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Analysis of Physiochemical Processes During Solder Aging

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Analysis of Physiochemical Processes During Solder Aging

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Abstract

A model describing the diffusion-controlled growth of multiple intermetallic layers and the displacement of the interfaces between layers was developed and implemented in a one-dimensional computer code based on the method-of-lines. The model can accommodate cases involving: (1) finite initial layer thicknesses, (2) rate-limiting interfacial reactions, (3) multiple and variable diffusion coefficients, and (4) finite material boundaries. Additionally, the effects of nucleation can be modeled empirically. A transformation of the spatial coordinate circumvented the need to remesh the growing and (or) shrinking layers. Results from the one-dimensional code were verified by comparing the numerical results with analytical solutions for simple systems involving two, three, and five layers. The computer code was applied to analysis of the intermetallic layer growth that occurred in solder aging experiments that were done with 100Sn and 63Sn-37Pb solders. The analyses indicated that intermetallic layer growth was consistent with a bulk diffusion mechanism involving Cu and (or) Sn and variable diffusion coefficients that should reflect some enhanced diffusion during early layer growth. The enhanced diffusion can be accounted for empirically using a variable diffusion coefficient that is a function of layer thickness.

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1.0 INTRODUCTION

The reliability of mechanical and electronic systems can be acutely dependent on the integrity of the soldered joints used in those assemblies. During manufacture and service, intermetallic layers form in reaction zones between the dissimilar materials forming the joints. The thermal and mechanical properties of the intermetallic layers can be substantially different from those of the initial bulk materials. The properties of the intermetallic layers can seriously affect wetting characteristics during manufacturing, and reliability during service in a variety of defense applications and commercial products in the electronics, automotive, aircraft, nuclear, and power industries.

Because thermal cycling during service can result in chemical and physical changes to the microstructure of the joint (for example, intermetallic layer growth), analytical models describing these changes can contribute to improved predictions of the long-term reliability of soldered joints. The objective of this work was to develop the numerical tools for analyzing the multiphase, multicomponent, solid-state diffusion-controlled growth of intermetallic layers in soldered joints during service conditions and aging experiments at elevated temperatures. The software provides a new analysis capability for materials researchers to examine intermetallic growth mechanisms and for design engineers to predict manufacturing defects and joint failure during service for a wide variety of defense and commercial applications involving both traditional and advanced materials. Although the work primarily addresses processes occurring during service conditions after joints are formed, the modeling capabilities developed are amenable to analysis of several physiochemical processes associated with formation of soldered joints.

The research approach consisted of two complementary tasks: (1) the development of mathematical models and numerical tools and (2) the acquisition of data for intermetallic layer growth in solder-aging experiments using copper substrates with tin and tin-lead solders. The experimental work was intended to provide data for determining relevant mechanisms, in addition to chemical diffusion, that should be included in the models and software, and to quantitatively describe mechanisms and rates in real solder systems of interest in both defense and commercial applications.

The formation and growth of intermetallic layers is shown schematically in Fig. 1. In general, contact between a metal substrate and molten solder results in a chemical reaction between the metal and constituents of the solder, and an intermetallic phase nucleates at the metal-solder interface. As the reaction continues, nuclei coalesce to form a continuous intermetallic layer. If the solder material is an alloy, such as 63Sn-37Pb, the reaction of one constituent (for example Sn) to form the intermetallic layer causes the composition of the solder to become richer in the non-reacting constituent (Pb). The non-reacting constituent may precipitate as a dispersed phase in the bulk solder or may form a second layer ahead of the intermetallic layer. Growth of the intermetallic layer is controlled by the interfacial reaction rate and the diffusion of reacting constituents in the layer. Growth of the intermetallic layer continues while the temperature of the system remains elevated, but becomes negligible as the material cools to ambient temperature. During service, solder joints may be heated due to thermal cycling of the assemblies in which the joints are used. Reaction between solder and substrate can occur, causing additional growth of the intermetallic layer. Furthermore, additional intermetallic phases can nucleate and form layers adjacent to the initial layer.

After an intermetallic layer forms, growth of that layer is described by the diffusion equations

Figure 1. Schematic diagram illustrating (a) initial solder and substrate, (b) intermetallic nucleation, (c) intermetallic layer growth and dispersed phase precipitation, and (d) growth of an additional intermetallic layer.

for the reacting constituents in the layer, the reaction rates at each interface bounding the layer, and the equations for the displacement of each interface bounding the layer. Mathematical problems of this type are often referred to as "Stefan Problems" and have been discussed by several authors. The preponderance of the literature addresses heat transfer problems. An introduction to moving boundary, or Stefan, problems is given in the books by Carslaw and Jaeger (1959), Jost (1960), and Crank (1975). A much more detailed discussion of the analytical theory and existence of solutions to Stefan problems is given by Friedman (1964) and by Rubenstein (1971). Analytical solutions also are discussed in Friedman (1959 and 1960), Miranker and Keller (1960), and Cannon, et al. (1967). Various numerical approaches have been developed and are summarized by Crank (1988). These include front-tracking, front-fixing, and enthalpy methods, which are primarily applied to one-dimensional, one- or two-phase heat transfer problems. The enthalpy method has been particularly useful in heat transfer problems, for example, Shamsunder and Sparrow (1975), Meyer (1973), and Meyer (1978). The latter also discusses some transformation techniques applicable to limited classes of problems. Unfortunately, in mass transfer problems, an analogy to the enthalpy method does not exist, because the latent heat does not have an analogy in mass transfer. The use of an "immobilizing" transformation for a two-phase solidification problem is discussed by Duda, et al. (1975), and an application of the method-of-lines to one-dimensional, single-phase problems is discussed in Meyer (1977). Additional applications in which Stefan problems occur are discussed in Ockendon and Hodgkins (1974) and Chadam and Rasmussen (1993a and 1993b).

A numerical model describing the diffusion-controlled growth of multiple intermetallic layers and the displacement of the interfaces between layers was developed and implemented in a one-dimensional computer code based on the method-of-lines. A transformation of the spatial coordinate circumvented the need to remesh the growing and (or) shrinking layers. The model can accommodate problems involving: (1) finite initial layer thicknesses, (2) rate-limiting interfacial reactions, (3) multiple and variable diffusion coefficients, and (4) finite material boundaries. Additionally, the effects of nucleation can be modeled empirically. Results from the one-dimensional code were verified by comparing the numerical results with analytical solutions for simple systems involving two, three, and five layers. The analytical solutions assumed zero initial thickness of the intermetallic layers, although the process being modeled begins with small but finite thickness of these layers. The initial thickness of the layers is represented in the computational solution.

The computer code was applied to the analysis of the intermetallic layer growth that occurred in solder aging experiments performed with 100Sn and 63Sn-37Pb (wt.%) solders. The Cu-63Sn-37Pb couples were aged for 1 to 400 days at temperatures of 70, 100, 135, and 170° C. The Cu-100Sn samples were aged similarly, except that heat treatments were also performed at 205° C. Two intermetallic layers, Cu₃Sn and Cu₆Sn₅, were observed for both alloys. Some differences between the thickness of the intermetallic layers that formed with 100Sn and 63Sn-37Pb solders were also observed. At higher temperatures, the total thickness of the layers was similar, but the ratio of the thicknesses of the Cu₃Sn to the Cu₆Sn₅ layer differed. The analyses indicated that intermetallic layer growth was consistent with a bulk diffusion mechanism involving Cu and (or) Sn and variable diffusion coefficients that probably should reflect some enhanced diffusion during early layer growth. Furthermore, the enhanced diffusion can be accounted for empirically using a variable diffusion coefficient that is a function of layer thickness. A satisfactory explanation for the difference between intermetallic layer growth that was observed with 100Sn and 63Sn-37Pb

solder was not established. With 63Sn-37Pb samples, diffusion in the Cu_6Sn_5 layer appeared to be faster than with 100Sn samples. The higher diffusion rates may have been due to trace amounts of Pb perturbing the layer crystal structure, but the presence of Pb could not be confirmed by electron microprobe analysis or Auger depth profiling.

In the text below, the experimental procedures used in the solder aging experiments are described, and the data for intermetallic layer growth in the Cu-100Sn and Cu-63Sn-37Pb systems are summarized. Then, the physiochemical mechanisms controlling intermetallic layer growth, the governing equations used in this work, and the model for intermetallic layer growth are described. Next, implementation of the model in a computer code, and verification of the numerical calculations are discussed. The model and code are then applied to the analysis of the solder aging experiments, and the results of that analysis are summarized. Finally, some opportunities for further model and code development are discussed, and some conclusions are offered.

2.0 EXPERIMENT

The experimental work focused primarily on aging the solder joint to provide data for variations in intermetallic layer thickness as a function of time and temperature. Experiments with copper substrates and 63Sn-37Pb solder were performed with aging times of 1 to 400 days and temperatures of 70, 100, 135, and 170° C. Experiments with copper substrates and 100 Sn solder were also performed with aging times of 1 to 400 days and temperatures of 70, 100, 135, 170, and 205° C. The procedures for these experiments and the results obtained are summarized below.

Additional work was done to develop techniques for depositing “marker layers” that could be used in Kirkendall-type experiments (Jost, 1960) to observe interface displacement, rather than intermetallic layer growth, which was measured in the solder-aging experiments. Development of the technique for depositing marker layers is summarized in Appendix A, and although a suitable technique was found, the effort required to develop the technique was substantial and precluded further pursuit of the Kirkendall-type experiments.

2.1 Procedure

The procedures for experiments with 100Sn and 63Sn-37Pb solders were analogous except that with 100Sn solder, aging temperatures were extended to 205° C. The procedure with 63Sn-37Pb solder is described below.

The substrates used in the study were oxygen-free, high-conductivity (OFHC) copper tabs measuring 0.635 x 0.635 x 0.159 cm. The composition of the copper was verified by atomic emission spectroscopy, induced couple plasma (AES-ICP) analysis to be less than: 40 ppm Ag, 50 ppm Al, 10 ppm Mg, 30 ppm Mn, and 10 ppm Si (with a relative error of ± 100 %). The tabs were punched from sheet stock and a small hole drilled into one of the corners. The surface from which the thickness values were measured was optically polished. Next, a steel wire was secured to the hole which allowed for handling of the specimen.

The intermetallic layer thickness data were obtained by hot dipping the copper specimen into a bath of 63Sn-37Pb solder. The eutectic temperature of the solder is 183° C; however, the working temperature was slightly higher, 215° C. The tabs were solvent degreased, and then coated with a water soluble, organic acid flux. Next, each specimen was completely immersed into the solder bath for 5 s with the optically polished face down and parallel to the surface of the solder so that, upon removal, a large accumulation of solder formed on the surface. This procedure provided for an effectively infinite source of both the solder and the copper for intermetallic growth. Finally, the specimens were cleaned of flux residues.

