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INFLUENCE OF THE FABRICATION CONDITIONS ON THE p⁺-TaSi₂/POLY-Si GATE QUALITY

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Résumé - L'effet des différentes étapes de fabrication du poly-siliciure sur la qualité résultante du dopage p⁺ de la grille est examiné. En particulier on étudie la diffusion de bore en corrélation avec la contamination en oxygène. Le rôle de l'interface TaSi₂/poly-Si et de l'atmosphère de recuit sur la redistribution du bore est montré.

Abstract - The effect of different polycide fabrication steps on the resulting p⁺ gate quality is investigated. In particular the boron diffusion is studied in correlation with the oxygen contamination. The role of the quality of the TaSi₂/poly-Si interface and of the annealing atmosphere on the boron redistribution are pointed out

1 - INTRODUCTION

The TaSi₂/poly-Si bilayers constitute a well suited system for the fabrication of p⁺-gates /1/. Nevertheless controversial results concerning the boron redistribution after implantation into the TaSi₂ layer have been reported /1,2/, questioning the p⁺ TaSi₂ polycide quality, i.e. the p⁺ work function ϕ_{ms} reproducibility. Earlier experiments suggested a correlation of the boron profiles with the oxygen distribution in TaSi₂ /3/. Recently, evidence for TaB₂ formation after high dose implantation ($2 \times 10^{16} \text{cm}^{-2}$) has been reported /4/. However, the effect of oxygen contamination in TaSi₂ or Ta-B compound formation on the B outdiffusion from TaSi₂ into the poly-Si is yet unclear.

In this paper we first discuss the effects of an interfacial oxide on the B transport across the TaSi₂/poly-Si interface and point out the importance of poly-Si surface cleaning prior to TaSi₂ deposition for good p⁺ polycide behavior. Next, the role of oxygen on the B diffusion within the silicide is discussed. Finally the effects of annealing ambient on the B redistribution in the p⁺ TaSi₂ polycide are investigated.

2 - EXPERIMENTAL

MOS-capacitors (0.01 to 8 mm²) with TaSi₂/poly-Si gates were fabricated on n- or p-type (100) Si. The gate oxide thickness was 25 nm. The poly-Si cleaning prior to TaSi₂ sputtering consisted of a buffered HF dip and an in-situ sputter cleaning with Ar ions. The silicide films were prepared by Ta and Si cosputtering to a thickness of 200 nm. The Ta-Si sandwich thickness was $\approx 1.3 \text{nm}$. After TaSi₂ deposition on poly-Si (100 nm) the wafers were covered with a CVD-oxide (100 nm). B was implanted either into TaSi₂ or into poly-Si prior to Ta-Si sputtering. The B doses used were $5 \times 10^{15} \text{cm}^{-2}$ and $2 \times 10^{16} \text{cm}^{-2}$. Finally the samples were annealed at 900°C in a N₂ or Ar/H₂ atmosphere.

SIMS profiles of B and oxygen were recorded using oxygen (O₂⁺) and cesium (Cs⁺) primary ions, respectively. For Capacitance-Voltage C(V) measurements the oxide cap was removed. SIMS measurements were done before and after the oxide cap etching; the SIMS results for samples with and without oxide cap did not differ. B implantation standards for poly-Si and TaSi₂ were used to calibrate the B SIMS signal. Unimplanted polycide samples were used to determine the background SIMS signal. For B (O⁺ primary ions) the background was $3-5 \times 10^{17} \text{cm}^{-3}$ in TaSi₂ and $2-4 \times 10^{16} \text{cm}^{-3}$ in poly-si. The oxygen profiles (Cs⁺ primary ions) were limited by a background of about 10³ counts.

3 - RESULT AND DISCUSSION

3.1 - TaSi₂/POLY-Si INTERFACE

Earlier experiments with TaSi₂ on mono-Si have shown, that interfacial oxides impede B transport from TaSi₂ into mono-Si /5/ for T≤900°C. At the barrier brakes down /2,5/, but this temperature is not compatible with modern CMOS processes. In order to elucidate the role of interfacial oxide on poly-Si on the B transport across the TaSi₂/poly-Si interface at 900°C, samples with and without the poly-Si surface cleaning were investigated.

The B profiles for the samples with appropriate poly-Si surface cleaning and with interfacial oxide are shown in Fig. 1. The B profiles can be roughly divided into a region of low B diffusivity (the implanted region), and a tail region of fast diffusing B (grain boundary enhanced diffusion) /1,6/. There is no significant difference between the low diffusivity regions of both type of samples. In contrast the tail regions differ significantly (Fig. 1). For the clean interface sample the flat tail saturates at $1-2 \times 10^{19} \text{cm}^{-3}$ /1,6/. For the sample with interfacial oxide the B concentration in the tail region continuously decreases. Additionally a pronounced peak is observed at the TaSi₂/poly-Si interface. In the poly-Si the B doping is very low. At the poly-Si/gate-oxide interface the B concentration is as low as or lower than $1 \times 10^{17} \text{cm}^{-3}$.

