

**INVESTIGATION OF RU THIN FILMS
PREPARED BY CHEMICAL VAPOR DEPOSITION
AS BOTTOM ELECTRODES FOR MEMORY
APPLICATIONS**

Sang Yeol Kang, Ha Jin Lim, Cheol Seong Hwang and
Hyeong Joon Kim
Seoul National University, School of Materials Science
and Engineering
San #56-1, Shillim-dong, Kwanak-ku, Seoul, 151-742,
Korea

The requirement for the chemical vapor deposition (CVD) of metal electrodes for the Ta₂O₅ and (Ba,Sr)TiO₃ [BST] capacitors for dynamic random access memory (DRAM) devices becomes more critical as the storage node height increases more than 0.3 μm with a 0.1-μm spacing between them[1]. One candidate for use as an electrode is Ru, because it has excellent characteristics, such as low resistivity, good susceptibility to dry etching, and a conductive oxide phase of RuO₂[2]. Ru bottom electrode is located between TiN diffusion barrier and high-k dielectric layer. Post-annealing on the capacitor induces to roughen Ru surface and to form TiOx layer at the Ru/TiN interface and the changes affect on the electrical property of the capacitor. Therefore, the stability of the two interfaces, Ru/TiN and high-k layer/Ru, should be necessarily investigated. In this study, we deposited Ru thin films using Cyclopentadienylpropylcyclopentadienyl ruthenium (II) [RuCp(i-PrCp)], and investigated the stability of Ru/TiN and high-k layer/Ru during annealing. Figure 1 shows a schematic diagram of the MOCVD system which consists of a vertical warm wall reactor, a resistive substrate heater 6" in diameter, the bubbler system and the DLI system. Ar gas was used as a carrier gas as well as a diluent gas, and O₂ gas was introduced into the reactor in order to eliminate carbon incorporation into the film and also to enhance the decomposition of the metalorganic precursors. To deposit Ru-Ti alloy films, Ti(O-iPr)₄ was injected to the reactor through the DLI system during the delivery of RuCp(i-PrCp) by the bubbler system

Figure 2 shows variation of electrical resistivity of Ru-Ti film(30nm). At the high Ti source flow rate, the film seems to be an insulator due to networking of TiO₂ phases in the film. Figure 3 shows variation of surface roughness of Ru(30nm) films deposited with various Ti source flow rate and their N₂-annealed (at 700°C) films. In Figure 3, we can find out the addition of Ti prevents roughening of the films' surface below 0.15ccm of Ti source addition. It is believed that at the small amount of Ti content, TiOx phases distributed at the grain boundaries of Ru film, suppress the grain growth of the Ru grains during annealing. Figure 4 (a) and (b) show cross-sectional TEM images of the as-deposited and post-annealed under H₂ atmosphere at 400 °C followed by a N₂ atmosphere at 600 °C, respectively. The slightly light contrast effect on the TiN surface is due to the initially high oxygen concentration. It can be confirmed that post-annealing does not induce any structural change at the interface

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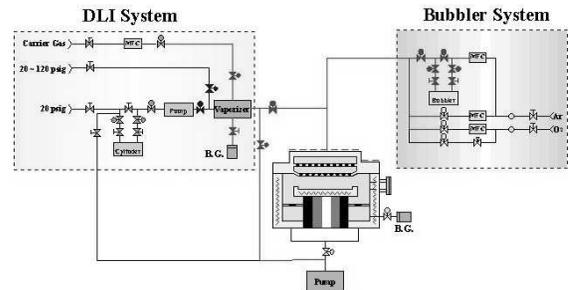


Figure 1. Schematic diagram of MOCVD system

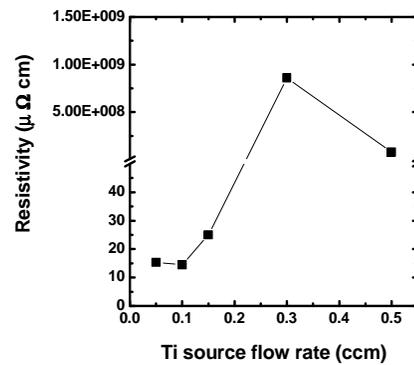


Figure 2. Variation of electrical resistivities of Ru-Ti films deposited with various Ti source flow rate.

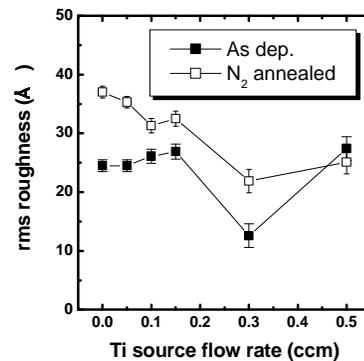


Figure 3. Variation of surface roughness of Ru-Ti films deposited with various Ti source flow rate

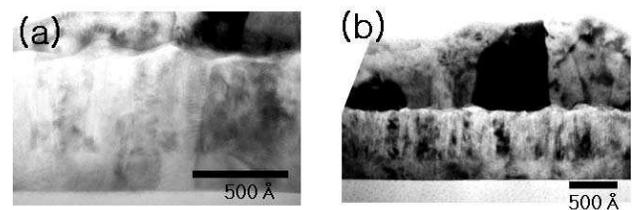


Figure 4. The cross-sectional TEM images of the Ru films deposited (a) at 375 °C with 50sccm of oxygen flow rate on TiN/TiO₂/SiO₂/Si substrate and (b) annealed under H₂ atmosphere at 400 °C and then N₂ atmosphere at 600 °C