High density plasma deposition of device quality silicon nitride. II. Effects of thickness on the electrical properties
Marie-Christine Hugon, Franck Delmotte, Bernard Agius, Eugene A. Irene

To cite this version:
Marie-Christine Hugon, Franck Delmotte, Bernard Agius, Eugene A. Irene. High density plasma deposition of device quality silicon nitride. II. Effects of thickness on the electrical properties. Journal of Vacuum Science and Technology B, 1999, 17 (4), pp.1430-1434. <10.1116/1.591100>. <hal-00903605>

HAL Id: hal-00903605
https://hal-iogs.archives-ouvertes.fr/hal-00903605
Submitted on 12 Nov 2013

HAL is a multi-disciplinary open access archive for the deposit and dissemination of scientific research documents, whether they are published or not. The documents may come from teaching and research institutions in France or abroad, or from public or private research centers.

L’archive ouverte pluridisciplinaire HAL, est destinée au dépôt et à la diffusion de documents scientifiques de niveau recherche, publiés ou non, émanant des établissements d’enseignement et de recherche français ou étrangers, des laboratoires publics ou privés.
High density plasma deposition of device quality silicon nitride. II. Effects of thickness on the electrical properties

M. C. Hugon, F. Delmotte,* and B. Agius
Laboratoire Charles Fabry de l’Institut d’Optique (URA D0014), 91403 Orsay, France
E. A. Irene
Department of Chemistry, University of North Carolina, Chapel Hill, North Carolina 27599-3290

(Received 28 October 1998; accepted 9 April 1999)

Dielectric behavior of SiN$_x$ films, fabricated by microwave electron cyclotron resonance discharge, has been studied as a function of film thickness on the basis of the current–voltage and the capacitance–voltage characteristics. In the thickness range (20 nm < d < 80 nm), the resistivity and the critical field for SiN$_x$ were found not to be sensitive to the film thickness (d) and which was opposite to strong dependence of the dynamic dielectric constant $\varepsilon_d$ on thickness. To explain the $\varepsilon_d$ behavior as a function of d, a model based on trapped space charge effects is proposed. The dominant mode of electronic conduction, determined from $I-E^{1/2}$ curves and Arrhenius plots of leakage current, appears to be Poole–Frenkel emission only for thicker films (d > 20 nm). Finally, the spatial profile of fixed charges reveals that SiN$_x$/Si interface has a much greater concentration of defects than the bulk film. © 1999 American Vacuum Society. [S0734-211X(99)00704-0]

I. INTRODUCTION

SiN$_x$ thin films have important applications in microelectronics, optoelectronics, optics, and hard surface coatings. Presently, in the ultralarge scale integration (ULSI) circuit applications, as device dimension shrink into the submicron range, a proportional decrease in the gate dielectric thickness is indispensable.\textsuperscript{1} For the development of InP based microelectronics, the formation of a high quality dielectric insulator is key to technology. As a consequence, low temperature deposition is required to prevent thermal decomposition of the InP surface.\textsuperscript{2}–\textsuperscript{5} These different requirements have stimulated considerable research on the plasma deposition of dielectric thin films in particular the use of high density plasmas (HDPs).\textsuperscript{6}

With film thicknesses below 20 nm now commonplace, control of the film physical and electrical properties becomes very critical. Only a few papers dealing with the electrical characteristics of SiN$_x$/Si structures have been published,\textsuperscript{4,7} even though the behavior of SiN$_x$ devices strongly depends on the film’s electrical properties, especially when the film thickness is less than 20 nm. For example, Poole–Frenkel emission is known to be the dominant conduction mechanism for silicon nitride thin films deposited at high temperatures (800–1000 °C) by low pressure chemical vapor deposition (LPCVD)\textsuperscript{8} or by radio frequency plasma enhanced CVD (PECVD).\textsuperscript{9,10} However, only a few papers\textsuperscript{11} have described the electrical properties of SiN$_x$ films deposited by electron cyclotron resonance (ECR)-PECVD, and it is not particularly obvious whether the conduction mechanism will be the same when the thickness varies from nanometers to hundreds of nanometers. Therefore, understanding of the conduction mechanism for such thin films is necessary.

