HEAT-RELEASE KINETICS OF AMMONIUM PERCHLORATE IN PRESENCE OF CATALYSTS AND FUEL**

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† This work was sponsored by the Office of Naval Research, Department of the Navy, under Contract Nonr-3415(00), Authority NR 092-507/10-13-64.
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IN PRESENCE OF CATALYSTS AND FUEL*†

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March 9, 1967

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The rate of heat release of ammonium perchlorate-fuel or ammonium perchlorate-catalyst or both under adiabatic condition was studied as a function of temperature. In the presence of catalysts, the autocatalytic feature associated with nucleation of pure ammonium perchlorate disappears and the adiabatic reaction kinetics follows Arrhenius-type behavior. Studies made with ammonium perchlorate-fuel show that measurable heat release occurs at temperatures below the phase transition temperature of ammonium perchlorate and that the rate of heat generation per mass of material is greater than that of pure ammonium perchlorate. The rate of exothermic decomposition of pure ammonium perchlorate is unaffected by an increase in pressure. This observation suggests that the heat release is controlled by homogeneous or heterogeneous reactions in the condensed phase.
Introduction

Measurements of the decomposition kinetics of ammonium perchlorate (AP) were described in a previous publication. In the measurement studies, the rate of heat release under adiabatic conditions was determined as a function of temperature and related to the rate of reaction and nucleation. These determinations have now been extended to AP samples containing catalysts and fuel. Several measurements have also been carried out of the kinetics of exothermic decomposition of AP at elevated pressures to explore the effect of total gas pressure on the reaction.

Experimental Details

As described in detail in Reference 1, the apparatus used in the adiabatic experiments employs a sample in the form of a pressed wafer (30mm in diameter, 4mm in thickness) prepared by pressing about 5 gram of AP-additive mixture at ~ 25000 psi. By this procedure, a hard, durable wafer was produced with a porosity of about 4 to 5 percent in the case of pure AP. The samples containing added fuel were prepared by cutting out a desired dimension of casted AP-carboxy-terminated polybutadiene (AP-CTPB) propellant. A small radial hole near the center of the sample was drilled in each wafer to accommodate a glass sheathed chromel-alumel thermocouple (3 mil wire). The wafer was then clamped between two electrically heated plates that were held in position by spring-loaded bolts. During the course of an experiment, the temperature of the heating plates was continually adjusted to match within less than $0.1^\circ$C that of the interior of the sample, i.e., adiabatic conditions. The sample thermocouple emf was recorded on a strip-chart recorder, while the emf difference between the block and sample thermocouples was displayed
on the indicating meter of a microvolt amplifier and kept within the prescribed limits by manual control of the electric power fed to the heating plates.

In a typical experiment, the block assembly, composed of heating plates and sample, was placed in a small furnace held at a temperature of about 250°C. The sample was brought to a temperature slightly above 240°C by supplying electrical power to the heater plates. The power was then reduced to establish within the assembly a constant temperature in the range 240°-250°C. Several minutes were generally required for this process. As the sample temperature began to rise above that of the heating plates due to exothermic decomposition, electrical power was supplied to the heater plates at a rate sufficient to maintain a uniform temperature throughout the sample. The sample temperature history was recorded until the experiment was terminated due to an excessive rate of temperature rise (more than 100°C/min). For the high pressure studies, a steel vessel was employed that could be heated and pressurized to 200 psi.

The reagent grade crystalline AP was ball milled and sieved into four fractions with a particle diameter range: (a) less than 43μ, (b) 43-61μ, (c) 61-86μ, and (d) 88-124μ. The material of a specified particle size range was then mixed with the desired amount of powdered additive, fuel, or catalysts or both in a mortar and pestle, and pressed into a pellet. The compositions of the mixtures studied are summarized in Table 1. The composition of the casted AP-CTPB propellant was 75wt% AP/25wt% binder. The AP particle size spectrum for this sample was bimodal, ranging from 2 to 200μ. The particle size of the additives (Fe₂O₃, Cr₂O₃, and Norit A) was estimated from microscopic examination.

**Experimental Results and Discussions**

Examination of the experimental data was based on the theoretical analysis of adiabatic reaction kinetics involving a first order reaction. According to this analysis, the rate of temperature rise (dT/dt) can be related to the effective activation energy E for the process: 
\[
dT/dt = p e^{-E/RT}
\]
where the constant p is given by 
\[
p = QNA/pc,
\]
and Q represents
the molar heat of reaction, \( N \) the molar density, \( A \) the preexponential coefficient of the Arrhenius rate equation, \( \rho \) the mass density, and \( C \) the specific heat at constant pressure. A typical example of the experimental data for the catalyzed decomposition of AP is shown in Figure 1. The results for a series of additives are summarized in Table 2.

