STUDY OF ELECTRONIC TRANSPORT AND BREAKDOWN
IN THIN INSULATING FILMS

Walter C. Johnson
PRINCETON UNIVERSITY
Department of Electrical Engineering
Princeton, New Jersey 08540
Telephone: (609) 452-4621

1 June 1977

SEMI-ANNUAL TECHNICAL REPORT NO. 3

Approved for public release; distribution unlimited.

Prepared for:

NIGHT VISION LABORATORY
U. S. Army Electronics Command
Fort Belvoir, Virginia 22060

Sponsored by:

DEFENSE ADVANCED RESEARCH PROJECTS AGENCY

DARPA Order No. 2182
Program Code. No. D10
Contract DAAG53-76-C-0059
Effective Date: 17 November 1975
Expiration Date: 17 November 1977

The views and conclusions contained in this document are those of the authors and should not be interpreted as necessarily representing the official policies, either expressed or implied, of the Defense Advanced Research Projects Agency of the U. S. Government.
**Study of Electronic Transport and Breakdown in Thin Insulating Films**

Walter C. Johnson  
Telephone: (609) 452-4621

**Performing Organization Name and Address**  
Princeton University  
Department of Electrical Engineering  
Princeton, New Jersey 08540

**Controlling Office Name and Address**  
Defense Advanced Research Projects Agency  
1400 Wilson Boulevard  
Arlington, Virginia 22209

**Approved for public release; distribution unlimited.**

**Keywords**  
Insulating Films  
Electronic Transport in Insulators  
Charge Trapping in Insulators  
Dielectric Breakdown  
Silicon Dioxide

**Abstract**  
Recent progress is reported in an ongoing program of studies of high-field charge-carrier injection, transport, trapping, and dielectric breakdown in thin insulating films. The investigations reported here include the generation of interface states in the Si-SiO₂ system by ionizing radiation and by high-field stress, the high-field generation of electron traps in SiO₂, a method for measuring interface-state densities at low temperatures, and studies of high-field effects in the Al₂O₃-Si system.
# TABLE OF CONTENTS

<table>
<thead>
<tr>
<th>Section</th>
<th>Page</th>
</tr>
</thead>
<tbody>
<tr>
<td>1. Introduction</td>
<td>1</td>
</tr>
<tr>
<td>2. Radiation-Induced Interface-State Formation in MOS Capacitors</td>
<td></td>
</tr>
<tr>
<td>(J. J. Clement collaborating)</td>
<td></td>
</tr>
<tr>
<td>2.1. Introduction</td>
<td>3</td>
</tr>
<tr>
<td>2.2. Description of Samples and Apparatus</td>
<td>4</td>
</tr>
<tr>
<td>2.3. Measurement of Interface-State Densities</td>
<td>4</td>
</tr>
<tr>
<td>2.4. Influence of Holes and Temperature on Interface-State Formation</td>
<td>6</td>
</tr>
<tr>
<td>2.5. Temperature Dependence of the Interface-State Formation</td>
<td>10</td>
</tr>
<tr>
<td>2.6. Summary</td>
<td>10</td>
</tr>
<tr>
<td>3. A Study of High-Field Effects on MOS Capacitors</td>
<td></td>
</tr>
<tr>
<td>(C. Jenq collaborating)</td>
<td></td>
</tr>
<tr>
<td>3.1. Introduction</td>
<td>13</td>
</tr>
<tr>
<td>3.2. Low-Temperature Ledge (LTL) Technique for Measuring Interface-State Densities</td>
<td>13</td>
</tr>
<tr>
<td>3.3. Generation of Interface States at 90°K During Internal Photoinjection After High-Field Stress</td>
<td>18</td>
</tr>
<tr>
<td>(A). Introduction</td>
<td>18</td>
</tr>
<tr>
<td>(B). Experimental Evidence</td>
<td>18</td>
</tr>
<tr>
<td>(C). Discussion</td>
<td>21</td>
</tr>
<tr>
<td>3.4. Some Properties of Electron Traps Generated in SiO₂ by High-Field Stress</td>
<td>23</td>
</tr>
<tr>
<td>(A). Introduction</td>
<td>23</td>
</tr>
<tr>
<td>(B). Samples and Experiment</td>
<td>24</td>
</tr>
<tr>
<td>(C). Theory for Obtaining Capture Cross Section</td>
<td>24</td>
</tr>
<tr>
<td>(D). Results</td>
<td>26</td>
</tr>
<tr>
<td>(E). Discussion</td>
<td>26</td>
</tr>
<tr>
<td>(a) Comments on the Method of Obtaining Capture Cross Section</td>
<td>26</td>
</tr>
<tr>
<td>(b) Comments on the Results</td>
<td>30</td>
</tr>
<tr>
<td>(F). Summary</td>
<td>31</td>
</tr>
<tr>
<td>3.5. Generation of Interface States After High-Field Stress at 90°K</td>
<td>31</td>
</tr>
<tr>
<td>(A). Introduction</td>
<td>31</td>
</tr>
<tr>
<td>(B). Samples and Experiments</td>
<td>31</td>
</tr>
<tr>
<td>(C). Results</td>
<td>32</td>
</tr>
<tr>
<td>(D). Discussion</td>
<td>36</td>
</tr>
</tbody>
</table>
4. Investigation of High-Field Effects in Al-Al₂O₃-Si Capacitors (O. Bar-Gadda collaborating)
   4.1. Samples ........................................... 41
   4.2. Experimental Procedures ........................... 41
   4.3. Results and Discussion ............................ 41

5. Corona and MIS Studies of High-Field Effect in Al₂O₃
   (S. S. Li collaborating)
   5.1. Introduction ........................................... 50
   5.2. Samples and Experimental Apparatus ................. 50
   5.3. Constant-Current Negative Corona Charging .......... 50
   5.4. MIS Experiments ....................................... 52
   5.5. Capture Cross Section and Concentration of
        Electron Traps ........................................ 54
   5.6. Discussion ............................................ 56

References
A. Publications, Reports, and Doctoral Dissertations Resulting
   from Work Done Under This Program ....................... 59
B. Other References ........................................... 61
1. INTRODUCTION

We report here on recent progress in an ongoing program of research directed toward a basic understanding of the electronic properties of thin insulating films and of the interfaces of such films with semiconductors and metals. Of particular interest are the high-field properties, including charge-carrier injection through the interfaces, electronic transport through the insulator, charge-carrier trapping and recombination at the interfaces and in the insulator, the high-field generation of interface states and trapping centers, and the mechanisms leading to dielectric breakdown. The objective of the program is to provide a rational basis for the choice of materials, processing methods and treatment of the insulating films in order to obtain the desired performance and reliability. The insulating films under study at the present time are silicon dioxide, aluminum oxide, and silicon nitride on silicon substrates. The techniques and apparatus that we have developed under this program are, moreover, immediately applicable to the study of other types of insulating films and substrates.

We have previously reported on many of the results of the studies made under this program,1-29 and additional reports and papers are in preparation. The organization of this report is as follows:

The generation of interface states in the Si-SiO$_2$ system is a matter of great concern. In a previous report$^{25}$ we showed that after high-field stress at liquid nitrogen temperature, the interface states did not make their appearance until the sample had been warmed toward room temperature. In Ch. 2 of this report, J. J. Clement presents recent results obtained on radiation-induced interface states in the Si-SiO$_2$ system. By irradiation of the structure at 87 K, he finds that the nucleation of interface states is apparently associated with the presence of holes at the interface, but he finds further that even if the holes are annihilated by injecting electrons, the interface states appear when the sample is warmed. The interface states in his HCl-steam grown oxides are found to be generated in three temperature ranges: (1) near 140°C, (2) in the range 240-290°C, and (3) at approximately 333°C.

In Ch. 3, C. Jenq presents new results on high-field effects in the Si-SiO$_2$ system. He has previously reported$^{27}$ his observation that electron traps are created in SiO$_2$ at fields $\geq$ 7 MV/cm, and in this report he gives his results on the rate of generation of the traps and on their capture cross sections. The high-field generation of electron traps may
have important consequences in short-channel IGFETs operating with large electric fields in the vicinity of the drain. In addition, Mr. Jenq gives recent results on the high-field generation of interface states, and he shows that electron photoinjection has an influence on interface-state generation. His techniques require the measurement of interface-state densities at liquid nitrogen temperatures, and for this purpose he has developed a low-temperature ledge (LTL) method for this measurement.

In Ch. 4, O. Bar-Gadda presents recent results obtained in a study of high-field effects in Al-Al₂O₃-Si structures. Electron trapping takes place near the negative electrode and positive charging takes place near the positive electrode, thus increasing the electric field in the central region of the insulator. Breakdown is observed to take place near the edge of the field plate, perhaps implying the importance of an edge effect.

In Ch. 5, S. S. Li gives the results of both corona and MIS studies of the Al₂O₃-Si structure. He confirms Mr. Bar-Gadda's observations of electron trapping near the negative electrode and positive charging near the positive electrode, and offers the results of photoinjection experiments to show that the concentration of electron traps is of the order of \(10^{18} \text{ cm}^{-3}\) and that the electron trapping cross section is approximately \(3 \times 10^{-13} \text{ cm}^2\). The observed positive charge is attributed to either hole injection from the positive electrode or field emission from neutral centers in the oxide rather than to impact ionization.
2. RADIATION-INDUCED INTERFACE-STATE FORMATION IN MOS CAPACITORS
   (J. J. Clement collaborating)

2.1. Introduction

Previous investigations into the process of interface-state formation in MOS devices exposed to ionizing radiation have shown that the direct interaction of radiation at the Si-SiO₂ interface is not needed in order to produce interface states; however, interface-state formation is dependent on the transport of radiation-produced holes to the interface. In one study, it was determined that the major part of the interface state generation occurs after the holes have had time to reach the interface.

