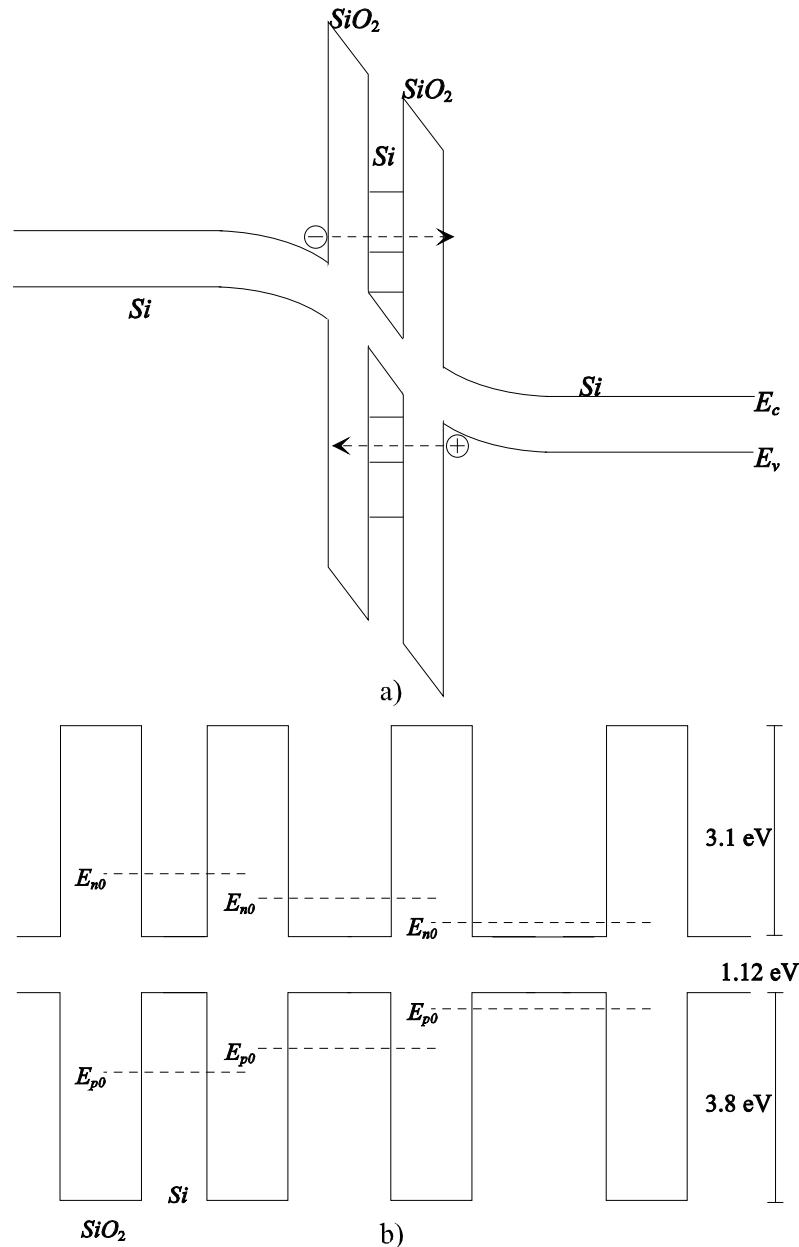


Figure 1.1 Schematic illustration of energy band structure of a) RTD b) silicon/silicon dioxide heterostructure. E_{p0} and E_{n0} represent the ground state energies in the potential wells.

Another major problem has to do with the fact that silicon dioxide is an amorphous material, and the average distances between silicon atoms in silicon lattice and in silicon dioxide are different. Because of the 'lattice mismatch' it is difficult to grow single

crystalline silicon on top of the oxide without holes through the oxide or in some other way so that the epitaxial layer can come into direct contact with the substrate. After that high temperature treatment is also usually needed to transform multicrystalline material into single crystal. Some experiments have, however, succeeded in growing single crystal silicon on SiO_x , where the oxygen content is about 20 at.% or less ($x < 0.25$)¹⁴. This, of course, means that the oxide properties are changed.

There has been intensive research on the outermost limit of the downscaling of silicon dioxide films. A decrease in device size, and the oxide thickness leads to an increase in the tunneling current. Tunneling can have a significant effect on the transistor operation. The stress induced leakage current (SILC) is considered to be the major limiting factor for the downscaling of tunnel oxides. If the devices operate at elevated temperatures, SILC increases. Present estimates are that the film thickness can not go much below 2.2 nm (1 V supply voltage)¹⁵. After that the reliability decreases too much. Various fabrication procedures and the set-ups for experimentation result, however, slightly different properties. Device structures can also be modified to improve performance without extended downscaling. One of the ways of tackling the problems occurring in the ultrathin region has, in fact, been to change the MOSFET structure rather than just scale down the dimensions¹⁶. The drain and source wells can be made deeper, which decreases sheet resistance. Smaller gate decreases leakage current and increases drain current. Nonuniform channel doping in both vertical and lateral directions can be used to minimize the short channel effects, while low temperature CMOS and double gate MOSFETs can potentially lead to the outermost limit of silicon scaling⁹. SOI and SiGe devices offer performance and power advantages over bulk CMOS without channel length scaling. The demands of the rapidly developing semiconductor industry will, however, still require even smaller dimensions and thinner silicon dioxide films.

2 MOS Structure

The fabrication of good quality insulator layers requires a good molecular structure and the complete absence of impurities: there may be no free bonds, traps or other defects that might affect the oxide charge or conductivity. The interface between silicon and silicon dioxide should also be as abrupt as possible, and there should be no interface states. Electron mobility, for instance, has been shown to correlate directly with oxide/channel interface roughness¹⁷.

The basic MOS structure (Figure 2.1) consists of a metal (often aluminum) or heavily doped polysilicon layer, insulator (silicon dioxide or silicon dioxide and silicon nitride layers) and doped silicon.

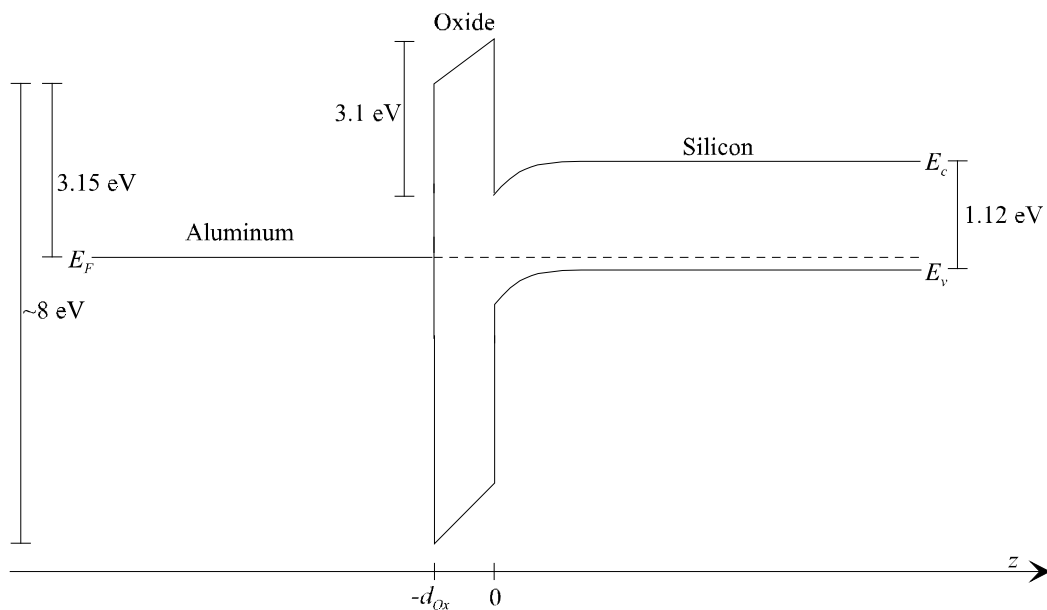


Figure 2.1 Energy band structure of an aluminum/silicon dioxide/p-type silicon structure with zero bias.

Oxide defects are traditionally divided into four general classes¹⁸:

- fixed oxide charge
- mobile ionic charge
- interface-trapped charge
- oxide trapped charge

Interface traps communicate directly with the underlying silicon, whereas oxide traps and fixed charge do not. There are also so-called ‘border traps’¹⁹, a term applied to traps lying within about 3 nm of the silicon/silicon dioxide interface and in communication with the silicon. These are near-interfacial oxide traps that can either rapidly or slowly exchange charge with the silicon. They play an important role in determining low frequency (1/f) noise levels in MOS transistors. Border traps can act like either interface traps or bulk oxide traps, depending on the bias and the time scale. They include so-called ‘slow states’²⁰, ‘anomalous positive charge’²¹ and rechargeable E’ centers²². It must be noted that all defects in ultrathin oxides may act as border traps, since the film thickness is on the 3-nm scale, or even lower.

Impurities also have an impact on energy band discontinuity at the silicon/silicon dioxide interface²³. They induce changes in the band structure and have been found to bring about a decrease in conduction band off-set and an increase in valence band off-set. The evidence suggests these changes are almost identical in thermal oxide²³. The magnitude of the energy gap is thus not changed, but the bands are bended. The variations in the energy band structure between the bulk oxide and the interface region depend of course on the impurity concentration and the oxide quality. In some conventional dry oxidation experiments (below 800 °C) a typical value for the off-set shift has found to be about 0.2 eV, on both silicon (100) as on silicon (111) surfaces^{24,25}. This value can be reduced, however, by using ultra clean processes²³.

2.1 Silicon/Silicon Dioxide Interface

Almost all silicon wafers that are used in microelectronics are single crystalline. Silicon dioxide, on the other hand, is amorphous. The average distances between atoms also differ from one another. Once the silicon surface is oxidized, the interface between these two materials can no longer be perfect. The volume per silicon atom in silicon dioxide is about twice that in silicon crystal, which means that silicon atoms are inevitably emitted from the interface to release stress during oxidation²⁶. The atomic scale structure of the interface between crystalline silicon and its amorphous oxide

remains controversial²⁷. However, a very high degree of perfection has been achieved in practice, which is evidenced by the very low density of defects observed at the interface.

Depending on the oxidation procedure interface roughness between silicon and silicon dioxide layer has been reported to be up to 1 - 1.6 nm^{28,29}. Roughness is also affected by surface orientation and film thickness. On (100) surfaces there is less roughness than on (111) surfaces, but it increases with increasing thickness (film thickness 1 nm, roughness about 0.06 nm; film thickness 2 nm, roughness 0.14 nm³⁰). On (111) surfaces, however, the range of roughness depends hardly at all on thickness (about 0.3 nm³⁰), but the horizontal size of the protrusions increases with film thickness.

Several assumptions have been made about the possible interfacial structure of the silicon/silicon dioxide interface. Non-reacted silicon - silicon bonds have been observed at the interface by means of Auger spectroscopy. On this basis it can be assumed that there is at least one monolayer of silicon dioxide into which some amorphous silicon is mixed³¹. Auger analysis has also shown that there is an interface layer in which there is less oxygen and usually some silicon clusters³². The same effect has been reported on the basis of ellipsometric measurements³³, where the change in the refractive index shows lower oxygen contents. There have also been made some measurements of the structural nature of silicon/silicon dioxide interface (oxide thickness 5 - 27 nm) by using infrared absorption associated with oxygen vibrational modes²⁹. These measurements indicated that the interface difference is mainly due to stoichiometric changes, not to stress or densification.

The interface region width depends not only on the substrate and oxide layer fabrication procedure, but also on the measurement techniques used. This is because there are in fact two kinds of distinct regions near the interface: compositional and structural transition layers³⁴. The compositional transition occurs abruptly within one silicon atomic layer, which due to cross linking consists of intermediate oxidation states³⁵, such as Si¹⁺, Si²⁺ and Si³⁺. The structural transition, by contrast, takes a much greater extent from the bulk silicon to bulk silicon dioxide. On the silicon side it consists of one or two silicon monolayers in which the atomic arrangement differs from the bulk³⁶. Computer simulations of oxygen transport into silicon and the formation of bonds

indicate that within the first monolayer there are extra silicon atoms due to stress, before oxygen diffuses into the silicon network^{27,37}. The same simulations provide a partial explanation for the low interface defect densities²⁷: during oxidation the oxygen atoms can momentarily be bonded to three silicon atoms, which allows the interface to evolve without leaving dangling bonds.

On the silicon dioxide side the transition layer can be as wide as 1 nm, but there remains some uncertainty about its structure. This may have to do with the difficulty of preparing uniform ultrathin oxide layers³⁴. It has been proposed that the main structural difference between the bulk and interfacial region is the difference in the intertetrahedral Si-O-Si bond angle (Figure 2.2), otherwise the structure is the same corner-sharing tetrahedron network (i.e. continuous random network of 4-, 6-, 7- and 8-member rings of SiO₄ tetrahedra, joined by bridging oxygens³⁸). In calculations where experimental XPS data have been compared with the calculated energy levels of the structure as in Figure 2.2, the calculated value for the reduced Si-O-Si bond angle has been about 135° - 140°; the intratetrahedral bond angle used was 109.47°³⁴. In the oxygen interface region, within about 3 nm of the interface, the distribution changes so that oxide consists of Si₂O₃, SiO and Si₂O³⁸. Another possibility is that the coordination for the corner-sharing tetrahedron network is changed, and the intratetrahedral bond angle is not reduced. Then there are overcoordinated oxygen centers (O₃⁺, probably associated with O⁻)³⁹.

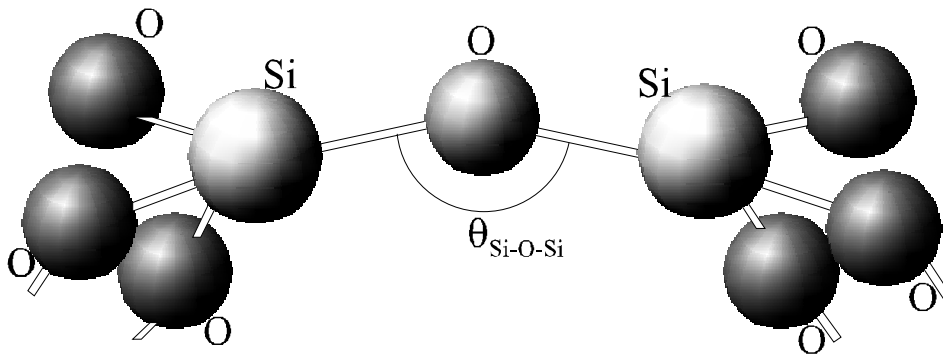


Figure 2.2 Schematic illustration of SiO₄ tetrahedras joined by bridging oxygen.

The energy band transition from silicon to silicon dioxide cannot either occur without some transition region. Calculated values for the valence and conduction bands indicate

that it takes about 0.4 - 0.5 nm before the energy gap corresponds to the bulk values, even for an ideal interface between silicon and crystalline silicon dioxide⁴². This is consistent with simulations²⁷ indicating that the stoichiometry of the first 0.5 nm of oxide is close to SiO.

Results have been published according to which there might be some crystalline silicon dioxide near the interface, maintaining an epitaxial relation with the substrate⁴⁰. However, it is estimated that the concentration of these crystallites is low (possibly a few percent) and they are probably unstable⁴¹. Thus, the possibility to form a totally crystalline transition layer is highly dependent on the oxidation process. One very prominent suggestion is that in an optimal case, the transition layer between crystalline silicon and amorphous silicon dioxide could be an ordered, about 0.5 nm-thick crystalline oxide, tridymite (Figure 2.3), a stable bulk form of SiO₂. This stability is assumed to decrease, however, when the layer thickness exceeds 1.5 nm⁴². There is also some evidence from transmission electron microscopy (TEM) studies which supports this suggestion⁴³.

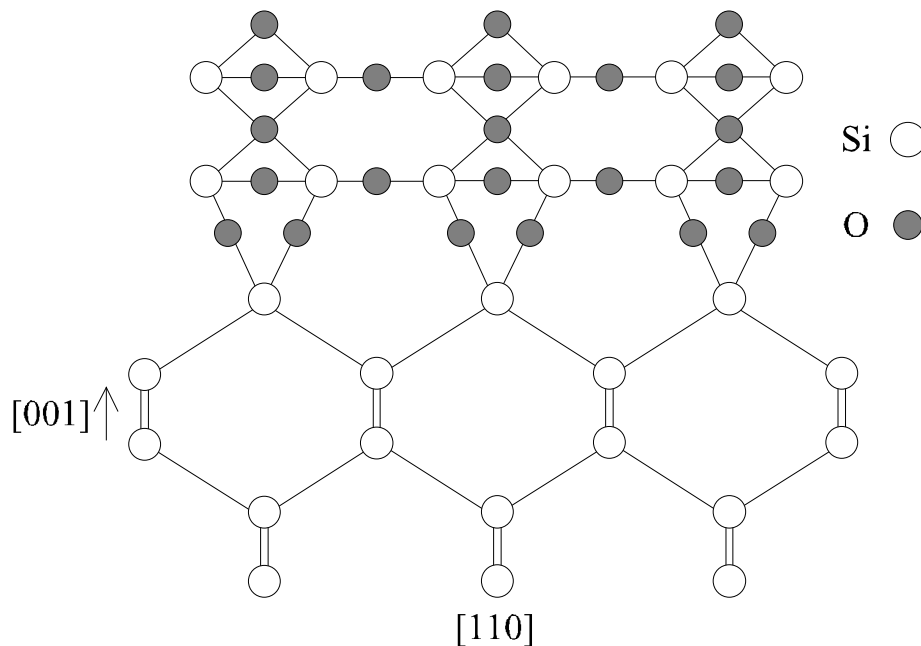


Figure 2.3 Schematic illustration of silicon/silicon dioxide tridymite structure⁴³.

First-principle calculations^{44,45} have proven that in ideal structures the interfaces between silicon and tridymite can be fully relaxed, so that there will be no electron

states in the silicon gap. In these calculations it has been assumed that a tridymite is attached to the silicon (100) surface and the strain is localized to a transition region at the interface and does not persist in the bulk oxide⁴⁵. The width of the transition region, 0.5 nm, accommodates the required change in silicon density. The amount of suboxide of 1.5 monolayers is compatible with the best experimental results. The structure corresponds to a local energy minima, but there might also be other minimas that are even lower⁴⁵. Therefore in order to fabricate such crystals, the process has to be very accurately controlled.

2.2 Tunneling, Leakage, Breakdown and Defect Generation

As was mentioned earlier, tunneling through the oxide becomes significant when the film thickness decreases to about 3 nm. This may cause problems in device operation and also during characterization. In the ultrathin region the direct tunneling current increases by about one order of magnitude when the thickness of the silicon dioxide layer decreases by one nanometer. For the direct tunneling (Figure 2.4a) the current density can be evaluated from the approximate formula⁴⁶

$$J_{DT} = \frac{q^2(2\Phi_M - V_{ox})}{4\pi h d_{ox}^2} \exp\left[-\frac{4\pi d_{ox}}{h} \sqrt{qm^*(2\Phi_M - V_{ox})}\right], \quad (2.1)$$

where q is unit charge, h Planc constant, m^* electron effective mass, Φ_M the barrier height at the metal/oxide interface, V_{ox} the voltage across the oxide and d_{ox} is the oxide thickness.

The so-called Fowler-Nordheim (FN) current might cause more trouble (Figure 2.4b). The magnitude of it can be evaluated by using the following approximate formula⁴⁷

$$J_{FN} = \frac{q^3 V_{ox}^2}{8\pi h \Phi_M d_{ox}^2} \exp\left[-\frac{8\pi \Phi_M d_{ox}}{3hq V_{ox}} \sqrt{2m^* \Phi_M}\right], \quad (2.2)$$

In addition, this tunneling current oscillates to some extent as a function of voltage. This could, in theory, affect the measured impedance. However, the risk is not very great as the oscillation is quite modest and starts at relatively high voltages. Often the oxide field in ultrathin oxides exceeds the breakdown limit even before the Fowler-Nordheim oscillation has started. It has been estimated⁴⁸ that direct tunneling dominates for oxides thinner than 4.2 nm and FN-tunneling for oxides over 5.1 nm. This phenomenon, however, has been used to determine the oxide thickness as the oscillation period is dependent on the tunneling barrier thickness⁴⁹. FN-current causes also some stress on the oxide, producing an oxide charge that becomes unstable upon continuing electrical measurements⁵⁰. This further causes instability, possibly making it more difficult to define the material parameters.

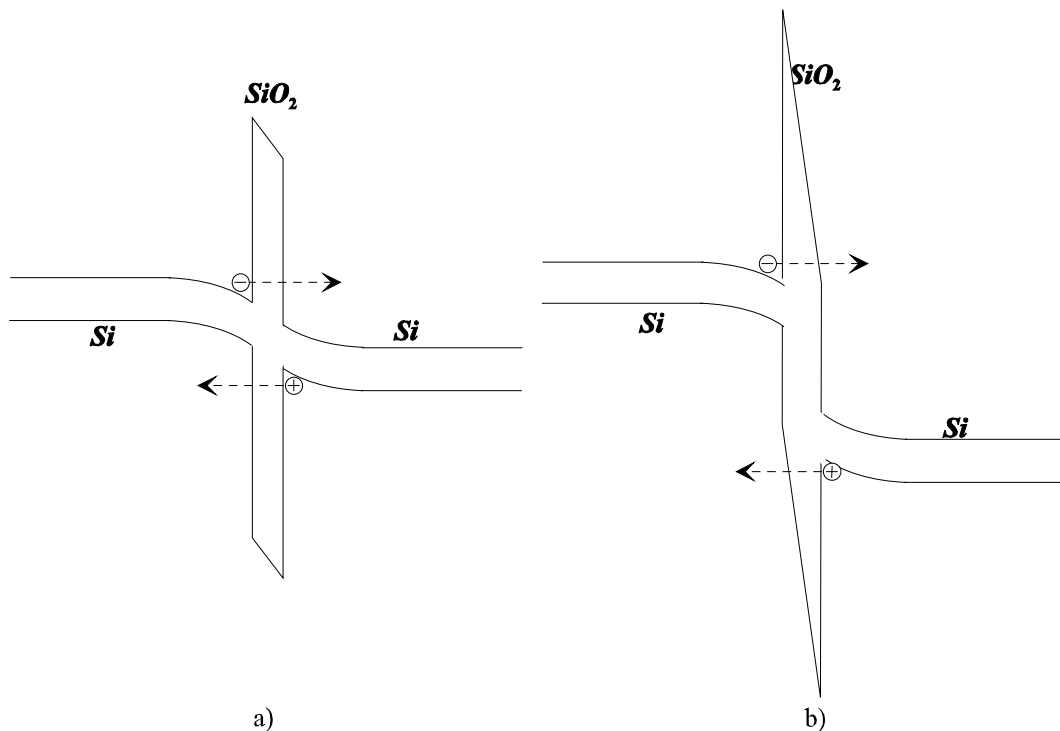


Figure 2.4 Tunneling in an MOS structure: a) direct tunneling b) Fowler-Nordheim tunneling.

In real devices tunneling is not the only mechanism that allows current passing through the oxide films. If the difference between the theoretical tunneling current and the measured value is significant, this means the structure is non-ideal. Impurities increase the conductivity of the oxide and form extra energy states, increasing the probability of tunneling. Near the interfaces the molecular structure is not as perfect as in bulk

material, which also affects the energy band structure. That may significantly reduce the potential barrier height. Various kinds of lattice defects may create conductive channels through the insulator. Because the dimensions are small, any kinds of impurities may diffuse, or spike, through the layers, especially during high temperature processing. The probability of a major defect occurring within the device obviously increases with increasing device area. This means that also the leakage current increases faster than the area. One example of this phenomenon is shown in Figure 2.5. We measured the forward current density in a larger area device to be considerably higher than in a smaller one, even though on reverse bias the situation was the opposite which indicates that the smaller device was not particularly good.

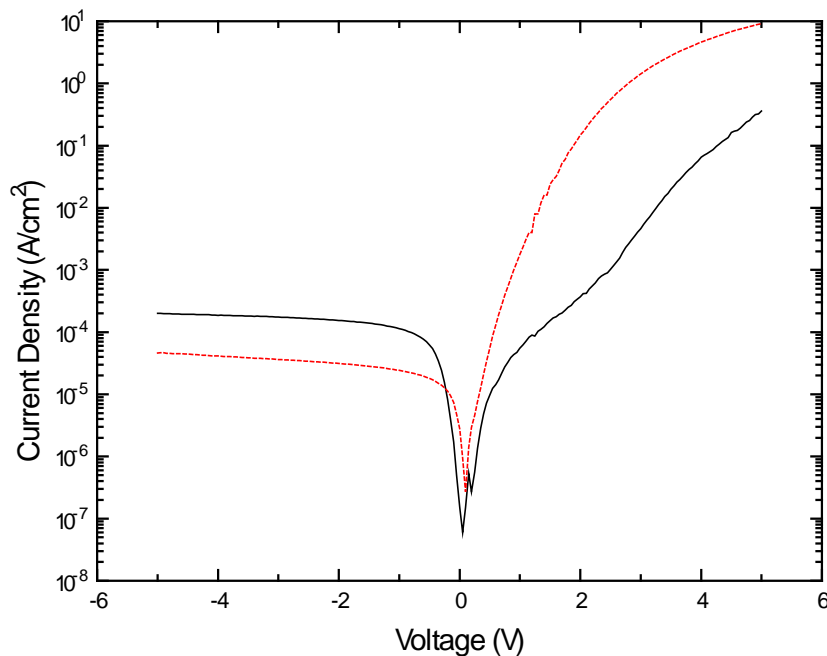


Figure 2.5 Measured average current density through MOS structures with ultrathin silicon dioxide layer; device area $(100 \mu\text{m})^2$ solid black line, $(500 \mu\text{m})^2$ dashed red line.

The geometry of the device can play quite an important role, particularly when the structures are small. The charge tends to increase and the electric field to get higher towards the edges and in the corners, which means that a significant proportion of the current may in fact be going through the edges of the device rather than evenly through the whole device. At the interfaces the likelihood of impurities occurring is greater, which may also increase conductivity. An example of this can be seen in our measurements presented in the Figure 2.6.

