Micro-porosity Formation on the Underside of Plasma Sprayed Nickel Splats: A Microstructural Study

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Abstract:
The gas (micro) porosity formation in splats produced by atmospheric plasma spraying was microscopically studied through a new approach. The aim was to prove hydrogen content in plasma gas mixture propagates micro-porosity to the splat-substrate interface. Nickel powder was sprayed under argon plus 0, 7, 10, and 13 vol.% hydrogen as a secondary plasma gas on a stainless steel substrate, and the obtained microstructures of the underside splats were investigated by a scanning electron microscope. It was shown that by increasing hydrogen concentration in plasma gas mixture, more micro-pores formed in the microstructures, while pores content reached to a saturation level at higher hydrogen concentrations.

Keywords: Atmospheric plasma spray; Micro-porosity formation; Microstructure; Hydrogen gas content; Nickel splats.

Introduction

Atmospheric plasma spraying (APS) is one of the most versatile and rapid techniques to deposit protective metal or nonmetallic coatings onto the surface of components to enhance its properties under severe conditions, improve surface porosity of implants, fabricating low porosity metallic scaffolds and bioceramic coatings on the surface of bioinert metals. The fundamental of the process is heating up the powders above their melting point in a plasma jet and accelerate them toward a substrate. The coating will be formed by the build-up of splats resulting from the impact, flattening, and solidification of melted droplets on the substrate [1]. The coating important properties such as level and distribution of porosity are strongly affected by the in-flight particle conditions [2]. The process has long been considered to be an effective rapid solidification processing, since it allows instant quenching during coating buildup. The microstructures of plasma-sprayed deposits are also ultimately based on the solidification of many individual molten droplets [3]. APS coatings structure usually contain some level of porosity, up to ~10%, some unmelted or partially melted particles, fully melted and deformed “splats,” metastable phases, and oxides.

In APS, a plasma gas mixture, which consists of a primary plasma forming gas which is usually argon and a secondary gas, such as hydrogen or helium [1, 4], is introduced into the plasma torch. A high intensity direct current (DC) arc is
generated between the tip of the cathode and the cylindrical anode. A feedstock powder is then injected into high enthalpy, high temperature and velocity jet of partially dissociated and ionized gases emerging from the nozzle. The role of the secondary gas is to enhance the thermal conductivity of plasma for effective heating and melting of the injected powders.

Due to the high temperature and surface area of the particles, any changes in the process parameters can result in significant changes in the particle properties and consequently in the microstructure of the coating [1]. Inspection of the underside of splats often reveals the presence of small gas pores. These pores could be either due to heterogeneous nucleation of absorbed gases in the melt or the result of gas or vapor entrapment under the splat [5]. The size of these pores, however, depends on the mechanism by which they were formed. Entrapment of gas will generate pores which are large (> 1 micron) whereas heterogeneous nucleation generates micro-pores (<< 1 micron). In this paper, we are investigating the latter case, i.e., the heterogeneous nucleation on the underside of the splats.

During its flight towards the substrate, a molten droplet entrains and dissolves plasma gases and the surrounding air. Upon the high velocity impact (>100 m/s) a high transient hydrostatic pressure (approaching 1GPa) established that may rise within ~10³–10⁷ s under the droplet [56]. During the depressurization, there will be a point when the absorbed gases will be in supersaturated state and will nucleate. Micro (gas) porosity is generally caused by the evolution of gases during the solidification process. Generally, the solubility of gases in the solid is significantly less than that in the liquid state, and the gases may, therefore, be nucleated during solidification [7]. The solubility of gases in liquids increases nonlinearly with temperature [8]. For nickel and some other metals, a discontinuous and abrupt decrease is observed in the solubility at its melting point of 1453°C (Fig. 1). This will result in a significant generation of micro-pores [8, 9, 10, 11] level of porosity depends on the amount of dissolved gases in the liquid, the type of the metal, chemical kinetics, and the surface tension [7].

The major objective of the present work was to study the effect of hydrogen concentration in the plasma gas mixture on micro-porosity formation on the underside of nickel splats. Commercially pure nickel powder was selected as a model in our experiments since it is main component of many nickel based superalloys used in various high temperature applications. During its flight, the molten nickel particle will absorb some of the hydrogen present in the jet. Upon impact on the substrate, heterogeneous nucleation during solidification is expected to result in generation of micro-pores. This should lead to better understanding and control of micro-porosity formation in the splats, where porosity can be either undesirable or beneficial [12].

**Material and Methods**

Nickel powder (type 12313, Westaim Corp., Edmonton, Alberta, Canada) was used as feedstock material. The initial powder was sieved to make more uniform particle size distribution with the range -45+25 µm. Coatings 200-300 µm thick were deposited on a roughened (0.74µm ±0.04 ) stainless steel substrates. Roughening was done by 120 grid sand paper manually. In order to suppress splashing of the impacting molten droplets, and time providing for heterogeneous nucleation of micro-pores during the rapid solidification of splats, the substrate temperature was kept at 300°C by an electric heater. The substrate temperature was measured with a K-type thermocouple.

