

Available online at www.sciencedirect.com

SciVerse ScienceDirect

Scripta Materialia 66 (2012) 741-744



www.elsevier.com/locate/scriptamat

The effect of a concentration gradient on interfacial reactions in microbumps of Ni/SnAg/Cu during liquid-state soldering

Y.S. Huang, a H.Y. Hsiao, a Chih Chena, and K.N. Tu^b

^aDepartment of Materials Science and Engineering, National Chiao Tung University, Hsinchu 30010, Taiwan, ROC ^bDepartment of Materials Science and Engineering, University of California at Los Angeles, Los Angeles, CA 90095-1595, USA

> Received 1 November 2011; revised 21 January 2012; accepted 23 January 2012 Available online 28 January 2012

We report new findings in Ni/SnAg–solder/Cu microbumps having a reduced solder thickness in the range of 40– $10~\mu m$; the growth rate of the intermetallic compounds depends strongly on the solder thickness. In the Ni/40 μm solder/Cu samples, the compound which grew on the Ni side grew slightly faster than that on the Cu side. However, the growth reverses as the solder thickness decreases below 20 μm due to the change in Cu and Ni concentration gradients in the solder. © 2012 Acta Materialia Inc. Published by Elsevier Ltd. All rights reserved.

Keywords: Diffusion; Intermetallic compounds; Soldering; Copper; Nickel

The microelectronics industry is currently making advances in three-dimensional integrated circuit (3-D IC) technology. However, the packaging of 3-D IC is problematic because of metallurgical reactions in microbumps [1]. In the 3-D IC packaging, the diameter of the solder joints can be reduced to 10 µm; this is known as microbumps [2]. The volume of such joints is 1000 times smaller than that of solder joints in flip-chip technology, which have a diameter of 100 µm. At the same time, the contact area is 100 times smaller, but the diffusion or reaction distance is only 10 times less. The reduced size of the microbump has introduced new and challenging yield and reliability concerns, because the temperature and time of processing remains unchanged. A critical concern is the rapid saturation of the concentration of Cu and Ni in the solder due to dissolution of Cu and Ni from under-bump metallization (UBM), and also the effect of dissolution on the rate of ternary (Cu, Ni)₆Sn₅ intermetallic compound (IMC) formation in the solder joint [3–8].

In the literature, several reports have addressed the metallurgical reactions in the Ni/Pb-free solder/Cu systems in flip-chip technology. Wang and Liu [9] and Teng and Liu [10] reported that the IMCs on the Ni side grew faster than those on the Cu side for a solder joint of thickness of 150 μ m. Shao et al. [11] also reported that the Cu atoms diffused to the Ni side to form ternary

Cu–Ni–Sn IMCs in a flip-chip solder joint. This is because the flux of Ni atoms to the Cu end is smaller than that of Cu atoms to the Ni end because the solubility of Cu in the solder is higher. Recently, Li et al. [12] studied interfacial reactions in the Cu/10 µm Sn/Cu system and found that the Cu–Sn IMCs grew from both sides at approximately the same rate. The entire Sn layer was transformed into Cu–Sn IMCs after reflowing at 340 °C for 40 min. The key finding, as mentioned above, is that the Cu and Ni atoms in the solder can significantly affect the growth of the interfacial IMCs on both sides of the joint. Hence we expect that the effect will be even more serious in the reduced size of microbumps. However, no reports have been published on this subject yet.

In this study, we fabricated an Ni/SnAg solder/Cu structures with solder thicknesses of 40, 20 and 10 µm. The samples were reflowed at 260 °C for various times. We found that the growth rate of the IMC increases on the Cu side and decreases on the Ni side with decreasing solder thickness. Only the ternary (Cu, Ni)₆Sn₅ IMC was found. Compositional analysis of the solder and of the IMCs was performed in order to propose a plausible mechanism for this interesting observation.

