

MOCVD GROWTH OF Pr₂O₃ HIGH-K GATE DIELECTRIC FOR SILICON: SYNTHESIS AND STRUCTURAL INVESTIGATION

Raffaella Lo Nigro,^a Roberta G. Toro,^b Graziella Malandrino,^b Vito Raineri,^a and Ignazio L. Fragalà^b

^aIMM, sezione di Catania, CNR, Stradale Primosole n 50, 95121 Catania, Italy. Tel:++39-095-591912, Fax: ++39-095-7139154.

E-mail: raffaella.lonigro@imm.cnr.it

^bDipartimento di Scienze Chimiche, Università di Catania, and INSTM, UdR Catania, Viale A. Doria 6, 95125 Catania, Italy. Tel.: ++39-095-7385055, Fax: ++39-095-580138

E-mails: gmalandrino@dipchi.unict.it, lfragala@dipchi.unict.it

Over the past two decades, the growth of oxide films on silicon has been investigated to fabricate highly integrated circuits. The aggressive scaling down of complementary metal-oxide semiconductor technology calls for identifying high K dielectrics to replace SiO₂ gate oxide, whose thickness is now approaching the quantum tunneling limit (about 10-15 Å).¹ One obvious way to reduce tunneling, while maintaining the required levels of gate oxide capacitance, is to substitute the SiO₂ gate oxide with thicker alternative materials having higher values of dielectric constant.² In recent years, Pr₂O₃³ has received much attention because of the possible use as alternative gate dielectrics in CMOS devices. Physical vapour deposition methods have been successfully used to grow Pr₂O₃ films on silicon substrates,⁴ while, to date, there are no studies on chemical vapour deposition methods.

We report on the results of a recent study on the deposition of praseodymium oxides thin films on silicon substrates by metal-organic chemical vapor deposition (MOCVD). Two different Pr(III) β-diketonate precursors have been investigated as candidates for Pr metal source and the deposition conditions have been carefully selected because of a large variety of possible PrO_{2-x} (x= 0-0.5) phases. Praseodymium oxide films have been deposited in an hot-wall MOCVD reactor at 750°C deposition temperature. A new praseodymium precursor has been synthesised, Pr(hfa)₃diglyme (H-hfa=1,1,1,5,5,5-hexafluoro-2,4-pentadione, diglyme= CH₃O(CH₂CH₂O)₂CH₃) and studied as praseodymium source. The fluorinated precursor yields an oxyfluoride phase, PrOF (figure 1), even under a water saturated oxygen flow. The Pr(tmhd)₃ (H-tmhd= 2,2,6,6-tetramethyl-3,5-heptandione) precursor has been investigated as praseodymium source as well, and depending on the oxygen partial pressure, different praseodymium oxides have been obtained. PrO₂ films (figure 2) have been formed using P_{O₂}=2 torr, while Pr₆O₁₁ and Pr₂O₃ films (figure 3) have been deposited using P_{O₂}=0.7 and 10⁻⁵ torr, respectively. In particular, the θ-2θ scan of Pr₂O₃ films deposited on Si (100) substrate, shows the formation of the Pr₂O₃ hexagonal random phase. Their complete structural and morphological characterization has been

carried out by X-ray diffraction (XRD), transmission electron microscopy (TEM), and atomic force microscopy (AFM). Finally, preliminary electrical measurements point to MOCVD as a reliable growth technique to obtain good quality Pr₂O₃ films.

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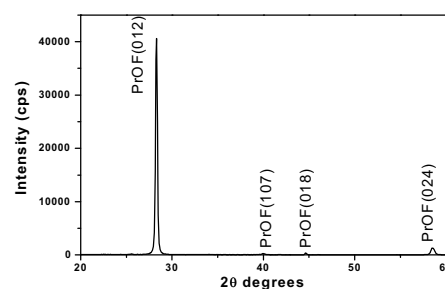
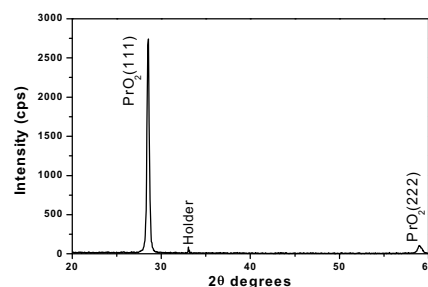


Figure 1. XRD spectrum of a PrOF film deposited on Si(100) substrate at 750°C using the Pr(hfa)₃diglyme



precursor.

Figure 2. XRD spectrum of a PrO₂ film deposited on Si(100) substrate at 750°C using the Pr(tmhd)₃ precursor at P_{O₂}= 2 torr.

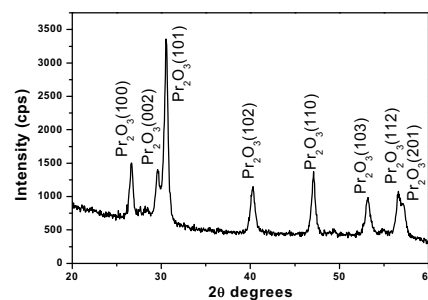


Figure 3. XRD spectrum of a Pr₂O₃ film deposited on Si(100) substrate at 750°C using the Pr(tmhd)₃ precursor at P_{O₂}= 10⁻⁵ torr.