Direct (Seedless) Cu Deposition on W: Implications for ULSI Processing

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Direct electrodeposition of Cu onto W without the use of a Cu seed layer is reported. The seedless electrodeposition of Cu on W or other reactive barrier metals is of considerable importance in ULSI processing, and has broader implications for control of metal nucleation and growth at the liquid/solid interface. Depositions were carried out in 0.05 M CuSO₄ in H₂SO₄ supporting electrolyte (deaerated) at pH 1 at cathodic potentials (< -0.2 V vs Ag/AgCl): conditions at which tungsten oxides are thermodynamically unstable with respect to clean W metal[1]. CV data (fig. 1) indicate a Cu deposition peak at ~ -0.24 mV/Ag/AgCl. Pulsed data (fig. 2), indicate a nucleation phase followed by diffusioninduced decrease in the cathodic current. Deposited films are shiny, and pass the "scotch tape" test. Preliminary XPS and EDX data indicate that films are pure Cu, without observable S or other contamination.

Under UHV conditions, the presence of a monolayer of oxide will prevent Cu from growing conformally ("wetting") the copper surface and induce poor adhesion[2]. It is not yet understood whether Cu nucleation will be governed in an analogous fashion in the more complex electrodeposition environment. In addition, the effects of specifically adsorbed anions and electrochemical potential on nucleation behavior remain to be explored.

This talk will discuss electrochemical, SEM and XPS data for electrodeposition on W in comparison with corresponding data on other barrier surfaces. Results will be discussed in light of expected differences for Cu electrodepositon on reactive vs unreactive substrates in aqueous environments.

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References:

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Figure 1; CV data for Cu electrodeposition on clean tungsten Figure 1; CV data for Cu electrodeposition on clean tungsten





Fig. 2 W pulsed in CuSO4=0.05M/pH=1H2SO4 for 10s

