Infrared brazing of Ti$_{50}$Ni$_{50}$ shape memory alloy using pure Cu and Ti–15Cu–15Ni foils

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Abstract

Infrared brazing of Ti$_{50}$Ni$_{50}$ using two brazing filler metals was investigated in the study. Three phases, including Cu-rich, CuNiTi ($\Delta$) and Ti(Ni,Cu), were observed in the Ti$_{50}$Ni$_{50}$/Cu/Ti$_{50}$Ni$_{50}$ joint after brazing at 1150 °C. The Cu-rich phase was rapidly consumed in the first 10 s of brazing, and the eutectic mixture of CuNiTi and Ti(Ni,Cu) phases were subsequently observed in the joint. Samples brazed for longer time resulted in less CuNiTi and more Ti(Ni,Cu) phases in the joint. The existence of CuNiTi phase deteriorated the shape memory effect of the joint, but Ti(Ni,Cu) could still preserve shape memory behavior even alloyed with a large number of Cu. Therefore, higher shape recovery ratio was observed for specimens brazed for a longer time period. Extensive presence of Ti$_2$(Ni,Cu) phase was observed in Ti$_{50}$Ni$_{50}$/Ticuni$^b$/Ti$_{50}$Ni$_{50}$ joint upon brazing the specimens up to 1150 °C. The bending test could not be performed due to the inherent brittleness of Ti$_2$(Ni,Cu) matrix. Moreover, the stable Ti$_2$(Ni,Cu) phase was difficult to be removed completely by increasing either brazing time and/or temperature.

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1. Introduction

The importance of TiNi-based shape memory alloys (SMAs) is increasing due to their superior behavior in shape memory effect, superelasticity, and high corrosion resistance compared with many other SMAs [1–6]. The maturity of various manufacturing processes plays an important role in the applications of TiNi-based SMAs, and rolling, drawing as well as joining have been extensively studied [7–12].

Brazing is an economical fabrication method of complex assemblies among all joining methods [13]. Although there is no melting of the base metal during brazing, the heating can still affect the properties of the materials being joined [13]. It will be highly preferred that minimum heating is introduced during brazing in order to avoid deterioration of the base metal. The rapid infrared joining technique has been originally developed at the University of Cincinnati for high temperature materials [14]. Infrared brazing is a novel technique featured with rapid thermal cycle in comparison with the traditional furnace brazing [15]. With the aid of infrared energy generated by heating a tungsten filament in a quartz tube, the infrared furnace can provide rapid heating rate up to 3000 °C/min [15]. Therefore, the infrared brazing is highly suitable for studying the mechanism of early-stage reaction kinetics in the joint, and many reports are already available in literatures [16–19]. Infrared brazing has become an important method in studying the microstructural evolution of the brazed joint.

The primary challenge in brazing TiNi-based SMAs is to preserve its shape memory characteristics across the joined interface. Conventional furnace brazing with slow heating rates is probably not suitable in bonding TiNi-based SMAs, because the base metal experiences elevated temperatures for a long time. The reaction of Ti with other elements at elevated temperatures may deteriorate many featured properties of TiNi-based SMAs. Consequently, the investigation of brazing TiNi SMA is very limited [11].
It is reported that Ni atoms can be replaced by a large number of Cu atoms in Ti$_{50}$Ni$_{50}$ alloy and Ti$_{50}$Ni$_{50-x}$Cu$_x$ ($x \leq 30$ wt.%) SMAs can still exhibit shape memory behavior [20,21]. Consequently, both pure Cu and Ti$-15$Cu$-15$Ni foils are selected as brazing filler metals in joining Ti$_{50}$Ni$_{50}$ SMA. It is well known that the properties of the bonding interface between filler metal and base metal are strongly related to the interfacial microstructure. The morphology, quantity and species of the phases at the interface play crucial roles in evaluating the brazed joint. The main purpose of this study is aimed to infrared brazing Ti$_{50}$Ni$_{50}$ SMA using two filler metals, pure Cu and Ti$-15$Cu$-15$Ni, respectively. With the aid of fast infrared heating rate, the early-stage of microstructural evolution in the joint will be studied extensively. The shape recovery character of an infrared-joined Ti$_{50}$Ni$_{50}$ SMA is also discussed.