A B pile-up is a characteristic of interfaces between thick oxides and silicide /7,8/. Fig. 2 shows an example of B accumulation at the oxide-cap/TaSi₂ interface. By increasing the implantation energy the B profile peak is moved deeper into TaSi₂, and the B accumulation near the oxide cap (100nm) is separated from the implanted region. In the case of interfacial oxide the resulting B profile exhibits a similar B pile-up at the TaSi₂/poly-Si interface (Fig. 1, curve b), although in this case the oxide thickness is only 3-5nm. Furthermore, the characteristic B distribution in the tail region for the TaSi₂/poly-Si bilayers (Fig. 1, curve a) is altered. In the sample with interfacial oxide the B profile within the TaSi₂ layer strongly resembles those obtained for TaSi₂ between two thick oxide layers /7/. The fast diffusing B accumulates in TaSi₂ near and at the TaSi₂/poly-Si interface inducing a B concentration decrease in the tail region deeper in the TaSi₂ layer.

C(V) measurements confirmed the SIMS results. For the sample with interfacial

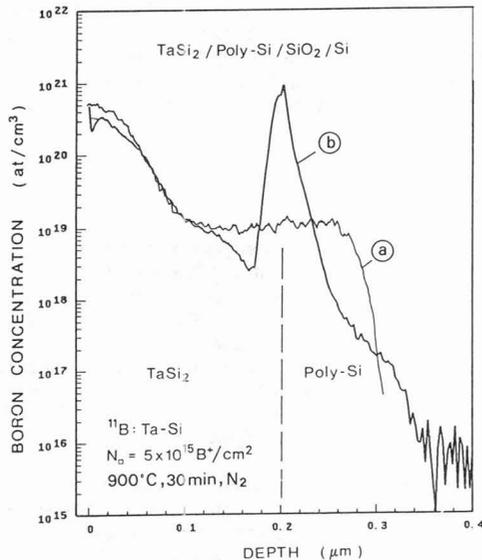


Fig. 1

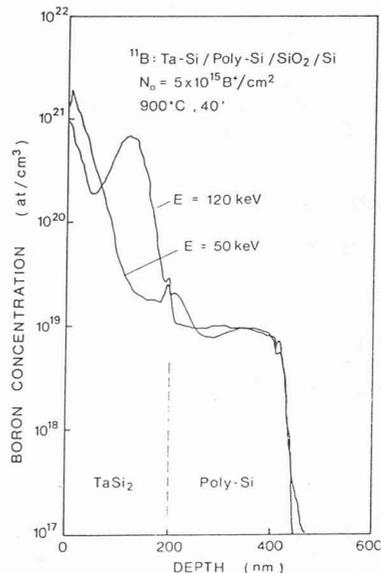


Fig. 2

Fig. 1 Boron profiles after annealing at 900°C in N₂ for samples a) with poly-Si surface cleaning and b) with interfacial oxide.
 Fig. 2 Boron profiles after annealing in N₂ for two implantation energies. Boron was implanted into the as-deposited silicide.

oxide the C(V) characteristic are shifted to more negative gate voltage values indicating a lower $p^+ - \phi_M$. This consistent with the insufficient doping of the underlying poly-Si from SIMS measurements.

3.2 - OXYGEN CONTAMINATION

Fig. 3 shows the oxygen profiles for the samples with poly-Si cleaning and with interfacial oxide. The corresponding B profiles are shown in Fig. 1. The oxygen concentration in the as-deposited TaSi₂ layer has a constant value throughout the layer and the poly-Si is largely oxygen free. For annealed samples with poly-Si surface cleaning a decrease of the oxygen concentration towards the oxide/TaSi₂ and TaSi₂/poly-Si interfaces is observed. Additionally, oxygen diffuses into the poly-Si increasing its oxygen content by several orders of magnitude. For the sample with interfacial oxide the oxygen concentration remains quite constant in TaSi₂ and increases near the oxide/TaSi₂ interface. The poly-Si remains practically oxygen free. The interfacial oxide appears as a small oxygen peak at the TaSi₂ interface. Apparently an interfacial oxide also hinders the oxygen diffusion across the interface.

The oxygen profiles for samples with poly-Si cleaning with and without B implantation were alike. Moreover for a B dose increase from $5 \times 10^{15} \text{cm}^{-2}$ to $2 \times 10^{16} \text{cm}^{-2}$ no dose effect related to the oxygen distribution is found. It follows that neither the presence of B nor the implantation affects the oxygen distribution in the polycide. A similar oxygen distribution is also expected in the case of B implantation into the poly-Si prior to TaSi₂ deposition. In this way implantation effects within TaSi₂ are avoided. As shown in Fig. 4, a widely uniform B concentration is obtained and again a B pile-up near the oxide/TaSi₂ interface is observed. The oxygen concentration near the oxide/TaSi₂ interfaces can decrease, increase or remain at a constant value

