A Study on Dynamic Properties of Composite Metal Foams

by Afsaneh Rabiei and Vincent H. Hammond

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A Study on Dynamic Properties of Composite Metal Foams

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14. ABSTRACT
Composite metal foam is a new class of closed cell metal foam offering higher energy absorption compared to the monolithic material at approximately one-third of the density. The superior energy absorption capability of composite metal foam makes this material a potential candidate for structural components such as automotive crash box and crumple zones. However, a systematic study on the dynamic properties of composite metal foams is necessary before they can be utilized in vehicular structures. It has been reported that the energy absorption of other metal foams will increase by increasing the strain rate up to 4× at impact speeds of up to 200 m/s. In this study, the dynamic properties of composite metal foams under various impact speeds are obtained through a variety of experimental techniques. Subsequently, there results are compared to those published in the literature for other foams tested at various impact rates.

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1. Introduction

Composite Metal Foam (CMF) is a cellular material developed at North Carolina State University consisting of a random loose arrangement of hollow metal spheres embedded in a metallic matrix. Quasi-static compression testing shows a uniform deformation behavior without the presence of localized collapse bands, resulting in a high plateau strength in the range of 50–150 MPa depending on the material and processing methods (1). Composite metal foams have also been tested extensively under compression-compression fatigue, loading-unloading compression, and 4 point bending (2–8). However, their behavior at higher loading rates has never been tested.

Excluding a singular exception (9), most of the studies on high strain rate loading of metal foams in the literature report an increased yield and plateau strength as the loading rate increases for foams made from aluminum (10–12), steel (13), and hollow sphere structures (14, 15). The strain rate effect on metal foams is explained as a combination of compressed air trapped inside the cells during impact, micro-inertial effects of the cell walls, and strain rate sensitivity of the cell wall material (12), although some differences exist in the literature as to which of these effects is dominant. A few studies have tested Al foams at higher speeds up to 210 m/s, and report an additional shock phenomena that dominates the foam’s behavior above a critical velocity (16, 17). This “critical” velocity appears to depend on foam properties, such as density (17), and for some foams it is reported as low as 50–55 m/s (18). In this study, steel composite foams, processed using powder metallurgy technique with a variety of spheres sizes, are tested at several loading rates up to 28 m/s using servo-hydraulic machine and Split Hopkinson Pressure Bar apparatus.

2. Experimental Procedure/Materials and Method

2.1 Materials and Processing

Steel composite foam specimens with various sphere sizes of 2.2-, 4- and 5.2-mm outer diameter were produced by powder metallurgy technique. The hollow spheres were produced at Hollomet in Dresden, Germany and their chemical composition and wall thickness are shown in table 1 along with chemical properties of the matrix powder. The matrix material for all static and dynamic compression specimens is 316L stainless steel powder produced by North American Hoganas High Alloys LLC with particle size sieved to −325 mesh (95%) and −200/+325 mesh (5%).
Table 1. Physical properties and chemical composition of hollow spheres and the matrix material.

<table>
<thead>
<tr>
<th>Composite Foam Component</th>
<th>Chemical Composition Weight-Percent</th>
<th>Sphere Properties</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>Fe  C  Mn  Si  Cr  Ni  Mo</td>
<td>Wall thickness (μm)</td>
</tr>
<tr>
<td>5.2 mm spheres</td>
<td>balance 0.87 0.07 0.34 17.09 12.60 2.12</td>
<td>275</td>
</tr>
<tr>
<td>4.0 mm spheres</td>
<td>0.58–0.69 0.15–0.07 1.14–0.32 17.34–16.48 12.28–12.42 2.28–2.11</td>
<td>225</td>
</tr>
<tr>
<td>2.2 mm spheres</td>
<td>balance 0.68 0.13 0.82 16.11 11.53 2.34</td>
<td>110</td>
</tr>
<tr>
<td>Matrix material</td>
<td>balance 0.030 2.00 1.00 16.00–18.00 10.00–14.00 2.00–3.00</td>
<td>NA</td>
</tr>
</tbody>
</table>

