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Research Division
NATIONAL RESEARCH CORPORATION
70 Memorial Drive
Cambridge 42, Massachusetts

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Semi-annual Technical Summary Report
January 1, 1963 - June 30, 1963

THERMODYNAMIC PROPERTIES
OF
BIMETALLIC COMPOUNDS (U)

Mr. Ludwig Fasolino El. 4-5400 Ext. 320

DOWNGRADED AT 3 YEAR INTERVALS
DECLASSIFIED AFTER 12 YEARS
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Director
Advanced Research Projects Agency
The Pentagon, Room 3D-159
Washington 25, D. C.
Attn: Advanced Propellant Chemistry Office

ARPA Order Number: 23-61
Project Code Number: 3910
Contract Date: 15 September 1961
Expiration Date: 14 November 1963
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Approved by: Allen L. Kilbanov
Program Director

Reviewed by: John V. E. Hansen
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Research Division

July 15, 1963

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INTRODUCTION

Of interest to the propellant field are several bimetallic materials whose thermochemical characteristics are required for reliable thermodynamic calculations.

This laboratory is devoted toward determining the heats of formation of selected bimetallic compounds by separately measuring the heats of solution (or reaction) of the compound and its separate constituents.

Those materials which are easily reacted with a suitable solvent are studied by a closed bomb solution calorimeter. Higher temperatures may be required for less easily dissolved materials. Materials which cannot feasibly be dissolved must be studied by other techniques. One technique gaining popularity is that of fluorine combustion. The high reactivity of fluorine provides a means of carrying out a reaction whose generated heat may be measured. This laboratory is equipped with both the solution and fluorine combustion calorimeters, thus enabling the study of a large number of compounds.
SUMMARY

It was shown that the determination of the heat of formation of aluminum dodecaboride (AlB_{12}) by solution calorimetry was not feasible. The method which would most likely be successful in studying the compound is that of fluorine combustion. An apparatus was constructed to handle fluorine and proceed with the combustion of the materials under investigation in a fluorine atmosphere. The combustion calorimeter has been calibrated and is ready for the combustion experiments.

The solution calorimeter has been successfully employed in determining the heat of formation of lithium aluminum hydride. Heats of solution of lithium, aluminum, and lithium aluminum hydride in hydrochloric acid were measured. From this data, the heat of formation of the compound was calculated to be $-26.63 + 0.31$ Kcal/mole.

The operation of the solution calorimeter at elevated temperatures ($75^\circ$C) was necessary to measure the heat of reaction of aluminum hydride with hydrochloric acid. This was accomplished with satisfactory precision. This work is presently nearing completion.
I FEASIBILITY OF SOLUTION CALORIMETRY AS APPLIED TO THE ALUMINUM - BORON SYSTEM

Thirty solvent systems were investigated as possible calorimetric fluids at two levels of temperature, 30° and 75°C. Acidic potassium periodate at the higher temperature satisfactorily dissolved powdered boron but was without effect on the aluminum dodecaboride. Of the other 29 solvent systems, none was found to be suitable. It was, therefore, concluded that the thermochemical study of the aluminum-boron system could not feasibly be undertaken by solution calorimetric methods.

Consideration of the other possible techniques has indicated that combustion calorimetry utilizing fluorine as the oxidizer may be the best approach for this particular system.

II A. ASSEMBLY OF THE FLUORINE COMBUSTION CALORIMETER

The decision to undertake the thermochemical study of the aluminum-boron system by fluorine combustion techniques necessitated additional equipment to do so. A special combustion bomb was purchased and a manifold was constructed for charging and discharging the bomb. Figure 1 shows the schematic of the manifold for handling the fluorine gas. In this system the bomb will be pressurized to the desired level, sealed by its valves, removed and
transferred to the calorimeter in which the combustion will be carried out.

II B. ELECTRICAL CALIBRATION OF FLUORINE COMBUSTION CALORIMETER

A series of electrical calibrations was performed utilizing a 23.21 A Cupron heater attached to the electrodes within this bomb. To facilitate the transfer of heat from within the bomb to the calorimeter bucket, 100 ml of water from the bucket was placed within the bomb immersing a major portion of the heater. Since the fluorine had not been delivered as scheduled, the calibrations were made with nitrogen gas within the bomb. The specific heats of these gases are nearly identical and no correction was required. The fluorine gas has since been delivered and combustion runs are planned in the near future. Table I shows the results of the electrical calibration.

III HEAT OF FORMATION OF LiAlH₄

During the initial planning of the fluorine combustion work, instructions were received from ARPA to determine the heat of formation of a compound being prepared by Reaction Motors, Li₂AlH₅. It was suggested that a preliminary study be made on the analog of this compound, LiAlH₄, to demonstrate the capabilities of our experimental method. The heat of formation of LiAlH₄ was determined by measuring the heats of solution of lithium, aluminum, and lithium.
aluminum hydride in 4.0 N hydrochloric acid. The reactions were carried out in a closed bomb, thus confining the generated hydrogen. The heat of formation was found to be -26.63 ± 0.31 kcal/mole. The details of this work were written up in a special report and distributed on April 26, 1963, and therefore, will not be repeated here. The Reaction Motors compound, Li₂AlH₅, has not been received to date. As soon as this is received, its heat of formation will be determined.