Figure 1(a) shows a schematic drawing of the sample structure and the bonding setup. The structure consists of two parts. The lower part is fabricated using Si wafer as the substrate, 20 μ m Cu as the UBM and of 9, 19 or 60 μ m Sn2.3Ag as the solder cap. The upper part is fabricated using Si wafer as the substrate, 3 μ m electroplated

^{*} Corresponding author. E-mail: chih@cc.nctu.edu.tw

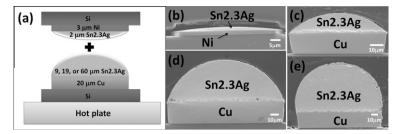


Figure 1. (a) Schematic diagram of the sample cross-sectional structure and bonding process of the Ni/SnAg/Cu microbumps. Cross-sectional SEM images for (b) 10 μm thick, (c) 19 μm thick and (d) 60 μm thick SnAg solder on 20 μm electroplated Cu UBM. (e) Cross-sectional SEM images for 2 μm thick SnAg solder on 3 μm electroplated Ni UBM.

Ni as the UBM, and 2 μm Sn2.3Ag as the solder cap. Figure 1(b) shows the cross-sectional scanning electron microscopy (SEM) image of the upper part of the sample. Figure 1(c)–(e) shows the cross-sectional SEM image of the lower part of the sample. Both parts of the samples were reflowed (the first reflow) at 260 °C for 1 min to ensure the formation of the solder cap and its firm attachment to the Cu or Ni. The bonding and reflowing process shown in Figure 1(a) was then carried out. The two parts were aligned, placed in face-to-face contact and reflowed at 260 °C for 3 min (the second reflow). By joining the parts in Figure 1(b), (c) and (d) to the part in Figure 1(e), we fabricated the Ni/SnAg/Cu structures with three SnAg solder thicknesses, of 10, 20 and 40 µm, respectively. These three samples are denoted as Ni/10 µm solder/Cu, Ni/20 μm solder/Cu, and Ni/40 μm solder/Cu in this paper.

The as-bonded samples were further reflowed (the third reflow) at 260 °C for 0, 5, 10 and 20 min in order to investigate the growth of IMC on both the Cu/solder and Ni/solder interfaces. Cross-sectional samples were polished and observed using field-emission SEM. The compositions of solder and IMC in the joint were analyzed by field-emission electron probe X-ray microanalysis (EPMA), using a JXA-8800M microanalyzer with a resolution of 0.01 at.%.

In this paper, the reflow time represents the total reflow time of three reflows, i.e. it includes the 1 min reflow after the electroplating of the solder, the 3 min reflow during bonding and the additional reflow to grow the IMC. Thus the reflow times under investigation are actually 4, 9, 14 and 24 min.

The growth behavior of interfacial IMCs in the Ni/ 40 μm solder/Cu appears similar to that in the Ni/ 150 μm thick Sn/Cu structure reported in the literature [9]. Figure 2(a) and (b) shows cross-sectional SEM back-scattered images of the Ni/40 µm solder/Cu samples after reflows at 260 °C for 4 and 24 min, respectively. The compositional analysis by EPMA indicates that the IMCs grown on both the top and bottom sides are (Cu, Ni)₆Sn₅. Figure 2(a) shows the cross-sectional SEM image for the Ni/40 µm solder/Cu sample after the 4 min reflow, which is the as-bonded sample. The (Cu, Ni)₆Sn₅ on the Cu side is of the scallop-type, with an average thickness of about 3.48 µm, whereas that on the Ni side is needle-like, with an average thickness of about 1.99 μm. Figure 2(b) shows the microstructures of the Ni/40 µm solder/Cu sample after the 24 min reflow. The average thickness of the (Cu, Ni)₆Sn₅ on the

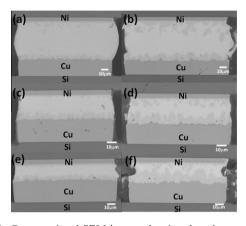


Figure 2. Cross-sectional SEM images showing the microstructure of the samples. (a) As-bonded with the 40 μ m thick solder. (b) Reflow for 24 min with the 40 μ m thick solder at 260 °C; (c) As-bonded with the 20 μ m thick solder. (d) Reflow for 24 min with the 20 μ m thick solder at 260 °C. (e) As-bonded with the 10 μ m thick solder. (f) Reflow for 24 min with the 10 μ m thick solder at 260 °C.

Cu side grew to $6.10 \,\mu\text{m}$. The average thickness of the (Cu, Ni)₆Sn₅ on the Ni side grew to $5.58 \,\mu\text{m}$. Some of the IMC on the Ni side spalled into the SnAg solder.