It has been shown by other workers that holes in thermally grown silicon dioxide are subject to a temperature-activated, field-dependent transport process through the oxide. Thus, at liquid nitrogen temperatures and for low values of applied electric field, the holes are essentially immobile and are trapped through the oxide near the point of creation. However, it has been shown that the motion of holes through the oxide can be enhanced by applying higher electric fields and/or by illuminating the device with light having photon energies in the range from 1.6 to 2.0 eV.

Although the transport of holes through thermally grown silicon dioxide has been studied at low temperatures, past studies of radiation-induced interface-state generation have been carried out almost exclusively at room temperature. In the investigation reported here, the irradiation of the MOS device was done at around 87°C, for several reasons. In addition to the intrinsic significance of how the interface-state density is affected by irradiation at this low temperature, we are better able to control the movement of the radiation-induced holes through the oxide to the Si-SiO₂ interface. Also, this procedure allows us to observe the effect of temperature on the interface-state formation process. As is shown in Sections 2.4 and 2.5 of this report, we find that irradiation of the MOS structure with soft X-rays produces negligible change in the number of interface states at 87°C. However, an increase in interface states is observed when the sample is subsequently warmed. This temperature effect is similar to the one that we reported previously for interface states generated in the Si-SiO₂ system by high fields. We present evidence that only a temporary presence of holes at the interface is required to start the process of
interface-state generation. We observe that interface states are generated in this (HCl-steam) oxide in three temperature ranges: (1) near 140°K, accounting for approximately half of the interface-state generation; (2) in the range 240-290°K; and (3) at approximately 333°K. We suggest that the generation process is electrochemical in nature.

2.2. Description of Samples and Apparatus

The MOS capacitors used in these experiments had n-type (100) silicon substrates with 5-10 ohm-cm resistivity. On this an HCl-steam oxide was thermally grown at 875°C to a nominal thickness of 1000Å. Following a 10-minute anneal at 950°C in nitrogen, semi-transparent aluminum field plates were evaporated. The samples were fabricated at RCA Laboratories through the courtesy of K. M. Schlesier.

The sample was mounted on the cold finger of a vacuum cryostat, making it possible to cool the sample with liquid nitrogen. A resistive heater mounted on the sample stage allowed control of the temperature. The sample chamber was evacuated by an oil diffusion pump. The ionizing radiation used in these experiments was soft X-rays which were generated by 5 kV electrons striking a molybdenum target.

The sample chamber was fitted with a fused silica window for illumination of the sample. Ultra-violet light for the internal photoemission of electrons was provided by a Bausch and Lomb high-intensity monochromator with a Xenon lamp source. Light of lower energy was provided by a Bausch and Lomb tungsten-halide lamp source. Capacitance measurements were made with a Boonton model 72A capacitance meter operating at 1 MHz. Current measurements were made with a Keithley Model 610C electrometer.

2.3. Measurement of Interface-State Densities

Three methods of interface-state measurement were utilized in these experiments. At 333°K the technique utilizing the quasistatic C-V curve as described by Kuhn and Castagne was employed. At 87°K, a measure of the concentration of interface states in the central region of the bandgap was obtained by using the ledge phenomenon found in the low-temperature high-frequency C-V curves. An example of this effect for a typical sample is shown in Fig. 2.1. In this figure, Curve A is the high-frequency C-V curve obtained at 87°K by ramping the bias from accumulation into deep depletion at a fairly fast rate (1 V/sec in this example). The sample is then
Fig. 2.1. Typical low-temperature high-frequency C-V curves for a fresh MOS capacitor used in this study. Curve A is the deep depletion curve obtained by ramping the bias from accumulation into deep depletion. After temporary illumination of the sample, ramping back up to accumulation results in curve B. The width of the ledge indicated by $\Delta V_{AB}$ is used to compute the number of interface states (Eq. 2.1).
temporarily illuminated with visible light, thus supplying minority carriers to fill the interface states. Following this, the bias is ramped toward accumulation, resulting in Curve B. The ledge present in this curve is due to majority carriers filling the interface states in the central region of the bandgap. As is shown in Sec. 3.2 of this report, the section EF of Curve B is essentially parallel to the section CD of Curve A, and the voltage difference, $\Delta V_{AB}$, between the two curves in this region can be used to determine the number of interface states in the central portion of the bandgap according to the formula:

$$N_i = C_{ox} \frac{\Delta V_{AB}}{q} \quad (cm^{-2}) \quad (2.1)$$

where $C_{ox}$ is the oxide capacitance per $cm^2$ and $q$ is the magnitude of the electronic charge. As is also shown in Sec. 3.2 of this report, the central region of the bandgap in which the interface states make the transition at 90°K is approximately 0.7 eV wide in silicon, and so the average concentration of interface states in $cm^{-2}eV^{-1}$ in this central bandgap region is obtained by dividing $N_i$ ($cm^{-2}$) by 0.7 eV.

The Gray-Brown shift of the flatband voltage as the sample is cooled below room temperature provides another method for measuring the number of interface states present in the upper part of the band gap.

The foregoing three methods complement each other and in addition they serve to check each other in their regions of overlap.

2.4. Influence of Holes and Temperature on Interface-State Formation

We have reported previously that if an Si-SiO$_2$ structure is subjected to high field stress at low temperature (91°K), the generation of interface states is observed only after the sample has been warmed toward room temperature. It is well known that if an MOS structure is exposed to ionizing radiation, the buildup of trapped holes is accompanied by an increase in interface-state density. The purpose of these experiments was to examine further the connection between the trapped holes and the interface states. For this purpose we used soft X-rays to produce hole-electron pairs in the oxide at 87°K, following which the electrons were drifted out by application of a moderate electric field, leaving the holes behind. The holes could then be transported to one interface or the other by means of a
larger electric field together with excitation by visible light. The holes were then annihilated by photoinjected electrons, after which the sample was warmed to observe the generation of interface states. The results of these experiments are summarized in Table I, as follows:

<table>
<thead>
<tr>
<th>Column I</th>
<th>identifies the sample. A fresh sample was used each time. All fresh samples were essentially identical.</th>
</tr>
</thead>
<tbody>
<tr>
<td>Column II</td>
<td>shows the number of interface states, ( N_i \text{ cm}^{-2} ), as determined on the fresh sample at 870K by the low-temperature ledge method, using Eq. 2.1.</td>
</tr>
<tr>
<td>Column III</td>
<td>shows the gate voltage used during a 20-minute irradiation with soft X-rays at 870K. This bias will quickly drift the radiation-produced electrons to the positive electrode but will produce very little transport of holes in the oxide at this temperature. Some of the holes produced in the thin hole-trapping layer near the Si-SiO(_2) interface may be trapped there. Also, the holes produced in the bulk of the SiO(_2) will have some initial motion before they are immobilized, and with a positive gate bias this motion will bring some holes into the hole-trapping region. With a negative gate bias, some of the radiation-produced holes in the substrate may drift into the interfacial region. Thus we expect some trapping of holes near the interface for either polarity of gate voltage, but, at these fields, and without light assistance, the main body of holes in the bulk will be essentially immobile at 870K.</td>
</tr>
<tr>
<td>Column IV</td>
<td>shows the gate bias which was applied for three hours at 870K to transport the radiation-produced holes through the bulk of the oxide. The transport of holes during this period was assisted by illuminating the sample with tungsten light which was filtered through a water cell and a yellow filter. After drift of the holes through the bulk of the oxide, essentially all of the holes, both those remaining in the bulk and those trapped at the interface, were annihilated by an internal photoinjection of electrons at 870K.</td>
</tr>
<tr>
<td>SAMPLE</td>
<td>$N_i$ (cm$^{-2}$) INITIAL</td>
</tr>
<tr>
<td>--------</td>
<td>--------------------------</td>
</tr>
<tr>
<td>1</td>
<td>$2.31 \times 10^{11}$</td>
</tr>
<tr>
<td>2</td>
<td>$2.55 \times 10^{11}$</td>
</tr>
<tr>
<td>3</td>
<td>$2.47 \times 10^{11}$</td>
</tr>
<tr>
<td>4</td>
<td>$2.39 \times 10^{11}$</td>
</tr>
</tbody>
</table>

Table I. Radiation-induced generation of interface states. (See text for explanation)
UV light with a photon energy of approximately 5 eV was used for this purpose.

Column V shows the density of interface states at 87°K after the foregoing processes had been completed. Comparison with Col. I shows that the number of interface states remained essentially unchanged throughout the operations of X-irradiation, drift of most of the holes to one interface or the other, and subsequent annihilation of the holes.

Column VI shows the effect of warming the sample and holding it at room temperature for approximately 12 hrs, after which the sample was again cooled to 87°K so that a direct comparison of \( N_1 \) could be made using the same technique as before.

Column VIII shows the increase in interface states that resulted from each of the four sequences of operations.

We can make the following observations from the results shown in Table I. Firstly, from Col. V we see that none of the sequences resulted in an appreciable increase in interface states so long as the sample was held at 87°K. The interface states appeared only during or after warmup. Secondly, the four different sequences show appreciable differences in Col. VII, as follows: On Sample 1, which had no X-irradiation and served as a control sample, the increase in \( N_1 \) after warming was so small as to be negligible. On Sample 2, which was irradiated with positive gate bias but which had no subsequent drift of holes to either interface, the increase in interface states was a moderate one which might have been associated with holes that were produced near the interface and trapped there. On Sample 3, where the holes produced in the bulk were transported to the gate, the increase in interface states was similarly moderate in value. The difference between Samples 2 and 3 is within experimental error and may not be significant. On Sample 4, however, where the holes in the bulk were transported to the interface, the increase in interface states is quite appreciably greater.
Thus the generation of interface states appears to be associated with holes at the interface, and the states do not make their appearance until the sample is warmed. However, since the holes were annihilated before the sample was warmed, merely the temporary presence of holes at the interface appears to be sufficient to start the interface-state generation process.

