Surface modification of resistance welding electrodes by electro-spark deposited composite coatings
Part II. Metallurgical behavior during welding

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Abstract

Electrodes with monolithic TiC\(_p\)/Ni coatings and multi-layer Ni/(TiC\(_p\)/Ni)/Ni coatings were used to resistance spot weld Zn-coated sheet steel to investigate metallurgical behaviour of the coatings during welding. Scanning electron microscopy, energy-dispersive X-ray analysis and X-ray diffraction were employed to characterize the microstructure of coatings, reactions of electrodes with Zn-coating and alloy layer formation. The results showed that molten Zn penetrated TiC\(_p\)/Ni coatings via the cracks that were present within as-coated TiC\(_p\)/Ni coating, starting from the first weld. Additional cracks continually formed in the coating during welding due to action of the welding force on the low toughness coating, resulting in formation of a granular loose overlay at the outer surface which were easily detached and stuck onto the work sheet. On the contrary, cracks could be rarely found within Ni/(TiC\(_p\)/Ni)/Ni coating until 100 welds or more were made, and much fewer cracks formed up to 400 welds, compared to the TiC\(_p\)/Ni coating. With Ni/(TiC\(_p\)/Ni)/Ni coating on the electrode surface, alloying between copper alloy and molten Zn as well as pitting (erosion) of the electrode tips were remarkably reduced, and hence, a slower growth rate of tip diameter was observed.

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1. Introduction

The use of zinc-coated steels has attracted much attention over the past decade owing to their good corrosion resistance and relatively low cost. Resistance spot welding is the primary method for joining sheet steel components in the automotive industries. However, the zinc coating has resulted in serious welding problems due to its lower electrical resistance and melting temperature, and accelerated degradation of resistance welding electrode tips. This leads to a drastic reduction in the electrode life. A short electrode life limits the rate of production due to the need for frequent electrode dressing and/or electrode changes. Electrode degradation during resistance welding of coated steels has been the subject of many studies, in which, for example, developments in electrode materials and design of electrode have been explored in an effort to improve weldability and increase electrode life \([1–10]\). It is commonly believed that Zn coatings will react with the electrodes to form brass alloys, causing electrode sticking due to local bonding between the electrode and workpiece. Fracture of these local bonds can lead to stripping of parts of the electrode surface causing progressive erosion and/or pitting which eventually causes growth of the tip diameter and, hence, the failure of the electrode.

On the other hand, coating the electrode surfaces has been suggested as a method by which to extend electrode life. For example, Dong and Zhou \([11]\) have shown that an electrode with electro-spark deposited TiC\(_p\)/Ni composite coating can extend the life of micro-resistance welding electrodes. Their tests found the coating increases tip life by approximately 70% by reducing the amount of local bonding between electrode and Ni sheet. More recently, in addition, the use of the TiC\(_p\)/Ni
composite coating for electrode life improvement in resistance spot welding of hot-dip galvanized zinc coated steel has also been reported and shown that TiCp/Ni coating acts as a barrier against erosion and alloying of electrode by molten Zn [12]. The TiCp/Ni composite coating, however, needs further improvement as a barrier layer due to the presence of cracks within the coating and delamination at the coating–substrate interface [13]. A modified coating approach has been described in Part I of the present work, where it was demonstrated that a dense and crack-free Ni/(TiCp/Ni)/Ni coating could be deposited on electrode surfaces by a multi-layer electro-spark deposition process [13].

In the present work, the metallurgical behavior during resistance welding of Zn-coated steel using both monolithic TiCp/Ni and multi-layer Ni/(TiCp/Ni)/Ni deposited electrode coatings has been investigated.

