

Nonvolatile Memory with Multi-Stack Nanocrystals as Floating Gates

Kamran Afshari

Electrical Engineering, University of California-Los Angeles

NNIN REU Site: Cornell NanoScale Science & Technology Facility, Cornell University

NNIN REU Principal Investigator: Prof. Edwin Kan, Electrical & Computer Engineering, Cornell University

NNIN REU Mentor: Tuo-Hung Hou, Electrical & Computer Engineering, Cornell University

Contact: kafshari@ieee.org, kan@ece.cornell.edu

Abstract

Nonvolatile memory technologies are focusing on devices with longer retention, faster read and write, higher bit density, improved endurance, and low-voltage program/erase characteristics. Nanocrystal (NC) memories are promising for realizing high-density nonvolatile storage with the inherent advantage of low-voltage operation [1]. Recently carbon molecules in the form of fullerenes (C_{60}), known as “bucky balls,” also have been incorporated in non-volatile memory devices [2] with its advantages of mono-disperse nature and molecular size. In this project, various memory structures with self-assembled multi-stacked gold, platinum, and C_{60} nanocrystals as floating gates have been fabricated and characterized. In two-layer nanocrystal structures, the C_{60} bottom layer acts as an additional barrier to prevent charge back-tunneling from the upper layer, improving retention time without a commensurate penalty in program time. The upper nanocrystal layer functions as additional charge storage to provide sufficient memory window.

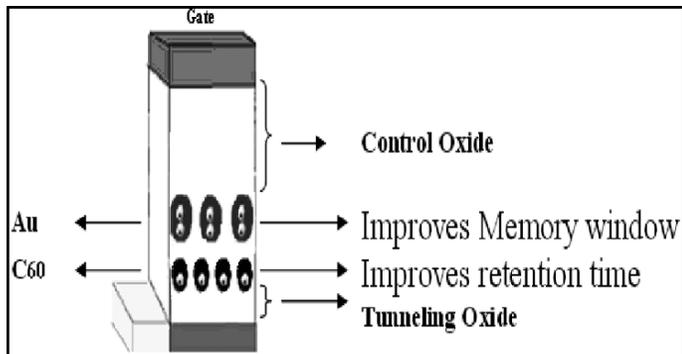


Figure 1: Schematic of the double-layer nanocrystal memory.

Introduction

Figure 1 shows the cross section of a double-layer nanocrystal memory device. By applying enough positive bias across the gate, electrons will quantum-mechanically tunnel through the thin tunneling oxide. If the lower-layer nanocrystal is designed so that it is energetically unfavorable for charge storage due to its small size, electrons will step through the lower nanocrystal layer, and get stored in the second layer nanocrystal. After taking away the applied potential, the energy penalty for electrons in the second layer to tunnel back into the first layer is larger enough, again due to the large charging energy associated with the small size. The first-layer nanocrystal acts as a part of tunneling barrier to provide prolonged retention time. The same concept applies to the hole storage when negative bias is applied. Charging nanocrystals with electrons and holes can require different gate biasing due to different properties of nanocrystals, which also contribute to different retention time characteristics.

The goal was to engineer the best structure such that one can obtain a memory device with an optimal memory window and retention time. This required the optimization of the tunneling oxide thickness, nanocrystal diameter and the choice of a nanocrystal material.

Experimental Procedure

P-type wafers with a doping level of 10^{15} cm^{-3} were used in this study. The device isolation was achieved by the local oxidation isolation (LOCOS). 2-3 nanometers of tunneling oxide was grown on the active region in a diluted oxygen ambience using an atmospheric-pressure furnace. Each wafer was named separately.

Sample 1 (S_1) was the control device without any floating gate, sample 2 (S_2) had a single-layer gold (Au) nanocrystal as a floating gate, S_3 had a double-layer Au nanocrystal floating gate, S_4 had a single-layer C_{60} nanocrystal floating gate, S_5 had a double-layer C_{60} and Au nanocrystal floating gate, and S_6 had the same structure as S_5 but with a thinner thickness of tunneling oxide.

The Au nanocrystals were deposited through e-gun evaporation while the C_{60} nanocrystals through the thermal evaporation. For those samples with double-layer nanocrystals, 2-3 nm of evaporated silicon dioxide (SiO_2) was inserted between two layers. After the floating-gate formation, 20-30 nm of control oxide was deposited through a plasma-enhanced chemical vapor deposition (PECVD), followed by chromium and aluminum evaporation as the control gate. Finally the control gate was patterned to finish the device fabrication. The device was now measured, and memory window and retention time for different configurations were compared.