2.5. Temperature Dependence of the Interface-State Formation

In order to obtain information on the temperature dependence of the interface-state formation process, the following experiment was performed. An MOS capacitor sample was cooled to 87°K and, with the gate biased at +20 V, was exposed to soft X-rays for 30 minutes. The sample, with its temperature held at 87°K, was then biased at +35 V for 3 hours while illuminated with light from a tungsten lamp filtered through a water cell and a yellow filter. As described previously, this procedure transports the holes through the bulk of the oxide to the Si-SiO$_2$ interface. Following this, electrons were photoinjected into the oxide to recombine with the remaining holes. The sample was next warmed to a particular temperature and held there for one hour, then it was cooled to 87°K and the width of the ledge of the low-temperature high-frequency C-V curve was measured for computation of $N_1$ (Eq. 2.1). The sample was heated (and cooled to 87°K) in successive steps to temperatures of 120, 140, 160, 190, 220, 240, 262, 290, 333°K. The total increase in the number of interface states following the last step was $7.1 \times 10^{10} \text{ cm}^{-2}$ as computed from the net change in the width of the ledge. The percentage of this increase which was observed following each step is shown in Fig. 2.2. There appear to be three peaks of interface-state generation, one near 140°K, a broad peak centered at about 270°K, and a third peak at 333°K. It is interesting to note that the peak below 160°K accounts for approximately half of the increase in interface states and that the peaks above 220°K account for the remaining half.

2.6. Summary

1) MOS capacitors exposed to ionizing radiation at 87°K show negligible interface generation for periods up to 3 hrs.

2) Upon warming, interface states are formed. In our HCl-steam grown samples, approximately half of the states are formed near 140°K. The remainder are generated in two temperature ranges: from 240-290°K, and near 333°K.
Fig. 2.2. Generation of interface states vs. temperature after exposure to soft X-rays at 87 K. The sample was (a) heated to 120 K, held for 1 hr, then cooled to 87 K, then cooled to 140 K, held for 1 hr, then cooled to 87 K for measurement of interface-state density and similarly for the successively higher temperatures shown by the circles.
3) The interface-state generation takes place despite the fact that the holes, with which their nucleation appears to be associated, had been annihilated by recombination with photoinjected electrons before the warmup process was initiated.

The requirement of thermal energy for the formation of the interface states suggests that the phenomenon may not be purely electronic but may be electrochemical in nature. Some suggestions in this regard have been made by C. T. Sah.48
3. A STUDY OF HIGH-FIELD EFFECTS ON MOS CAPACITORS
   (C. Jenq collaborating)

3.1. Introduction

In this chapter we present the results of a further study of the
effects of high-field stress on MOS capacitors. Our techniques require
the use of liquid-nitrogen temperatures where the conventional methods of
determining interface-state densities can not be employed. In Sec. 3.2 we
present a method that we have developed for overcoming this difficulty.
In the final sections, 3.3 - 3.5, we describe our recent results on the high-
field generation of interface states and of bulk electron traps in the
Si-SiO₂ system.

3.2. Low-Temperature Ledge (LTL) Technique for Measuring Interface-State

Densities

The usual methods for measuring interface-state densities require
the charge in the interface states to be in equilibrium with the applied
voltage, and this condition can not be achieved within reasonable amounts
of time at lowered temperatures. Brown and Gray⁵⁰ observed a ledge
phenomenon in the C-V curves at liquid-nitrogen temperatures which they
attributed to the presence of interface states, and Goetzberger and Irwin⁵¹
related the width of the ledge to the density of interface states in the
central region of the bandgap. We have analyzed the effect in greater
detail and have made use of the phenomenon in our low-temperature studies
of interface states. The low-temperature ledge (LTL) method is analyzed
and described in the following paragraphs.

Figure 3.1 shows the C-V curves of a MOS capacitor measured at 91₀K
(curves 1 and 2) and an ideal deep-depletion C-V curve calculated for the
temperature of 91₀K (curve 3). Curve 1 was taken by sweeping the bias at
-1.0V/sec from accumulation into deep-depletion in the dark. At Point A
the sample was illuminated with visible light to generate minority carriers.
This caused the capacitance to increase to Point B. The light was cut off,
whereupon the capacitance dropped to Point C. The gate voltage was then
ramped up at 1.0V/sec, producing Curve 2.⁵¹ From now on we shall call
Curve 1 the deep-depletion curve and Curve 2 the light-assisted curve. The
Fig. 3.1. C-V curves at 90°K. Curve 1: Deep-depletion C-V curve. Curve 2: Sweep up after temporary illumination. Curve 3: Ideal deep-depletion C-V curve.
features of the curves will be explained in the following paragraphs.

The emission time constant for an electron captured by an interface state with an energy level $E_s$ is given by \( \tau_{en} = \frac{1}{v_{th} \sigma n_i} \exp(E_i - E_s)/kT \)

where $v_{th}$ is the thermal velocity electrons, $\sigma$ is the electronic capture cross section, $n_i$ is the intrinsic carrier density, and $E_i$ is the intrinsic level of the semiconductor. At $90^0$K, by taking $\sigma$ to be $10^{-15}$ cm$^2$ and $v_{th}$ to be $10^7$ cm/sec, $\tau_{en}$ is found to be 0.1, 1, 10, 100 and 1000 sec when the interface state is at 0.18, 0.19, 0.21, 0.23 and 0.24 eV below the conduction band, respectively. Hence a slight change of 0.01 eV in the energy level of the interface state will result in a change of $\tau_{en}$ of an order of magnitude. During the period when the deep-depletion curve (Curve 1 in Fig. 1) was taken, we may then assume that the electrons which occupied the interface states with an energy level 0.20 eV or more away from the conduction band edge would not be emitted into the conduction, while if the interface state was within 0.20 eV the electrons would have time to be emitted from the states. Note that this approximation will result in an error of not more than 0.02 eV. The possibility that the interface states can capture holes is unlikely due to the negligible generation rate of holes at $90^0$K.

Thus the electrons which are captured by the interface states with energy levels greater than 0.20 eV away from the conduction band will be frozen in the states. If the capacitor is laterally uniform, the part of the C-V curve below this point should be parallel to the ideal deep-depletion curve.\textsuperscript{21} The voltage shift $\Delta V_{AI} = V_A - V_I$ between the lower part of the two curves is due to the oxide charges plus the charges frozen into the interface states. The states which are more than 0.20 eV away from the conduction band edge will remain in their more negative state, and those closer than 0.20 eV will be in their more positive state owing to their emission of electrons. Let $N_{sa}$ be the total number of acceptor-type interface states which are 0.20 eV away from both the conduction and the valence band edges, $N_{sav}$ be those which are within 0.20 eV of the valence band edge, and $N_{sdc}$ be the total number of donor-type interface states that are within 0.20 eV of the conduction band edge. Also, let $Q_{ox}$ denote the
equivalent oxide charge (referred to as the SiO$_2$-Si interface). Then, we have from the above discussion,

$$\Delta V_{AI} = V_A - V_I = q \frac{d_{ox}}{\epsilon_{ox}} \left( \frac{q}{q + N_{sa} + N_{sav} - N_{sdc}} \right)$$

(3.1)

where $d_{ox}$ is the oxide thickness, $\epsilon_{ox}$ is the dielectric permittivity of the oxide, and $q$ is the magnitude of the electronic charge.

In the case of the light-assisted curve before the sample is ramped up (Point C of Fig. 1), the SiO$_2$-Si interface region is flooded with holes generated by the light. The interface states are then all occupied by holes. When the gate voltage is swept toward accumulation, the capacitance remains almost constant at a magnitude higher than the thermal equilibrium value because of the presence of the excess holes at the interface. The constancy of the capacitance indicates that the width of the depletion layer has remained constant. The external charging current during this period is caused by the reduction of the excess holes in the interface region. The reduction process can be either the recombination of the excess holes with the electrons in the conduction band through the assistance of the bulk traps or the interface states, or the diffusion of the holes out of the depletion layer. With a sweep rate of 1 V/sec, the recombination current was calculated to be at least 12 orders of magnitude less than the charging current $\frac{dV}{dt}$. Hence, these excess holes diffuse out of the interface region at a rate determined by the ramping speed. When the number of holes remaining is negligible, the depletion layer then starts to respond to the change of gate voltage and the capacitance begins to increase (Point D) as a consequence of the reduction of the depletion width. As the capacitance increases, the surface potential is reduced and the electron concentration in the conduction band is increased. Eventually at Point E the recombination current from the electrons in the conduction band and the holes in the interface states is large enough to dominate the charging current. The section between Points D and E of Curve 2 will then be parallel to the corresponding section of the ideal deep-depletion curve, provided the sample is uniform. Here the voltage shift $\Delta V_{DI} = V_D - V_I$ between the two sections of curves is due to the oxide charge and the changes in interface state charge. For those states which are more than 0.20 eV above the valence band, the holes previously captured will be frozen in, while the states that are
within 0.20 eV from the valence band edge will be in their more negative state owing to the emission of holes. Let \( N_{\text{sd}} \) denote the number of donor-type states which are more than 0.20 eV away from both the conduction and valence bands, \( N_{\text{sd}} \) denote those within 0.20 eV from the conduction band, and \( N_{\text{sav}} \) denote the acceptor-type states within 0.20 eV from the valence band. Then we have:

\[
\Delta V_{\text{DI}} = V_D - V_I = q \frac{d}{\varepsilon_{\text{ox}}} \left( \frac{Q_{\text{ox}}}{q} - N_{\text{sd}} - N_{\text{sd}} + N_{\text{sav}} \right) \tag{3.2}
\]

From Eqs. (3.1) and (3.2) we have

\[
\Delta V_{\text{AD}} = V_A - V_D = q \frac{d}{\varepsilon_{\text{ox}}} (N_{\text{aa}} + N_{\text{sd}}) \tag{3.3}
\]

or

\[
N_{\text{aa}} + N_{\text{sd}} = \frac{1}{q \frac{d}{\varepsilon_{\text{ox}}}} \Delta V_{\text{AD}} \tag{3.4}
\]

Equation (3.4) gives the interface-state density in cm\(^{-2}\) for those states which are 0.20 eV away from both the conduction and valence bands, i.e., in the mid 0.7 eV of the band gap. Goetzberger and Irwin \(^{51}\) and Gray \(^{46}\) first observed the possibility of using the ledge of the light-assisted curve to measure the interface state density, but the ledge was not unambiguously defined.

