University of Utah Fundamentals of Microfabrication

Electro Chemical Deposition: Principles, Methods and Applications

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Electrochemical Processes Electro Deposition

- Introduction
- Experimental Setup
- Importance of Electro Deposition
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- Electroless and Displacement Depositions
- Equilibrium Electrode Potentials
- Factors Affecting the Electrodeposits
- Micro and Nano-scale Structures

Introduction

What is Electro deposition? Electro deposition is the process of coating a thin layer of one metal on top of a different metal to modify its surface properties. Done to achieve the desired electrical and corrosion resistance, reduce wear&friction, improve heat tolerance and for decoration. E.D. is a surface coating method that forms an adherent layer of one metal on another.

Electroplating Setup



Electroplating Applications

Automotive parts Printed circuitry and electrical contacts General engineering components Gold-Silver wares and Jewelry Musical Instruments and Trophies Soft metal gaskets Anti-seize bearings Decorative door, light & bathroom fittings Production of micro parts for MEMS

Electroforming

Electroforming is a process used for making metallic articles with tight dimensional tolerance



By depositing a metal into or on to a mold or mandrel, a free-standing metal object is made

Useful for the production of originals and making exact copies of originals

Need for Electroforming

- Accuracy of reproduction
- Production of foils and mesh-products
- Screen-printing cylinders
- Manufacture of complex shaped objects
- Manufacture of large or small products
- Manufacture of molds

Electroforming Applications

- Micro components and prostheses
- Complex wave guides
- Metal bellows
- Reflectors, Nose cones
- Heat exchangers, micro filters
- Decorative ware

When combined with lithography, electroforming is extremely useful for making micro parts, and overcomes the difficulty of traditional machining



Electrochemical Deposition

Understanding the relationships between microstructure and mechanical properties is one of the main goals of metallurgy and materials science. This allowed the tailoring of materials properties by varying the composition and the structure. The fascinating field of electro deposition allows one to tailor the surface properties of a material or to form the entire part by electroforming. To explore new materials for a variety of applications.

Faraday's Law

The amount of electrochemical reaction that occurs at an electrode is proportional to the quantity of electric charge (Q) passed through the cell.

The weight of a product of electrolysis is W, then W=ZQ (Z is the electrochemical equivalent) Since Q=It, it follows that W=ZIt

Production of one gram equivalent of a product at the electrode (Weq) in a cell needs 96487 Coulombs

Electrochemical Equivalence

The electrochemical equivalent of a metal Z(M) is the weight in grams produced or consumed by one coulomb of charge

Weq = A / n
 When Q = 1,
$$W_{(Q=1)} = Z$$

 Weq = 96487 Z
 Z = Weq / 96487 = Weq / F

 Z = A / nF
 W = ZQ = AQ / nF



Water Structure At The Interface

The presence of excess charge on metal causes ion redistribution and reorientation of water dipoles.



Requisites for Electroplating MEMS Applications

A conducting surface as a seed layer is required. Clean substrates, solutions, and containers, etc. Even low impurity levels would result in poor-quality films.

Good resist adhesion on the substrate is required. The plating solution must have easy access to recesses, and be compatible with the resist and substrate materials.

Uniform current distribution on the substrate surface.

Typical Composition of Nickel Baths

	Watts	Sulfamate
Material	Quantity	Quantity
Nickel Sulfate	250 g/L	
Nickel Sulfamate		350 g/L
Nickel Chloride	5 g/L	25 g/L
Boric Acid	25 g/L	80 g/L
Ferrous Sulfate	8 g/L	
Additives	3 g/L	15 ml/L

Influencing factors in Electro deposition

The morphology and composition of electrodeposits vary significantly, and depend on:

Current density

The nature of the anions or cations in the solution

Bath composition and temperature

Solution concentration

Power supply current waveform

The presence of impurities

Physical & chemical nature of substrate surface

Pulse and Pulse-Reverse Plating

In dc-plating, constant current is used, and the rate of arrival of metal ions depends on their diffusion coefficient (electrode-to-part spacing and agitation). In PC and PRC, a modulated current waveforms are used to get a better leveling of the deposit, and to minimize the porosity, contamination, etc. The morphology of some metal and alloy deposits were found to be superior to the dc-plated deposits. Complex current wave forms can be generated by using pulse rectifier: unipolar and bipolar pulses.