Nickel powders were deposited by SG-100 DC plasma torch (Praxair/TAF) sprayed under four different compositions of plasma gases of pure argon and its mixtures with 7, 10, and 13 vol. % of hydrogen. All experiments were done at
120 mm spray distance. During experiments the torch input power was changed in a manner so that particle temperatures remained the same for different H$_2$ content to provide similar condition for the nickel particles.

Schematic representation of the experimental set up is represented in Fig. 2.

![Fig. 2 Schematic of experimental set up](image)

The operating parameters of the torch are listed in Table 1.

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<td>79</td>
<td>52</td>
<td>8</td>
<td>52</td>
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In-flight particle conditions during experiments, i.e., temperature, diameter, and velocity upon impact shown in Table 2, were measured with a DPV-2000 monitoring system (Tecnar Ltée, Montreal, Quebec, Canada).

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<th>Sample #</th>
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<th>III</th>
<th>IV</th>
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<tbody>
<tr>
<td>Vol.% of H$_2$ in plasma gas mixture</td>
<td>0</td>
<td>7</td>
<td>10</td>
<td>13</td>
</tr>
<tr>
<td>Diameter μm</td>
<td>29 ±4.9</td>
<td>28 ±4.7</td>
<td>27 ±4.6</td>
<td>28 ±4.5</td>
</tr>
<tr>
<td>Temperature °C</td>
<td>2360 ±84</td>
<td>2339 ±92</td>
<td>2403 ±89</td>
<td>2326 ±91</td>
</tr>
<tr>
<td>Velocity m/s</td>
<td>125 ±12</td>
<td>102 ±16</td>
<td>119 ±17</td>
<td>93 ±15</td>
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</tbody>
</table>

The splats were removed from the substrates and their underside, i.e., the interface between the splat and the substrate, were studied under scanning electron microscope (SEM) with EDS attachment TM-3000 from Hitachi. The hydrogen content of splats was also measured, by Instrumental Gas Analysis (IGA) (Horiba EMGA-621W). Depending on the Relative Standard Deviation (RSD) obtained, 3 to 5 analytical measurements were done on each sample.

**Results**

**Nickel Particle Feedstock**

Fig. 3 shows the morphology of the nickel powder feedstock.

![Fig. 3 Nickel powder particles used for plasma spraying](image)

The nodularity of the particles were measured by image analysis and an average aspect ratio of 1.29 ± 0.18 (about 30% deviation from perfect sphere) were obtained. The better nodularity of the nickel particles and size uniformity provide the enhanced flow ability of the powder into the plasma jet, and higher chance for them to be at similar conditions upon impact [13].

**Porosity on Splat Microstructures**

Fig. 4 shows the microstructure of nickel splat-substrate interface sprayed in pure argon at two different magnifications. Although this case corresponds to pure argon, some micro-porosities are still formed. Our measurements of in-flight particle temperature showed that the particles were superheated above the nickel melting point of 1455°C up to about 2400 °C. The atmosphere around the nickel particle usually contains nitrogen, oxygen, and some moisture. The moisture dissociates at high temperatures and thus can be a source of hydrogen which is easily dissolved in molten droplets with high surface area. Besides, the initial nickel powder was produced by precipitation from a solution...
of nickel ammonium sulphate by the introduction of hydrogen gas at high pressure and temperature. Hence, it was possible that some hydrogen dissolve in nickel during production. After particle impact on the substrate, pressure within the drop raises significantly and instantaneously. This is followed by high cooling rate in a range $10^6–10^9$ °C/s and a fast drop in pressure. Both of these effects result in nucleation of the dissolved gases. Solidification of the splat then shapes the nucleated gases into micro-pores. It should be noted that the focus of the present study was only on the formation of micro-pores (as shown by arrows in Fig. 4), and the large size pores (> 1 micron) which form due to incomplete filling of the intersplat gaps or due to gas entrapment were not studied.

Fig. 4 The microstructure of the splats produced with no H$_2$ content gas in the plasma atmosphere mixture in different magnifications; A) 800X and B) 2000X

Fig. 5 illustrates the EDS pattern on a single splat underside. Besides the Ni peaks, some oxygen is also observed, this can be attributed to oxidation during flight and cooling on the substrate.

Fig. 5 The EDS spectrum on a single splat produced with no H$_2$ content gas in the plasma atmosphere mixture showing nickel and oxygen peaks

Fig. 6 illustrates the microstructure of nickel splat interface sprayed in argon with 7% H$_2$ at different magnifications. Compared to the pure argon case, a higher amount of microporosity is scattered throughout the sample (Fig. 6A). In a single splat (Fig. 6B) it was observed that addition of hydrogen to the plasma has considerably increased the number of micro-pores compared to the pure argon case.

Fig. 6 The microstructure of the splats produced with argon 7% H$_2$ in the plasma atmosphere mixture in different magnifications; A) 800X and B) 2000X

Fig. 7 Shows the EDS pattern for a single splat prepared with argon 7% H$_2$ in the plasma gas mixture.