Both spheres and powder are placed into a mold and vibrated at a frequency of 20 Hz for ~50 min to achieve a dense packing between the spheres and matrix followed by sintering at 1200 °C in a vacuum hot press. More details of the powder metallurgy processing are available elsewhere (6). All sintered specimens were cut into blocks with a square cross section (24 × 24 mm or 36 × 36 mm) using a Buehler Isomet 4000 linear precision saw. Sample sizes for mechanical testing were chosen such that at least 6 spheres are present in the cross-section to avoid any edge effects and all samples have a width to length aspect ratio of 1.75. Thin slices of foam samples were also cut for scanning electron microscopy imaging (SEM).

2.2 Structural Analysis

The surfaces of thin slice samples were ground using progressive sanding paper followed by polishing with a progression of diamond slurries. Digital images were taken of the foam surface macrostructure as well as individual hollow spheres from each batch. A Hitachi S-3200 Environmental SEM equipped with Energy Dispersive Spectroscopy (EDS) was used to take higher magnification images of the composite foam’s microstructure using accelerating voltage of 20–30 keV, and EDS was performed at several locations to determine the chemical composition of various components in the microstructure. Image analysis software (Image-J) was used to calculate the percentage of the matrix area occupied by porosity.

2.3 Quasi-Static and Dynamic Compression Testing Procedure

Both quasi-static and low speed dynamic compression tests were performed on a servo-hydraulic high rate test system with 300-mm crush depth at velocities up to 8 m/s and load capacity from 60,000–100,000 lb dependent on the velocity. Crosshead displacement is recorded by a Linear Variable Differential Transformer (LVDT) that indicates the actuator position. Tests were performed on multiple specimens of each composite foam at four different crosshead displacement speeds: 0.01, 0.1, 1, and 8 m/s. One test at each speed was also performed on a steel hollow sphere foam (HSF) made by Hollomet with average cell diameter of 2.45 mm and
specimen dimensions 23 × 23 × 40 mm for comparison. High speed video was recorded from each test at 500–10000 fps depending on impact speed.

### 2.4 Split Hopkinson Pressure Bar

Another batch of stainless steel composite foam samples was tested using Split Hopkinson Pressure Bar (SHPB) technique. Cylindrical specimens were manufactured through the powder metallurgy technique described previously using 2.2-mm diameter stainless steel spheres and 316L stainless steel powder, similar to those explained in section 2.1. The diameter of the SHPB specimens was 19 mm with 9.53-mm length for an L/D ratio of 0.5. The SHPB used 19-mm diameter aluminum incident and transmitted bars and a striker bar 177.8-mm long. Air pressure on the gas gun ranged from 50–100 psi to achieve strain rates up to 2770 s⁻¹ (26.4 m/s). Data was reduced to calculate the stress-strain material response through the 2-wave analysis method and high speed video was recorded at 96,000 fps.

### 3. Results and Discussion

#### 3.1 Structural Properties

Figure 1 shows digital and SEM images of the composite foam samples used in this study. As can be seen, there is a uniform distribution of spheres in the matrix in all samples with some leftover pores in the matrix which is the nature of composite foams processed using powder metallurgy technique. Using image analysis, the porosity was measured to be in the range of 10% to 13% for all samples.

The (SEM-EDS) of the sphere wall and the matrix showed the diffusion of the various alloying elements in the 316L matrix such that sharp compositional gradients are minimized (table 2). Similarly, although not directly measured, the sharp gradient in carbon content for the spheres (minimum of 0.6 weight-percent) relative to the matrix (0.03 weight-percent) supports the outward diffusion of carbon from the spheres into the matrix. This combined elemental diffusion helped balance the compositional differences between the spheres and the matrix and lowered the carbon content of spheres. As a result, the hardness of the composite foam was reduced while their ductility and energy absorption was improved.
Figure 1. (A), (B) and (C) Digital images of the composite foam with 2.2-, 4- and 5.2-mm sphere diameters, (D), (E) and (F) corresponding SEM images of the area highlighted with the box in each image respectively.