Reproducible results were obtained with samples containing 2.9wt\% CC* and 4.8wt\% CC (Table 2); however, the data obtained were less reproducible than similar data for pure AP, because of (a) difficulty in establishing isothermal starting conditions, (b) sample fracture during decomposition, and (c) nonuniform distribution of catalyst throughout the sample. In the presence of 4.8wt\% \( \text{Cr}_2\text{O}_3 \) in the sample, ignition occurred at about 280\(^\circ\)C so that the adiabatic studies were restricted to a small temperature interval.

The addition of carbon caused fragmentation of the samples when the rates of temperature rise exceeded 30 deg/min. Similar but less reproducible results were obtained on substitution of Cabot carbon for Norit A carbon. Studies of the decomposition rates of three carbon- and catalyst-containing mixtures† (Norit A) were attempted: \( \text{AP/C (97\%/2\%)} \), \( \text{AP/CC/C (95/2\%/2\%)} \), and \( \text{AP/\text{Cr}_2\text{O}_3/C (95/2\%/2\%)} \). No definitive data were obtainable for these ternary carbon-containing mixtures since ignition occurred near 260\(^\circ\)C, corresponding to a rate of temperature rise of only a few degrees/min.

Hematite (\( \text{Fe}_2\text{O}_3 \)), a well known burning rate promoter, was examined as a catalyst for AP decomposition. The results obtained for a mixture containing 4.8wt\% \( \text{Fe}_2\text{O}_3 \) extend to a rate of temperature of 100 deg/min. It is apparent that catalytic additives, such as metal oxides, alter the course of decomposition of \( \text{NH}_4\text{ClO}_4 \) and cause ignition or explosion of the material. In the presence of sufficient concentration of such additives, the autocatalytic feature of the decomposition of pure \( \text{NH}_4\text{ClO}_4 \) disappears. In such cases, the rate of decomposition follows Arrhenius-type behavior even at the lowest temperatures examined.

---

* Copper chromite.
† All values in weight percent.
The experimental heat-release rates from the propellant mixture AP-CTPB bring out the interesting fact that the activation energy for the propellant mixture is nearly the same as that previously obtained for pure AP. We may therefore conclude that the rate-controlling process for decomposition is the same in both cases, but the net amount of heat released per mass of material decomposed is greater because of the presence of added CTPB. Some of the intermediates or products resulting from the decomposition of AP are interacting with the added organic fuel. Apparently, such a reaction with heat generation occurs already at temperatures below the transition temperature of AP (240°C) as demonstrated by the curve in Figure 2. Since exothermic heat generation by homogeneous or heterogeneous reaction in the condensed phase takes place at significantly lower temperatures than in the case of pure AP, the presence of binder should enhance ignition and combustion of such a propellant.

To determine the influence of total gas pressure on the decomposition of AP, the adiabatic measurements were extended to elevated pressures. As shown in Figure 3, the rate of exothermic decomposition of pure AP is unaffected by a more than tenfold increase in pressure. The absence of a pressure effect suggests that, at least for pure AP, the heat release observed in our experiments is controlled by reactions in the solid phase. Heterogeneous reactions at the gas/solid interface would be expected to be influenced by pressure except in the case of high surface density of adsorbate—i.e., a condition in which the rate of reaction on the surface itself rather than the transport of reactant to the surface is rate-limiting. In the presence of catalysts, a different reaction model may prevail. As a matter of fact, the apparent correlation between burning rate of AP/catalyst propellants and catalytic activity for ammonia oxidation strongly indicates the importance of heterogeneous reactions in ignition and combustion.

The implications of these condensed-phase, exothermic reactions on propellant properties are currently under investigation. A measure of their contribution can be seen from the data in Table 2, in which we have computed the heat flux generated by exothermic chemical reaction at a temperature of 8000K, representative of the surface temperature of an AP propellant during steady-state burning and ignition.
References