Figure 3 plots the relationship between the thickness of (Cu, Ni) $_6$ Sn $_5$ and the reflow time of this set of samples (Ni/40 µm solder/Cu), using solid triangular dots. It indicates that the thickness of the (Cu, Ni) $_6$ Sn $_5$ on the Cu side is slightly thicker than that on the Ni side. However, the slope or the growth rate appears faster for the IMC on the Ni side. The thickness of the IMCs on the Ni side increases from 1.99 to 5.58 µm after the 24 min reflow, whereas it increases from 3.84 to 6.10 µm on the Cu side.

The IMC growth rate changed in the samples with the reduced thickness of solder. Figure 2(c) and (d) shows cross-sectional SEM images of the Ni/20 μ m solder/Cu samples after reflow at 260 °C for 4 and 24 min, respectively. The compositional analysis indicates that the interfacial IMC on both sides are also (Cu, Ni)₆Sn₅. Figure 2(c) shows the microstructure of Ni/20 μ m solder/Cu sample after the 4 min reflow, which is the asbonded sample. The (Cu, Ni)₆Sn₅ on the Cu side is of the scallop-type, with an average thickness of 3.58 μ m, whereas that on the Ni side appears to be needle-like, with an average thickness of 1.57 μ m. Compared with the results in Figure 2(a), the thickness of the needle-like (Cu, Ni)₆Sn₅ on the Ni side is thinner than that in the

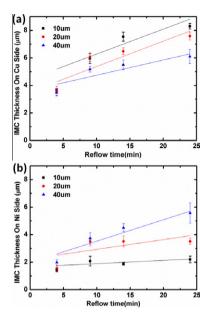


Figure 3. The measured $(Cu,Ni)_6Sn_5$ IMC (a) on the Cu side and (b) on the Ni side for the three sets of samples.

Ni/40 μ m solder/Cu sample after the 4 min reflow. As the reflow time increased, the IMC on the Cu side grew faster than that on the Ni side, as shown in the Figure 2(d). After the 24 min reflow, the average thickness of the (Cu, Ni)₆Sn₅ on the Cu side is 7.59 μ m, whereas it is only 3.51 μ m on the Ni side.

Figure 3 also plots the measured IMC thickness as a function of reflow time of this set of samples (Ni/20 μm solder/Cu), using solid circular dots. The IMC on the Cu side continued to grow thicker as the reflow time increased. However, the IMC on the Ni side did not grow obviously after the 9 min reflow, with the average thickness remaining at 3.51 μm for the three samples reflowed after 9, 14 and 24 min. The growth rate of the IMC on the Cu side becomes faster than that on the Ni side, which is the reverse of that in the Ni/40 μm solder/Cu sample.

It is more intriguing that the IMC on the Ni side is further inhibited as the solder thickness is reduced to 10 μm. Figure 2(e) shows the as-bonded Ni/10 μm solder/Cu sample. It is quite surprising that the IMC on the Ni side appears to be layer-type, with an average thickness of only 1.42 μm. The composition of the IMC remains (Cu, Ni)₆Sn₅ on both sides. After reflowing for 24 min, some of the IMC on the Cu side bridged the IMC on the Ni side, as shown in Figure 2(f). The IMC on the Cu side grew rapidly to 8.30 μm while that on the Ni side grew slowly, to only 2.20 μm.

Figure 3 also shows the measured IMC thickness as a function of reflow time for the set of Ni/10 µm solder/Cu samples, using solid square dots. The difference in IMC thickness between the Cu side and the Ni side in this set appears to be the largest among the three sets of samples in this study.

The data in Figure 3 indicate that the thickness of the $(Cu, Ni)_6Sn_5$ on the Cu side increases with decreasing solder thickness, and the trend becomes more obvious at a reflow time longer than 14 min. However, for the IMC on the Ni side, the average thickness decreases with

decreasing solder thickness, as shown in Figure 3(b). The results suggest that more Ni atoms may have diffused to the Cu side in the samples with the thinner solder because the growth of the IMC requires the supply of Ni. This point will be discussed later.