For a MOS capacitor biased in the regime where the change in the capacitance is due solely to the change of the depletion capacitance of the substrate, we have the following relationship (for a n-type sample):

\[
\left( \frac{C_{\text{ox}}}{C} \right)^2 = 1 - \left( \frac{2 C_{\text{ox}}^2}{qN_D \varepsilon_s} \right) (V_G - \Delta V) \tag{3.5}
\]

where \( C_{\text{ox}} \) is the oxide capacitance, \( N_D \) is the donor concentration, \( \varepsilon_s \) is the dielectric permittivity of the substrate, \( V_G \) is the gate voltage, and \( \Delta V \) is the voltage shift due to the combination of metal-semiconductor work-function difference, oxide change, and interface-state change.

If \( \left( \frac{C_{\text{ox}}}{C} \right)^2 \) is plotted against \( V_G \) for the lower part of Curve 1 and the section between D and E of Curve 2, then two parallel straight lines with
Fig. 3.2. Plot of $(C_{ox}/C)^2$ vs. gate voltage for the sample of Fig. 3.1. The solid parts of Curves 1 and 2 correspond to Curves 1 and 2 of Fig. 3.1.
the same slope should be obtained as in Fig. 3.2. The displacement between the two lines offers an alternate way of determining $\Delta V_{AD}$ for use in Eq. (3.4).

3.3. Generation of Interface States at 90°K During Internal Photoinjection After High-Field Stress.

3.3(A). Introduction

As was described in the last report, electron traps, not initially present, were found to exist in the oxide layers of MOS capacitors after the samples had been subjected to high-field stress. It was also found that if two samples were subjected to the same high-field treatment at low temperature but received different quantities of photoinjected electrons (so that one sample had more field electron traps than the other), the two still showed essentially the same amount of interface-state generation after they had been warmed up to room temperature. We still find this to be a valid, reproducible result; however, more recently, we have also observed that if the interface-state density is measured by the LTL method before warming the sample, additional interface states, apparently related to the photoinjection of electrons, are indicated by the test. This phenomenon is discussed in the following section.

3.3(B). Experimental Evidence

The samples used in the following discussion were RCA HCl-steam grown $\text{SiO}_2$ films on (100) $\text{Si}$ substrates with $5-10$ $\text{kΩcm}$ resistivity. The gates were semi-transparent aluminum.

Figure 3.3 shows three sets of curves. Set 1 consists of the initial 90°K deep-depletion curve and the light-assisted curve of the fresh sample. Set 2 is a similar set of curves taken after high-field stress, with field plate positive, at 7.4 MV/cm for 30 min at 90°K, and Set 3 are the curves taken after internal photoinjection from the $\text{Si}$ substrate at an interface field of approximately 1.0 MV/cm for 3 hrs. The initial interface-state density in the mid 0.7 eV region of the bandgap is $2.52 \times 10^{11}$ cm$^{-2}$ calculated by the LTL method. The deep-depletion curve in Set 2 is slightly distorted, but the distance between the deep-depletion and the light-assisted curves remains approximately the same, and the interface-state density is
Fig. 3.3. C-V curves taken at 90°K. Set 1: Initial curves. Set 2: After high-field stress at 90°K. Set 3: After internal photoinjection at 90°K.
thus about the same as before. From Set 3, taken after internal photo-
injection, the interface-state density is $3.06 \times 10^{11}$ cm$^{-2}$, approximately
a 20% increase.

Further evidence is provided by the results of the following
experiment. One of the samples was high-field stressed, with field plate
positive, at 7.2 MV/cm for 60 min at 90°K. It was then warmed up and
allowed to sit at room temperature for 17 hours with the gate short
circuited to the substrate. At the end of this period the high frequency C-V
curve was taken, which is Curve 1a in Fig. 3.4. The sample was then cooled
to 90°K and the deep-depletion and light-assisted curves were taken. These
are Curves 1b and 1c of Fig. 3.4. The sample was now subjected to internal
photo-injection, using a photon energy of 5 eV. The injection field was
first set at 1 MV/cm with the substrate negative. After the C-V curves
had shifted to an almost saturated position, the injection field was reduced
to 0.25 MV/cm. When the C-V curves had again shifted to an almost
saturated position, the internal photo-injection was terminated. Curves
2b and 2c are the 90°K C-V curves taken after the internal photo-injection.
The sample was then warmed to 66°C, and the high frequency C-V curves were
taken. This is Curve 2a of Fig. 3.4. The horizontal distance between the
parallel portions of Curves 2b and 2c (width of ledge) is greater than the
corresponding distance between Curves 1b and 1c, indicating again an in-
creased number of interface states after internal photo-injection.

3.3(C). Discussion

When one attempts to measure the electron capture cross section of
the electron traps by monitoring the C-V shift as a function of the total
amount of photoinjected electrons, one finds the accompanying generation of
interface states to be a very undesirable feature. Fortunately, by examining
the relations between the solid and the dashed curves in Fig. 3.4, we are
still able to find the C-V shift that is due only to the oxide charge. In
Fig. 3.4, Curve 2a is almost exactly parallel to Curve 1a. The near
parallelism indicates that the shift $\Delta V_a$ between Curve 1a and Curve 2a is
due mainly to the increase in the oxide charge. If $\Delta V_a$ is compared with
$\Delta V_b$, we find that these two quantities are very nearly the same. This
relationship was found also to hold with other samples which were subjected
to similar treatment. From Eq. (2.1) we know that the shift $\Delta V_b$ is due to
Fig. 3.4. Sample was first high-field stressed at 7.3 MV/cm for 60 min at 90\(^\circ\)K. Curve 1a: After warming to 66\(^\circ\)C. Curves 1b and 1c: Deep-depletion and light-assisted curves at 90\(^\circ\)K. Curves 2b and 2c: After cooling to 90\(^\circ\)K and internal photoinjection of electrons. Curve 2a: After warmup to 66\(^\circ\)C.
an increase in either the equivalent oxide charge \( Q_{ox} \) or the interface states \( (N_{sa}, N_{sav}, \text{ or } N_{adc}) \). Now since \( \Delta V \) is due mostly to the increase in the oxide charge, and \( \Delta V_a = \Delta V_b \), we may conclude that \( \Delta V_b \) is also due mainly to the increase in the oxide charges.

The physical mechanism which is responsible for the aforementioned interface-state generation is still under investigation.

3.4. Some Properties of Electron Traps Generated in SiO\(_2\) by High-Field Stress

3.4(A). Introduction

We have reported previously on our observations that electron traps are generated in silicon dioxide by high-field stress \((E \approx 7 \text{ MV/cm})\). The traps are deep, the optical depth being over 4 eV; thus they are different in character from the 2.4-eV electron traps which were studied by DiMara, Feigl, and Butler\(^{60}\) and which have been tentatively attributed to sodium incorporated into the SiO\(_2\) during growth.\(^{61}\) Furthermore the traps, before the capture of electrons, do not produce an observable positive charging of the oxide, and so they appear also to be different from the positive electron trapping centers, studied by Ning and Yu,\(^{55}\) Ning, Osburn, and Yu,\(^{62}\) and Ning,\(^{63}\) which have been observed in fresh SiO\(_2\) samples and which can be diminished in number by an anneal in forming gas. By the same token, they appear to be different from the radiation-induced positive electron trapping centers studied by Aitken and Young\(^{64}\) and by Aitken, DiMaria, and Young.\(^{65}\)

In a recent paper, Harari\(^{66}\) has reported on high-field generation of electron traps, perhaps similar to ours, in ultrathin (30-300 ˚) films of silicon dioxide, and he suggests that the high-field breakdown of these thin films may be initiated by the generation and charging of these traps. One may speculate on whether the high-field generation of electron traps may be a contributing factor in the electron trapping in IGFET insulators that results in a threshold instability as reported by Ning, Osburn, and Yu.\(^{67,68}\)

Short-channel IGFETS operating with high electric fields should be particularly susceptible to this difficulty. In the present chapter we give some recent results on the measurement of electron capture cross sections of high-field-generated electron traps, and we show some preliminary results on the trap generation rates.
3.4(B). Samples and Experiment

The samples used in these experiments were RCA HCl-steam grown SiO$_2$ films with a thickness of about 1950 Å. The substrates were n-type (100) silicon having 5-10 Ω-cm resistivity. The front contacts of the capacitors were semi-transparent Al gates.

Because of the extremely long minority-carrier response time in the Si substrate for these samples, the quasi-static and high-frequency C-V curves were taken when the sample was warmed up to a temperature of 65-67°C. This allows a faster ramping rate to be used in taking the quasi-static C-V curves. Also, the noise-induced errors were greatly reduced.

