FUNDAMENTAL STUDIES OF ROCKET COMBUSTION CHEMISTRY AND COMBUSTION INSTABILITY

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This report provides a summary of results obtained over the past year in fundamental studies of the chemistry of solid rocket combustion and of combustion instability in liquid rocket combustion. A coordinated research program has been initiated to study separate aspects of the complex multidimensional, multiphase and multicomponent reacting flow characteristic of rocket combustion. The focus of the combustion chemistry studies is on flame reactions of gas phase species formed as decomposition products near the surface of the propellant. These flames typically involve a variety of fuel species and nitrogen oxides as oxidizers. The focus of the combustion instability studies is the effect of atomization and vaporization processes on the oscillatory conditions characteristic of combustion instability. The research objectives, progress and research in progress in each of these areas are summarized in the report.
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A. FUNDAMENTAL STUDIES OF ROCKET COMBUSTION CHEMISTRY

A.1 RESEARCH OBJECTIVES

The combustion of solid rocket propellants and other energetic materials is a complex multidimensional and multiphase process involving a wide variety of chemical species. The very high pressure and temperature conditions of practical rocket combustion chambers are at present inaccessible by most conventional diagnostic techniques. The study of these coupled phenomena in situ, therefore, has not been possible in sufficient detail to develop a complete understanding of the chemistry and physics of the combustion process. The objective of most recent studies of the combustion of these materials has been to study separate aspects of the overall process in an effort to provide a comprehensive understanding of the combustion mechanism. This study is one component of that coordinated investigation and has as its focus the gas phase reactions associated with the combustion of these solid fuels.

The decomposition of many of these solid energetic materials during combustion leads to the formation of gaseous hydrocarbon fuel species and oxides of nitrogen which serve as oxidizers (ref. 1 and 2). The reactions of these decomposition products above the propellant surface lead to a gaseous flame which can provide heat which is transferred back to the propellant surface and can thereby influence the burning rate. The purpose of this paper is to summarize the current status of studies we have undertaken of model gas phase flames associated with the combustion of nitramine bases solid rocket propellants. These studies consist of measurements of the structure of stable and unstable species concentration profiles and temperature in laminar, premixed, flat flames of fuel/NO$_2$ mixtures at low pressure. The experimental measurements are then compared to calculations of the concentration profiles using a one dimensional flame code which models the transport processes and chemistry of the flame. The transport processes include species diffusion and thermal conduction through the flame and the chemistry is modeled by a detailed chemical kinetic reaction mechanism.

The basic mechanism used for the modeling work is a subset of the 331-reaction mechanism of Volponi and Branch (ref. 3). Reactions removed were those involving species with low concentrations, including condensed phase molecules, and molecules containing more than three carbon atoms. The resulting mechanism contains 272 reactions. This mechanism was the basis for all of the modeling described in this paper. The flame code used to solve the one dimensional flame equations was by Kee et al. (ref. 4).

A.2 RESEARCH APPROACH AND RECENT RESULTS

A.2.1 FLAMES SUPPORTED BY N$_2$O AS OXIDIZER

Several recent studies have focused on measuring the temperatures and species concentrations in flames supported by N$_2$O. Habeebullah et al. (ref. 5), Zabarnick (ref. 6), and Vandooren et al. (ref. 7) studied CH$_4$ reacting with N$_2$O. Habeebullah measured stable species concentrations using probe sampling, radical species concentrations using laser-induced fluorescence (LIF), and flame temperatures using thermocouples and LIF. Zabarnick used LIF to measure stable and radical species concentrations and flame temperatures. Dindi et al. (ref. 8) studied CO reacting at N$_2$O, and used probe sampling to measure stable species concentrations, LIF to measure unstable species concentrations, and thermocouples to measure flame temperatures.

We have modeled the structure of all of these flames using our 272 step reaction mechanism. It was
found that in order to have best agreement with the full range of data represented by these results, the rates of several of the reactions needed to be adjusted as indicated in Table 1. The most significant of these changes was for the N\textsubscript{2}O decomposition which was adjusted to within the upper range of the literature data for this reaction. This change was needed to model the methane flame data correctly. The other reactions listed in Table 1 were adjusted to provide good agreement for the CO/N\textsubscript{2}O flame data. The results of the modeling for the most important cases studied are given below for the flames with N\textsubscript{2}O as oxidizer.

CO-N\textsubscript{2}O Flame Structure

Low pressure, laminar, premixed flames of CO-N\textsubscript{2}O have been stabilized over a rectangular flat-flame burner (ref. 8). Laser-induced fluorescence spectroscopy was used to establish the absence of CN, CH, NH, NH\textsubscript{2} and OH in these flames. Gas chromatographic sample analysis was used to determine the CO, CO\textsubscript{2}, N\textsubscript{2}O, NO, N\textsubscript{2} and O\textsubscript{2} concentration profiles for three CO-N\textsubscript{2}O flames having equivalent ratios of 1.00, 1.32, 1.50. Lean flames could not be stabilized. Temperature profiles for all three flames were measured using R type thermocouples. Measured temperature profiles were corrected for radiation losses. These flames are considerably lifted above the burner and contain a single luminous zone. The main feature of these flames is the absence of any reactive intermediates except oxygen atom. We have eliminated reactions involving the element H and reduced the 272 step mechanism to 27 reactions. This mechanism was then used to model these CO-N\textsubscript{2}O flame data.

A combined "elementary reaction contribution" and "sensitivity" analysis showed that only four of the 27 elementary reactions used in the original kinetic mechanism were important in the production of or the consumption of major species in the flame. A comparison of the calculated concentration profiles using the original 27-reaction and the four-reaction mechanisms for all three CO-N\textsubscript{2}O flames studied in this investigation showed that the difference between the two profiles was always less than 1% of the total concentration. Hence we propose the following four-step mechanism for the CO-N\textsubscript{2}O flames:

\[
\begin{align*}
\text{CO} + \text{N}_2\text{O} & \rightarrow \text{CO}_2 + \text{N}_2 \\
\text{N}_2\text{O} + \text{M} & \rightarrow \text{N}_2 + \text{O} + \text{M} \\
\text{N}_2\text{O} + \text{O} & \rightarrow \text{N}_2 + \text{O}_2 \\
\text{N}_2\text{O} + \text{O} & \rightarrow \text{NO} + \text{NO}
\end{align*}
\]

The first of the four reactions listed above is the most important reaction for CO-N\textsubscript{2}O flames. It accounts for almost all the CO consumption and nearly 90% of the N\textsubscript{2}O consumption. The second reaction (N\textsubscript{2}O+M→N\textsubscript{2}+O+M) is an important reaction for kineticists. It plays a key role in various environments for NO\textsubscript{x} formation or decomposition. It is also used for generating O atoms in studies of elementary oxidation reactions. The last two reactions of the four-step mechanism listed above control the concentration profiles of O\textsubscript{2} and NO.

