Detonability Study of Liquid Hydrazine

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ABSTRACT

This paper presents the results of a study conducted at NASA White Sands Test Facility (WSTF) on the shock detonability of liquid hydrazine. Liquid hydrazine is the propellant used in the propulsion modules for Space Station Freedom. The focus of this study was to investigate the shock sensitivity of liquid hydrazine subjected to a hypervelocity projectile impact. This study provides the minimum power density needed for the initiation of homogenous liquid hydrazine, and thus provides the information needed to conduct a meaningful impact test on liquid hydrazine's shock detonability.

BACKGROUND

WSTF, in the past, has performed tests that attempted to shock-initiate liquid hydrazine and found no evidence of hydrazine reaction (see references 1 and 2). The most recent test on the shock detonability of liquid hydrazine was conducted by Science Applications International Corporation (SAIC) for Eglin Air Force Base (see reference 3). The SAIC test suggested that there was some evidence of hydrazine reaction, although the details of their results were poorly substantiated. The following is a short synopsis of the previous tests:

1) Condensed Phase Detonation Studies, WSTF # 90-24354, dated September 28, 1990. 887 g of C-4 was detonated on top of a stainless steel tube (4 in. x 10 in. long) filled with liquid hydrazine. The liquid hydrazine did not detonate or sustain a reaction.

2) Demonstration of Hazardous Hypervelocity Test Capability, TR-692-001, dated September 24, 1991. A 1/8-in. aluminum projectile was shot with a velocity of 6.1 km/sec at a 300-ml stainless steel vessel filled with liquid hydrazine. The liquid hydrazine did not detonate or sustain a reaction.

3) Fuel Tank Explosion Lethality, SAIC 91-5425-SH, dated April 1991. A 100-g cylindrical projectile was shot with a velocity of 5.0 km/sec at an aluminum, 100-mm-diameter spherical
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vessel filled with liquid hydrazine. The test results indicated that some reaction of liquid hydrazine occurred, although not enough evidence was gathered to emphatically verify that a detonation occurred.

Given the differences in these results, this study was conducted in an attempt to investigate in detail the shock stimuli that would be necessary to achieve an appreciable hydrazine reaction. The methodology employed was one used by C. L. Mader in his success with modeling homogeneous energetic materials using Arrhenius kinetic parameters determined from laboratory thermal stability experiments. Mader was very successful at numerically reproducing the shock initiation of nitromethane observed in experiment.

PURPOSE

An energetic material is one which decomposes exothermically, i.e., with the release of heat. By this definition liquid hydrazine, a monopropellant, is an energetic material and therefore should detonate given the proper shock stimuli. This study investigates fundamental information on the appropriate stimulus needed to achieve a detonation. Specifically, this study provides the minimum power density needed for the detonation of homogenous liquid hydrazine. The term power density implies the pressure and time duration sustained by a projectile in order to provide sufficient heat for attaining the critical temperature for a detonable condition. The result of this study is a proposed experiment aimed at reproducing the most realistic situation in which a hypervelocity projectile impact might initiate homogenous liquid hydrazine.

APPROACH

The initial effort of this study was to investigate, in detail, previous tests conducted by others. Hydrodynamic models of each experiment were calculated to determine the pressure and duration delivered to the liquid hydrazine. Then, as chemical kinetic parameters were determined, reactive hydrodynamic models were constructed to determine if the results of the experiments could be duplicated. If successful, a numerical model would be formulated to design an experiment that would replicate actual conditions that might exist for a titanium tank filled with liquid hydrazine on board Space Station Freedom.

The previous tests were modeled using the hydrocodes SIN, TDL, and ZEUS. The employment of these codes is shown in the flow diagram below. The following list expands on the approach as summarized in the flow diagram.

1. Determine the unreacted equation of state (EOS) of liquid hydrazine.

For accurately determining the variables of other shock states, the EOS of any substance is usually required, being the equation that bridges the gap with mass, momentum, and energy. This is an experimental plane in which the shock velocity and particle velocity (or free surface velocity) are measured. The proposed method for determining the EOS was one used by McQueen and Rice of Los Alamos National Laboratory. The test method uses an impedance
matching technique in which a projectile with a known EOS impacts a target with an unknown EOS at a fixed impact velocity. By measuring the shock velocity and knowing the initial density of the known material, one can determine the Rayleigh line, that will intersect the projectile shock hugoniot at one point. By repeating the procedure for different impact velocities and determining the different slopes of the Rayleigh line, one can construct the shock hugoniot of the unknown material. A minimum of six shots at different velocities are needed to determine the EOS.