2. Experimental procedures

Ti$_{50}$Ni$_{50}$ in atomic percent was used as the base metal in this study. The master alloy was prepared by vacuum arc remelting using high-purity pellets under protective argon atmosphere. The TiNi ingot was firstly cut into approximate 10 mm $\times$ 5 mm $\times$ 1 mm specimens. Brazed surfaces of each TiNi specimen were polished by a SiC paper of 600 grit, and subsequently cleaned well in an ultrasonic bath prior to infrared brazing. Pure copper foil and Ticuni$^{\circ}$ foil of thickness 25 $\mu$m as well as 50 $\mu$m were used as the brazing filler metals. Ticuni$^{\circ}$ foil is a commercial product of Wesgo company, and its chemical composition in weight percent is 70Ti$-15$Cu$-15$Ni. Ticuni$^{\circ}$ is a clad-laminated brazing filler metal [22]. It has an inner Cu$_{50}$Ni$_{50}$ (wt.%) alloy layer and two pure outer Ti layers. The solidus and liquidus temperatures of Ticuni$^{\circ}$ foil are 910 and 960 $^\circ$C, respectively.

The brazing filler foil was sandwiched by two pieces of Ti$_{50}$Ni$_{50}$ base metal. To enhance infrared absorption of the brazed specimen, all specimens were clamped between two graphite plates, and a thermocouple was kept in contact with the specimen as shown in Fig. 1 to measure the temperature. An ULVAC SINKO-RIKO RHL-P610C infrared furnace with an Ar flow rate of 200 cm$^3$/min was used throughout the experiment. Table 1 summaries all process variables used in infrared brazing. The specimen was cut by a low speed diamond saw. A standard metallographic procedure was applied to the specimen before the microstructural observation. Their cross-sections were first ground by SiC papers, and subsequently polished by 0.3 $\mu$m alumina powder. Kroell’s reagent with 2 ml HF, 4 ml HNO$_3$ and 100 ml H$_2$O was selected as an etching solution. The etched cross-sections of the brazed specimens were examined using a Philips XL30 scanning electron microscope (SEM) with energy dispersive X-ray spectroscopy (EDS). Quantitative chemical analysis was also performed using a JEOL JXA-8600SX electron probe microanalyzer (EPMA) equipped with a wavelength dispersive spectrometer (WDS) operated at 15 kV with a probe current of 100 nA.

The shape memory effect (SME) was measured by the bending test to evaluate the shape recovery ratio of infrared brazed joint [23]. Fig. 2 displays a schematic diagram of the bending test. The initially flat brazed Ti$_{50}$Ni$_{50}$/Cu/Ti$_{50}$Ni$_{50}$ and Ti$_{50}$Ni$_{50}$/Ticuni$^{\circ}$/Ti$_{50}$Ni$_{50}$ specimens were machined into a size of approximately 20 mm $\times$ 5 mm $\times$ 0.5 mm. Then, it was bent into an angle $\theta_i$ at the temperature of 77 K (liquid nitrogen). After the temperature was raised to various temperatures of up to 130 $^\circ$C, $\theta_i$ was recovered to $\theta_f$, which was temperature dependent. The shape recovery ratio of the joint was determined by $(\theta_i - \theta_f)/\theta_i$. The detailed measurement of the shape-recovery characteristics has been presented elsewhere [23]. A pure Ti$_{50}$Ni$_{50}$ specimen with the same dimension of the above infrared brazed sample was also tested for the purpose of comparison.