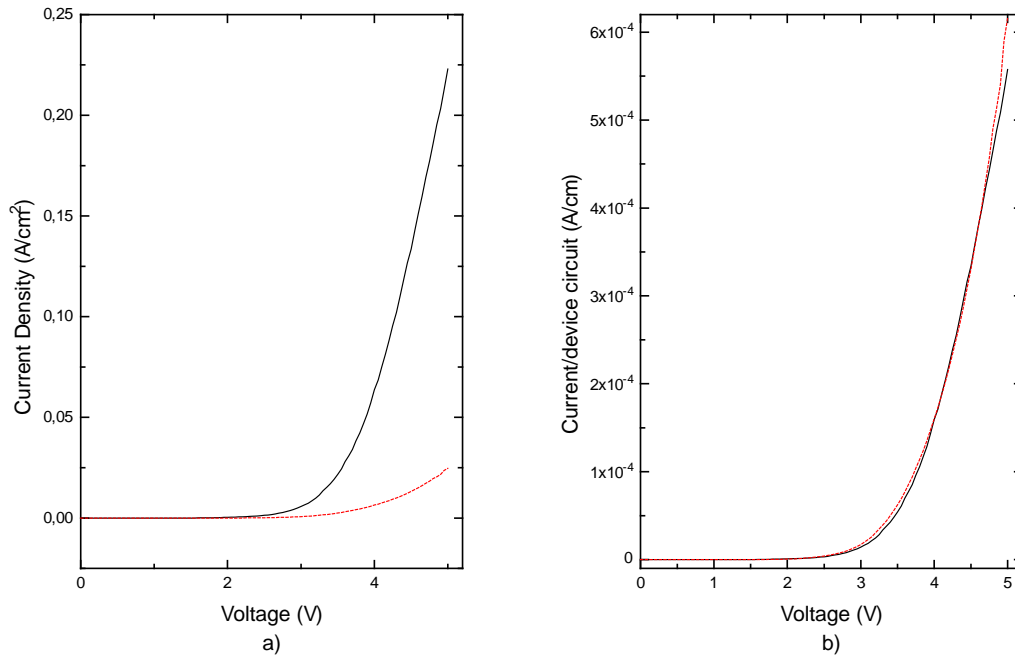


Figure 2.6 Measured a) average current density b) current divided by the length of device periphery, through two MOS structures. (Solid black line: Device area $(100\ \mu\text{m})^2$. Dashed red line: Device area $(1\ \text{mm})^2$.)

When the tunneling current through ultrathin oxides is measured and compared to simulated ones, it is possible to see indications of trap assisted tunneling components in addition to direct tunneling⁵¹. It is interesting that even when the number of traps is eliminated as far as possible by careful processing, trap assisted tunneling can still be seen in ‘thicker’ oxides, but when the film thickness drops below 2 nm, the only mechanism seems to be direct tunneling⁵¹. Tunneling straight through the film is then therefore energetically more favorable compared to via-traps tunneling. The other possibility, i.e. a sharp decrease in the amount of traps, does not appear very plausible. However, when the film thickness is so small the current level becomes significant anyhow even under normal device operating conditions.

One of the most important material parameters of insulators is the breakdown voltage, or breakdown field. Dielectric breakdown of MOS devices is important both for reliability considerations⁵² and in the determination of suitable operation voltages. It is of particularly great significance in the context of ultrathin insulators. Conventional, ‘thick’ insulator layers made from silicon dioxide have a breakdown field of about

10 MV/cm⁴⁶. This may seem quite high, but expressed in other units it is 1 V/nm. It follows that when the MOS insulator is only 2 – 3 nm, the maximum voltage that can be used across the oxide is no more than a couple of volts. It has also been noted that the strength and stability of these ultrathin oxide films is often not as good as the properties of bulk oxide. This fact must be taken into account in device design.

In ultrathin layers, the breakdown effect differs from the one seen when measuring thicker layers. Because the interface region, for instance, represents a significant portion of ultrathin oxide films, this has an impact on such properties as reliability. It is often possible to see a phenomenon called ‘soft breakdown’, i.e. a ‘smaller’ breakdown occurs before the actual breakdown point. It has been suggested that when the bias exceeds the soft breakdown value, electron transport through the film is mainly due to multiple tunneling via generated electron traps⁵³. The breakdown effect in general has been related to electron trapping⁵⁴, hole trapping⁵⁵ and interface state generation⁵⁶. There are some results according to which interface state generation during voltage stress is quite comparable in thermal oxides and N₂O oxides, indicating that these state generations are not necessarily the main reason behind the dielectric breakdown⁵⁷. In constant current injection stress measurements, no widening of the breakdown points (as a result of the spreading of thermal faults) has been observed⁵³. If, however, the current is not limited, it is possible to see a roughening effect. TEM studies⁵⁸ have found that in breakdown spots the silicon melts, deoxidization occurs and the interface is locally shifted. It has been reported that the breakdown is due to a buildup of positive charge at the cathode interface in localized areas⁵⁵. This, it is assumed, consists of holes generated by impact ionization in the oxide and drifted to the cathode. This increases the local field and, through barrier lowering, the current density, and further accelerates the buildup until a very brief runaway process brings the oxide to destructive breakdown.

When holes are trapped near the interface and electrons are subsequently injected, recombination occurs. From the energy released in this process, interface states may be formed²¹. It has been found that the density of these states is dependent on the number of recombination events, but independent of the average oxide field, and therefore the electron energy. Superimposed upon this effect is the generation of interface states due to hot electrons above the heating threshold (about 1.5 MV/cm). Because of this the

defect generation rate also increases sharply ²¹.

Thin oxides stressed at high voltages have increased low level pre-tunneling leakage currents ⁵⁹. The leakage seems to be localized, not uniform. It is assumed to be due to tunneling through stress-generated positive centers ⁶⁰, or local lowering of tunneling barrier due to the generation of local defects ⁶¹. Higher pre-tunneling currents and lower breakdown distributions have been related to oxygen deficiencies at the silicon/silicon dioxide interface ⁶². The intrinsic oxide decomposition reaction $\text{Si} + \text{SiO}_2 \rightarrow 2\text{SiO}$ at the interface is supposed to be nucleated at existing defect sites prior to the growth of physical voids. These defects can become electrically active during annealing at temperatures lower (e.g. below 900 °C in UHV) than needed for void formation. Unless sufficient O₂ is available, this can lead to low field dielectric breakdown ⁶².

It has been found that an increase in pre-tunneling currents correlates with the number of traps generated within the oxide by high voltage stress ⁵⁹. Low field leakage currents introduced in the thin oxide during high field stress are related to defects produced by hot electron transport ⁶³. The generation of neutral electron traps is assumed to be the dominant cause of this phenomenon. Atomic hydrogen exposure causes similar leakages, supporting the conclusion that currents are related to hydrogen induced defects ⁶³. Oxide trap distribution is relatively uniform, and trap density increases as the cube root of the fluence of electrons passing through the film ⁵⁹. The trapped electrons in stressed oxides seem to be located primarily at border traps: it has been estimated that the centroid of the trapped positive charge lies as close as 1.2 nm to the silicon/silicon dioxide interface ⁶⁴. Capacitance-voltage (CV) characteristics analysis of thin oxides stressed at Fowler-Nordheim region have also shown slow traps quite near the interface (charge centroid located up to 2.5 nm from the interface) ⁶⁵. The interface trap density increases as the square root of the stress fluence ⁵⁹. The high electric fields near the interface and/or reduced barriers to local tunneling, or charge injection caused by hole trapping, may trigger border trap formation as a natural response to increasing local electric fields ⁶⁴. It is, however, also possible to see an annealing effect of the defects with increasing temperature ⁶⁶.

3 Fabrication of Thin Silicon Dioxide Films

3.1 Cleaning of the Substrate Surface

On the surface of silicon wafers, along with possible contamination, there is a native oxide layer whose thickness is typically a couple of nanometers. This native oxide has to be completely removed in order that a good quality thin oxide film can be produced on silicon surface. It is also required that the sample surface is flat on an atomic scale, because the thin oxide morphology seems to reflect the original surface of the wafer⁶⁷. It is also important that the substrate, its cleaning procedures and other processing stages are carefully selected. It has been shown⁶⁸ that oxide defects in thin films can be associated with substrate defect: the defect density on Czochralski -silicon can be nearly one order of magnitude larger than that for Float-Zone silicon. However, these defects can be removed by a sacrifice oxidation procedure, in which the oxide thickness is at least 0.4 μm . Etching of the substrate surface does not alone seem to suffice⁶⁸.

Usually the wafers are first cleaned by using some chemical cleaning procedure, such as so-called RCA cleaning:

- 5 minute ultrasonic in acetone
- 5 minute isopropyl alcohol
- 10 minutes in $\text{H}_2\text{O}:\text{NH}_4\text{OH}:\text{H}_2\text{O}_2$ (5:1:1) at 80 °C
- 10 minutes in $\text{H}_2\text{O}:\text{HCl}:\text{H}_2\text{O}_2$ (5:1:1) at 80 °C

Better results have been achieved by using⁷⁷:

- 10 minutes rinsing in deionized water
- twice 5 minutes in methyl alcohol agitation
- 15 minutes boil in trichloroethylene
- twice 5 minutes in methyl alcohol agitation
- rinsing in water
- 10 minutes in 69% HNO_3 in 130 °C
- 10 s dips in 2.5% HF followed by rinsing in water until the surface becomes

hydrophobic

- 10 minutes in 25% NH₄OH: 30% H₂O₂:H₂O (1:1:3) in 90 °C
- 10 s dip in HF
- rinsing in water
- 10 minutes in 37% HCl:H₂O₂:H₂O (3:1:1) in 90 °C
- 10 minutes rinsing in water

Chemical cleaning procedures are used to remove all unwanted impurities from the surface. The resulting surface is usually of a good enough quality for the fabrication of conventional semiconductor devices. After chemical cleaning, it is possible to passivate the surface by HF treatment, for instance, which will prevent reactions with oxygen in the air or some other oxidizing species in the environment. This passivation, however, is seldom perfect and even at best will not last for long. Recently it has also been demonstrated⁶⁹ that the presence of a chemical oxide barely affects growth kinetics or uniformity.

The best way to remove the oxide is to do it *in situ*. This means it is done immediately before the beginning of oxidation and in the same processing equipment, reducing the risk of sample contamination. Excellent results have been reported with this method (e.g. thickness control of the layers fabricated and interface smoothness)¹⁷. The most common methods of *in situ* native oxide removal are plasma etching or the heating of the sample in an oxygen free environment (in UHV or in inert gas, depending on the process equipment). Since plasma etching is expected to roughen the surface, the heating procedure is often preferred. During heating following reaction will occur⁷⁰



and reaction products will desorb from the surface. *In situ* plasma cleaning with hydrogen has also been found to be suitable for silicon MBE processes (e.g. ref.⁷¹).

Studies of oxygen desorption from silicon surfaces in UHV conditions have found that the desorption starts inhomogeneously, mainly at defect sites⁷². Voids are therefore formed at these sites. Low co-ordination number silicon monomers transform to be free

mobile monomers, which diffuse (within the voids) and are incorporated into a reactive oxide complex at the silicon/oxide boundary. The reactive SiO_x complex and the addition of silicon atoms can then form $\text{SiO}(\text{g})$, which desorbs from the surface⁷². This reaction consumes silicon, which means there will form a pit into the silicon surface at the defect site.

An atomically flat silicon surface can be produced by flashing (heating rapidly) the substrate to $1200\text{ }^\circ\text{C}$ ⁷³. Even though the silicon melting point is around $1400\text{ }^\circ\text{C}$, its surface may start (incompletely) to melt at around $1150 - 1200\text{ }^\circ\text{C}$ in ultrahigh vacuum conditions; there may occur some sublimation of silicon even at lower temperatures (though still above $1000\text{ }^\circ\text{C}$)⁷⁴. The use of high temperatures seems to smoothen the surface. Interface roughness has been measured both from RCA cleaned and from samples that were RCA cleaned and flashed at $1200\text{ }^\circ\text{C}$ in UHV before thermal oxidation at $900\text{ }^\circ\text{C}$ ⁷⁵. It was found that if only chemical methods are used to clean the substrate, roughness increases as the layer thickness decreases from about 10 nm , reaching its maximum at a layer thickness of 4 nm . In UHV cleaned samples, however, roughness decreases with decreasing layer thickness. It is, however, not always practical or even possible to use such high temperatures. It has been shown that at these temperatures oxygen vacancies and charge traps are formed and there might exist some gettering which roughens the silicon/silicon dioxide interface⁷⁶.

In our case the processing equipment is a UHV chamber. Thus the native oxide can be removed simply by heating the sample. We can verify the desorption of the silicon dioxide layer by monitoring the surface by RHEED (Reflection High Energy Electron Diffraction): when the surface turns into a clean, monocrystalline silicon surface, the RHEED pattern becomes clear. Even a thin native oxide layer totally prevents the pattern to be formed. We have used two different native oxide removal procedures. The first consists simply in heating the sample to $875\text{ }^\circ\text{C}$ at $10^{-10} - 10^{-9}\text{ mbar}$ ¹. The native oxide is then removed in less than 15 minutes. The second procedure we have used has been shown to produce an excellent surface, entirely clean and flat by removing the native oxide by keeping the sample at $850\text{ }^\circ\text{C}$ and etching the oxide off by a low silicon flux⁷⁷. When some silicon is introduced into the surface in gas form, the reaction $\text{Si} + \text{SiO}_2(\text{s}) \rightarrow 2\text{SiO}(\text{g})$ happens on the surface, not in the silicon/silicon dioxide interface. For this reason bulk silicon consumption should be lower than in conventional methods.

This probably explains why this method produces quite flat surfaces, even though the temperature is not that high. In fact, it has been suggested that even 780 °C may be enough, given the background pressure of $7 \cdot 10^{-11}$ mbar⁷³ and the chamber pressure of $1.5 \cdot 10^{-9}$ mbar during silicon deposition. In our experiments²⁻⁴ we have used this lower temperature and seen that the native oxide is removed quite quickly, within one minute.

3.2 Thermal Oxidation

Thermal oxidation is the most common, and also the simplest way to produce silicon dioxide layers. In microelectronics device processing the best silicon/silicon dioxide interfaces are fabricated by thermal oxidation of (100) silicon⁷⁸. When the sample is heated in an oxygen-containing atmosphere, its surface becomes oxidized. The oxygen is normally introduced in gas form ('dry oxidation'), but sometimes water vapor ('wet oxidation') is used. It is also possible to use oxygen plasma, which increases the oxidation rate, but in this case the process is more sensitive and it might increase interface roughness and defect density (e.g.⁷⁹). This is because the plasma may also etch the surface.

The main process parameters affecting the oxidation rate, and also the oxide and interface quality, are temperature and oxygen concentration. An increase in either one of these also increases the oxidation rate and the maximum layer thickness that can be achieved. Oxidation temperature is the most important process parameter affecting oxide quality; the hotter the better. Oxidation at high temperatures minimizes interface roughness and produces the best quality material. In practice this is also seen in decreasing interface trap density⁸⁰.

The most usual reactions



are also reversible. If the desorption rate from the surface is high enough, this may increase surface roughness. The presence of hydrogen during oxidation, or annealing

oxides in a hydrogen-containing atmosphere, is usually considered to have beneficial effects on oxide quality. The addition of hydrogen has the beneficial property of rendering charge traps inactive⁴⁶. One must, however, be careful in adjusting the process, because there is evidence that the introduction of extra hydrogen into the oxide may also aid in the creation of interface traps⁸¹.

During the early stages of thin oxide growth the interface between silicon and its thermal oxide undergoes atomic scale roughening⁸². This reaches its peak and then begins to diminish as growth proceeds. This is also an indication of a nonuniform oxidation process: the growth of about 1 nm protrusions can be explained by a reduction in the interface reaction rate brought about by local stress effects⁸². At temperatures above about 950 - 975 °C, oxide films grow stress-free, which is attributed to the 'viscous flow' of the oxide⁸³. The temperature limit above which this is possible is called the 'viscous flow point'. Relaxation of the silicon dioxide layer by this viscous flow and the transition to diffusion controlled growth act to eliminate roughness. This is reported to peak at a layer thickness of about 4 nm⁸². Another defect-density-maximum has also been found in Czochralski silicon wafers⁸⁴, with the oxide defect density increasing with increasing film thickness between 5 and 40 nm, and decreasing above 80 nm. The dependence of defect density on thickness is probably much affected by the substrate, impurity concentrations and oxidation procedure: for instance the '60 nm defect maximum' has not been found when using Float-Zone substrates⁸⁴. It has also been noted that even thick oxides with seemingly low oxide defect densities do in fact contain many small defects. These, too, may have serious adverse effects on oxide reliability. This quality decrease is probably caused by oxygen molecules diffusing through the oxide⁸². With increasing thickness, the diffusion slows down. At the beginning, on the other hand, there is less oxide structure to be damaged in the first place.

First-principle calculations of stresses affecting the process have shown that a uniform oxide layer can be obtained with any thickness by thermal oxidation once a uniform surface oxide layer has been formed²⁶. Therefore the preparation of the initial surface oxide is crucial for obtaining a uniform surface oxide layer with atomically controlled thickness.

The 'rapid thermal oxidation' (RTO) process is in widespread use in the fabrication of thin oxide films. Typically this is done by oxidizing silicon with O_2 or by using N_2O . The results have been excellent, and the processes are now so refined that thickness can be controlled to the accuracy of a few Angstroms. On the thickness scale of 2 - 5 nm, thickness can be controlled within 10%⁸⁵. The temperatures used are typically between 700 - 1200 °C, and the oxidation time is around one minute. After the oxidation of ultrathin layers⁸⁵ the surface state densities can be as high as $10^{12} \text{ eV}^{-1}\text{cm}^{-2}$ but they can be reduced by annealing to the level of $10^{10} \text{ eV}^{-1}\text{cm}^{-2}$. In the RTO method, too, increasing temperature decreases leakage current and fixed charge density, but it has also been found to decrease breakdown voltage⁸⁶. This may be due to pinholes or to defects created by the high temperature processing into the oxide.

It is predicted that the interface quality declines when the oxidation temperature is below the viscous flow point of the oxide⁸². Using temperatures above this point helps to avoid mechanical failure effects in IC-processing. High temperatures, which are usually needed to fabricate good quality layers, also increase the oxidation rate and therefore make it harder to adjust the film thickness. However, good results have been achieved by using ultra clean processes and low partial pressures of oxygen in some inert gases like argon⁸⁷, or low pressure oxidation⁸⁸. These not only help to achieve more precise thickness control, but also a thickness uniformity may be improved¹⁷. This does not, however, mean that oxidation times would not be much shorter than in conventional thermal oxidation. The oxide properties also tend to differ, sometimes quite considerably, when the oxidation parameters are changed⁸⁹.

There has also been some research to study the use of temperatures lower than in RTO (in practice usually above 650 °C⁹⁰) in the fabrication of ultrathin silicon dioxide layers. However, at temperatures below 1000 °C there may occur stress-induced dislocations and edge cracking. The problems have been tackled by reducing oxidation times to no more than a few seconds³¹. On the other hand, oxidation at lower temperatures can produce layers comparable to high temperature oxides. As well as giving improved thickness control, low temperature processing also prevents degradation of device performance due to the redistribution of impurities and the formation of stress-induced defects in silicon⁹¹. There still are, however, problems with low temperature oxidation: the oxidation rate is slow, and the oxidation also saturates

too early to form a layer thick enough for microelectronics purposes. To increase the film growth rate, and especially to make it possible to fabricate slightly thicker layers, additional techniques have been developed. It has been demonstrated⁹² that by using high pressure oxidation (10 bar), the temperature can be lowered to 650 °C and the oxide still shows excellent characteristics. The use of activated oxygen and low pressure also makes it possible to use low temperatures (300 - 500 °C) and still produce high quality properties⁹³.

A typical way of increasing the growth rate is to use plasma instead of oxygen gas (or N₂O gas or water vapor). This allows the oxidation temperature to be dropped by a couple of hundred degrees, say to 600 °C^{94,95}, or if no more than about two nanometers are needed, even as low as 400 °C⁹⁶. This is due to the smaller size and increased reactivity of oxygen atoms as compared to the ‘traditional’ oxidizing molecules, molecular oxygen or water. To carry out oxidation at room temperature, one possibility is to use a low energy ion beam⁹⁷ or a multipolar electron cyclotron resonance source for plasma generation⁹⁸. One method that has been used to fabricate high quality silicon dioxide films at room temperature (up to 11 nm) is oxidation by O₂ cluster ion beams⁹¹. In this case less damage will be caused by the beam, since the acceleration energy is divided between the single atoms of the cluster. It follows that the atom interactions occur near the surface. Some annealing is needed after this process, but it has been found that 30 minutes at 400 °C should be enough⁹¹.

Nitrided Oxides

N₂O has recently become an increasingly popular oxidizing gas. The main benefits achieved with N₂O in oxidation, or post oxidation nitridation of conventionally oxidized layers, are the improved breakdown properties^{52,99}. There is less degradation under hot electron stress and resistance to hot carrier induced interface state generation also improves with increasing N concentration^{100,101}. It has been reported that increased nitrogen concentration in the silicon/silicon dioxide interface increases interface state densities, but decreases the tunneling current. This current reduction has been estimated to be about one decade at any film thickness from 2.5 to 5 nm¹⁰². This, it is assumed, is due to the increase in the area of the tunneling barrier (Figure 3.1). The growth rate is also significantly lower, helping with process adjustment^{31,100}.

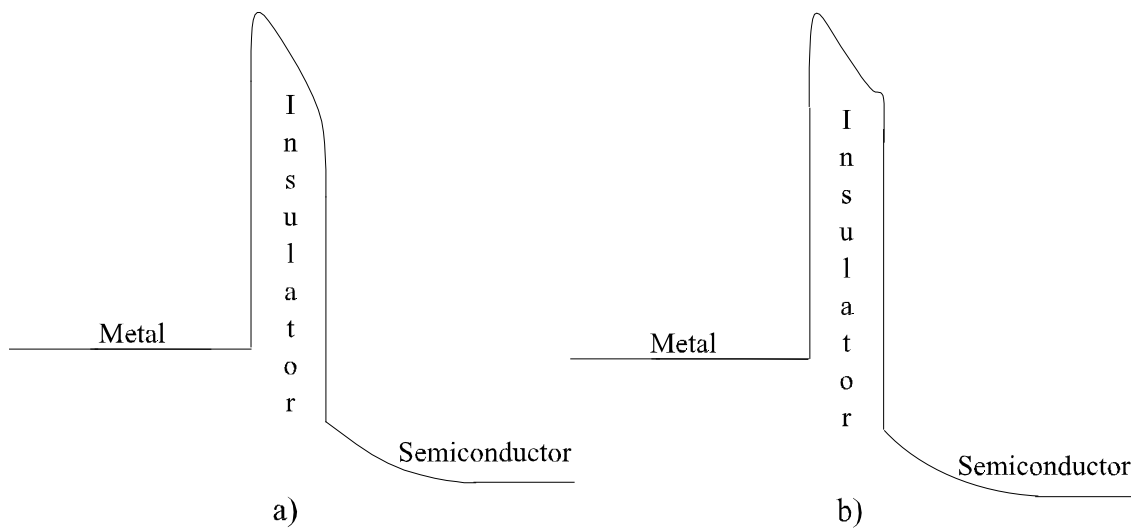


Figure 3.1 The conduction band structure of MOS structure with a) silicon dioxide b) silicon dioxide with nitrided interfacial layer tunneling barrier.

There are also some results which suggest that interface state generation during voltage stress is more or less comparable in thermal oxides and N_2O oxides⁵⁷. However, immunity to impurity state generation seems to be better with N_2O oxides under stress⁵². Improved radiation hardness has also been shown¹⁰³. This is the result of both suppressed electron trapping and enhanced hole detrapping due to interfacial nitrogen incorporation⁵². Impurity diffusion is decreased because the barrier properties to impurity penetration are better^{104,100} and the mobility under high normal field in the MOSFET channel is higher^{100,101,105}. It has been concluded that this is due to a combination of surface roughness scattering and the buried-channel nature of carrier conduction¹⁰⁵. Peak mobility, however, decreases with the increase in nitrogen concentration, at least as far as NMOS is concerned¹⁰¹. In PMOS even peak mobility has been found to improve¹⁰¹. However, the fixed oxide charge and interface state densities also increase with increasing nitrogen, and it is estimated that the optimal nitrogen concentration is about 3%¹⁰¹. There is also some evidence that oxynitride film thickness may be more homogenous than in silicon dioxide films¹⁰⁶.

3.2.1 Deal-Grove Model

In the conventional, so-called Deal-Grove model used for oxidation rate calculations in thermal oxidation^{46,107}, an oxygen flux creates an oxygen concentration C_0 on the surface. Oxygen diffuses through the existing silicon dioxide layer (thickness z) into the silicon. When the oxygen concentration in the silicon/silicon dioxide interface is C_S , the flux is defined as

$$F_1 = D \frac{dC}{dz} \approx D \frac{C_0 - C_S}{z}, \quad (3.1)$$

where D is the diffusion coefficient. The oxygen molecules coming to the silicon/silicon dioxide interface react with silicon at a rate that is proportional to the concentration. This requires the flux

$$F_2 = kC_S. \quad (3.2)$$

When

$$F = F_1 = F_2, \quad (3.3)$$

the oxidation rate is

$$\frac{dz}{dt} = \frac{F}{C_1} = \frac{C_0}{C_1} \frac{D}{z + D/k}, \quad (3.4)$$

where C_1 is the concentration of oxidizing molecules in the silicon dioxide. The time needed to grow a layer of thickness z can be calculated from equation (3.4). The result is

$$t = \frac{1}{B} \left[z^2 - d_0^2 + A(z - d_0) \right], \quad z > d_0, \quad (3.5)$$

where d_0 is the oxide thickness at the beginning. The parameters $B/A \equiv 2kC_0/C_1$ and

$B \equiv 2DC_0/C_1$, which are usually referred to as linear and parabolic rate constants, possess the temperature dependence $\exp(-E_A/k_B T)$ ⁴⁶.