One alternative to the above method was to make some assumptions about homogenous hydrazine and input these parameters into a code called SEQS (Solid Equation of State). This code will generate single shock hugoniots in the temperature vs specific volume plane, pressure vs particle velocity plane, and shock velocity vs particle velocity plane. The results of these computations are shown in Appendix A.

2. Determine the reactive hugoniot equation of state for the detonation products.

A description of the expansion isentrope for the detonation products is needed to determine the C-J pressure and velocity. The code used to determine this information was the BKW (BeckerKistiakowski-Wilson) code. This code performs a chemical equilibrium balance and generates an expansion isentrope, a curve for the reaction products in the pressure vs specific volume plane. The results of this computation are shown in Appendix B.

3. Perform thermal stability experiments.

Thermal stability experiments are needed to determine the chemical reaction rate, products, critical temperatures, and activation energy for liquid hydrazine. These quantities are constants for the Arrhenius rate law for burn, which is needed to perform hydrodynamic calculations in the SIN and TDL codes. The information for this section was taken from tests performed by the New Mexico Institute of Mining and Technology (NMT) in Socorro, New Mexico. Their results are shown in Appendix C.

4. Determine the minimum power density for steady-state detonation.

One-dimensional hydrocode calculations were performed using SIN. This code is used to model explosive flow using one-dimensional Lagrangian, reactive hydrodynamics. The SIN code was the primary workhorse for modeling the previously conducted tests and in determining the detonability of liquid hydrazine for the proposed experiment. After determining the conditions (i.e. the power density requirement) for attaining a steady-state detonation, a two-dimensional Lagrangian, reactive hydrocode calculation was performed to determine the effects of geometry on the release wave attenuation. The code used to perform these calculations was TDL. The two-dimensional calculations were augmented by calculations using the Zeus code. This code provided a check on the shock pressures generated in the liquid hydrazine. The results of these calculations are presented in the results section.
Flow Diagram for Explosion Response of Liquid Hydrazine
RESULTS

The tests performed by other investigators were numerically modeled with the following assumptions. A shock travels into the homogenous hydrazine, compressing and heating the propellant. The shock heating results in chemical decomposition that accelerates exponentially. The reaction begins at the rear boundary, because it has been hot the longest, and a detonation wave propagates at the C-J state of the shock propellant. This type of bulk heating or thermal initiation analysis has been successfully performed by C. L. Mader for homogenous energetic materials using the Frank-Kamenetskii equation with the Arrhenius chemical kinetics. Using this type of analysis, each test was modeled to investigate the shock stimuli provided in each situation and to formulate the minimum power density that would be required for steady-state detonation.

Condensed Phase Detonation Studies

This test was modeled using the SIN code with the chemical kinetic parameters provided by NMT. The result of this calculation indicated that the shock wave generated by the C-4 explosive was not sufficient to generate a hydrazine reaction. The shock pressure into the hydrazine was 150 kbar with a duration of approximately 2 microseconds. Figure 1 in Appendix D shows the shock pressure generated into the liquid hydrazine and Figure 2 shows that no reaction occurred.

Demonstration of Hazardous Hypervelocity Test Capability

This test was modeled using the TDL code with the same chemical kinetics as above and the Zeus code with no chemical kinetics. Zeus calculations were performed to provide a better representation of the shock wave generated by the sphere impacting the cylindrical vessel. This model shows that the release waves attenuate the shock wave into the liquid hydrazine very quickly, thus rendering an ineffective shock wave in the hydrazine. The pressure and duration, as calculated by Zeus, were 83 kbar and approximately .8 microseconds, which is in agreement with the TDL calculations. Figure 3 shows the shock pressure generated into the liquid hydrazine. The results of the TDL code calculation revealed that the pressure generated by the aluminum projectile was not sufficient to generate a hydrazine reaction. Figure 4 shows the initial projectile impact and Figure 5 shows that no hydrazine reaction occurred.