<table>
<thead>
<tr>
<th>Temperature ($^\circ$C)</th>
<th>Ti$<em>{50}$Ni$</em>{50}$/Cu/Ti$<em>{50}$Ni$</em>{50}$ (s)</th>
<th>Ti$<em>{50}$Ni$</em>{50}$/Ticuni$^{\circ}$/Ti$<em>{50}$Ni$</em>{50}$ (s)</th>
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<tbody>
<tr>
<td>960</td>
<td>–</td>
<td>12</td>
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<tr>
<td>960</td>
<td>–</td>
<td>30</td>
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<tr>
<td>960</td>
<td>–</td>
<td>120</td>
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<tr>
<td>960</td>
<td>–</td>
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<td>1150</td>
<td>60</td>
<td>–</td>
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<tr>
<td>1150</td>
<td>300</td>
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</table>
3. Results and discussion

3.1. Infrared brazing of TiNi/Cu/TiNi Joint

The heating of substrate may result in changes of its properties, particularly if the alloy is heated above its annealing temperatures. Where mechanical properties are obtained by heat treatment, e.g. quench and temper, they may be altered by the brazing operation. In contrast, the alloy in the annealed condition will generally experience no appreciable change due to brazing [24]. It is expected that brazing has little effect on the mechanical properties of TiNi substrate. Additionally, one of the significant
base metal interactions which have been used in determining the behavior of brazed joint is alloying. The extent of interaction varies greatly depending on the compositions of the base metal, the brazing filler metal and thermal cycle [24]. A short thermal cycle can usually minimize the alloying effect during brazing. Compared with the traditional furnace brazing, infrared brazing is featured with a very rapid thermal cycle, so higher infrared brazing temperature will do less damage to the base metal. However, high brazing temperature can greatly speed up the microstructural evolution of the brazed joint in the study.

The melting point of pure Cu is 1084.9 °C, so 1150 °C was the chosen as the brazing temperature throughout the experiment. Fig. 3 displays backscattered electron images (BEIs) of the interface infrared-brazed specimen at 1150 °C for 2, 10, 30, 60 and 300 s. Numbers 1–7 shown in the figure are the chemical analyses of the selected areas, and the results are listed in Table 2. According to Table 2, there are three phases observed in the joint, including: Cu-rich phase, CuNiTi (Δ) phase and Ti(Ni,Cu) phase [25]. Residual Cu can be initially found in the joint, but it is completely consumed in 10 s as shown in Fig. 3(a) and (b). In Fig. 3(a), the central Cu-rich phase (marked by 1) is alloyed with minor Ti and Ni, and a CuNiTi phase (marked by 2) is intermediated between the Cu-rich phase and Ti50Ni50 base metal. The dissolution of Ti50Ni50 base metal into molten Cu-rich braze results in the formation of CuNiTi phase. The ternary CuNiTi (Δ) intermetallic compound with a stoichiometry Cu30Ni36.5Ti33.5 has been reported in the literature [25]. The microstructure displayed in Fig. 3(b) exhibits the eutectic structure of CuNiTi and Ti(Ni,Cu). It was reported that high solubility of Cu was found in Ti50Ni50, with Ni being replaced by Cu [20,21].

Fig. 4 shows the isothermal section of Cu–Ni–Ti system at 800 °C in atomic percent [25]. According to Fig. 4, TiNi has a huge solubility for Cu, and it is consistent with our experimental result. There are two major regions between TiNi and Cu in Fig. 4, including a region mixed with

Table 2
Chemical analyses of the selected areas in Fig. 3

<table>
<thead>
<tr>
<th>No.</th>
<th>Composition (at.%)</th>
<th>Remarks</th>
</tr>
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<tbody>
<tr>
<td>1</td>
<td>4.7Ti–2.6Ni–92.7Cu</td>
<td>Cu-rich phase</td>
</tr>
<tr>
<td>2</td>
<td>34.8Ti–35.5Ni–29.7Cu</td>
<td>CuNiTi (Δ) phase</td>
</tr>
<tr>
<td>3</td>
<td>34.8Ti–38.1Ni–27.0Cu</td>
<td>CuNiTi (Δ) phase</td>
</tr>
<tr>
<td>4</td>
<td>48.2Ti–40.8Ni–11.0Cu</td>
<td>Ti (Ni,Cu) phase</td>
</tr>
<tr>
<td>5</td>
<td>37.4Ti–39.0Ni–23.5Cu</td>
<td>CuNiTi (Δ) phase</td>
</tr>
<tr>
<td>6</td>
<td>48.2Ti–44.1Ni–7.7Cu</td>
<td>Ti (Ni,Cu) phase</td>
</tr>
<tr>
<td>7</td>
<td>50.6Ti–48.2Ni–1.2Cu</td>
<td>Ti (Ni,Cu) phase</td>
</tr>
</tbody>
</table>