Thermal aging of the solder-coated copper test samples was performed in air furnaces with temperature stabilities of ± 0.5 ° C. The Cu-63Sn-37Pb couples were annealed at 70, 100, 135, and 170° C for time periods of between 1 and 400 days (± 30 minutes). Upon completion of the annealing treatment, each specimen was cut in half; one half was mounted and polished for metallographic examination and analysis. The other half of the sample was retained for archival purposes.

2.2 Results

The composition of the intermetallic layers, as a function of position, was determined by electron microprobe analysis (EMPA). At higher temperatures, two Cu-Sn intermetallic layers, Cu_3Sn and Cu_6Sn_5 , formed in both Cu-100Sn and Cu-63Sn-37Pb samples. A micrograph illustrating the two Cu-Sn intermetallic layers that formed in a Cu-100Sn sample aged for 40 days at 205°C is shown in Fig. 2. The corresponding electron microprobe results are shown in Fig. 3. The Cu_3Sn layer forms adjacent to the Cu (bottom layer), and the Cu_6Sn_5 adjacent to the Sn (top layer). Similarly, a micrograph illustrating the two Cu-Sn intermetallic layers that formed in a Cu-63Sn-37Pb sample aged for 400 days at 170°C is shown in Fig. 4. In addition to the two Cu-Sn layers, a Pb-rich Sn layer also formed as illustrated in the corresponding electron microprobe data shown in Fig. 5. Again, the Cu_3Sn layer formed adjacent to the Cu (bottom layer). The Cu_6Sn_5 formed between the Cu_3Sn and the Pb-Sn layer that formed adjacent to the bulk solder containing a Pb-rich dispersed intermetallic precipitate. At lower temperatures, only one Cu-Sn intermetallic layer, Cu_6Sn_5 , formed in both Cu-100Sn and Cu-63Sn-37Pb samples.

The microprobe traces were made perpendicular to the layer (using 0.5 micron steps for the case of one layer composition). However, the spatial resolution of the concentration profile was a circle of 3-micron diameter about the target point caused by the interaction volume of the electron beam and the subsequent x-ray emissions. The elemental concentrations were listed as a function of distance along the trace. Through the change in the elemental concentrations, the boundaries of the individual intermetallic layers were designated; the sample point midway between the boundary values was used to designate the layer composition for that particular trace. Ten such traces were made per test sample in order to establish the composition of the layers(s). The traces were extended into the solder field, which together with point analysis, were used to determine the extent of Cu dissolved into the Pb- and Pb-rich phases as well as any response by the composition of those phases to the aging conditions.

Figure 2. Micrograph from Cu-100Sn sample aged for 40 days at 205°C.

Figure 3. Electron microprobe results showing composition of intermetallic regions, Cu_3Sn and Cu_6Sn_7 , in a Cu-100Sn sample aged for 40 days at 205°C.

Figure 4. Micrograph from Cu-63Sn-37Pb sample aged for 400 days at 170° C.

Figure 5. Electron microprobe results showing composition of intermetallic layers, Cu_3Sn , Cu_6Sn_5 , and Pb-rich Pb-Sn, in a Cu-63Sn-37Pb sample aged for 400 days at 170° C.

Assessment of the intermetallic layer growth was determined by quantitative image analysis. A study was performed to determine an optimum number of thickness measurements that would satisfy data accuracy, while also limiting the effort required on the part of the analyst. It was determined that a minimum of 40 data points would be adequate for the study. Therefore, a procedure was developed whereby four regions along the solder-copper interface were imaged at a magnification of 1000X. Ten measurements were made per image. In those cases for which two compositions comprised the total intermetallic layer, the thickness measurements were performed on the *inner* layer (Cu_3Sn) as well as on the *total* layer. The thickness of the outer layer (Cu_6Sn_5) was determined by subtraction of the inner layer thickness from the total value. The absolute error of the thickness measurement technique was 0.5 microns. The thickness data for each annealing time and temperature condition was represented by the mean of the data and one standard deviation of the mean.

The mean thicknesses of the Cu_3Sn and Cu_6Sn_5 layers determined by image analysis of the Cu-100Sn samples are summarized in Table 1 as a function of aging temperature and time. The numbers in parentheses represent one standard deviation. The thicknesses of the Cu-63Sn-37Pb samples are summarized in Table 2. At a given temperature and time, the total thickness of the intermetallic layers in the Cu-100Sn and Cu-63Sn-37Pb samples are similar. However, the ratio of the thicknesses of the Cu_3Sn layer to the thickness of the Cu_6Sn_5 layer in the Cu-100Sn samples are generally larger than the ratio for the Cu-63Sn-37Pb samples.

Table 1: Intermetallic Layer Thicknesses for Cu-100Sn

Temp. (C)	Time (days)	Cu_3Sn Layer (μm)	Cu_6Sn_5 Layer (μm)	Total (μm)
ambient	0	0	0.79 (0.28)	0.79 (0.28)
70	2	0	0.84 (0.51)	0.84 (0.51)
	10	0	1.42 (0.75)	1.42 (0.75)
	25	0	0.87 (0.28)	0.87 (0.28)
	100	0	0.90 (0.32)	0.90 (0.32)
	200	0	0.83 (0.38)	0.83 (0.38)
	350	0	1.02 (0.27)	1.02 (0.27)
100	2	0	0.82 (0.35)	0.82 (0.35)
	10	0	0.94 (0.43)	0.94 (0.43)
	25	0	1.51 (0.52)	1.51 (0.52)

Table 1: Intermetallic Layer Thicknesses for Cu-100Sn

Temp. (C)	Time (days)	Cu ₃ Sn Layer (μm)	Cu ₆ Sn ₅ Layer (μm)	Total (μm)
100	100	0	1.98 (0.54)	1.98 (0.54)
	200	1.11 (0.46)	0.91 (0.71)	2.02 (0.73)
	350	2.21 (0.70)	2.16	4.36
135	2	0	1.38 (0.31)	1.38 (0.31)
	10	1.4 (0.4)	0.86 (0.37)	2.3 (0.6)
	25	1.8 (0.6)	1.1 (0.8)	2.9 (1.0)
	100	3.1 (0.9)	5.9	9.0
	200	4.1 (1.0)	8.6 (7.7)	12.7 (7.5)
	350	5.7 (1.2)	10.2 (6.4)	16.0 (7.0)
170	1	1.6 (0.4)	0.9 (0.4)	2.5 (0.5)
	4	2.5 (0.6)	1.6 (0.6)	4.1 (0.6)
	10	3.6 (0.8)	2.9 (0.9)	6.5 (1.1)
	40	3.5 (1.1)	5.3 (2.7)	10.8 (2.6)
	100	7.7 (1.4)	7.3 (2.6)	15.0 (2.3)
	150	9.7 (1.8)	7.5 (2.3)	17.4 (2.1)
	200	11.1 (1.5)	8.8 (2.1)	19.9 (2.1)
	400	14.6 (2.8)	12.7 (6.2)	27.3 (5.8)
205	1	2.7 (0.5)	2.3 (0.8)	4.9 (0.5)
	4	4.5 (0.5)	3.9 (1.1)	8.4 (0.8)
	10	7.0 (1.2)	5.2 (1.7)	12.2 (1.2)
	40	11.6 (1.2)	10.1 (2.0)	21.7 (1.4)
	200	26.2 (2.4)	16.3 (2.9)	42.4 (1.4)

Table 2: Intermetallic Layer Thicknesses for Cu-63Sn-37Pb

Temp. (C)	Time (days)	Cu ₃ Sn Layer (μm)	Cu ₆ Sn ₅ Layer (μm)	Total (μm)
ambient	0	0		
70	2	0	1.3 (0.3)	1.3 (0.3)
	6	0	1.3 (0.3)	1.3 (0.3)
	10	0	1.4 (0.3)	1.4 (0.3)
	25	0	1.4 (0.3)	1.4 (0.3)
	51	0	1.7 (0.5)	1.7 (0.5)
	90	0	1.7 (0.4)	1.7 (0.4)
	150	0	1.9 (0.4)	1.9 (0.4)
	200	0	1.7 (0.4)	1.7 (0.4)
	300	0	2.1 (0.5)	2.1 (0.5)
	351	0	1.9 (0.5)	1.9 (0.5)
	400	0	2.0 (0.5)	2.0 (0.5)
100	2	0	1.5 (0.4)	1.5 (0.4)
	6	0	1.5 (0.2)	1.5 (0.2)
	10	0	1.7 (0.3)	1.7 (0.3)
	25	0	2.0 (0.3)	2.0 (0.3)
	51	0	2.3 (0.4)	2.3 (0.4)
	90	0	3.0 (0.5)	3.0 (0.5)
	150	0	3.7 (0.5)	3.7 (0.5)
	200	0	4.0 (0.9)	4.0 (0.9)
	300	0	4.4 (0.9)	4.4 (0.9)
	351	0	4.2 (0.5)	4.2 (0.5)
	400	0	4.8 (0.9)	4.8 (0.9)

Table 2: Intermetallic Layer Thicknesses for Cu-63Sn-37Pb

Temp. (C)	Time (days)	Cu ₃ Sn Layer (μm)	Cu ₆ Sn ₅ Layer (μm)	Total (μm)
135	2.96	0	2.2 (0.6)	2.2 (0.6)
	5.2	0	2.4 (0.3)	2.4 (0.3)
	10.3	0	2.9 (0.5)	2.9 (0.5)
	25	0	3.7 (0.5)	3.7 (0.5)
	50.4	0	5.0 (0.8)	5.0 (0.8)
	135	1.1 (0.4)	6.5 (0.8)	7.6 (1.2)
	150	1.4 (0.4)	7.7 (0.9)	9.1 (1.3)
	205	3.1 (0.7)	7.3 (1.0)	10.4 (1.7)
	300.3	2.9 (0.5)	8.8 (1.3)	11.7 (1.3)
	400	4.8 (0.7)	9.0 (2.0)	13.3 (1.6)
170	1	0	2.8 (0.3)	2.8 (0.3)
	4.08	0	3.5 (0.5)	3.5 (0.5)
	10	1.4 (0.4)	4.3 (1.1)	5.7 (1.5)
	10	1.7 (0.3)	4.0 (1.0)	5.7 (0.8)
	40	3.5 (0.7)	6.9 (2.1)	10.4 (2.8)
	60	4.2 (0.9)	7.7 (2.5)	11.9 (3.4)
	98	5.3 (1.0)	8.5 (3.0)	18.8 (4.0)
	149	6.5 (1.5)	11.7 (2.6)	18.2 (4.1)
	200	8.1 (1.7)	12.9 (2.8)	21.0 (4.5)
	262	8.6 (1.6)	15.1 (2.7)	23.7 (4.3)
	311	9.7 (1.4)	17.2 (4.7)	26.9 (6.1)
	402.7	11.1 (1.8)	16.2 (4.5)	27.3 (3.6)

3.0 THEORY

The growth of intermetallic layers occurs by displacement of the interfaces between layers that results from diffusion and interfacial reaction of the constituents of the layers. Analytical solutions to the diffusion and interface displacement equations are available for analyzing simple systems in which: (1) diffusion of the constituents can be represented by Fick's law with a constant diffusion coefficient; (2) interfacial reactions are sufficiently fast so that local chemical equilibrium essentially exists at both sides of each interface, and constituent concentrations at both sides can be considered constant, although generally they are not equal; (3) all layers grow simultaneously and have negligible initial thicknesses; (4) constituent diffusion and interface displacement are not hindered by finite material boundaries; and (5) diffusion and layer growth can be considered one-dimensional. The objective of the present modeling and code development effort was to provide numerical tools necessary for the analysis of more complex systems and for predicting intermetallic growth during service in which: (1) diffusion of constituents occurs by multiple mechanisms having variable diffusion coefficients; (2) interfacial chemical reactions are slow enough to be rate-limiting; (3) all layers do not grow simultaneously, due to different nucleation rates; (4) initial film thicknesses are not negligible; (5) constituent diffusion and interface displacement are hindered by finite material boundaries; (6) the temperature and, therefore, the diffusion coefficients vary substantially during layer growth; and (7) constituent diffusion and layer growth occur in more than one dimension.