Characteristics of Unipolar Currents

To characterize a train of current pulses, three parameters need to be known

- The cathodic peak pulse current density (Jc)
- The cathodic pulse length (tc)
- The interval between pulses (tp)

The average current density Jav is measured,

and it is expressed for PC as: $J_{av} = (J_c \ x \ t_c)/(t_c + t_p)$ The duty cycle (T) which represents the portion of the time in each cycle when the current is ON.

It is defined by T = tc/(tc + tp); $J_{av} = J_c x T$

Characteristics of Bipolar Currents

To characterize a train of pulse-reverse current waveform, four parameters need to be known.

- The cathodic peak pulse current density (Jc)
- The cathodic pulse length (tc)
- The anodic pulse time (ta)
- The anodic current density (Ja)

 $J_{av} = (J_c x t_c + J_{ata}) / (t_c + t_a)$ Tprc = (J_c x t_c - J_{ata}) / (J_c t_c)

Effects of Pulsed Current on the Deposit Properties

PC affects the mechanism of electro-crystallization, which in turn, controls the physical and mechanical properties of the deposit.

Nucleation rate of the deposit ∝ current density. Use of high J_{av} pulses can produce deposits with reduced porosity, and finer grains through the desorption of impurities and renucleation of deposits with the formation of new, smaller crystal grains. Grain size depends on the duty cycle and the bath. Low C and N content, deposits with low stress, increased ductility and electrical conductivity.

Structure and Properties of Deposits

The crystalline structure formed as a result of E.D. depends on a competition between rates of new crystalline formation and existing crystal growth.

Several factors in plating process influence the crystal structure and grain size of the deposits.

Plating Bath Control Using Ampere -Time Instruments

Plating is done by following a quantity called amperage which indicates the total amount of current that has passed through a plating bath. A change in bath's resistance will change amperage. The amount of metal deposited depends on three factors:

Amperage, Plating time, Current efficiency of the bath -Ampere-time x C.E. = mg of metal deposited -Thickness (μ") = mg of metal x Factor plated area (in²) -Ampere-time x C.E. = Thickness Area factor

Current Efficiency

Current efficiency is the ratio between the actual amount of metal depositing (or dissolved) M_a to that calculated theoretically from Faraday's law M_t in %. C.E. = $(M_a / M_t) \ge 100$ C.E. indicates the faction of total current that generates desired products. The overall amount of chemical change produced by a given quantity of electricity can be determined. Cathode & anode efficiencies - In an ideal situation, cathode efficiency will be equal to anode efficiency.

Cathode Efficiency

The ratio of the weight of metal actually deposited to the weight that would have resulted if all the current had been used for depositing it is called the cathode efficiency.

Cathode efficiency in plating depends on:

Electrolyte or bath

Concentration of chemical component

pH and agitation

Current density

The current efficiency in Ni plating is close to 100%, and in Cr plating it is approximately 20%

Macro Throwing Power

The ability of a bath to produce deposits of more or less uniform thickness on cathodes having macroscopic irregularities is termed Macro-throwing power. Ni plating (also other acidic Cu and Zn baths) show poor throwing power. If the C.E. values are close to 100% at low and high current density values, then the macroscopic irregularities on the cathode will lead to non-uniform deposits.

Alkaline baths have better macro throwing power, since the metal ions are present as complex ions.

Uniform Electrodeposit Through The Use of Complex Baths

- In most complex baths, the deposition potentials are amenable to hydrogen evolution which competes with metal deposition such that C.E. falls as current density is increased. This results in a more uniform deposit on cathodic macro irregularities.
- When the ions are complexed, they encounter high concentration polarization (CP). If the CP is high, the micro throwing power is rather poor.
- The ability to produce a deposit over a surface including recesses is called covering power.