Table 2. Composition of various features in 2.2-mm sphere CMF before and after processing.

<table>
<thead>
<tr>
<th>Feature</th>
<th>Fe</th>
<th>C</th>
<th>Mn</th>
<th>Si</th>
<th>Cr</th>
<th>Ni</th>
<th>Mo</th>
</tr>
</thead>
<tbody>
<tr>
<td>Before processing</td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Hollow spheres</td>
<td>Balance</td>
<td>0.68</td>
<td>0.13</td>
<td>0.82</td>
<td>16.11</td>
<td>11.53</td>
<td>2.34</td>
</tr>
<tr>
<td>316L powder</td>
<td>Balance</td>
<td>0.03</td>
<td>2.00</td>
<td>1.00</td>
<td>16–18</td>
<td>10–14</td>
<td>2–3</td>
</tr>
<tr>
<td>After processing</td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Sphere wall</td>
<td>Balance</td>
<td>NA</td>
<td>0.21</td>
<td>0.77</td>
<td>15.54</td>
<td>12.49</td>
<td>1.92</td>
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<tr>
<td>Sphere/matrix interface</td>
<td>Balance</td>
<td>NA</td>
<td>0.23</td>
<td>0.283</td>
<td>16.46</td>
<td>13.85</td>
<td>1.57</td>
</tr>
<tr>
<td>Matrix</td>
<td>Balance</td>
<td>NA</td>
<td>0.07</td>
<td>0.31</td>
<td>16.73</td>
<td>14.17</td>
<td>1.12</td>
</tr>
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</table>

Figure 2A presents a digital image of the outer shell of all spheres. As can be seen, the 2.2-mm spheres have a very smooth surface while some features can be seen on the surface of 4- and 5.2-mm spheres. Despite the increased contact area between the matrix and spheres due to the greater surface roughness, these spheres have not bonded well to the matrix (figure 2B). This effect caused some spheres to be debonded from the matrix during mechanical loading. Further investigation is ongoing to change the matrix powder size for better filling those surface features on sphere walls, and to provide optimum bonding at the interface of the matrix with the sphere wall.
Figure 2. Surface features of hollow spheres (A), and the presence of limited debonding at the interface (B).

Table 3. Compositional changes for selected elements after processing for foams with indicated sphere size.

<table>
<thead>
<tr>
<th>Sphere Size</th>
<th>Feature</th>
<th>Fe</th>
<th>Mn (Δ%)</th>
<th>Cr (Δ%)</th>
<th>Mo (Δ%)</th>
</tr>
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<tbody>
<tr>
<td>5.2 mm spheres</td>
<td>Sphere wall</td>
<td>Balance</td>
<td>0.10 (42.9%)</td>
<td>16.56 (–3.2%)</td>
<td>1.71 (–19.4%)</td>
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<td>Sphere/matrix interface</td>
<td>Balance</td>
<td>0.19</td>
<td>15.99</td>
<td>1.62</td>
</tr>
<tr>
<td></td>
<td>Matrix</td>
<td>Balance</td>
<td>0.26 (–87%)</td>
<td>15.59 (–8.3%)</td>
<td>0.84 (–66.4%)</td>
</tr>
<tr>
<td>4 mm spheres</td>
<td>Sphere wall</td>
<td>Balance</td>
<td>0.30 (172.8%)</td>
<td>16.25 (–4.0%)</td>
<td>1.75 (–20.3%)</td>
</tr>
<tr>
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<td>Sphere/matrix interface</td>
<td>Balance</td>
<td>0.30</td>
<td>15.93</td>
<td>1.80</td>
</tr>
<tr>
<td></td>
<td>Matrix</td>
<td>Balance</td>
<td>0.38 (–81%)</td>
<td>15.07 (–11.4%)</td>
<td>2.18 (–12.8%)</td>
</tr>
<tr>
<td>2.2 mm spheres</td>
<td>Sphere wall</td>
<td>Balance</td>
<td>0.18 (38.5%)</td>
<td>16.05 (–0.4%)</td>
<td>1.95 (–16.7%)</td>
</tr>
<tr>
<td></td>
<td>Sphere/matrix interface</td>
<td>Balance</td>
<td>0.23</td>
<td>15.57</td>
<td>1.87</td>
</tr>
<tr>
<td></td>
<td>Matrix</td>
<td>Balance</td>
<td>0.31 (–84.5%)</td>
<td>15.4 (–9.5%)</td>
<td>2.14 (–14.4%)</td>
</tr>
</tbody>
</table>
3.2 Mechanical Properties

