

Accurate Determination of Ultrathin Gate Oxide Thickness and Effective Polysilicon Doping of CMOS Devices

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Abstract— We present an efficient and accurate method to characterize the physical thickness of ultrathin gate oxides (down to 25 Å) and the effective polysilicon doping of advanced CMOS devices. The method is based on the model for Fowler–Nordheim (F–N) tunneling current across the gate oxide with correction in gate voltage to account for the polysilicon–gate depletion. By fitting the model to measured data, both the gate oxide thickness and the effective poly doping are unambiguously determined. Unlike the traditional capacitance–voltage (C – V) technique that overestimates thin-oxide thickness and requires large area capacitor, this approach results in true physical thickness and the measurement can be performed on a standard sub-half micron transistor. The method is suitable for oxide thickness monitoring in manufacturing environments.

THE gate oxide thickness (T_{OX}) of CMOS transistors has been scaled down constantly with the reduction in gate length. With oxides approaching atomic dimensions, it is imperative we formulate methodologies to monitor and precisely control T_{OX} . Additionally, for modeling purposes, it is important to accurately determine the electrically active doping concentration of the polysilicon gate (N_P). However, the standard methods for extracting T_{OX} from capacitance–voltage (C – V) measurements of large area MOS capacitors are no longer accurate for thin oxides ($T_{OX} < 60$ Å) and consistently result in values larger than real physical thickness. This discrepancy between the electrical and physical/optical characterization techniques is due to the effects of finite thickness of inversion/accumulation layer (including quantum effects) and polysilicon depletion [1]–[4]. Until device simulators can incorporate these effects, there exists an urgent need to identify and characterize the difference between the electrical and physical values of T_{OX} . Even though the electrical T_{OX} can suffice for device modeling and design, for process design, physical T_{OX} is required. Here, we report an accurate and efficient approach for determining physical T_{OX} and effective N_P from measured tunneling current across the oxide.

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The Fowler–Nordheim (F–N) tunneling current density J_{FN} is given by [5]

$$J_{FN} = CE_{OX}^2 \exp[-\beta/E_{OX}] \quad (1)$$

where E_{OX} is the electric field across the oxide and C and β are constants; $C = 9.92 \times 10^{-7} \text{ AV}^{-2}$ and $\beta = 2.635 \times 10^8 \text{ V cm}^{-1}$ [6]. J_{FN} is a unique function of E_{OX} , which is related to oxide voltage V_{OX} . V_{OX} is a function of the applied gate voltage V_G (substrate grounded) and is given by

$$V_{OX} = E_{OX}T_{OX} = V_G - V_{FB} - \psi_S - V_P \quad (2)$$

where ψ_S is the Si surface potential, V_{FB} is the flat-band voltage, and V_P is the voltage drop in the poly. V_{FB} and V_P are given by

$$V_{FB} = \Phi_{MS} - Q_o/C_{OX} \cong \Phi_{MS} \quad (3)$$

$$V_P = \frac{\epsilon_{ox}^2 E_{OX}^2}{2q\epsilon_s N_P} \quad (4)$$

where $q\Phi_{MS}$, in eV (q being electron charge), is workfunction difference between poly and Si, Q_o is effective charge at Si/SiO₂ interface, C_{OX} is oxide capacitance, and ϵ_{ox} and ϵ_s are permittivities of SiO₂ and Si, respectively. For thin, good-quality gate oxides, $Q_o/C_{OX} < 0.01 \text{ V}$ ($Q_o/q < 10^{10}/\text{cm}^2$), and is negligible compared to Φ_{MS} . Φ_{MS} itself depends on the conduction type of both Si and poly. In this study, n⁺-poly nMOSFET's and p⁺-poly pMOSFET's are used. Substrate doping (N_{SUB}) for both n and pMOSFET is $2 \times 10^{17}/\text{cm}^3$. Assuming that poly has same band structure as Si and is degenerately doped, $\Phi_{MS} = -1.0 \text{ V}$ and $+1.0 \text{ V}$ for n and pMOSFET, respectively, at this N_{SUB} doping level. For V_P expression in (4), depletion approximation is used with uniform N_P [3]. Maximum possible V_P is 1.12 V (Si bandgap at 300 K) due to pinning of the Fermi level E_F at the band edges E_C and E_V . Substituting (4) in (2), we get