A summary of the modeling and corresponding code development work is given below. The equations governing the basic phenomena controlling intermetallic growth are described first. Analytical and numerical solutions of the equations are then discussed, and the use of the numerical model is illustrated by application to experimental data for the Cu-100Sn and the Cu-63Sn-37Pb systems. This work primarily addresses the long-term growth of intermetallic layers that were previously formed by short-term contact between a metal substrate and molten solder.

This work is based on a one-dimensional, multi-layer system involving a binary set of interacting constituents, such as Cu and Sn, and in each layer, the diffusion of either constituent can be rate-controlling. The model consists of a basic set of material balance equations for each layer and displacement equations for the interfaces on each side of the layer. The material balance equations describe bulk diffusion of the rate-controlling constituent in each layer. In addition to bulk diffusion, the material balance equations can accommodate terms for additional diffusion mechanisms, such as grain boundary diffusion, and terms for rate-limiting interfacial reactions. Material balance equations can also be included to describe diffusion of trace constituents as appropriate.

A substrate-solder system in which $I-1$ intermetallic layers form and grow between the substrate and bulk solder is shown schematically in Fig. 6. Let the substrate form the left boundary of the system, and solder the right boundary. Let $i = 1, 2, 3, \dots, I-1$ denote the sequence of intermetallic layers growing from left to right. That is, $i = 1$ denotes the layer adjacent to the right edge of the substrate ($i = 0$), and $i = I-1$ denotes the layer adjacent to the left edge of the solder ($i = I$).

Figure 6. Schematic diagram of intermetallic layer growth between substrate and solder.

Furthermore, let T denote the temperature, t denote time, x denote position with respect to an origin fixed in space, $\delta_i(t)$ the position of the interface between layer $i-1$ and layer i , and $\Delta_i(t)$ the thickness of layer i

$$\Delta_i(t) = \delta_{i+1}(t) - \delta_i(t) \quad (1)$$

Then, for $\delta_i(t) < x < \delta_{i+1}(t)$, the material balance for the rate controlling constituent in each layer is given by

$$\frac{\partial w_{A_i}}{\partial t} = \frac{\partial}{\partial x} (D_{A_i} \frac{\partial w_{A_i}}{\partial x}) \quad (2)$$

where w_{A_i} denotes the mass fraction of the rate-controlling constituent in layer i , and D_{A_i} denotes the diffusion coefficient, which can be a function of x , t , w_{A_i} , and T , and can be specified accordingly. In general, the temperature T can be a function of x and t , which also can be specified.

When constituent diffusion and interface displacement are not affected by finite material boundaries, and the rate of interfacial chemical reaction, R_i , is sufficiently fast, local chemical equilibrium exists at the interface, and the concentrations of the diffusing constituents in layer i are essentially constant at $\delta_i(t)$. Therefore, at $\delta_i(t)$, the boundary condition for Eq. 2 is

$$(w_{A_i})_{\delta_i^r} = \text{constant} = (w_{A_i}^*)_{\delta_i^r} \quad (3)$$

and if local equilibrium exists at $\delta_{i+1}(t)$, the boundary condition is

$$(w_{A_i})_{\delta_{i+1}^l} = \text{constant} = (w_{A_i}^*)_{\delta_{i+1}^l} \quad (4)$$

where $(w_{A_i}^*)_{\delta_i^r} \neq (w_{A_i}^*)_{\delta_{i+1}^l}$, and superscripts l and r denote values at the left and right sides of the interface, respectively. The initial condition for Eq. 2 is

$$w_{A_i}(x, 0) = f_i(x) \quad (5)$$

where $f_i(x)$ is known or assumed.

When the rate of interfacial reaction, R_i , is slow enough to preclude local equilibrium at the interface, the constant concentration boundary conditions given by Eqs. 3 and 4 must be replaced by flux-type boundary conditions given by

$$-\rho_i D_{A_i} \left(\frac{\partial w_{A_i}}{\partial x} \right)_{\delta_i^r} = R_i^r \quad (6)$$

$$-\rho_i D_{A_i} \left(\frac{\partial w_{A_i}}{\partial x} \right)_{\delta_{i+1}^l} = R_{i+1}^l \quad (7)$$

where ρ_i denotes the mass density of layer i . Furthermore, when constituent diffusion encounters a finite material boundary, the boundary condition for Eq. 2 becomes the zero flux condition at that boundary.

The displacement of each interface is given by

$$\left[\rho_{i-1} (w_{A_{i-1}})_{\delta_i^l} - \rho_i (w_{A_i})_{\delta_i^r} \right] \frac{d\delta_i}{dt} = -\rho_{i-1} D_{A_{i-1}} \left(\frac{\partial w_{A_{i-1}}}{\partial x} \right)_{\delta_i^l} + \rho_i D_{A_i} \left(\frac{\partial w_{A_i}}{\partial x} \right)_{\delta_i^r} \quad (8)$$

The initial condition for Eq. 8 is

$$\delta_i (t = 0) = \delta_i^0 = \text{constant} \quad (9)$$

In the numerical analyses discussed below, the interfacial reaction rates were expressed by generic linear-driving-force expressions of the form

$$R_i^r = k_i^r \rho_i \left[(w_{A_i}^*)_{\delta_i^r} - (w_{A_i})_{\delta_i^r} \right] \quad (10)$$

and

$$R_{i+1}^l = k_{i+1}^l \rho_i \left[(w_{A_i}^*)_{\delta_{i+1}^l} - (w_{A_i})_{\delta_{i+1}^l} \right] \quad (11)$$

where k_i^r and k_{i+1}^l are usually empirical rate constants. Equations 10 and 11 are useful for examining order of magnitude effects.

Depending on the magnitude of the right side of Eq. 8, the left side can cause numerical problems when the term in brackets approaches zero, since $d\delta/dt$ can become large and physically unreasonable. These problems can be circumvented with an alternate form of Eq. 8 derived using the interface $\delta_i(t)$ as the frame of reference for diffusion in the region $\delta_i(t) < x < \delta_{i+1}(t)$. This alternate approach is summarized in Appendix B, where equations analogous to Eqs. 1-9 are given. The disadvantages in using $\delta_i(t)$ as the frame of reference are that the physical interpretation of the diffusion coefficient is difficult, and the approach would be very difficult to extend to two dimensions.

4.0 NUMERICAL APPROACH

4.1 Equation Transformation

Section 1.0 discusses different approaches to solving Stefan problems, particularly in the heat transfer arena. One advantage of the enthalpy method is that the need to explicitly track interface boundaries is circumvented. Unfortunately, there is no enthalpy analogy in problems of mass transfer. A difficulty with moving boundary solutions is remeshing the growing or shrinking layers. Remeshing increases computational time and can introduce error as the solution from one mesh is interpolated onto an updated mesh. In the method described in this report, a transformation of the spatial coordinate, x , given by

$$\bar{x}_i = \frac{x - \delta_i(t)}{\Delta_i(t)} \quad (12)$$

is used to circumvent the need to remesh the growing or shrinking layers. Then, for $\delta_i < x < \delta_{i+1}$, or $0 < \bar{x}_i < 1$,

$$\frac{\partial w_{A_i}}{\partial t} = \frac{1}{\Delta_i^2} \frac{\partial}{\partial \bar{x}_i} \left(D_{A_i} \frac{\partial w_{A_i}}{\partial \bar{x}_i} \right) + F(\bar{x}_i, \delta_i, \Delta_i) \frac{\partial w_{A_i}}{\partial \bar{x}_i} \quad (13)$$

where

$$F(\bar{x}_i, \delta_i, \Delta_i) = \left(\frac{1}{\Delta_i} \frac{d\delta_i}{dt} + \frac{\bar{x}_i}{\Delta_i} \frac{d\Delta_i}{dt} \right) \quad (14)$$

At $x = \delta_i^r(t)$, or $\bar{x}_i = 0$,

$$\left[\rho_{i-1}(w_{A_{i-1}})_{\bar{x}_{i-1}=1} - \rho_i(w_{A_i})_{\bar{x}_i=0} \right] \frac{d\delta_i}{dt} = - \frac{\rho_{i-1} D_{A_{i-1}}}{\Delta_{i-1}} \left(\frac{\partial w_{A_{i-1}}}{\partial \bar{x}_{i-1}} \right)_{\bar{x}_{i-1}=1} + \frac{\rho_i D_{A_i}}{\Delta_i} \left(\frac{\partial w_{A_i}}{\partial \bar{x}_i} \right)_{\bar{x}_i=0} \quad (15)$$

and at $x = \delta_{i+1}^l(t)$, or $\bar{x}_i = 1$,

$$\left[\rho_i(w_{A_i})_{\bar{x}_i=1} - \rho_{i+1}(w_{A_{i+1}})_{\bar{x}_{i+1}=0} \right] \frac{d\delta_i}{dt} = - \frac{\rho_i D_{A_i}}{\Delta_i} \left(\frac{\partial w_{A_i}}{\partial \bar{x}_i} \right)_{\bar{x}_i=1} + \frac{\rho_{i+1} D_{A_{i+1}}}{\Delta_{i+1}} \left(\frac{\partial w_{A_{i+1}}}{\partial \bar{x}_{i+1}} \right)_{\bar{x}_{i+1}=0} \quad (16)$$

During intermetallic growth, the initial rates of diffusion often appear to be higher than the rates at later times. To empirically account for those effects, the following variable diffusion coefficient $\bar{D}_{A_i}$ was used

$$\bar{D}_{A_i} = D_{A_i} \left(1 + \frac{D'_{A_i}}{D_{A_i}} \exp(-\lambda_i \Delta_i) \right) \quad (17)$$

where D_{A_i}' is an early-time diffusion coefficient that decays with the layer thickness $\Delta_i(t)$, and λ_i is the decay constant.