2. Experimental

Electrodes were coated by electro-spark deposition either using TiCp/Ni sintered composite rod (42 vol.% of TiC particle and 21 wt.% of Ni with addition of Mo and Co) to apply a monolithic TiCp/Ni layer or using TiCp/Ni and Ni rods to make a three-layer Ni/(TiCp/Ni)/Ni coating, as described in Part I. The thickness of the coatings was around 35μm. Welding tests of electrodes with and without coating were carried out using a 250kV A single-phase AC spot welding machine. Hot dip galvanized (HDG60G) mild steel (0.7mm thick with 0.01mm thick Zn-coating on both side) was employed as worksheets. Welding test parameters used in this work are listed in Table 1 in order to investigate and compare the degradation process of electrodes with and without coating under same initial heat generation condition. As can be seen from Table 1, the welding current for coated electrodes was reduced to obtain a similar starting nugget diameter, which means that coated electrodes can generate the same heat at a lower current as that generated using uncoated electrodes at a higher current. Under such conditions, the growth rate of electrode tip diameter during welding was monitored as an evaluation parameter of electrode performance, determined by a carbon imprint technique in which a copy paper together with a carbon paper were put on the surface of a sheet steel to record the shape of electrodes top surface when electrodes contact with the sheet steel under pressure [12]. In order to investigate the metallurgical behavior of coatings during welding, after certain numbers of welds, tip surfaces and cross-sections of sacrificed electrodes, as well as worksheet weld surfaces were examined using a JEOL JSM-840 scanning electron microscope (SEM) equipped with energy-dispersive spectrometer (EDS). Phase identification was conducted using a Siemens D500 X-ray diffractometer (XRD) with Cu Kα radiation operating at 30kV and 40mA.

3. Results

3.1. Electrode tip growth during welding

Electrode tip growth has recently been suggested as the dominant process that determines electrode life when resistance spot welding Zn-coated steels [3]. It is also known that in terms of electrode wear rate, the change in the electrode tip diameter with increasing number of welds is more important than the absolute value of electrode face diameter itself [6]. Accordingly, the rate of tip diameter growth of electrodes without and with monolithic TiCp/Ni coating or multi-layer deposited Ni/(TiCp/Ni)/Ni coating during welding was first investigated in the present work. As shown in Fig. 1, the tip diameter growth rate of the electrode with monolithic TiCp/Ni coating was similar to that of the electrode without coating before 100 welds. However, the former was much less than the latter when number of welds exceeded 100. On the contrary, the electrode with multi-Ni/(TiCp/Ni)/Ni coating showed smaller tip diameter growth rate than that with monolithic TiCp/Ni coating up to 400 welds.

3.2. XRD analyses

X-ray diffraction analysis was performed on the electrode tip surface before and after welding testing. Strong peaks of TiC were found from the X-ray diffraction pattern of electrode with as-deposited monolithic TiCp/Ni coating, as shown in Fig. 2. After 400 welds, however, the intensity of TiC peaks significantly reduced, indicating loss of TiCp/Ni coating during welding due to sticking and removal to the sheet. In addition, a

![Fig. 1. Growth of electrode tip diameter as a function of weld number.](image-url)
the TiCP/Ni coating was similar to a brazing process that
metallurgically integrated and dense TiCP/Ni coating was no longer present, leaving non-continuous and separate loose TiCP/Ni coating islands on the surface of the substrate. The thickness of Cu–Zn alloy layer increased to 30 μm, and cracks had formed within Cu–Zn alloy layer due to its brittleness.

3.4. Metallurgical phenomena of multi-layer deposited Ni/(TiCP/Ni)/Ni coating

The surface photographs of electrode with multi-Ni/(TiCP/Ni)/Ni coating after 1 weld are shown in Fig. 11, along with the EDX element maps of Ti, Ni and Zn. Comparing to Figs. 4 and 5, Zn was found randomly deposited on the surface of multi-Ni/(TiCP/Ni)/Ni coating, no penetration of molten Zn into the coating could be seen owing to the absence of cracks within the coating. After 10 welds, the multi-Ni/(TiCP/Ni)/Ni coating was still dense and did not show any cracking. However, cracks were detected within the Ni/(TiCP/Ni)/Ni coating after welding.

β-Cu–Zn alloy was observed, resulting from the reaction between Zn and electrode copper alloy substrate. Fig. 3 exhibits the XRD patterns of the electrode with multi-layer Ni/(TiCP/Ni)/Ni coating before and after 400 welds. Different from the monolithic TiCP/Ni coating, the predominant phases of the as-deposited Ni/(TiCP/Ni)/Ni coating were Ni and TiC. After 400 welds, the intensity of Ni remarkably decreased, indicating loss of the top Ni layer. However, Ni/(TiCP/Ni) coating was still maintained on the electrode surface, as evidenced by strong TiC peaks. Moreover, there was no evidence that any reaction of the copper alloy substrate with the molten Zn had taken place. On the other hand, Zn and Fe–Zn compound were present on the electrode surfaces after welding.