$$E_{OX} = \frac{1}{A} \left[-T_{OX} + \sqrt{T_{OX}^2 + 2A(V_G - V_{FB} - \psi_S)} \right]$$

where

$$A = \frac{\epsilon_{ox}^2}{q\epsilon_s N_P} \quad (5)$$

which is valid for the range where V_P varies with V_G . After saturation of V_P , E_{OX} is obtained from the general expression

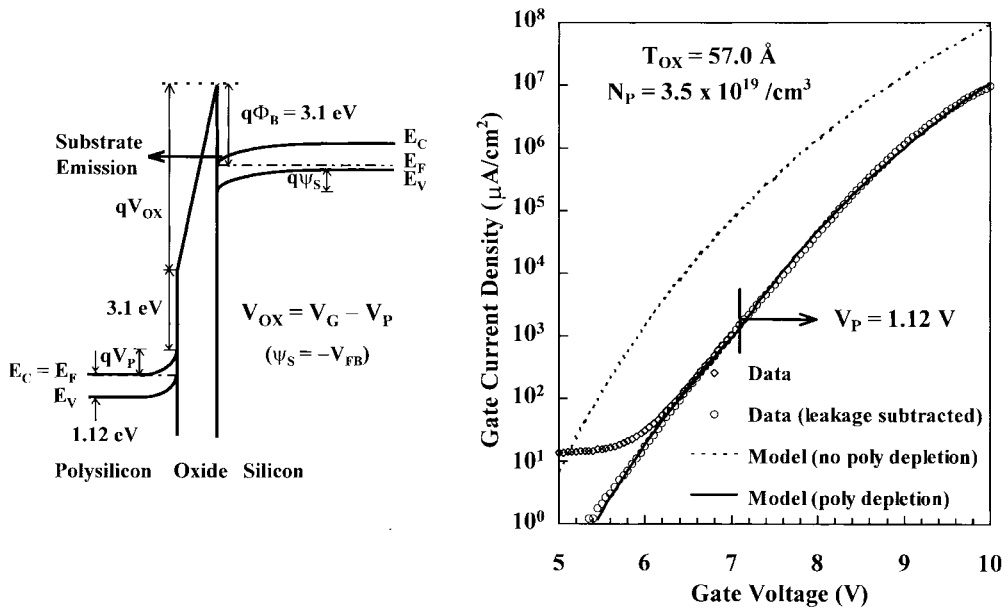


Fig. 1. Schematics of band diagram and fitting results for a $0.35\text{-}\mu\text{m}$ n^+ -poly nMOSFET biased under a positive V_G , resulting in F–N tunneling of electrons from the substrate. The model with poly depletion matches with the data.

in (2). Using T_{OX} and N_P as fitting parameters E_{OX} can be calculated for the complete F–N bias range and used to predict or match the tunneling current.

Fig. 1 shows the schematics of band diagram and fitting results for a nMOSFET biased with positive V_G (source, drain, and substrate grounded) such that the substrate is inverted and emits electrons. The positive V_G results in depletion of poly and its extent depends on V_G . The maximum possible $V_P = 1.12$ V is shown. For F–N tunneling bias range, ψ_S is a constant since E_F at the Si surface is pinned at E_C , as shown. For $N_{SUB} = 2 \times 10^{17}/\text{cm}^3$, $\psi_S = +1$ V which is equal to $-V_{FB}$. In fact, for n^+ -poly nMOSFET, if E_F at the Si surface is pinned at E_C and Q_o/C_{OX} is negligible, then $\psi_S = -V_{FB}$, regardless of N_{SUB} . Thus, from (2), $V_{OX} = V_G - V_P$. As seen, J_{FN} predicted by the model with poly depletion matches well with data. Fitting was first performed for the range of V_G where V_P is saturated and T_{OX} is the only variable. A difference of 0.5 Å was distinguishable and for the device shown, the best fit is obtained for $T_{OX} = 57.0$ Å. Then, fixing T_{OX} and varying N_P , fitting was performed for lower V_G as shown in Fig. 2. A difference of $0.5 \times 10^{19}/\text{cm}^3$ in N_P is easily discernible. The best fit is realized for $N_P = 3.5 \times 10^{19}/\text{cm}^3$.