4.2 Implementation

The computer code ILaG (Intermetallic Layer Growth) solves the equations of Section 4.1 that describe intermetallic layer growth and mass fraction distribution in a multicomponent system. The numerical approach adopted to solve these equations, known as the method of lines (MOL) (Hyman, 1979), uncouples the spatial and temporal discretization of the partial differential equations. In this application, the spatial discretization is represented using a finite difference approximation, which results in a set of coupled, first-order, ordinary differential equations (ODE) in time. These are solved in turn by an efficient library ODE solver that determines internally the time step required to maintain stability. The implicit solution procedure of the ODE solver, DEBDF, uses a variable order (one through five) backward difference formulation (Shampine and Watts, 1980; Hindmarsh, 1981). This method has been successfully used in a number of applications (e.g. Baer, et al., 1986; Gross, et al., 1993). Baer's implementation for solving the reactive diffusion equations associated with thermal ignition of energetic materials was used as the basis for the current application. This implementation includes an adaptive meshing scheme (Baer, et al., 1986) that automatically resolves steep mass fraction gradients and will return to a coarser grid as the gradient relaxes. ILaG reads an ASCII input file that defines the case-specific mesh, material, and solution parameters. The code produces an output file that repeats the problem setup and reports the resulting mass fraction distribution and interface locations at the user-defined times. Two other ASCII output files are produced for use in plotting mass fraction profiles and layer thickness. A flow chart of the major functions within ILaG is given in Appendix C. The format of the required ASCII input file is described in Appendix D. A general discussion of the code's functions and the adaptation of the equations to the solution procedure follows.

After the user input file has been read (USERIN) and the solution parameters initialized, the initial mesh is generated (GRIDIN) for use with adaptive meshing. Mesh adjustments include positioning a node close to the zone boundaries and ensuring a smooth transition in the mesh from zone to zone. The distance between two nodes cannot vary by more than a factor of two than the distance to their neighboring nodes, a further restriction for mesh smoothness. The adaptive meshing routine, RFE1P1D, examines the initial conditions and default mesh adding nodes where needed based on the tolerances and maximum levels of refinement defined in line 4 of the user input file (see Appendix D). The meshing algorithm first performs *a posteriori* error analysis to produce a cumulative relative solution error. If this error is greater than RTOLREF, refinement is sought; if the error is less than RTOLREC, combining two adjacent cells is considered. Default values for RTOLREF and RTOLREC are 1.0×10^{-4} and $0.5 \times \text{RTOLREF}$, respectively. The default values may be invoked by setting RTOLREF and/or RTOLREC to zero in the input file. The default maximum level of refinement is six and again may be invoked by a zero value in the input file. Initial conditions, along with grid information, are written to the output file. Solution procedure parameters are then initialized, including the initial time step size which is some fraction of the maximum time step supplied in the input file.

Next, the transient solution procedure begins. The time step size is increased by ten percent until it reaches the specified maximum time step size. The current solution and mesh are passed to

RFE1PID for possible adaptation. Coordinate transformations are then applied to the modified mesh. After defining the time at which a new solution is needed, the ODE solver is called. DEBDF makes use of a subroutine (FUNC) external to the library package to evaluate the system of differential equations. DEBDF takes intermediate time steps to maintain stability, passing a value of time and the solution vector at that time to FUNC which returns the time derivatives. The time derivatives use a finite difference approximation to compute the required gradients. Two time derivatives, $d\delta_i/dt$ and $d\Delta_i/dt$, are used in computing the mass fraction time derivative. These derivatives, denoted by $C_{i,1}$ and $C_{i,2}$, respectively, in Eq. 18, are computed using a finite difference approximation at the end of a time step and then held constant during the subsequent time step. This approach is used because DEBDF is much more efficient if the Jacobian matrix is banded. For convenience, the subscript A denoting a particular constituent has been assumed in the equations below. Also, let $i = 0, 1, 2, \dots$ denote the layers of the system, and $n = 1, 2, \dots, N$ denote the nodal points within a layer so that $w_{i,n}$ is the mass fraction in layer i of the n^{th} node. The mass fraction time derivatives defined by Eq. 13 are calculated according to the following approximation

$$\dot{w}_{i,n} = \frac{1}{\Delta_i} \frac{[q_{n+1/2} - q_{n-1/2}]}{\Delta \bar{x}_n} + \frac{1}{\Delta_i} [C_{i,1} + \bar{x}_{i,n} C_{i,2}] \left[\frac{w_{i,n+1} - w_{i,n}}{\bar{x}_{i,n+1} - \bar{x}_{i,n}} \right] \quad (18)$$

where

$$q_{n+1/2} = \frac{\bar{D}_i}{\Delta_i} \left[\frac{w_{i,n+1} - w_{i,n}}{\bar{x}_{i,n+1} - \bar{x}_{i,n}} \right] \quad (19)$$

$$q_{n-1/2} = \frac{\bar{D}_i}{\Delta_i} \left[\frac{w_{i,n} - w_{i,n-1}}{\bar{x}_{i,n} - \bar{x}_{i,n-1}} \right] \quad (20)$$

$$\Delta \bar{x}_n = \frac{1}{2} [\bar{x}_{i,n+1} - \bar{x}_{i,n-1}] \quad (21)$$

Flux conditions are applied at the boundaries of each layer. The flux is defined to be zero at the outside boundaries, that is, at the left edge of the substrate and the right edge of the solder (Fig. 1). The flux at internal boundaries (or interfaces) is defined by the rate of reaction described in Eqs. 10 and 11. Note that the flux at the right side of interface i is the flux at the left edge of layer i and likewise, the flux at the left side of interface $i+1$ is the flux at the right edge of layer i . The finite difference implementation of this boundary flux yields the following time derivative for the mass fraction at the left end of a layer

$$\dot{w}_{i,1} = \frac{1}{\Delta_i} \frac{q_{3/2} + R_i^r/\rho_i}{\Delta \bar{x}_0} + \frac{1}{\Delta_i} [C_{i,1} + \bar{x}_{i,1} C_{i,2}] \left[\frac{w_{i,2} - w_{i,1}}{\bar{x}_{i,2} - \bar{x}_{i,1}} \right] \quad (22)$$

and for the right end

$$\dot{w}_{i,N} = \frac{1}{\Delta_i} \frac{q_{i,N-1/2} + R_{i+1}^l / \rho_i}{\Delta \bar{x}_N} + \frac{1}{\Delta_i} [C_{i,1} + \bar{x}_{i,N} C_{i,2}] \left[\frac{w_{i,N} - w_{i,N-1}}{\bar{x}_{i,N} - \bar{x}_{i,N-1}} \right] \quad (23)$$

where

$$\Delta \bar{x}_0 = \bar{x}_{i,1} + \frac{1}{2} [\bar{x}_{i,2} - \bar{x}_{i,1}] \quad (24)$$

$$\Delta \bar{x}_N = 1.0 - \bar{x}_{i,N} + \frac{1}{2} [\bar{x}_{i,N} - \bar{x}_{i,N-1}] \quad (25)$$

Presently, only the internal interface locations are allowed to move, that is, the outside edges of the solder and substrate are considered fixed. The time derivatives for the interface locations defined by Eq. 15 are approximated as

$$\dot{\delta}_i = \frac{-\frac{\rho_{i-1} \bar{D}_{i-1}}{\Delta_{i-1}} \left[\frac{w_{i-1,N} - w_{i-1,N-1}}{\bar{x}_{i-1,N} - \bar{x}_{i-1,N-1}} \right] + \frac{\rho_i \bar{D}_i}{\Delta_i} \left[\frac{w_{i,2} - w_{i,1}}{\bar{x}_{i,2} - \bar{x}_{i,1}} \right]}{\rho_{i-1} w_{i-1,N} - \rho_i w_{i,1}} \quad (26)$$

Control is returned to the main program from DEBDF with the solution vectors at the specified time. Results are printed and/or written according to criteria defined by the user in the input file. The values of $C_{i,1}$ and $C_{i,2}$ are updated, time is incremented, and the procedure repeated until the user-specified maximum time or number of time steps is exceeded.

4.3 Verification

If D_{A_i} and ρ_i have constant values in each layer, if the boundary conditions given by Eqs. 3 and 4 apply, and if δ_i^0 in Eq. 9 is negligible, then Eqs. 2 and 8 can be solved analytically for each layer as discussed by Jost (1960). In each layer, w_{A_i} is given by

$$w_{A_i} = (w_{A_i})_{\delta_i^0} - \frac{H_i}{\rho_i} \left[\frac{\text{erf}(\gamma_i) - \text{erf}(\eta_i)}{\text{erf}(\gamma_i) - \text{erf}(\gamma_{i+1} \sqrt{\phi_{i+1}})} \right] \quad (27)$$

where

$$\eta_i = \frac{x}{2\sqrt{D_{A_i} t}} \quad (28)$$

$$\phi_i = \frac{D_{A_i}}{D_{A_{i-1}}} \quad (29)$$

and

$$H_i = \rho_i \left[(w_{A_i})_{\delta_i^r} - (w_{A_i})_{\delta_{i+1}^l} \right] \quad (30)$$

The displacement of each interface is given by

$$\delta_i(t) = 2\gamma_i \sqrt{D_{A_i} t} \quad (31)$$

where

$$\gamma_i = \frac{1}{\sqrt{\pi\phi_i}} \frac{1}{h_i} \left[\frac{\sqrt{\phi_i} H_i \exp(-\gamma_i^2)}{\operatorname{erf}(\gamma_i) - \operatorname{erf}(\gamma_{i+1} \sqrt{\phi_{i+1}})} - \frac{H_{i-1} \exp(-\phi_i \gamma_i^2)}{\operatorname{erf}(\gamma_{i-1}) - \operatorname{erf}(\gamma_i \sqrt{\phi_i})} \right] \quad (32)$$

and

$$h_i = \rho_{i-1} (w_{A_{i-1}})_{\delta_i^l} - \rho_i (w_{A_i})_{\delta_i^r} \quad (33)$$

Equations 27 through 33 are useful for data analysis and for verifying numerical techniques. It should be noted that terms involving γ_i , γ_{i-1} , and γ_{i+1} appear in Eq. 32. Therefore, the equations for γ_i must be solved simultaneously, and since the equations are nonlinear, a numerical solution is necessary. For a system involving growth of $I-1$ intermetallic layers, the γ_i in Eq. 32 satisfy the following inequalities

$$\gamma_i < 0 \quad (34)$$

$$\gamma_i < \gamma_{i+1} \sqrt{\phi_{i+1}} \quad (35)$$

when $i = 1, 2, 3 \dots I$, and

$$\gamma_{I+1} > 0 \quad (36)$$

In this work, the equations for γ_i were solved using the symbol manipulation code Mathematica (Wolfram, 1991). For each γ_i , Eq. 32 was used as the source term for an ordinary differential equation of the form

$$\frac{d\gamma_i}{dt} = A_i + B_i \gamma_i \quad (37)$$

so that the long-time solutions of the differential equations approach the roots of the original equations for γ_i .