CH\textsubscript{4}-N\textsubscript{2}O Flame Structure

Laminar, premixed flat flames of CH\textsubscript{4} with N\textsubscript{2}O have been stabilized and studied at 50 torr (ref. 5). This study represents the first nearly complete study of the structure and kinetics of CH\textsubscript{4}-N\textsubscript{2}O flames including stable and unstable species measurements and detailed chemical kinetic modeling. Three flames were investigated, with slightly fuel rich, near stoichiometric and lean mixtures. Flame modeling has been done using the 272 step reaction mechanism and the reaction mechanism evaluated.
The results of the flame calculations and the experiments are given in Figure 1. The comparison shows good quantitative and qualitative agreement between the measured and calculated profiles especially for stable species. The modeling simulation clearly predicts the general flame structure and species concentration profiles. For the flame radicals, the theoretical results are in good agreement with the experimental results for NH, CN and OH. The computed maximum for the CH radical is shifted in the flame. The exact prediction of the radical concentrations using flame modeling is difficult owing to the high reactivity of these radicals, especially CH.

The reaction mechanism starts with the thermal decomposition of N\textsubscript{2}O to N\textsubscript{2} and O which is the initial reaction for radical formation in the flame. The oxygen atom formed from this reaction reacts with H\textsubscript{2} to form a pool of H and OH radicals which then propagate the rest of the mechanism. The results of the sensitivity analysis also shows that N\textsubscript{2}O is not totally decomposed to N\textsubscript{2}, but it forms some NH and NO molecules directly through reaction with H atoms. The mechanism also shows that methane has a long induction period before being consumed mainly through radical reactions (especially with OH and H) to form methyl (CH\textsubscript{3}) intermediate. Once methyl intermediate is formed it has two reaction paths to follow. The first path is to form formaldehyde (CH\textsubscript{2}O) which then starts a reaction path to form CO\textsubscript{2}. The major reaction for CO conversion to CO\textsubscript{2} in this path is through its reaction with OH radical. The second path that CH\textsubscript{4} follows is to form CH\textsubscript{3} intermediate which then forms CH through radical reactions. The CH formed through this path reacts with NO from N\textsubscript{2}O decomposition to produce NH or CN radicals. This reaction forms the first link in hydrocarbon-nitrogen interactions. The high CN and NH concentration in the flame is attributed to this reaction. However, the reaction of nitrous oxide (N\textsubscript{2}O) with H atoms is another important source for the formation of these two radicals (CN and NH). The final products in the exhaust gas were mainly N\textsubscript{2}, H\textsubscript{2}O, CO and CO\textsubscript{2} with small amounts of NO.

A.2.2 FLAMES SUPPORTED BY NO\textsubscript{2} AS OXIDIZER

Five flames supported by NO\textsubscript{2} as the oxidizer have been studied recently and were tested against our 272 step reaction mechanism. Volponi and Branch (ref. 3 and 9) studied H\textsubscript{2} and C\textsubscript{2}H\textsubscript{2} reacting with NO\textsubscript{2} in an argon diluent. They measured stable species concentrations using probe sampling, radical species concentrations using LIF, and flame temperatures using thermocouples and LIF. Branch et al. (ref. 10) studied two different flames supported by NO\textsubscript{2}, CH\textsubscript{4}/NO\textsubscript{2}/O\textsubscript{2} and CH\textsubscript{2}O/NO\textsubscript{2}/O\textsubscript{2}. They measured stable species concentrations using probe sampling, unstable species concentrations using LIF, and flame temperatures with a thermocouple. Zabarnick (ref. 11) also studied the CH\textsubscript{4}/NO\textsubscript{2}/O\textsubscript{2} flame, using LIF to measure flame temperatures and stable and radical species concentrations. The results of our current modeling of the hydrogen and the acetylene flames are most indicative of the reaction mechanisms involved and are discussed below.

H\textsubscript{2}-NO\textsubscript{2} Flame Structure

Measurements of the composition of stable and unstable species and temperature in laminar, premixed, flat flames of H\textsubscript{2}-NO\textsubscript{2}-Ar have been made and compared to the structure calculated with a flame code including detailed chemical kinetics (ref. 9). No previous detailed flame structure measurements and chemical kinetic modeling of this flame have been presented. Similar measurements and calculations are reported for a companion H\textsubscript{2}-O\textsubscript{2}-Ar flame in order to provide a comparison to a previously well characterized flame. We have modeled these flame measurements using the 272 step reaction mechanism with the carbon species removed. The resulting 87 step mechanism gives good agreement to the experimental data as shown in Figure 2.
In contrast to the H$_2$O-O$_2$-Ar flame, the rate and sensitivity calculations for H$_2$, NO$_2$, H$_2$O and OH in the H$_2$-NO$_2$-Ar flame show that species net reaction rates are usually dominated by a single reaction. Almost all of the H$_2$ consumption and H$_2$O formation is by OH+H$_2$=H$_2$O+H. The consumption of NO$_2$ and formation of NO is almost entirely by NO$_2$+H=NO+OH. The sum of these two reactions gives the global reaction of the flame H$_2$+NO$_2$=NO+H$_2$O. There is some formation of O$_2$ early in the flame by NO$_2$+O=NO+O$_2$, followed by O$_2$ consumption by O$_2$+H=O+OH. The OH is a balance of formation by NO$_2$+H=NO+OH and consumption by OH+H$_2$=H$_2$O+H.

The reactions with greatest negative sensitivity for H$_2$ and NO$_2$ have the highest positive sensitivity for NO and H$_2$O. The reactions with the largest positive sensitivity for H$_2$ and NO$_2$ likewise have the largest negative sensitivity for NO and H$_2$O. The OH is primarily sensitive to its major formation and consumption reactions.

