Aluminum-Scandium Alloys: Material Characterization, Friction Stir Welding, and Compatibility With Hydrogen Peroxide

(MSFC Center Director’s Discretionary Fund Final Report, Project No. 04–13)

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<thead>
<tr>
<th>Acronym</th>
<th>Description</th>
</tr>
</thead>
<tbody>
<tr>
<td>Al</td>
<td>aluminum</td>
</tr>
<tr>
<td>AOL</td>
<td>active oxygen loss</td>
</tr>
<tr>
<td>CDDF</td>
<td>Center Director’s Discretionary Fund</td>
</tr>
<tr>
<td>Cr</td>
<td>chromium</td>
</tr>
<tr>
<td>Cu</td>
<td>copper</td>
</tr>
<tr>
<td>FSW</td>
<td>friction stir welding</td>
</tr>
<tr>
<td>HAZ</td>
<td>heat-affected zone</td>
</tr>
<tr>
<td>HNO(_3)</td>
<td>nitric acid</td>
</tr>
<tr>
<td>H(_2)O(_2)</td>
<td>hydrogen peroxide</td>
</tr>
<tr>
<td>Hyper-X</td>
<td>Hypersonic-X</td>
</tr>
<tr>
<td>IMC</td>
<td>isothermal microcalorimeter</td>
</tr>
<tr>
<td>ISTAR</td>
<td>integrated system test of an air-breathing rocket</td>
</tr>
<tr>
<td>Mg</td>
<td>magnesium</td>
</tr>
<tr>
<td>Mn</td>
<td>manganese</td>
</tr>
<tr>
<td>MSFC</td>
<td>Marshall Space Flight Center</td>
</tr>
<tr>
<td>NaOH</td>
<td>sodium hydroxide</td>
</tr>
<tr>
<td>O(_2)</td>
<td>oxygen</td>
</tr>
<tr>
<td>RBCC</td>
<td>rocket-based combined cycle</td>
</tr>
<tr>
<td>Sc</td>
<td>scandium</td>
</tr>
<tr>
<td>Ti</td>
<td>titanium</td>
</tr>
<tr>
<td>TM</td>
<td>Technical Memorandum</td>
</tr>
<tr>
<td>Zn</td>
<td>zinc</td>
</tr>
<tr>
<td>Zr</td>
<td>zirconium</td>
</tr>
</tbody>
</table>
NOMENCLATURE

$C$  \quad$\text{initial fraction H}_2\text{O}_2$

$C_1$  \quad$\text{initial concentration (H}_2\text{O}_2 \text{ stability)}$

$d_1$  \quad$\text{density at initial temperature}$

$W_1$  \quad$\text{initial net weight (active oxygen loss); initial flask weight (H}_2\text{O}_2 \text{ stability)}$

$W_2$  \quad$\text{final net weight (active oxygen loss); final flask weight (H}_2\text{O}_2 \text{ stability)}$
1. INTRODUCTION

This Technical Memorandum (TM) describes the development of high-strength aluminum (Al) alloys for fuel tanks and air frame applications that are chemically compatible with hydrogen peroxide (H$_2$O$_2$) propellant. These new high-strength alloys could represent an enabling material technology for the development of NASA’s next generation of Hypersonic-X (Hyper-X) vehicles, where flight weight reduction is a critical requirement. These Hyper-X vehicles are being studied as air-breathing hypersonic research vehicles that feature a lifting body configuration with a rocket based combined cycle (RBCC) propulsion system.\textsuperscript{1} This task supports three main Engineering Directorate’s thrust areas at Marshall Space Flight Center (MSFC): (1) Advanced cryogenic tank area for materials’ compatibility with H$_2$O$_2$ systems, including metallic liners and permeation barriers, (2) advanced structures and materials area for lightweight and high specific strength Al alloys, and (3) advanced manufacturing in the space transportation area for welding of advanced Al alloys compatible with H$_2$O$_2$ propellant.

As shown in figure 1, this work directly supports the product line for NASA’s Hyper-X flight test air-breathing vehicles such as the X–43B, which is an integrated system test of an air-breathing rocket (ISTAR) flight demonstrator.\textsuperscript{2} The flight-type ground test engine sized specifically for the X–43B uses JP–7 fuel and 90 percent concentrated liquid H$_2$O$_2$ as an oxidizer. When highly concentrated H$_2$O$_2$ is allowed to decompose in the combustion chamber, the H$_2$O$_2$ rapidly decomposes into steam (water) and oxygen (O$_2$) and releases a large amount of heat that can be used as an oxidizer in a bipropellant mode, for thrust in monopropellant mode, and as the working fluid for pressurization in a turbine drive. Currently, conventional 1060 and 5254 materials are the baseline Al alloys for H$_2$O$_2$ fuel tanks and integral structures, but their yield strengths are relatively low at 17 ksi. Being non-heat-treatable alloys, these materials may not be suitable for advanced lightweight tanks or airframes due to low specific strengths.
Figure 1. Artist rendering of the air breathing X–43B Hyper-X vehicle.