In this article, we report the study of the electrical properties of SiN$_x$ films deposited using the distributed ECR (DECR) plasma method at floating temperature, on silicon substrates, as a function of the film thickness in the 7–80 nm range. This DECR plasma process has proved to be one of the most suitable “soft” deposition techniques to obtain device quality insulator thin films with good electrical properties at very low substrate temperatures (below 150 °C). In view of the increasing interest in SiN$_x$ interlayers in gate dielectric structures, the aim of this article is to study the electrical properties of SiN$_x$ films by means of current–voltage ($I$–$V$), current–temperature ($I$–$T$), and capacitance–voltage ($C$–$V$) measurements. We focus essentially on the positive fixed charge distribution and on the determination of dominant conduction mechanisms.

II. EXPERIMENT

The DECR equipment used in this study was described previously.\textsuperscript{12,13} Briefly, it consists of a 400 mm diam chamber equipped with a loadlock; the base pressure in both chambers is below 5 × 10$^{-5}$ Pa. The microwave power (2.45 GHz) is coupled to the plasma through 14 antennas evenly distributed at the periphery of the chamber and permanent magnets (875 G) are placed near each antenna outside the chamber.

SiN$_x$ films were deposited on (100) $n$-type (5–6 × 10$^{15}$ cm$^{-3}$) silicon substrates. Before loading into the chamber, the substrates were subjected to conventional RCA cleaning.\textsuperscript{14} This cleaning procedure ended with a 1 min rinse in diluted HF, a 10 min rinse in de-ionized (DI) water and being blown dry with nitrogen.

The deposition process parameters were the same throughout this study: the pressure and microwave power were 0.3 Pa and 1500 W, respectively, and the N$_2$/SiH$_4$ flow ratio was fixed to 19 (19 sccm N$_2$ and 1 sccm SiH$_4$). From our previous results, these process parameters allow us to optimize the physical and electrical properties of SiN$_x$ thin
films.5,13,15 These films display a homogeneous composition over a large thickness range (7–100 nm). However, the film stoichiometry near the SiNₓ/Si interface (N/Si = 1.52 ± 0.05) is higher than in the film bulk. From a previous spectroscopic ellipsometry study, an interface layer with a lower index than that in the bulk film is necessary for modeling the SiNₓ/Si structure.16

For the electrical measurements, Al dots of 0.152 mm² were evaporated through a shadow mask. The metal–insulator–semiconductor (MIS) structures were then annealed for 30 min at 450 °C in forming gas (Ar +10%H₂). The electrical properties were characterized with (I–V) measurements carried out with a HP 4140B picoammeter at various temperatures within the range of 150–500 K, the structure being biased in the accumulation regime. High frequency (1 MHz) C–V characteristics, performed at room temperature with an impedance analyzer (HP 4192 A), were used to estimate the density of the fixed charges in the film.

III. RESULTS AND DISCUSSION

A. Fixed charge distribution

In our previous study of SiNₓ/Si structures,16 we showed evidence for the presence of an interface layer with a higher N/Si ratio than the bulk film and having a lower refractive index than the bulk SiNₓ film. In the present study our C–V characteristics show a large flat band voltage shift (ΔV_{FB}) toward negative values which is indicative of fixed positive charges in the SiNₓ films. The ΔV_{FB} values indicate about 10¹³ cm⁻² positive charges assuming that the charges are at the interface of the SiNₓ/Si capacitor or in the bulk of the SiNₓ layer.13,15 This high level of positive charges has been previously observed in similar films.9,10 Figure 1 displays ΔV_{FB}−Φ_{MS} as a function of film thickness (d) where Φ_{MS} is the work function difference between the Al gate and the semiconductor. As ΔV_{FB} is due to an uniform fixed charge distribution, we would expect that ΔV_{FB} varies like d². Since Fig. 1 shows a linear relationship, we conclude that the fixed charges are not distributed uniformly in the film. If we assume that the fixed charges are mainly located at the SiNₓ/Si interface where the film is nitrogen rich, the charge distribution [Q_f(x)] could be described by the following:

