

Ultra-Shallow Junction Formation by AsH₃ Adsorption Method

Y.H. Song, S.Pidin, T.Kurabayashi¹, H.Kurino and M.Koyanagi

Department of Machine Intelligence and System Engineering,

¹ Department of Mechatronics and Precision Engineering,

Tohoku University, Aramaki, Aoba-ku, Sendai 980-77, Japan

Phone: +81-22-217-6909; Fax: +81-22-217-6907; E-mail: song@sd.mech.tohoku.ac.jp

1. Introduction

Table 1 shows the technical comparison for the recent N-type doping methods. Even though the technical originality of several doping methods, process feasibility and controllability are still questionable in sub 0.1 μm MOSFET fabrication. In particular, low energy and new implantation method cannot avoid a thermal annealing to cure the implantation damage. AsH₃ adsorption in GSMBE provides ultra-shallow junction with wide temperature margin which means good controllability, and no extra annealing and simple integration which mean good process feasibility in sub 0.1 μm MOSFET fabrication.

2. Experimental

A GSMBE apparatus is shown schematically in Fig.1. The base pressure of the main chamber was less than 3×10^{-10} Torr. AsH₃ gas diluted to 10% concentration by H₂ gas was used as the N-type doping gas for junction formation. The standard process sequence was carried out as shown in Fig.2. After thermal cleaning, 10SCCM AsH₃ gas was injected into main chamber at the variable temperatures and adsorption times for the arsenic doping.

3. Results and discussion

Fig.3 shows the typical arsenic SIMS depth profile for AsH₃ adsorption. Very shallow (X_j ; 300 Å) and heavy doping (C_p ; above $1 \times 10^{20} \text{cm}^{-3}$) profile was obtained at 850°C, 5 min doping condition. Fig.4 shows the temperature effect of AsH₃ adsorption. The As peak concentrations and junction depths ($\sim 10^{18} \text{cm}^{-3}$)

showed the saturation trend at above 550°C. The stable junction characteristics can be achieved at the temperature range from 700°C to 950°C as shown in Fig.4. These experimental results were quite well matched to arsenic coverage simulation data of Fig.5-(1) [1]. Fig.6 shows the change of arsenic concentration at the increasing adsorption times. It was found that arsenic concentrations reached to some maximum values and then went to decrease with increasing doping time. Initial stage of increasing concentration can be explained from the simulation data of adsorption time dependence (Fig.5-(2)) but the second stage of decreasing concentration cannot be well understood. This mechanism will be verified from the further experiments. The higher C_p (peak concentration) and longer peak time for one of 850°C compared with 700°C can be attributed to higher surface desorption and diffusivity. Fig.7 shows in-situ annealing effect at 850°C. As shown in Fig.7, increasing annealing time gave negative effect for the sheet resistance, which means that extra annealing is not necessary for junction formation. Finally, Fig.8 shows the strong correlation between arsenic concentrations and junction depths.

4. Conclusion

From the results of junction characteristics, it is suggested that AsH₃ adsorption method is very effective for N-type ultra-shallow junction formation.

References

- [1] L.Kipp, The American Physical Review B, Vol. 50, N8, 15 August 1994, p.5448 - 5455.

method	controllability	process and scaling feasibility	simple integration	damage and defect free
SPD	bad	good	good	must anneal
ion implant	good	good	very good	must anneal
doped epi.	good	bad	fair	need anneal /epi -defect
delta doping	fair	bad	fair	must anneal
inversion ST	fair	very bad	fair	skip anneal
AsH3 doiping	good	good	very good	skip anneal

Table1. N-type doping technology comparison.
SPD means solid phase diffusion method.

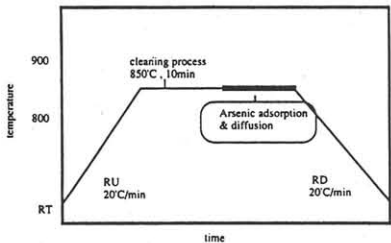


Fig. 2. Standard process sequence for AsH3 adsorption.