The numerical solution of Eqs. 2 and 8 was compared with the analytical solution for systems involving 2, 3, and 5 layers. The analytical solution is based on zero initial thickness of the inter-metallic layers and local chemical equilibrium between constituents at the interfaces between layers. The numerical solutions for the same conditions were in good agreement with the analytical results, as indicated in Fig. 7, which compares the analytical and numerical results for a five-layer system. The parameter values used in the calculations shown in Fig. 7 are given in Table 3. The material density was assumed constant and equal in all layers.

Figure 7. Comparison of analytical and numerical solutions for a five-zone system.

Table 3: Parameter Values for Comparison of Analytical and Numerical Solutions

Layer	$(w_{A_i})_{\delta_i^r}$	$(w_{A_i})_{\delta_{i+1}^l}$	$D_i \times 10^5$ (cm ² /day)	γ_i
1	1.0	1.0	0.0	---
2	0.8	0.7	3.0	-0.56894
3	0.6	0.5	3.0	-0.197728
4	0.4	0.2	3.0	0.0861818
5	0.1	0.1	0.0	0.838057

5.0 APPLICATIONS

In this work, the principal application of numerical modeling was in the analysis of the experimental data discussed previously for the Cu-100Sn and Cu-63Sn-37Pb systems. Detailed analyses using numerical modeling were performed to compare the experimental data for intermetallic growth with results calculated assuming growth was controlled by bulk diffusion of Cu or Sn. The analysis for Cu-100Sn is discussed first, followed by that for Cu-63Sn-37Pb.

The analyses for the Cu-100Sn system were performed using a four-layer model that consisted of Cu, Cu₃Sn, Cu₆Sn₅, and Sn layers, denoted by subscripts 1, 2, 3, and 4, respectively. Since the experimental data indicated that diffusion of Sn into the Cu substrate and diffusion of Cu into the bulk Sn were negligible, two diffusion equations (Eq. 2) were required, one for each of the Cu₃Sn and Cu₆Sn₅ layers, and three interface displacement equations, one each for the interface $\delta_1(t)$ between Cu and Cu₃Sn, $\delta_2(t)$ between Cu₃Sn and Cu₆Sn₅, and $\delta_3(t)$ between Cu₆Sn₅ and Sn. However, the experimental data did not indicate which constituent controlled bulk diffusion in the intermetallic layers. Since the choice of constituent does not alter the overall effect of bulk diffusion, the analyses were based on Cu being the diffusion-controlling constituent. Local equilibrium was assumed at each side of $\delta_1(t)$, $\delta_2(t)$, and $\delta_3(t)$. The constant concentration boundary conditions given by Eqs. 3 and 4 were then evaluated from the Cu-Sn phase diagram and are shown in Table 4, which also includes the mass density of each layer. The initial thicknesses of the intermetallic layers were taken from the experimental data for one day of heat treatment (1.6 and 0.9 microns for Cu₃Sn and Cu₆Sn₅, respectively).

Table 4: Interfacial Concentrations and Mass Densities

Layer	$(w_{A_i})_{\delta_i^r}$	$(w_{A_i})_{\delta_{i+1}^l}$	ρ_i (g/cm ³)
Cu (1)	1.000	1.000	8.9
Cu ₃ Sn (2)	0.623	0.605	9.0
Cu ₆ Sn ₅ (3)	0.410	0.400	8.4
Sn (4)	0.000	0.000	7.3

The diffusion coefficients D_{A_i} in both intermetallic layers were initially assumed constant. The experimental data for layer growth at times from 1 to 40 days were then used to determine values for D_{A_i} that reasonably reproduced the experimental data. Using those values, growth of the Cu₃Sn and Cu₆Sn₅ layers was predicted for times to 400 days and compared with the corresponding experimental data. Analogous calculations were then repeated using the experimental data from 1 to 100, 1 to 150, 1 to 200, and 1 to 400 days to determine the values for D_{A_i} . These results are summarized in Figs. 8 and 9 for the Cu₃Sn and Cu₆Sn₅ layers, respectively, at 170° C. The solid portion of each curve indicates the time interval in which the experimental data were used to evaluate the diffusion coefficients, and the dashed portion indicates the time interval in which the thickness was predicted. The diffusion coefficients corresponding to each curve for Cu₃Sn and Cu₆Sn₅ are shown in Table 5. The results generally indicate that the experimental data are consistent with a bulk diffusion mechanism, but the diffusion coefficient is probably not con-

Figure 8. Intermetallic layer thicknesses calculated using a constant diffusion coefficient for Cu_3Sn at 170°C .

Figure 9. Intermetallic layer thicknesses calculated using a constant diffusion coefficient for Cu_6Sn_5 at 170°C .

stant. During early times, diffusion appears to be faster relative to later times. Similar effects have been reported previously. Kofstad (1966) used the term “enhanced diffusion” to indicate effects which could be attributed to grain boundary diffusion, residual strain, and poor crystallization during early layer growth.

Table 5: Constant Cu Diffusion Coefficients for the Cu-100Sn Layers at 170° C

Time Interval from Day 1 to Day	$D_A (\text{Cu}_3\text{Sn}) \times 10^8$ (cm ² /day)	$D_A (\text{Cu}_6\text{Sn}_5) \times 10^7$ (cm ² /day)
40	9.0	1.5
100	7.5	1.2
150	6.8	1.0
200	6.5	1.0
400	6.0	0.9

To empirically account for the effects of enhanced diffusion, an analogous series of calculations was performed using a variable diffusion coefficient $\overline{D}_{Ai}$ defined by Eq. 17. The numerical results are compared with the experimental data in Figs. 10 and 11. The corresponding values for $\overline{D}_{Ai}$, D_{Ai} , and λ_i are given in Table 6. The agreement between experimental and calculated values is considerably improved. This improvement is further illustrated in Figs. 12 and 13 which compare the numerical results obtained using a constant diffusion coefficient and the variable coefficient given by Eq. 17.

Table 6: Variable Cu Diffusion Coefficients for the Cu-100Sn Layers at 170° C

Layer	Time Interval from Day 1 to Day	$D_A \times 10^8$ (cm ² /day)	$D'_A \times 10^7$ (cm ² /day)	$\lambda_i \times 10^3$ (cm ⁻¹)
Cu ₃ Sn	100	3.5	1.6	3.5
	200	4.5	1.4	3.8
	400	4.3	1.4	3.8
Cu ₆ Sn ₅	100	4.0	1.8	1.9
	200	4.0	1.8	2.1
	400	5.0	1.7	2.1

Figure 10. Intermetallic layer thicknesses (100Sn solder) calculated using a variable diffusion coefficient (Eq. 17) for Cu_3Sn at 170°C .

Figure 11. Intermetallic layer thicknesses (100Sn solder) calculated using a variable diffusion coefficient (Eq. 17) for Cu_6Sn_5 at 170°C .

Figure 12. Comparison of Cu_3Sn layer thicknesses calculated using constant and variable (Eq. 17) diffusion coefficients at 170°C

Figure 13. Comparison of Cu_6Sn_5 layer thicknesses calculated using constant and variable (Eq. 17) diffusion coefficients at 170°C .

Similar analyses were performed with the data from the experiments at 135 and 205° C, and the results also indicated that layer growth is consistent with a bulk diffusion mechanism involving a variable diffusion coefficient that reflects some enhanced diffusion during early layer growth. A comparison between the computational and experimental data for 135° C is shown in Figs. 14 and 15 for the Cu₃Sn and Cu₆Sn₅ layers, respectively, and for 205° C in Figs. 16 and 17. The values for D_{Ai} , D'_{Ai} , and λ_i corresponding to the computational results shown in Figs. 14 through 17 are given in Table 7. The computational results at 135° C require some additional explanation. Initially ($t = 1$ day), only the Cu₆Sn₅ layer was present. But, during the period between 2 and 10 days the Cu₃Sn layer nucleated, coalesced, and began to grow. The net effect of which was a "sudden" decrease in the thickness of the Cu₆Sn₅ layer and the "sudden" appearance of a Cu₃Sn layer. This effect was accounted for numerically by choosing an initial nucleation time ($t = 6$ days), a negligibly small initial Cu₃Sn layer thickness (0.2 microns), and large values for D_{Ai} , and λ_i so that growth of the Cu₃Sn layer was extremely fast during the first 10 days and then decayed rapidly to the long time rates observed experimentally. To obtain better agreement with the experimental data at early times, non-equilibrium was assumed at the Cu₃Sn-Cu₆Sn₅ interface during nucleation and coalescence. The flux-type boundary condition given by Eqs. 6 and 10 was used for the Cu₆Sn₅ layer at δ_2^r ($k_2^r = 2.8 \times 10^{-4}$ cm/s).

Table 7: Comparison of Variable Cu Diffusion Coefficients for the Cu-100Sn Layers

Layer	Temperature (°C)	$D_A \times 10^8$ (cm ² /day)	$D'_A \times 10^7$ (cm ² /day)	$\lambda_i \times 10^3$ (cm ⁻¹)	D'_A/D_A
Cu ₃ Sn	135	1.0	8.0	20.0	80.0
	170	4.3	1.4	3.8	3.3
	205	29	12.0	5.0	4.1
Cu ₆ Sn ₅	135	2.0	8.0	3.8	4.0
	170	5.0	1.7	2.1	3.4
	205	30.0	14.0	2.6	4.7

Analysis of the data from the Cu-63Sn-37Pb experiments was analogous to analysis of the data from the Cu-100Sn samples discussed above, except that the data were from experiments performed at 170, 135, and 100° C. The Pb-rich Pb-Sn layer was neglected because Cu was not detected in the layer. Again, the results indicated that layer growth was consistent with a bulk diffusion mechanism involving a variable diffusion coefficient that reflects some enhanced diffusion during early layer growth. However, the numerical values for the diffusion coefficients were significantly different from the values obtained for the Cu-100Sn samples at the same temperature. This is somewhat unexpected, since the bulk composition of Cu₃Sn and Cu₆Sn₅ layers appear similar in both the Cu-100Sn and Cu-63Sn-37Pb samples. A comparison between the

Figure 14. Intermetallic layer thicknesses (100Sn solder) calculated using a variable diffusion coefficient (Eq. 17) for Cu_3Sn at 135°C .