Fig. 4. The isothermal section in the Cu–Ni–Ti system at 800 °C in atomic percent [25].
Ti(Ni,Cu) and CuNiTi (NiTi + Δ), and a region mixed with CuNiTi and Cu-rich phase (Δ + γ). The dissolution Ti₅₀Ni₅₀ substrate into the molten braze is strongly related to the final microstructure of the joint. With the increment of brazing time, both the diffusion of Cu atoms into Ti₅₀Ni₅₀ substrate and Ni atoms into the braze are greatly enhanced. The formation of CuNiTi phase in the braze results in the disappearance of Cu-rich phase in the joint, so the eutectic Ti(Ni,Cu) and CuNiTi phase, instead of eutectic CuNiTi and Cu-rich phase, is observed in the experiment. Additionally, the amount of CuNiTi phase in the joint is gradually decreased with the increment of brazing time, because the TiNi substrate has a huge solubility of Cu. The consumption of Cu from the CuNiTi phase cause shrinkage of CuNiTi in the infrared brazed joint as illustrated in Fig. 3(c)–(e).

According to the experimental observations, it is expected that CuNiTi phase can be completely dissolved into Ti₅₀Ni₅₀ substrate if the brazing time is long enough. The dissolution of the braze alloy (Cu), dissolution of the Ti₅₀Ni₅₀ into the Cu-rich melt, saturation of the melt, isothermal solidification of the molten braze alloy and homogenization of the brazed joint through diffusion of the solute atoms into the base metal [26]. It is similar to the process of transient liquid phase bonding (TLPB) [26–28]. The melting of Cu during infrared brazing is very rapid as demonstrated in Fig. 3(a). In contrast, the isothermal solidification and homogenization are not completed during infrared joining. Blue proposes an estimation of the time for isothermal solidification at various specified joining temperatures [26]. However, there is no further calculation in this research due to insufficient diffusion data of Cu and Ti atoms in Ti₅₀Ni₅₀ substrate. Accordingly, it needs further study in the future.

The infrared brazing Ti₅₀Ni₅₀ using pure Cu consists of several major stages, including: melting of the braze alloy (Cu), dissolution of the Ti₅₀Ni₅₀ into the Cu-rich melt, saturation of the melt, isothermal solidification of the molten braze alloy and homogenization of the brazed joint through diffusion of the solute atoms into the base metal [26]. It is similar to the process of transient liquid phase bonding (TLPB) [26–28]. The melting of Cu during infrared brazing is very rapid as demonstrated in Fig. 3(a). In contrast, the isothermal solidification and homogenization are not completed during infrared joining. Blue proposes an estimation of the time for isothermal solidification at various specified joining temperatures [26]. However, there is no further calculation in this research due to insufficient diffusion data of Cu and Ti atoms in Ti₅₀Ni₅₀ substrate. Accordingly, it needs further study in the future.

Table 3 summaries the result of the bending test. The shape recovery ratio of the original Ti₅₀Ni₅₀ strip is 95% at 80 °C, and it increases to 99% as the specimen heated to 130 °C. Higher recovery ratio can be obtained for all specimens heated to a higher temperature. According to Table 3, the recovery ratio of the specimen brazed for 300 s is better than that of specimens brazed for 60 s. There are two different thickness of the Cu foils used in the test, including 50 and 25 μm. Both thicknesses of the Cu fillers demonstrate similar tendency of the test.