3.3. SEM observation of monolithic TiCP/Ni coating

Figs. 4 and 5 show surface photographs of an electrode with monolithic TiCP/Ni coating after 1 weld, along with the EDX element maps of Ti, Cu, Fe and Zn. It is worth noting that only 1 weld sufficed to cause local loss (pitting) of TiCP/Ni coating (Fig. 4) and penetration of molten Zn via cracks into the coating (Fig. 5). Fig. 6 illustrates a low-magnification surface photograph of an electrode with monolithic TiCP/Ni coating after 100 welds. It can be clearly seen from Fig. 6 that much TiCP/Ni coating had been removed, allowing molten Zn and/or Zn–Fe to directly react with Cu to form brass alloys. The high-magnification SEM photographs of the electrode surface after 100 welds demonstrated that the initial cracks within the TiCP/Ni coating had been sealed by penetrated and solidified Zn containing Fe (Fig. 7). The sealing of cracks in the TiCP/Ni coating was similar to a brazing process that might prevent TiCP/Ni coating from removal due to sticking. On the other hand, comparing the diameter of islands within as-deposited TiCP/Ni coating (~250 μm), the most important finding was that the diameter of islands reduced to 50–80 μm after 100 welds, indicating ongoing cracking of the coating due to its low toughness and action of welding force upon each weld event. When welding was continued to 400 welds, the diameter of uncracked islands decreased to 30–60 μm, and cracks were again found between sealed Zn–Fe alloy and TiCP/Ni coating, as shown in Fig. 8.

Fig. 9 shows the cross-section of an electrode with TiCP/Ni coating after 100 welds. The average thickness of TiCP/Ni coating decreased to about 20 μm from its original thickness (approximately 35 μm). It is interesting to note that the TiCP/Ni coating could be divided into two different regions. (i) The region close to the outer surface marked by “A” was granular and loose in which some cracks had been formed. (ii) The region adjacent to copper alloy substrate marked by “B” was relatively dense as compared with region “A.” But some grooves that had been sealed by Zn–Fe alloy existed within the region “B.” In addition, Fig. 9 clearly shows that Zn mainly penetrated through the cracks formed in the granular TiCP/Ni layer and the grooves in the dense layer to react with copper to form a 12-μm-thick Cu–Zn alloy layer underneath the TiCP/Ni coating (layer “C” in Fig. 9). Fig. 10 illustrates the cross-section of an electrode with TiCP/Ni coating after 400 welds. The integrated and dense TiCP/Ni coating was no longer present, leaving non-continuous and separate loose TiCP/Ni coating islands on the surface of the substrate. The thickness of Cu–Zn alloy layer increased to 30 μm, and cracks had formed within Cu–Zn alloy layer due to its brittleness.
Fig. 4. SEM surface image of electrode with TiCp/Ni coating after 1 weld showing loss of coating (a) and corresponding EDX element maps of Ti Kα (b), Cu Kα (c) and Zn Kα (d).

Fig. 5. SEM surface image of electrode with TiCp/Ni coating after 1 weld showing penetration of molten Zn containing Fe (a) and corresponding EDX element maps of Ti Kα (b), Fe Kα (c) and Zn Kα (d).
Fig. 6. Low-magnification SEM surface image of electrode with TiCp/Ni coating after 100welds showing removal of coating (a) and corresponding EDX element maps of Ti K$_{α}$ (b), Cu K$_{α}$ (c) and Zn K$_{α}$ (d).

Fig. 7. High-magnification SEM surface image of electrode with TiCp/Ni coating after 100welds (a) showing sealing of cracked coating by penetrated Zn–Fe (indicated by arrows) and corresponding EDX element maps of Ti K$_{α}$ (b), Fe K$_{α}$ (c) and Zn K$_{α}$ (d).
Fig. 8. SEM surface image of electrode with TiCp/Ni coating after 400 welds (a) showing cracking at coating-sealed Zn–Fe interface and corresponding EDX element maps of Ti Kα (b), Fe Kα (c) and Zn Kα (d).