The importance of the reaction H$_2$+NO$_2$=HNO$_2$+H to the H$_2$-NO$_2$-Ar flame mechanism is observed in the sensitivity and analysis for H$_2$, NO$_2$, NO, H$_2$O and OH. This reaction provides the most important initiation step. It also has a significant effect on the OH profile by producing H early in the flame and promoting the formation of OH by NO$_2$+H=NO+OH. The HNO$_2$ is subsequently consumed by several reactions. Using a mechanism that contains only the three reactions

\[
\begin{align*}
\text{NO}_2 + \text{H} &= \text{NO} + \text{OH} \\
\text{OH} + \text{H}_2 &= \text{H}_2\text{O} + \text{H} \\
\text{H}_2 + \text{NO}_2 &= \text{HNO}_2 + \text{H}
\end{align*}
\]

gives a flame model that compares to within 5% of the model using the entire reaction mechanism.

C$_2$H$_2$-NO. Flame Structure

Volponi and Branch (ref. 9) studied C$_2$H$_2$ reacting with NO$_2$ in an argon diluent at a pressure of 25 torr. We modeled this flame using the 272 step reaction mechanism and the measured temperature profile. The modeling results using this mechanism are close to those Volponi and Branch obtained using the full, 331-reaction mechanism from which the 272-reaction mechanism was derived. This indicates that modeling results are not significantly affected using the smaller mechanism.

Measured and modeled mole fraction profiles for major species are shown in Figure 3. Discrepancies can be seen near the burner surface, as the modeling overpredicts the surface mole fractions of unreacted species and underpredicts the surface mole fraction of product species. Agreement is good, however, in the general trends shown in the profiles, and in the final mole fractions obtained in the flame. The modeling was used to find the principal reactions involved in each species' production/consumption. The principal reaction consuming C$_2$H$_2$ is C$_2$H$_2$+OH=C$_2$H+H$_2$O; the principal reaction consuming NO$_2$ is NO$_2$+H=NO+OH; the principal reaction producing NO is NO$_2$+H=NO+OH; the principal reaction producing O$_2$ is 2NO$_2$=2NO+O$_2$; the principal reaction producing CO$_2$ is NO$_2$+CO=NO+CO$_2$; and the principal reaction producing H$_2$O is OH+C$_2$H$_2$=C$_2$H+H$_2$O. The importance of radicals in breaking down the unreacted species is evident.

Comparisons were also made between modeled and measured mole fractions for the radical species OH and CN. There were discrepancies in the calculated and measured trends for these species. The principal reaction involving OH production is NO$_2$+N=NO+OH, and the principal reaction involving CN production is HCN+OH=CN+H$_2$O, and the principal reaction involving CN production is HCN+OH=CN+H$_2$O.
Volponi and Branch (ref. 3) also modeled C₂H₂ reacting with O₂ with Argon gas as a diluent at a pressure of 25 torr. We modeled this flame using the 272 step mechanism and the measured temperature profile. The modeling's accuracy was on a par with what was obtained for the CH₄/N₂O flame. Modeling was generally in good agreement with experimental data for major species concentrations in the flame, with the exception of H₂O, whose concentration was slightly underpredicted. Modeling and experiments agreed in the general trends for the concentration of radical species OH and CH, but modeling overpredicted the peak concentrations of both species by a factor of two. As with the CH₄/NO₂ flame, our modeling results compared well with those Volponi and Branch obtained using the 331-reaction mechanism.

A.2.3 FLAMES SUPPORTED BY NO AS OXIDIZER

Zabarnick (ref. 6) studied a CH₄/NO/O₂ flame at 63 torr, using LIF to measure temperature and stable and radical species concentrations. This flame was modeled using the measured temperature profile and the 272 step mechanism.

The measured NO concentration profile showed a more pronounced drop-off to a steady-state value than modeling predicted. Also, the final calculated value of NO in the flame is 50% higher than what was measured; NO was the only stable species Zabarnick performed measurements on in this work.

Modeling and experimental results were compared for radical species. The predicted peak concentration of CN leads the experimentally measured peak by about 1 mm. The predicted OH concentration showed a drop-off late in the flame that was not detected experimentally. The predicted NH peak led the measured peak by about .5 mm. The predicted CH peak led the measured peak by about 1 mm also. Note that modeling now leads the experimental data for the location of peak radical concentrations, while for the CH₄/N₂O flame also measured by Zabarnick (ref. 6) modeling predictions generally lagged the experimental data.

A.2.4 BURNING VELOCITY OF FUEL/NOₓ LAMINAR PREMIXED FLAMES

The 272 step reaction mechanism was used to model the burning of free-standing flames supported by N₂O and NOₓ and the modeled flame speed was compared to the experimental results obtained by Parker and Woltford (ref. 12) using the bunsen burner technique (Table 2). Also shown are the burning velocities obtained with the Miller and Bowman (ref. 13) mechanism without the modifications discussed in the present investigation. The results indicate that generally good agreement is obtained between the modeling and the experimental data and that the 272 step mechanism gives better results than earlier mechanisms. The burning velocity of the flames with N₂O are significantly higher than those with NOₓ as oxidizer. Two of the calculations are discussed below in more detail in order to indicate the importance of fuel and oxidizer decomposition reactions in flame propagation.

Table 1 gives the calculated and measured results for the burning rate of a CH₄/N₂O flame. The calculated flame velocity is 90 cm/sec, very close to the measured velocity of 105 cm/sec. In addition, for this flame we were able to perform analysis on what reactions contribute most strongly to a given species's production/consumption. It was found that the major reactions leading to the decomposition of CH₄ are the following, in decreasing order of their rate of CH₄ consumption:
\[
\begin{align*}
\text{CH}_4 + H &= \text{CH}_3 + \text{H}_2 \\
\text{CH}_4 + \text{OH} &= \text{CH}_3 + \text{H}_2\text{O} \\
\text{CH}_4 + O &= \text{CH}_3 + \text{OH}
\end{align*}
\]

The major reactions leading to the breaking down of N\textsubscript{2}O were found to be, in decreasing order of the amount of N\textsubscript{2}O they consume:

\[
\begin{align*}
\text{N}_2\text{O} + H &= \text{H}_2 + \text{OH} \\
\text{N}_2\text{O} + M &= \text{N}_2\text{O} + M
\end{align*}
\]

The importance of the radicals H, OH and O are clearly evident in the propagation of this flame.