The Deal-Grove model assumes that there always exists an initial oxide layer through which the oxidizing molecules have to penetrate and that this diffusion process is the main factor limiting the growth (*diffusion limited growth*). The parabolic growth behavior presented in equation (3.5) has been verified down to an oxide thickness of 5 nm ⁸⁷. As the thickness of the insulator layer needed for MOS devices decreases below 20 nm, some experiments ¹⁰⁸ (thermal oxidation in atmospheric pressure and temperatures around 1000 °C) have, however, indicated that the Deal-Grove model tends to underestimate the initial growth rate. Therefore an extension has been developed to this model ¹⁰⁸. Here the parameter k from equation (3.2) assumes the form

$$k \rightarrow rC_{\text{gr}}, \quad (3.6)$$

where C_{gr} is the concentration of so-called *growth sites* in the silicon/silicon dioxide interface. These are suggested to be sites formed by silicon suboxides, silicon clusters or interstitial atoms that can react with the oxidizing species.

The growth site concentration changes due to a generation mechanism (which is assumed to have an Arrhenius-type temperature dependence) and to recombination, in which the dominating factors are the concentration of these sites and the amount of the silicon atoms. The atom concentration is constant, therefore

$$\frac{dC_{\text{gr}}}{dt} = G_0 e^{-E_A/k_B T} - \frac{1}{\tau} C_{\text{gr}}, \quad (3.7)$$

and since we are talking about ultrathin layers, the growth rate is about the same as the linear rate constant

$$\left. \frac{dz}{dt} \right|_{x=\text{small}} \approx \frac{B}{A} = r \frac{C_0}{C_1} C_{\text{gr}} \approx e^{-E_A/k_B T} (l_0 + a_0 e^{-t/\tau}). \quad (3.8)$$

On this basis the initial growth rate is exponential rather than constant. Deductions made from simulations of quantum molecular dynamics suggest that there might indeed be a very fast oxidation step that precedes the diffusion limited growth. It has been estimated that the oxide layer has to be at least about 0.7 nm thick before the oxidation becomes diffusion limited ¹⁰⁹.

3.3 Other Fabrication Methods

One of the most common ways of fabricating thin silicon dioxide layers in microelectronics is by the means of *chemical vapor deposition* (CVD). In this method the oxide, and especially the electrical quality of the silicon/silicon dioxide interface, is not as good as in thermal oxidation, but the use of post oxidation annealing has produced some good results. In fact results have been published according to which the MOSFET transconductance can be even higher than in devices in which the gate oxide is fabricated by thermal oxidation. One possible reason for this is the smoother silicon/silicon dioxide interface, which is defined only by the original high quality silicon surface ¹¹⁰. It is also possible that the explanation lies in some resultant tensile stress in the oxide ¹¹¹. The higher oxide charge may also have some effect, rejecting carriers from the interface and thus decreasing the interfacial roughness scattering.

The *plasma enhanced CVD* (PECVD) method is widely used in microelectronics, not least on account of its low processing temperatures, usually below 350 °C. Using this method, silicon dioxide layers have been fabricated ¹¹² (at 250 °C) in which after annealing the surface state densities are in the range of $1 - 4 \cdot 10^{10} \text{ cm}^{-2} \text{ eV}^{-1}$. These values are comparable to thermally grown layers. Hydrogenated silicon oxide films have also been fabricated by PECVD at temperatures as low as 15 - 150 °C ¹¹³. However, this tends to produce a low oxygen concentration ($\text{SiO}_x\text{:H}$, $x < 2$) and a porous film structure. This precludes applications in devices, but it is still possible to take advantage of the fact that the layer will be easily oxidized ¹¹³.

In plasma assisted processes there is the risk that the excited oxidizing species may damage the substrate surface upon impact. For this reason there has been quite a lot of

interest in the *low pressure CVD* (LPCVD; e.g. 0.13 mbar¹¹⁴ or ≤ 0.01 mbar¹¹⁵) process. This process gives a slower growth rate and thus allows for easier control of the thickness of ultrathin layers. In this process the silane and oxygen are mixed prior to entering the reactor, and deposition is achieved by a pyrolytic process at the substrate surface¹¹⁴. In *remote plasma enhanced CVD* (RPECVD) the process temperature can be lower. The activation energy in the process is also much lower, about one order of magnitude, than in thermal oxidation⁹³. It must also be noted that if higher temperatures are used in RPECVD, some thermal oxidation will occur and affect film growth⁹³.

As-grown chemical oxides are uniform, have a low density, and contain a variety of defects such as Si-H bonds, dangling bonds, voids and OH groups, both in the oxide and at the interface. Therefore in order to reach gate oxide quality, it is necessary to have high temperature (800 - 1000 °C) process stages¹¹⁶. For example absorbed hydrocarbon contaminants dissociate and become chemically incorporated into thin oxides as additional silicon oxide, carbide, hydride and hydroxyl species. In UHV conditions these persist up to the SiO₂ desorption temperature (up to 850 - 1000 °C) so that once formed, these defects will be present in the final device structure¹¹⁶.

Deposited (LPCVD at 420 °C, as well as RPECVD at 300 °C) thin oxides show enhanced Fowler-Nordheim tunneling currents compared with thermal oxides, even after some high temperature (750 °C 40 minutes N₂-O₂¹¹⁷) annealing. They also exhibit very large electron trapping, with the traps reside close to the silicon/silicon dioxide interface¹¹⁷. It has also been reported that after annealing layer quality (oxide fixed charge and surface state density) can be comparable to thermal oxides, although the measured electron barrier height is still significantly lower¹¹⁴.

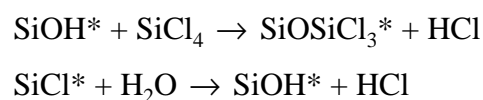
To maximize the benefits achieved by the low temperature CVD process and to combine those with the high quality interface properties achieved with thermal oxidation, a two-step process has also been used¹¹⁸. Here a low trap-density silicon/silicon dioxide interface has been fabricated by plasma assisted oxidation. After this the 'bulk oxide' is made by RPECVD deposition of silicon dioxide with device quality properties. The first stage, which produces an oxide film about 0.5 nm thick, also removes carbon residuals from the surface and prevents nitrogen incorporation at the interface.

As mentioned earlier in chapter 3.2, nitrogen incorporation into the oxide improves some insulator properties. CVD can also be used to grow these nitrided oxide films. The difference between these silicon dioxide and nitrided films are quite similar to thermal oxides: above 3% the positive oxide charge increases linearly with the nitrogen concentration, and above 4% there is also a sharp decline in mobility ¹¹⁹.

The silicon surface can also be oxidized *chemically*, for instance simply by keeping it in a hydrogen peroxide solution. This gives a very slow oxidation process, however, and it is only possible to produce very thin layers. This oxidation procedure has been tested by oxidizing monohydride terminated atomically flat silicon (111) wafers in an ultrapure 30% H₂O₂ solution at room temperature ¹²⁰. The first bilayer was oxidized within 30 minutes and then the process saturates; after two months only 60% of the next bilayer has been oxidized. This is obviously not a practical procedure for producing layers for most device applications: the tunneling probability is too high (chapter 2.2). Also, the average composition of the first bilayer is about SiO_{1.2}, and hence the layer quality does certainly not correspond to the properties of SiO₂.

Another way to fabricate thin silicon dioxide layers on silicon is to use the method of *anodic oxidation*. By using NH₄OH:H₂O ¹²¹ or 0.1M HCl ¹²², for instance, as electrolyte, it is possible to grow layers 1 - 10 nm thick. The resulting oxide thickness is strongly dependent on the visible illumination intensity ¹²³, which can be used to adjust the process.

The *atomic layer epitaxy* (ALE, also Atomic Layer Deposition (ALD) or Atomic Layer CVD) method has also been used to fabricate thin silicon dioxide layers. One of the binary reaction sequences used is (asterisks indicate the surface species) ¹²⁴



As each of the half-reactions is self-limiting, growth is controlled layer by layer. The resulting surfaces are very flat ¹²⁵, as flat as the original silicon substrate surfaces. The measured deposition per one processing cycle (at 330 °C and 1 - 13 mbar) is only

0.11 nm. This is quite interesting as one silicon dioxide monolayer should be more than two times that. This might mean that the layer is strongly ordered. The 'growth rate', i.e. the thickness of one deposited layer, decreases with increasing temperature (at least between 330 and 530 °C¹²⁵), which correlates with the thermal stability of the SiOH* surface functional groups¹²⁶. The other reaction used in ALE¹²⁷ is based on the deposition of CH₃OSi(NCO)₃ and H₂O₂. This process can be performed at room temperature with a deposition rate of 0.2 nm/cycle. After 100 cycles surface roughness is only ±1.0 nm, but the measurements indicate that the layers formed are silicon rich (SiO_{1.73}).

Thermal oxidation produces higher quality material than any other method, but all of these other methods are nonetheless usable, if not for MOSFETs or tunneling devices then at least for other purposes. Poor quality oxide may even result in higher intensity light emission for instance. Although the method of sputtering, for example, is quite a 'rough' way of making such fine structures as nanometer scale silicon/silicon dioxide devices, it has been used to fabricate some luminescent structures. The results seem very encouraging, but the usability of these layers in electronic devices remains to be established. In photoluminescent devices the non-homogeneity of the silicon dioxide and the grain boundaries may even serve to improve the luminescence. The main benefits of sputtering are its low costs and low processing temperatures. On the other hand, the interface state densities tend to be high and the breakdown fields too low¹²⁸. One of the methods employed in an attempt to improve the layer quality has been to carry out the sputtering in an oxygen-containing atmosphere and to use some post oxidation rapid thermal annealing (RTA). RTA has been found to drop the interface state densities to the level of $1.8 \cdot 10^{12} \text{ eV}^{-1} \text{ cm}^{-2}$ and by post metallization anneal (350 °C) to $3.5 \cdot 10^{11} \text{ eV}^{-1} \text{ cm}^{-2}$ ¹²⁹. A lowering of the sputtering pressure also seems to produce better layers.

3.4 Annealing

Device processing always involves in practice some post oxidation thermal treatments. Annealing after oxidation decreases the amount of oxide charge and the silicon/silicon

dioxide interface trap density. There are two basic annealing procedures: low temperature annealing, which typically continues for tens of minutes, and rapid thermal annealing at high temperatures.

Although the general preference is for high temperatures, it has been shown that temperatures as low as 300 - 400 °C are sufficient to decrease the surface state densities. Values between $1 \cdot 10^{10}$ and $5 \cdot 10^{10} \text{ cm}^{-2} \text{ eV}^{-1}$ have been measured after annealing from samples fabricated by the PECVD method ^{112,115}.

When studying high temperature annealings, it has been observed that a few minutes of annealing (at 900 °C ²⁸) reduces trap density exponentially as a function of time. Interface roughness, especially on the (100) surface, has also been reported to decrease dramatically during post oxidation annealing. In RTA experiments ¹²⁹ an increase in annealing time (50 - 500 s) has been found to be more effective in decreasing interface trap densities than an increase in annealing temperature (700 °C - 1000 °C). Oxygen defects and traps are removed ⁸⁶, and the strain between the silicon and oxygen bonds is probably also relieved. There is also some evidence that an increase in temperature decreases leakage current. The underlying mechanism may be an increase in barrier height. If, however, the temperature rises above 1000 °C, the barrier height might on the contrary decrease ⁸⁶.

Dealing as we are with ultrathin layers and processing in high vacuum conditions, high temperature processing gives rise to certain problems. Diffusion or spiking may quite easily cause impurities to penetrate into or even through these films. It has also been noted ¹³⁰ that voids are formed in about 1 nm thick layers when annealed at 750 °C. This is caused by the reaction $\text{Si} + \text{SiO}_2 \rightarrow 2\text{SiO}(\text{g})$ in the silicon/silicon dioxide interface ¹³¹. Because this reaction also consumes silicon, the silicon surface is lower in the void spot ¹³². It has been suggested ¹³³ that these voids start to form at defect sites. Layers fabricated with different processes, therefore, should have different void densities. In fact, it has been observed that on layers grown by rapid thermal oxidation, the average void distance is only half the corresponding distance on thermal oxides ¹³⁴. This is probably due to the fact that higher temperatures are used in RTO which as mentioned serves to improve the oxide quality.

It has been proposed¹³⁵ that the primary result of (inert atmosphere) high temperature (above 950 °C) annealing might be the creation of oxygen vacancies or oxygen vacancy related complexes, like oxygen vacancy center/silicon interstitial complex¹³⁶, in the silicon dioxide. In addition, there are also oxygen related donors in the silicon. These defects result from the outdiffusion of oxygen from the silicon dioxide network into the silicon¹³⁶. The driving force behind this mechanism is the chemical potential difference between oxygen in silicon and silicon dioxide^{135,136}. These oxide defects appear to be the major source of enhanced hole trapping, and it has been concluded that they extend from the interface into the bulk of the oxide. This suggests that they should affect not only the density of bulk oxide traps, but also the border and interface trap densities¹³⁵. The mechanism behind thermal post oxidation interface degradation, which begins when temperatures above 640 °C are used, has been identified as permanent paramagnetic point interface defect creation (so-called P_b defect: trivalent interfacial Si backbonded to three Si atoms in the substrate: •Si≡Si₃¹³⁷). This reveals that interface breakdown mechanism on an atomic scale is interfacial Si-O bond rupture¹³⁸. However, it seems that this process is reversible even below 400 °C⁸¹.

The atmosphere in which the annealing is done has quite a major impact, especially at lower temperatures. Annealing Si/SiO₂ structures in an ambient containing hydrogen at relatively low temperatures (typically 250 - 450 °C) is an effective way of reducing the surface state density⁸¹. Low temperature annealing in nitrogen, by contrast does not seem to have any effect on the surface state density⁸¹. Elimination of traps like trivalent Si centers (Si dangling bonds), involves the chemical saturation of the free bonds by forming an Si-H configuration. At 500 °C and above, these bonds tend to dissociate and the interfacial trap density increases again⁸¹.

Differences have also been found between argon and deuterium annealing, for instance. Annealing at 500 °C in argon was not found to cause any changes in the interface trap density, whereas in deuterium the density was clearly decreased⁸⁰. Annealing in a hydrogen containing ambient has been a common method of passivating silicon interface, because this allows the hydrogen ions to passivate the dangling bonds of the SiO_x in the interface region. However, a positive charging of silicon dioxide layers has been found while annealing Si/SiO₂/Si structures at temperatures above 500 °C. This has been assumed to be due to the formation of positively charged oxygen centers¹³⁹. In

contrast to hydrogen, so called water-related species, Si-OH or loosely bound H₂O, in the bulk oxide are undesirable because they act as electron traps¹⁴⁰. Bulk electron trapping, especially water-related traps, can be reduced by low temperature annealing in a hydrogen ambient, but this process is also reversible⁸¹. Hole trapping is not reduced to the same level as in subsequent low temperature hydrogen annealing, but on the other hand there is also no reversibility⁸¹. It has also been shown that annealing in a helium atmosphere can yield better films than similar annealings in vacuum, for instance¹⁴¹. In particular, the electrical integrity and the density of hole traps are clearly improved. These effects seem to be linearly dependent on the helium pressure, at least as the pressure is between 0.1 and 22 bar¹⁴¹.

Another method that has been used is microwave annealing in N₂O at high pressure⁹⁴. This significantly improves the oxide quality, although more than one hour is needed to decrease the plasma oxidized sample interface state density.

Sometimes annealing after metallization is either preferred or needed. A metal commonly used on top of the oxide film is aluminum, although in MOSFET manufacturing highly doped polysilicon is more popular. The aluminum consumes silicon dioxide, causing spiking of aluminum into the oxide⁴⁶. Studies of metal film evaporation and the metal/oxide interface have shown¹⁴² that (in UHV, substrate at room temperature) the aluminum first clusters around each surface O and then grows Al₂O₃ by reducing oxide to SiO_x (x < 2), leaving excess silicon at the interface. An unannealed metal/silicon dioxide interface, aluminum evaporated at room temperature, can be atomically abrupt¹⁴³. Annealing, however, can start the oxidation - reduction reaction. This proceeds linearly with time. The predominant structure of the reaction products is most likely to be SiO₂/Al₂O₃/Si/Al in which the interfacial width can be varied by changes in processing¹⁴³. Annealing of MOS structures with aluminum gate in a neutral ambient has produced similar results as post oxidation annealing in a hydrogen containing ambient¹⁴⁴. The reaction between the metal and hydroxyl groups at the oxide surface is assumed to liberate active (probably atomic) hydrogen, which eliminates the traps¹⁴⁴.

While post oxidation annealing processes are required to reduce the oxygen charge, they must not at the same time degrade the dielectric breakdown characteristics of the

oxide¹⁴⁵. Oxygen deficient annealing in vacuum tends to lead to the formation of electrical defects in the oxide. This phenomenon appears to correlate with the initial stages of oxide decomposition. It has been found that high temperature anneals enhance hole trapping in thermally grown silicon dioxide films. Breakdown can be prevented by annealing in controlled O₂ ambient at pressures above critical values, which are similar to the SiO vapor pressure¹⁴⁵. In this way the decomposition can be suppressed by reoxidation of the volatile SiO reaction product.

4 Electrical Characterization of Ultrathin Insulators

There are several characterization methods that give information about thin films and interfaces. The focus here is on electrical characterization, which provides useful information particularly for the fabrication of electronic devices. Conventional capacitance-voltage (CV) characterization techniques cannot, however, be used in the separation of positive and negative trapped-charge densities, as they sense the total charge, only ⁶⁴.

The Poisson equation of ideal MOS structure (Figure 2.1) gives the relation between charge and potential. As mentioned before our samples can be quite far from being ideal. The non-idealities can, however, be taken into account in the characterization by separating their effect from the measurements or some of them can also be excluded by careful measurements and fabrication processes.

One-dimensional equations can be used as valid approximation when the area of the MOS structure is large compared to the thickness of the insulator. Then

$$\frac{d^2V}{dz^2} = -\frac{dE}{dz} = \frac{q}{\epsilon_{Si}} [N_A^-(z) - N_D^+(z) + n(z) - p(z)], \quad (4.1)$$

where, ϵ_{Si} is the dielectric constant of the silicon, q is unit charge, n is electron and p hole density, z is position co-ordinate perpendicular to the interface, and N_A^- and N_D^+ are the densities of the ionized acceptor and donor atoms. V and E denote the potential and electric field.

From equation (4.1), when the potential and electric field far from the silicon/silicon dioxide -interface are defined as zero, a relation can be derived for electric field ⁴⁶:

$$\epsilon_{Si} E^2(z) = 2 \left[k_B T (p(z) - p_0 + n(z) - n_0) + q (N_A^- - N_D^+) V(z) \right] \quad (4.2)$$

when

$$n(z) \approx n_0 e^{qV/kT} \wedge p(z) \approx p_0 e^{-qV/kT} \quad (4.3)$$

and n_0 and p_0 are the carrier concentrations in thermal equilibrium, far from the interface. Then at the silicon/silicon dioxide -interface

$$V_s = V(z=0) = \frac{kT}{q(N_A^- - N_D^+)} [p_0 - p(V_s) + n_0 - n(V_s)] + \frac{\epsilon_{Si}}{2q(N_A^- - N_D^+)} E_s^2 . \quad (4.4)$$

If there is no charge in the oxide, the surface field in the silicon is

$$E_s = E(z=0+) = \frac{\epsilon_{ox}}{\epsilon_{Si}} E(z=0-) = \frac{\epsilon_{ox}}{\epsilon_{Si}} \frac{(V_g - V_{fb} - V_s)}{d_{ox}} . \quad (4.5)$$

In this equation V_g is the gate voltage (potential on the metal), V_{fb} is the flat band voltage and d_{ox} is the thickness of the oxide. The bending of the energy bands, and consequently the charge densities, can be numerically calculated from equations (4.3), (4.4) and (4.5)¹⁴⁶. It can be seen from Figure 4.1, that the band bending in accumulation is relatively modest.

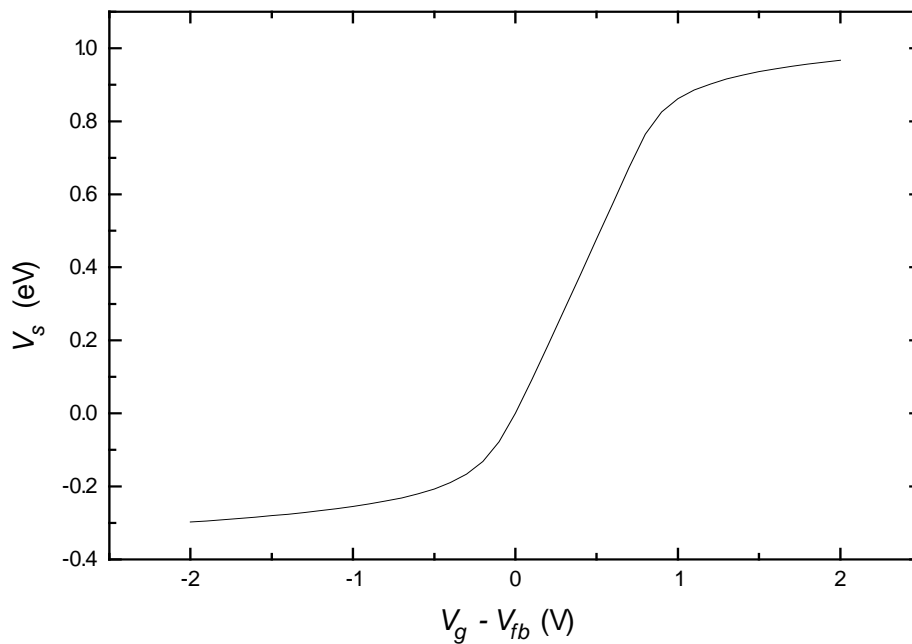


Figure 4.1 Potential at the oxide/silicon interface of an MOS structure ($N_A = 10^{16} \text{cm}^{-3}$, $d_{ox} = 2 \text{ nm}$) as a function of gate voltage.

4.1 Oxide Charge

The most usual way to define the density of a fixed charge in the insulator layer is to calculate it on the basis of the flat-band voltage shift.

If the insulator is charged, then the electric field will not stay constant, like assumed in equation (4.5). If it is assumed that there exists a Dirac delta function (δ) like interface charge (N_s) in the silicon/silicon dioxide interface and that otherwise the charge density in the insulator is uniform (N_{ox}), then the Poisson equation (4.1) in the oxide is (structure like in Figure 2.1)

$$\frac{dE(z)}{dz} = -\frac{d^2V}{dz^2} = \frac{q}{\epsilon_{ox}} [N_{ox} + N_s \delta(0)], \quad z \in [-d_{ox}, 0]. \quad (4.6)$$

Then the electric field in oxide is

$$E(z) = \frac{q}{\epsilon_{ox}} [N_{ox}z + N_s] + \frac{\epsilon_{Si}}{\epsilon_{ox}} E(0+), \quad z \in [-d_{ox}, 0] \quad (4.7)$$

and potential

$$V(z) = -\frac{q}{\epsilon_{ox}} \left[\frac{1}{2} N_{ox} z^2 + N_s z \right] - \frac{\epsilon_{Si}}{\epsilon_{ox}} E(0+)z + V_s, \quad z \in [-d_{ox}, 0]. \quad (4.8)$$

In the metal/oxide interface the potential is defined by the gate potential

$$V(-d_{ox}) = V_g - \Delta V_{fb} - \frac{1}{q} \left[\Phi_{Met} - \Phi_{Si} - \frac{E_g}{2} - k_B T \ln \left(\frac{N_A}{n_i} \right) \right] \quad (4.9)$$

for p-doped and

$$V(-d_{ox}) = V_g - \Delta V_{fb} - \frac{1}{q} \left[\Phi_{Met} - \Phi_{Si} - \frac{E_g}{2} + k_B T \ln \left(\frac{N_D}{n_i} \right) \right] \quad (4.10)$$

for n-doped silicon. n_i is intrinsic carrier concentration, Φ_{Met} is the work function of metal and Φ_{Si} the work function of silicon. E_g is the width of the energy gap and

$$V_{fb} = \Delta V_{fb} + V_{bi}, \quad (4.11)$$

in which the V_{bi} is flat band voltage in an ideal MOS structure. The shift in the gate potential caused by the oxide and interface charge is

$$\Delta V(-d_{ox}) = -\frac{q}{\epsilon_{ox}} \left[\frac{1}{2} N_{ox} d_{ox}^2 + N_s d_{ox} \right], \quad (4.12)$$

and therefore the relation between the flat band shift and the charge in the oxide and silicon/silicon dioxide interface is

$$N_s + \frac{1}{2} N_{ox} d_{ox} = -\frac{\epsilon_{ox} \Delta V_{fb}}{q d_{ox}}. \quad (4.13)$$

4.2 Impedance of Thin Oxide MOS Structure

It is extremely difficult to produce reliable measurements when the measured oxide is very thin. It has been observed, for instance¹⁴⁷, that in MOS structures, the gate capacitance is typically less than 90% of the ‘traditional’ oxide capacitance, and for ultra thin oxide layers and relatively low doped polysilicon gates ($< 10^{20} \text{ cm}^{-3}$) it can even drop below 75%. For thin oxides then, the effective electrical thickness exceeds the physical thickness of the layer. This means that the established measurement techniques and analyses used in more traditional MOS characterization are not necessarily valid any more. It follows that the traditional formulas need to be developed if we intend to achieve better reliability in MOS characterization.

Some problems are caused in the situation where there is significant leakage through the insulator layer and where it is not possible to exclude all of the series resistance from the measurements. Equivalent circuit for this case is shown in Figure 4.2 .

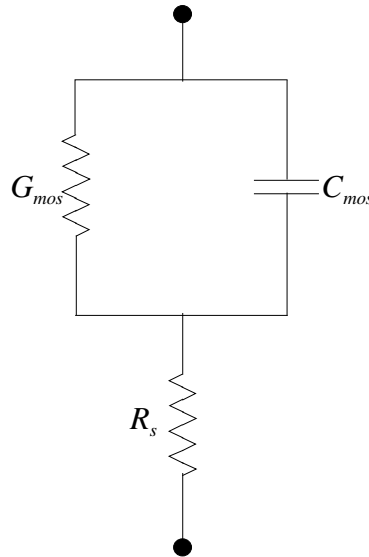


Figure 4.2 Equivalent circuit of the MOS structure with leakage (G_{mos}).