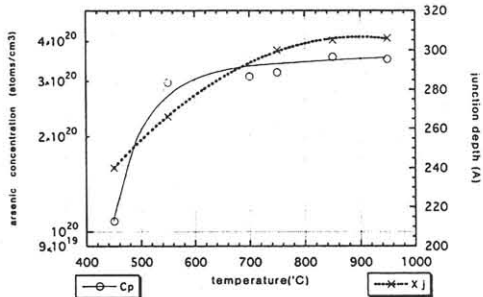


Fig. 4. Temperature dependence of junction characteristics. The only 5min adsorption without extra annealing was done.

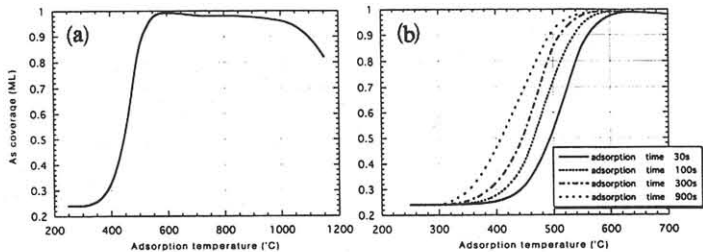


Fig. 5. Arsenic coverage simulation data using a model with Langmuir adsorption and first-order desorption reaction, curve(a) is temperature dependence at adsorption time 300s, (b) is adsorption time dependence at the variable temperature.

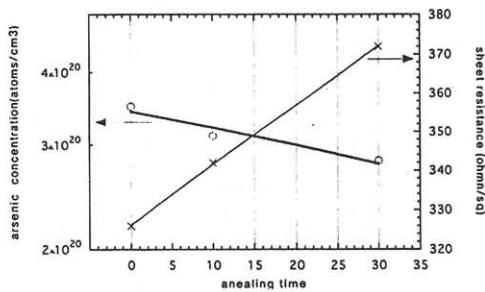


Fig. 7. Annealing effect at 850°C. Adsorption time was 5 min.

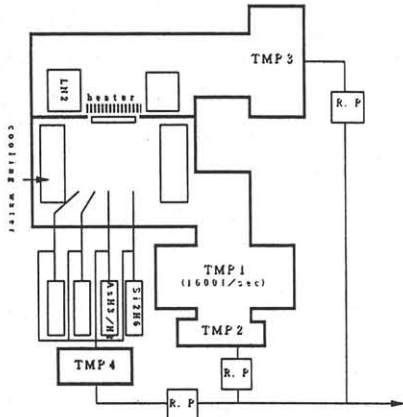


Fig. 1. Schematic diagram of gas source MBE.

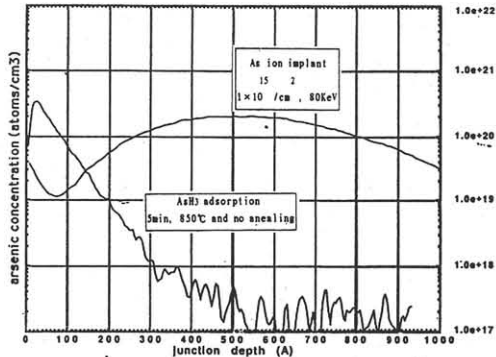


Fig. 3. SIMS depth profile of AsH3 adsorption. Arsenic ion implantation for the comparison was done.

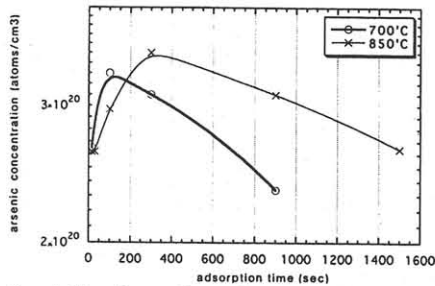


Fig. 6. AsH3 adsorption time dependence of arsenic concentration at temperatures of 850 and 700°C.

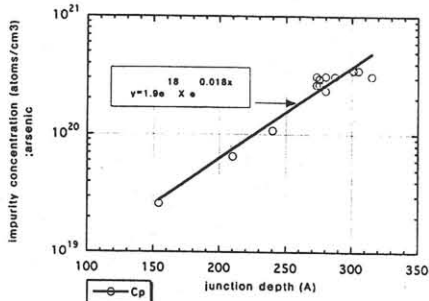


Fig. 8. Correlation result for Cp vs Xj. It showed good exponential relation with R. F of 0.92.