The EPMA analysis revealed that there are concentration gradients of Cu and Ni across the solder layers. The three locations analyzed inside the solder joints were: the solder near the Ni UBM; the middle of the solder; and the solder near the Cu UBM. All the analyses were performed at locations where no IMC is presented. The average value was obtained by averaging four data points around the three locations. In the solder of the Ni/40 μ m solder/Cu sample, the Ni concentration is 0.20 ± 0.09 , 0.11 ± 0.09 and 0.05 ± 0.05 wt.% at the three positions, respectively. The Cu concentration is 0.81 ± 0.07 , 0.57 ± 0.12 and 0.80 ± 0.19 wt.% at the three positions, respectively. The average concentration of Cu is 0.72 wt.% and that of Ni is 0.12 wt.%.

When the solder thickness was reduced to 20 μm , we found that the average concentrations of the Cu and Ni increased. The measured Ni concentration is $0.36\pm0.085,\ 0.26\pm0.04$ and 0.10 ± 0.07 wt.% at the three positions, respectively. The Cu concentration is measured to be $0.93\pm0.10,\ 0.81\pm0.21$ and 1.48 ± 0.21 wt.% at the three positions, respectively. The average Cu and Ni concentrations in this sample are 1.07 and 0.24 wt.%, respectively.

The measured Cu and Ni concentrations in the solder increase further as the solder thickness decreases to 10 μm . The measured Ni concentration is 1.27 ± 0.14 , 0.673 ± 0.06 and 0.47 ± 0.13 wt.% at the three positions, respectively, whereas the Cu concentration increases to $1.05\pm0.13,\ 1.20\pm0.16$ and 1.67 ± 0.21 wt.% at the three positions, respectively. The average Cu and Ni concentrations in this sample are 1.30 and 0.80 wt.%, respectively. Note that the values in the thin solder samples are higher than those of the thicker solder samples. The results also clearly indicate that there exists an Ni concentration gradient inside the solder, decreasing from the Ni side to the Cu side.

Figure 4(a) and (b) shows plots of the measured Cu and Ni concentration in the solder at locations, respectively, closer to the IMC near the Cu UBM and the Ni UBM as a function of solder thickness. By comparing Figure 4(a) and (b), we can see clearly a concentration gradient of Ni in the samples, especially in the 10 µm thick solder samples, which decreases from 1.27 wt.% near the Ni side to 0.47 wt.% near the Cu side. There could also be a concentration gradient of Cu in the sample, but, if so, it is much smaller. This is because of the formation of (Cu, Ni)₆Sn₅ on both the Cu and Ni UBMs. Since the IMC is an equilibrium compound with Cu, the Cu solubility near the IMC should be the same. Thus we expect a small or no concentration gradient of Cu in the solder. On the other hand, the ratio of Cu and Ni in the IMC may differ slightly between the IMC on the Cu side and that on the Ni side. This effect may influence the solubility slightly, but the effect on the Cu concentration gradient will be small.

Chen et al. [13] reported that the Cu concentration in the solder has a significant effect on the interfacial IMC formation between the Sn–Cu solder and the Ni UBM.

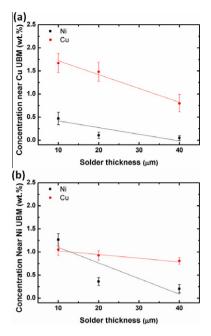


Figure 4. (a) The measured Cu and Ni concentrations in the solder close to the Cu UBM as a function of solder thickness. (b) The measured Cu and Ni concentration in the solder close to the Ni UBM as a function of solder thickness.

When the Cu concentration in the Sn-Cu solder increases to 0.6 wt.%, the stable ternary IMC changes from (Ni, Cu)₃Sn₄ to (Cu, Ni)₆Sn₅, the latter being more stable than the former. In the present study, most of the measured Cu concentrations in the solder are higher than 0.6 wt.% in all three sets of samples. Thus, the stable IMC is (Cu, Ni)₆Sn₅ on both the Cu and Ni sides. On the Ni side, no obvious layer of (Ni, Cu)₃Sn₄ could be observed. Note that the measured Cu concentration at the Cu end in the 10 µm thick sample is higher than the saturation solubility of Cu in SnAgCu solder at 260 °C, which is 1.54 wt.% [14]. The Cu concentration at this location is larger than the saturation solubility. This is most likely due to the effect of the Gibbs-Thomson potential, which increases the equilibrium concentration of solute around small precipitates, if we assume that there are many small ternary compound particles near the Cu UBM due to spalling or precipitation. On the other hand, the Ni saturation solubility in eutectic SnAg solders is 0.28 wt.% at 250 °C [6]. Most of the measured Ni concentrations are also larger than the solubility limit, suggesting that many Ni atoms have precipitated out as tiny Cu-Ni-Sn IMC in the solder.

