Abstracts of Papers Presented at

The 25th International Field Emission Symposium

July 17-22, 1978

Albuquerque, New Mexico

Local Organizing Committee:	Steering Committee:
J. A. Panitz, Chairman J. L. Fowler J. E. Houston G. L. Kellogg G. L. Kuswa F. L. Vook R. J. Walko	J. H. Block Fritz Haber Institute, Berlin (West) S. S. Brenner U. S. Steel Corp., Monroeville, Pennsylvania M. Drechsler University of Marseille, Marseille, France
	Gert Ehrlich University of Illinois, Urbana, Illinois
	R. Gomer University of Chicago, Chicago, Illinois
Hosted by:	A. J. Melmed (Symposium Secretary) National Bureau of Standards, Washington, D.C.
Sandia Laboratories Albuquerque, New Mexico	M. J. Southon Cambridge University, Cambridge, England

Abstracts of the contributions in the various topics may be found grouped according to the indicated pages:

	Page
Biological Imaging	365
Surface Adsorption	366
E. W. Müller Memorial Medal Competition	368
Field Emission and Ion Sources	372
Surface Chemistry	374
Field Desorption Mass Spectrometry	375
Field Emission	377
Mass Transport and Electrical Breakdown	378
Metallurgy	37 9
Atom-Probe	382
Emission Spectroscopies	386
Field-electron and Ion Emission	387
Controlled Thermonuclear Fusion and Ion Implantation	389

LOW-FIELD DESORPTION IMAGING OF PROTEINS*

J. A. Panitz⁺, Ivar Giaever**

†Sandia Laboratories^{††}, Albuquerque, New Mexico; ***General Electric Re*search and Development Center, Schenectady, New York 12301

A novel low-field (< 1 V/ $^{\circ}$) desorption technique is described for imaging the contour of protein molecules placed on the apex of field-emitter tips. Image contrast relies on a distinct and measurable difference in the desorption field of a species of low ionization potential physisorbed on the molecule, and on the adjacent tip surface below 80 K. Preliminary results are described, using benzene to image Bovine Serum Albumin (BSA) and Immuno-Gamma-Globulin (IgG). Although the imaging procedure appears to be feasible, extreme difficulty in reproduceably placing the proteins on the tip apex, in the desired concentration, has been encountered. These difficulties together with future implications of the technique will also be discussed.

*This article sponsored by the U.S. Department of Energy, Office of Basic Energy Sciences, under Contract AT(29-1)789.

++A U. S. Department of Energy Facility.

SURFACE ADSORPTION

SINGLE ATOM SELF DIFFUSION ON NICKEL*

Raymond T. Tung⁺, William R. Graham⁺⁺

[†]Department of Physics, ^{††}Department of Metallurgy and Materials Science, Laboratory for Research on the Structure of Matter, University of Pennsylvania, Philadelphia, Pennsylvania 19104

Past FIM studies of surface diffusion have been limited to 5d and 4d transition metal substrates. The importance of extending this powerful technique for the study of the atomic mechanisms of surface interactions to the 3d transition series metals is clearly evident. In this paper we present the results from self diffusion studies on several of the low index planes of nickel. We also include some discussion of the specimen preparation techniques which are required to produce nickel surfaces of sufficient perfection for surface diffusion studies.

*Work supported by NSF Grant DMR-75-23254 and by NSF-MRL Grant 76-00678.

BINDING SITES AND BINDING STATES OF $\ensuremath{\mathtt{Sn}}$ on $\ensuremath{\mathtt{W}}$

O. Nishikawa

Department of Materials Science and Engineering, Graduate School of Science and Engineering, Tokyo Institute of Technology, O-okayama, Meguro-ku, Tokyo 152 Japan

The binding sites of tin atoms on a tungsten surface are studied by superposing the image of a pseudomorphic tin layer upon the image of a substrate tungsten surface. Superposed micrographs indicate that the lateral locations of the tin atoms correspond to the locations of unoccupied tungsten lattice sites above the surface.

Variations in field emission current with increasing evaporation field are also studied for the (011), (111), (112) and (114) planes. Evaporation of adatoms can be noticed by a sharp increase or decrease in the emission current at the field strength corresponding to the binding state of the adatoms. The binding states of the tin atoms on these planes are discussed.

FIELD ION MICROSCOPY OF GALLIUM ON TUNGSTEN*

Robert J. Culbertson, T. Sakurai, T. T. Tsong

Physics Department, Pennsylvania State University, University Park, Pennsylvania 16802

The field evaporation of Ga from a W emitter was investigated using a magnetic sector atom probe field ion microscope. A continuous supply of Ga was provided by adjusting both tip temperature and electric field. The Ga coating of a W tip was achieved by dipping the tip in liquid Ga at 325 K in air. A clean W and cap was obtained by fieldevaporating the tip at 78 K. By raising the temperature to 300 K Ga atoms diffused from the shank of the tip and field-evaporated from the tip apex at fields $(2V/\Re)$ well below the evaporation field of W (5.7 V/\Re). The rate of Ga evaporation could be controlled over a wide range by varying either the