A free-standing \text{C}_2\text{H}_4 flame at a pressure of 0.1 atm was modeled and the calculated flame speed was compared to that measured by Parker and Wolfhard (ref. 10). The results are included in Table 2, and as can be seen, the agreement is excellent for this flame. It was difficult to obtain a converged solution modeling many NO\textsubscript{2}-supported flames, so there are not more cases reported at this time. It is possible that the larger mechanism used to model this flame aided in obtaining better agreement with experimental data than was obtained in several of the cases of N\textsubscript{2}O-supported flames. More cases would need to be run, however, to draw conclusions on the relative ability of the two mechanisms to model free-standing flames supported by NO\textsubscript{2} and N\textsubscript{2}O.

Modeling results for this flame showed that the top four reactions contributing to the decomposition of \text{C}_2\text{H}_4 in the flame were the following:

\[
\begin{align*}
\text{C}_2\text{H}_4 + \text{OH} &= \text{C}_2\text{H}_3 + \text{H}_2\text{O} \\
\text{C}_2\text{H}_4 + M &= \text{C}_2\text{H}_2 + \text{H}_2 + M \\
\text{C}_2\text{H}_4 + H &= \text{C}_2\text{H}_3 + \text{H}_2 \\
\text{C}_2\text{H}_4 + O &= \text{CH}_3 + \text{HCO}
\end{align*}
\]

The top four reactions leading to the decomposition of NO\textsubscript{2} were found to be:

\[
\begin{align*}
\text{NO}_2 + N &= \text{NO} + \text{OH} \\
\text{NO}_2 + \text{CO} &= \text{NO} + \text{CO}_2 \\
\text{NO}_2 + M &= \text{NO} + O + M \\
\text{NO}_2 + \text{OH} &= \text{HO}_2 + \text{NO}
\end{align*}
\]

Initiation of the chemistry in these flames can be seen to be closely tied to reactions producing radical species.

\section{A.2.5 SUMMARY AND CONCLUSIONS}

We have completed detailed comparisons of calculations and measurements of the structure and burning velocity of fuel/NO\textsubscript{2} mixtures using a 272 step reaction mechanism. The comparison between the calculated flame structure and the experimental flame structure for stable species was found to be very good for fuel/N\textsubscript{2}O flames and good for fuel/NO\textsubscript{2} flames. The concentration profiles for radical species were found to be generally well represented qualitatively but not well represented quantitatively. It was concluded that, despite some remaining difficulties with the reaction mechanism, it appeared to be reliable.
in describing the overall combustion behavior of a wide range of fuel and oxidizer mixtures.

The most important reactions of the oxidizer (N\textsubscript{2}O or NO\textsubscript{2}) are with H atoms and, to a lesser extent, with CO. Reaction of either oxidizer with H is a chain propagating reaction, in contrast with the chain branching reaction of H with O\textsubscript{2} which is of equal importance in fuel oxidation by O\textsubscript{2}. In addition, the reaction of NO\textsubscript{2} with H is slower than N\textsubscript{2}O with H. Finally, the reaction of N\textsubscript{2}O with CO can be significant both in consumption of CO and formation of CO\textsubscript{2}. This situation is again in contrast with the oxidation chemistry of systems by O\textsubscript{2} in which the conversion of CO to CO\textsubscript{2} is almost entirely by reaction of CO with OH. The difference between the use of N\textsubscript{2}O or NO\textsubscript{2} as oxidizer is that the former produces primarily N\textsubscript{2} while the latter produces NO. The subsequent slow reduction of NO to N\textsubscript{2}, even when it is thermodynamically favored, accounts for the most striking difference between the two oxidizers. The most important effects of the oxidizer, therefore, are that the nitrogen oxides are less effective chain carriers and lead to slower reaction rates compared to O\textsubscript{2}.

The oxidation reactions of CH\textsubscript{4} in the presence of NO\textsubscript{2} is generally similar to the oxidation by O\textsubscript{2}. Chain propagating and branching reactions lead to the formation of H, O and OH and these species progressively abstract hydrogen and partially oxidize CH\textsubscript{4} to CH\textsubscript{2}O. The CH\textsubscript{2}O is then converted largely to CO through the intermediate HCO. Subsequent oxidation of CO to CO\textsubscript{2} is by reaction with NO\textsubscript{2} of OH as mentioned above. If the fuel is CH\textsubscript{2}O instead of CH\textsubscript{4}, the latter stages of this chain become dominant. If the fuel is C\textsubscript{2}H\textsubscript{2} instead of CH\textsubscript{4}, then reactions of CH\textsubscript{2} become more important and CH\textsubscript{2}O becomes less important.

Rate constants for the reactions involved in the hydrocarbon chemistry derived from previous studies are generally successful in the description of the transformation of CH\textsubscript{4} to CO.

The third major aspect of fuel/NO\textsubscript{2} flame chemistry is the interaction of H, C and N containing species. Reaction of CH\textsubscript{4} species with NO to form HCN and subsequent reaction of HCN, CN or NCO with NO lead to the conversion of NO to N\textsubscript{2}. This process is obviously of most importance when NO\textsubscript{2} or NO is the oxidizer rather than when N\textsubscript{2}O is oxidizer, since in the latter case N\textsubscript{2} is formed directly. This scheme also shows the essential features of the oxidation process in the case where HCN is formed as a fuel during the decomposition of the energetic solid.

A.3 REFERENCES


3. Volponi, J.V., and M.C. Branch, "Flame Structure of C\textsubscript{2}H\textsubscript{2}-O\textsubscript{2}-Ar and C\textsubscript{2}H\textsubscript{2}-NO\textsubscript{2}-Ar Laminar Premixed Flames," 24th Symposium (International) on Combustion, in press.


A.4 RECENT PAPERS AND PRESENTATIONS FROM THIS AND PREVIOUS AFOSR SUPPORT


7. J.V. Volponi and M.C. Branch, "Flame Structure of C_2H_2-O_2-Ar and C_2H_2-NO_2-Ar Laminar Premixed Flames," *24th Symposium (International) on Combustion*, The Combustion Institute, Pittsburgh, in press.