The basic measurement required for the characterization of insulator layers is impedance (or admittance) measurement. Our HP 4192A LF measurement unit^{148,149} calculates parallel conductance and capacitance as well as series resistance and capacitance from measured values. The measurement unit gives us the *measured* admittance

$$Y_m = G_m + i\omega C_m, \quad (4.14)$$

or

$$Z_m = R_m + iX_m = \frac{1}{Y_m} . \quad (4.15)$$

As the total impedance of the circuit (Figure 4.2) is

$$Z = R_s + \frac{1}{G_{mos} + i\omega C_{mos}} = \frac{1}{Y}, \quad (4.16)$$

it follows that when this is assumed to be equal with the measured values

$$C_{mos} = \frac{1}{\omega} \text{Imag} \left[\left(\frac{1}{Y_m} - R_s \right)^{-1} \right]. \quad (4.17)$$

or

$$C_{mos} = \frac{C_m}{(1 - R_s G_m)^2 + (R_s \omega C_m)^2}, \quad (4.18)$$

It is, however, hard to define the real value of series resistance. If it is assumed that $G_{mos} = 0$ (i.e. there is no leakage), then

$$R_s (G_{mos} = 0) = \frac{G_m}{G_m^2 + (\omega C_m)^2} \quad (4.19)$$

and

$$C_{mos} (G_{mos} = 0) = -\frac{1}{\omega X_m}. \quad (4.20)$$

The main difficulty in characterization is presented by combining series resistance and leakage current. If the leakage current is low, equation (4.20) gives quite a good approximation of real oxide capacitance. If the leakage is high, then the real value of the capacitance is closer to the admittance measurement (equation (4.14), example in Figure 4.3). Then it is not possible to calculate exact values for layer capacitance, but only upper and lower limits.

Series resistance is usually taken into account by giving it a constant value. This is calculated from the measured accumulation capacitance and conductance based on

equation (4.19), assuming that the leakage current is low. The most important values are those obtained for accumulation, providing as they do the most reliable picture of the insulator layer capacitance and conductance. The high frequency depletion layer capacitance can be ignored in accumulation. In accumulation, however, the leakage is higher, which increases the error of the calculated series resistance. The series resistance is also not constant, but increases with reverse bias, i.e. with increasing depletion layer width and a decreasing amount of carriers near the oxide/silicon -interface. This change is relatively small, however, and in practice can usually be ignored.

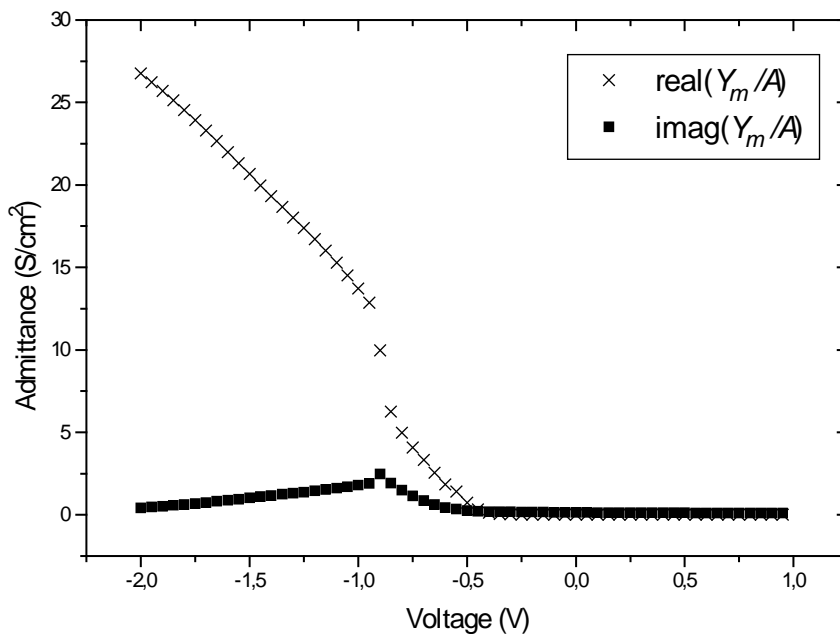


Figure 4.3 Measured admittance from an MOS structure (oxide thickness 3.9 nm, frequency 1 MHz). Squares are the imaginary and crosses the real part of the admittance.

However, the equivalent circuit described is unable to present one important phenomenon that can be seen in the AC measurements of MOS structures: it does not take into account the interface effects, mainly interface traps and charge. In high frequency measurements the depletion region capacitance can not be neglected. A better equivalent circuit of the system is presented in Figure 4.4.

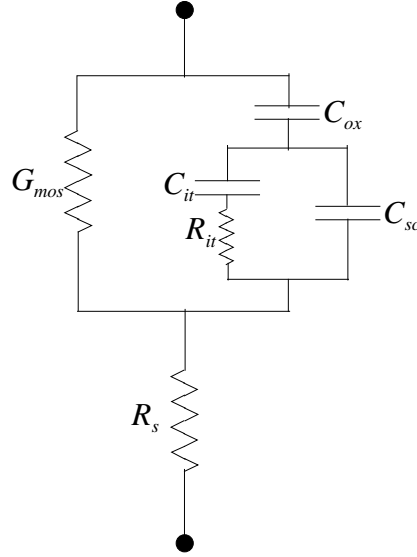


Figure 4.4 Equivalent circuit of MOS structure. G_{mos} is the leakage conductance through the structure, C_{ox} is the metal-oxide-semiconductor capacitance, C_{it} is the capacitance of the interface traps, R_{it} is their resistance, C_{sc} is the depletion region capacitance and R_s is the series resistance.

As described in ref. ¹⁵², the effect of traps can be presented by a capacitor (C_{it}) in a series with a resistor (R_{it}). The amount of interface traps is characterized by a parameter called the *surface state density* (D_{it}) which is directly proportional to the interface state capacitance C_{it} .

$$D_{it} \propto C_{it} . \quad (4.21)$$

Because it takes some time the traps to change their occupancy the resistor, R_{it} , must also be included.

The impedance of the circuit in Figure 4.4 is then

$$Z = R_s + \frac{1}{G_{mos} + \frac{1}{Z_s - i(\omega C_{ox})^{-1}}} . \quad (4.22)$$

where

$$Z_s = \begin{cases} Z_{it} = R_{it} - \frac{i}{\omega C_{it}} , & \text{in accumulation} \\ \frac{Z_{it}}{1 + iZ_{it}\omega C_{sc}} , & \text{in depletion} \end{cases} . \quad (4.23)$$

The depletion region capacitance C_{sc} is defined by

$$C_{sc} = \frac{\epsilon_{Si} A}{d_{sc}} . \quad (4.24)$$

The width of the depletion region d_{sc} is usually derived by using the assumption that there is a region in which all donors and acceptors are ionized and there are no free carriers. Outside this region there prevails charge neutrality. By using the Poisson equation (4.1), one gets

$$d_{sc} = \sqrt{\frac{2\epsilon_{Si} V_s}{q |N_A^- - N_D^+|}} . \quad (4.25)$$

A maximum length is usually defined for the depletion region. This is reached when depletion turns into inversion, i.e. when the density of minority carriers at the interface exceeds the majority doping concentration

$$\max(d_{sc}) = \sqrt{\frac{4\epsilon_{Si} k_B T \ln \left| \frac{N_A^- - N_D^+}{n_i} \right|}{q^2 |N_A^- - N_D^+|}} . \quad (4.26)$$

4.3 Oxide Thickness

The basic formula for the calculation of oxide thickness from an MOS structure can be derived from equation (4.5). On this basis, the ideal differential low frequency capacitance of an ideal MOS structure is

$$C = \frac{dQ}{dV_g} = A \frac{d\varepsilon_{Si} E_s}{dV_g} = \frac{\varepsilon_{ox} A}{d_{ox}} \left(1 - \frac{dV_s}{dV_g} \right) = \frac{\varepsilon_{ox} A}{d_{ox} + \frac{\varepsilon_{ox}}{\varepsilon_{Si}} \frac{dV_s}{dE_s}}, \quad (4.27)$$

where A is the area of the capacitor. By using equations (4.4), (4.5) and (4.27) it is possible to derive the capacitance formula even further:

$$C = \frac{\varepsilon_{ox} A}{d_{ox} + \frac{\varepsilon_{ox} E_s}{q(N_D - N_A - p_s + n_s)}}. \quad (4.28)$$

Based on equation (4.27), the oxide thickness is ¹⁵⁰

$$d_{ox} = \varepsilon_{ox} \left(\frac{A}{C} - \frac{1}{\varepsilon_{Si}} \frac{dV_s}{dE_s} \right). \quad (4.29)$$

When the MOS structure is biased into accumulation V_s should saturate, and consequently its derivative should approach zero. This can also be seen from equation (4.28), as the density of the surface majority carrier increases exponentially with the surface potential, faster than the surface field (equation (4.5)). Then we get the familiar equation for the relation between capacitance and insulator thickness:

$$C_{acc} \approx \varepsilon_{ox} \frac{A}{d_{ox}}. \quad (4.30)$$

C_{acc} is the saturated accumulation capacitance of the MOS structure.

The first problem is how to define the real oxide capacitance. First of all, the series resistance does not remain constant even during one measurement, because the bias voltage affects the depth of the depletion layer and the interface charge. For the measurement of thickness, the accumulation capacitance is more important. The problem is that the leakage current peaks in accumulation, thus increasing the error in series resistance given by equation (4.19). More reliable values can be obtained by

fitting the whole measured curve with the formula presented in equation (4.28) instead of using equation (4.30) and the estimated accumulation value of the capacitance. If leakage is high and series resistance significant, the measured capacitance can be as shown in Figure 4.5. The correct series resistance can be deduced from the slope of the capacitance curve in the accumulation region.

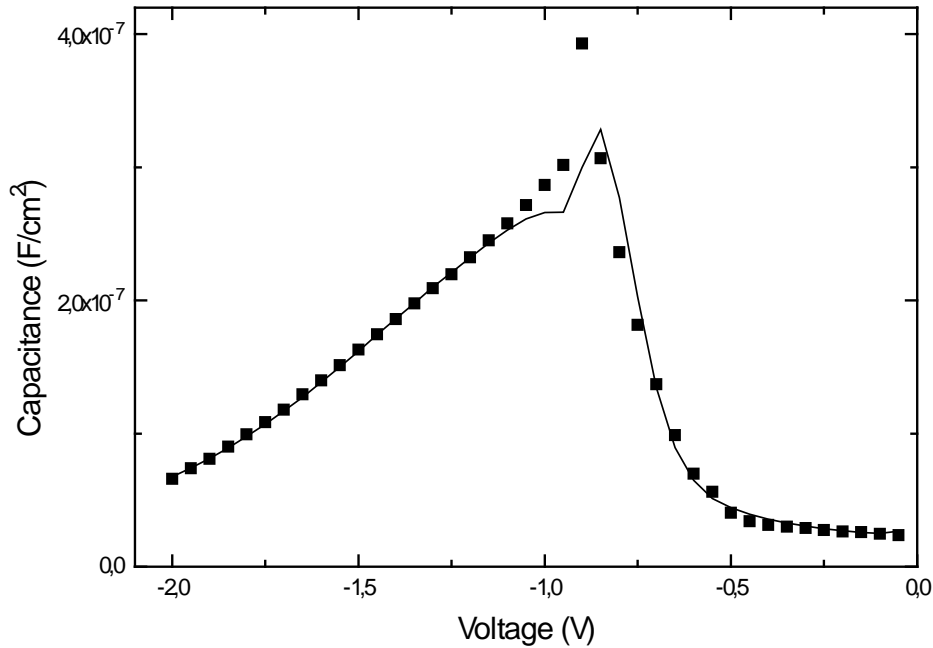


Figure 4.5 Measured (squares) and calculated (solid line) capacitance of the same MOS structure as in Figure 4.3. The calculation is based on the following parameters: Oxide thickness 3.9 nm, series resistance 310 Ω , doping density $4 \cdot 10^{15}$ $1/\text{cm}^3$, and flat band voltage -0.68 V.

The calculated value for resistance in Figure 4.5 is quite high. However, this is mainly due to the fact that the ‘bulk contact’ was on the other side of the wafer. This means that the measurement current had to penetrate the whole high resistivity wafer. When the wafer thickness is around 400 μm and the pad size is $(100 \mu\text{m})^2$, the substrate resistivity has to be only about 7 Ωcm and an approximate formula⁴⁶ gives series resistance of 300 Ω , which does not include contact resistances.

One of the difficulties in the measurement of ultrathin layers is the finite thickness of the interface charge. It is estimated that the classical extrapolation method of capacitance-voltage characteristics gives film thickness values that are 0.3 - 0.5 nm

larger than those produced by quantum mechanical analysis of the structure¹⁵. The main reason for this is said to be that in quantum capacitance calculations, the inversion charge has to be calculated on the basis of the wave functions of every sub-band forming into the interface when the energy bands are bended. For this reason the oxide thickness given by equation (4.30) is really the average distance of the accumulation charge in silicon from the interface charge between the oxide and the metal. Since the accumulation capacitance also does not totally saturate when the bias increases, it seems that the oxide layer becomes thinner in direct proportion to increasing voltage. Equation (4.28) takes this into account, which means that this effect can be avoided by numerical fitting of admittance measurements.

4.4 Surface State Density

Surface state density is a very important factor in MOSFET structures, since the current flows along the silicon/silicon dioxide interface. If the density is too high, MOSFETs will not work at all. The problem can not be avoided entirely, but in a good MOS process it is possible to achieve state densities around $10^{10} \text{ eV}^{-1} \text{ cm}^{-2}$.

The capacitance caused by interface traps (C_{it}) can be calculated on the basis of the equivalent circuit presented in Figure 4.4. For the forward biased, i.e. accumulation, case

$$C_{it} = \frac{C_{ox}}{\omega C_{ox} \text{Imag} \left\{ \left[\left(\frac{1}{Y} - R_s \right)^{-1} - G_{mos} \right]^{-1} \right\} - 1}, \quad (4.31)$$

and for the reverse biased depletion case

$$C_{it} = \frac{1}{\text{Imag} \left\{ iC_{sc} - \left[\frac{i}{C_{ox}} - \omega \left(\left(\frac{1}{Y} - R_s \right)^{-1} - G_{mos} \right)^{-1} \right]^{-1} \right\}}. \quad (4.32)$$

When the trap capacitance is known, the interface trap density can be calculated by using equations (4.14) and (4.21).

However, the voltage dependent impedance measurement on a single frequency does not give a very reliable value for the surface state density as the traps tend to be ‘slow’. It takes time for them to fill in. The existence of several separate energy levels, with slightly differing properties, also underscores the need to develop the model further.

In the so-called conductance technique measurement of surface state densities^{151,152}, impedance has to be measured as a function of voltage as well as frequency. In actual structures the physical situation is not as simple as in Figure 4.2, but this only describes the ‘lumped parallel equivalent circuit’ of a more realistic one, which is shown in Figure 4.6¹⁵². This represents the situation where there exist several trap levels rather than just one. Since the small voltage oscillation in the measurements mainly affects the traps near the Fermi level (within a few $k_B T$), the bias voltage can be used to determine the energy level of the traps that are seen. This makes it possible to measure the surface state density as a function of energy allowing us to determine the state distribution in the energy gap.

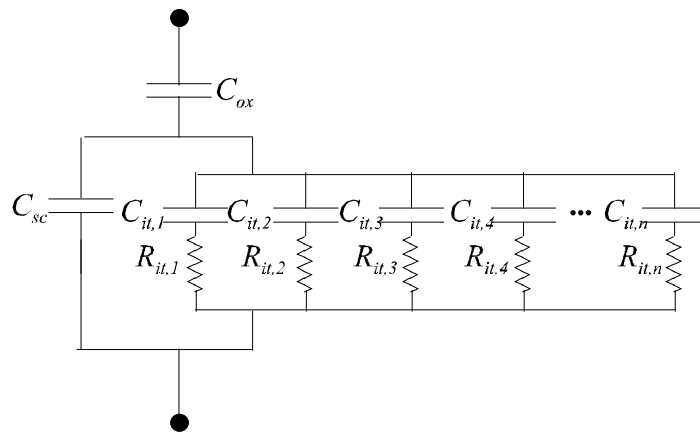


Figure 4.6 Equivalent circuit of the MOS capacitor in depletion for a distribution of single-level interface traps (leakage G_{mos} excluded).

The surface state density at a defined energy level can be calculated by using the relation ¹⁵²

$$D_{it} = \frac{1}{q^2 A} f_b \frac{\text{Real}(Y_{it}(\omega))}{\omega} \Big|_{\max}, \quad (4.33)$$

where A is capacitor area, f_b is material dependent parameter, and

$$Y_{it} = \frac{1}{\left[(Z - R_s)^{-1} - G_{mos} \right]^{-1} + i(\omega C_{ox})^{-1}} \quad (4.34)$$

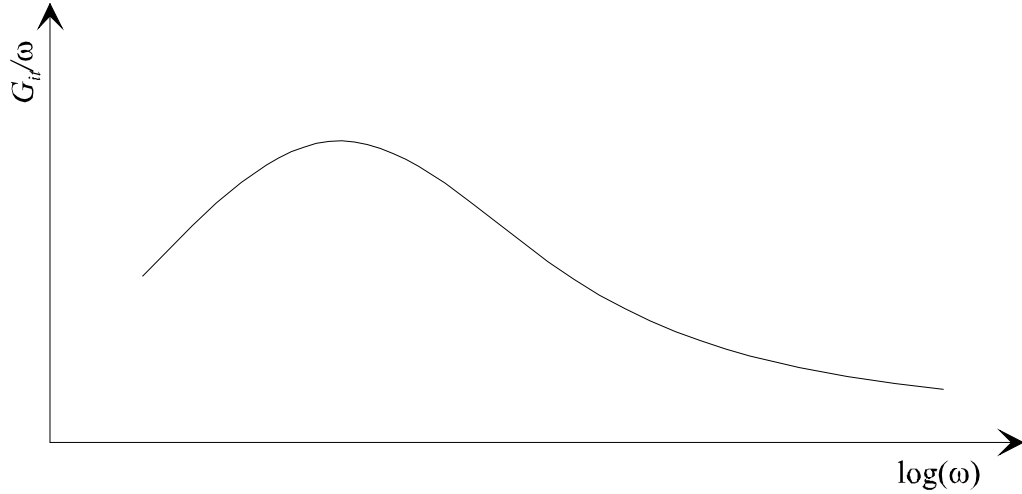


Figure 4.7 A schematic illustration of trap conductance as a function of angular frequency.

The state potential that corresponds to a given bias voltage is

$$V_{trap} = \begin{cases} \frac{E_g}{2q} + \frac{kT}{q} \ln\left(\frac{N_A}{n_i}\right) + V_s(V_g) + \frac{E_v}{q}, & \text{for p-doped semiconductor} \\ \frac{E_g}{2q} - \frac{kT}{q} \ln\left(\frac{N_D}{n_i}\right) + V_s(V_g) + \frac{E_v}{q}, & \text{for n-doped semiconductor} \end{cases}, \quad (4.35)$$

where E_g is the semiconductor energy bandgap and interface potential V_s like in equation (4.4).

This model is quite commonly used. However, it has been criticized for not including so-called 'slow traps', as there is some evidence that there are two populations of defects at the silicon/silicon dioxide interface²⁰. The measurements are also time-consuming because the admittance has to be measured as a function of both voltage and frequency. This was nonetheless the method I decided to use in an attempt to get a more reliable picture of the state densities in the gap as a function of energy.

5 Ultrahigh Vacuum Plasma Oxidation (UPO)

In ref. ¹ we have utilized plasma oxidation of silicon in ultra high vacuum. Since the best silicon/silicon dioxide -interface is fabricated by thermal oxidation of (100) silicon ⁷⁸, the samples used in our experiments were ordinary p-type (100) silicon wafers. The use of epitaxial silicon surfaces or high quality Float-Zone wafers would probably decrease defect densities and the impurity diffusion, as they are usually of better quality and have lower doping levels. We nevertheless decided to use ordinary substrates as they were, in order to imitate actual device processing.

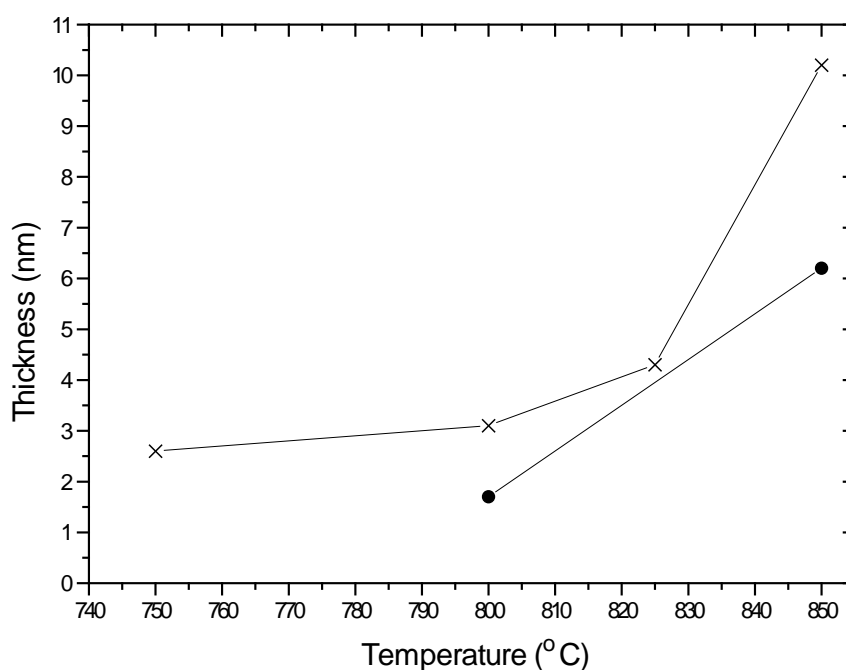


Figure 5.1 Oxide thickness after 20 minute oxidation as a function of temperature (Crosses; plasma on. Circles; without plasma).

We opted to use plasma instead of molecular oxygen because that gave a higher growth rate (Figure 5.1) and increased maximum layer thickness. These are due to higher reactivity and also faster diffusion, resulting from the smaller size of the oxidizing species, which we hoped would also help to reduce the final amount of free bonds.

5.1 Equipment

The equipment used was a modified two chamber VG Semicon V80M molecular beam epitaxy (MBE) system. The sample temperature can be adjusted by a resistive heater and verified by an optical pyrometer between 500 and 900 °C. The input oxygen is fed through an inlet that is controlled by a mass flow controller. The gas is activated in a plasma cell ¹⁵³ (Figure 5.2). Typical values for the RF-field used are 265 - 300 W. The background pressure is around 10^{-10} - 10^{-9} mbar and during growth the pressure in the chamber is $3 \cdot 10^{-5}$ mbar and the oxygen mass flow about 1.2 sccm. The growth chamber houses cells for silicon, doping materials and several metals including aluminum. There is also a Staib Instrumente RHEED System EK-2035-R that can be used for *in situ* monitoring of the substrate surface and a Sycon Instruments STC-200 Deposition Rate Controller.

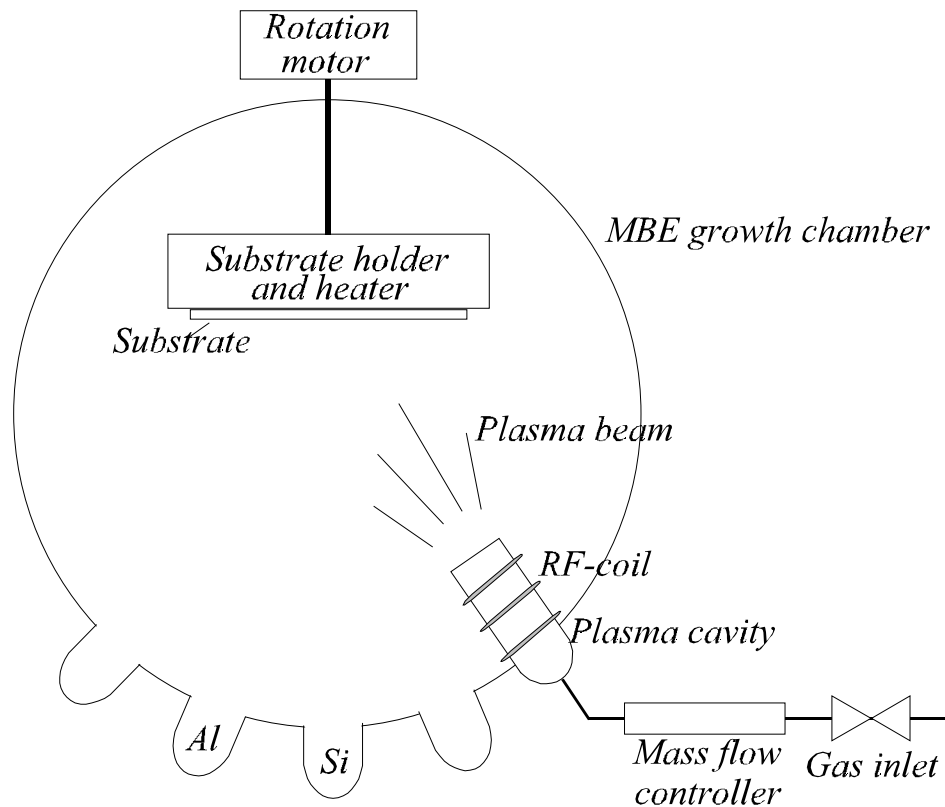


Figure 5.2 Diagram of the plasma assisted MBE growth apparatus.