1.1 Hydrogen Peroxide Impact on Design of Propulsion Systems and Storage Tanks

H$_2$O$_2$ has a water-like appearance but is very chemically active as an oxidizer, and is highly soluble in water at all proportions; the usual commercial forms are from 3 to 30 percent aqueous solution. For rocket propulsion applications, high concentrations of H$_2$O$_2$ solutions of >70 percent are used since they are stable at room temperature. When properly contained in chemically compatible vessels, H$_2$O$_2$ has a very weak decomposition rate. A concentration of 90 percent grade H$_2$O$_2$ was selected for the X–43B vehicle, with an active O$_2$ content of 42.3 percent. Table 1 shows some selected physical properties for highly concentrated H$_2$O$_2$. When highly concentrated H$_2$O$_2$ comes in contact with incompatible materials like heavy metals or various organic compounds, or mixes with certain impurities, it produces O$_2$ gas and decomposition heat. The decomposition rate increases about 2.2 times for each 10 °C increase in temperature from 20 to 100 °C. Table 2 shows the effect of heat on the decomposition rate. At near room temperature, the rate of decomposition is ≈1 percent per year. This rate increases exponentially to 1 percent per week if the temperature is allowed to rise to 66 °C (151 °F), and could result in a rapid pressure increase in a closed system.

Indications of H$_2$O$_2$ decomposition are: pressure buildup by activation of pressure relief valves, visible rapid bubbling, temperature increase, gas or steam evolution, and possible explosion when pressure relief is inadequate. Pressure buildup will occur very rapidly in a closed system and excessive pressure buildup can result in ruptured storage tanks, fuel tanks, or feed lines. Using H$_2$O$_2$ could have an impact on propulsion system and storage tank designs. Figure 2 shows the X–43B vehicle configuration layout for the H$_2$O$_2$ tank and the JP–7 fuel tank.
Table 1. Selected physical properties of highly concentrated H\textsubscript{2}O\textsubscript{2}.

<table>
<thead>
<tr>
<th>Properties</th>
<th>H\textsubscript{2}O\textsubscript{2} Concentration (wt. %)</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>70%</td>
</tr>
<tr>
<td>Active oxygen (%)</td>
<td>32.9</td>
</tr>
<tr>
<td>Specific gravity @ 25 °C</td>
<td>1.28</td>
</tr>
<tr>
<td>Boiling point (°C)</td>
<td>125.4</td>
</tr>
<tr>
<td>Freezing point (°C)</td>
<td>-40.3</td>
</tr>
<tr>
<td>Flash point (°C)</td>
<td>None</td>
</tr>
<tr>
<td>Autoignition in air (°C)</td>
<td>316</td>
</tr>
<tr>
<td>Vapor pressure (mmHg)</td>
<td>10.1</td>
</tr>
</tbody>
</table>

Table 2. Effect of heat on decomposition rate of H\textsubscript{2}O\textsubscript{2}.

<table>
<thead>
<tr>
<th>Temperature</th>
<th>Rate of Decomposition</th>
</tr>
</thead>
<tbody>
<tr>
<td>°C</td>
<td>°F</td>
</tr>
<tr>
<td>22</td>
<td>72</td>
</tr>
<tr>
<td>66</td>
<td>151</td>
</tr>
<tr>
<td>103</td>
<td>218</td>
</tr>
</tbody>
</table>

Figure 2. X–43B configuration layout for the H\textsubscript{2}O\textsubscript{2} and JP–7 fuel tank.
Using any \( \text{H}_2\text{O}_2 \) system would affect the design of certain types of Hyper-X vehicle’s propulsion system, such as onboard fuel tanks, valves, and pumps, which must be pacified to enhance chemical compatibility. All valves must be compatible and have relief capability with no isolated segments in the feed system to trap any \( \text{H}_2\text{O}_2 \). \( \text{H}_2\text{O}_2 \) is not an intrinsically hazardous fluid, but it must be contained in a pacification system that is well engineered to remove excessive heat and pressure from the storage and feed system due to decomposition and high-temperature effects. Pacification of storage tanks and feed lines is usually accomplished through various acid baths and exposure to varying degrees of \( \text{H}_2\text{O}_2 \). An \( \text{H}_2\text{O}_2 \)-based system will impact overall system complexity because of feed system architecture.

1.2 Classifications for Material Compatibility With Hydrogen Peroxide

According to FMC corporation’s technical data from *Bulletin 104*, materials should be classified into categories based on their contemplated types of use. Therefore, all materials are not required to be suitable for indefinite storage, because in applications requiring only short-time contact with \( \text{H}_2\text{O}_2 \), materials with a lesser degree of compatibility can be employed. To facilitate the material selection on this basis, FMC has developed four categories of material classifications as follows:

- **Class 1**: Unlimited use materials—can be used for long-term contact with \( \text{H}_2\text{O}_2 \) as storage containers.