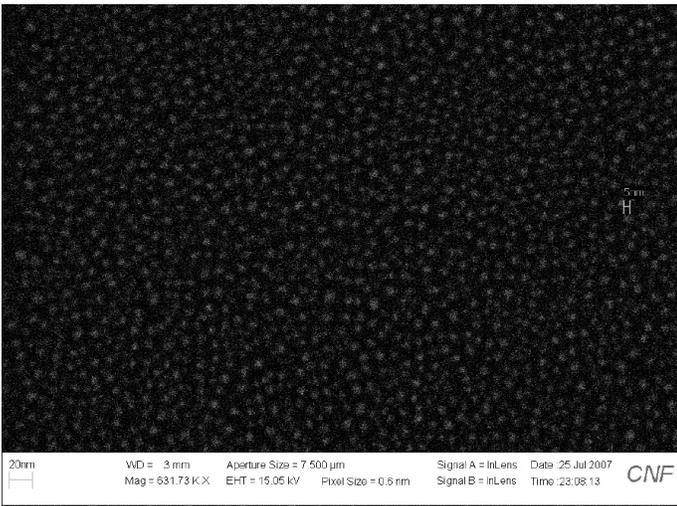


Figure 2: SEM image of Au nanocrystals on SiO₂ surface.

Results and Conclusions

After 1.2 nm Au deposition on the tunneling oxide, the minimization of local energy at the Au/SiO₂ interface reshaped the thin film into discontinuous and spherical Au nanocrystals, as shown in the scanning electron microscopy (SEM) image (Figure 2). The diameter of each nanocrystal was about 5 nm.

To measure the memory window, we performed multiple capacitance-voltage (CV) sweeps from inversion to accumulation and back again. The observed hysteresis of the CV curves indicated memory effect through charge storage. Figure 3 represents the memory window obtained from a double-layer C₆₀ and Au floating gate.

We then measured the retention time for each configuration, and as demonstrated in Table I, we compared the values to obtain the optimal configuration for a memory device. Based on the results

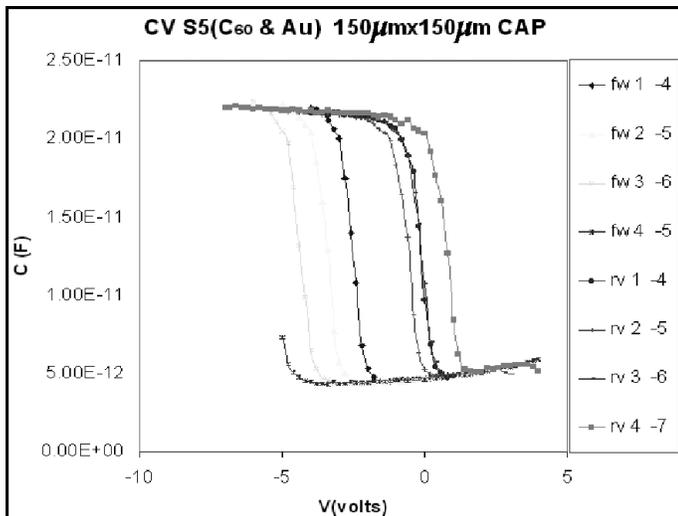


Figure 3: Memory window for a double-layer C₆₀/Au NC memory.

we concluded that double-layer nanocrystal floating gates with C₆₀ as the bottom layer and Au as the upper layer would give the best memory device in terms of memory window and retention time. Since mono-dispersed C₆₀ has a diameter of about 1 nm, it performed best in blocking back-tunneling of electrons and holes from the upper layer to the channel in this project.

Future Work

More tests will be performed to study the behavior of these memory devices under short-pulse program/erase operations. The retention time will be further improved by incorporating high-κ material as the tunneling dielectric. Finally, the control oxide thickness can be scaled to eliminate short channel effects in the sub-micron gate-length devices.

Acknowledgements

This work has been supported by Cornell NanoScale Science & Technology Facility, National Nanotechnology Infrastructure Network Research Experience for Undergraduates Program, and the National Science Foundation. I would like to thank Edwin Kan, Tuo-Hung Hou, Daniel Ruebusch, CNF staff, and the rest of the Kan group for their guidance and support.

References

- [1] Hanafi, H.I.; Tiwari, S.; Khan, I. , “Fast and long retention-time nano-crystal memory,” Electron Devices, IEEE Transactions on Volume 43, Issue 9, Sept. 1996 Page(s):1553-1558.
- [2] Hou, Tuo-Hung; Ganguly, Udayan; Kan, Edwin C, “Programmable molecular orbital states of C60 from integrated circuits,” Appl. Phys. Lett., vol. 89, 253113 (2006).

Run#1 Tunneling oxide = 2.41 nm

Run#2 Tunneling oxide = 2.72nm

Voltage sweep (-2 to 5 V)

Sample #	Nanocrystal Layer/s	Run#1		Run#2	
		Mem-win(V)	Reten t (S)	Mem-win	Reten t
S1	Control	0	0	0	0
S2	Au	5.8	400	5.4	NA
S3	Au & Au	NA	NA	NA	NA
S4	C60	2.4	240	2.8	250
S5	C60 & Au	3	1100	4.1	5100
S6	C60 & Au	4.7	500	4.4	5000

Table I: Summary of memory window and retention time.