5.2 Electrical Characterization

The MOS structures (see chapter 5.4.1 for a description of fabrication and structure) were characterized by using an HP4155A Parameter Analyzer for DC measurements and an HP4192A Impedance Analyzer for admittance measurements. Both of these units were connected to the same probe station. Layer thickness was defined by using voltage dependent capacitance measurement, at 1 MHz frequency. Oxide charge was calculated from the flat band shift (chapter 4.1). Surface state densities were calculated from frequency dependent conductance measurements¹⁵² by using the relation in equations (4.33) and (4.34). Series resistance was defined by comparing the theoretical capacitance of an ideal structure (equation (4.28)) and the measured high frequency admittance (equation (4.17)). Leakage current was assumed to be equal to the DC current measurement.

Great care and caution has to be exercised in the characterization of these ultrathin layers. When the thickness of the silicon dioxide layer is below 3 nm, the tunneling current increases and becomes significant: the effect of both the tunneling current and that of series resistance has to be taken into account. Dielectric breakdown also occurs at fields of about 10 MV/cm⁴⁶, which means only one volt across one nanometer of silicon dioxide. In our measurement system series resistance was only a couple of ohms, which made the calculations easier. At the same time this meant that the measurement itself had to be done very carefully, without using too high voltages, otherwise the current density could rise to a level where it might begin to create defects.

5.3 Oxidation Rate

In reactions involving silicon and oxygen there occurs both oxidation and etching of silicon: this applies particularly to low-pressure conditions. The higher the temperature and the lower the pressure, the faster the etching. The critical conditions between these two processes have been studied¹⁵⁴, and an exponential dependence between temperature and critical pressure has been found. In their experiments the temperature

was between 890 and 1150 °C and pressure from 0.07 to 70 mbar. It was noted that etching and growth rates were equal when the pressure is

$$P_{cr}(T) = P_0 e^{-E_A/kT}. \quad (5.1)$$

where $P_0 = 5.9 \cdot 10^{12}$ bar and $E_A = 3.83$ eV. These parameters seemed to be independent of silicon doping and surface crystal orientation¹⁵⁵.

However, this formula was tested in a different temperature and pressure range than we are using, which means it cannot be used as such. The best way to find the proper process parameters is by way of experimentation.

5.3.1 Experiment

The samples used in the oxidation rate determination¹ were ordinary p-type (100) silicon wafers, with a resistivity of about 10 Ωcm. These were cleaned using a standard chemical cleaning process (5 minute ultrasonic in acetone and isopropyl alcohol, 10 minutes in H₂O:NH₄OH:H₂O₂ (5:1:1) at 80 °C, 10 minutes in H₂O:HCl:H₂O₂ (5:1:1) at 80 °C). The native oxide was removed *in situ* in the MBE UHV chamber by keeping the samples at 875 °C for 15 minutes. The surface was monitored by RHEED to check that the oxide had been fully removed.

During oxidation the substrate temperature was between 600 and 850 °C as determined by an optical pyrometer. The growth times varied from 20 to 30 minutes. In all our experiments the oxygen flow was about 1.3 sccm. The background pressure was around 10⁻⁸ mbar, and during growth the pressure in the chamber was 2.5·10⁻⁵ mbar.

The oxide thicknesses were measured by an ellipsometer using the fixed value of the refractive index $n = 1.465$, typical of silicon dioxide (wavelength 632.8 nm). All the results are averaged from ten measurements, which varied by less than about 0.1 nm. It is known, however, that the refractive index of very thin silicon dioxide films differs from the value of thicker layers³³, which may give rise to inaccuracies. Also, the

manufacturer of the ellipsometer does not guarantee the accuracy of the results below 1 nm. The ellipsometric measurements were, however, compared with results from capacitance measurements as well as tunneling current measurements¹⁵⁶ and were found to be in close agreement (differences within 0.1 nm, the film thicknesses being 2 - 3 nm).

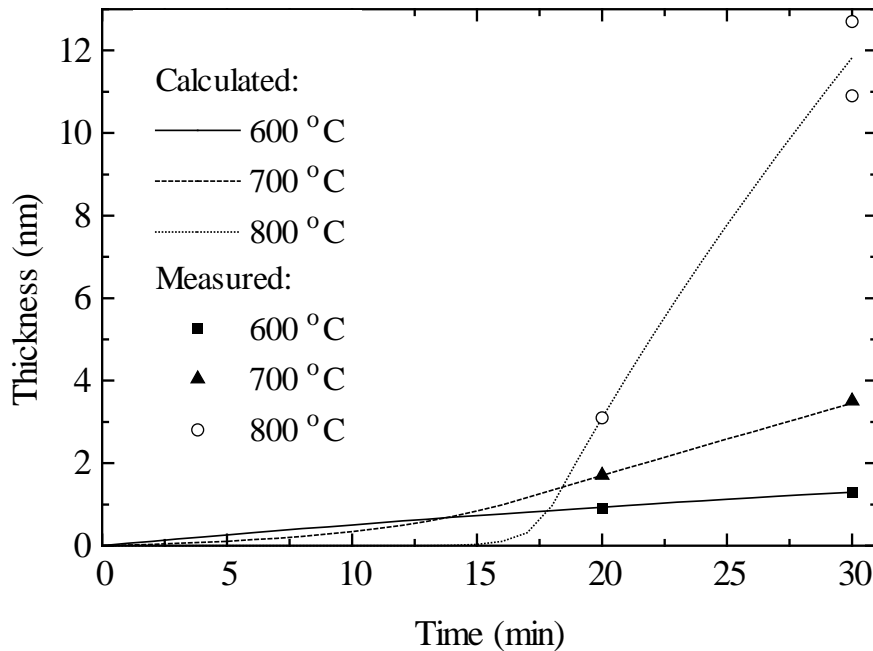


Figure 5.3 Measured thickness of the silicon dioxide layer as a function of time at various temperatures¹. The calculations are based on equations (5.3) and (5.4) which are fitted with the measurements.

As we can see in Figure 5.3, the average growth rate is very slow for the first 15 minutes, less than 0.1 nm/min^{-1} . It then increases very sharply, reaching about 1 nm/min at $800 \text{ }^\circ\text{C}$. This indicates that the sticking coefficient of oxygen molecules is smaller on a pure silicon surface than it is on a silicon dioxide surface, like in ref.¹⁵⁷. This contradicts the models presented in chapter 3.2.1. This behavior has been seen also in some other thermal oxidation experiments¹⁵⁸. We observed an incubation period at the onset of the oxidation procedure at temperatures from room temperature through to $1000 \text{ }^\circ\text{C}$. Initially, there were no signs at all of any oxidation, but it only started after a certain period of time. The duration of this interval seemed to decrease with increasing temperature, and is supposed that it is sensitive to the nature of surface termination and the presence of contamination. Growth behavior is therefore not as simple and

straightforward as described by equation (3.8). There may be considerable variation in initial growth, depending on the surface quality. We model this process using only one parameter, i.e. effective sticking coefficient, to describe all the surface effects and reactions ¹.

Since the sticking coefficients of oxygen are presumably different for silicon and silicon dioxide surfaces, the oxygen surface concentrations C_{Si} and C_{Ox} also differ from each other. When the oxygen flux starts to ‘see’ the surface as a pure silicon dioxide surface, the critical thickness d_{cr} has been reached and $C_0 \rightarrow C_{Ox}$. Below critical thickness, we simply interpolate for the average surface concentration. Thus we obtain ¹

$$C_0 = \begin{cases} C_{Si} + \frac{C_{Ox} - C_{Si}}{d_{cr}} z, & \text{when } z < d_{cr} \\ C_{Ox}, & \text{when } z \geq d_{cr} \end{cases}. \quad (5.2)$$

Solving equations (3.1) - (3.4) and (5.2), we get

$$t = \begin{cases} \left\{ \frac{2d}{B(1-r)} \left[z + \left(\frac{A}{2} - \frac{rd_{cr}}{1-r} \right) \ln \left[1 + \left(\frac{1}{r} - 1 \right) \frac{z}{d_{cr}} \right] \right] \right\}, & \text{when } z < d_{cr} \\ \frac{1}{B} [z^2 - d_{cr}^2 + A(z - d_{cr})] + \tau, & \text{when } z \geq d_{cr} \end{cases}, \quad (5.3)$$

where

$$\tau = t(d_{cr}) = \frac{2d_{cr}}{B(1-r)} \left[d_{cr} - \left(\frac{A}{2} - \frac{rd_{cr}}{1-r} \right) \ln(r) \right]. \quad (5.4)$$

The parameter

$$r = C_{Si}/C_{Ox}, \quad (5.5)$$

is called the relative sticking coefficient. If $r < 1$, what it seems to be for good quality silicon surfaces, the initial growth rate is slow but increases to its maximum when the

thickness reaches around 1 nm¹.

The initial growth rate should decrease with increasing temperature¹⁵⁹. Because the linear rate constant tends to increase with temperature⁴⁶, the relative sticking coefficient has to decrease even faster. From equations (5.3) and (5.4), it can be derived that

$$\left. \frac{dz}{dt} \right|_{t=0} = \frac{B}{A} r . \quad (5.6)$$

The value of r is approximately the same as the ratio between the initial growth rate and the growth rate after $z > d_{cr}$, as

$$\left. \frac{dz}{dt} \right|_{t=\tau} = \frac{B}{A + 2d_{cr}} . \quad (5.7)$$

The decrease in r with increasing temperature can be derived then from a comparison of these growth rates at different temperatures. This means that the relative sticking coefficient r is much lower than 1. It is a difficult task to measure the initial growth behavior, but our results suggest it is also possible that r could even reach values over one at lower temperatures. In other words the sticking probability would be higher on silicon than on silicon dioxide surfaces¹ and should increase with decreasing temperature.

Our measurements¹ indicate that the critical thickness d_{cr} is about 1 nm for our samples when the oxidation temperature is over 700 °C, and seems to increase slowly with declining temperature. This can be interpreted to mean that the average layer thickness increases at the moment when the entire sample surface is finally covered with dioxide. This implies a deterioration of layer quality. If the growth starts from separate spots and continues in a island form until they unite, it can be assumed that bigger the d_{cr} is, higher the islands get. Consequently there is more roughness. Another possible explanation is that at the lower temperatures, the initial layer quality is in some other way so poor that the reactions differ from those which are happening on high quality silicon dioxide surface. A thicker layer is needed before the reactions begin to approximate those occurring on silicon dioxide surfaces.

As can be seen in Figure 5.4, the growth rate is closely dependent on temperature. For maximum thickness control, it is better to use lower temperatures, because otherwise the growth rate, when the critical thickness is exceeded, will increase too much.

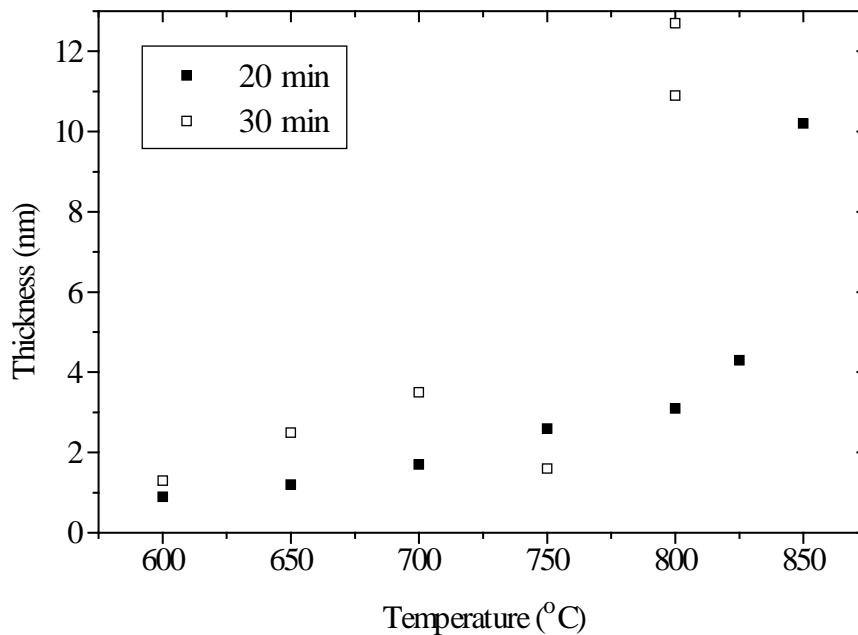


Figure 5.4 Thickness of the silicon dioxide layer as a function of temperature after 20 and 30 minutes of processing ¹.

Figure 5.5 shows a typical thickness profile of a silicon dioxide layer. The measurements were taken in four directions from the center. The thickness seems to be more or less uniform, which means that the processing parameters and conditions used are about right. The processing temperature does not seem to have any significant effect on thickness uniformity. The minor variations in thickness can be partly explained by the fact that in the MBE, the molecule flux is bigger in the center of the beam. There is, however, considerable variation in the results from different wafers, which can be explained by temperature changes on the wafer. Owing to the structure of the resistive heating element and the sample holder, the temperature is usually somewhat higher in the center of the wafer, and the same goes for growth velocity. The condition of the heating element is, however, crucially important: when it starts to corrode before reaching breaking point, part of the film will be clearly thinner because the sample will not be evenly heated (Figure 5.6).

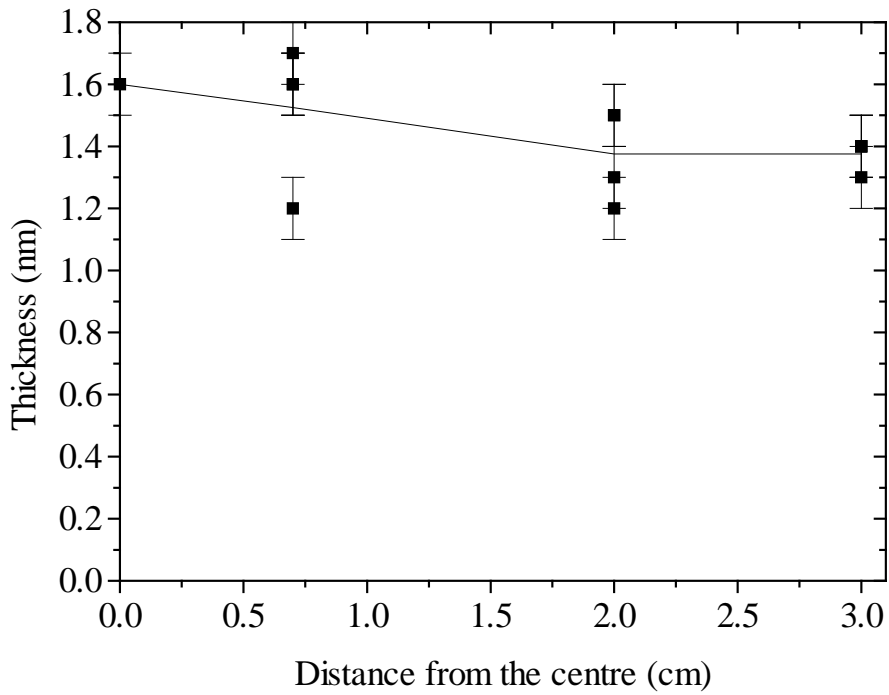


Figure 5.5 Thickness profile of a silicon dioxide layer processed at 750 °C for 30 minutes. The solid line is averaged from the measured values from four directions from the center. The measurements are presented as squares.¹

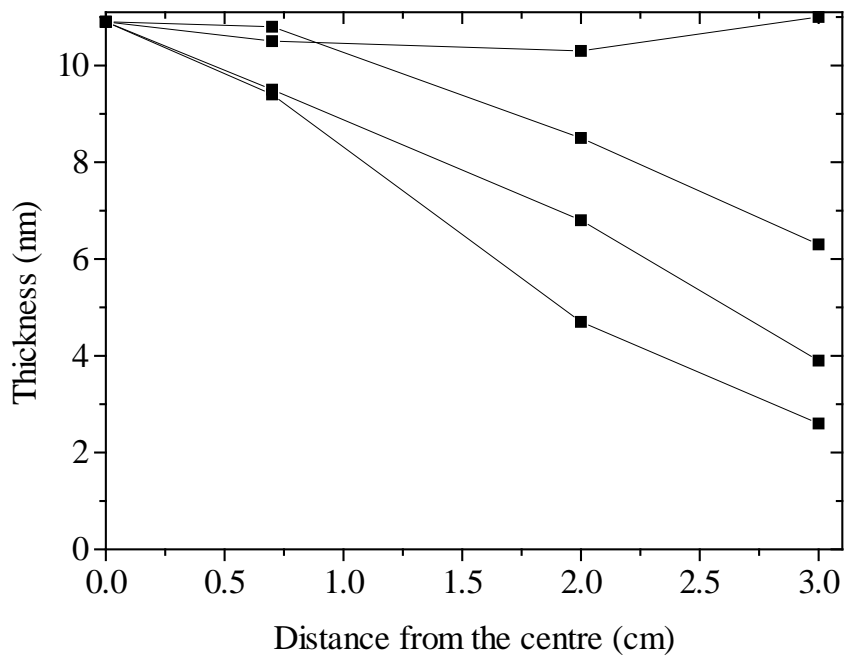


Figure 5.6 Thickness profile of a silicon dioxide layer in four directions from the center. The sample was processed at 800 °C for 30 minutes with a corroded heating element.¹

5.4 Room Temperature Oxidation

As has been pointed out earlier, higher processing temperatures produce better quality silicon dioxide layers. However, in a UHV system it is not possible to use very high temperatures, because otherwise the desorption rate may exceed the oxidation rate. Equipment limitations also restrict the temperatures available at below 900 °C. In our experiments ¹ we have noticed that below 700 °C, oxidation almost saturates after only a few nanometers. This means that the useful process window is quite narrow. To resolve this problem we have tried a completely opposite approach, i.e. oxidation at room temperature, which is a more ‘gentle’ process. TEM measurements (Figure 5.7) indicate that the process we are introducing seems to produce an extremely flat interface and homogenous layer thickness ¹².

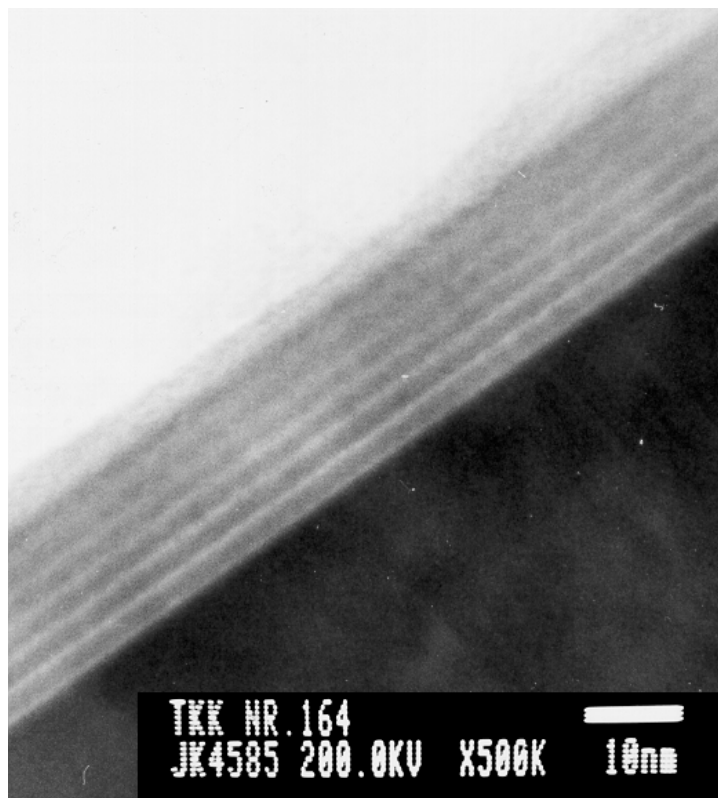


Figure 5.7 TEM picture of a silicon/silicon dioxide heterostructure.

5.4.1 Experiment

Three-inch boron doped (100) silicon wafers ($N_A = 0.5 - 1 \cdot 10^{16} \text{ cm}^{-3}$) were used as substrates. These were thermally dry oxidized at 1000°C to form a 100-nm thick insulator layer. $100 \mu\text{m} \times 100 \mu\text{m}$ openings were etched using photolithography and buffered HF. After an HF dip, samples were inserted in the MBE system, and the sample temperature was raised to 780°C . A low silicon flux was then used to etch the native oxide layer off from the openings. The temperature used has been reported to be the lowest possible for purposes of producing an atomically flat surface⁷³. Oxide removal was verified by monitoring the surface by RHEED. The temperature was then allowed to drop to room temperature. The actual oxidation was performed by feeding pure oxygen gas through a plasma cell in which the RF power was 265 W. The pressure in the chamber during oxidation was $3 \cdot 10^{-5}$ mbar, and the oxygen mass flow was 1.2 sccm.

In our experiments¹² we have found that the maximum layer thickness that can be reached at room temperature is 1 - 1.1 nm. In order to produce thicker layers, the procedure we applied was to evaporate a thin, about 0.6 nm, silicon layer on the silicon dioxide surface and to repeat the oxidation. The silicon layer thickness was defined using the Sycon Instruments STC-200 Deposition Rate Controller. Each processing cycle increased the layer thickness by one nanometer. We used 10 minutes for one oxidation cycle to make sure that all of the vaporized silicon had been oxidized. Once the desired thickness had been achieved, an aluminum layer of about 100 nm was vaporized on top of the sample. No post oxidation thermal treatments were used. After removal from the vacuum chamber, the test structures were patterned by using photolithography and wet chemical etching (Figure 5.8). The bulk contacts were made by using GaIn eutect, so that no sintering was needed. The MOS structures (capacitor area $(100 \mu\text{m})^2$) were then characterized¹⁵² as described earlier in chapter 5.2.

The method was used to produce two to five nanometer thick silicon dioxide layers. As can be seen in Figure 5.9, the measured and the expected values are reasonably close to one another². After two oxidation cycles, the measured values varied from 2.0 to 2.2 nm (calculated error is 0.3 nm, mainly due to the leakage current), after three cycles

from 3.0 to 3.3 nm (error 0.1 nm), and after five cycles from 4.9 to 5.2 nm (error 0.1 nm). The measurements were taken randomly from different places across the samples. The plasma oxidized silicon surface was first thermally oxidized and then the silicon dioxide layer was partly chemically etched off, which might have an impact on impurity concentrations on the surface and in the oxide. The results might have been even better if the silicon surface had been epitaxial instead of the wet etched and cleaned sample we used.

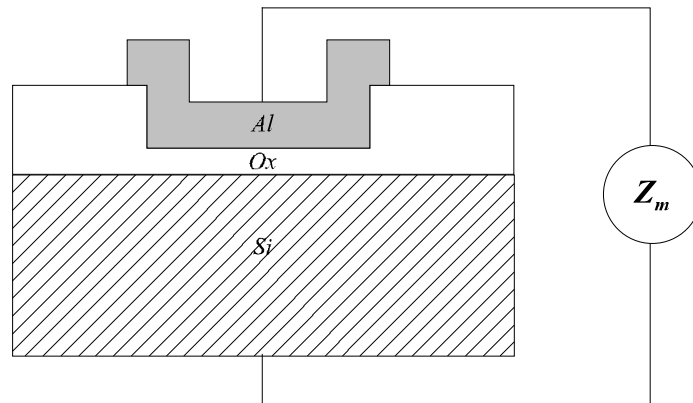


Figure 5.8 Schematic illustration of the MOS structure.

The results confirm our previous results according to which oxidation saturates after 1 - 1.1 nm and the thickness uniformity is quite good. The variation that can be seen in these figures is mainly within error limits, but the quality of the sample surface may also play a role.

The surface state density of these MOS structures seems to be quite good². Figure 5.10 shows that it is slightly above $10^{11} \text{ eV}^{-1} \text{ cm}^{-2}$ in the mid-gap. An increase in surface state densities can be seen in 2 nm oxide layer samples around 0.7 eV above the valence band. This might be caused by the impurity energy levels of oxygen and silicon (0.72 and 0.78 eV in silicon⁴⁶). Broken Si-O bonds, caused among other things by annealing, can be seen in the appearance of a localized peak of interface states at 0.8 eV. The dangling bonds tend to form a peak at 0.3 eV above the valence band¹⁶⁰. However, the situation improves significantly when the layer thickness increases to 3 nm. Because of the fabrication method used, the silicon/silicon dioxide interface should always be similar regardless of the layer thickness. At room temperature it is unlikely that there will be many changes in the interface structure because of oxygen diffusion or strain

when new oxide layers are added to the structure. This phenomenon may also be caused by impurities from the metal, or spiking of the aluminum, which tend to decrease with increasing oxide thickness.²

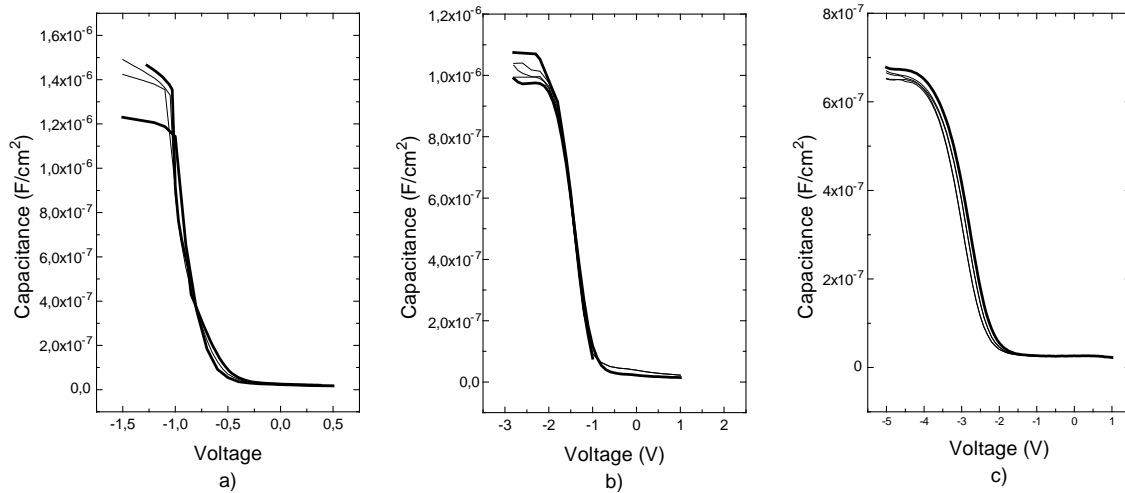


Figure 5.9 Measured capacitance of MOS samples fabricated with a) two, b) three and c) five oxidation cycles (frequency 1 MHz). Thick lines represent the calculated error limits.²

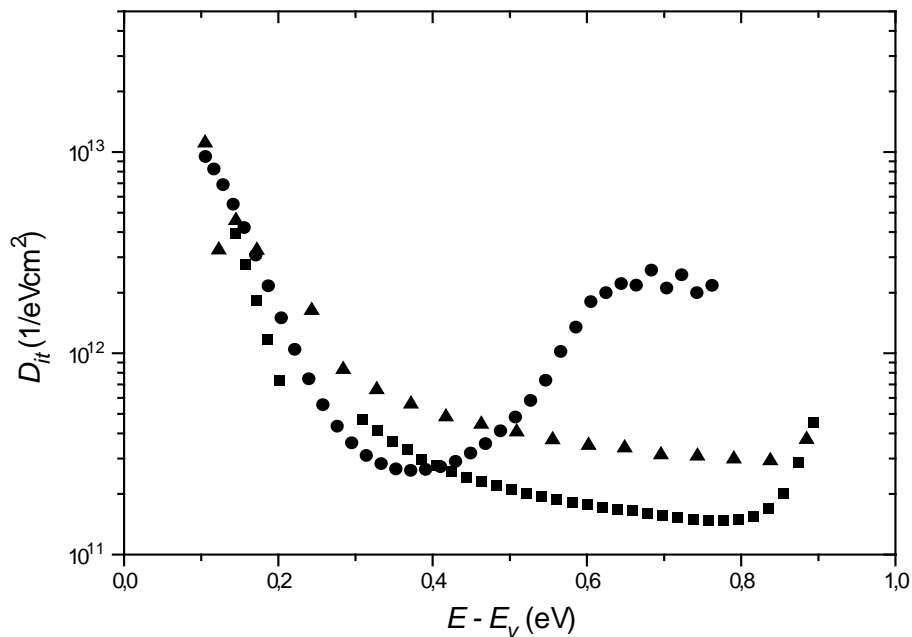


Figure 5.10 Surface state densities in MOS samples: oxide thickness 2.2 nm (circles), 3.3 nm (squares) and 5.0 nm (triangles)².