A.5 PERSONNEL

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2. Hai-Ming Tsai, Ph.D. Candidate, University of Colorado, Boulder.

3. Joseph Cor, Ph.D. Candidate, University of Colorado, Boulder.

4. Andy Wilson, Undergraduate Research Affiliate, University of Colorado, Boulder.

5. Matthew Fuller, Undergraduate Research Affiliate, University of Colorado, Boulder.
TABLE 1. Rate constants used in the present 272 Step Reaction Mechanism which differ from those of Volponi and Branch (Ref. 3). Units are mole, cm³, sec, K, and Cal/mole.

<table>
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<th>Reaction</th>
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<td>N₂O+OH=N₂+HO₂</td>
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<td>10000</td>
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<td>28200</td>
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<tr>
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<td>0</td>
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<td>2.00e12</td>
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TABLE 2. Comparison of measured and calculated flame speed of fuel/NOₓ mixtures.

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<th>MEASURED</th>
<th>CALCULATED</th>
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<td>M/B</td>
<td>This Work</td>
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<td>227</td>
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<td>CH₁/N₂O</td>
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<tr>
<td>C₂H₂/N₂O</td>
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<td>128</td>
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<tr>
<td>C₂H₆/N₂O</td>
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<td>94</td>
</tr>
<tr>
<td>C₃H₆/N₂O</td>
<td>62.5</td>
<td>61</td>
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(a) All flame speeds given in cm/sec at 0.10 atm and for a stoichiometric mixture.
(b) Parker & Wolfhard (Ref. 12).
(c) Calculation using mechanism of Miller & Bowman (Ref. 13).
(d) Calculation using the present 272 step mechanism.

Figure 1. Comparison of CH₁/N₂O flame structure data of Habeebullab et al. (Ref. 5) to calculations of flame structure using 272 step reaction mechanism.
Figure 2. Comparison of H$_2$/NO$_x$ flame structure data of Volponi and Branch (Ref. 9) to calculations of flame structure using 272 step reaction mechanism.

Figure 3. Comparison of C$_2$H$_6$/NO$_x$ flame structure data of Volponi and Branch (Ref. 3) to calculations of flame structure using 272 step reaction mechanism.
B. FUNDAMENTAL STUDIES OF ROCKET COMBUSTION INSTABILITY

B.1 RESEARCH OBJECTIVES

Rocket engine manufacturers rely heavily on empirical and actual test data during the design stage of rocket engines in order to prevent the occurrence of instabilities during flight operation. This is clearly an expensive approach. Thus, the long term objective of our research is to develop predictive models that will help rocket designers build high performance, stable liquid rocket engines.

A complex interaction of several processes such as atomization of the primary fuel jet, secondary atomization of the resulting drops, vaporization, mixing and combustion are responsible for liquid rocket engine instability. Since atomization is the first mechanism in the chain of events leading to eventual combustion and possibly combustion instability, our near term objective is to understand the influence of atomization on the stability properties of rocket engines. The focus of our work is to elucidate and model the response of atomization under controlled modulated conditions.

B.2 BACKGROUND

Combustion instability that is characterized by large amplitude combustion chamber pressure oscillations is a commonly observed feature in many liquid propelled rocket engines. Severe engine damage could result due to mechanical failure and/or enhanced heat transfer to the chamber. Liquid fuel injected into a rocket engine combustion chamber undergoes a sequence of processes prior to exiting the chamber in the form of products of combustion. A typical scenario leading to combustion instability is as follows: an initial chamber pressure disturbance induces fluctuations in atomization rate. This results in variation of propellant droplet size distribution, which affects vaporization, mixing and reaction rates, inducing fluctuations in the engine burning rate. This process can feed energy back into the pressure field, thus closing the loop. If proper phase relationship is achieved during these processes, amplification of initial disturbances can occur, leading to combustion instability. Integration of all the physical processes involved into a comprehensive model of engine stability is not probable, unless each process is well understood.

Injectors in liquid rocket engines are exposed to relatively high amplitude pressure and velocity field oscillations (acoustic waves). These oscillations have a pronounced effect on the atomization of liquid propellant. Sufficient evidence from past work and our present research reveals that the structure and characteristics of atomization under unsteady conditions (as would occur during a combustion instability) deviate considerably from atomization behavior under steady conditions. Thus, the dynamic behavior of liquid propellant atomization is of major importance, especially when atomization produces very small droplets.

Miesse (1955) showed that significant acoustic interactions can be achieved between a low speed jet and local, high amplitude sound fields. Reba and Brosilow (1960) conducted the first experiment on liquid jet behavior perturbed by a longitudinal acoustic wave created by a siren (their work is also reported by Torda and Schmidt, 1964). Buffum and Williams (1967) investigated the behavior of a liquid jet subject to a transverse standing acoustic field. They studied the periodic oscillations of the liquid jet position in space and argued that this phenomenon might explain combustion instability in liquid rocket engines with impinging jet type injectors. Jet oscillatory deviations cause fluctuations in impingement point and geometrical injection parameters, which can induce variations in the atomization rate and pattern. Their
results showed a strong correlation between the acoustic field and the jet behavior for forcing amplitudes of 160 dB and frequencies up to 500 Hz.

Heidmann (1965) studied acoustic field and jet interaction in a two-dimensional circular combustor. He found a strong correlation between the acoustic field and the oscillatory jet breakup behavior. Ingebo (1966) studied the behavior of liquid jet atomization perturbed by longitudinal acoustic waves. A siren periodically obstructed the chamber exit and created longitudinal acoustic perturbations. Streak photography showed reversal of the gas flow during instability. High speed photography provided the first measurement of drop size distribution under unsteady and hot environment. He showed that under resonant conditions, atomization and vaporization rates are significantly enhanced. The jet breakup length was one sixth compared with the unforced case. Much finer droplet size was obtained (62 microns in resonant forcing conditions and 217 microns for unforced cases). The length for complete vaporization decreased from ~50 cm to 5 cm. More recently, Lecourt and Foucaud (1987) studied the stability of several injectors using the forcing method developed by Ingebo. Due to extreme operating conditions (1000 psia), only pressure-time traces were recorded.

