An Auger electron spectroscopic investigation of the electron states of copper alloys

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Abstract. LVV Auger transitions of copper alloys show a feature due to a band-like states, the shape and intensity of which depend on the composition. The energy separation between this feature and the LMM peak increases progressively with Cu concentration.

Keywords. Auger transitions; electron states; copper alloy.

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1. Introduction

Unlike x-ray photoelectron spectroscopy (XPS), Auger electron spectroscopy (AES) has not been employed extensively to investigate electron states of metal alloys. AES should be specially useful in examining perturbations of the d-bands of transition metal alloys and to observe features due to band-like density of d-states. We have investigated several alloys of copper, both crystalline and glassy, and have obtained encouraging results.

2. Experimental

X-ray photoelectron spectra and x-ray induced Auger electron spectra were recorded using an ESCA 3 Mark II spectrometer (VG Scientific Ltd., UK), fitted with a sample preparation chamber, a gas handling manifold and Ar+ etching facility (Rao et al 1979). The Au (4f7/2) signal at 83.7 eV was used for energy calibration of the spectrometer.

Cu-Zr metaglasses were prepared by rapid quenching from the melt (Sen et al 1984). Cu-Al and Cu-Mn alloys were prepared by induction melting under vacuum. Electron beam melting was employed for the Cu-Pd system and the remaining alloys were prepared in an induction furnace in argon atmosphere.

All the samples were first cleaned and polished mechanically before introduction into the vacuum (2 x 10^-10 Torr) of the sample preparation chamber of the spectrometer. Subsequently, the samples were cleaned by scraping them in situ with a stainless steel file. Surface compositions of samples were determined by XPS. All the studies were carried out at room temperature (298K).
3. Results and discussion

In figure 1, we show the L_{3}VV region of Cu in Cu metal and glassy Cu_{50}Zr_{50}. The L_{3}M_{45}M_{45} Auger structure appearing between 912 and 923 eV, clearly shows five distinct features (marked A, B, C, D and E) similar to those noticed by earlier workers (Antonides et al 1977). These features arise from quasi-atomic two-hole excitations in the d-level and double ionization processes. We have decomposed these multiplets in terms of five gaussian peaks by a least-squared-error procedure after subtracting the background. This analysis shows that the central intense peak (C) contributes 45% of the total intensity in Cu metal and 42–44% in the other alloys studied, with the exception of the two metglasses, Cu_{50}Zr_{50} and Cu_{30}Zr_{70}, where the contribution of peak C is only 30–32%. The intensity of C increases up to 48% on depositing potassium on Cu metal. Such variations in the relative intensities of the component peaks suggest that the L_{3}M_{45}M_{45} transition of Cu in these alloys involves band-like states. Besides the L_{3}M_{45}M_{45} structure, figure 1 also shows a broad feature (F) between 924 and 930 eV. We assign this feature to the band part of the L_{3}M_{45}M_{45} spectrum. This assignment finds support from the studies of Weightman and Andrews (1979) as well as of Sawatzky (1977).

We have investigated the F peak around 927 eV, assigned to band-like density of d-states convoluted with itself, in several crystalline alloys of Cu with Mn and Ni as well as in the metglasses with Zr and find that its shape and intensity change from alloy to alloy (see figure 2a). In Cu-Zr alloys, the feature becomes narrower with decreasing Cu concentration, suggesting an increased atomic impurity-like behaviour of Cu in the matrix of Zr. These changes are likely to arise from a narrowing of the Cu(3d) level due

![Figure 1. X-ray induced L_{3}VV (L_{3}M_{45}M_{45}) Auger spectra of Cu in Cu metal and Cu_{50}Zr_{50} metglass. The high energy side of the signal has been expanded to show the band part of L_{3}VV transition.](image-url)
to increase in 3d-4d overlap. In Cu-Ni alloys however, the intensity of the feature decreases with decreasing Cu concentration with no evidence of s-band localization. We have found similar changes in Cu-Al and Cu-Mn alloys.

In figure 1, we also see a dip (marked G) in intensity around 931 eV appearing between the rising parts of the band-derived L_3M_45M_45 feature (F) and the smoothly decaying part of the L_3M_45M_45 transition. We may point out here that G would have appeared at L_1 binding energy, had there been no decaying tail of the L_2M_45M_45 transition. The presence of the latter makes G appear as a dip and pushes it to lower kinetic energy, the magnitude of the energy shift depending on the intensity of the band part of the L_3M_45M_45 transition. The energy difference between G and the most intense peak C, $E_G-E_C$, decreases progressively with the decreasing Cu concentration irrespective of the alloy systems (figure 2b). It is noteworthy that the Cu-Zr system exhibits a more pronounced decrease in $E_G-E_C$ compared to other alloys. This is in accord with the fact that the Cu(3d) signal in xps shifts away from $E_F$ when alloyed with Zr.
In Cu-Pd alloys, we find that the Pd($M_{45}N_{45}N_{45}$) transition energy shifts to lower kinetic energy by $\sim 1.5$ eV in going from Pd to Cu$_{92}$Pd$_8$ indicating an increase in the $U_{\text{eff}}$ value for dilute Pd alloys. This observation suggests a narrowing of the Pd(4d) level on dilution.

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References