The leakage current due to tunneling through these ultrathin films is significant and increases exponentially, and therefore it is sometimes quite hard to spot the dielectric

breakdown. However, the breakdown field of these ultrathin layers seems to be somewhere above 10^7 V/cm (Figure 5.11a). This is about the same that has been quoted in the literature for silicon dioxide⁴⁶. In addition, there may occur a so-called 'soft breakdown' in fields above 10^6 V/cm⁵³. This can be seen in sudden increases in film conductivity (Figure 5.11b). These are rarely destructive, but they may badly interfere especially with AC measurements. The leakage current itself (Figure 5.12a) seems to be at the same level or slightly higher than the theoretical (equations (2.1) and (2.2)^{47,46}) tunneling current. The other measurement (Figure 5.12b) is from a sample in which the alignment was slightly off. The edge of the metal is partly on the thin oxide. In these cases a significant part of the current seems to go through the edges of the device: the current is more dependent on the length of the device periphery than on its area.

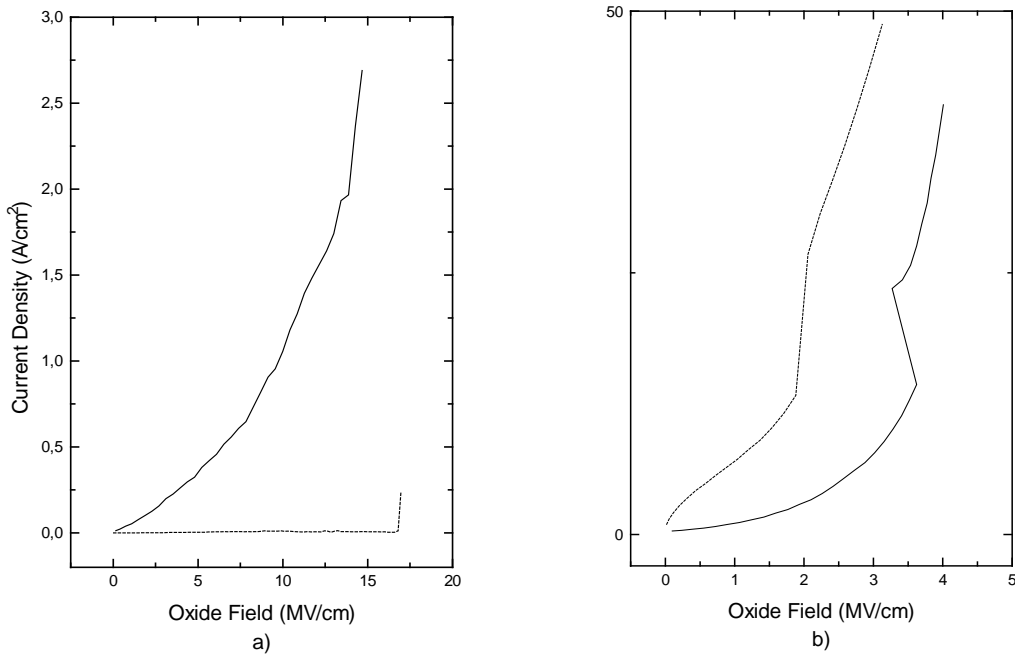


Figure 5.11 Measured currents through ultrathin silicon dioxide layers ($d_{ox} \approx 2 - 2.5$ nm).²

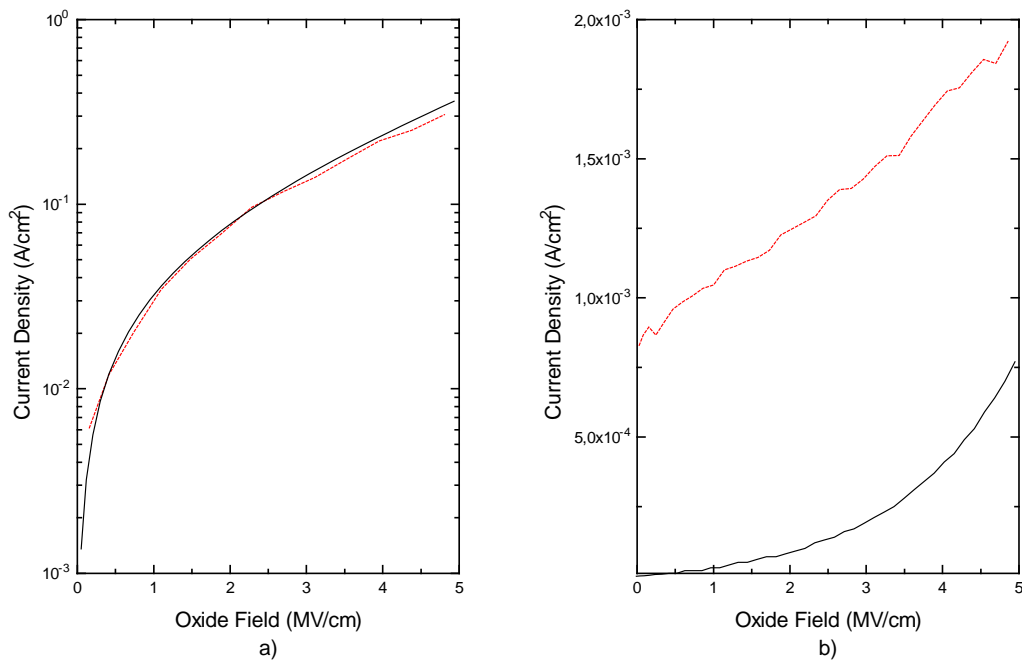


Figure 5.12 Calculated (solid black line) and measured (dashed red line) tunneling current through a) 2.2 nm and b) 3 nm ultrathin silicon dioxide layers as a function of oxide field. ²

5.5 Effect of Oxidation Temperature on the Silicon Dioxide Quality

Given the key impact of temperature on oxide quality, we also studied the effect of oxidation temperature. The range of temperatures covered was from room temperature to 800 °C.

5.5.1 Experiment

The substrates were p-type, 10 Ωcm, (100) silicon wafers supplied by Okmetic Ltd. These were thermally dry oxidized at 1000 °C in order to grow a 100-nm thick insulator layer. The thermal oxide was patterned using photolithography and buffered HF. Samples were cleaned using a chemical cleaning process (5 minute ultrasonic in acetone and isopropyl alcohol, 10 minutes at 80 °C in H₂O:NH₄OH:H₂O₂ (5:1:1) and

H₂O:HCl:H₂O₂ (5:1:1)).

After the samples had been inserted into the UHV growth chamber the native oxide was removed *in situ* by raising the temperature to 780 °C and allowing a low silicon flux etch the oxide off from the openings. Oxide removal was verified by monitoring the surface by RHEED as described earlier.

UHV plasma oxidation was performed by feeding pure oxygen gas through a plasma cell in which RF power was 265 W. The pressure in the chamber during oxidation was about 5·10⁻⁵ mbar, and the oxygen flow was 1.2 sccm. The background pressure in our MBE chamber was around 10⁻¹⁰ mbar. In high temperature oxidation the substrate temperatures were adjusted by the resistive heater. The room temperature samples were fabricated as described in chapter 5.4.1. No post oxidation annealing was applied to the samples.

At temperatures below 850 °C and in a UHV environment, the initial oxidation rate on a clean, good quality silicon surface is about the same regardless of temperature until the silicon dioxide layer thickness exceeds 1 nm. This may take a relatively long time, up to 20 minutes¹. From that point on the effect of temperature becomes significant, with the oxidation rate increasing with increasing temperature. At room temperature oxidation virtually stops after one nanometer², when around 800 °C it reaches the maximum growth rate after that¹. In the higher temperature oxidation process (600 - 800 °C) the layer thickness can be defined by adjusting the oxidation time, although below 650 °C the oxidation is really slow. It may take as long as 30 minutes to grow a 2-nm thick oxide at 600 °C.

Once the predetermined silicon dioxide layer thickness was achieved, i.e. 2 - 5 nm, an aluminum layer of about 100 nm was evaporated on top of the sample. Following the removal of the sample from the vacuum chamber, the test structures were patterned by using photolithography and wet chemical etching (Figure 5.8). The area of the devices is about (100 μm)².

The quality of oxide usually improves with increasing oxidation temperature. The best quality ultrathin layers have been fabricated by thermal oxidation at around 1000 °C, or

by using some post oxidation annealing (above 350 °C, for instance ¹⁶¹). However, the equipment we used limited oxidation temperatures to below 900 °C. The use of high temperatures is also prohibited by the ultra high vacuum environment; if the temperature is too high, the surface will be etched rather than oxidized ¹⁵⁴. In our experiments ¹ we have found that the native oxide layer is vaporized from the silicon surface at 870 °C. Oxide evaporation from the surface increases surface roughness, at least if the temperature is below 800 °C ⁷⁷.

Studies of the quality of these 2 – 5 nm thick UPO layers have shown that the difference between layers oxidized at 600 - 800 °C is quite small. The measured surface state densities in the mid-gap are between 0.4 - 3 10^{11} eV⁻¹cm⁻² (Figure 5.13). Flat band voltages are about -0.55 V (Figure 5.14). This corresponds to an interface charge of about $2 \cdot 10^{12}$ cm⁻². Given that we are dealing with extremely small dimensions, even minor changes in the measured values have a relatively major impact on the calculated parameters. Therefore absolute values are uncertain, but these values can still be used for purposes of qualitative comparison between different fabrication methods.

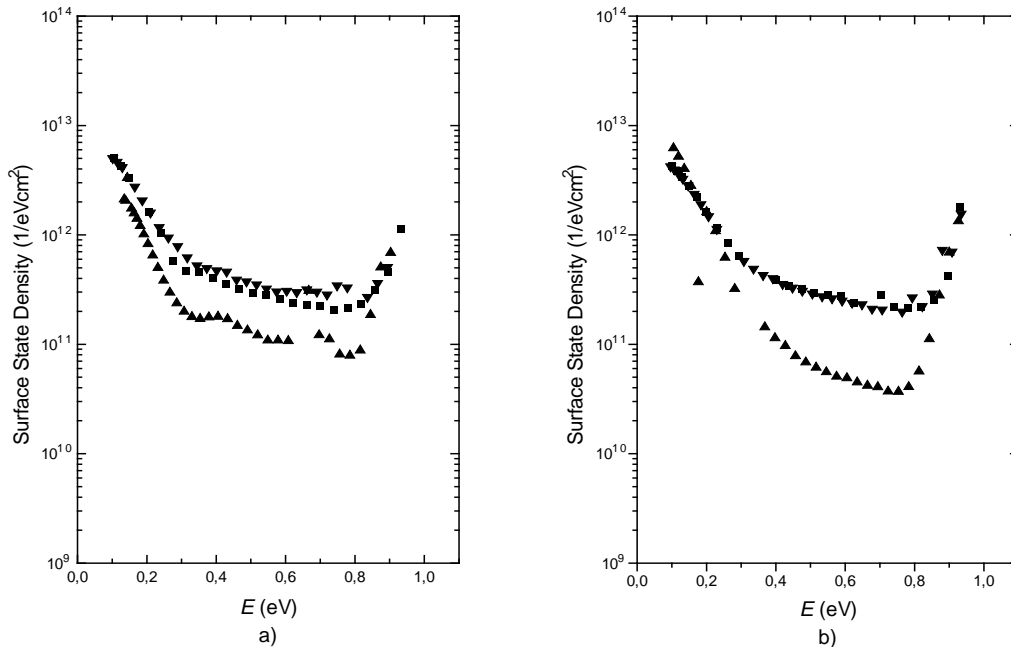


Figure 5.13 Surface state densities from samples plasma oxidized at a) 700 °C (oxide thickness: up triangles 2.5 nm, squares 3 nm and down triangles 3.4 nm) and b) 800 °C (oxide thickness: up triangles 2 nm, squares 4.2 nm and down triangles 4.4 nm). ⁴

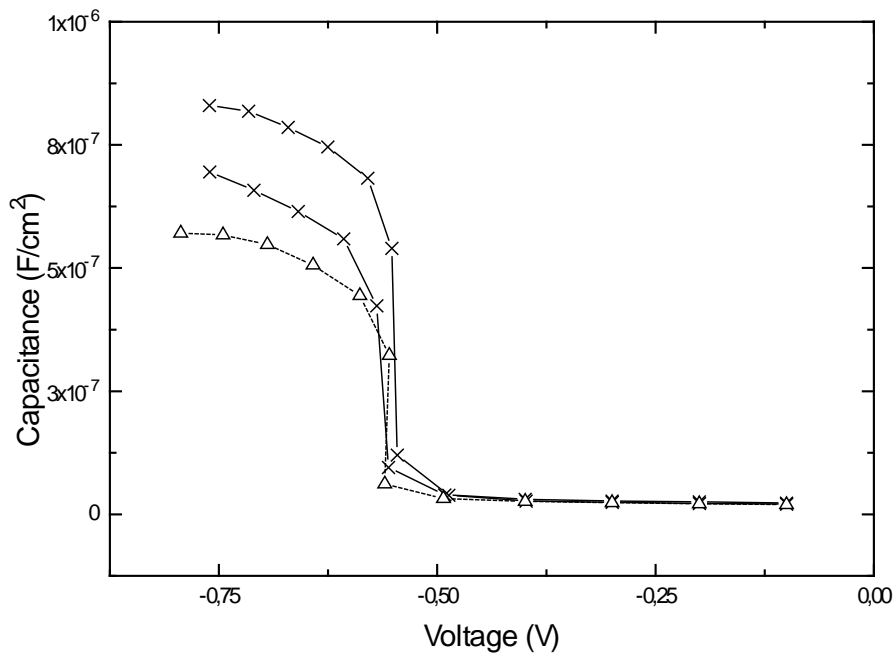


Figure 5.14 Capacitance as a function of gate voltage from samples plasma oxidized at 700 °C (crosses) and 800 °C (triangles).⁴

The interface state densities seem to be more or less the same regardless of the temperature that is used in oxidation⁴ (Figure 5.15 and Figure 5.16). The lowest surface state densities are usually measured from samples with oxide layers of about 3 nm or little below. If the oxide thickness is below 2.5 nm, the average surface state densities tend to be slightly higher than the values measured from 5 nm samples. This phenomenon might also be due to increased leakage through the oxides which disturbs the measurements. In some of these samples a peak can be seen at about 0.7 - 0.8 eV above valence band (Figure 5.16). This corresponds to the oxygen and silicon impurity states in silicon (0.72 and 0.78 eV, respectively⁴⁶), which are usually identified as being caused by dangling bonds. We have previously seen this phenomenon in room temperature oxides (chapter 5.4.1), but sometimes it can also be found after high temperature oxidation. The surface state densities presented in Figure 5.16 are measured from samples fabricated in the same way; the only difference between them is in oxide thickness, measuring a few Angstrom. We have yet been unable to establish why the peak that is seen in the other curve can sometimes be detected and sometimes not. However, it seems that the probability of it occurring increases when the layer thickness decreases to below 3 nm.⁴

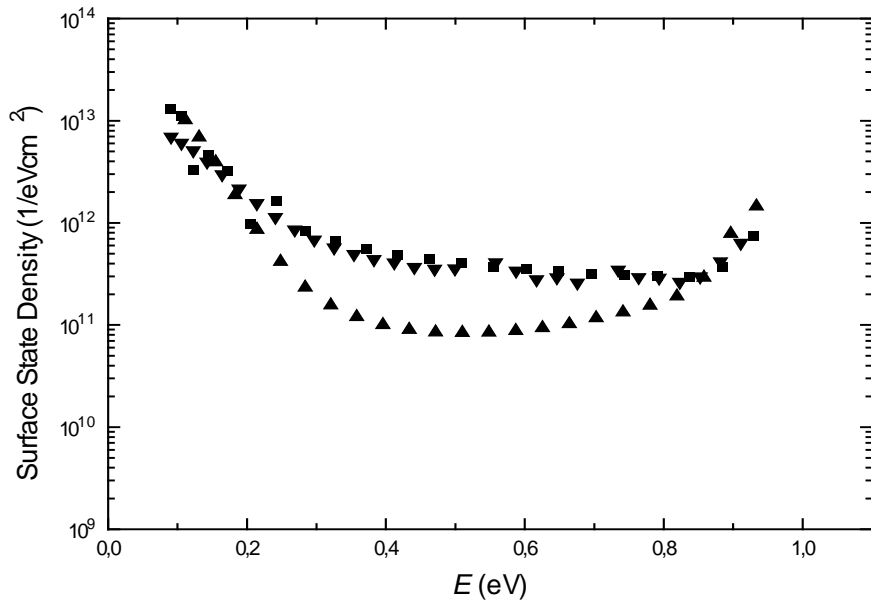


Figure 5.15 Surface state densities from samples plasma oxidized at room temperature (oxide thickness: down triangles 2 nm, up triangles 3 nm and squares 5 nm).⁴

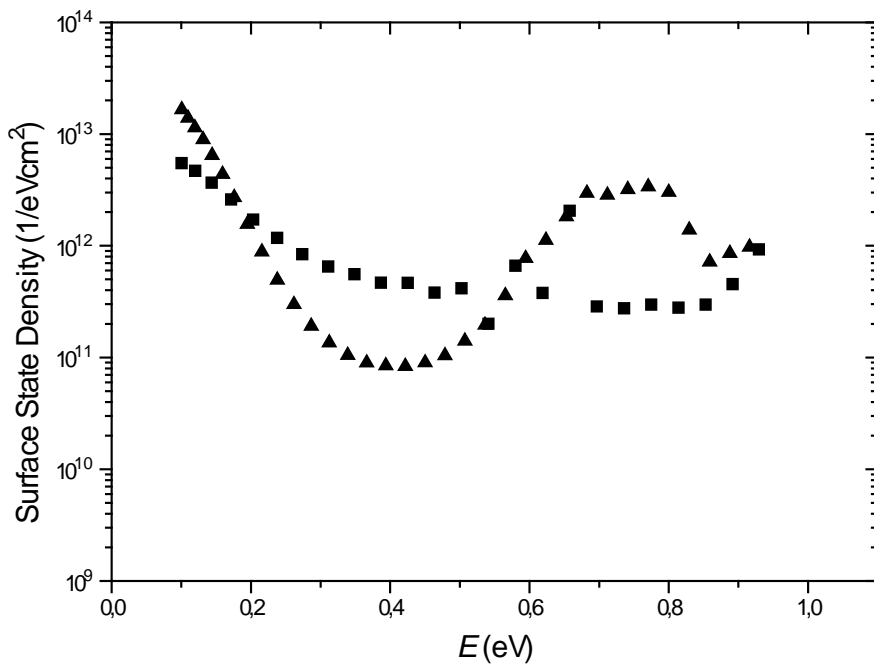


Figure 5.16 Surface state densities from two MOS samples with about 3 nm silicon dioxide layers, both plasma oxidized in 700 °C.⁴

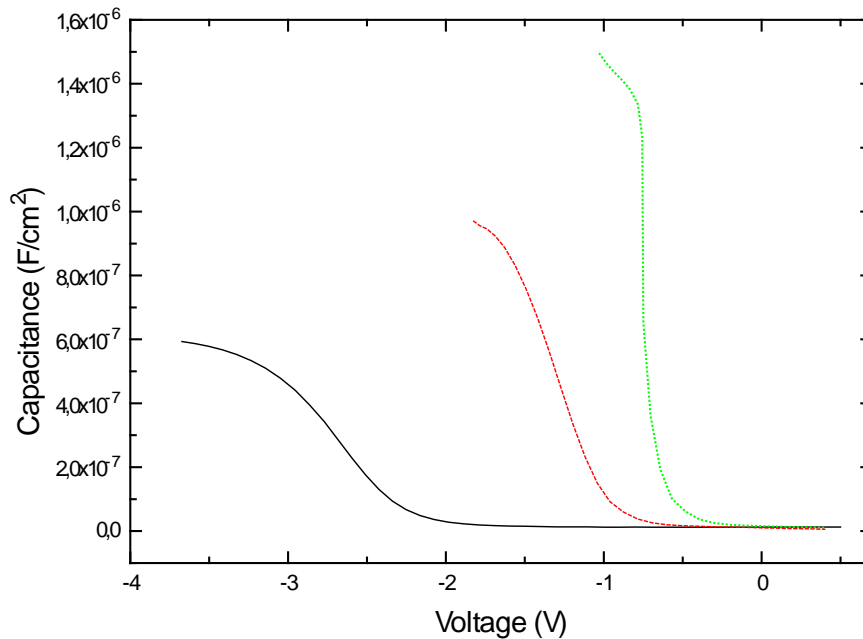


Figure 5.17 Capacitance as a function of gate voltage from samples plasma oxidized at room temperature. (Oxide thickness: solid black line 5 nm, dashed red line 3 nm and dotted green line 2 nm).⁴

The oxide layer thickness has a major influence on the flat band voltage if the oxidation is performed at room temperature (Figure 5.17). No such dependence can be seen in high temperature oxide measurements (Figure 5.14). The oxide charge can be calculated from the flat band shift as described in chapter 4.1. Assuming that the interface charge is constant in every sample, the extrapolation of the thickness dependence of the charge gives negative values for the interface charge, at least compared to the high temperature oxides. The oxide itself, on the other hand, seems to be highly positively charged. It seems that unit charge density can reach values as high as 10^{19} cm^{-3} .⁴

5.6 The Effect of Post Oxidation Annealing

As was discussed earlier self-terminating room temperature plasma oxidation of silicon seems to be quite an interesting tool that can be used in the *in situ* fabrication of silicon/silicon dioxide heterostructure devices². Annealing is a standard procedure in the improvement of oxide quality. The fabrication of silicon/silicon dioxide

heterostructures also requires certain thermal treatments in order to turn the amorphous or multicrystalline silicon into single crystalline material.

The use of a UHV MBE system limits the use of post oxidation high temperature treatments. Even at 870 °C the oxide is vaporized from the surface¹. As the metal we are using is aluminum, it is also necessary to perform the post metallization heat treatment at low temperatures; about 350 °C is preferred for ultrathin oxides¹⁶¹. Excessive temperatures increase the impurity diffusion and the spiking effect. We studied the effect of two different post oxidation annealing procedures on the electrical characteristics of thin oxide MOS structures.

5.6.1 Experiment

Three-inch boron doped, 20 mΩcm, (100) silicon wafers were used as substrates. These were thermally dry oxidized at 1000 °C to make a 100-nm thick insulator layer. Openings through this layer were etched using photolithography and buffered HF. After an HF dip, samples were inserted in the VG Semicon V80M MBE system and the sample temperature was raised to 780 °C. Oxide removal was verified by monitoring the surface by RHEED. The temperature was then allowed to drop to room temperature. The actual oxidation was performed as described in chapter 5.4.1. RF power was 265 W. The pressure in the chamber during oxidation was $3 \cdot 10^{-5}$ mbar, and the oxygen mass flow was 1.2 sccm. An aluminum layer of about 100 nm was evaporated *in situ* to form top contact.

The results presented are measured from samples that were either post oxidation (pre metallization) annealed at 750 °C in UHV for 15 minutes, or post metallization annealed at 300 °C in air. For comparison a set of samples were fabricated without any post oxidation thermal treatments³. After removal from the vacuum chamber, the test structures (capacitor area $(100 \mu\text{m})^2$) were patterned by using photolithography and wet chemical etching (Figure 5.8).