Figure 3 presents the stress-strain curve of composite foams with various sphere sizes tested under various loading speeds. Figures 4A and B compare the yield strength and modulus of elasticity of all samples at various speeds. As can be seen, both the yield and plateau strength of all composite foams is decreased by increasing the spheres diameter whereas an increase in the loading speed increased the yield and plateau strength of the material. As the result, the highest strength belonged to the composite foams made out of the 2-mm spheres tested under 25 m/s loading speed. In addition, a distinct increase is also observed in the plateau stress when the response of the composite foams is compared with a sintered sphere-only perform of similar material and sphere dimensions. This shows the importance of the matrix in bonding the spheres together and reinforcing the sphere walls under loading.

![Stress-strain curve for samples with various sphere sizes under various loading speeds.](image)

It is notable that an improvement of 80% to 140% is observed in the yield strength of the composite foam samples made with 2-mm hollow spheres and tested under SHPB. This is in agreement with previous reports that an increase in the loading speed can increase the yield and plateau strength of metal foams. Comparing the modulus of elasticity of the samples tested in servo-hydraulic testing machine shows a decrease in modulus with increasing loading speed from 0.01 to 8 m/s. However, results from those tests using Hopkinson Bar techniques show a completely opposite relationship, namely an increase in modulus with increasing loading speed.
This can be related to the large compliance in the servo-hydraulic testing machine, particularly at the higher loading speeds. The compliance is also responsible for the wavy stress-strain curve in figure 3 at higher impact rates.

4. Conclusions

- Composite metal foams consisting of hollow steel spheres in 316L stainless steel matrix have been produced using powder metallurgy techniques.
- Microstructural analysis indicated that the hollow spheres were uniformly distributed in the matrix. However, some residual porosity (10%–13%), as well as some matrix-sphere debonding, was also observed.
- Elemental analysis indicated that manganese and nickel diffused from the matrix into the sphere wall. In addition, the large difference in carbon content for the spheres compared to the matrix suggests the outward diffusion of carbon from the spheres into the matrix. As a result, the hardness of the foam is reduced while the ductility and energy absorption was improved.
- High strain rate testing indicated that an increase in sphere diameter resulted in a reduction in the yield and plateau strength for the composite foam. For constant sphere diameter, increasing the loading speed acted to increase the yield and plateau strengths.
- In keeping with other studies reported in the literature, an increase in the yield and plateau strength was observed with increasing loading speed for the metal foam. In addition, the elastic modulus was also found to increase with loading speed in the Hopkinson Bar experiments.
5. References

List of Symbols, Abbreviations, and Acronyms

<table>
<thead>
<tr>
<th>Acronym</th>
<th>Description</th>
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<tbody>
<tr>
<td>CMF</td>
<td>Composite Metal Foam</td>
</tr>
<tr>
<td>EDS</td>
<td>Energy Dispersive Spectroscopy</td>
</tr>
<tr>
<td>HSF</td>
<td>hollow sphere foam</td>
</tr>
<tr>
<td>LVDT</td>
<td>Linear Variable Differential Transformer</td>
</tr>
<tr>
<td>SEM</td>
<td>scanning electron microscopy imaging</td>
</tr>
<tr>
<td>SHPB</td>
<td>Split Hopkinson Pressure Bar</td>
</tr>
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<td>ORGANIZATION</td>
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<td>ARMY RESEARCH LABORATORY 4600 DEER CREEK LOOP APG MD 21005</td>
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