Fuel Tank Explosion Lethality

This test was modeled using the SIN code with the same chemical kinetics as above and the Zeus code with no chemical kinetics. The results of the SIN code calculation revealed that the shock wave generated by the aluminum projectile was sufficient to attain some decomposition of hydrazine. One reason a higher shock pressure was generated into the liquid hydrazine as compared to the other experiments was the impedance matching of materials adjacent to the liquid hydrazine. Figures 6 and 7 represent a pictorial view of the shock impedance matching of the SAIC test and the Condensed Phase Detonation Study. The pressure and duration as calculated by SIN was 285 kbar and approximately 5 microseconds. Figure 8 shows the
shock pressure generated into the liquid hydrazine and Figure 9 shows the amount of decomposition. The pressure and duration as calculated by Zeus was 267 kbar and approximately 1.5 microseconds, which is in agreement with the SIN calculations. Figure 10 shows the shock pressure generated into the liquid hydrazine. Another reason there was more hydrazine decomposition in this test as compared to the other tests was that the projectile geometry sustained the shock pressure for a longer duration before the release wave reduced its magnitude.

Proposed Experiment

The goal of this study was to use the information gathered from analyzing the previous tests to formulate a test situation in which the minimum power density requirement was met, replicating the actual hydrazine tank conditions existing on board Space Station Freedom. This experiment was created from a numerical model which simulated a steel slug impacting a titanium vessel filled with liquid hydrazine. The projectile was designed to be a cylinder to provide the sustained pressure necessary to generate enough shock heating. This experiment was also designed such that the projectile velocities could be achieved with WSTF's 1-in. light gas gun. The desired projectile velocity was between 7-7.5 km/sec. This experiment was modeled using SIN and TDL with the same chemical kinetics as above and the Zeus code with no chemical kinetics. The result of the SIN and TDL model revealed that the shock wave generated by the steel slug was sufficient to achieve a substantial amount of hydrazine reaction. The shock pressure into the hydrazine, as calculated by SIN, was 600 kbar with a duration of approximately 2.4 microseconds. Figure 11 shows the shock pressure generated into the liquid hydrazine and Figure 12 shows the amount of decomposition. Figure 13 shows the amount of hydrazine decomposition as calculated by TDL.

The pressure and duration, as calculated by Zeus, was 600 and approximately 1 microsecond, which is in agreement with the TDL calculations. Figure 14 shows the shock pressure generated into the liquid hydrazine.

Table 1 is presented as a comparison of the results for all tests and the proposed experiment. The relative ranking is based on the power density requirement for homogenous materials, which is

\[ P^2 \tau \]

This implies the pressure, \( P \) applied for a stock duration of \( \tau \).
CONCLUSIONS

Numerical reactive models of the WSTF tests and the SAIC test successfully reproduced their respective test results. The WSTF tests and the SAIC test emphasized that under particular shock loading conditions a minimum power density is required to achieve a hydrazine reaction. These models suggest that there is increased shock heating as a function of power density applied to the liquid hydrazine for each case. Analyzing previously conducted experiments provided a platform from which the proposed experiment has been suggested. This analysis has provided the most realistic power density required to achieve hydrazine reaction in a titanium tank. Future work should consider tests in which many parameters in reactive modeling are unknown for propellants. This should also include a program of determining the equations of state of propellants used by NASA and other agencies, and a program of enhancing a data base of kinetic parameters for various propellants specifically for inputs into reactive hydrodynamic models. Finally, the proposed experiment should be carried out to verify the results of the current modeling effort. This should not be looked at as a pass or fail experiment but only as a tool for verifying the suspected parameters which were calculated. This experiment needs to employ enough detailed instrumentation so that there would be sufficient information on the extent of the hydrazine reaction and to verify model parameters for future tests.

ACKNOWLEDGMENT

The authors are indebted to C. L. Mader of Mader Consulting Company, Honolulu, Hawaii, who spent many hours of his time in providing us with his knowledge in numerical modeling of explosives.

REFERENCES:


APPENDIX A
Solid Equation of State

Solid Equation of State Calculation for ** Hydrazine

Us = 1.500000E-01 + 1.500000E+00 Up from po to 1.050000E+00 megabars

The Initial Density is 1.0000000000E+00 g/cm$^3$

The Compressibility is 5.0000000000E+01

The Linear Coefficient of Expansion is 6.0000000000E-05

The Initial Temperature is 3.0000000000E+02

The Heat Capacity is 1.0000000000E+00

The Volume Increalaent is 2.0000000000E+00

The Temperature fit is between 1.000000E-04 and 5.000000E-01 megabars

\[ \ln(T) = 5.69480500000E+00 -3.62450400000E-01 \ln(V) -1.88700200000E+00 \ln(V)^2 -4.40056400000E+00 \ln(V)^3 + 1.74659500000E+00 \ln(V)^4 \]