- **Class 2**: Repeated use in short-time contact materials—can be used for either transient contact with the \( \text{H}_2\text{O}_2 \) prior to storage or limited contact prior to use. Such contact is not to exceed 4 hr at 72 °C (160 °F) or 1 wk at 22 °C (70 °F)—typically used for valves and pumps in transfer lines and feed tanks.

- **Class 3**: Short-time contact materials—can be used for repeated contact, but no single contact period should exceed 1 min at 72 °C (160 °F), or 1 hr at 22 °C (70 °F), because sufficient contamination of the \( \text{H}_2\text{O}_2 \) may render it unsuitable for storage. Many class 3 materials indicate satisfactory service at room temperature. However, the material should be pretested prior to use.

- **Class 4**: Not recommended materials—will cause excessive decomposition of \( \text{H}_2\text{O}_2 \) even on short-time contact and yield corrosion or deterioration on products to form an impact-sensitive mixture with concentrated \( \text{H}_2\text{O}_2 \).
2. EXPERIMENTAL

2.1 Alloy Development

The current baseline material recommended for long-term storage of highly concentrated class 1 (>90 percent) H₂O₂, is 5254 Al alloy, which is wrought and non-heat-treatable Al-magnesium (Mg) alloy. The 5000 series Al-Mg alloys are good candidate materials for structural applications due to their low density, reasonably good weld ability, and excellent corrosion resistance. However, similar to most Al-Mg systems, the 5254 alloy does not have adequate specific strength and density for advanced aerospace applications. Table 3 shows the compatibility assessment of selected alloys for H₂O₂ storage tank applications. As a general rule, the higher strength alloys tend to be incompatible with H₂O₂ for long-term storage applications.

Table 3. Compatibility assessment of selected alloys for H₂O₂ storage tanks.

<table>
<thead>
<tr>
<th>Current Materials for Tankage</th>
<th>Temper</th>
<th>Yield Strength (ksi)</th>
<th>H₂O₂ Compatibility Class</th>
</tr>
</thead>
<tbody>
<tr>
<td>1060</td>
<td>H14</td>
<td>13</td>
<td>1</td>
</tr>
<tr>
<td>1100</td>
<td>H14</td>
<td>17</td>
<td>1</td>
</tr>
<tr>
<td>2024</td>
<td>T6</td>
<td>57</td>
<td>3</td>
</tr>
<tr>
<td>Al–Li 2195</td>
<td>T81</td>
<td>66</td>
<td>2</td>
</tr>
<tr>
<td>2219</td>
<td>T81</td>
<td>50</td>
<td>3</td>
</tr>
<tr>
<td>5254</td>
<td>H112</td>
<td>17</td>
<td>1</td>
</tr>
<tr>
<td>6061</td>
<td>T6</td>
<td>40</td>
<td>2</td>
</tr>
<tr>
<td>7075</td>
<td>T6</td>
<td>73</td>
<td>4</td>
</tr>
</tbody>
</table>

The initial goal was to develop a new experimental Al-Mg-based alloy that has the same class 1 compatibility rating with H₂O₂, but with a significant improvement in yield strength by a factor of 2–3 times more than 5254. This strategy was designed to modify the chemistry of 5254 alloy by adding scandium (Sc) and zirconium (Zr), which was not perceived to act as catalysts to decompose liquid H₂O₂. Fine Al₃Sc precipitates are known to be coherent with the Al matrix and are expected to contribute to the alloy strength through dislocation-particle interactions. Scandium also effectively increases the recrystallization temperature for Al. However, simultaneous addition of Sc and Zr has been shown to synergistically promote much higher strengths than either Sc or Zr additions produce alone. Figure 3 shows the synergistic strengthening effect of Sc and Zr additions for Al.
Analyses for the catalytic effects of Sc, Zr, copper (Cu), zinc (Zn), manganese (Mn), titanium (Ti), and Mg additions to the Al matrix have shown that these elements may yield only a minor effect on the rate of decomposition for $\text{H}_2\text{O}_2$ if they are added in relatively small amounts without surpassing their maximum solid solution in the Al matrix composition. Such elements have been added to several experimental Al-Mg alloys in appropriate amounts. Ingots were cast and rolled into thin sheet metals and test coupons were machined from sheet metals for long-term exposure and mechanical properties testing with $\text{H}_2\text{O}_2$ (fig. 4). In addition, the ability to weld the new alloys using friction stir welding (FSW) has also been explored.

Figure 3. Synergistic strengthening effect of Sc and Zr additions.

Figure 4. Typical microstructures for experimental alloys: (a) cast and (b) cold rolled at x 200 magnification.
2.2 Candidate Materials

Table 4 shows the chemical compositions of four candidate Al-Mg-Sc-bearing alloys, including the 5254 alloy as a baseline reference material. The rationale for selecting these candidate materials is explained as follows:

• Successfully developed a new experimental Al-Mg-chromium- (Cr-) based alloy named RX 5000 by adding Sc and Zr.

• Strengthened RX 5000 by Mg atoms in solid solution and by cold working in the H112 conditions, which increased yield strength to 42 ksi—nearly 2.5 times stronger than the 17 ksi yield strength of conventional 5254 alloy.