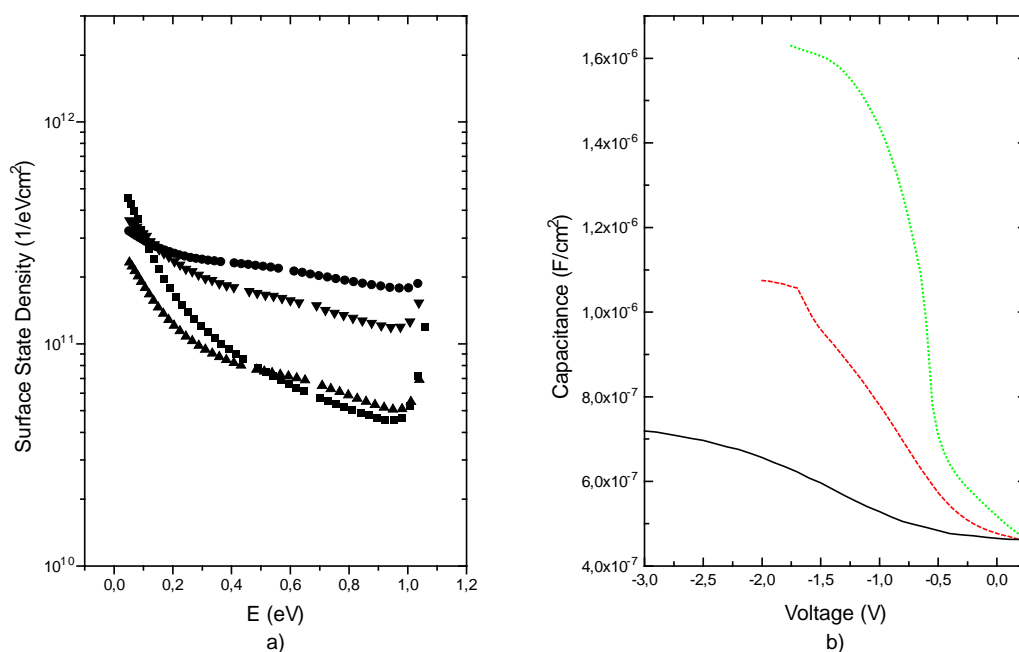


Figure 5.18 Measured a) surface state densities (squares 2 nm, up triangles 3 nm, down triangles 4.6 nm and circles 5 nm oxide film) and b) high frequency (1 MHz) capacitances (dotted green line 2 nm, dashed red line 3 nm and solid black line 5 nm oxide) from samples fabricated without any post oxidation annealing.³

As can be seen in Figure 5.18a, it is possible to produce fairly good surface state densities, below $10^{11} \text{ eV}^{-1}\text{cm}^{-2}$ in the mid-gap, even without any annealing. The oxide charge, on the other hand, may rise to quite a high level⁴. It can be seen in Figure 5.18b, that the flat band voltage shifts as the oxide thickness changes, which means that there exists some charge not only at the interface but also in the oxide. The measured flat band voltage is about 0.8 V in the 2 nm oxide samples, for 5 nm oxide it can even exceed 2 V. Our measurements indicate³ that the oxide charge density can be as high as 10^{19} cm^{-3} (unit charge in a volume) in room temperature UHV oxides and that there exists some negative interface charge.

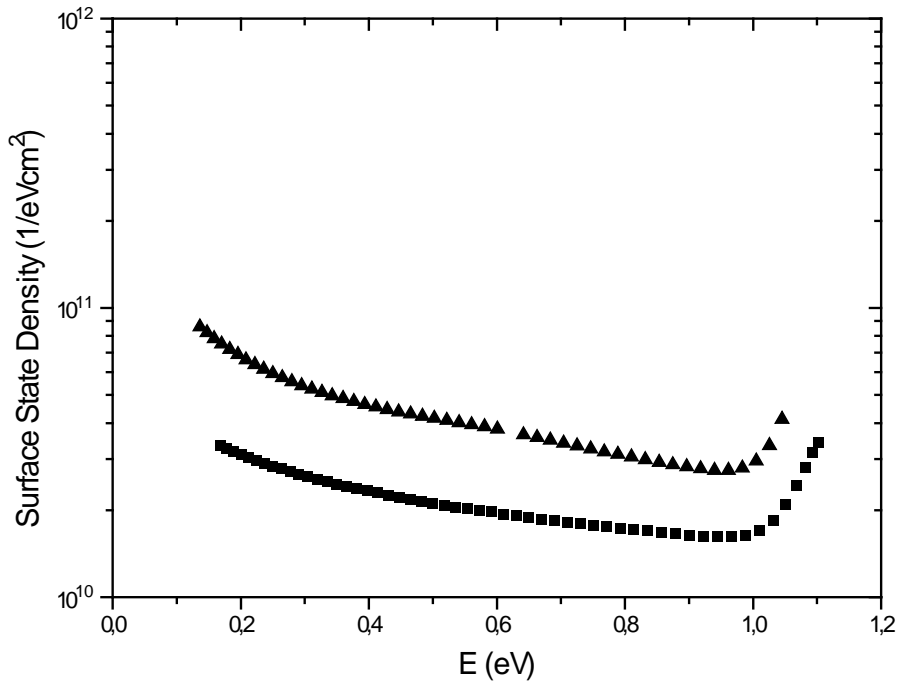


Figure 5.19 Measured surface state density after annealing at 750 °C (triangles) and 300 °C (squares). Film thickness around 4 nm.³

After annealing the surface state densities are still about the same or slightly better than before it (Figure 5.19). Even 300 °C seems to be enough to improve surface state density. The oxide charge, on the other hand, changes more (Figure 5.20b). It can be almost completely removed by annealing at 750 °C. At the same time the interface charge shifts in a positive direction. It is more difficult to analyze the effect of post metallization annealing. The depletion region capacitance seems to behave quite well: in every sample the capacitance curve starts to rise at the same voltage value (Figure 5.20a). The accumulation capacitance, by contrast, only reaches its saturation value at rather high voltages. This may be caused by impurities from the metal or from the furnace, which act as a moving charge. Adjustment of the annealing time could help the results. An encouraging observation was that neither of these annealings decrease the layer thickness.³

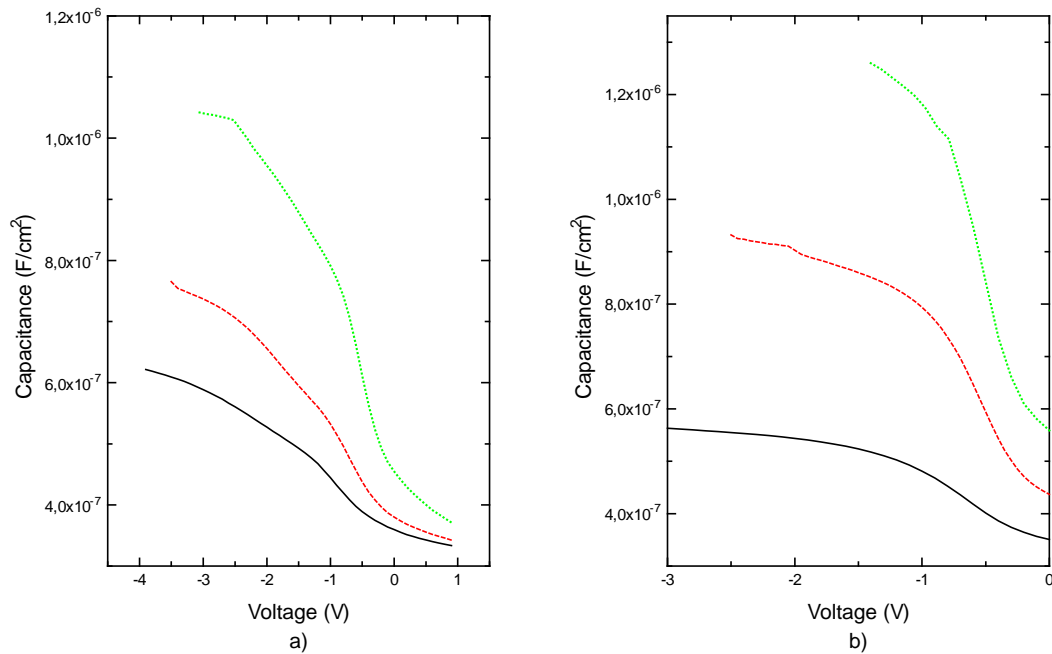


Figure 5.20 Measured high frequency capacitances after annealing at a) 300 °C and b) 750 °C³ (oxide film thicknesses from 2.6 to 5.8 nm).

5.7 Evaporation Technique

A major drawback of the room temperature process is that it is quite slow. The maximum thickness that can be reached is no more than about one nanometer¹². Therefore if thicker layers are required, the oxidation process will have to be repeated a sufficient number of times, with a thin, no more than 0.6 nm, silicon layer evaporated on top of the sample between each oxidation cycle². We have experimented with a method in which oxidation and silicon evaporation are performed simultaneously. This significantly reduces the processing time. A quite similar approach has recently also been used elsewhere. In the so-called low-temperature low-pressure PEE (Plasma-enhanced Evaporation) process, argon is fed into a plasma source as the working gas for the discharge, while oxygen is let directly into the chamber^{162,163}. The cracked oxygen molecules then oxidize the silicon that is simultaneously evaporated. The surface state densities reached are at the level of $10^{11} - 10^{12} \text{ eV}^{-1} \text{ cm}^{-2}$ and the oxide charge $10^{11} - 10^{12} \text{ cm}^{-2}$. The material density, however, is presumed to be low¹⁶². The relative

dielectric constant (ϵ_{ox}) can reach values as high as 8 - 10, which can be assumed to be due to the oxide structural change and, as the net charge is low, local polarization.

5.7.1 Experiment

The substrates used in this study were p-type, 10 Ω cm, (100) silicon wafers. Samples were cleaned using a chemical cleaning process (5 minute ultrasonic in acetone and isopropyl alcohol, 10 minutes at 80 °C in H₂O:NH₄OH:H₂O₂ (5:1:1) and H₂O:HCl:H₂O₂ (5:1:1)). The processing tool used was the same modified VG Semicon V80M molecular beam epitaxy (MBE) system as employed earlier¹⁵³. In this process silicon is electron beam evaporated from a cell after the oxygen plasma has been turned on. The oxygen is fed through an inlet, controlled by a mass flow controller. The gas is activated in a plasma cell by using a 265 W RF-field.

Once the samples had been inserted into the UHV growth chamber the native oxide was removed *in situ* by raising the temperature to 780 °C and allowing a low silicon flux etch the oxide off. Oxide removal was verified by monitoring the surface by RHEED.

Great care has to be exercised when working with plasma. Although it increases the process rate, it can also on the other hand damage the surface. One way to avoid this is to use some remote plasma-enhanced process (e.g. ¹⁶²) in which a working gas is fed into the plasma source and process gas directly into the chamber. In our case we fed the oxygen directly through the plasma cell, while a silicon flux was generated by e-beam evaporation into the chamber. This simplifies the process and the equipment set-up. To avoid creating damage, we opted to use a slow room temperature process in a UHV chamber which decreases the energies of the oxygen molecules colliding with the surface. Also, we first grew a one nanometer thick silicon dioxide layer by oxidizing the silicon wafer surface by oxygen plasma at room temperature as described in chapter 5.4. This was done not just to form a protective layer but to improve the quality of the silicon/silicon dioxide interface. Another benefit is that an oxide layer even as thin as 1 nm acts as a diffusion barrier at room temperature¹⁶⁴. This allows for more accurate adjustment of layer thickness. This first layer was formed by feeding pure oxygen gas

for ten minutes through a plasma cell in which RF power was 265 W. The pressure in the chamber during oxidation was about $5 \cdot 10^{-5}$ mbar, the oxygen flow was 1.2 sccm.

As soon as the first oxide layer is complete, silicon evaporation is started. The oxide thickness is defined by adjusting the silicon flux and evaporation time. The silicon evaporation rate was defined by using a Sycon Instruments STC-200 Deposition Rate Controller. The silicon flux was 0.07 nm/s. Some previous samples have been characterized by Auger measurement, which has indicated that the use of these parameters produces silicon dioxide.

The samples were annealed in UHV conditions at 750 °C for 15 minutes. Some reference samples were fabricated without any annealing. On top of the silicon dioxide layer 100 nm of aluminum was evaporated. The test structures were then patterned by using photolithography and wet chemical etching (Figure 5.21). The area of the devices from which the presented results have been measured is about $(100 \mu\text{m})^2$.

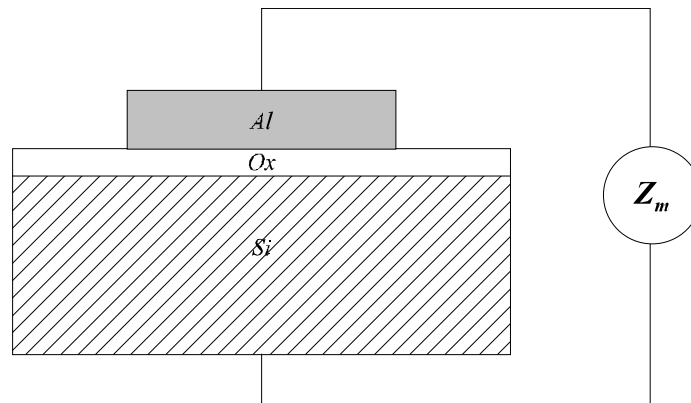


Figure 5.21 Schematic illustration of the MOS structure.

Our experiments have shown that the process described produces silicon dioxide layers with surface state densities of around $1 \cdot 10^{11} \text{ eV}^{-1} \text{ cm}^{-2}$ in the mid-gap even without any annealing (Figure 5.22). This result is the same as that achieved with the ultra high vacuum plasma oxidation process², which was used for the oxidation of the first nanometer thick layer. However, the oxide charge tends to be quite high, which can be seen from the flat band voltage shift in the high frequency capacitance measurements (Figure 5.23). Values of around 10^{13} cm^{-2} have been measured.

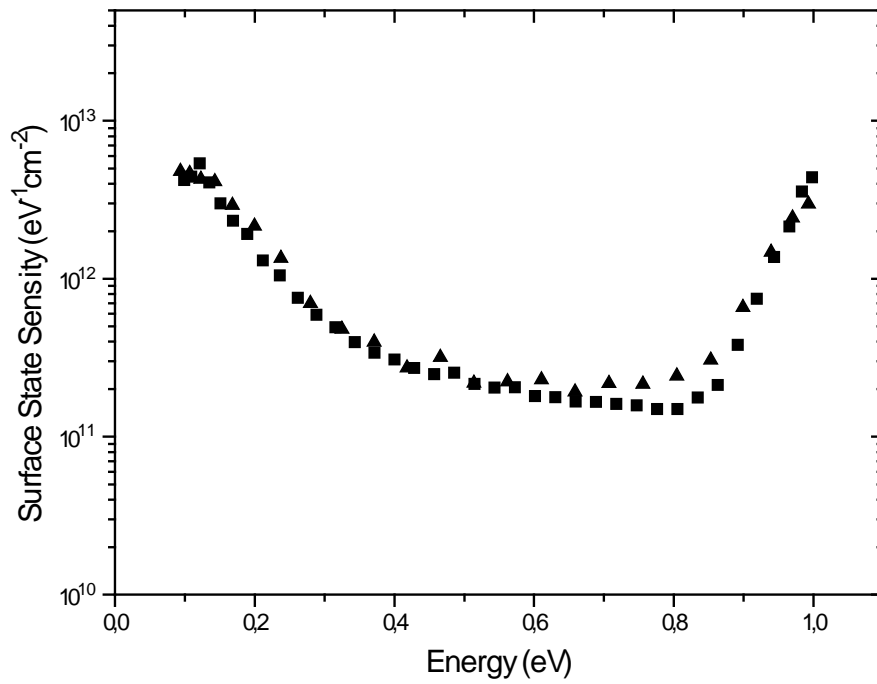


Figure 5.22 Surface state densities from MOS sample fabricated at room temperature without annealing.

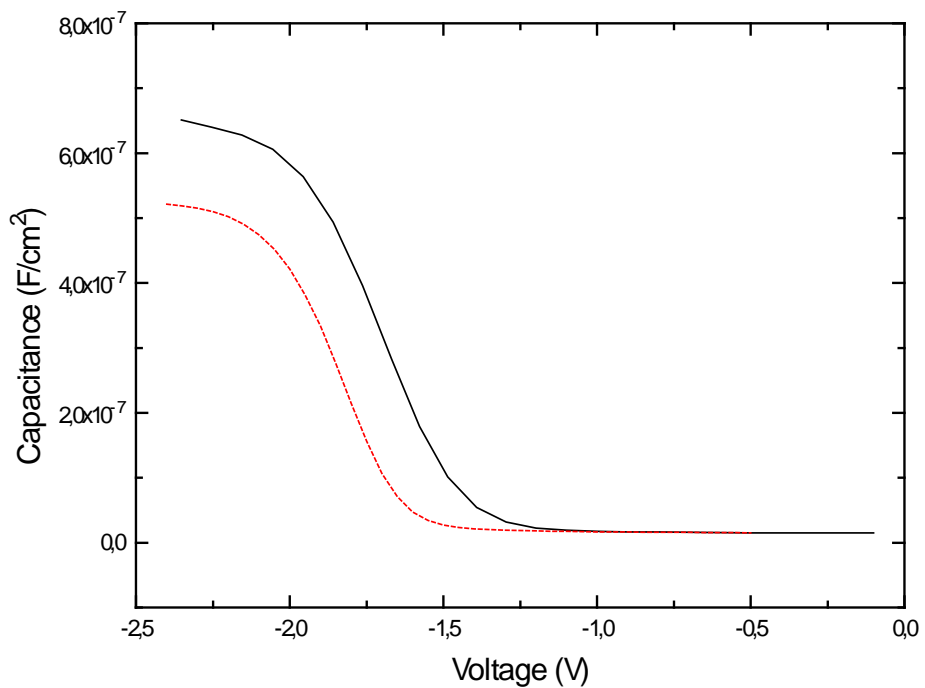


Figure 5.23 High frequency capacitance from thin oxide MOS samples fabricated at room temperature without annealing. Oxide thickness is 4.75 nm (solid black line) and 6 nm (dashed red line).

The oxide charge can be reduced by annealing at 750 °C (Figure 5.24) which may also decrease the surface state density but does not have any effect on layer thickness, as our previous results have shown (chapter 5.6). After annealing, however, there may appear a small peak around 0.8 eV above the valence band in the surface state densities measurements (Figure 5.25). This can be interpreted as an indication of breakage of some silicon - oxygen bonds in the interface area ¹⁶⁰. However, lower temperatures or longer annealing times might improve the results.

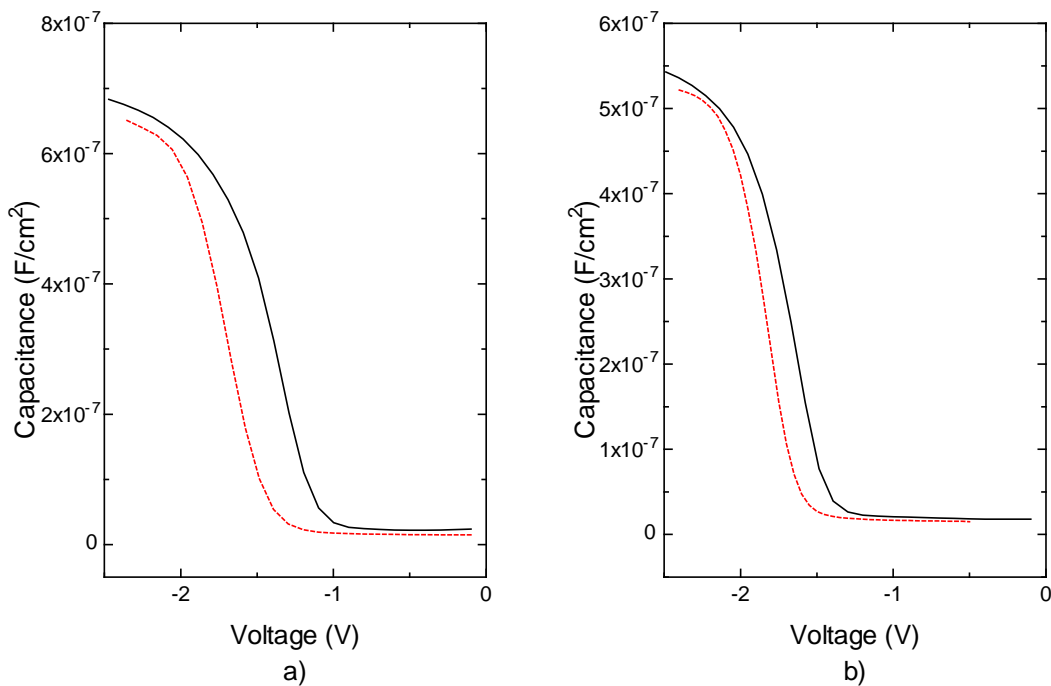


Figure 5.24 High frequency capacitance from thin oxide MOS samples (dashed red line before and solid black line after annealing). Oxide thickness is a) 4.75 nm and b) 6 nm.

The silicon dioxide layer growth rate was estimated on the basis of the silicon flux. It was assumed that when the flux is 0.07 nm/s, it takes about 10 seconds to grow 1 nm of silicon dioxide. However, the measured electrical thickness of the layers exceeded the predictions. It seemed that the first two nanometers grew quite rapidly, after which the growth rate saturated and came quite close to what was predicted.

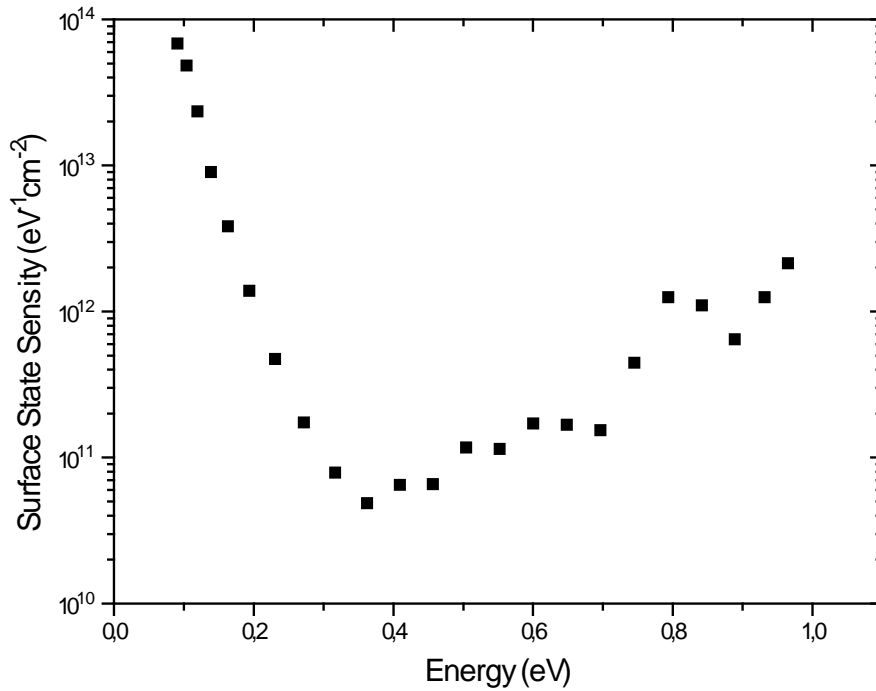


Figure 5.25 Surface state densities from sample fabricated at room temperature and annealed at 750 °C.

There is a problem then with the adjustment of layer thickness. In the case of layers that are no more than a few nanometers thick, 2-nm onset is much too large. It is possible that the first nanometer does not prevent the diffusion of silicon monoxide after all, but this does not sound right as it stops the oxygen diffusion². Also, the surface state density does not increase during the process, which indicates that the interface probably remains more or less unchanged. The most probable explanation then is that the shutter mechanism and the evacuation pumping in the growth chamber are not efficient enough. It follows that there occurs some oxidation/growth before and after the actual growth phase, which increases the layer thickness. This effect can probably be eliminated simply by equipment development.

6 Conclusions

We have fabricated 1 - 10 nm thick silicon dioxide layers by plasma oxidation of silicon in ultra high vacuum (UHV). The UHV chamber is a very clean environment where both temperature and molecule fluxes are easy to control. The main benefit of the room temperature MBD (Molecular Beam Deposition) method is that it allows us to fabricate versatile nanosize silicon-based heterostructures *in situ*.

In the temperature range 700 - 850 °C, oxidation is initially very slow. Once an uniform layer has developed on the surface, the growth rate increases. At this stage the increasing temperature also acts to accelerate the growth rate. During initial growth the temperature dependence is probably the exact opposite. In thermal oxidation the oxide quality improves with increasing oxidation temperature. Our MBE system, however, limits the oxidation temperatures to below 850 °C. Because of the oxidation rate, on the other hand, we have to use temperatures over 650 °C to achieve thicknesses over 2 nm, otherwise the oxidation procedure has to be performed gradually one nanometer at a time. Our measurements predict that in order to get a good quality as-grown layer with this process, the oxidation temperature should be at least 700 °C.

Since high temperatures are not always desirable in processing, we have also studied room temperature oxidation. The room temperature ultrahigh vacuum plasma oxidation method has been proven to produce good quality ultrathin silicon dioxide layers, with quite accurately controllable layer thickness, relatively high breakdown fields and low surface state densities. It is predicted, that as higher temperatures smoothen the surface, lower temperatures do not create as much roughness in the first place. At the room temperature the oxidation saturates at around 1 nm which means that the fabrication process can be so designed that it is fully automated, even if the system incorporates no thickness measurement of amorphous layers. Also, processing does not require high temperatures. The only exception is native oxide removal, where we use a temperature of 780 °C. However, even this can be avoided by using chemical etching and passivation of the sample surface. The precise effects of this on layer quality remain to be seen, though.

The ultrahigh vacuum plasma oxidation method seems to produce quite homogenous silicon/silicon dioxide interface state densities, regardless of the oxidation temperature. In layers oxidized at room temperature, however, the oxide charge tends to rise to quite a high level. Some kind of annealing should therefore be used to decrease the oxide charge. Our measurements showed that post oxidation UHV annealing at 750 °C and post metallization annealing at 350 °C in air slightly decrease the surface state densities. The oxide charge, which is quite high in these oxides, can be decreased by UHV annealing. The effect of post metallization annealing on the oxide charge is less clear. This is probably due to impurity diffusion. Neither of the annealings used decrease the oxide layer thickness. This backs up our experiments which show that the desorption of silicon dioxide from the silicon surface starts quite abruptly at 870 °C under a background pressure of 10^{-10} mbar¹. This means that temperatures at least up to 750 °C can safely be used when there is no protective layer covering the silicon dioxide film.