Figure 15. Intermetallic layer thicknesses (100Sn solder) calculated using a variable diffusion coefficient (Eq. 17) for Cu_6Sn_5 at 135°C .

Figure 16. Intermetallic layer thicknesses (100Sn solder) calculated using a variable diffusion coefficient (Eq. 17) for Cu_3Sn at 205°C .

Figure 17. Intermetallic layer thicknesses (100Sn solder) calculated using a variable diffusion coefficient (Eq. 17) for Cu_6Sn_5 at 205°C .

computational and experimental results for 170° C is shown in Figs. 18 and 19 for Cu₃Sn and Cu₆Sn₅, respectively, for 135° C in Figs. 20 and 21, and for 100° C in Fig. 22 for Cu₆Sn₅ (Cu₃Sn did not form at 100° C). The values for D_{Ai} , D'_{Ai} , and λ_i corresponding to the computational results shown in Figs. 18 through 22 are given in Table 8.

Table 8: Comparison of Variable Cu Diffusion Coefficients for Cu-63Sn-37Pb Layers

Layer	Temperature (°C)	$D_A \times 10^8$ (cm ² /day)	$D'_A \times 10^7$ (cm ² /day)	$\lambda_i \times 10^3$ (cm ⁻¹)	D'_A/D_A
Cu ₃ Sn	100	---	---	---	---
	135	0.7	0.4	20.0	5.7
	170	3.7	8.0	20.0	21.6
Cu ₆ Sn ₅	100	0.1	0.22	3.9	22.0
	135	2.0	1.6	3.7	8.0
	170	13.0	3.6	3.8	2.8

The values of D_A obtained for the Cu-100Sn and Cu-63Sn-37Pb samples at 135 and 170° C are compared in Table 9. In the Cu₃Sn layer, the value of D_A for the Cu-63Sn-37Pb samples at both temperatures is about 70 to 80 percent that of the value for the Cu-100Sn samples. In the Cu₆Sn₅ layer, the value of D_A for the Cu-63Sn-37Pb sample at 135° C is the same as the value for the Cu-100Sn sample. However, the value for the Cu-63Sn-37Pb sample at 170° C is much larger than the value for the Cu-100Sn sample.

The differences in the values of D_A obtained with Cu-100Sn and Cu-63Sn-37Pb samples result from the differences in layer growth shown in Tables 1 and 2, and primarily from the differences in the ratios of the thickness of the Cu₃Sn layer to the thickness of the Cu₆Sn₅ layer, which are shown in Table 10.

Table 9: Comparison of D_A Obtained from Cu-100Sn and Cu-63Sn-37Pb Samples

T (°C)	Intermetallic Layer Thickness (microns)	Solder	D_A (cm ² /day) $\times 10^8$
135	Cu ₃ Sn	100-Sn	1.0
		63Sn-37Pb	0.7
	Cu ₆ Sn ₅	100Sn	2.0

Table 9: Comparison of D_A Obtained from Cu-100Sn and Cu-63Sn-37Pb Samples

$T (^{\circ}\text{C})$	Intermetallic Layer Thickness (microns)	Solder	D_A (cm ² /day) x 10 ⁸
135	Cu ₆ Sn ₅	63Sn-37Pb	2.0
170	Cu ₃ Sn	100-Sn	4.3
		63Sn-37Pb	3.7
	Cu ₆ Sn ₅	100-Sn	5.0
		63Sn-37Pb	13.0

Table 10: Ratio of Cu₃Sn to Cu₆Sn₅ Layer Thickness

T (°C)	Time (days)	Cu-100Sn Samples	Cu-63Sn-37Pb Samples
70	---	---	---
100	350	1.02	0.00
135	350	0.56	
	400		0.53
170	400	1.15	0.69
205	200	1.6	---

Figure 18. Intermetallic layer thicknesses (63Sn-37Pb solder) calculated using a variable diffusion coefficient (Eq. 17) for Cu_3Sn at 170°C .

Figure 19. Intermetallic layer thicknesses (63Sn-37Pb solder) calculated using a variable diffusion coefficient (Eq. 17) for Cu_6Sn_5 at 170°C .

Figure 20. Intermetallic layer thicknesses (63Sn-37Pb solder) calculated using a variable diffusion coefficient (Eq. 17) for Cu_3Sn at 135°C .

Figure 21. Intermetallic layer thicknesses (63Sn-37Pb solder) calculated using a variable diffusion coefficient (Eq. 17) for Cu_6Sn_5 at 135°C .

Figure 22. Intermetallic layer thicknesses (63Sn-37Pb solder) calculated using a variable diffusion coefficient (Eq. 17) for Cu_6Sn_5 at 100°C .

Arrhenius plots of the values for D_A in the Cu_3Sn and Cu_6Sn_5 layers formed with the Cu-100Sn samples are shown in Figs. 23 and 24, respectively, and with the Cu-63Sn-37Pb samples in Figs. 25 and 26. The values for D_o and Q in the expression

$$D_A = D_o \exp \left(-\frac{Q}{RT} \right) \quad (38)$$

are given in Table 11 and were calculated using the data at the two highest temperatures (lowest values of $1/T$), because nucleation of Cu_3Sn occurred very early during layer growth.

Table 11: Arrhenius Parameters

Solder	Intermetallic Layer	D_o (cm^2/day)	Q (kcal/mol)
Cu-100Sn	Cu_3Sn	1.0×10^4	23.2
	Cu_6Sn_5	1.9×10^3	21.6
Cu-63Sn-37Pb	Cu_3Sn	10	17.2
	Cu_6Sn_5	330	19.2

An apparent reason for the differences in layer growth and, therefore, diffusion coefficients obtained with Cu-100Sn and Cu-63Sn-37Pb samples is that diffusion in the Cu_6Sn_5 layer is more rapid in the Cu-63Sn-37Pb samples than in the Cu-100Sn samples. The results at 170°C (Table 9) certainly indicate rapid diffusion in the Cu-63Sn-37Pb sample, although the results at 135°C appear to contradict rapid diffusion. However, nucleation of Cu_3Sn in the Cu-63Pb-37Sn sample occurred relatively late at 135°C and could have perturbed layer growth enough to preclude accurate analyses with the current numerical model, since nucleation processes were not specifically included in the model development. Furthermore, some additional considerations also indicate that diffusion in the Cu_6Sn_5 layer is faster in the Cu-63Sn-37Pb samples: (1) at 100°C , Cu_3Sn nucleated at long times in the Cu-100Sn samples but did not nucleate in the Cu-63Sn-37Pb samples; (2) also at 100°C , prior to nucleation of Cu_3Sn , the Cu_6Sn_5 layer in the Cu-63Sn-37Pb samples grew faster than the layer in the Cu-100Sn samples; and (3) at 135°C , the Cu_3Sn layer nucleates much earlier in the Cu-100Sn samples than in the Cu-63Sn-37Pb samples.

The mechanism for rapid diffusion in the Cu_6Sn_5 layer of the Cu-63Sn-37Pb samples was not determined. An obvious possibility is a perturbation caused by trace amounts of Pb in the Cu_6Sn_5 layer; however, Auger depth profiling did not detect Pb in the Cu_6Sn_5 layers.

Figure 23. Arrhenius plot for D_A in Cu_3Sn layer (Cu-100Sn).

Figure 24. Arrhenius plot for D_A in Cu_6Sn_5 layer (Cu-100Sn).

Figure 25. Arrhenius plot for D_A in Cu_3Sn layer (Cu-63Sn-37Pb).

Figure 26. Arrhenius plot for D_A in Cu_6Sn_5 layer (Cu-63Sn-37Pb).

6.0 RECOMMENDATIONS FOR FURTHER WORK

The model and code developed in this work provide tools for analyzing growth of multiple intermetallic layers in one-dimensional systems in which layer growth is primarily controlled by bulk chemical diffusion and (or) interfacial reactions involving two layer-forming constituents. In both commercial and research applications, these conditions often will be satisfied. However, more complex systems are becoming increasingly of interest, and several avenues for further work exist, as illustrated in Fig. 27. These principally consist of extending the current model to include expressions for additional mass transfer mechanisms, equations for multicomponent systems of diffusing, layer forming species, and multidimensional mass transfer and layer growth. These extensions are discussed in more detail below.

Expressions for several mass transfer mechanisms could be added to the current model. *The most immediately useful addition would be Arrhenius expressions for the diffusion coefficients $D = D_0 \exp(-Q/RT)$ and interfacial reaction rate constants $k = k_0 \exp(-Q/RT)$, which would permit analysis of nonisothermal aging experiments and the prediction of intermetallic growth under nonisothermal service conditions for product reliability analyses.* Other mass transfer mechanisms which can be added to the current model and could be important in reliability analyses and basic materials science and engineering studies include: (1) *multiple chemical diffusion mechanisms (such as substitutional, interstitial, etc.),* (2) *concentration and stress dependent diffusion coefficients,* (3) *grain boundary diffusion,* (4) *stress-induced diffusion,* and (5) *thermally-induced diffusion.* These mechanisms, particularly (2-5), can be especially important in analyzing or predicting intermetallic layer growth in situations involving thermal and (or) mechanical cycling during which large temperature or stress gradients exist and may be time-dependent.

While the current model is limited to diffusion of two layer forming species, the governing equations can be expanded to *include diffusion of additional trace constituents (say one to three species).* This extension would be useful in examining the possible effects of contaminants on intermetallic layer growth and the formation of interfacial defects due to contaminant accumulation, which can have substantial impact on joint integrity and reliability.

Although the emphasis in this work has been on aging studies and intermetallic growth during service, the model can also be applied to intermetallic growth during several processing applications, which will often involve a melt layer. For analyses involving a melt layer, the model should incorporate *temperature-dependent interfacial concentrations (to account for a temperature dependent solubility of the diffusing species in a melt layer).* *Heat conduction equations with latent heat and energy of reaction terms* also should be added to account for solidification and energy generation due to intermetallic formation. These additions would also be valuable for modeling the combustion of gasless pyrotechnics such as thermites.

Also, further work can be done to develop *approximate expressions for incorporating the effects of phase nucleation and coalescence; enhanced diffusion during early layer growth could be addressed in more detail, and for ionic systems, the interaction between the electric field and the diffusing ionic species* could be included. Since the mechanisms for enhanced diffusion are somewhat obscure, a more detailed treatment of this phenomena must be approached with complementary experimental work. The effects of nucleation, particularly when examining criteria for when nucleation will occur, will require a similar approach.

Figure 27. Schematic diagram showing recommendations for future work.