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FIGURE 3

[Graph showing temperature vs. volume with a sharp decrease at a certain volume]
FIGURE 4
APPENDIX B
BKW Equation of State

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A FORTRAN BKW Calculation for the Explosive Hydrazine for Garcia- Large NH3 Covolume

The Number of Elements is 2

The Number of Gas Species is 4

The Number of Solid Species is 0

The BKW Equation of State Parameters are
- Alpha= 5.000000E-01
- Beta= 1.600000E-01
- Theta= 4.000000E+02
- Kappa= 1.0909780E+01

The Composition of the Explosive is
- 2.000000E+00 Moles of n
- 4.000000E+00 Moles of h

The Density of the Explosive is 1.011000E+00, g/cm³

The Molecular Weight is 3.2045280E+01 grams

The Heat of Formation at 0 deg K is 1.0500000E+04 Calories per Formula Weight

The Solid (Cowan) Equation of State parameters V₀, AS, BS, CS, DS, ES, ES_

The Input Detonation Product Elemental Composition Matrix

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</table>
A FORTRAN BKW Calculation for the Explosive Hydrazine for Garcia- Large NH3 Covolume

The Computed CJ Pressure is 1.1229070E-01 megabars

The Computed Detonation Velocity is 7.5930330E-01 cm/microsecond

The Computed CJ Temperature is 3.4224570E+02 Degrees Kelvin

The Computed CJ Volume 7.9857050E-01 cc/gm of Explosive

The Computed Gamma is 4.1908430E+00

The Volume of the Gas is 8.53009E+00 cc/mole 3.00002E+00

Moles of Gas

Solid Volume in cc/cm

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<th>the I C, Heat For, Covolume</th>
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The BKW Hugoniot for the Detonation Products of Hydrazine for Garcia- Large NH3 Covolume

Pressure = 5.0000000E-01 Volume = 5.6719090E-01 Temperature = 1.1385820E+03 Shock Velocity = 1.0767470E+00 Particle Velocity = 4.5930810000E-01

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Specie  No of Moles
n2     1.0000040E+00
h2     2.0000130E+00
nh3    1.0000000E-08
h      1.0000000E-08

Pressure = 2.0000000E-01  Volume = 7.0006750E-01  Temperature = 4.7937030E+02  Shock Velocity = 8.2276270E-01  Particle Velocity = 2.4043740000E-01

Specie  No of Moles
n2     1.0000040E+00
h2     2.0000130E+00
nh3    1.0000000E-08
h      1.0000000E-08

A BKW Isentrope thru BKW CJ Pressure for Hydrazine for Garcia- Large NH3 Covolume

\[
\ln(P) = -2.7180350E+00 -3.6537590E+00 \ln(V) -6.0775230E-01 \ln(V)^2 \\
8.7331600E-01 \ln(V)^3 -4.5526350E-01 \ln(V)^4
\]

\[
\ln(T) = 6.1689300E+00 -1.5809880E+00 \ln(V) -9.8901980E-01 \ln(V)^2 \\
8.3671260E-01 \ln(V)^3 -6.9144960E-01 \ln(V)^4
\]

\[
\ln(E) = -1.3238010E+00 4.8534600E-01 \ln(P) + 4.598300E-02 \ln(P)^2 \\
7.3650140E-03 \ln(P)^3 -1.1714560E-03 \ln(P)^4
\]

The constant added to energies was 1.0000000E-01

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FIGURE 7
APPENDIX C
Thermal Stability Experiments

DETERMINATION OF HYDRAZINE DECOMPOSITION PARAMETERS

Jimmie Oxley; James Smith; Hongtu Feng; Evan Rogers; Nancy Gilson

New Mexico Institute of Mining and Technology
October 23, 1993

Accurate activation energy and pre-exponential factor values are essential for the SIN and TDL code calculations on hydrazine detonability. In this work we have determined these parameters over as wide a temperature range as possible.

Experimental Techniques

Hydrazine was transferred from the storage cylinder provided by Lockheed into a 5 mL septum cap vial inside a nitrogen-flushed glove bag. Inside the inert-atmosphere bag, hydrazine liquid was allowed to slowly fill the vial which was then capped with a screw-top septum prior to removal from the glove bag.