\[ Q_f(x) = \begin{cases} \rho_0 & \text{for } 0 < x < d_1, \\ 0 & \text{for } x > d_1, \end{cases} \]

where d₁ is the characteristic length that describes the fixed charge location. Thus, ΔV_{FB} varies linearly with d² only for 0 < x < d₁ [Eq. (1)] and ΔV_{FB} should display a linear dependence with d for x > d₁ [Eq. (2)]. This charge profile is in better agreement with the results in Fig. 1 and provides further evidence that the SiNₓ/Si interface and SiNₓ bulk film are different.

B. Thick films

The current density versus applied electric field strength (J–E_{appl}) characteristics for thick SiNₓ films (78.3, 58.3, 40.5, and 21.4 nm) are shown in Fig. 2(a), and these data were obtained using an electric field ramp rate (r) of 5 × 10⁻³ MV/cm s. For low electric fields (<4 MV/cm), the
different $J - E_{\text{appl}}$ characteristics are similar, and from this initial (low field) region of the $J - E_{\text{appl}}$ data, we determine the film resistivity ($\rho$) from the slope of the $J - E_{\text{appl}}$ curve at low voltages as well as the critical field ($E_c$) which is defined as the field required for a 1 nA cm$^{-2}$ increase of the current density through the Al gate. The values of $\rho$ and $E_c$ for different thicknesses are reported in Table I. For films with thickness in the range of 20–78 nm, only small variations of $\rho$ and $E_c$ are observed and values of $0.7 \times 10^{16}$–1.7 $\times 10^{16}$ Ohm cm and 3.2–3.5 MV/cm, respectively, are obtained for $\rho$ and $E_c$, which are two electrical properties that describe the bulk behavior of SiN$_x$ films. As was the case for the film composition, $\rho$ and $E_c$ are independent of the film thickness.

For high electric fields (>4 MV/cm), the shape of the $J - E_{\text{appl}}$ data is different, as is shown in Figs. 2(a) and 2(b), and is directly related to the conduction mechanism in SiN$_x$ films. Figure 2(b) shows that ln$J$ is linear with $E_{\text{appl}}^{1/2}$ which suggests that the carrier transport process in SiN$_x$ films is via Poole–Frenkel emission, as is commonly observed, and the current–voltage characteristics can be expressed as

$$J = J_0 \exp \left( -\frac{q}{kT} \left( \phi - \beta \sqrt{E_{\text{appl}}} \right) \right),$$

where $J_0$, $\phi$, and $\beta$ are, respectively, the current density, the Poole–Frenkel barrier height, and a constant that depends on trap density. $E_{\text{appl}}$ is equal to $V/d$ where $V$ and $d$ are the applied voltage and the dielectric film thickness, respectively. $\beta$ is given by

$$\beta = \left( \frac{q}{\pi \varepsilon_0 \varepsilon_d} \right)^{1/2},$$

where $\varepsilon_0$ and $\varepsilon_d$ are, respectively, the dielectric constant of free space and the dynamic dielectric constant of SiN$_x$ films. Figure 2(b) shows that for all the film thickness studies there is a linear relationship between ln$J$ and $E_{\text{appl}}^{1/2}$. The dynamic dielectric constant is determined directly from the slope of the characteristic ln$J$ vs $E_{\text{appl}}^{1/2}$ and Eqs. (3) and (4). The results are reported in Table I. These results show that $\varepsilon_d$ is a strong function of film thickness while $\rho$, $E_c$, and the atomic ratio N/Si remain constant in this thickness range (20–78 nm). This suggests that the conduction mechanism changes with the thickness. For the determination of $\varepsilon_d$, it has been incorrectly assumed that the electric field is constant in SiN$_x$ films and equal to $E_{\text{appl}}$. In fact, it is well known that injected carriers (electrons in our case) from Si are trapped in the nitride film and a space charge is built up. Therefore, the electric field is different at the SiN$_x$/Si interface and in the bulk nitride film. The distribution of occupied electronic traps, $N_T$, is the result of the trapping and emission rates for electrons. Andrews et al. assumed that $N_T$ is an exponentially decreasing function of distance from the injecting interface and given as