• RX 5000 is completely compatible with class 1, 90 percent $\text{H}_2\text{O}_2$.

<table>
<thead>
<tr>
<th>Chemical Composition (wt. %) of candidate alloys.</th>
</tr>
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<tbody>
<tr>
<td><strong>Alloy</strong></td>
</tr>
<tr>
<td>-----------</td>
</tr>
<tr>
<td>5254</td>
</tr>
<tr>
<td>RX 5000</td>
</tr>
<tr>
<td>C557</td>
</tr>
<tr>
<td>7X0X</td>
</tr>
<tr>
<td>7X11</td>
</tr>
</tbody>
</table>

A search for an existing alloy with similar composition to RX 5000 in commercial production yielded the C557 alloy, which is Al-Mg-Sc-Zr with a slightly higher Mn content than the RX 5000 alloy, without Cr. The chemistry of the C577 rolled sheet is very similar to the Russian alloy 1535 (table 4). The C557 alloy was produced by Alcoa® for commercial scale ingots with the H116 heat treatment to ensure good stress corrosion resistance. Several C577 test panels were later received from Alcoa for FSW and $\text{H}_2\text{O}_2$ compatibility testing at MSFC.

The characterization of Sc-bearing Al-Mg alloy C577 for aerospace applications was previously documented in the literature for strength-toughness behaviors. As previously stated, the C557 is also completely compatible with class 1, 90 percent $\text{H}_2\text{O}_2$.

Because RX 5000 and C557 alloys are non-heat-treatable, further attempts to increase strength would be limited without further development and/or characterization of heat-treatable Al-Mg-Sc-Zr alloys with the additions of either Zn and/or Cu. Surface Treatment Technologies has characterized two heat-treatable alloys named 7X0X and 7X11 (T. Langan, Presentation at Technical Interchange Meeting, 2003). These two heat treatable alloys are completely compatible with class 1, 90 percent $\text{H}_2\text{O}_2$. 
3. RESULTS AND DISCUSSION

3.1 Mechanical Properties

Table 5 shows comparisons of typical mechanical properties of four candidate Al-Mg-Zr-Sc-bearing alloys, and displays the properties for conventional 5254 alloy as the baseline reference material. 7X0X and 7X11 wrought, heat-treatable alloys are more superior in tensile and yield strengths than the 5254 alloy by a large margin; followed by a moderate increase in strength from non-heat-treatable alloys, C557 and RX 5000. An important point of reference is that the yield strength and apparent fracture toughness values of C577 are within 10 percent of established values for 2024-T3 sheet.

On the other hand, C557 and RX 5000 are inherently more corrosion resistant than 7X0X and 7X11 because they do not contain secondary strengthening precipitates from Mg, Zn, and Cu.

Comparisons between 7X0X and 7X11 show a minor increase in strength for the 7X11 alloy, due to the addition of Cu. With the T6 condition, the 7X11’s yield strength is \( \approx 4.8 \) times higher than the conventional 5254. Since yield and tensile strength are important basic material design properties, figure 5 shows the yield and tensile strengths of all four candidate materials as compared to 5254 baseline alloy.

3.2 Chemical Compatibility With Hydrogen Peroxide

FMC personnel conducted chemical compatibility tests for each type of alloy at 66 °C for 7 days in a water bath with controlled temperature, and each set of coupons ran in duplication. The approximate surface area of each coupon is \( \approx 1.75 \text{ in}^2 \). About 86 mL of \( \text{H}_2\text{O}_2 \) was used for each run, which made the surface area-to-\( \text{H}_2\text{O}_2 \) volume ratio roughly equal 0.33 in\(^{-1} \). This coupon was immersed in 0.25 percent sodium hydroxide (NaOH) for 20 min and then immersed in 45 percent nitric acid (HNO\(_3\)) for 1 hr.
and finally conditioned in 35 percent H₂O₂ for 24 hr before the compatibility test in 90 percent H₂O₂. Appendix A shows the procedure used by FMC for determining the compatibility of materials with H₂O₂ by immersion in liquid and vapor at controlled temperatures. Appendix B shows the procedure for determining the stability of H₂O₂. Figure 6(a) shows test coupons immersed in a typical test vial that contains 90 percent H₂O₂. Figure 6(b) shows the isothermal microcalorimeter (IMC), which is another method used to determine decomposition rate. The IMC determines the difference between the background thermal energy of the H₂O₂ before and after the sample is added in the vial. The IMC was used at MSFC to determine the active oxygen loss (AOL) percent of the H₂O₂.