The experimental procedure was as follows. The initial quasi-static and high-frequency C-V curves of a fresh sample were first taken at 66°C. The sample was then cooled to 90 K and the deep-depletion and light-assisted curves were taken. High-field stress with field plate positive was then applied to the sample. After this treatment, another set of low-temperature C-V curves were taken. The sample was warmed up to 66°C where a set of C-V curves (both quasi-static and high frequency) were taken. The sample was immediately cooled down to 90 K and the C-V curves were measured. The sample was then warmed up and allowed to set at room temperature until the interface state density increased to its saturation value (see the next section). The sample was then cooled to 90 K again. Electrons were internally photoinjected using 5 eV photons and an oxide field of approximately 1 MV/cm (with substrate negative). The shift of the deep-depletion curve was monitored to obtain a measure of the oxide charge. After the deep-depletion curve had shifted to an almost saturation position, the internal photoinjection was continued with a reduced oxide field of about 0.25 MV/cm until the final saturation of the shift was accomplished. (The reduced field was used at the end because of the possibility that the capture cross section may be reduced at high fields). After the internal photoinjection process, the sample was warmed up. Another set of 66°C and 90 K C-V curves were then taken.

3.4(C). Theory for Obtaining Capture Cross Section

By assuming that only a single species of trap is present in the oxide, and assuming that no detrapping is occurring during the photoinjection procedure, the following equation can be written to describe the trapping process:
\[
\frac{dn_t}{dt} = \sigma v_n n (N_t - n_t) \tag{3.6}
\]

where \(\sigma\) is the capture cross section of the traps, \(v_n\) is the velocity of the free electrons, \(n\) is the density of the free electrons, \(n_t\) is the density of the trapped electrons, and \(N_t\) is the density of traps.

If only a small portion of the injected electrons are trapped, the injection current, \(j\), is related to \(n\) as follows:

\[
j = q v_d n \tag{3.7}
\]

where \(v_d\) is the drift velocity of the electrons. With a field of 1 MV/cm at 900 K, \(v_d\) can be expected to reach a saturation value which is somewhat larger than the thermal velocity of the electrons at 900 K, and, to a good approximation, we can use \(v_d\) in Eq. (3.6). Then, from Eqs. (3.6) and (3.7) we have

\[
\frac{dn_t}{dt} = \sigma j (N_t - n_t)/q \tag{3.8}
\]

The solution to this equation is

\[
n_t(t) = N_t [1 - \exp(-\sigma N_{inj})] \tag{3.9}
\]

where

\[
N_{inj} = q^{-1} \int_0^t j d t
\]

If we assume that the centroid of \(n_t(t)\) remains the same as \(N_t\), then Eq. (3.9) is equivalent to

\[
\Delta V(t) = \Delta V_{sat} [1 - \exp(-\sigma N_{inj})] \tag{3.10}
\]

where \(\Delta V(t)\) is the C-V shift due to the presence of the trapped charges \(n_t(t)\) in the oxide, and \(\Delta V_{sat}\) is the shift when all the traps \(N_t\) are filled.

From Eq. (3.10) we have

\[
\ln \left(1 - \frac{\Delta V(t)}{\Delta V_{sat}}\right) = -\sigma N_{inj} \tag{3.11}
\]
3.4(D). **Results**

Figure 3.5 shows a plot of \( \ln(1 - \frac{\Delta V_{t}(t) / \Delta V_{sat}}{\Delta V_{sat}}) \) against \( N_{inj} \) for three samples. Each had been high-field stressed for 30 min at fields of 7.2, 7.3, and 7.4 MV/cm respectively. The three sets of data agree well with each other and plot reasonably well on a straight line. The capture cross section determined from these data by use of Eq. (3.11) is \( \sigma = 8.4 \times 10^{-14} \text{ cm}^2 \).

In Fig. 3.6 we show a similar plot for three samples which were all high-field stressed at 7.2 MV/cm for periods of 10 min, 20 min, and 60 min respectively. The data here show somewhat greater differences than those of Fig. 3.5 and indicate capture cross sections that range from \( 8.4 \times 10^{-14} \) to \( 9.5 \times 10^{-14} \text{ cm}^2 \), with the larger cross sections consistently corresponding to smaller stress times.

The values of \( \Delta V_{sat} \) obtained at \( E = 7.2 \text{ MV/cm} \) were used to calculate the corresponding values of electron trap density, \( N_{t} \text{ cm}^{-2} \), for various times of stressing. The results, which are plotted in Fig. 3.7, suggest that the generation of traps tends to saturate as the time of stress increases. The effect of the magnitude of the field at constant stress time is being investigated.

3.4(E). **Discussion**

(a) **Comments on the Method of Obtaining Capture Cross Section**

In using Eq. (3.11) we assume that the measured saturation voltage shift of the deep-depletion curve is the true saturation shift which corresponds to complete trap filling. In fact, the measured \( \Delta V_{sat} \) can be expected to be at least slightly less than the true \( \Delta V_{sat} \). The result of this error, as can be analyzed from Eq. (3.11), is to make the \( \ln(1 - \frac{\Delta V_{t}(t) / \Delta V_{sat,measured}}{\Delta V_{sat}}) \) vs \( N_{inj} \) plot curve up relative to the ideal straight line. The discrepancy between the two curves is given in dimensionless form by:

\[
\Delta = \frac{\Delta(\Delta V_{sat})\Delta V(t)}{(\Delta V_{sat})^2 (1 - \frac{\Delta V(t)}{\Delta V_{sat}})}
\]

where \( \Delta V_{sat} \) is the true saturation voltage, \( \Delta(\Delta V_{sat}) \) is the difference between the true and the measured saturation voltages. The discrepancy is smaller when \( \Delta V(t) / \Delta V_{sat} \) is smaller. With a fixed \( \Delta(\Delta V_{sat}) / \Delta V_{sat} \), the initial portion of the measured curve will always give a better reading of
Fig. 3.5. Log plot of \(1 - \frac{\Delta V(t)}{\Delta V_{sat}}\) vs. \(N_{inj}\) for samples subjected to high-field stress for 30 min at fields of 7.2 MV/cm, 7.3 MV/cm, and 7.4 MV/cm respectively.
Fig. 3.6. Log plot of \(1 - \frac{\Delta V(t)}{\Delta V_{\text{sat}}})\) vs. \(N_{\text{inj}}\) for samples subjected to a fixed field stress of 7.2 MV/cm for periods of 10, 20 and 60 min respectively.

\[\sigma = (9.0 \pm 0.6) \times 10^{-14} \text{ cm}^2\]
Fig. 3.7. Number of electron traps generated by high-field stress as a function of stress time. The stress field was 7.2 MV/cm.
the capture cross section.

Another possible reason for nonlinearity of the measured curve may be the existence of two or more species of traps with different capture cross sections. If there are two non-interacting species of traps with capture cross section \( \sigma_1 \) and \( \sigma_2 \), then we have the following equation.

\[
1 - \frac{\Delta V(t)}{\Delta V_{\text{sat}}} = r_1 \exp(-\sigma_1 N_{\text{inj}}) + r_2 \exp(-\sigma_2 N_{\text{inj}})
\]

where \( r_1 = N_{t1}/(N_{t1} + N_{t2}) \), \( r_2 = N_{t2}/N_{t1} + N_{t2} \). Here \( N_{t1} \) and \( N_{t2} \) are the total trap densities of Species 1 and 2, respectively. This equation shows clearly that the effect of the second species of trap is to curve the \( \ln(1 - \Delta V(t)/\Delta V_{\text{sat}}) \) vs \( N_{\text{inj}} \) plot upward. If \( \sigma_1 \) is greater than \( \sigma_2 \) by, say, an order of magnitude, then the effect of the second term will remain small as long as \( N_{\text{inj}} \) is not large. Here we again find that the initial portion of the measured curve provides a better reading for capture cross section. If we are to find the values of both \( \sigma_1 \) and \( \sigma_2 \), we have to be able to increase \( N_{\text{inj}} \) by at least two more orders of magnitude, a procedure which may not be feasible because of limitations on intensity of the UV source.

Finally, instead of Eq. (3.11) there is an equivalent form of Eq. (3.6) which avoids the requirement for determining \( \Delta V_{\text{sat}} \). However, this form requires the injection current to be constant, a requirement which is quite difficult to achieve with the present experimental setup.

(b) Comments on the Results

As was shown in Fig. 3.5, the capture cross section seems to be constant over the range of field strengths used there. However, Fig. 3.6 shows small but appreciable differences in cross section for different stress times. More experimental data are required to determine whether this is due to experimental error or is a true effect.

A value of \( 8.4 \times 10^{-14} \text{ cm}^2 \) for a capture cross section might be taken to indicate that the capturing center is coulomb-attractive and therefore positive in polarity before capture of an electron. No negative C-V shift corresponding to such a positive charge was found however, on a sample after it was high-field stressed at a field slightly below 7.3 MV/cm.
One way to explain the apparent coulombic nature of the trapping center would be to postulate the existence of compensating pairs of charges with opposite signs. If this is true, the negative member of the pair might possibly serve as a hole trap.

3.4(F). Summary

The capture cross section of the electron traps generated by high-field stress at 90°K was found to be between \(8.4 \times 10^{-14} \text{ cm}^2\) and \(9.5 \times 10^{-14} \text{ cm}^2\). The density of the generated traps tends to saturate as the stress time increases. At 7.2 MV/cm, the density of electron traps appeared to saturate at about \(7 \times 10^{10} \text{ cm}^{-2}\) at stress times of the order of 50 min.

3.5. Generation of Interface States After High-Field Stress at 90°K

3.5(A). Introduction

Various investigators have reported the generation of interface states in MOS capacitors after high-field stress at room temperature and after negative-bias stress in the temperature ranges of 25°C - 125°C and 250° - 450°C. We have reported previously that if the high-field stressing was performed at low temperature (91°K), the generation of interface states was not observed immediately, but instead the interface states made their appearance only after the sample was warmed. A further study of this subject is described in the following sections of this report.