Fig. 3 shows that with the same J_{FN} model and constant $N_P = 3.5 \times 10^{19}/\text{cm}^3$, good match is obtained with data for devices fabricated with same process and target T_{OX} ranging from 60 Å to 25 Å. The oscillations in data for $T_{OX} = 36.5$ Å are due to electron wave interference which is T_{OX} dependent [7], [8]. Recently, these oscillations have been used to extract T_{OX} [8]. However, the technique proposed in this letter is simpler and can be used to monitor T_{OX} variation in manufacturing. T_{OX} characterization results by three different methods are compared in Fig. 4. For optical measurements, Thermo-Wave's Opti-Probe 6131 measurement system, which uses a Beam Profile Ellipsometer to measure ultrathin film thickness, was used. For the C – V data, inversion value is

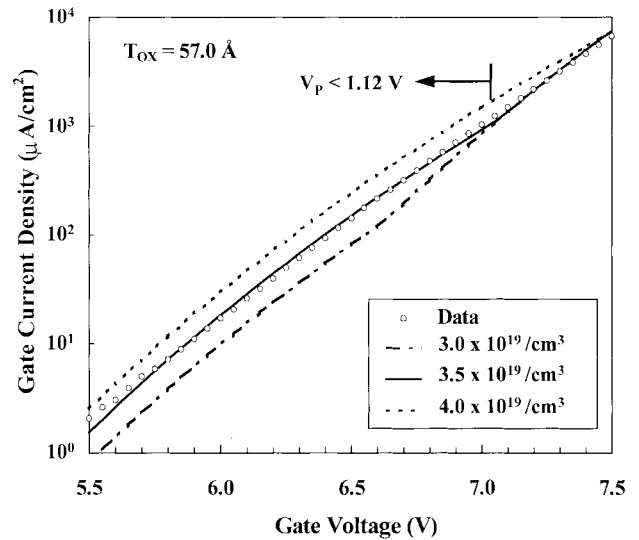


Fig. 2. Sensitivity of the F–N tunneling current model to variation in polysilicon doping in the low current ($V_P < 1.12$ V) range. The oxide thickness is kept constant at the value (57 Å) obtained from fitting the model to the high current ($V_P = 1.12$ V) range.

used at $E_{OX} \cong 2.5$ MV/cm. To reduce the direct-tunneling leakage current, the substrate was floating during the measurement and the source and drain were grounded. It should be noted that the optical measurements were performed on test wafers and hence, there is an inherent process variation. Still, physical values obtained from J_{FN} match well with optical measurement results, whereas the electrical values obtained from C – V consistently overestimate the thickness.

The case of a pMOSFET biased with positive V_G (substrate emission) is different from the nMOSFET. The Positive V_G results in accumulation of the p^+ -poly, and $V_P = 0$ since E_F is pinned at E_V . Consequently, N_P is not a fitting parameter. Si substrate is also in accumulation with $E_V = E_F$ at the

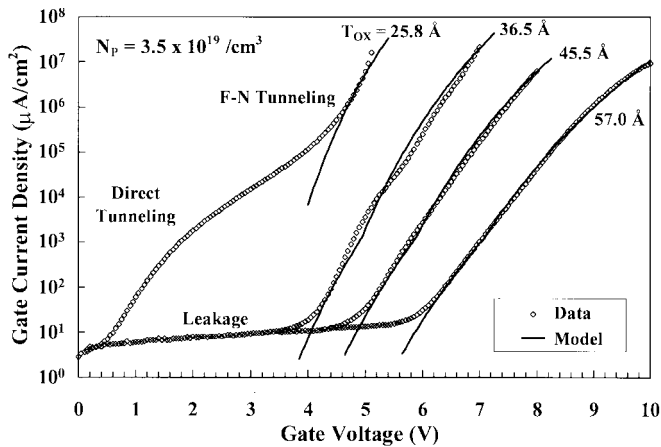


Fig. 3. Substrate emission F-N tunneling current density characteristics for $0.35\ \mu\text{m}$ n^+ -poly nMOSFET's fabricated with same process but different oxide thickness. With constant poly doping concentration ($3.5 \times 10^{19}/\text{cm}^3$), model matches with data for all the thickness.