Fig. 9. SEM cross-section of electrode with TiCp/Ni coating after 100 welds (a) showing granular and loose layer (A), dense layer (B) and Cu–Zn alloy layer (C) and EDX element maps of Ti Kα (b), Cu Kα (c) and Zn Kα (d).
Fig. 10. SEM cross-section of electrode with TiC<sub>p</sub>/Ni coating after 400 welds (a) and EDX element maps of Ti K<sub>x</sub> (b), Fe K<sub>x</sub> (c) and Zn K<sub>x</sub> (d).

Fig. 11. SEM surface image of electrode with multi-layer Ni/(TiC<sub>p</sub>/Ni)/Ni coatings after 1 weld (a) and EDX element maps of Ti K<sub>x</sub> (b), Ni K<sub>x</sub> (c) and Zn K<sub>x</sub> (d).
to 100 welds (Fig. 12). In addition, similar to the monolithic TiCp/Ni coating, molten Zn containing Fe penetrated into the coating through the newly formed cracks (Fig. 12). It is worth noticing that a large percentage of Ni/(TiCp/Ni) coating still covered the electrode surface and the diameter of newly formed islands between cracks was relatively large (150–200 μm) compared to that of the TiCp/Ni coating at 100 welds (Fig. 7). When welding to 400 welds, it was found that the diameter of islands decreased to 100–180 μm and width of cracks also increased, revealing continuous cracking of Ni/(TiCp/Ni)/Ni coating during welding. However, the Ni/(TiCp/Ni)/Ni coating still remained on the surface even after it cracked.

Fig. 13 shows the cross-section of an electrode with Ni/(TiCp/Ni)/Ni coating after 100 welds. Except for rarely observed cracking within the coating, the coating was dense and adhered to the substrate. Zn was mainly present on the coating surface and did not show much diffusion into the substrate to form Cu–Zn alloy. Consistent with surface observation, SEM cross-section of electrode with Ni/(TiCp/Ni)/Ni coating after 400 welds demonstrated that the coating started to develop cracking (Fig. 14). In addition, delamination at coating–substrate interface was also visible (Fig. 14). As a result, Zn was found to penetrate and diffuse into the coating mainly through the cracks and delamination. However, different from the monolithic TiCp/Ni coating, the multi-layer Ni/(TiCp/Ni)/Ni coating did not exhibit the granular and loose structure up to 400 welds and showed only limited cracking and delamination. Particularly, Zn seemed to have more difficulty to diffuse through the Ni layer than TiCp/Ni coating. Consequently, no visible Cu–Zn alloy layer could be found. Table 2 summarizes the difference in metallurgical behaviours of monolithic TiCp/Ni coating and multi-layer Ni/(TiCp/Ni)/Ni coating before and after certain numbers of welds and clearly shows the benefits of multi-Ni/(TiCp/Ni)/Ni coating in preventing the coating from cracking and the substrate from alloying with Zn.

EDX analyses were used to determine the amount of coating materials remaining on the surface of the electrodes and the amount of electrode materials including coating and copper substrate removal to the sheet. As shown in Table 3, consistent with the XRD and SEM results, although the TiCp/Ni coating had much higher TiC content than the multi-layer Ni/(TiCp/Ni)/Ni coating prior to welding, it showed a much faster loss rate during welding due to the formation of a granular loose coating which is easily pulled off by adhesion to the workpiece.

4. Discussion

It is well known that resistance spot welding electrodes must possess (1) high electrical conductivity to minimize electrode heating; (2) high thermal conductivity to dissipate heat from the contact area between the electrode and the worksheet; (3) high hot strength to resist deformation caused by the application of high welding forces; and (4) ability to provide resistance to alloying. As a part of the electrode material after deposition, the composite coatings have the advantages of (3) and (4) mentioned above, especially acting as barrier layer between molten Zn and copper alloy substrate to arrest formation of Cu–Zn alloy [12]. This benefit of composite coating is similar to the
formation of Al\textsubscript{2}O\textsubscript{3} layer on the top surface of an aluminum oxide dispersion strengthened copper, which prevents the Cu–Zn alloy layer from spalling and suppresses the deposition of Zn–Fe on the electrode face and hence alloying between Zn and copper [14].