Outside the glove bag, hydrazine was quantitatively transferred (5 uL) from the nitrogen-filled vial into melting-point capillary tubes (1.8-mm o.d. by 6-cm long) using a 10 uL syringe. To avoid build-up of negative pressure in the septum vial, the syringe was filled with 5 uL of nitrogen gas which was injected into the septum vial, just prior to the removal of 5 uL of hydrazine. The melting-point capillary, into which the hydrazine was transferred, was flushed with argon until the hydrazine was added. At that point the inert gas flush was switched to helium, and the melting-point capillary was flame sealed. [The reason for the change in the inert gas fill was that although argon is a denser gas and provides a more reliable inert blanket, its retention time (GC) under the experimental conditions overlapped with that of nitrogen. Helium is, of course, invisible to in gas chromatography (GC) since it is the carrier gas.]

The capillaries of hydrazine were heated isothermally in a molten metal (Wood's metal) bath. At specific times samples were removed, and the progress of the decomposition was monitored by quantifying the evolved nitrogen gas up to that time. A Varian 3600 gas chromatograph (GC) equipped with a Haysep DB column, a thermal conductivity detector (TCD), and helium carrier gas (20 mL/min.) was used. only two decomposition gases were observed: N2 with retention time about 7 minutes and NH, retention about 14 minutes. When quantification of decomposition of gases was desired, peak areas of each gas were compared against calibration curves for nitrogen and ammonia standards. When the extent of decomposition was being assessed, the peak area of the nitrogen evolved after a given time of isothermal heating was ratioed against the peak area of nitrogen evolved
Results and Discussion

Rate constants for hydrazine decomposition were found over a 130°C temperature range using isothermal methods. Rate constants determined and the goodness of the linear-regression fit are shown in Table I. (Individual first-order plots are provided in the Appendix.) Using the rate constants obtained at ten temperatures, an Arrhenius plot was constructed (Fig. 1) and activation energy and frequency factor were calculated over the entire temperature range (26.8 kcal/mol and $6.51 \times 10^6$ sec$^{-1}$) and over the lower four temperatures (35.0 kcal/mol and $3.61 \times 10^{10}$ sec$^{-1}$) where there appeared to be a slight change in slope. Comparison to previously reported data is shown in Table II.$^{1,2}$

Initial plans to quickly examine hydrazine thermal stability by scanning differential scanning calorimetry (DSC) were abandoned after several attempts produced irreproducible results. Decomposition kinetics obtained by scanning DSC analysis are considered inferior to those obtained by isothermal methods, but the DSC method is usually quicker. In the case of hydrazine, kinetics obtained by DSC have been published (Table II). However, after numerous attempts, it was deemed a waste of time and money to pursue this method. Because of the volatility of hydrazine, the hydrazine sample must be in a pressure tight container. The pressure-tight vessels we commonly use are flame sealed microcapillaries (about 1 cm long). For hydrazine, this meant the flame used to seal the capillaries came quite close to the sample and evidently produced varying amounts of decomposition in various samples.

The decomposition gases were examined at both the high (275°C) and low (190°C) end of the temperature range examined. The relative ratio of nitrogen to ammonia produced at these temperatures did not significantly vary (Fig. 2). Therefore, despite the slight kink in the Arrhenius plot between 250 and 240°C, we cannot claim that the mechanism changes over this temperature range.

$$3 \text{ N}_2\text{H}_4 \rightarrow \text{ N}_2 + 4 \text{ NH}_3$$

References


<table>
<thead>
<tr>
<th>Table I</th>
<th>Rate Constants of $N_2H_4$ Decomposition</th>
</tr>
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<tbody>
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<td>°C</td>
<td>$k$ sec$^{-1}$</td>
</tr>
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<td>6.55E-07</td>
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<tr>
<td>200</td>
<td>2.54E-06</td>
</tr>
<tr>
<td>220</td>
<td>1.26E-05</td>
</tr>
<tr>
<td>240</td>
<td>4.25E-05</td>
</tr>
<tr>
<td>250</td>
<td>4.49E-05</td>
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<td>260</td>
<td>7.06E-05</td>
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<td>270</td>
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<td>3.21E-04</td>
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<td>314</td>
<td>5.58E-04</td>
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Table I Rate Constants of $N_2H_4$ Decomposition

<table>
<thead>
<tr>
<th>Table II</th>
<th>Arrhenius Parameters of $N_2H_4$ Decomposition</th>
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<tr>
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<td>Isothermal (this work)</td>
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<td>Temp Range (°C)</td>
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Table II Arrhenius Parameters of $N_2H_4$ Decomposition
Figure 9 Hydrazine Arrhenius Plot
(Appendix C, Fig. 1)