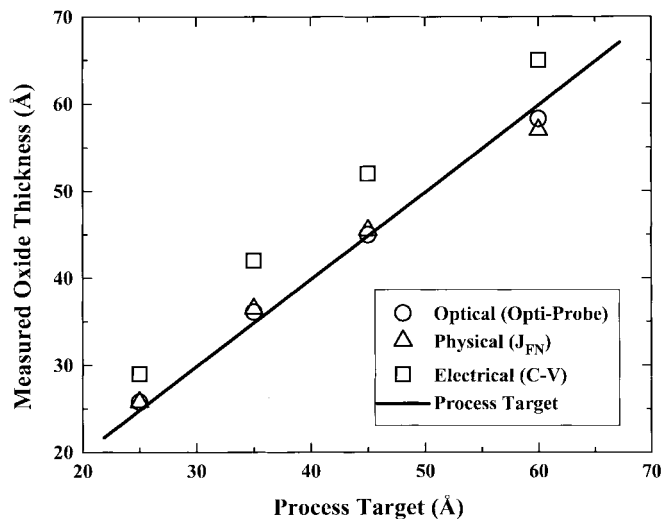


Fig. 4. Oxide thickness characterized by three different measurement techniques. Physical values, obtained from F-N tunneling current data, are consistent with the optical results, whereas the electrical values, obtained from $C-V$ data, are an overestimate.

surface. Hence, $\psi_S = 0.12\ \text{V}$ and from (2), $V_{\text{OX}} = V_G - 1.12$. Like the nMOSFET, for F-N tunneling bias range in a p^+ -poly pMOSFET, this result is independent of N_{SUB} . Consistent with optical measurements, $T_{\text{OX}} = 58.3\ \text{\AA}$ was obtained by fitting the model to data. This result is also consistent with the nMOSFET substrate emission case discussed earlier. Both the n^+ -poly nMOSFET and the p^+ -poly pMOSFET devices are on the same wafer and the difference of $1.3\ \text{\AA}$ in the results is within the limits of processing variation and the measurement error of this technique. Similar results were obtained for nMOSFET gate emission.

The results so far indicate that J_{FN} model for a nMOSFET substrate emission can be used to accurately characterize T_{OX} and N_P . We further developed this method into a wafer level electrical test technique to monitor T_{OX} during manufacturing. This technique can work even with both T_{OX} and N_P varying, as is expected in a manufacturing environment, if a sufficiently high fixed current is chosen (for example $1\ \text{A}/\text{cm}^2$) such that E_F on the n^+ -poly side is pinned at E_V . The V_G corresponding to this fixed high current is measured. T_{OX} is then calculated from the measured V_G using (2) as band bending in poly is saturated with fixed $V_P = 1.12\ \text{V}$ and poly doping is no longer required for the calculation. This technique was used to monitor T_{OX} variation at different die locations on a wafer. The sensitivity was found to be $0.1\ \text{\AA}$ and maximum error due to the oscillations in data was $\pm 1.5\ \text{\AA}$ [9].

In summary, the J_{FN} model has been shown to accurately characterize true physical thickness of ultrathin gate oxides ($60\ \text{\AA}$ to $25\ \text{\AA}$) after correction is made in V_G to account for poly depletion. Also, the electrically active poly doping concentration is obtained by fitting the model to n^+ -poly nMOSFET substrate emission data. A single point measurement of $J_{\text{FN}} = 1\ \text{A}/\text{cm}^2$ results in V_G being linearly proportional to T_{OX} and has been used to monitor T_{OX} variation on a wafer. The sensitivity is $0.1\ \text{\AA}$, with a maximum error of $\pm 1.5\ \text{\AA}$.

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