![Graph showing strengths of candidate materials compared to 5254 baseline alloy](image)

**Figure 5.** Strengths of candidate materials compared to 5254 baseline alloy.

The results of the H₂O₂ compatibility test are shown in table 6, which was performed using procedures outlined by FMC in appendices A and B. Four types of Al-Mg-Sc-bearing alloys were tested, plus a fifth type of sample C577 that was taken from an FSW joint. For each type of material, two coupons were required to conduct the test. A blank control test was also run in parallel for each type of alloy for monitoring the system contributions to the results. Changes in weight and concentration of the H₂O₂ solution were measured in the presence of coupons to calculate relative AOL percent. In table 6, the ΔAOL percent represents the measured difference between the sample’s AOL and the blank’s AOL, which displays the true effect of the coupon on H₂O₂ decomposition.
Figure 6. (a) Vial contains 90 percent grade H$_2$O$_2$ and test coupons, and (b) isothermal microcalorimeter is used to determine the H$_2$O$_2$ decomposition rate.

Table 6. H$_2$O$_2$ compatibility assessment of current class 1 materials.

<table>
<thead>
<tr>
<th>Run No.</th>
<th>Coupon ID</th>
<th>H$_2$O$_2$ Start Weight (g)</th>
<th>H$_2$O$_2$ Weight Loss (g)</th>
<th>Final Assay (%)</th>
<th>AOL (g)</th>
<th>AOL (%)</th>
<th>∆AOL (%)</th>
<th>Coupon Weight Loss (g)</th>
<th>Stability (%)</th>
</tr>
</thead>
<tbody>
<tr>
<td>G0380–138–1</td>
<td>Blank–1</td>
<td>119.48</td>
<td>0.036</td>
<td>90.21</td>
<td>0.097</td>
<td>0.192</td>
<td>–</td>
<td>–</td>
<td>99.65</td>
</tr>
<tr>
<td>G0380–138–2</td>
<td>Blank–2</td>
<td>119.85</td>
<td>0.034</td>
<td>90.21</td>
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<td>0.19</td>
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<td>–</td>
<td>99.16</td>
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<td>G0380–138–3</td>
<td>RX 5000–1</td>
<td>119.71</td>
<td>0.053</td>
<td>90.2</td>
<td>0.115</td>
<td>0.226</td>
<td>0.035</td>
<td>0</td>
<td>98.24</td>
</tr>
<tr>
<td>G0380–138–4</td>
<td>RX 5000–2</td>
<td>119.99</td>
<td>–0.214</td>
<td>89.9</td>
<td>0.166</td>
<td>0.326</td>
<td>0.135</td>
<td>0</td>
<td>99.16</td>
</tr>
<tr>
<td>G0380–138–5</td>
<td>C557–1</td>
<td>119.64</td>
<td>0.444</td>
<td>89.87</td>
<td>0.464</td>
<td>0.913</td>
<td>0.722</td>
<td>0.0001</td>
<td>98.24</td>
</tr>
<tr>
<td>G0380–138–6</td>
<td>C557–2</td>
<td>120.44</td>
<td>0.427</td>
<td>89.9</td>
<td>0.438</td>
<td>0.856</td>
<td>0.665</td>
<td>0.0002</td>
<td>98.24</td>
</tr>
<tr>
<td>G0380–146–1</td>
<td>Blank–3</td>
<td>120.82</td>
<td>0.156</td>
<td>89.9</td>
<td>0.268</td>
<td>0.524</td>
<td>–</td>
<td>–</td>
<td>99.79</td>
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<tr>
<td>G0380–146–2</td>
<td>Blank–4</td>
<td>122.34</td>
<td>0.058</td>
<td>89.94</td>
<td>0.209</td>
<td>0.403</td>
<td>–</td>
<td>–</td>
<td>–</td>
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<tr>
<td>G0380–146–3</td>
<td>7X0X–1</td>
<td>119.29</td>
<td>0.073</td>
<td>90.05</td>
<td>0.151</td>
<td>0.297</td>
<td>–0.248</td>
<td>–0.0001</td>
<td>99.51</td>
</tr>
<tr>
<td>G0380–146–4</td>
<td>7X0X–2</td>
<td>119.88</td>
<td>–0.009</td>
<td>89.91</td>
<td>0.196</td>
<td>0.384</td>
<td>–0.161</td>
<td>–0.0001</td>
<td>99.51</td>
</tr>
<tr>
<td>G0376–20–1</td>
<td>Blank–5</td>
<td>119.7</td>
<td>–0.5</td>
<td>90.23</td>
<td>0.113</td>
<td>0.222</td>
<td>–</td>
<td>–</td>
<td>99.62</td>
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<tr>
<td>G0376–20–2</td>
<td>Blank–6</td>
<td>120</td>
<td>0</td>
<td>90.67</td>
<td>0.06</td>
<td>0.157</td>
<td>–</td>
<td>–</td>
<td>–</td>
</tr>
<tr>
<td>G0376–20–3</td>
<td>7X11–1</td>
<td>119.4</td>
<td>0.1</td>
<td>90.58</td>
<td>0.173</td>
<td>0.34</td>
<td>0.151</td>
<td>–0.0003</td>
<td>99.29</td>
</tr>
<tr>
<td>G0376–20–4</td>
<td>7X11–2</td>
<td>121.1</td>
<td>0.1</td>
<td>90.58</td>
<td>0.175</td>
<td>0.339</td>
<td>0.15</td>
<td>–0.0004</td>
<td>–</td>
</tr>
<tr>
<td>G0376–20–5</td>
<td>FSW C557–1</td>
<td>120.9</td>
<td>0.1</td>
<td>90.6</td>
<td>0.165</td>
<td>0.319</td>
<td>0.13</td>
<td>–0.0003</td>
<td>99.57</td>
</tr>
<tr>
<td>G0376–20–6</td>
<td>FSW C557–2</td>
<td>119.6</td>
<td>0</td>
<td>90.56</td>
<td>0.141</td>
<td>0.277</td>
<td>0.088</td>
<td>–0.0003</td>
<td>–</td>
</tr>
</tbody>
</table>