Extension of the governing equations to ternary systems of the layer forming species could have broad applications in the joining of advanced materials, such as braze joints involving metals and ceramics. However, since the chemistry and geometry of intermetallic formation in ternary systems is often complex, care would be needed in formulating equations and defining criteria for their application. Also, *volume averaging techniques should be investigated for application to complex geometries involving dispersed and (or) very irregularly shaped intermetallic phases.* One probable situation is a very irregularly shaped intermetallic layer between two otherwise relatively uniform bulk materials. Here, volume averaging could be applied to an interfacial region, which is then coupled to the bulk regions using appropriately derived boundary conditions. A second probable situation involves a dispersed phase having a relatively simple geometry. Here, the interface is treated as in the current model, and volume averaging would be applied to the bulk material.

Development of a 2-D code based on the method of lines appears feasible. The extension to complex 2-D, and 3-D, geometries would probably require finite element techniques. The extension to at least 2-D would have considerable value with respect to the geometries that could be modeled as well as addressing phenomena, such as nucleation and density change accompanying intermetallic formation, that cannot adequately be addressed in 1-D. Nucleation and coalescence is inherently at least a 2-D process, although simplifying assumptions can always be attempted. Likewise, density changes that occur as a result of intermetallic formation usually require at least a 2-D description. Density changes will also cause stress and strain effects and are inherently coupled with the mechanical properties of the intermetallic layers and bulk materials. The effects of density changes can severely impact joint integrity and reliability. Density changes can result in severe stress, such as expansion of a growing layer at an internal corner, and can also result in void formation, which seriously affects the fracture toughness of brittle materials such as ceramics.

To adequately model the effects of density changes, a *2-D intermetallic growth code would have to be loosely coupled to a solid mechanics code.* In addition to analyzing the effects of density change due to reaction, the coupling would provide the means for analyzing the combined effects of intermetallic growth, thermal stress, and work failure (fatigue), which could be particularly important to the reliability of joints in systems involving thermal and mechanical cycling, especially if large temperature and stress gradients occur.

Finally in terms of analyzing experimental data, an extremely valuable addition to the current code and to any codes developed in future work would be a parameter estimation routine to determine the parameters giving the "best" fit to experimental data. In fact the most immediate progress would be made by extending the current code to include Arrhenius expressions for the diffusion coefficients and interfacial rate constants and then coupling the resulting code with a parameter estimation routine.

7.0 CONCLUSIONS

A model describing the diffusion-controlled growth of multiple intermetallic layers and the displacement of the interfaces between layers was developed and implemented in a one-dimensional computer code based on the method-of-lines. The model can accommodate problems, which cannot be modeled analytically, involving: (1) finite initial layer thicknesses, (2) rate-limiting interfacial reactions, (3) multiple and variable diffusion coefficients, and (4) finite material boundaries. Additionally, the effects of nucleation can be modeled empirically. A transformation of the spatial coordinate circumvented the need to remesh the growing and (or) shrinking layers. Results from the one-dimensional code were verified by comparing the numerical results with analytical solutions for simple systems involving two, three, and five layers. The computer code was applied to analysis of the intermetallic layer growth that occurred in solder aging experiments that were done with 100Sn and 63Sn-37Pb solders. The Cu-63Sn-37Pb couples were aged for 1 to 400 days at temperatures of 70, 100, 135, and 170° C. The Cu-100Sn samples were aged similarly, except that annealing treatments were also done at 205° C. Two intermetallic layers, Cu₃Sn and Cu₆Sn₅, were observed. Some differences between the thickness of the intermetallic layers formed with 100Sn and 63Sn-37Pb solders were observed. At higher temperatures, the total thickness of the layers were similar, but the ratio of the thicknesses of the Cu₃Sn to the Cu₆Sn₅ layer differed significantly.

The analysis of the experimental data demonstrated that numerical modeling can be a valuable tool in the investigation of intermetallic layer growth. The results indicated that intermetallic layer growth was consistent with a bulk diffusion mechanism involving Cu and (or) Sn and variable diffusion coefficients that reflected enhanced diffusion during early layer growth. The enhanced diffusion can be accounted for empirically using a variable diffusion coefficient that is a function of layer thickness. Such effects are important when using data from short term isothermal experiments to evaluate diffusion coefficients to be used for predicting long-term layer growth under nonisothermal service conditions. The effects of enhanced diffusion are also important when trying to predict the nucleation of additional phases during growth of the original layers. A satisfactory explanation for the difference between intermetallic layer growth that was observed with 100Sn and 63Sn-37Pb solder was not established. Diffusion in the Cu₆Sn₅ layer appeared to be faster with 63Sn-37Pb samples than with 100Sn samples. The faster diffusion may have been due to trace amounts of Pb perturbing the layer, although the presence of Pb could not be confirmed by Auger depth profiling and electron microprobe analysis.

Several commercially relevant avenues for further work exist. These principally consist of extending the current model to include expressions for additional mass transfer mechanisms, equations for multicomponent systems of diffusing, layer forming species, and multidimensional mass transfer and layer growth. Expressions for several mass transfer mechanisms could be added to the current model. The most immediate progress would be made by extending the current code to include Arrhenius expressions for the diffusion coefficients and interfacial rate constants and then coupling the resulting code with a parameter estimation routine.

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APPENDIX A: Marker Layers

The use of marker layers at the solder-substrate interface would allow for determining the absolute movement of the various interfaces during growth of the intermetallic compound layer. The marker is typically a thin film layer deposited at the interface and which does not react with either of the members of the diffusion couple (in this case, the solder and the substrate material) so that the position of the original interface can be maintained as a reference point. Marker layers have been used in diffusion studies of vapor deposited multilayer thin films, the marker layers, themselves, being thin films, as well (Tu and Thompson, 1982). Diffusion studies between bulk materials have used refractory metal wires to denote movement of the interface(s) (Wheeler, 1965). The use of marker layers in solder-substrate diffusion couples would provide a method of determining the diffusion mechanism(s) responsible for the growth of interfacial intermetallic layers.

The selection of the marker layer material was based upon several requirements. First, the material had to be deposited by evaporation techniques to thicknesses of greater than one micron (which is relatively thick by thin-film evaporation standards) in order to be visible for later reference. A non-precious metal material was favored, due to the extent of lost metal by this technique. The metal had to be adherent to the substrate metal (oxygen-free, high-conductivity OFHC copper) and remain so after deposition. Therefore, the residual stresses in the film, at those thicknesses, had to be minimal. Secondly, the layer had to be wetted by the molten solder, yet not be dissolved by the solder. Finally, the marker layer should not react during the solid state diffusion process so that the interface position would remain accessible.

Preliminary trials to find a suitable marker film were performed on OFHC copper tabs 6.35 x 6.35 x 1.60 mm. One surface of the tab was optically polished. Two tabs were cleaned by: (1) solvent degreasing; (2) 1-minute etch in 1:1 HCL:H₂O solution; and (3) rinsed with water and alcohol. Next, the substrate was coated with BlackstoneTM 642 flux and immersed into a pot of molten 100Sn at 270° C to provide the fused coating. Finally, the samples were cleaned of any flux residues.

The durability of the marker layer to heat treatments used in the solid state aging studies had to be assessed. Therefore, one of the coated tabs was air annealed at 170° C for 10 days and the other at 205° C for the same period. Upon completion of the annealing process, the tabs were cross sectioned across the marker layer, and one-half was mounted and polished for metallographic examination of the marker film.

Several marker layer systems were initially identified, primarily from barrier layer studies on copper. The first two systems included: (1) 800nm TiN + 2000nm Cu (top), and (2) 200nm TiN + 1000nm Ni + 2000nm Cu. The TiN and TiN + Ni formed the marker layers, respectively, while the copper layer served as the solderable finish in order to promote wetting by the molten tin. The TiN layer was sputtered deposited; the Ni and Cu layers were evaporation deposited. The two layer systems demonstrated excellent adhesion to the copper. The samples were then electroplated with copper (0.038 mm) to form the second layer to which the tin would wet. In this case, the marker layers would have been buried and themselves, not be contacted by the molten tin. The markers survived the electroplating process. However, the marker film exhibited cracking after hot dipping in the molten bath, prior to aging.

The next two systems that were examined included: (1) 3nm Al + 1000nm Cu, and (2) 3000nm Al. The films adhered very well to the copper. However, they did not survive the hot dipping operation.

Film systems were developed using nickel as the marker layer, owing to its reduced propensity for dissolution onto the molten tin. Two systems were attempted: (1) 1500nm Ni + 3000nm Au, and (2) 3000nm Ni + 3000nm Au. The gold served to protect the nickel layer from excess oxidation, thereby maintaining its solderability. The latter system did not survive the deposition process; the nickel layer spalled from the copper surface due to an excess of residual stresses (which are common in thick, nickel films). The system (1) showed no spalling. When the samples were hot dipped in solder and later cross sectioned (prior to the aging treatment), the markers were clearly visible with excellent wetting by the tin (i.e., no voids were present). This system was developed too late to be used in an aging study. However, the selected marker system is *1500nm Ni + 3000nm Cu*.

Lastly, fixtures and aperture masks were designed and fabricated which would allow for the deposition of the marker strips on copper plates measuring 38 x 38 x 3.2 mm in order to fabricate a large number of samples at the same time. In addition, a technique was developed to fabricate said samples with a controlled amount of tin coating. The specific tin coating thicknesses were 7.6 microns, 20 microns, 25 microns, and 41 microns. This technique would provide the quantitative growth data of the intermetallic layer as a function of available tin from which to further refine the computer models.