Heidmann and Groeneweg (1969) studied the transfer function between pressure oscillations and atomization rate. They argued that if the in-phase transfer function is positive, unstable combustion might occur. The justification for this approach comes from the Rayleigh criterion. Different types of transverse acoustic oscillations (travelling, standing, radial transverse mode and vortex mode, characterized by a steady tangential velocity component) were investigated experimentally and analytically. They found that variations in the peak values of the response factors are caused by the velocity sensitivity of the atomization process, and a velocity perturbation, in-phase with pressure, elevated the peak value and is destabilizing to a combustion system. However, an out-of-phase velocity perturbation reduced the peak value of the response factor and is stabilizing. They predicted that potentially the maximum average response for the traveling mode is larger than for either the standing or radial modes. Both their analytical and experimental results indicated that a steady vortex velocity had an amplifying effect and increased the peak value of the response factor by an order of magnitude.

Overall, very little is known about the dynamics of atomization under modulated conditions. Current interest in combustion instabilities of liquid rocket engines and the advent of modern diagnostics has initiated our study of atomization behavior under oscillatory environment. In our work external forcing is used to simulate combustion induced pressure and velocity fluctuations. The advantage is a complete control over frequency, and amplitude, so that linear and nonlinear forcing regimes can be investigated. Our study focuses on the phase and amplitude relationship between forcing and atomization response. Although these quantities have been shown to be important in theoretical studies on the possible coupling between atomization and acoustic waves leading to combustion instability, they have never been measured. This knowledge can lead to the use of linear stability theory to clarify the role of atomization in high frequency combustion instability.

B.3 DESCRIPTION OF EXPERIMENTAL FACILITY

As one of the objectives of this research program is to provide useful atomization information for the design of actual injectors, it is important to give special attention to similarity parameters. Moreover, it is extremely difficult to model a real rocket engine because of the severe conditions encountered in an actual combustion chamber. High pressure and intense heat transfer rates precludes accurate measurement of quantities such as pressure, temperature and droplet size distribution. Propellants used in the rocket
industry are usually toxic and hypergolic. The approach adopted is to identify dimensionless numbers, which govern atomization and try to maintain them, or at least operate in the proper range. Four dimensionless numbers, liquid jet Reynolds number

$$Re_l = \frac{\rho_l V_l d_p}{\mu_l}$$

Weber number based on gas phase properties

$$We_g = \frac{\rho_g (V_g - V_l) d_p^2}{\sigma_l}$$

\(\mu_l/\mu_g\) and \(\rho_l/\rho_g\) are used to characterize atomization. Here, the symbols \(\mu\) and \(\rho\) represent dynamic viscosity and density. The subscripts \(l\) and \(g\) indicate liquid and gas. However, the definition of the quantities which are chosen to calculate the Reynolds and Weber numbers varies in the literature.

The turbulence level in the liquid phase, as the jet emerges from the nozzle is important because it characterizes the amplitude of the disturbances on the jet surface, which are likely to be amplified by dynamic interactions with the surrounding gas. This justifies our definition of the Reynolds number. The Weber number based on gas phase properties takes into account aerodynamic forces, the main mechanism for jet breakup and atomization at high velocities. Although no single mechanism can explain the entire process of atomization, aerodynamic interactions between the liquid jet and the surrounding gas provide the main destabilizing agent (Reitz and Bracco, 1986).

A survey of studies on coaxial injectors has been completed. The operating range of relevant results is reported in a \(Re_l - We_g\) diagram in Figure 1. The two horizontal lines correspond to theoretical results from Ranz (1958) and Miesse (1955), who predicted that atomization regime lies beyond a critical \(We_g\). Our approach is to operate in the atomization regime and come as close as possible to the configuration under simulation by a proper choice of simulants.

**B.3.1 Simulation Selection**

Since real atomizer injection velocities are very high, they cannot be increased to maintain similarity. A survey of different propellant combinations revealed that the best which can be done is to maintain the value of liquid and gas velocity. In order to maintain geometrical similarity, we have retained the inner post diameter (Figure 2). Therefore, the dimensionless numbers indicate that the simulant selection is governed by the following two equations.

$$\left(\frac{\rho_l}{\mu_l}\right)_{\text{actual}} = \left(\frac{\rho_l}{\mu_l}\right)_{\text{model}}$$

$$\left(\frac{\rho_g}{\sigma_l}\right)_{\text{actual}} = \left(\frac{\rho_g}{\sigma_l}\right)_{\text{model}}$$
Tables 1 and 2 show the values of $p_i/p_t$ and $p_f/o_i$ for different combinations for both hot and cold run simulations of $L_O_2/GH_2$ rocket propellants. For cold flow testing, the combination Freon 113, R13B seems adequate, if atomization data on $L_O_2/GH_2$ is to be obtained. For hot flow tests, one sees that for all simulants (hydrogen peroxide, ethanol) the value of $p_i/p_t$ falls short by a factor of at least 2.

In our case, we chose to use a combination of water and air mixed with refrigerant for cold flow runs. For experiments with combustion, the ethanol/air mixed with oxygen combination has been selected.

B.3.2 Combustor Design

The size of the combustor chamber has to be chosen carefully to ensure almost complete combustion before the nozzle exit. In particular, considerations using parameters such as $L^*$ or $t_{rd}$ helped us in our task. The quantity $L^*$ is a ratio of chamber volume to throat area. Values of $L^*$ must be chosen in accordance with the fuel-oxidizer choice. The fluid residence time in the chamber is denoted by $t_{rd}$. Also, the selected excitation system places a bound on the size of the chamber, because large amplitude modulations are possible only if the chamber volume is small enough.

The combustion chamber design is shown in Figure 3. The uni-element coaxial combustor has transverse dimensions of 5 cm x 5 cm. The chamber is designed to withstand a pressure of 10 atm absolute. This limitation comes from the quartz windows (1.25 cm thick). The chamber length can be varied from 28 to 43 cm by moving the exit nozzle location. The adjustable length allows for the study of various combustion efficiencies and longitudinal acoustic modes. The length of the chamber dictates the residence time of the reactants and hence the amount consumed, or in essence the combustion efficiency. Two quartz windows, 25 x 5 cm, on opposite sides of the chamber provide full optical access from the injection plane to 25 cm downstream. The operating pressure of the chamber for given fuel and oxidizer mass flow rates can be varied by using exit nozzles of different throat area. The chamber is mounted vertically to eliminate liquid accumulation during cold flow tests. The injector head is made of a 5 x 5 x 7 cm stainless steel block, as shown in Figure 2. The injector head can easily be modified to accommodate other types of atomizers such as an impinging jet. The center post of the injector is machined to allow for smooth reduction of liquid fluid passage area. The mounting of the center post can be adjusted to achieve the desired recess relative to the injection plane. Center posts of different design and diameter can be mounted into the injector block. The injector used in this study has a center post with inside diameter of 1 mm with a 4 mm width of the surrounding annulus.