The dissolved Ni atoms may affect the Cu saturation solubility in the solder on the Ni side. Wang and Liu [9] studied the interfacial reaction of Cu/150 µm Sn/Ni samples. They found that the IMC on the Ni side grew thicker than on the Cu side. They proposed that the saturation solubility of Cu may decrease from 1.1 to 0.6 wt.% when Ni atoms are present in the solder, resulting in a Cu concentration gradient across the Sn layer.

When the thickness of the solder is reduced, the trend of ternary IMC formation is reversed. We propose a mechanism for this as follows. When the thickness of solder is reduced to 10 µm in a microbump, the formation of IMC is different from that in flip-chip solder joints. The major influence comes from the effect of solder thickness on the solubility and concentration gradient of Cu and Ni in the solder. The reduced thickness tends to increase the solubility and leads to the formation of (Cu, Ni)₆Sn₅ on both sides of the solder joint. As we have mentioned above, since the IMC is an equilibrium compound with Cu, it will greatly reduce the concentration gradient of Cu across the solder joint, but not the Ni concentration gradient. The latter is the driving force of diffusion of Ni to the Cu side. It is natural to expect a faster growth of the IMC on the Cu side, provided the supply of Ni is unrestricted.

In summary, the interfacial reactions in the liquid state have been studied in Ni/solder/Cu microbumps with 10, 20 and 40 μm thick SnAg solders. The IMC growth rate depends significantly on the thickness of the solder. The IMC thickness on the Cu side increases with decreasing solder thickness with a fixed reflow time. However, the IMC thickness on the Ni side decreases as the solder thickness is reduced for a fixed reflow time. Compositional analysis by EPMA indicates that the Cu and Ni concentrations in the solder increase with decreasing solder thickness. Furthermore, there is a concentration gradient of Ni, but not Cu, across the microbumps, especially in the 10 μm thick solder samples. We propose that this is the mechanism that enhances the growth of the ternary IMC on the Cu side.

The financial support from the National Science Council, Taiwan, under contract NSC 98-2221-E-009-036-MY3, is acknowledged.

- [1] J.C. Lin, W.C. Chiou, K.F. Yang, H.B. Chang, Y.C. Lin, E.B. Liao, J.P. Hung, Y.L. Lin, P.H. Tsai, Y.C. Shih, T.J. Wu, W.J. Wu, F.W. Tsai, Y.H. Huang, T.Y. Wang, C.L. Yu, C.H. Chang, M.F. Chen, S.Y. Hou, C.H. Tung, S.O. Jeng, D.C.H. Yu, IEDM, IEEE International (2010) 2.1.1–2.1.4.
- [2] K.N. Tu, Microelectron. Reliab. 51 (2011) 517.
- [3] H.F. Hsu, S.W. Chen, Acta Mater. 52 (2004) 2541.
- [4] J. Shen, Y.C. Chan, S.Y. Liu, Acta Mater. 57 (2009) 5196.
- [5] M.S. Park, R. Arroyave, Acta Mater. 58 (2010) 4900.
- [6] C.P. Huang, C. Chen, C.Y. Liu, S.S. Lin, K.H. Chen, J. Mater. Res. 20 (10) (2005) 2772.
- [7] M. Kerr, N. Chawla, Acta Mater. 52 (2004) 4527.
- [8] M.O. Alam, Y.C. Chan, J. Appl. Phys. 98 (2005) 123527.
- [9] S.J. Wang, C.Y. Liu, Scripta Mater. 55 (2006) 347.
- [10] H.W. Tseng, C.Y. Liu, Mater. Lett. 62 (2008) 3887.
- [11] T.L. Shao, T.S. Chen, Y.M. Huang, C. Chen, J. Mater. Res. 19 (2004) 3654.
- [12] J.F. Li, P.A. Agyakwa, C.M. Johnson, Acta Mater. 59 (2011) 1198.
- [13] W.T. Chen, C.E. Ho, C.R. Kao, J. Mater. Res. 17 (2002) 263.
- [14] K. Zeng, K.N. Tu, Mater. Sci. Eng. R 38 (2002) 55.