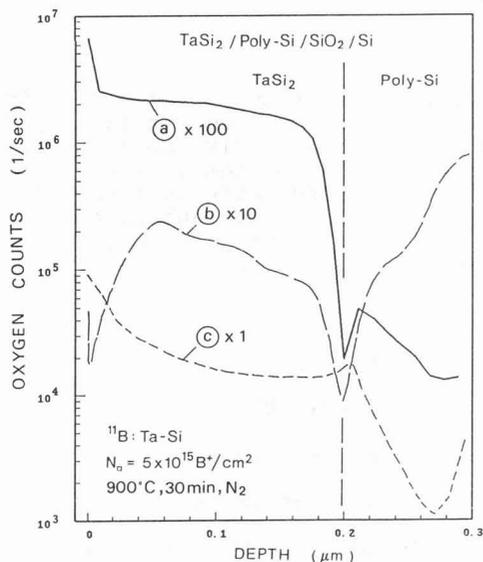


Fig. 3

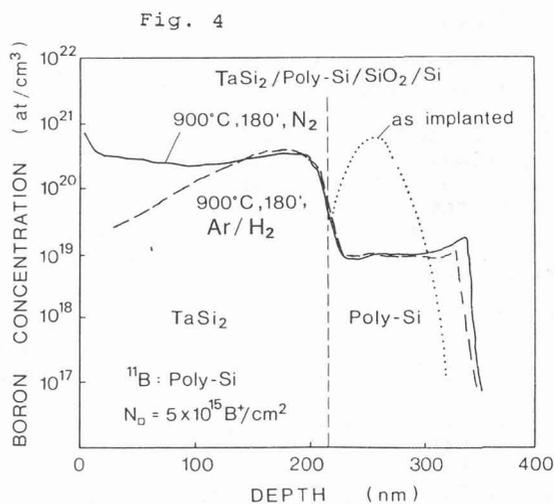


Fig. 4

Fig. 3 Oxygen profiles for a) as-deposited TaSi₂, and after annealing at 900°C in N₂ for samples b) with poly-Si surface cleaning and c) with interfacial oxide.

Fig. 4 Boron profiles in case of boron implantation into poly-Si prior to TaSi₂ deposition: a) as-implanted and after annealing in b) N₂ and c) Ar/H₂.

though in either case B accumulates near the oxide/TaSi₂ interfaces. This suggests that oxygen does not significantly affect the B redistribution.

3.3 - ANNEALING AMBIENT

For samples annealed in Ar/H₂ dopant loss has been detected in spite of a 100nm oxide cap /3/. In order to investigate the dependence of the B redistribution on the annealing ambient (Ar/H₂ or N₂) independently of implantation effects in TaSi₂, B was implanted into undoped poly-Si prior to TaSi₂ deposition. The same poly-Si surface cleaning was used as described before.

Typically for CMOS processes, after the gate fabrication step the remaining 900°C processing time is 180-250 min. Fig. 4 shows the resulting B profiles after annealing in Ar/H₂ or in N₂ at 900°C for 180 min. After annealing most of the implanted B diffuses into the TaSi₂ layer. If annealed in N₂ a high and uniform B concentration ($\approx 2 \times 10^{20} \text{cm}^{-3}$) is achieved. For Ar/H₂ anneals a considerable dopant loss occurs in spite of the 100nm oxide cap. This is due to the increase of the B diffusivity in oxide by several orders of magnitude in case of H₂ containing annealing ambients /9/. Such a B loss to the oxide cap is only possible if a sufficient B supply is assured by the TaSi₂ layer. This suggests that the grain boundary diffusion is the dominant mechanism for the B redistribution even for long annealing times. Other mechanisms such as Ta-B compound formation do not seem to significantly affect the B redistribution. Nevertheless this does not rule out that B in TaSi₂ partly exists as TaB₂.

A rough estimate gives that approximately half of the implanted dose is lost if annealed in Ar/H₂. Although for B dose $\geq 5 \times 10^{15} \text{cm}^{-2}$ the polycide p⁺- ϕ_M remains unchanged (B concentration in poly-Si $\approx 1 \times 10^{19} \text{cm}^{-3}$). But, on the other hand, the B contamination of the furnace might be of concern. The poly-Si doping does not depend on the B dose for doses $\geq 5 \times 10^{14} \text{cm}^{-2}$ and for low B doses the resulting B redistribution is quite independent of the implantation scheme /6/. If low doses are used, however, subsequent processing must avoid H₂ containing anneals in order to assure p⁺- ϕ_M reproducibility.

4 - CONCLUSION

In summary we have shown that a native oxide constitutes an effective diffusion barrier for B. Therefore the cleanliness of the TaSi₂/poly-Si interface is a crucial factor for reproducible p⁺-TaSi₂ polycide gates. The content and distribution of oxygen in the polycide does not affect the B diffusion. Annealing at 900°C in H₂ containing ambients leads to dopant loss in spite of a thick oxide cap. In case of long annealing times and/or of low B doses the dopant loss would reduce the p⁺-doping of the p⁺ polycide, i.e. of the underlying poly-Si layer, deteriorating the MOS properties.

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