B.3.3 Excitation Method

Acoustic compression drivers are used to excite transverse modes in the combustion chamber. The experimental configuration is shown in Figure 4. This kind of perturbation is chosen because, in many high frequency combustion instability cases, the coupling between combustion and chamber transverse acoustic modes have been recorded. The use of acoustic excitation devices in studies in the field of active control of combustion instability has proved to be successful. In particular, very high pressure excitation levels can be obtained.

Most acoustic drivers operate adequately in the frequency range from 1000 Hz to 6000 Hz. Moreover, many high frequency combustion instabilities are encountered in this range. Hence, the excitation frequency domain for transverse excitation is chosen as above. This sets the chamber transverse dimension to under 5 cm. It is also desirable to excite the chamber at a frequency different from any of its characteristic frequencies, in order to avoid resonance. Thus, we can exercise complete control on the
amplitude of modulation. High forcing pressure amplitudes can be obtained by tuning the length of the ducts connecting the compression chambers to the combustion chamber.

B.4 DIAGNOSTIC TECHNIQUES

Diagnostic techniques aimed at studying this configuration are under development. In particular, the time variation of quantities such as atomization rate, jet breakup length, length for complete atomization, under forcing and non-forcing conditions are of interest. Large and small scale fluid mechanical fluctuations and their phase with respect to external excitation are also important in better understanding the dynamic behavior of atomization. The task is difficult because the frequency range of interest is above 1000 Hz. Currently, phase locked, back-lighted photography and Schlieren are used. Diagnostic systems like the ones in Figures 5 and 6 can give access to the aforementioned interesting quantities. The signal from the acoustic drivers is used to periodically trigger a light source. An optical arrangement focuses the light onto a region of the atomizing jet. The resulting image is captured by a continuous high speed camera. Phase averaging is then used to minimize cycle-to-cycle variations. The images are then processed to extract jet breakup length and characteristic time for complete atomization. By shifting the phase of the light trigger with respect to the excitation signal, phase sensitive images of the atomizing jet are obtained.

Currently, the response of the chamber pressure and localized heat release in the combusting region to an acoustic perturbation is under investigation. In particular, the amplitude and phase of the response relative to the perturbation is being analyzed. A piezoelectric transducer mounted on the chamber wall measures the chamber pressure response. This type of transducer is capable of measuring high frequency and low amplitude acoustic waves. The localized heat release is measured by collecting the light emitted by C2 radicals, which are only present in the high temperature combusting regions of the flow. The measurement is local in the sense that an optical system only gathers light from a very small region in the combustion region. The emitted light is collected and focused onto a photomultiplier tube for signal generation. In this manner, a relative measure of the localized heat release can be obtained for comparison between excited and unexcited conditions.

In addition, laser back scattering and extinction diagnostic systems are under development, as shown in Figure 7. These systems are capable of measuring the atomization response to an imposed perturbation. The idea is to focus a He-Ne laser beam into the initial atomizing region near the inner post tip, the scattered light is then collected by a series of PMT detectors and the extinction of the laser beam is measured by a photo diode. The intensity of the laser beam received by the photo diode is directly proportional to the amount of fluid which has passed through it. Spectral analysis of the signals can then give information concerning the modification of frequency content of the atomization process due to acoustic perturbations.

B.5 RESULTS

B.5.1 Cold Flow Results

A series of pressure measurements inside the chamber have been conducted in order to obtain the fluctuating pressure and velocity field generated by the excitation system. The measurements indicate that transverse acoustic field strengths of up to 156 dB, and acoustic velocity fluctuations of up to 4.2 m/s at the injector location can be produced.
A numerical code using linear acoustic equations has been developed to model the pressure field in the test section. The equations are solved using a second order accurate finite difference scheme. Input includes the acoustic driver boundary conditions and a temperature map of the system to account for the speed of sound distribution. Output includes the acoustic modes for the system, and resulting pressure and velocity fields. The code will be enhanced to include flow within the chamber to better model real conditions.

A set of experiments has been carried out to determine the effect of acoustic forcing on the global behavior of the fuel spray. These measurements are difficult due to the accumulation of fluid on the windows of the chamber. The results indicate that under out-of-phase forcing conditions, there is an increase in the volume of fuel droplets drawn into the recirculating region at the top of the chamber. Visually, the level of fluid accumulated on the window rises with excitation. Further experiments are planned to elucidate the physical phenomenon and obtain precise quantitative results.

B.5.2 Hot Flow Results

The combustion/excitation system has been improved and made ready for experimentation. Smooth reliable combustion, as well as integration with the excitation system has been obtained. The operating limits involving reactant parameters and chamber geometrical settings have been explored. In addition, the diagnostic Schlieren imaging technique has been refined to generate high quality(contrast) imaging of the flow during hot runs, as shown in Figure 8.

Recent experimentation has been conducted to determine the effect of equivalence ratio and excitation amplitude upon the time variation of the chamber pressure between forced and unforced conditions. The gas phase and liquid phase velocities were held constant at 92 m/s and 15 m/s, respectively. The equivalence ratio was varied by the addition of pure oxygen to the gas phase resulting in a range from 3.8 to 7.4. These procedures were carried out in order to keep the atomization characteristics of the system constant. The enrichment of the gas phase with oxygen increased the gas phase velocity very little. Moreover, the aerodynamic properties of oxygen are essentially similar to air. Thus, atomization parameters including Weber number and density ratio were held relatively constant. However, the static chamber pressure varied with equivalence ratio. This was caused by the use of the same size chamber exit nozzle for all the experiments. A variety of exit nozzles are under construction in order to decouple the chamber pressure from equivalence ratio. As an example, for an experiment with an equivalence ratio near one, which is a high heat release and hence high chamber pressure case, a larger exit nozzle will be used to reduce the chamber pressure so that it will remain constant.