IV HEAT OF SOLUTION OF ALUMINUM HYDRIDE

Upon completion of the lithium aluminum hydride work and prior to the delivery of the fluorine gas (difficulty encountered in acquiring the requested cylinder size) a request was made and granted to attempt to measure the heats of solution of Dow's aluminum hydride, Dowane-1451, and/or Olin Mathieson's aluminum hydride, Olane-58. Samples were obtained from the respective companies and preliminary beaker tests were made to determine the conditions of reaction which would allow a meaningful calorimetric study. Both materials were found to react similarly. The conditions finally adopted were as follows:

Solvent = 6.0N hydrochloric acid
Catalyst = chloroplatinic acid (6 drops, 10% solution)
Temperature = 75°C
Under these conditions, 0.03 gram samples were found to react smoothly and completely within 20 minutes. The final solution was clear with no sign of unreacted material.

The solution calorimeter was then modified to operate under these conditions. A tantalum liner was inserted in the bomb to prevent a side reaction between the solvent and inner bomb walls. The crushing rod and baffle plates were also constructed of tantalum. A 41Al Veco thermistor was used as the temperature measuring device. Its resistance is approximately 10,000 ohms at 25°C and 1800 ohms at 75°C. The calorimeter bucket liquid and the jacket liquid were changed from water to Bayol-F oil. An immersion heater was placed in the jacket to control its temperature. It was found that the calorimeter jacket could be controlled to ± 0.005°C at 75°C as read on the Beckmann thermometer. The calorimeter bucket and bomb temperature had a linear leak rate of approximately $10^{-4}$°C min$^{-1}$. The thermistor sensitivity is approximately $5 \times 10^{-5}$°C.

The system was electrically calibrated and the results are shown in Table II. The heats of solution of aluminum, Olane-58, Dowane-1451 have been measured. The final solutions have always been clear with no residue remaining. Several more runs are required to complete the series. The heats of formation of these compounds will then be calculated. A special report will soon be issued giving the details of this work.
### TABLE I

**ELECTRICAL CALIBRATION OF FLUORINE COMBUSTION CALORIMETER**

<table>
<thead>
<tr>
<th>Run No.</th>
<th>( L_{\text{cal}} )</th>
<th>( \Delta R ) ohms</th>
<th>( E_{\frac{1}{2} R} ) cal ohm (^{-1} )</th>
</tr>
</thead>
<tbody>
<tr>
<td>6</td>
<td>1284.5</td>
<td>36.528</td>
<td>35.150</td>
</tr>
<tr>
<td>8</td>
<td>1258.1</td>
<td>35.868</td>
<td>35.093</td>
</tr>
<tr>
<td>9</td>
<td>1282.2</td>
<td>36.431</td>
<td>35.194</td>
</tr>
<tr>
<td>10</td>
<td>1417.1</td>
<td>40.291</td>
<td>35.171</td>
</tr>
<tr>
<td>13</td>
<td>1246.0</td>
<td>35.478</td>
<td>35.120</td>
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<tr>
<td>15</td>
<td>1235.4</td>
<td>35.191</td>
<td>35.106</td>
</tr>
<tr>
<td>16</td>
<td>1240.1</td>
<td>35.318</td>
<td>35.113</td>
</tr>
</tbody>
</table>

**Mean**

35.135

**Uncertainty (2σ)**

0.028

0.079
## TABLE II

ELECTRICAL CALIBRATION OF SOLUTION CALORIMETER AT 75°C

<table>
<thead>
<tr>
<th>Run No.</th>
<th>$E_{2}$ cal</th>
<th>$\Delta R$, ohms</th>
<th>$E - E_{2}/\Delta R$, cal ohm$^{-1}$</th>
</tr>
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<tbody>
<tr>
<td>1A</td>
<td>285.94</td>
<td>8.609</td>
<td>33.214</td>
</tr>
<tr>
<td>2A</td>
<td>285.89</td>
<td>8.750</td>
<td>32.673</td>
</tr>
<tr>
<td>3A</td>
<td>281.50</td>
<td>8.629</td>
<td>32.623</td>
</tr>
<tr>
<td>4A</td>
<td>215.46</td>
<td>6.556</td>
<td>32.865</td>
</tr>
<tr>
<td>5A</td>
<td>297.69</td>
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<table>
<thead>
<tr>
<th>Mean</th>
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</thead>
<tbody>
<tr>
<td>Uncertainty (2σ)</td>
<td></td>
<td>.220</td>
</tr>
<tr>
<td>%</td>
<td></td>
<td>.66</td>
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