The silicon dioxide layers fabricated by evaporating silicon in a low pressure oxygen plasma ambient seem to have the same surface state densities as layers fabricated by the room temperature ultrahigh vacuum plasma oxidation method. The main benefit achieved with this process is the increased growth rate. It also allows the oxide thickness to be varied over a wider range. The main drawback is the high oxide charge, but that can be lowered by annealing.

In our studies we have tested the UPO method to produce high quality silicon dioxide films. The benefits of this method are good thickness control, a pure growth environment, highly accurate material fluxes, and the possibility of *in situ* device processing. However, the as-grown layers, especially those oxidized at room temperature have a very high oxide charge, even though the interface state density is reasonably low. The charge can affect on the leakage and the film stability in stressed conditions can weaken. Our preliminary experiments have shown that the layer quality can be improved by annealing, but the process needs to be further developed. Because various oxide fabrication procedures worldwide have been able to produce very high quality films, the usefulness of our UPO method remains controversial in conventional IC fabrication. Various thermal oxidation methods (most notably rapid thermal oxidation) have proven to be accurate and reliable and to produce an extremely high oxide quality. Nonetheless there do remain other purposes for our process. Accurately

controlled system can supply much information about the oxidation process itself. It also can be used in device fabrication and then the most logical choice seems to be the fabrication of the silicon/silicon dioxide heterostructures. For this purpose the process has to be carefully adjusted and studied because the interface quality has to be even better than in MOSFETs so that really separate quantum states may be formed. The possibility of making fully crystalline structures seems to be particularly interesting. The most probable crystalline form of SiO_2 in low pressures in the temperature range 870 - 1470 °C is tridymite¹⁶⁵. However, processing will require a deep understanding of the oxidation mechanisms and the parameters affecting these mechanisms. In addition, the substrates used have to be of a very high quality and low resistivity, with epitaxial buffer layers to form an atomically flat surface. It is likely that even quite small changes in oxidation temperatures or oxidation rates will produce different oxide species. Surface roughness causes local tensions during oxidation and in this way prevents homogenous film formation. Traditionally hydrogen has been used to passivate dangling bonds, but this is probably not the best method to use when fabricating quantum devices because it can affect the interface charge as well as interface state generation. Nitrogen incorporation into the oxide also has many beneficial properties, but it can have an adverse effect on interface quality.

References

- 1 T. Majamaa, V.-M. Airaksinen, J. Sinkkonen, *Appl. Surf. Sci.* 107 (1996), 172
- 2 T. Majamaa, O. Kilpelä, S. Novikov, *Appl. Surf. Sci.*, 136 (1998), 17
- 3 T. Majamaa, O. Kilpelä, S. Novikov, J. Sinkkonen, *Appl. Surf. Sci.* 142 (1999), 351
- 4 T. Majamaa, O. Kilpelä, *Physica Scripta T79* (1999), 259
- 5 W. Kiunke, E. Hammerl, I. Eisele, *J. Crystal Growth* 127 (1993) 73
- 6 H. Gossner, I. Eisele, L. Risch, *Jpn. J. Appl. Phys.* 33, Part I (1994) 2423
- 7 K. Watanabe, S. Kimura, T. Tatsumi, *Jpn. J. Appl. Phys.* 38 (1999) L1055
- 8 A. Strass, P. Bieringer, W. Hansch, V. Fuenzalinda, A. Alvarez, J. Luna, I. Martil, F.L. Martinez, I. Eisele, *Thin Solid Films* 349 (1999) 135
- 9 Y. Taur, P.A. Buchanan, W. Chen, D.J. Frank, K.E. Ismai, S.-H. Lo, G.A. Sai-Halasz, R.G. Viswanathan, H.-J. C. Wann, S.J. Wind, H.-S. Wong, *Proc. IEEE* 85 (1997), 486
- 10 S. Tiwari, F. Rana H. Hanfi, A. Hartstein, E. F. Crabbé, K. Chan, *Appl. Phys. Lett.* 68 (1996), 1377
- 11 P. Masson, G. Ghibaud, J.L. Autran, P. Morfouli, J. Broni, *Electr. Lett.* 34 (1998), 1977
- 12 S. Novikov, J. Sinkkonen, O. Kilpelä, S.V. Gastev, *J. Vac. Sci Technol. B*, 15 (1997) 1471
- 13 A.C. Seabaugh, C.-C. Cho, R.M. Steinhoff, T.S. Moise, S. Tang, R.M. Wallace, E.A. Beam, Y.-C. Kao, R.C. Bowen, W.R. Frensley, K. H. Park, Y. Okuno, 2nd International Workshop on Quantum Functional Devices, Matsue, Japan, May 23-25 1995
- 14 M. Hacke, H.L. Bay, S. Mantl, *Thin Solid Films*, 280 (1996), 107
- 15 J.H. Stathis, D.J. DiMaria, *Microelectronic Engineering*, 48 (1999), 395
- 16 H.S. Momose, M. Ono, T. Yoshitomi, T. Ohguro, S. Nakamura, M. Saito, H. Iwai; *IEEE Transactions on Electron Devices*, 43 (1996), 1233
- 17 A. Chin, W.J. Chen, T. Chang, R.H. Kao, B.C. Lin, C. Tsai, J.C.-M. Huang, *IEEE Electr. Dev. Lett.* 18 (1997) 417
- 18 B.E. Deal, *IEEE Trans. El. Dev.* ED-27 (1980), 606
- 19 D.M. Fleetwood, *IEEE Trans. Nucl. Sci.* NS-39 (1992) 269
- 20 M. J. Uren, S. Collins, M.J. Kinton, *Appl. Phys. Lett.* 54 (1989), 1448

-
- 21 D.A. Buchanan, D.J. DiMaria, *J. Appl. Phys.* 67 (1990) 7439
- 22 M.E. Zvanut, F.J. Feigl, W.B. Fowlerm, J. K. Rudra, P.J. Caplan, E.H. Poindexter,
J.D. Zook, *Appl. Phys. Lett.* 54 (1989), 2118
- 23 T. Ohmi, M. Morita, T. Hattori: *The Physics and Chemistry of SiO₂ and the Si-
SiO₂ Interface*; Plenum (1988), 413
- 24 H. Nohira, T. Hattori, *Appl. Surf. Sci.* 117/118 (1997), 119
- 25 H. Nohira, A. Omura, M. Katayama, T. Hattori, *Appl. Surf. Sci.* 123/124 (1998),
546
- 26 H. Kageshima, K. Shiraishi, *Phys. Rev. Lett.* 81 (1998), 5936
- 27 A. Pasquarello, M.S. Hybertsen, R. Car, *Nature* 396 (1998), 58
- 28 X. Chen, J.M. Gibson, *Appl. Phys. Lett.* 70 (1997), 1462
- 29 Devine, *Appl. Phys. Lett.*, 68 (1996), 3108
- 30 T. Hattori, H. Nohira, K. Takahashi, *Microelectronic Engineering*, 48 (1999), 17
- 31 N. Gonon, J. Ciavatti, C. Plossu, B. Balland, D. Barbier, A. Gagnaire,
Microelectronics Journal 24 (1993) 401
- 32 J. S. Johanssen, W. E. Spicer, *J. Appl. Phys.* 47 (1976), 3028
- 33 C. A. Fenstermaker, F. L. McCracking, *Surf. Sci.* 16 (1965), 85
- 34 K. Hirose, H. Nohira, T. Koike, K. Sakamo, T. Hattori, *Phys. Rev. B* 59 (1999)
5617
- 35 D-A. Luh, T. Miller, T.C. Chiang, *Phys. Rev. Lett.* 79 (1997), 3014
- 36 R. Haight, L.C. Feldman, *J. Appl. Phys.* 53 (1983), 4884
- 37 A. Pasquarello, *Microelectronic Engineering*, 48 (1999), 89
- 38 F. J. Grunthaner, P.J. Grunthaner, R.P. Vasques, B.F. Lewis, J. Maserjian, *Phys.
Rev. Lett.* 43 (1979), 1683
- 39 G. Hollinger, E. Bergigna, H. Chermette, F. Himpsel, D. Lohez, M. Lannoo, M.
Bensoussan, *Philos. Mag. B* 55 (1987), 735
- 40 Y. Iida, T. Shimura, J. Harada, S. Samata, Y. Matsuhita, *Surf. Sci.* 258 (1991),
235
- 41 I. Takashi, T. Shimura, J. Harada, *J. Phys.: Condens. Matter* 5 (1993), 6525
- 42 C. Kaneta, T. Yamasaki, T. Uchiyama, T. Uda, K. Terakura, *Microelectronic
Engineering*, 48 (1999), 117
- 43 A. Ourmazd, D.W. Taylor, J. A. Rentschler, J. Bevk, *Phys. Rev. Lett.* 59 (1087),
213

-
- 44 A. Pasquarello, M.S. Hybertsen, R. Car, *Phys. Rev. Lett.* 74 (1995), 1024
- 45 A. Pasquarello, M.S. Hybertsen, R. Car, *Appl. Phys. Lett.* 68 (1996), 625
- 46 S. M. Sze: *Semiconductor Devices: Physics and Technology* (Wiley, New York, 1985)
- 47 M. Lenzlinger, E.H. Snow, *J. Appl. Phys.* 40 (1969) 278
- 48 M. Hiroshima, T. Yasaka, S. Miyazaki, M. Hirose, *Jpn. J. Appl. Phys.* 33 (1994), 395
- 49 S. Zafar, Q. Liu, E.A. Irene, *J. Vac. Sci. Technol. A* 13 (1995), 47
- 50 A. Scarpa, A. Paccagnella, G. Ghidini, A. Vianello, E. Zanoni, *Solid-State Electronics* 41 (1997), 935
- 51 A. Ghetti, E. Sangiorgi, T. Sorsch, I. Kizilyalli, *Microelectronic Engineering*, 48 (1999), 31
- 52 J. Ahn, A. Joshi, G.Q. Lo, D.-L. Kwong, *IEEE El. Dev. Lett.* 13 (1992), 513
- 53 M. Depas, T. Nigam, M.M. Heyus; *IEEE Transactions on Electron Devices*, 43 (1996) 1499
- 54 E. Harari, *J. Appl. Phys.* 49 (1978), 2478
- 55 S. Holland, I.C. Chen, T.P. Ma, C. Hu, *IEEE El. Dev. Lett.* 5 (1984) 302
- 56 J.J. Tzou, C.C. Yao, R. Cheung, H. Chan, *IEEE El. Dev. Lett.* 7 (1986), 446
- 57 M.-S. Liang, S. Haddad, W. Cox, S. Cagnina, *IEDM Tech. Dig.* (1986) 394
- 58 S. Ikeda, M. Okihara, H. Ucida, N. Hirashita, *Jpn. J. Appl. Phys.* 36 (1997), 2561
- 59 D.J. Dumin, J.R. Maddux, *IEEE Trans. El. Dev.* 40 (1993), 986
- 60 J. Maserjan, N. Zamani, *J. Vac. Sci. Techn.* 20 (1982), 743
- 61 P. Olivo, T.N. Nguyen, B. Ricc6, *IEEE Trans. El. Dev.* 35 (1988), 2259
- 62 G.W. Rubloff, K. Hofmann, M. Liehr, D.R. Young, *Phys. Rev. Lett.* 58 (1987), 2379
- 63 D.J. DiMaria, E. Cartier, *J. Appl. Phys.* 78 (1995), 3883
- 64 D.M. Fleetwood, P.S. Winokur, L.C. Riewe, R.A. Reber Jr, *J. Appl. Phys.* 84 (1998), 6141
- 65 M. Kerber, U. Schawalke, G. Innertsberger, *Microelectronic Engineering*, 48 (1999), 147
- 66 G. Tao, R. C6p, J. Dijkstra, F. Kuper, R. Verhaar, *Microelectronic Engineering*, 48 (1999), 419
- 67 Y. Homma, M. Suzuki, N. Yabumoto, *J. Vac. Sci & Technol. A*10 (1992), 2055

-
- 68 M. Itsumi, F. Kiyosumi, *Appl. Phys. Lett.* 40 (1982), 496
- 69 P. Mertens, T. Bearda, M. Houssa, L.M. Loewenstein, I. Cornelissen, S. De Gendt, K. Kenis, I. Teerlinck, R. Vos, M. Meuris, M.M. Heyns, *Microelectronic Engineering*, 48 (1999), 199
- 70 Y. Kobayashi, Y. Shinoda, K. Sugii, *Jpn. J. Appl. Phys.* 29 (1990), 1004
- 71 W. Hansh, E. Hammerl, W. Kiunke, I. Eisele, J. Raum, E. Beck, *Jpn. J. Appl. Phys.* 33, Part I (1994) 2263
- 72 K.E. Johnson, P.K. Wu, M. Sander, T. Ergel, *Surf. Sci.* 290 (1993), 213
- 73 G.D. Wilk, Y. Wei, H. Edwards, R.M. Wallace, *Appl. Phys. Lett.*, 70 (1997), 2288
- 74 Y. Homma, H. Hibino, T. Ogino, N. Aizawa, *Physical Review B*, 55 (1997), R10237
- 75 M. Niwa, T. Kouzaki, K. Okada, M. Udagava, R. Sinclair, *Jpn. J. Appl. Phys.* 33 (1994), 388
- 76 R.A.B. Devine, W.L. Warren, J.B. Xu, I.H. Wilson, P. Paillet, J.-L. Leray, *J. Appl. Phys.* 77 (1995), 175
- 77 J.L. Hansem, S.Y. Shiryaev, E.V. Thomsen, *Journal of Crystal Growth*, 157 (1994) 317
- 78 T. Ohmi, K. Matsumoto, K. Nakamura, K. Makihara, J. Takano, K. Yamamoto, *Appl. Phys. Lett.* 62 (1993), 405
- 79 M. Tabakomori, H. Ikoma, *Jpn. J. Appl. Phys.* 36 (1997), 5409
- 80 H. Fukuda, M. Yasuda, T. Iwabuchi, S. Kameko, T. Ueno, I. Ohdomari, *J. Appl. Phys.* 72 (1992), 1906
- 81 L. Do Tanh, P. Balk, *J. Electrochem. Soc.* 135 (1988), 1797
- 82 A.H. Carim, R. Sinclair, *J. Electrochem. Soc.* 134 (1987), 741
- 83 E.P. EerNisse, *Appl. Phys. Lett.* 35 (1979), 8
- 84 M. Itsumi, O. Nakajima, N. Shiono, *J. Appl. Phys.* 72 (1992), 2185
- 85 M. Depas, R.L. Van Meirhaeghe, F. Cardon, *Solid-State Electronics* 37 (1994) 433
- 86 G. Eftekhari, *J. Electrochem. Soc.* 140 (1993) 787
- 87 T. Ohmi, M. Morita, A. Teramoto, K. Makihara, K. S. Tseng, *Appl. Phys. Lett.*, 60 (1992), 2126
- 88 A.C. Adams, T.E. Smith, C.C. Chang, *J. Electrochem. Soc.* 127 (1980) 1787

-
- 89 R.R Razouk, B.E. Deal, *J. Electrochem. Soc.* 126 (1979) 1573
- 90 K.H. Lee, W.H. Liu, S.A. Campbell, I. Banerjee, *J. Electrochem. Soc.* 140 (1993) 501
- 91 M. Akizuki, J. Matsuo, S. Ogasawara, M. Harada, A. Doi, I. Yamada, *Jpn. J. Appl. Phys.* 35 (1996), 1450
- 92 V. Ramgopal Rao, W. Hansch, S. Mahapatra, D.K. Sharma, J. Vasi, T. Grabolla, I. Eisele, *Microelectronic Engineering*, 48 (1999), 223
- 93 T. Fuyki, S. Muranaka, H. Matsunami, *Appl. Surf. Sci.* 117/118 (1997), 123
- 94 P.C. Chen, K.Y.J. Hsu, H.L. Hwang, J.Y. Lin, *J. Appl. Phys.* 76 (1994), 5508
- 95 S. Kimura, E. Murakami, T. Warabisako, H. Sunami, T. Tokuyama, *IEEE Electr. Dev. Lett.* 7 (1986) 38
- 96 R. Bartels, A.G. Aberle, R. Hezel, 14th European PVSEC 1997, Barcelona
- 97 S.S. Todorov, E.T. Fossum, *J. Vac. Sci. Technol. B*, 6 (1988) 466
- 98 K.T. Sung, W.Q. Li, S.H. Li, S.W. Pang, P.K. Bhattacharya, *Electronic Letters*, 29 (1993) 277
- 99 H. Fukuda, S. Murai, T. Endoh, S. Nomura, *J. Appl. Phys.* 81 (1997) 1825
- 100 H. Hwang, M.-Y. Hao, J. Lee, V. Mathews, P.C. Fazan, C. Dennison, *Solid-State Electronics*, 36 (1993), 749
- 101 W.L. Hill, E.M. Voge., V. Misra, P.K. McLarty, J.J. Wortman, *IEEE Trans. El. Dev.* 43 (1996), 15
- 102 H. Yang, H. Niimi, Y. Wu, G. Lucovsky, J. Reister, J. Rowe, *Microelectronic Engineering*, 48 (1999), 307
- 103 Y.-L. Wu, J.-G. Hwu, *IEEE El. Dev. Lett.* 14 (1993), 1
- 104 J.S. Cable, R.A. Mann, J.C.S. Woo, *IEEE El. Dev. Lett.* 12 (1991), 128
- 105 J.S. Cable, J.S.C. Woo, *IEEE Trans. El. Dev. Lett.* 39 (1992) 607
- 106 B. Yuwono, T. Schloesser, A. Gschwandtner, G. Innersberger, A. Grassl, A. Olbrich, W.H. Krautschneider, *Microelectronic Engineering*, 48 (1999), 51
- 107 B. E. Deal, A. S. Grove, *J. Appl. Phys.* 36 (1965), 3770
- 108 S. Dimitrijević, H.B. Harrison, *J. Appl. Phys.*, 80 (1996), 2467
- 109 A.A. Demkov, O. F. Sankey, *Phys. Rev. Lett.* 83 (1999) 2038
- 110 S.S. Ang, S.R. Wilson, *J. Electrochem. Soc.* 134 (1987) 1254
- 111 R. Lathlaen, D.A. Diehl, *J. Electrochem. Soc.*, 116 (1969), 620
- 112 Z. Chen, K. Yasutake, A. Doolittle, A. Rohatgi, *Appl. Phys. Lett.* 63 (1993), 2117

-
- 113 W-S. Liao, S-C. Lee, *J. Appl. Phys.* 80 (1996), 1171
- 114 S.S. Ang, Y.J. Shi, W.D. Brown, *J. Appl. Phys.* 73 (1993) 2397
- 115 G.G. Fountain, R.A. Rudder, S.V. Haltangadv, R.J. Markunas, *J. Appl. Phys.* 63
(1988) 4744
- 116 A.P. Gurevich, M.K. Weldon, Y.J. Chabal, R.L. Opila, J. Sapjeta, *Appl. Phys.
Lett.* 74 (1999), 1257
- 117 V. Ramgopal Rao, W. Hansch, H. Baumgärtner, I. Eisele, P.K. Sharma, J. Vasi, T.
Grabolla, *Thin Solid Films* 296 (1997), 37
- 118 T. Yasudda, Y. Ma, S. Habermehl, G. Lucovsky, *Appl. Phys. Lett.* 60 (1992), 434
- 119 P. Masson, P. Morfouli, J. Autran, J. Wortmann, *Microelectronic Engineering*, 48
(1999), 211
- 120 U. Neuwald, A. Feltz, U. Memmert, R.J. Behm, *J. Appl. Phys.* 78 (1995), 4131
- 121 J.A. Bardwell, K.B. Clark, D.F. Mitchell, D.A. Bisailion, G.I. Sproule, B.
MacDougall, M.J. Graham, *J. Electrochem. Soc.* 141 (1993), 2135
- 122 J.A. Bardwell, N. Draper, P. Schmuki, *J. Appl. Phys.* 79 (1996), 8761
- 123 J.A. Bardwell, P. Schmuki, G.I. Sproule, D. Landheer, D.F. Mitchell, *J.
Electrochem. Soc.* 142 (1995), 3933
- 124 O. Sneh, M.L. Wisem A.W. Ott, L.A. Okada, S.M. George, *Surf. Sci.* 334 (1995)
135
- 125 J.W. Klaus, A.W. Ott, J.M. Johnson, S.M. George, *Appl. Phys. Lett.* 70 (1997)
1092
- 126 O. Sneh, S.M. George, *J. Phys. Chem.* 99 (1994) 4639
- 127 S. Morishita, Y. Uchida, M. Matsumura, *Jpn. J. Appl. Phys.* 34 (1995) 5738
- 128 K. Hara, Y. Suzuki, Y. Taga, *Jpn. J. Appl. Phys.* 18 (1979), L2027
- 129 W.K. Choi, C.K. Choo, Y.F. Lu, *J. Appl. Phys.* 80 (1996) 5837
- 130 Y. Wei, R.M. Wallace, A.C. Seabaugh, *Appl. Phys. Lett.* 69 (1996), 1270
- 131 J. Lander, J. Morrison, *J. Appl. Phys.* 33 (1962), 2098
- 132 R. Tromp, G. W. Rubloff, P. Balk, F.K. LeGoues, J. Van Loenen, *Phys. Rev. Lett.*
55 (1985), 2332
- 133 M. Liehr, J. E. Lewis, G.W. Rubloff, *J. Vac. Sci. Technol.* A5 (1987) 1559
- 134 Y. Wei, R.M. Wallace, A.C. Seabaugh, *Appl. Phys. Lett.* 69 (1996) 1270
- 135 W.L. Warren, D.M. Fleetwood, M.R. Shaneyfelt, J.R. Schwank, P.S. Winokur,
R.A.B. Devine, D. Mathiot, *Appl. Phys. Lett.* 64 (1994), 3452

-
- 136 R.A.B. Devine, D. Mathiot, W.L. Warren, D.M. Fleetwood, B. Aspar, Appl. Phys. Lett. 63 (1993), 2926
- 137 A. Stesmans, V.V. Afanas'ev, Microelectronic Engineering, 48 (1999), 113
- 138 A. Stesmans, V.V. Afanas'ev, Phys. Rev. B 54 (1996), R11129
- 139 W.L. Warren, K. Vandheusden, J.R. Scwank, D.M. Fleetwood, P.S. Winokur, R.A.B. Devine, Appl. Phys. Lett. 68 (1996), 2993
- 140 A. Hartstein, D.R. Youn, Appl. Phys. Lett. 38 (1981), 631
- 141 A. Stesmans, V.V. Afanas'ev, Microelectronic Engineering, 48 (1999), 131
- 142 R.S. Bauer, R.Z. Bachrach, L.J. Brillson, Appl. Phys. Lett. 37 (1980), 1006
- 143 M.H. Hecht, R.P. Vasques, F.J. Grunthner, N. Zamari, J. Maserjian, J. Appl. Phys. 57 (1985), 5256
- 144 P. Balk, The Electrochem. Soc. Meeting, Buffalo, NY, Oct. 10-14, 1965
- 145 K. Hofmann, G. W. Rubloff, D.R. Young, J. Appl. Phys. 61 (1987), 4584
- 146 T. Majamaa: Computing the Charge in the Inversion Layer of MOSFET; (in Finnish) Special Project in Materials Physics, Department of Technical Physics, Helsinki University of Technology (1992), p 27+7
- 147 K.S. Krisch, J.D. Buedo, L. Mauchanda, IEEE El. Dev. Lett. 17 (1996), 521
- 148 HP4192A Impedance Analyzer Instructions Manual
- 149 J. Ritola: Impedanssispektroskopia- ja kapasitanssimittausjärjestelmä; Sovelletun fysiikan erikoistyö, Teknillisen fysiikan osasto/Tkk (1994),pp 18+49
- 150 C.-Y. Hu, D. L. Kencke, S. Banerjee, B. Bandyopadhyay, E. Ibok, S. Garg, Appl. Phys. Lett. 66 (1995), 1638
- 151 E.H. Nicollian, A. Goetsberger, Bell Syst. Tech. J. 46 (1967), 1055
- 152 E.H. Nicollian, J.R. Brews: MOS (Metal Oxide Semiconductor) Physics and Technology; John Wiley & Sons (1982) 906
- 153 V.M. Airaksinen, J. Kaitila, H. Niemi, J. Lahtinen, J. Saarilahti, Physica Scripta T54 (1994), 205
- 154 F.W. Smith, G. Ghidini, J. Electrochem. Soc: Solid State Science and Technology, 129 (1982), 1300
- 155 Y. Mii, S. Rishton, Y. Taur, D. Kern, T. Lii, K. Lee, K.A. Jenkins, D. Quinlan, T. Brown Jr, D. Danner, F. Sewell, M. Polcari, IEEE Electr. Dev. lett. 15 (1994), 28
- 156 T. Tuurnala, T. Majamaa, Physica Scripta T69 (1997), 310
- 157 M.P. D'Evelyn, M.M. Nelson, T. Engel, Surf. Sci., 186 (1987), 75

-
- ¹⁵⁸ H. Z. Massoud, *Solid-State Electronics*, 41 (1997), 929
- ¹⁵⁹ V.D. Borman, E.P. Gusev, Y.Y. Lebedinski, V.I. Troyan, *Phys. Rev. B*, 49 (1994), 5415
- ¹⁶⁰ S.A. Nelson, H.D. Hallen, R.A. Buhrman, *J. Appl. Phys.* 63 (1988), 5027
- ¹⁶¹ P. Lundgren: *Electrically Active Defects in Metal - Tunnel Oxide - Silicon (MTOS) Devices* (Department of Solid State Electronics, Chalmers University of Technology, 1994)
- ¹⁶² A. Neubecker, P. Bieringer, W. Hansch, I. Eisele, *J. Crystal Growth* 157 (1995), 201
- ¹⁶³ A. Strass, W. Hansh, A. Neubecker, P. Bieringer, A. Fischer, I. Eisele, *Vacuum* 48 (1997) 701
- ¹⁶⁴ T. Majamaa, V.-M. Airaksinen, S. Novikov, *Physica Scripta*, T69 (1997), 215
- ¹⁶⁵ C. R. Helms, E. H. Poindexter, *Rep. Prog. Phys.* 57 (1994) 791