$$N_T = N_0 \exp \left( -\frac{x}{x_0} \right),$$

where $N_0$ is the trap density and $x_0$ is the characteristic length that describes electronic occupation traps. The electric field distribution $E(x)$ is deduced by integration of the Poisson equation:

$$E = E_0 + E_T = 1 - \exp \left( -\frac{x}{x_0} \right),$$

where $E_0$ represents the electric field without charge in SiN$_x$ film, and $E_T$ is the static dielectric constant of the nitride film. Integrating the Poisson equation a second time yields the total voltage through the structure:

$$V = \int_0^d E(x) dx = (E_0 + E_T) d + E_T x_0 \left[ \exp \left( -\frac{d}{x_0} \right) - 1 \right].$$

From Eq. (7), the applied field can then be expressed as

$$E_{\text{appl}} = \frac{V}{d} = (E_0 + E_T) + \frac{E_T x_0}{d} \left[ \exp \left( -\frac{d}{x_0} \right) - 1 \right].$$

Close to the SiN$_x$ film/Si substrate interface, the traps are filled and a high negative space charge is present. Further into the dielectric, a point is reached where the traps are emptied ($N_T = 0$) due to field assisted emission [the Poole–Frenkel (PF) mechanism]. This occurs at a field $E_{\text{PF}}$ equals to

$$E_{\text{PF}} = E_0 + E_T - \frac{E_T x_0}{d} \left[ \exp \left( -\frac{d}{x_0} \right) - 1 \right].$$

If we assume that $d \gg x_0$, $E_{\text{PF}}$ can be written as

$$E_{\text{PF}} = E_{\text{appl}} + \frac{E_T x_0}{d}.$$

Finally, Eq. (3) should be written as

$$J = J_0 \exp \left( -\frac{q}{kT} \left( \phi - \beta_{\text{eff}} \sqrt{E_{\text{PF}}} \right) \right),$$

where $\beta_{\text{eff}}$ is given by an equation similar to Eq. (4):
where \( (\varepsilon_d)_{\text{eff}} \) is the effective dynamic dielectric constant of the film. From Eq. (10), \( E_{PF} \) decreases with increasing thickness. Therefore, for a given \( E_{\text{appl}} \), the current density is lower for thick films. This result is confirmed by the \( J-E_{\text{appl}} \) characteristics in Figs. 2(a) and 2(b), respectively. From Eq. (11), \( \ln J \) is proportional to \( E_{PF}^{1/2} \). Since we observe a linear relationship between \( \ln J \) and \( E_{\text{appl}}^{1/2} \), we deduce that \( E_{PF} \) is proportional to \( E_{\text{appl}} \) (\( E_{PF} = \alpha E_{\text{appl}} \), \( \alpha \) being a constant that decreases with increasing \( d \)). Using Eqs. (3) and (11), we find that the apparent \( \beta \) depends on nitride thickness:

\[
\frac{d \ln J}{d(E_{\text{appl}}^{1/2})} = \frac{q}{kT} \beta_{\text{eff}} = \frac{q}{kT} \frac{\beta_{\text{eff}}}{\alpha^{1/2}}.
\]

(12)

Figure 3 shows a linear relationship between the \( C_{\text{max}}/A \) ratio and \( 1/d \) where \( C_{\text{max}} \) and \( A \) are, respectively, the capacitance in the accumulation regime and the Al dot area. Thus we can assume that the static dielectric constant \( \varepsilon_S \) does not depend on film thickness. Furthermore, we can assume that \( \beta_{\text{eff}} \) is also independent of thickness (20–78 nm), since we observe no significant changes in \( \mu, \varepsilon_C, \) and \( \varepsilon_S \). Using Eq. (13), it is found that \( \beta \) decreases with increasing \( d \) and therefore the apparent \( \varepsilon_d \) increases with \( d \). Table I confirms this conclusion for film thicknesses in the range of 20–78 nm.