3.5(B). Samples and Experiments

Three types of samples were used in this study. All had (100) n-type substrates. Samples RCA-1 and RCA-2 had HCl-steam grown oxides with thicknesses of approximately 2500 Å and 1950 Å, respectively, on silicon substrates having respective resistivities of 1-2 Ω-cm and 5-10 Ω-cm. The third type, Sample BTL, had a 3% HCl dry-grown oxide with a thickness of about 920 Å on a silicon substrate with a nominal resistivity of 5 Ω-cm. The Al gate for RCA-1 was about 1000 Å thick, and the samples designated RCA-2 and BTL had semitransparent Al gates.

The experimental procedures were as follows. A set of room-temperature (for RCA-1) or 66°C (for RCA-2 and BTL) C-V curves (quasi-
static and high frequency) were taken on the fresh sample, and the sample was then cooled to $90 \pm 1^\circ \text{K}$. The low temperature C-V curves (deep-depletion and light-assisted) were taken, after which the sample was high-field stressed at a chosen field for a certain period of time. Following the high-field stress, a set of low-temperature C-V curves were recorded. The sample was then warmed up to room temperature (RCA-1) or $66^\circ \text{C}$ (RCA-2 and BTL), and a set of room-temperature or $66^\circ \text{C}$ C-V curves was taken. The sample was immediately cooled down to $90^\circ \text{K}$ to take the corresponding low-temperature C-V curves. After this process the sample was warmed and allowed to sit at room temperature. A set of room-temperature (or $66^\circ \text{C}$) and low-temperature C-V curves were taken after certain periods of time.

In some cases, internal photoinjection of electrons was performed at $90^\circ \text{K}$ immediately after the high-field stress treatment and before the first warm-up.

3.5(C). \textbf{Results}

Table I shows the interface-state densities (as obtained by the LTL method) of an RCA-1 sample before and after high-field stress at $91^\circ \text{K}$ with a field of $7.6 \text{ MV/cm}$ for 22 min. $N_{ss}$ was found to be unchanged after the high-field stress if the sample was kept at $91^\circ \text{K}$. As the sample was warmed toward room temperature, $N_{ss}$ increased rapidly. When the sample was held at room temperature, $N_{ss}$ increased logarithmically with time. This is plotted in Fig. 3.8.

Two RCA-2 samples were high-field stressed at $90^\circ \text{K}$ with the same field of $7.3 \text{ MV/cm}$ for 30 min. Table II shows the history of $N_{ss}$ for both samples. Sample 4201 was warmed up to room temperature and kept at this temperature for about 33 hrs with the gate short circuited to the substrate. Sample 4201 was warmed up and held at $55^\circ \text{C}$ for 4 hrs with the gate short circuited (interface field nearly zero). Gate voltage was then applied so that the $\text{SiO}_2$-$\text{Si}$ interface field was about $1 \text{ MV/cm}$ for an additional 3 hrs. Again, $N_{ss}$ remained unchanged at $90^\circ \text{K}$ after the high-field stress. $N_{ss}$ increased by approximately 37\% over the initial $N_{ss}$ for both samples when they were first warmed to $66^\circ \text{C}$ for about 20 min (to take the $66^\circ \text{C}$ C-V curves). If the interface field was kept at approximately zero, $N_{ss}$ increased at a rate slower than the case of Table I regardless of whether the sample was at
<table>
<thead>
<tr>
<th>Sample condition</th>
<th>$N_{ss} \text{ (cm}^2\text{)}$</th>
<th>$\Delta N_{ss} \text{ (cm}^2\text{)}$</th>
</tr>
</thead>
<tbody>
<tr>
<td>Initial</td>
<td>$0.27 \times 10^{11}$</td>
<td></td>
</tr>
<tr>
<td>After high field stress</td>
<td></td>
<td></td>
</tr>
<tr>
<td>Before warm up</td>
<td>$0.27 \times 10^{11}$</td>
<td>$1.57 \times 10^{11}$</td>
</tr>
<tr>
<td>1 hr in room T</td>
<td>$1.84 \times 10^{11}$</td>
<td>$0.41 \times 10^{11}$</td>
</tr>
<tr>
<td>11 hrs in room T</td>
<td>$2.25 \times 10^{11}$</td>
<td>$0.09 \times 10^{11}$</td>
</tr>
<tr>
<td>33 hrs in room T</td>
<td>$2.34 \times 10^{11}$</td>
<td>$0.11 \times 10^{11}$</td>
</tr>
<tr>
<td>105 hrs in room T</td>
<td>$2.45 \times 10^{11}$</td>
<td></td>
</tr>
</tbody>
</table>

Table I. $N_{ss}$ obtained by the LTL method for stress and after high-field stress at $90^\circ K$ with 7.6 MV/cm for 22 min followed by warm-up to room temperature. Sample RCA-1.
Fig. 3.8. Increase of interface states as a function of time after the sample was high-field stressed, warmed, and held at room temperature.
<table>
<thead>
<tr>
<th>Sample No. 4200</th>
<th></th>
<th>Sample No. 4201</th>
<th></th>
</tr>
</thead>
<tbody>
<tr>
<td>Initial</td>
<td>$N_{ss}(\text{cm}^2) = 2.42 \times 10^{11}$</td>
<td>Initial</td>
<td>$N_{ss}(\text{cm}^2) = 2.32 \times 10^{11}$</td>
</tr>
<tr>
<td>20 min at 66°C</td>
<td>$N_{ss}(\text{cm}^2) = 3.33 \times 10^{11}$</td>
<td>20 min at 66°C</td>
<td>$N_{ss}(\text{cm}^2) = 3.18 \times 10^{11}$</td>
</tr>
<tr>
<td>33 hr at 75°C</td>
<td>$N_{ss}(\text{cm}^2) = 3.42 \times 10^{11}$</td>
<td>4 hr at 55°C</td>
<td>$N_{ss}(\text{cm}^2) = 3.23 \times 10^{11}$</td>
</tr>
<tr>
<td>$E_{\text{si}, 3} = 0 %$</td>
<td>$\Delta N_{ss}(\text{cm}^2) = 0.91 \times 10^{11}$</td>
<td>$E_{\text{si}, 3} = 0 %$</td>
<td>$\Delta N_{ss}(\text{cm}^2) = 0.05 \times 10^{11}$</td>
</tr>
<tr>
<td>$E_{\text{si}, 3} = 1 %$</td>
<td>$\Delta N_{ss}(\text{cm}^2) = 0.09 \times 10^{11}$</td>
<td>$E_{\text{si}, 3} = 1 %$</td>
<td>$\Delta N_{ss}(\text{cm}^2) = 0.22 \times 10^{11}$</td>
</tr>
</tbody>
</table>

Table II. $N_{ss}$ obtained by the LTIL method for two RCA-2 samples subjected to the same high field stress (6.3 MV/cm for 30 min) but with different warm-up conditions.
25°C or 55°C. This difference between the generation rates may be due to the difference between the stress fields (7.6 MV/cm for Table I, 7.3 MV/cm for Table II). If an interface field of 1 MV/cm is applied (Sample 4201), the increase of $N_{ss}$ is accelerated.

Table III shows the results obtained on a BTL dry-oxide sample which was high-field stressed at 900K with a field of 7.4 MV/cm for 30 min. Observation of the interface states by the LTL method showed no generation of interface states at 900K. The generation of states after warm-up is shown in Table III. The distribution of states within the silicon bandgap, $n_{ss}$ in cm$^{-2}$ eV$^{-1}$, is shown in Fig. 3.9 as calculated by Kuhn's method before stress, after warm-up to 65°C for 20 min, and after 14 hrs at 25°C. A prominent feature in Fig. 3.9 is the peak in $n_{ss}$ after the sample was first warmed up (Curve 2). This peak gradually reduced as the sample was held at room temperature. A similar peak in $n_{ss}$ was observed by Roitman in the interface states generated in the Si-SiO$_2$ system by electron-beam irradiation.

Figure 3.10 shows $n_{ss}$ (cm$^{-2}$ eV$^{-1}$) for the other BTL Sample. This sample was stressed at the same field and for the same time as the sample of Fig. 3.9, but internal photoinjection of electrons was performed before the same was warmed. This resulted in an appreciably reduced interface-state after warming. Also, the peak found in Fig. 3.9 is greatly reduced.

3.5(D). Discussion

The presence of holes at the SiO$_2$-Si interface has been suspected to be a source of interface states. We have found in this study, however, that even when the stress field is not high enough to cause impact ionization, so that after high-field stress only a net negative charge was detected in the oxide, interface states were still generated after the sample had been warmed up (Table II).

The general pattern for the generation of interface states after high-field stress at 900K can be summarized as follows. First, no generation can be observed for several hours at 900K after high-field stress. Secondly, the generation rate increases as the temperature is raised. For example, the generation of interface states was observed when one sample was warmed up to 168°C and then brought back immediately to 900K (see also J. J. Clement's results in Ch. 2 of this report). Thirdly, at room temperature the interface-
<table>
<thead>
<tr>
<th>Sample condition</th>
<th>$N_{ss}$ (cm$^2$)</th>
<th>$\Delta N_{ss}$ (cm$^2$)</th>
</tr>
</thead>
<tbody>
<tr>
<td>Initial</td>
<td>$0.51 \times 10^{11}$</td>
<td>0</td>
</tr>
<tr>
<td>After high field stress:</td>
<td></td>
<td></td>
</tr>
<tr>
<td>Before warm up</td>
<td>$0.51 \times 10^{11}$</td>
<td>$2.11 \times 10^{11}$</td>
</tr>
<tr>
<td>20 min at 65°C</td>
<td>$2.62 \times 10^{11}$</td>
<td>$0.09 \times 10^{11}$</td>
</tr>
<tr>
<td>14 hr at 25°C</td>
<td>$2.71 \times 10^{11}$</td>
<td></td>
</tr>
</tbody>
</table>

Table III. $N_{ss}$ obtained by the LTL method for a BTL sample which was high-field stressed at 7.4 MV/cm for 30 min and warmed.
Fig. 3.9. Plots of $n_{ss}$ (cm$^{-2}$ eV$^{-1}$) as a function of the energy below the Si conduction band edge for a BTL sample before the high-field stress (Curve 1), after stress at 900K with 7.4 MV/cm for 30 min followed by warming to 66°C (Curve 2), and after the sample had been at room temperature for 14 hours (Curve 3).
Fig. 3.10. Plots of $n_{ss}$ (cm$^{-2}$ eV$^{-1}$) as a function of the energy below the Si conduction-band edge for a BTL sample before it was field-stressed (Curve 1), after high-field stress followed by internal photoinjection, and warming to 66°C (Curve 2).
state density continues to increase slowly and follows approximately a linear function of \( \ln t \), as shown in Fig. 3.8. The generation rate can be accelerated by the assistance of a moderate field, as in Sample 4201 of Table II.