Unexcited chamber pressure recordings for equivalence ratios of 7.4 and 3.8 are shown in Figures 9 and 13, respectively. The power spectra of these signals, shown in Figures 10 and 14, indicate the presence of three frequency ranges. The first range is centered at 200 Hz and the other two ranges are centered at 1250 Hz and 2500 Hz. There is a slight increasing shift of these ranges as the equivalence ratio is reduced(approaches one). This is probably caused by the fact that as stoichiometric conditions are approached, the chamber experiences higher temperatures and thus higher sound speeds, and hence increased oscillation frequency.

The acoustic compression drivers were tuned to excite the chamber at 2950 Hz. This frequency was chosen because it corresponds to a large amplitude pressure fluctuation in the chamber (determined experimentally), and it is in the high frequency (screeching) range. Chamber pressure records under forced conditions, for equivalence ratios of 7.4 and 3.8 are shown in Figures 11 and 15, respectively. The power
spectra of these signals, shown in Figures 12 and 16, indicate a peak at the excitation frequency. This is expected because the combustion chamber, excitation system, and pressure transducer are in a closed system. The interesting result is that the observed peak at the excitation frequency increases as the equivalence ratio is decreased. The peak power at the excitation frequency versus equivalence ratio is displayed in Figure 17. This result is biased by the fact that the static chamber pressure also increased with decreasing equivalence ratio. As stated earlier, the elimination of this bias is underway.

The effect of excitation amplitude for a fixed equivalence ratio of 3.8, holding all other parameters constant, was examined. As shown in Figure 18, the peak power at the excitation frequency increases as the excitation amplitude is increased. Again, this was expected because the pressure transducer and excitation system are in a closed system.

Currently, work is under way to determine the effect of gas phase velocity, equivalence ratio, excitation frequency, chamber length, and injector post recess upon the time variation of chamber pressure and localized heat release (C2 emission) between forced and unforced conditions. Schlierien images are being acquired to analyze the fluid mechanical response to excitation, i.e (jet breakup length, length for complete atomization, vortical structure,...etc). The analysis includes an examination of the frequency content (power spectra) of the measured signals to determine if the acoustic forcing frequency can drive the physical process under investigation. The phase and amplitude response of the signals relative to the perturbation will also be obtained. This information is a direct measure of the (n,τ) parameters applied to (n,τ) correlations commonly used in industry to predict instability behavior. In addition, auto- and cross-correlations of these signals will be acquired to determine the periodicity and similarity of the measured signals and perturbation.

B.6 FUTURE WORK

Laser backscattering and extinction experiments are planned to obtain real time data on spray fluctuations in order to measure the frequency response of atomization to acoustic perturbation. Furthermore, a Phase Doppler system is under development in order to measure the fluctuation in drop size and drop velocity in the atomizing region. Thus, this system will provide precise quantitative measurements of real physical atomization parameters and give a more clear understanding of atomization response.

In addition, an image processing system has been acquired. This system will generate sharp, detailed images of the spray and flow structure. Phase averaging can then provide statistically reliable quantitative information concerning fluid mechanical response.

B.7 REFERENCES


3. Heidmann, M.F., "Oxygen-Jet Behavior During Combustion Instability in a Two-Dimensional


B.8 PAPERS AND PRESENTATIONS FROM THIS AFOSR SUPPORT


4. C. Huynh, A. Ghafourian, S. Mahalingam, and J.W Daily, "Dynamic Behavior of Atomizing Jets in


**B.9 SEMINARS AND PRESENTATIONS NOT INCLUDED ABOVE**


**B.10 PERSONNEL**

1. Akbar Ghafourian, Ph.D. candidate, University of Colorado, Boulder.

2. Can Huynh, visiting scholar, Ecole Centrale Paris, France.

3. Rom McGuffin, graduate student, University of Colorado, Boulder.

4. Franz-Josef Kahlen, visiting scholar, Technical University of Aachen, Germany.
Table 1: $\rho_l/\mu_l$ of Liquids

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<th>LOX (crit)</th>
<th>LOX (sat)</th>
<th>$H_2O_2$</th>
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<th>Jet A</th>
<th>Shell wax</th>
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Table 2: $\rho_g/\sigma_l$ of Different Gas Liquid Combinations

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Figure 1: $Re_l - We_p$ Diagram of Some Previous Experimental Studies on Coaxial Injectors
Figure 1: Picture of Combustion Chamber

Figure 2: Injector head, coaxial injector
Figure 4: Combustor Sketch for Transverse Mode Forcing
Figure 5: Diagram of Back Lighted Photography

Figure 6: Diagram of Schlieren Imaging System
Figure 7: Schematic of Laser Backscattering and Extinction System

Figure 8: Micrograph of a Hot Test
Figure 9: Unexcited chamber pressure, gas phase velocity = 93 m/s, liquid velocity = 15 m/s, equivalence ratio = 7.4

Figure 10: Power Spectrum, unexcited, gas phase velocity = 93 m/s, liquid velocity = 15 m/s, equivalence ratio = 7.4
Figure 11: Excited chamber pressure, gas phase velocity = 93 m/s, liquid velocity = 15 m/s, equivalence ratio = 7.4

Figure 12: Power Spectrum, excited, gas phase velocity = 93 m/s, liquid velocity = 15 m/s, equivalence ratio = 7.4
Figure 13: Unexcited chamber pressure, gas phase velocity = 93 m/s, liquid velocity = 15 m/s, equivalence ratio = 3.8

Figure 14: Power Spectrum, unexcited, gas phase velocity = 93 m/s, liquid velocity = 15 m/s, equivalence ratio = 3.8
Figure 15: Excited chamber pressure, gas phase velocity = 93 m/s, liquid velocity = 15 m/s, equivalence ratio = 3.8

Figure 16: Power Spectrum, excited, gas phase velocity = 93 m/s, liquid velocity = 15 m/s, equivalence ratio = 3.8
Figure 17: Peak Power Spectrum Amplitude at Excitation Frequency vs. Equivalence Ratio

Figure 18: Peak Power Spectrum Amplitude at Excitation Frequency vs. Excitation Amplitude