Strike Baths

- Strike bath is made by an extremely high current density for a very short time or in a specially formulated bath.
- In some cases, the required potential for metal deposition may not be reached in recesses and vias.
- Other processes such as hydrogen evolution or reduction of ions such as Fe3+ to Fe2+ may occur.
 In such cases, a preliminary strike bath deposit may be the solution.

Electroless Deposition



Advantages of Electroless Deposition

Mixed Potential Theory

No electrical contact is needed It is possible to plate both conductive and

insulating surfaces, provided the surfaces are first sensitized.

It is readily adaptable for three-dimensional coverage.

No field lines are present, and this enhances deposit uniformity

Metallizing Nonconductors

Certain parts or components whose functions are fully utilized only when the properties of both a metal and non-metal are combined.

- Generally a part is made of plastic or ceramic, and the metal added to its significant surfaces to impart specific metallic properties.
- 1. For electrical conductivity, as in PCB.
- 2. For metallic appearance, as in the buttons, door knobs, wheels in toys, etc.
- 3. For strength, as in woman's high heel shoes.

The catalytic reaction must be conducted in such a way that a homogeneous reaction between Mz+ and RED in the bulk of the solution is suppressed.

The equilibrium potential of the RED (Eeq, Red) must be more negative than that of the metal electrode (Eeq,M), so that the reducing agent can function as electron donor and Metal ion as electron acceptor.

Developing Good Adhesion Metal Deposition on Polymers

For metallizing nonconductors, their surfaces need to be mechanically roughened, or etched, or made hydrophilic. Nonconductors need pre-conditioning of their surfaces.

Polymers: ABS, Polyimides, Polysulfones, etc.

Cr₂O₃/H₂SO₄ (0.5:1.0); Neutralize in NaHSO₃ Fluorocarbons: Na metal in anhydrous NH₃ or THF Need to be treated further in sensitizing, nucleating (catalytic), accelerator solutions prior to plating in electroless solution.

Hydrogen Embrittlement

Hydrogen embrittlement is a generic term used to describe a variety of fracture phenomena having a common relationship to the presence of hydrogen in the metal as a solute or in the atmosphere as a gas. In general any process producing atomic H at a metal surface will induce considerable hydrogen absorption in that metal (ED, ELD, Acid Pickling, E-cleaning). A large fraction combines to form bubbles of gaseous or molecular hydrogen which is not soluble in metals.

Mechanism of Hydrogen Embrittlement

The mechanism of may change depending on the source of H and the nature of applied stress.

- 1. The non-hydride forming systems such as Fe&Ni alloys fail because H-decreases the atomic bonding. The fracture seems to be associated with H-induced plasticity in the vicinity of the crack tip.
- 2. Metals which form stable hydrides (Nb, Zr, or Ti) appear to fracture by a stress-induced hydride formation and cleavage mechanism.
- 3. Adsorption-decreased surface energy and high-pressure hydrogen gas bubble formations.

Effects of Hydrogen Embritllement

Corrosion reactions can generate H_2 , and choosing proper coating to prevent corrosion is important. Permeation of H through various protective coatings is considered to show the effectiveness of various barrier layers on minimizing the H egress. Problems as a result of H embrittlement/hydriding:

- Leakage from gas filled pressure vessels
- Breakage of aircraft components
- Blisters or fisheyes in Cu, Al, and Steel components
- Reductions in mechanical propeties of materials

Leveling of Deposit

Leveling is the progressive reduction of the surface roughness during deposition.

S.R. may be due to coarse mechanical polishing. Cathodic leveling results in a smooth deposit or a deposit of reduced roughness. Leveling occurs when metal is deposited in recessed areas than on peaks. Leveling is important in metal finishing industry and in electronic industry for deposition through masks. In electroforming of MEMS devices leveling means uniform deposition in micro profiles.

Types of Leveling

Leveling can be achieved in solutions either in the absence or presence of leveling agents.

There are two types of leveling:

- 1. Geometric leveling in the absence of additives.
- 2. True or electrochemical leveling in the presence of additives.