The value of the dynamic dielectric constant is quite high, especially for large film thicknesses (78.3 nm). Therefore, to confirm the Poole–Frenkel mechanism, it is necessary to examine the Arrhenius plots of leakage current for a constant electric field. Since the purpose of this article is not to find the charge distribution in SiNx films, the same current density, measured at room temperature, was chosen. Figure 4 displays Arrhenius plots of leakage current for two different film thicknesses (78.3 and 21.4 nm). At high temperature, the currents yield straight lines which strongly suggests that the electronic conduction corresponds to the Poole–Frenkel emission mechanism.
From the dependence of $\ln J$ vs $1/T$ in Fig. 4, the Poole–Frenkel barrier height $\Phi$ is found to be $1.2 \pm 0.2$ eV. In many silicon nitride films, the trap level $\Phi$ is in the range of 1–1.5 eV. Since evaluation of $\Phi$ by Poole–Frenkel emission has an error of $\pm 0.2$ eV, it is difficult to compare the conductivity of different silicon nitride films just by comparing the values of $\Phi$.

C. Thin films

Figure 5(a) and Table I show that, for thin films (<21 nm), there is a degradation of electrical properties. Contrary to Fig. 2(a), Fig. 5 shows that in the low field region the film characterizes are different, but this difference is essentially due to the electric field ramp rate ($r$) which is not the same (see Table I). In the low field region, the current strongly depends on the displacement current [$I = C(dV/dt)$] which is correlated to $r$ and depends slightly on the leakage current. As can be seen in Table I, the resistivity and critical field decrease with film thickness. This evolution could be correlated to the SiN$_x$/Si interface which plays a more important role for thin films. However, we observe a linear relationship between $\ln J$ and $E_{\text{appl}}^{1/2}$ which could mean that the current density is governed by Poole–Frenkel emission. The value of the dynamic dielectric constant determined from the slope of the characteristic $\ln J$ vs $E_{\text{appl}}^{1/2}$ is too high ($\varepsilon_d = 8.4$ for a thickness of 7 nm) to attribute the conduction mechanism to Poole–Frenkel emission. The current is not or not only limited by the bulk conduction through the SiN$_x$ film. An Arrhenius plot (Fig. 6) confirms this conclusion, and, contrary to Fig. 4, we observe a slight change of the current density as a function of the measurement temperature for thin films. Although the behavior is very similar to the Fowler–Nordheim emission process, the film is too thick (7 nm) to consider this conduction mechanism. The SiN$_x$/Si interface, which is more nitrogen rich and where the fixed charges are mainly located, now has more influence than the SiN$_x$ bulk film.

IV. CONCLUSION

In this article, a DECR plasma source has been used to deposit SiN$_x$ films, at floating temperature, with thicknesses in the range of 7–80 nm. For films thicker than 20 nm the influence of film thickness is minimal on the resistivity ($0.7–1.7 \times 10^{16} \Omega \text{cm}$) and on the critical field ($3.2–3.5 \text{MV/cm}$). For these films, the dominant conduction mechanism is Poole–Frenkel emission. The trap level, as determined from the temperature dependence of current density, is equal to $1.2 \pm 0.2$ eV. The assumption of a constant electric field throughout the structure is responsible for an apparent $\varepsilon_d$ variation. For thinner films (<20 nm), the electrical properties become degraded, particularly the critical field ($E_c = 1.45 \text{MV/cm for a 7 nm thick film}$). The conduction mechanism is not only determined by the bulk conduction; the SiN$_x$/Si interface also plays an important role. This interface presents properties which differ from those for the SiN$_x$ bulk film. We found, from $C–V$ characterization, that the fixed charges are mainly located at the SiN$_x$/Si interface.

In conclusion, the DECR process appears to be a promising technique by which to prepare gate dielectric films in a wide range of thicknesses.

ACKNOWLEDGMENTS

This work was supported by the Centre National de la Recherche Scientifique (GDR86) and France Telecom. One of the authors (E.A.I.) gratefully acknowledges the support of the National Science Foundation (NSF) Division of Materials Research.