Each candidate material has shown a stability value that is similar to the value from a blank test run >99.6 percent, which did not have any coupons in them. This is the baseline for the test results because H$_2$O$_2$ solutions may be slightly different from batch to batch, depending on the containers and varying temperature control from test to test. Blank samples served as references to eliminate all the noises. The results indicate that all candidate materials can be classified as class 1. FSW samples were also extracted from the welding surface area and tested for compatibility. The data also showed that an FSW joint has a stability of class 1, 99.57 percent. These data confirm the notion that FSW has
a potential for welding Al-Sc-bearing alloys with minimal material contamination that could affect the compatibility with $\text{H}_2\text{O}_2$ in the weld joint area.

### 3.3 Friction Stir Welding

FSW is a relatively new joining process that was originally developed and patented by the Welding Institute of Cambridge, England. Since 1993, FSW has been studied and demonstrated by researchers for Al alloys and metal matrix composites. FSW can be best described in conventional terms as a combination of extrusion and forging of metals at elevated temperatures. This process is considered a solid state process and does not require the need for gas shielding or filler metals. FSW consists of a rotating, nonconsumable pin tool that is slowly plunged into the bond line until the pin tool’s shoulder is in intimate contact with the work piece. As the tool rotates and moves forward along the bond line, the material on the bond line begins to heat and is forced to flow around the rotating tip to consolidate on the pin tool’s backside. This heat source is developed mainly because of local friction and plastic deformation while keeping the pin tool’s shoulder in intimate contact with the work piece at all times. Figure 7 shows FSW and the pin tool’s geometric design.

![Figure 7](image)

Figure 7. (a) FSW and (b) the pin tool’s geometric design.

Interestingly, FSW has the potential for welding Al-Sc alloys because the processing temperature occurs well below the metal’s melting point, thereby eliminating the undesirable chemical reactions, which could reduce material contaminations that may affect the compatibility with $\text{H}_2\text{O}_2$ in the weld joint area. Figure 8 shows a cross-sectional view of an FSW joint for C557 alloy. The hardness measurement was taken across the crown side of the weld zone as shown in figure 9. The yield and tensile strengths were slightly reduced in the heat-affected zone (HAZ) by overheating from the FSW process. However, the FSW strength efficiency of >90 percent is excellent when compared with relatively high-temperature fusion joining processes such as arc welding. Table 7 shows the FSW tensile strength and joint efficiency for candidate Al-Mg-Sc-bearing alloys. Because the FSW process does not require gas shielding or filler metals, welding joints have less contamination than arc welding (table 8).
Figure 8. Cross-sectional view of an FSW joint for C557 alloy.

Figure 9. HAZ across an FSW joint for C557 alloy.

Table 7. FSW tensile strength and joint efficiency.

<table>
<thead>
<tr>
<th>Candidate Alloy</th>
<th>Temper</th>
<th>Weld Joint Ultimate Tensile Strength (ksi)</th>
<th>Joint Efficiency (Weld Ultimate Tensile Strength/Base Metal Ultimate Tensile Strength) (%)</th>
</tr>
</thead>
<tbody>
<tr>
<td>7X11</td>
<td>T7</td>
<td>66.2</td>
<td>91.9</td>
</tr>
<tr>
<td>7X0X</td>
<td>T7</td>
<td>63.6</td>
<td>90</td>
</tr>
<tr>
<td>C557</td>
<td>H116</td>
<td>53.4</td>
<td>86</td>
</tr>
<tr>
<td>RX 5000</td>
<td>H112</td>
<td>48.6</td>
<td>88.3</td>
</tr>
</tbody>
</table>

Table 8. FSW results in low contamination at the welding joint.

<table>
<thead>
<tr>
<th>Alloy</th>
<th>H$_2$O$_2$ Stability After 7 Days @ 66 °C (%)</th>
<th>Compatibility Classification</th>
</tr>
</thead>
<tbody>
<tr>
<td>No sample</td>
<td>99.65</td>
<td>Reference Class 1</td>
</tr>
<tr>
<td>FSW C557 joint</td>
<td>99.57</td>
<td></td>
</tr>
</tbody>
</table>
4. SUMMARY AND CONCLUSIONS

The purpose of this work was to develop a new Al-Mg-based alloy that has the same class 1 compatibility rating with H$_2$O$_2$, similar to conventional 5254 alloy, but with a significant improvement in yield strength. Several Al-Mg-Sc-based alloys have been successfully developed and/or characterized and tested for H$_2$O$_2$ compatibility. Analysis for the catalytic effects of Sc, Zr, Cu, Zn, Mn, Ti, and Mg additions to the Al matrix has shown that these elements may yield only a minor effect on the decompose rate of H$_2$O$_2$, provided that they are added in relatively small amounts, without surpassing their maximum solid solution in the Al matrix.

