Transport via dopant-quantum-dots fabricated by thermal diffusion through nano-masks

^oDaniel Moraru^{1,*}, Arup Samanta¹, Le The Anh², Takeshi Mizuno¹, Hiroshi Mizuta^{2,3}, and Michiharu Tabe¹

¹Research Institute of Electronics, Shizuoka University ²School of Materials Science, Japan Advanced Institute of Science and Technology ³Nano Group, ECS, Faculty of Physical and Applied Sciences, University of Southampton, UK

E-mail: daniel@rie.shizuoka.ac.jp

Introduction

In recent years, we have been studying dopant-atom devices that work by single-electron tunnelling via single dopant atoms as quantum dots (QDs) [1]. These devices can offer a broad functionality range, also based on a different transport mechanism as compared with scaled-down conventional transistors. Recently, the field of single-dopant devices has been steadily developed and advanced through several breakthrough studies [2-6].

It should be noted, however, that, in most devices, the dopants were introduced in the channel with

conventional doping techniques, in random positions, and devices exhibiting single-dopant characteristics were picked up for more detailed study. Only a few works addressed directly the control of dopants, either in number, using single ion implantation [7], or in position, with atomic manipulation using scanning tunnelling microscope tips [8]. These are, however, state-of-the-art techniques, which are not fully suitable for CMOS fabrication.

In this work, we propose and demonstrate fabrication of dopant-based devices, with the dopant position controlled using precisely patterned doping masks.

Fabrication and electrical characteristics of selectively-doped SOI-FETs

The nanoscale channel of silicon-on-insulator (SOI) transistors [Fig. 1(a)] was selectively doped with phosphorus (P) within an area of ~30 nm in width, using a thermal diffusion process. The doping process is self-aligned, so that the dopant-quantum dots have a well-determined position relative to the leads, as shown in Fig. 1(b). Different from our previous works [1,4,6], doping concentration was higher ($N_D \cong 5 \times 10^{18} \text{ cm}^{-3}$), so average distance between P donors is <5 nm. This increases the probability of forming multiple-donor QDs within the selectively-doped area when several P donors are located near each other.

Electrical characteristics were measured for these devices, starting from low temperatures (T > 6 K), as shown in Fig. 2. Current peaks ascribed to tunnelling via dopant-QDs can be observed. The impact of the localized doping can be understood by comparison with uniformly-doped and nominally-undoped-channel FETs (not shown here), fabricated with the same fabrication process. The peaks also exhibit a noticeable substructure, which reflects the density-of-states inside the QD.

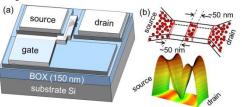


Fig. 1. (a) Structure of an SOI-MOSFET studied. (b) Top: FET channel selectively doped in nanoscale using thermal diffusion process. Bottom: possible potential landscape induced by localized donors.

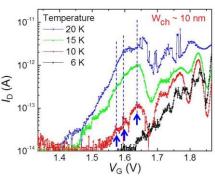


Fig. 2. I_D - V_G characteristics for a selectively-doped nano-FET at low T (T > 6 K). The low- V_G current peak reveals a clear sub-structure, with several inflections (as indicated by arrows), at the lowest temperatures.

These results provide information about the interaction between neighbouring dopant atoms, which may accelerate further development of dopant-based devices and technology.

Acknowledgments: The authors thank Y. Kuzuya, T. Nagasaka, and E. Hamid for their contributions in experiments. This work was partly supported by Grants in Aid for Scientific Research from MEXT Japan: 23226009, 25630144, and 2231085. References [1] D. Moraru *et al.*, Nanoscale Res. Lett. **6**, 479 (2011). [2] H. Sellier *et al.*, Phys. Rev. Lett. **97**, 206805 (2006). [3] Y. Ono *et al.*, Appl. Phys. Lett. **90**, 102106 (2007). [4] M. Tabe *et al.*, Phys. Rev. Lett. **105**, 016803 (2010). [5] M. Pierre *et al.*, Nature Nanotechnol. **5**, 133 (2010). [6] E. Hamid *et al.*, Phys. Rev. B **87**, 085420 (2013). [7] E. Prati *et al.*, Nature Nanotechnol. **7**, 443 (2012). [8] M. Fuechsle *et al.*, Nature Nanotechnol. **7**, 242 (2012).