However, there are many cracks within as-deposited monolithic Ti\textsubscript{Cp}/Ni coatings and delamination at coating–substrate interface. Consequently, molten Zn penetrates into the coating via those defects (Fig. 5) and local removal of Ti\textsubscript{Cp}/Ni coating due to sticking especially at delamination regions is also found from the first weld (Fig. 4). Continued welding causes further alloying between copper and penetrated Zn (layer “C” in Fig. 9) as well as cracking of the coating, resulting in the formation of surface granular layer (layer “A” in Fig. 9) that was then removed due to surface sticking (Fig. 10). It is thought that penetration of molten Zn has two different effects on coated electrodes. On one hand, a negative effect is accelerated formation of a Cu–Zn alloy layer underneath the coating, starting from very early stages of welding, which may accelerate degradation of electrodes during their use. On the other hand, as a positive effect, solidification of penetrated molten Zn (Fe) brazes the cracked Ti\textsubscript{Cp}/Ni coating together and helps adhesion of the coating to the substrate, which helps keep the coating on the surface to protect the Cu–Zn alloy from being lost to the sheet. As a result of these effects, electrodes with Ti\textsubscript{Cp}/Ni coating show a fast tip growth up to 100welds due to fast loss of the coating; whereas a slow tip growth process occurs after 100welds due to increased difficulty of loss of Zn-brazed coating (Fig. 1). A secondary effect is also possible due to reduced current density due to the growth of contact diameter and hence less heat generation. More importantly, however, the granular microstructure formed on the outer surface of the Ti\textsubscript{Cp}/Ni coatings...
Ni coating (Fig. 9), which might be caused by significant cracking of the coating due to thermal and impact fatigue, causes loss of the key feature of a dense coating as a barrier layer. Consequently, when sticking occurs, the granular TiCp/Ni coating will be easily removed to the worksheet surface. In addition, cracks are again formed at the TiCp/Ni coating-solidified Zn interface during continued welding probably due to continual mechanical and thermal impact on the electrode surface (Fig. 8). Progressive granulation and removal of the TiCp/Ni coating eventually results in almost all the coating being lost onto the worksheet surfaces. At this stage, the coated electrode behaves like an uncoated electrode.

The hardness of the as-deposited TiCp/Ni coating is about HV 1000. In general, the harder the composite, the lower the toughness. As a result, many cracks form within the as-deposited TiCp/Ni coating due to its low toughness and operational stress during deposition [13]. In addition, during welding, the TiCp/Ni coating further cracks upon mechanical and thermal impact, and eventually develops a fractured

**Table 2**
Summary of different metallurgical behaviors of TiCp/Ni coating and Ni/(TiCp/Ni)/Ni coatings after certain numbers of welds

<table>
<thead>
<tr>
<th>Coatings</th>
<th>Diameter of islands (μm)</th>
<th>Depth of Zn penetration and diffusion (μm)</th>
<th>Cu–Zn alloy layer (μm)</th>
<th>Density of coating</th>
</tr>
</thead>
<tbody>
<tr>
<td>TiCp/Ni coating</td>
<td>As-coated 250</td>
<td>50–80</td>
<td>25–35</td>
<td>Dense</td>
</tr>
<tr>
<td></td>
<td>100 welds 50–80</td>
<td>25–35</td>
<td>12</td>
<td>Granular</td>
</tr>
<tr>
<td></td>
<td>400 welds 30–60</td>
<td>28–45</td>
<td>30</td>
<td>Granular</td>
</tr>
<tr>
<td>Ni/(TiCp/Ni)/Ni</td>
<td>As-coated None</td>
<td>None</td>
<td>10–15</td>
<td>Dense</td>
</tr>
<tr>
<td>Ni coatings</td>
<td>100 welds 150–200</td>
<td>10–15</td>
<td>Little</td>
<td>Dense</td>
</tr>
<tr>
<td></td>
<td>400 welds 100–180</td>
<td>15–24</td>
<td>Little</td>
<td>Dense</td>
</tr>
</tbody>
</table>
granular microstructure. Such significantly cracked and granular TiCP/Ni coating is no longer able to act as a barrier layer to prevent Zn from penetration and alloying with copper.