The peak of \( n_{ss} \) in Fig. 3.9 suggests the possibility of a bond-breaking process during the high-field stress. The height of the peak is reduced while the width is increased as time goes on. This may indicate that a process of atomic rearrangement is taking place at the \( \text{SiO}_2-\text{Si} \) interface. Thermal energy may be required for the rearrangement.

From a comparison of Figs. 3.9 and 3.10 it would appear that the injection of electrons through the \( \text{SiO}_2-\text{Si} \) interface can anneal out a considerable portion of the potential defects which, when supplied with sufficient thermal energy, give rise to interface states. This further recommends the possibility of the aforementioned bond-breaking process. When electrons are injected through the interface, some of the broken bonds may become satisfied by capture of free electrons. Further study of the interface-generation phenomenon is indicated.
4. INVESTIGATION OF HIGH-FIELD EFFECTS IN Al-Al₂O₃-Si CAPACITORS
(O. Bar-Gadda collaborating)

We report here on the continuation of experiments relating to
charge storage and dielectric breakdown in Al₂O₃ on silicon substrates.

4.1. Samples

The samples used in this study were fabricated at Bell Laboratories,
and we thank G. E. Smith, D. M. Boulin, and E. Labate for their aid. The
substrates of the samples were (100) 4-6 ohm-cm silicon. Both n (P-doped)
and p (B-doped) substrates were included. CVD Al₂O₃ films were deposited
to thicknesses of 450-480 Å, using two deposition temperatures: 815°C
and 900°C. The samples had Al field plates approximately 1000 Å in
thickness.

As received, the MIS capacitors showed initial flatband voltages
in the range 1.0 ± 0.3 V, corresponding to an average fixed oxide charge of
\( \sim 2 \times 10^9 \) cm⁻².

4.2. Experimental Procedures

The following experiments were performed on these samples:

(1) The flatband voltage, \( V_{FB} \), vs. time \( t \) was measured for different
    applied fields, \( E_a \) for both field polarities and substrate types.

(2) The time to breakdown, \( t_B \), vs. applied field, \( E_a \), was measured
    for substrate biased in accumulation.

(3) C-V curves were recorded before and after breakdown.

(4) Oscillograms of the breakdowns were recorded.

(5) Samples were observed under an optical microscope after breakdown.

All experiments were carried out at room temperature. The voltages were
applied by either of two pulse generators. The first, used in the pulse-
width range \( \Delta t > 10 \) ms had a rise time of \( \sim 10 \) μs. The second, used in the
range \( \Delta t < 10 \) ms, had a rise time of \( \sim 30 \) ns.

4.3. Results and Discussion

Figure 4.1 shows the results of Procedure (1), as described above,
for samples biased in accumulation. In general, for the p-type samples,
\( \Delta V_{FB} \) tended to saturate at \( \sim 4 \) V, then decreased, the decrease being accompanied
by a large stretchout of the C-V curve. For the n-type samples, however, no saturation is apparent. The maximum flatband shift for positive bias on n-type is \( \sim 7 \) V.

For negative bias on n-type and positive bias on p-type, an incandescent light was used to illuminate the samples so as to maintain the inversion layer. Deep depletion would be an undesirable condition for studying the oxide, for two reasons: (a) Part of the applied field would appear across the depletion layer, and (b) impact ionization of carriers in the depletion layer would create hot carriers.

The results for illuminated samples biased in inversion were very similar to those shown in Fig. A1. Qualitatively, we can say that (i) the charging is faster in the n-type sample than in the p-type sample for a given field and polarity, (ii) the maximum \( \Delta V_{FB} \) for positive bias is greater for p-type than for n-type, (iii) saturation appeared to occur in all cases except for positive biased n-type at the higher bias voltages. In the case of negative bias, saturation was accompanied by large C-V stretchout followed by a decrease in \( \Delta V_{FB} \).

The injection mechanism cannot be inferred from the \( \Delta V_{FB} \) vs. \( t \) curve because nearly all types of injection will produce similar curves. For \( \text{Al}_2\text{O}_3 \), it is believed that direct tunneling of electrons to traps plays an important role because of the relatively low field (1-2 MV/cm) required for measurable injection current and charge trapping and because the measured density of electron traps (\( \sim 10^{18} - 10^{19} \) cm\(^{-3}\)) is high. We did not study the injection mechanism. It is unlikely that ionic current plays a role here because the flatband shift is positive for all polarities of applied field.

A possible model for the charging process is illustrated in Fig. 4.2. The field initially injects electrons from the Al into the oxide (Sketch a) where some of them are trapped (Sketch b). This reduces the injection field, decreasing the external current. The field near the positive contact increases, and this leads to a reduction in negative charge in the oxide (Sketch c) by a mechanism which is unclear at present. Electron emission from neutral centers or hole trapping near the Si interface are two possibilities. The sequence of events with positive gate bias is probably similar to the above, but the reduction in negative charge would
Fig. 4.1. \( \Delta V_t \) vs. log \( t \) for various gate voltages (as labeled).

Change in平板电压, $\Delta V$ vs. $t$.
not be observable because it would take place near the field plate and consequently would not result in an appreciable C-V shift.

Figure 4.3 is a plot of time-to-breakdown vs. applied field for the case of accumulation. The results indicate a threshold field for breakdown of $\sim 4.9 \text{ MV/cm}$ for p-type, and $\sim +4 \text{ MV/cm}$ for n-type. These fields are lower than the "intrinsic" field strengths measured for $\text{Al}_2\text{O}_3 (> 6 \text{ MV/cm})$, which suggests that the breakdowns are due to weak spots or, alternatively, that the maximum field in the oxide is significantly larger than the average field.

Figure 4.4 shows typical C-V curves before and after breakdown. For all cases except positive bias on n-type, a complete C-V curve from accumulation to inversion is traceable. For n-type, a nearly total breakdown is observed in which the oxide has been completely discharged. For the case of negative bias, there is not much change in the oxide charge or the C-V curve, and indeed the only electrical means of noting the breakdown is the orders-of-magnitude sudden increase in current. For the case of positive bias on illuminated p-type, two features are observed: (1) a small positive $\Delta V_{FB}$ and (2) an apparent increase in the capacitance in the weak accumulation regime.

We illustrate in Fig. 4.5(a) a typical voltage waveform taken of a breakdown on positively biased n-type. The breakdown discharge takes place with a time constant on the order of $1 \mu$s and is seen to be followed by a series of propagating breakdowns. Observation of the sample under an optical microscope reveals the presence of the breakdown areas. Features very similar to those reported for self-healing or self-quenched breakdowns are noted, such as the presence of a circular hole where breakdown is presumed to have evaporated the metal field plate. In addition we note that the breakdown spot displays an irregular star-shaped pattern with rays extending from it, possibly being filamental conduction paths or cracks on the field plate. A final important feature should be noted. In all cases of breakdown observed, the breakdown spots occurred very close to the edge of the field plate. This suggests that the field at the edges may be considerably higher than that in the rest of the oxide, and quite conceivably close to the intrinsic field strength. An edge effect would seem to be an important
(a) Initial band configuration on application of bias.

(b) After charging by electron trapping near Al and into the bulk.

(c) After reduction of flatband shift, possibly caused by electron emission from neutral centers in bulk, or hole trapping near the Si interface.

Fig. 4.2. Band diagram for negative bias
Fig. 4.3. Time to breakdown (s) vs. applied bias (volts on left, MV/cm on right).
(Polarity: - for p-type; + for n-type)
Fig. 49: Typical C-V curves obtained before bias, after bias but before breakdown, and after breakdown.
factor in the breakdown process. Finally, we note that high-field breakdown at the edges seems to imply the absence of pinholes or other severe weak spots in the film.
Fig. 4.5. Typical oscillogram and micrograph of breakdown
5. **CORONA AND MIS STUDIES OF HIGH-FIELD EFFECTS IN Al$_2$O$_3$**
(S. S. Li collaborating)

5.1. **Introduction**
Several investigators have observed that holes are generated and captured within the Al$_2$O$_3$ when the sample is exposed to X-rays or VUV radiation. Positive charging is also observed in Al$_2$O$_3$ after exposure to negative corona at high fields. In this chapter we present experimental evidence which indicates that the positive charging observed in Al$_2$O$_3$ is due to the trapping of holes.

We have also investigated electron trapping in Al$_2$O$_3$. For this purpose we have utilized an internal photoemission of electrons into the Al$_2$O$_3$ under a low positive field. If we assume that the electron traps are uniformly distributed through the bulk of the oxide and that all have the same cross section, our results indicate a trap concentration of $1.9 \times 10^{18}$ cm$^{-3}$ and an electron trapping cross section of $3 \times 10^{-13}$ cm$^2$.