According to the aforementioned result, specimens brazed for longer time result in less CuNiTi phase and more Ti(Ni,Cu) phase in the joint. It is reasonable to conclude that the existence of CuNiTi phase deteriorates the shape memory effect of the joint. Moreover, Ti(Ni,Cu) still preserves the shape memory behavior even though it is alloyed with Cu [20,21]. The amount of CuNiTi phase is greatly decreased in the joint as the brazing time is increased. Therefore, it is expected that

![Fig. 5. The SEIs and EPMA line scan profiles of the interface infrared brazed at 960 °C for (a) 12 s, (b) 120 s, and (c) 300 s.](image-url)
higher shape recovery ratio can be obtained for specimens brazed for a longer time period.

Specimens after bending test were subsequently examined by an SEM. It was confirmed that there was no cracks observed in the Ti50Ni50/Cu/Ti50Ni50 joint. Consequently, the use of pure Cu in brazing Ti50Ni50 has the potential in engineering applications.

3.2. Infrared brazing of Ti50Ni50/Ticuni8/Ti50Ni50 Joint

Fig. 5 displays the secondary electron images (SEIs) and EPMA line scan profiles of the interface infrared brazed at 960 °C for 12, 120 and 300 s. The brazed joint initially contains two phases, the matrix and the angular phase. Upon increasing the brazing time, only the matrix phase is left in the joint. Meanwhile, the line scan profiles demonstrate that the chemical composition of the braze becomes homogeneous upon increasing the brazing time. Chemical analysis using EPMA or EDS is necessary in order to further quantify these phases.

Fig. 6 displays SEM image and EDS chemical analyses of the specimen brazed at 960 °C for 30 s. It is noted that no interfacial reaction phase between Ti50Ni50 base metal and the braze alloy is observed. According to EDS analyses, the chemical composition of base metal is close to Ti50Ni50 as marked by 1 in Fig. 6. The base metal near the interface is alloyed with more Cu (marked by 2 in Fig. 6), and its stoichiometry is close to Ti50(Ni,Cu)50. Unlike aforementioned Ti50Ni50/Cu/Ti50Ni50 bonding, the current brazed joint is primarily comprised of Ti2(Ni,Cu) phase (marked by 3 in Fig. 6). Meanwhile, the chemical composition of the angular phase in the joint is still close to Ti2(Ni,Cu) as shown in Fig. 6 (marked by 4). The major difference between the matrix and the angular phase is the copper content in Ti2(Ni,Cu). The copper content in the matrix is lower than that in the angular phase. The Cu atoms can replace the Ni atoms in Ti2Ni phase, so Ti2(Ni,Cu) is formed during brazing.

Both Ti and Cu diffuse from braze alloy into Ti50Ni50, and Ni diffuses from Ti50Ni50 into braze driven by the concentration gradient of the joined alloys. According to Fig. 4, the isothermal section in the Cu–Ni–Ti system at 800 °C demonstrates that Ti2Ni phase can dissolve Cu up to 8 at% [25]. Additionally, there is no intermediate phase between Ti50Ni50 and Ti2Ni as displayed in Fig. 4. It is consistent with our experimental observations. The atomic composition of Ti–15Cu–15Ni is 74.8Ti–12.1 Cu–13.1Ni as marked by A in Fig. 4 [25]. Both Ti50Ni50 and Ti2Ni phases can dissolve Cu, so Ti(Ni,Cu) and Ti2(Ni,Cu) phases are observed in the experiment.
Fig. 7 shows the SEM images and EPMA chemical analyses of the Ti$_{50}$Ni$_{50}$/Ticuni$^\text{®}$/Ti$_{50}$Ni$_{50}$ joint at 1000 °C for 30 s. The result of EPMA chemical analyses is similar to that brazed at 960 °C for 30 s. Upon increasing the brazing temperature, the amount of angular phase in the joint decreases since the interdiffusion between the molten braze and Ti$_{50}$Ni$_{50}$ substrate is greatly enhanced at higher brazing temperatures. The consumption of Cu in the molten braze increases at 1000 °C. Consequently, the angular phase with higher Cu content is decreased. Fig. 8 shows the SEM images and EPMA chemical analyses of the Ti$_{50}$Ni$_{50}$/Ticuni$^\text{®}$/Ti$_{50}$Ni$_{50}$ joint at 1050 °C for 30 s. The angular phase is almost disappeared, and the width of Ti$_3$(Ni,Cu) matrix is further decreased. The depletion of Ti in the braze results in the diminishing of Ti$_2$(Ni,Cu) phase. Similar observation can be found for Ti$_{50}$Ni$_{50}$/Ticuni$^\text{®}$/Ti$_{50}$Ni$_{50}$ specimens brazed at 1100 °C for 30 s, as displayed in Fig. 9.