APPENDIX B: The Interface as the Frame of Reference

If $\delta_i(t)$ is used as the frame of reference for diffusion, then for $\delta_i(t) < x < \delta_{i+1}(t)$, the material balance (diffusion) equation corresponding to Eq. 2 is

$$\frac{\partial w_{A_i}}{\partial t} = \frac{\partial}{\partial x} (D_{A_i} \frac{\partial w_{A_i}}{\partial x}) + \frac{d\delta_i}{dt} \frac{\partial w_{A_i}}{\partial x} \quad (\text{B-1})$$

and at $\delta_i(t)$, the interface displacement equation corresponding to Eq. 8 is

$$\begin{aligned} & (\rho_{i-1}) (w_{A_{i-1}})_{\delta_i} \left(\frac{d\delta_{i-1}}{dt} \right) - \\ & \rho_{i-1} (w_{A_{i-1}})_{\delta_i} \left(\frac{d\delta_i}{dt} \right) = (\rho_{i-1}) D_{A_{i-1}} \left(\frac{\partial w_{A_{i-1}}}{\partial x} \right)_{\delta_i} + \\ & \rho_i D_{A_i} \left(\frac{\partial w_{A_i}}{\partial x} \right)_{\delta_i} \end{aligned} \quad (\text{B-2})$$

Let

$$\bar{x}_i = \frac{x - \delta_i(t)}{\Delta_i(t)} \quad (\text{B-3})$$

and

$$\Delta_i(t) = \delta_{i+1}(t) - \delta_i(t) \quad (\text{B-4})$$

Then, for $\delta_i < x < \delta_{i+1}$, or $0 < \bar{x}_i < 1$,

$$\frac{\partial w_{A_i}}{\partial t} = \frac{1}{\Delta_i^2} \frac{\partial}{\partial \bar{x}} (D_{A_i} \frac{\partial w_{A_i}}{\partial \bar{x}}) + F(\bar{x}_i, \delta_i, \Delta_i) \frac{\partial w_{A_i}}{\partial \bar{x}} \quad (\text{B-5})$$

where

$$F(\bar{x}_i, \delta_i, \Delta_i) = \frac{\bar{x}_i d\Delta_i}{\Delta_i dt} \quad (\text{B-6})$$

At $x = \delta_i(t)$,

$$\begin{aligned}
& \rho_{i-1} (w_{A_{i-1}})_{\bar{x}_{i-1}=1} \left(\frac{d\delta_{i-1}}{dt} \right) - \\
& \rho_{i-1} (w_{A_{i-1}})_{\bar{x}_{i-1}=1} \left(\frac{d\delta_i}{dt} \right) = \frac{\rho_{i-1} D_{A_{i-1}}}{\Delta_i} \left(\frac{\partial w_{A_{i-1}}}{\partial x} \right)_{\bar{x}_{i-1}=1} + \\
& \frac{\rho_i D_{A_i}}{\Delta_i} \left(\frac{\partial w_{A_i}}{\partial x} \right)_{\bar{x}_i=0}
\end{aligned} \tag{B-7}$$

When all $\delta_i(0) = 0$, Eqs. B-1 and B-2 can be solved analytically, similarly to Eqs. 2 and 8, using the constant concentration (local equilibrium at the interface) boundary conditions corresponding to Eqs. 3 and 4. For $\delta_i < x < \delta_{i+1}$, or $0 < \bar{x}_i < 1$,

$$w_{A_i} = (w_{A_i})_{\delta'_i} - \left[(w_{A_i})_{\delta'_i} - (w_{A_i})_{\delta'_{i+1}} \right] \frac{\text{erf}(\eta_i)}{\text{erf}(\gamma'_{i-1})} \tag{B-8}$$

where

$$\eta_i = \frac{x - \delta_i(t)}{2\sqrt{D_{A_i}t}} \tag{B-9}$$

$$\delta_i(t) = \delta_{i-1}(t) + \Delta_i(t) \tag{B-10}$$

$$\Delta_i(t) = \delta_i(t) - \delta_{i-1}(t) = 2\gamma'_i \sqrt{D_{A_i}t} \tag{B-11}$$

and

$$\begin{aligned}
\gamma'_i = & \frac{1}{\sqrt{\pi}} \left[\frac{(w_{A_{i-1}})_{\delta'_{i-1}} - (w_{A_{i-1}})_{\delta'_i}}{(w_{A_{i-1}})_{\delta'_i}} \right] \left(\frac{\exp(-\gamma'^2_i)}{\text{erf}(\gamma'_i)} \right) \\
& - \frac{1}{\sqrt{\pi}} \left[\frac{(w_{A_i})_{\delta'_i} - (w_{A_i})_{\delta'_{i+1}}}{(w_{A_{i-1}})_{\delta'_i}} \right] \left(\frac{\sqrt{\phi_i}}{\text{erf}(\gamma'_{i+1})} \right)
\end{aligned} \tag{B-12}$$

where ϕ_i is defined by Eq. 29. Since Eq. B-12 involves only γ_i and γ_{i+1} , the evaluation of γ_i is

much easier than in the case of Eq. 32. The numerical solution of Eqs. B-5 and B-7 was compared with the analytical solution given by Eqs. B-8 through B-12 for systems involving 2, 3, and 5 layers. The numerical results were in good agreement with the analytical results as shown in Fig. B-1, which compares the numerical and analytical results for a five-layer system. The parameter values used in the calculations shown in Fig. B-1 are given in Table B-1.

Figure B-1. Comparison of analytical and numerical solutions, using $\delta_i(t)$ as the frame of reference.

Table B-1: Parameter Values for Comparison of Analytical and Numerical Solutions, using $\delta_i(t)$ as the frame of reference

Layer	$(w_{A_i})_{\delta_i}$	$(w_{A_i})_{\delta'_{i+1}}$	ρ_i (g/cm ³)	$D_i \times 10^5$ (cm ² /day)	γ_i
1	1.0	1.0	1.0	0.0	---
2	0.8	0.7	1.0	3.0	-0.396825
3	0.6	0.5	1.0	3.0	0.126671
4	0.4	0.2	1.0	3.0	0.167156
5	0.1	0.1	1.0	3.0	0.519433

APPENDIX C: Program ILaG Flow Chart

APPENDIX D: ILaG Input File Description

The contents of the ASCII input file required by the user is defined below. Input is free-field format so that columns are unimportant but the number of fields per line can be important (i.e., Line 1 fields can be put on two lines however Line 2 fields cannot be appended to Line 1). Comments are allowable after the data required for a line has been supplied. A sample input deck is presented in Appendix E.

Lines 1 through 4 - these lines contain general problem definition and mesh information

Line 1 - four integers

- field 1: the geometry of the problem, currently must be 0
- field 2: the number of zones involved
- field 3: the number of components
- field 4: the index of the component that controls diffusion

Line 2 - $n_{\text{zone}}+1$ real numbers

- field 1: the coordinate of the left-most boundary of the domain
- field 2: the coordinate of the left-most internal interface
- .
- .
- field n_{zone} : the coordinate of the right-most internal interface
- field $n_{\text{zone}}+1$: the coordinate of the right-most boundary of the domain

Line 3 - n_{zone} integers

- field 1: the initial number of nodes in the left-most zone
- .
- .
- field n_{zone} : the initial number of nodes in the right-most zone

Line 4 - one integer, two real numbers

- field 1: the maximum level of mesh refinement (zero indicates the default value of six)
- field 2: mesh refinement tolerance (zero takes the default)
- field 3: mesh point removal tolerance (zero takes the default)

Lines 5 through 5+ n_{zone} - each line contains the diffusion coefficients for each component in the zone where zones are defined from left to right (i.e., zone 1 is the left-most zone)

Example: Line 5

- field 1: diffusion coefficient for component 1 in zone 1
- field 2: diffusion coefficient for component 2 in zone 1

.
.
field ncomp: diffusion coefficient for component ncomp in zone 1

Repeat the above line for each zone....

Next nzone-1 lines - each line contains the “fixed” concentration values for each component at the internal interface where internal interfaces are defined from left to right (i.e., internal interface 1 is the left-most interface)

Example: Line 1 of this set

field 1: the fixed concentration value on the left side of the first internal interface for component 1

field 2: the fixed concentration value on the right side of the first internal interface for component 1

field 3: the fixed concentration value on the left side of the first internal interface for component 2 field 4: the fixed concentration value on the right side of the first internal interface for component 2

.
.
field ncomp*2-1: the fixed concentration value on the left side of the first internal interface for the last component, ncomp

field ncomp*2: the fixed concentration value on the right side of the first internal interface for the last component, ncomp

Repeat the above line for each internal interface....

Next line - nzone real numbers defining the bulk density for each zone

field 1: the bulk density for zone 1 (left-most zone)

.
.
field nzone: the bulk density for zone nzone (right-most zone)

Next nzone lines - each line contains the initial concentration for each component in a zone

Example: Line 1 of this set

field 1: initial concentration of component 1 in zone 1

field 2: initial concentration of component 2 in zone 1

.
.
field ncomp: initial concentration of component ncomp in zone 1

Repeat the above line for each zone....

Next line - three real numbers, one integer to define simulation time and time step

field 1: initial time

field 2: final time

field 3: maximum time step (delta time)

field 4: maximum number of time steps

Next line - two real numbers to control output frequency

field 1: time of first write to print/plot file

field 2: increment of time to write to print/plot

APPENDIX E: Sample Input File

The following lines represent a sample input file. This particular case describes a 100%Sn-Cu system with two intermetallics as discussed in Section 5.0. The calculations are carried out to 400 days and are plotted in Figs. 10 and 11. A brief comment follows each input line; see Appendix D for a more detailed description.

0, 4, 2, 1	igeom(0=slab), nzone, ncomp, jcont
-1.0e-2, 0.0, 1.6e-4, 2.5e-4, 1.0e-2	xface (i.e., endpoints of zones)
10, 10, 10, 100	nnodes in each zone
0, 1.e-6, 0	ilevmx, rtolref, rtolrec ==> defaults
8.0e-7, 0.0	Dm for components in zone 1
0.0, 0.0	a for components in zone 1
0.0, 0.0	Dp for components in zone 1
1.0, 1.0	xk for components in zone 1
4.3e-8, 0.0	Dm for components in zone 2
0.0, 0.0	a for components in zone 2
1.4e-7, 0.0	Dp for components in zone 2
3.8e3, 1.0	xk for components in zone 2
0.5e-7, 0.0	Dm for components in zone 3
0.0, 0.0	a for components in zone 3
1.7e-7, 0.0	Dp for components in zone 3
2.1e3, 1.0	xk for components in zone 3
8.0e-7, 0.0	Dm for components in zone 4
0.0, 0.0	a for components in zone 4
0.0, 0.0	Dp for components in zone 4
1.0, 1.0	xk for components in zone 4
0.3, 0.0	kl (reaction rate coef) for components left end of zone 1
1.e-2, 0.0	kr (reaction rate coef) for components right end of zone 1
0.3, 0.0	kl (reaction rate coef) for components left end of zone 2
1.e-2, 0.0	kr (reaction rate coef) for components right end of zone 2
0.3, 0.0	kl (reaction rate coef) for components left end of zone 3
1.e-2, 0.0	kr (reaction rate coef) for components right end of zone 3
0.3, 0.0	kl (reaction rate coef) for components left end of zone 4
1.e-2, 0.0	kr (reaction rate coef) for components right end of zone 4

1.0, 0.623, 0.0, 1.0	wfix at left and right sides of first boundary for each component
0.605, 0.410, 1.0, 1.0	wfix at left and right sides of second boundary
0.400, 0.0, 1.0, 1.0	wfix at left and right sides of third boundary
8.9, 9.0, 8.37, 7.29	bulk density of each zone
1.0, 0.0	initial condition for zone 1
0.605, 1.0	initial condition for zone 2
0.4, 1.0	initial condition for zone 3
0.0, 1.0	initial condition for zone 4
1.0, 401.0, 0.1, 41000	initial time, final time, delta time, max steps
1.0, 5.0	time of first print/plot, increment of print/plot time

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