The preliminary test data indicate that all of these alloys are chemically stable (inert) when exposed to 90 percent H$_2$O$_2$. With the T6 condition, the 7X11 heat-treatable alloy’s yield strength is about 4.8 times higher than the conventional 5254 yield strength, while maintaining excellent H$_2$O$_2$ compatibility similar to the class 1 5254 alloy. Moreover, test data show that these alloys can be welded successfully using FSW.
APPENDIX A—PROCEDURE FOR DETERMINING THE COMPATIBILITY OF MATERIALS WITH HYDROGEN PEROXIDE BY IMMERSION IN LIQUID AND VAPOR AT CONTROLLED TEMPERATURES

A.1 Introduction

Compatibility testing and evaluation of materials with H₂O₂ is conducted by immersing material samples either in H₂O₂ for 4 wk at 30 °C (86 °F), if it cannot be tested at higher temperature, or 1 wk at 66 °C (150.8 °F). The percent loss of active O₂ is calculated from the decreased weight of oxide concentration and the decreased weight of the system. The AOL, together with an appraisal of the condition of the tested material, is used to govern the selection and recommendation of suitable materials for use with H₂O₂.

A.2 Sample Size

A 3 in × 0.5 in × 0.0625 in sample strip is normally used to provide an evaluation of both liquid and vapor phase attack. Surface-to-volume ratio is calculated for the wetted area only. A 1.5 in × 0.5 in × 0.0625 in sample strip will be used only for the material tested under a continuously wetted condition. A volume of 74 mL of liquid H₂O₂ will be used for this test, which yields an apparent sample surface area of 1 in² of test surface to 42.8 mL of H₂O₂. This approximates the 0.33 in²/in³ surface-to-volume ratio (0.33 in⁻¹) of the wetted surface of a standard drum containing 250 lb of H₂O₂. If it is necessary to test a sample that is not the standard 3 in × 0.5 in × 0.0625 in wetted surface, retaining this apparent surface-to-volume ratio will help interpret the results. For liquids, a 5-mL sample is used; for greases, a sample of ≈5 g is smeared on the inside of the test flask.

A.3 Apparatus

The following elements are used in the compatibility testing process:

- 125 mL pyrex kjeldahl flask
- 110 °C drying oven
- 66 °C (150.8 °F) at ±0.5 °C water bath
- 30 °C (86 °F) at ±0.5 °C water bath
- Aluminum foil to cover flasks
- Distilled water with conductivity of <2 μmho cm.

A.4 Cleaning and Passivation

A.4.1 Glassware

Prior to use, all glassware is immersed in a 10 percent NaOH solution for 1 hr at room temperature, rinsed with distilled water, immersed in a 10 percent HNO₃ solution for a minimum of 3 hr,
and finally rinsed with distilled water. The mouths of the flasks are then covered with clean aluminum foil and the flasks are dried in a 110 °C oven.

A.4.2 Metals

Aluminum samples are scrubbed with a warm detergent solution and then immersed at room temperature in N/15 NaOH for 15–20 min. Then they are washed and immersed for 45 min to 1 hr at room temperature in 45 percent HNO₃, and finally washed with distilled water, then pretreated with 35 percent H₂O₂ at 20–22 °C (68–72 °F) for 8–24 hr. Stainless steel samples are scrubbed with trichlor-ethylene to remove grease, rinsed with water, allowed to drip dry, immersed in 70 percent HNO₃ for 4–5 hr at room temperature, washed with clean, potable water, and finally washed with distilled water and pretreated with 35 percent H₂O₂. Special treatments for stainless steels that have not responded well to the above passivation technique are detailed in the FMC’s Bulletin 104.

A.4.3 Plastics

Plastics and elastomer samples are thoroughly scrubbed in 0.5 percent solution of Nacconal, a synthetic detergent, rinsed with distilled water, exposed to a 10 percent HNO₃ water solution at 20–22 °C for 1 hr, then pretreated with 35 percent H₂O₂ at 20–22 °C (68–72 °F) for 8–24 hr.

A.4.4 Samples Handling

During the final rinsing, gloves or tongs should be used so samples are not touched with bare fingers. It is convenient to wash strips on a pyrex funnel as a handling medium, taking care to wash all areas. The strip is dried between two sheets of filter paper at room temperature or in a 50 °C oven and then placed in a test flask, which is immediately covered with aluminum foil.

A.5 Sample Screening

Prior to quantitative testing, a new or untried material is immersed, after chemical pretreatment, in 75 mL of H₂O₂ at room temperature for 24 hr. Violent decomposition, combustion, solution, or dimensional changes are watched for. If no unusual action occurs, the sample is then subjected to a screening at 66 °C (150.8 °F) for 24 hr.