Hydrazine Arrhenius Plot

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<tr>
<th>°C</th>
<th>k sec(^{-1})</th>
<th>(R^2)</th>
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<tr>
<td>184</td>
<td>6.55e-07</td>
<td>0.97</td>
</tr>
<tr>
<td>200</td>
<td>2.54e-06</td>
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<tr>
<td>220</td>
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<td>300</td>
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<tr>
<td>314</td>
<td>5.58e-04</td>
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Fig. 1
Figure 10
Gas evolved at 190°C and Gas evolved at 275°C

Gas evolved at 190°C

Gas evolved at 275°C

Fig. 2
Figure 11
Hydrazine at 184°C HPLC Data and Hydrazine at 200°C HPLC Data

Hydrazine at 184°C
HPLC Data

Hydrazine at 200°C
HPLC Data
Figure 12
Hydrazine at 220°C HPLC Data and Hydrazine at 240°C HPLC Data

Hydrazine at 220°C
HPLC Data

Hydrazine at 240°C
HPLC Data
Figure 13
Hydrazine at 250°C HPLC Data and Hydrazine at 260°C HPLC Data

Hydrazine at 250°C
HPLC Data

Hydrazine at 260°C
HPLC Data
Figure 14
Hydrazine at 290°C HPLC Data and Hydrazine at 300°C HPLC Data
Figure 15
Hydrazine at 314°C HPLC Data
Figure 16
APPENDIX D
(Figure 1 of Appendix D)
Figure 1 Condensed Phase Detonation Study SIN calculation of the shock pressure into liquid hydrazine

Figure 1
Condensed Phase Detonation Study
SIN calculation of the shock pressure into liquid hydrazine
Figure 17
(Figure 2 of Appendix D)
Figure 2  Condensed Phase Detonation Study No hydrazine reaction

Figure 2
Condensed Phase Detonation Study
No hydrazine reaction
Figure 18
(Figure 3 of Appendix D)
Figure 3 Demonstration Hazardous Hypervelocity Test Zeus calculation of the shock pressure into the liquid hydrazine
Figure 19
(Figure 4 of Appendix D)

Figure 4  Demonstration of Hazardous Hypervelocity Test TDL calculation of projectile impacting the cylindrical vessel for the No hydrazine reaction.

Figure 4
Demonstration of Hazardous Hypervelocity Test TDL calculation of projectile impacting the cylindrical vessel for the No hydrazine reaction.
Figure 20
(Figure 5 of Appendix D)

Figure 5 Demonstration of Hazardous Hypervelocity Test TDL calculation of the shock pressure into the hydrazine. No hydrazine reaction.
Figure 21
(Figure 6 of Appendix D)
Figure 6 Shock matching curves for the SAIC Test
Figure 7
Shock matching curves for the Condensed Phase Detonation Study

Figure 7 Shock matching curves for the Condensed Phase Detonation Study
Figure 23
(Figure 8 of Appendix D)
Figure 8 SAIC Test
SIN calculation of the shock pressure into the liquid hydrazine
Figure 24
(Figure 9 of Appendix D)
Figure 9 SAIC Test
Hydrazine decomposition

Figure 9
SAIC Test
Hydrazine decomposition
Zeus calculation of the shock pressure into the liquid hydrazine
Figure 26
(Figure 11 of Appendix D)
Figure 11 Proposed Experiment
SIN calculation of the shock pressure generated into the liquid hydrazine
Figure 27
(Figure 12 of Appendix D)
Figure 12 Proposed Experiment
SIN calculation of the amount of hydrazine decomposition

Figure 12
Proposed Experiment
SIN calculation of the amount of hydrazine decomposition
Figure 28
(Figure 13 of Appendix D)
Figure 13 Proposed Experiment
TDL calculation of the amount of hydrazine decomposition

Figure 13
Proposed Experiment
TDL calculation of the amount of hydrazine decomposition
Zeus calculation of the shock pressure into the liquid hydrazine
### Table I

#### Reactive Analysis of Tests

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<th>DURATION (MICROSEC)</th>
<th>MASS FRACTION</th>
<th>PRESSURE (KBAR)</th>
<th>DURATION (MICROSEC)</th>
<th>POWER DENSITY (KB ~ 2-SEC)</th>
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Table III

(Table 1 of Appendix D)

Table 1 Reactive Analysis of Tests