On the contrary, the multi-layer Ni/(TiCP/Ni)/Ni coating contains more Ni than monolithic TiCP/Ni and, hence, shows a medium hardness around HV 500 [13]. As discussed in Part I of this work [13], increasing the Ni content, which is a metal with high ductility and toughness acting as an excellent binder for TiC composite, would increase the toughness of the overall Ni/ (TiCP/Ni)/Ni coating. As a result, a crack-free and continuous Ni/(TiCP/Ni)/Ni coating was obtained [13]. Consequently, upon welding, Ni/(TiCP/Ni)/Ni coating provided a better barrier action than the TiCP/Ni coating, especially before 100 welds (Figs. 12 and 13): with the dense Ni/(TiCP/Ni)/Ni coating maintained on the substrate surface, no formation of Cu–Zn alloy layer was possible. However, continuing to weld after 100 welds eventually resulted in diffusion and penetration of Zn into the Ni/(TiCP/Ni)/Ni coating. The composition (wt.%) of the central Ni/(TiCP/Ni)/Ni coating region at 400 welds (indicated by area A in Fig. 14) was determined by EDX to be 25.5Ti–29.7Ni–30.6Zn–12.9Cu. According to the Ni–Zn and Cu–Zn phase diagrams, some Zn might have reacted with Ni and/or Cu during the welding process to form Ni–Zn, Cu–Zn or Ni–Cu–Zn compounds, for example, NiZn (β phase) [15]. Such compound phases were not detected in XRD probably due to their low content below the detection limit of the XRD diffractometer (Fig. 3). The formation of brittle intermetallic compounds would decrease the toughness of the Ni/(TiCP/Ni)/Ni coating and, hence, would cause cracking of the coating after 100 welds (Figs. 12 and 13). However, much fewer cracks formed within the Ni/(TiCP/Ni)/Ni coating compared to the TiCP/Ni coating, and no granulation of the coating occurred up to 400 welds, demonstrating the better functioning of the former coating as a barrier to delay the formation of Cu–Zn alloy layers. It appears that the toughness of the coating is more critical than its hardness. A dense coating with relatively high toughness may have the ability to reduce or avoid cracking of the coating during welding and hence acts as a good barrier layer for a longer period.

From a different viewpoint, Tanaka reported a relatively long electrode life when resistance-welding Zn–Ni-coated sheet comparing to Zn-coated sheet, owing to disturbance of the formation of Zn–Cu–Fe alloy at the tip surface, thus delaying diffusion of Zn in the electrode [9]. Such beneficial influence of addition of Ni into a plated layer may also be significant when the coating contains high Ni content, such as the multi-layer Ni/ (TiCP/Ni)/Ni coating, showing no visible formation of Cu–Zn alloy and much slower diffusion of Zn into the coating (Fig. 14).

It has been shown that material loss from electrodes (i.e., pitting or erosion) is one of the primary reasons leading to the growth of electrode tips. As a result, electrodes with multi-layer Ni/(TiCP/Ni)/Ni composite coating demonstrated slower tip diameter growth rate than electrodes coated with monolithic TiCP/Ni owing to its dense and less cracked state as-deposited and, hence, much lower removal to the work sheets (Fig. 1).

5. Conclusions

1. Molten Zn penetrated into the monolithic TiCP/Ni coating via the cracks that were present within the as-deposited coating even from first weld.
2. The monolithic TiCP/Ni coating continued to crack during welding due to action of the welding force and its low toughness, resulting in the formation of a granular broken structure near the outer surface that was easily removed by worksheet adhesion and the formation of Cu–Zn alloy layers underneath the coating.
3. The multi-layer Ni/TiC–Ni/Ni coating showed much less cracking and, hence, formed a better barrier against Zn penetration and reaction with the copper electrodes.
4. With a multi-layer Ni/(TiCP/Ni)/Ni coating on the electrode surface, pitting (erosion) of electrode was remarkably reduced, and hence, a slower growth rate of tip diameter was observed.

References