A.6 Test Procedure

The prepared screened sample is placed in a passivated 100-mL kjeldahl flask, which is rinsed with a small volume of H₂O₂. The weight of the flask is measured to ±0.1 g. H₂O₂ of known strength (75 mL) is added to the flask, the Al covering replaced, and the flask again weighed. The initial weight of H₂O₂ solution is the difference in the two weights.

The flasks are placed in a constant temperature water bath for 1 wk at 66 °C (150.8 °F) and 4 wk at 30 °C (86 °F). The flask and its contents are then removed and cooled to room temperature, weighed, and a sample of the H₂O₂ withdrawn to determine concentration. These tests are run in duplication.
A.7 Results

The percent of AOL is calculated as follows:

\[
\text{Percent AOL} = \frac{(W_1 - W_2)}{(C W_1 \times 0.470) \times 100 \text{ percent}} ,
\]

where

- \(W_1\) = initial net weight
- \(W_2\) = final net weight
- \(C\) = initial fraction \(H_2O_2\).

Qualitative observations are also made and recorded concerning such effects as discoloration of the \(H_2O_2\) and apparent changes in the physical properties of the test material. Physical property changes include:

- For metals—any corrosion, staining, or surface change during or after test.
- For plastics—any blistering, swelling, distortion, changes in flexibility, color, transparency, or tear resistance.

A.8 Stability Test

Following the compatibility test, a sample of the remaining \(H_2O_2\) is removed and subjected to a stability test at 100 °C (212 °F) for 24 hr, according to the procedure presented in appendix B.
APPENDIX B—PROCEDURE FOR DETERMINING HYDROGEN PEROXIDE STABILITY

B.1 Introduction

This test is done to determine the storage stability of H₂O₂, and runs for 24 hr at 100 °C (212 °F). For materials evaluation, this test is run on the remaining H₂O₂ after the compatibility tests.

B.2 Apparatus

The following elements are used in the stability determining process:

• Constant temperature baths at 100 °C (212 °F) ±1 °C.
• Thermometer zero to 150 °C (302 °F) ±1 °C.
• 50-mL volumetric flask with fused 4-in neck extension.
• Distilled or deionized water and aluminum foil to cap the mouth of the flask.
• Laboratory balance.

B.3 Preparation of Test Flask

The flasks are filled with 10 percent NaOH solution for 1 hr at room temperature, rinsed with tap water and refilled with 35 percent H₂SO₄ for 3 hr at room temperature. The flasks are thoroughly rinsed with distilled water and filled with H₂O₂ for 4–6 hr. The H₂O₂ is poured out and the flasks are capped with aluminum foil without any rinsing.

B.4 Test Procedure for Concentration Above 35 Percent Hydrogen Peroxide

Measure exactly 50 mL of H₂O₂ of a known concentration into a cleaned and conditioned, special extended neck, 50-mL volumetric flask and weigh. Cap with glass cap or aluminum foil, immerse in the constant temperature bath at 100 °C (212 °F) and continue the test for 24 hr as above. After 24 hr, remove the flask from the bath and remove the cap and reweigh it on the balance used for the initial weighing. The difference is the AOL. The tests run in duplication. Normal stability of H₂O₂ is 98 to 99 percent. Samples having stability below 90 percent should be closely watched. If stability is below 80 percent, H₂O₂ should be dumped. Calculate the stability as follows:

\[
\frac{50(d_1)(C_1)(0.47) - (W_1 - W_2)}{50(d_1)(C_1)(0.47)} = \text{percent stability}
\]

where

\[
d_1 = \text{density at initial temp} \\
C_1 = \text{initial concentration} \\
W_1 = \text{initial weight} \\
W_2 = \text{final flask weight}.
\]
REFERENCES


Aluminum-Scandium Alloys: Material Characterization, Friction Stir Welding, and Compatibility With Hydrogen Peroxide

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Prepared by The Materials, Processes, and Manufacturing Department, Engineering Directorate

This Technical Memorandum describes the development of several high-strength aluminum (Al) alloys that are compatible with hydrogen peroxide (H\textsubscript{2}O\textsubscript{2}) propellant for NASA Hypersonic-X (Hyper-X) vehicles' fuel tanks and structures. The yield strengths for some of these Al-magnesium-based alloys are more than 3 times stronger than the conventional 5254-H112 Al alloy, while maintaining excellent H\textsubscript{2}O\textsubscript{2} compatibility similar to class 1 5254 alloy. The alloy development strategy is to add scandium, zirconium, and other transitional metals with unique electrochemical properties, which will not act as catalysts, to decompose the highly concentrated 90 percent H\textsubscript{2}O\textsubscript{2}. Test coupons are machined from sheet metals for H\textsubscript{2}O\textsubscript{2} long-term exposure testing and mechanical properties testing. In addition, the ability to weld the new alloys using friction stir welding has also been explored. The new high-strength alloys could represent an enabling material technology for Hyper-X vehicles, where flight weight reduction is a critical requirement.