Table 1. Composition of Mixtures Used in Adiabatic Studies

<table>
<thead>
<tr>
<th>Ammonium perchlorate*</th>
<th>Additive</th>
<th>Type</th>
<th>Percent by wt</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td></td>
<td>43-61; 88-124</td>
<td>copper chromite†</td>
</tr>
<tr>
<td></td>
<td></td>
<td>88-124</td>
<td>chromic oxide*</td>
</tr>
<tr>
<td></td>
<td></td>
<td>88-124</td>
<td>cupric oxide‡</td>
</tr>
<tr>
<td></td>
<td></td>
<td>43-61; 88-124</td>
<td>ferric oxide*</td>
</tr>
<tr>
<td>88-124</td>
<td></td>
<td>carbon (Norit A;</td>
<td></td>
</tr>
<tr>
<td></td>
<td></td>
<td>Cabot-Sterling VR)</td>
<td></td>
</tr>
<tr>
<td></td>
<td></td>
<td>88-124</td>
<td>carbon/copper</td>
</tr>
<tr>
<td></td>
<td></td>
<td>chromite</td>
<td></td>
</tr>
<tr>
<td>88-124</td>
<td></td>
<td>88-124</td>
<td>carbon/chromic oxide</td>
</tr>
</tbody>
</table>

* From Matheson, Coleman, and Bell (reagent grade).
† From Harshaw Chemical Company.
‡ From J. T. Baker Company.
Table 2. Results of Adiabatic Studies

<table>
<thead>
<tr>
<th>Additive</th>
<th>Wt percent</th>
<th>Ammonium perchlorate†</th>
<th>Additive</th>
<th>$\frac{dT}{dt}$ (deg/sec)</th>
<th>$\dot{q}^{**}$ cal/cm³ sec</th>
</tr>
</thead>
<tbody>
<tr>
<td>Copper chromite</td>
<td>2.9</td>
<td>43-61</td>
<td>~ 3.5‡</td>
<td>$10^{13.8} \exp (-40,000/RT)$</td>
<td>$8.4 \times 10^2$</td>
</tr>
<tr>
<td>Copper chromite</td>
<td>4.8</td>
<td>88-124</td>
<td>~ 3.5‡</td>
<td>$10^{16.5} \exp (-43,000/RT)$</td>
<td>$6.6 \times 10^4$</td>
</tr>
<tr>
<td>Copper chromite</td>
<td>4.8</td>
<td>43-61</td>
<td>~ 3.5‡</td>
<td>$10^{17.7} \exp (-46,000/RT)$</td>
<td>$1.7 \times 10^5$</td>
</tr>
<tr>
<td>Chronic oxide</td>
<td>4.8</td>
<td>88-124</td>
<td>&lt; 0.5</td>
<td>$10^{27.3} \exp (-68,000/RT)$</td>
<td>$6.6 \times 10^8$</td>
</tr>
<tr>
<td>Carbon (Norit A)</td>
<td>2.5</td>
<td>88-124</td>
<td>&lt; 0.5 to 30§</td>
<td>$10^{19.9} \exp (-52,500/RT)$</td>
<td>$4.2 \times 10^5$</td>
</tr>
<tr>
<td>Ferric oxide</td>
<td>4.8</td>
<td>88-124</td>
<td>&lt; 0.5</td>
<td>$10^{12.8} \exp (-34,300/RT)$</td>
<td>$4.2 \times 10^3$</td>
</tr>
<tr>
<td>Ferric oxide</td>
<td>4.8</td>
<td>43-61</td>
<td>&lt; 0.5</td>
<td>$10^{13.8} \exp (-36,600/RT)$</td>
<td>$8.3 \times 10^3$</td>
</tr>
</tbody>
</table>

* Average of 4 to 8 experiments.
† Particle size screened by mechanical sieving.
‡ Reference 2.
§ Nonspherical particles.
** Heat generation rate at 800°K. For these calculations a density of 1.6 g/cm³ and a specific heat of 0.4 cal/g deg were employed.
FIG. 1 THE ADIABATIC RATE OF DECOMPOSITION OF AN NH$_4$ClO$_4$/4.8wt% CC WAFER
FIG. 2 ADIABATIC DECOMPOSITION RATE OF AP-CTPB PROPELLANT
FIG. 3 ADIABATIC DECOMPOSITION RATE OF AP AT 1 AND 13.5 ATMOSPHERES (particle size 61-80μ)

OPEN SYMBOL 1 atm, AIR
CLOSED SYMBOL 13.5 atm, N₂
The rate of heat release of ammonium perchlorate-fuel or ammonium perchlorate-catalyst or both under adiabatic condition was studied as a function of temperature. In the presence of catalysts, the autocalytic feature associated with nucleation of pure ammonium perchlorate disappears and the adiabatic reaction kinetics follow Arrhenius-type behavior. Studies made with ammonium perchlorate-fuel show that measurable heat release occurs at temperatures below the phase transition temperature of ammonium perchlorate and that the rate of heat generation per mass of material is greater than that of pure ammonium perchlorate. The rate of exothermic decomposition of pure ammonium perchlorate is unaffected by an increase in pressure. This observation suggests that the heat release is controlled by homogeneous or heterogeneous reactions in the condensed phase.
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