5.2. **Samples and Experimental Apparatus**

The samples, which were fabricated at Bell Laboratories, had a 900°C pyrolytic Al$_2$O$_3$ film with a thickness of about 450Å. The substrate was n-type (100) silicon with 8-12 Ω-cm resistivity. The front Al gate of the MIS structure used in the photoinjection experiment was semi-transparent.

The set-up for the corona-charging experiment was described in Reference 23. The photoinjection experiments were performed with the sample mounted in a chamber which could be evacuated to a pressure of approximately 1 micron. The sample was exposed to UV light through a silica window. A Keithley 610B electrometer was used to monitor the external flow of charge. Charge storage in the insulator was determined from the flatband voltage of the MIS structure as measured at 1 MHz.

5.3. **Constant-Current Negative Corona Charging**

Negative corona was applied to charge the bare surface of the Al$_2$O$_3$ sample. The surface potential and flatband voltage as functions of the charging time are shown in Fig. 5.1 for three different current levels. Initially, the surface potential buildup and the flatband voltage decay are very fast. Negative flatband shift corresponds to the loss of
Negative Corona Current

1. \( J = 9 \times 10^{-8} \text{ Amp/cm}^2 \)
2. \( J = 3.75 \times 10^{-7} \text{ Amp/cm}^2 \)
3. \( J = 9.25 \times 10^{-7} \text{ Amp/cm}^2 \)

Fig. 5.1. The build-up of the surface potential and the flatband voltage decay as functions of the charging time for three different negative-corona current levels.
negative charge or the gain of positive charge. With high charging
current and long charging times, the flatband voltage falls lower than
-1 volt, indicating that a net positive charge exists within the oxide.

If the charged sample is exposed to low-level negative corona
for a time, the C-V curve will shift to the right side. This indicates
that part of the positive charge has been annealed out by the electrons
injected from the front surface.

When a fresh sample is thermally annealed in vacuum at 500°C
for one hour, no flatband shift is observed. This indicates that the
negative charges within the fresh sample may be due either to deeply
trapped electrons or to fixed negative charges. The positive charges
found after negative corona treatment can be reduced appreciably if
the surface charge is neutralized and the sample is annealed in vacuum
at 300°C for 50 minutes. A 3 V positive flatband shift is observed with
this treatment.

5.4. MIS Experiments

As is shown in Fig. 5.2, biasing the MIS capacitor to an
average field of -4.7 MV/cm for 10 minutes causes a positive flatband
shift and a large hysteresis that was not present in the C-V curves of
the fresh sample. As the sample biasing time is increased, the C-V
curves begin to shift to the left, indicating a loss of negative charge
or a gain of positive charge. This agrees with an observation made by
Tsujido.\(^7\) The hysteresis can be attributed to the injection of electrons
which neutralizes holes captured close to the interface. It appears that
positive charge trapping exists only after negative biasing. There are
several possibilities:\(^2\)

(a) Holes may be injected from the substrate.

(b) Electrons may tunnel out of neutral centers and leave positive
charge behind.

(c) Injected electrons from the front surface might obtain enough
energy to impact-ionize the lattice or neutral traps. The generated holes
may then be captured.
Fig. 5.2. C-V curves for the Al-Al₂O₃-Si capacitor. (1) Fresh sample. (2) After 10 min, -21 V. (3) After total of 30 min, -21 V. (4) After total of 60 min, -21 V. Also shown is a sketch diagram illustrating the charge storage.
Impact ionization will depend very much on the mean free path of the charge carriers in the oxide. The optical phonon scattering length is about 4–5 Å, which is one-sixth of that in SiO₂ and the optical phonon energy is about 0.1 eV. Taking this into account, we can see that the number of hot electrons will be quite limited. Also, it is probable that generated holes will suffer geminate recombination even in high fields. Therefore, impact ionization appears to be an unlikely source of holes.

From the above argument, we propose the following charge-trapping model for the Al-Al₂O₃-Si structure under negative high-field stress. Electrons can tunnel from the Al gate and be captured in the bulk of the Al₂O₃. Holes are injected from the substrate and are trapped. The centroid of the trapped electrons is close to the front surface, but the centroid of the holes is near the Al₂O₃-Si interface (Fig. 5.2). The electron and hole trapping reduces the field at both interfaces, thus enhancing the field in the bulk. This explains why the current decays with time, and will also be a factor in the breakdown of the oxide.

5.5. Capture Cross Section and Concentration of Electron Traps

The Al-Al₂O₃-Si sample was held at a bias of 1.2 MV/cm until the dark current fell below 10⁻¹¹ A/cm². Electrons were then photoinjected from the substrate into the Al₂O₃ by 5 eV UV light. The external current and the flatband shift were measured after each 30-sec interval of exposure.

It has been shown by Harari and Royce and by N. Johnson, using the results of photodepopulation experiments, that electron traps are distributed almost uniformly through the bulk of the oxide. Therefore in our analysis we assume that the traps are uniformly distributed and that all the traps have the same electron trapping cross section. Under the assumption that photodetrapping and field detrapping are small enough to be neglected, the equations which determine the trapped electron concentration, \( n_T(x,t) \), are

\[
\frac{\partial n_T(x,t)}{\partial t} = - \frac{3\lambda(x,t)}{\partial x}
\]

and
Fig. 5.3. Experimental and theoretical curves of $C_{ox} \Delta V_{FB} / q$ vs $N_{inj}$. All three solid curves are drawn for the same value of $N_p = 5.68 \times 10^{15}$ cm$^{-2}$. 

$\bar{x} = 180.5^\circ$ (See Fig. 5.4)
where \( \sigma \) is the electron capture cross section. The solution is

\[
\frac{3n_T(x,t)}{q} \frac{\partial}{\partial t} = \sigma j(x,t) [N_T - n_T(x,t)]
\]  

(5.2)

where \( n_T(x,t) \) is the number of injected electrons per unit area and is given by

\[
n_T(x,t) = \frac{N_n \left[ \exp(\sigma N_{inj}) - 1 \right]/\left[ \exp(\sigma N_{inj}) - 1 + \exp(\sigma N_T x) \right]}{\exp(a N_T x)}
\]  

(5.3)

where \( N_{inj} \) is the number of injected electrons per unit area and is given by

\[
N_{inj} = C_{ox} \frac{\Delta V_{FB}}{q} + \frac{Q_{ext}}{q}
\]  

(5.4)

Here \( Q_{ext} \) is the amount of charge that flows through the external circuit during the photoinjection process.

The effective trapping efficiency is defined by

\[
\eta_{eff} (N_{inj}) = \frac{d(C_{ox} \Delta V_{FB}/q)}{d N_{inj}}
\]  

(5.5)

Thus,

\[
\eta_{eff}(0) = 1 - \frac{1 - \exp(-\sigma N_T t_{ox})}{\sigma N_T t_{ox}}
\]  

(5.6)

From Eq. (5.5) we see that the initial slope of \( C_{ox} \Delta V_{FB}/q \) vs \( N_{inj} \) will give \( \eta_{eff}(0) \), which can be used to determine \( N_T \). In Fig. 5.3 we show the experimental data. These fit the theoretical calculation very well with the values \( N_T = 1.9 \times 10^{18} \text{ cm}^{-3} \) and \( \sigma = 3 \times 10^{-13} \text{ cm}^{2} \). Our measured trap density is consistent with the number of negative trapped charges, \( 10^{17} - 10^{18} \text{ cm}^{-3} \), measured by Mehta, Butler, and Feigl in their samples of \( \text{Al}_2\text{O}_3 \).

The centroid of the electron trapping vs \( N_{inj} \), as calculated from \( n_T \) and \( \Delta V_{FB} \), is plotted in Fig. 5.4. The centroid advances from \( 139 \) \( \AA \) to \( 181 \) \( \AA \) gradually until the photoinjection is stopped. The initial electron trapping efficiency is computed to be 92.2%.

5.6. Discussion

Trapped holes are observed after the \( \text{Al}_2\text{O}_3 \) has been under negative high-field stress. The small electron-optical phonon scattering length
Fig. 5.4. Calculated position of centroid vs $N_{\text{inj}}$.

\[
\sigma = 3 \times 10^{-13} \text{ cm}^2
\]

\[
N_T = 1.89 \times 10^{19} \text{/cm}^3
\]
leads us to believe that impact ionization is not the principal source of the trapped holes. The tunneling of holes from the substrate or an out-tunneling of electrons from neutral centers seem to be likely mechanisms. Electron trapping near the negative electrode and hole trapping near the positive electrode can account for the observed decay in current as time goes on.

Under the assumption of uniformly distributed traps and uniform trapping cross sections, we find the electron trap concentration to be $1.9 \times 10^{18}$ cm$^3$ in our samples. The electron trapping cross section is found to be $3 \times 10^{-13}$ cm$^2$. The magnitude of this cross section makes it appear likely that the traps are coulomb-attractive in nature.

We plan to study the electron injection mechanisms and expect also to investigate the spatial and energy distributions of the captured electrons.
REFERENCES

A. PUBLICATIONS, REPORTS, AND DOCTORAL DISSERTATIONS RESULTING FROM WORK DONE UNDER THIS PROGRAM


5. Z. A. Weinberg, W. C. Johnson, and M. A. Lampert, "Determination of the Sign of Carrier Transported Across SiO₂ Films on Silicon."
   (b) Special Report No. 2 (AFCRL-TR-74-0206), April 1974.

   (b) Special Report No. 3 (AFCRL-TR-74-0278), June 1974.


29. C. C. Chang and Walter C. Johnson, "Frequency and Temperature Tests for Lateral Nonuniformities in MIS Capacitors."
   (b) Accepted for publication by IEEE Trans. Electron Devices.

B. OTHER REFERENCES


76. Noble M. Johnson, private communication.