The bending test could not be performed due to the inherent brittleness of the infrared brazed joint. Ti$_{50}$Ni$_{50}$/Ticuni$^\text{®}$/Ti$_{50}$Ni$_{50}$ was broken at the joint after the specimen was bent. The direction of crack propagation was along the braze joint. It was resulted from the extensive presence of brittle Ti$_3$(Ni,Cu) matrix in the joint. In Ti$_{50}$Ni$_{50}$/Cu/Ti$_{50}$Ni$_{50}$ bonding, the joint was primarily comprised of eutectic Ti(Ni,Cu) and CuNiTi phase. It was reported that Ti(Ni,Cu) phase still preserved SMA even alloyed with ~30 wt% Cu [20,21]. However, Ti(Ni,Cu) phase was absent from Ti$_{50}$Ni$_{50}$/Ticuni$^\text{®}$/Ti$_{50}$Ni$_{50}$ joint after infrared brazing. Moreover, Ti$_3$(Ni,Cu) phase was still observed in Ti$_{50}$Ni$_{50}$/Ticuni$^\text{®}$/Ti$_{50}$Ni$_{50}$ for specimens infrared brazing up to 1150 °C. Consequently, the brittle and stable Ti$_3$(Ni,Cu) phase was difficult to be completely removed by increasing either brazing time or temperature.
4. Conclusions

1. Infrared brazing of Ti$_{50}$Ni$_{50}$/Cu/Ti$_{50}$Ni$_{50}$ was investigated in the study. Three phases, including Cu-rich, CuNiTi ($\Delta$) and Ti(Ni,Cu), were observed in the joint after infrared brazing at 1150 $^\circ$C. The Cu-rich phase was rapidly consumed within the first 10 s of brazing. Thereafter, eutectic mixture of CuNiTi and Ti(Ni,Cu) phases was observed in the bond. With the increase of brazing time, the amount of CuNiTi phase decreased due to the consumption of Cu in the joint.

2. The shape recovery ratio of the original Ti$_{50}$Ni$_{50}$ strip was 95% at 80 $^\circ$C and 99% at 130 $^\circ$C in the bending test. The shape recovery ratio of Ti$_{50}$Ni$_{50}$/Cu/Ti$_{50}$Ni$_{50}$ joint could reach 99% at 130 $^\circ$C for the specimen infrared brazed at 1150 $^\circ$C for 300 s. Samples brazed for longer time resulted in less CuNiTi phase and more Ti(Ni,Cu) phase in the joint. The existence of CuNiTi phase deteriorated the shape memory effect of the joint, but Ti(Ni,Cu) still preserved the shape memory behavior even alloyed with a large number of Cu.

3. Extensive presence of Ti$_2$(Ni,Cu) phase was observed in Ti$_{50}$Ni$_{50}$/Ticuni/Ti$_{50}$Ni$_{50}$ joint for specimens infrared brazed up to 1150 $^\circ$C for 30 s. The bending test could not be completed due to the inherent brittleness of Ti$_2$(Ni,Cu) matrix in the joint. Moreover, the stable Ti$_2$(Ni,Cu) phase was difficult to be removed by increasing either the brazing temperature or time.

Acknowledgements

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References