Fig. 7 The EDS spectrum on a single splat produced with 7%H$_2$ content gas in the plasma atmosphere mixture showing only nickel peaks
In comparison with Fig. 5, no oxygen peak was observed as a result of hydrogen presence, a feature that can be important in some sprayed coatings despite its porosity initiation effect.

Fig. 8 depicts the microstructure of nickel splat-substrate interface sprayed with argon plus 10% hydrogen.

In Fig. 8A larger pores are scattered throughout the sample and, as seen in Fig. 8B, it appears that some pores have merged together. This can cause hydrogen to be more easily released from the samples during cooling period.

Fig. 9 shows the microstructure of nickel splat-substrate interface sprayed with argon plus the extreme case of 13% H₂ in plasma atmosphere at two magnifications. Typically, in plasma spraying processes, 10%H₂ is added to argon as a secondary gas. The selected 13% was to enhance the effect of dissolved hydrogen on the formation of micro-pores.

Comparison Fig. 9 with 13% H₂ and Fig. 8 with 10% hydrogen shows the same trend in porosity formation. In fact, once the particles converted to molten droplets in plasma jet and during travelling to the substrate target, they exposed to hydrogen gas and due to higher solubility of H₂ in molten nickel (Fig. 1), hydrogen was dissolved, and during subsequent cooling and solidification on the substrate micro-pores were nucleated. During cooling and solidification, the rough surface of preheated stainless steel substrate of 300°C provides the nucleation sites, and micro-pores are formed. As the numbers and the density of these pores increase they join together and produce larger pores. This phenomenon also can affect the total H₂ content absorbed by the sample during the process hence higher rate of hydrogen release out of liquid nickel can be expected.

**Discussion**

In order to support this statement, the samples were analyzed for their hydrogen content and the results presented in Table 3. The table shows maximum, minimum, and average concentration of hydrogen in parts per million (ppm). Also, the results for the RSD% (Relative Standard Deviation), known as the coefficient of variance, is calculated. The RSD is a measure for the precision of average of results. For example, a standard deviation of 6% when average result is 40 would mean that the vast majority of results fall between 34 and 46. The relative standard deviation is widely used in analytical chemistry to express the precision and repeatability of an assay. It is calculated as the following:

\[
\text{RSD} = \left( \frac{\text{standard deviation of array X}}{\text{average of array X}} \right) \times 100
\]

As it can be seen, the sample produced under no hydrogen content in plasma gas mixture (I) still contains some hydrogen. As it was mentioned earlier, this could be the reason for lowest level of porosity seen in Fig. 4. Indeed the preheated-roughened stainless steel substrate provided nucleation sites for micro-pores to form in this

<table>
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<tbody>
<tr>
<td>Vol% of H₂ in plasma gas mixture</td>
<td>0</td>
<td>7</td>
<td>10</td>
<td>13</td>
</tr>
<tr>
<td>High Rep (ppm)</td>
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<td>164</td>
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</tr>
<tr>
<td>Low Rep (ppm)</td>
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<td>148</td>
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<td>170</td>
</tr>
<tr>
<td>Avg. (ppm)</td>
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<td>177</td>
<td>178</td>
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<td>5.1</td>
<td>10</td>
<td>3.8</td>
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</table>
sample. Higher RSD% in this sample can be an indication of random hydrogen pick up during powder production. In subsequent samples, higher amounts of hydrogen contents with lower RSD’s can be attributed to hydrogen pick up from plasma gas mixture. Consequently, higher level of porosity is observed on the corresponding microstructures. In the sample IV, the measured hydrogen content ranges to about the same as sample III, in which the microstructures are relatively similar corresponding to their hydrogen content (Fig.’s 8 and 9). It may be that in the latter case, hydrogen bubbles in liquid state merged together and released more H₂ gas as a result of high hydrogen saturation (Fig. 9).

Some other scattering in results obtained could be due to differences in process parameters and droplet properties imposed by plasma spraying route. According to the Table 2, only some minor differences in the impact conditions are observed between the four cases considered here. As it was previously mentioned, after helium, hydrogen has the highest thermal conductivity among gases. Addition of hydrogen will result in faster melting of the injected powder and, hence, more time to dissolve gases before reaching the substrate. In our results this effect reflected in enhancement of torch efficiency by increasing the hydrogen content of the plasma gas mixture despite keeping the particle temperature constant for the sake of similar conditions for the nickel particles throughout the experiments (Table 1).

**Conclusion**

The influence of hydrogen content in plasma gas mixture on formation of micro-pores on the underside of nickel splats was evaluated microscopically. It was shown microscopically that the higher the amount of hydrogen in the gas mixture, the size and the number of micro-porosities increase while the rate of increasing porosities reaches to a saturation level in higher hydrogen concentrated gas mixtures. All the tests in this work were done on commercially pure nickel, however, due to high differences in hydrogen solubility in liquid and solid states for most commercial metals and alloys, the results can be applicable for them as well.

**References**
