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FINAL REPORT  
OIL CONTAMINATION IN OXYGEN SYSTEMS

Contract NObs-94416  
Project No. SP013-08-14, Task 3917

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## FOREWORD

This report covers the work performed under NAVSHIPS Contract No. NObs-94416 (Project No. SF013-08-14, Task 3917). The NAVSEC contract monitor was Mr. Herman Kraut.

The objective of this program was to collect the necessary experimental data on ignition and flame propagation in an oil contaminated oxygen system in order to more exactly define the required level of cleanliness for such systems. The results are to be incorporated into military specifications for oxygen systems to define the required cleanliness.

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## INTRODUCTION

Under auspices of the Department of the Navy, NAVSEC, a study was undertaken to determine the compression ignition and flame propagation limits of hydrocarbons in a high pressure oxygen system.

Considerable work has been done concerning compression ignition and flame propagation in high pressure air systems, but very little has been done with pressurized oxygen systems. The limited work performed with oxygen did not employ conditions corresponding to the specific systems now in use, namely 3000 psi oxygen with 1/2" monel piping. The program reported herein was designed to study that specific area.

The study was primarily concerned with the determination of the acceptable levels of oil contamination (2190) on the internal surface of a 1/2" monel pipe below which compression ignition and flame propagation will not occur when pressurized with 3000 psi oxygen. It was also desired to determine that level of oil contamination where, upon ignition, a pressure rise will not exceed 25%.

The first phase of the program consisted of a literature survey to assemble and review all available information from previous studies in the area of compression ignition and flame propagation in high pressure air and oxygen systems. A summary of these earlier studies is presented in Section I.

Following the literature survey a test program was designed and initiated to accomplish the objectives of the contract. A description of the test apparatus used is presented in Section II, the testing procedure is described in Section III, and the results of the test program are presented in Section IV.

Throughout this report, the term "oil concentration" is used. This refers to the amount of oil coating, expressed as  $\text{mg/ft}^2$ , on the inside surface of the test pipe.



## SUMMARY

A test program was conducted to determine the ignition limits of oil contamination in 1/2" monel pipe when pressurized with 3000 psi oxygen. The oils investigated were 2190 lubricating oil and Habcoc #318 cutting oil. Thin films of these oils were applied to the internal surface of a 4 ft. section of 1/2" monel pipe. The pipe was then rapidly pressurized with 3000 psi oxygen. In addition, tests were made using oil droplets and varying the initial pipe wall temperature.

The test results showed little danger of compression ignitions at oil concentrations below 160 mg./ft<sup>2</sup> of 2190, or below 110 mg./ft<sup>2</sup> for Habcoc. Concentrations of both oils above 260 mg./ft<sup>2</sup> will ignite when rapidly pressurized with 3000 psi oxygen. These concentrations must be compared to the present oxygen clean standards of 0.013 mg./ft<sup>2</sup> maximum allowable hydrocarbon concentration.

The results also showed that ignition will not occur with oil droplets or puddles smaller than 100  $\mu$  in size and that a pressure rise greater than 25% will not occur below oil concentrations of 500 mg./ft<sup>2</sup>.

As a result of this test program, it is recommended that the oxygen clean standards be relaxed to 1.5 mg./ft<sup>2</sup>. The level of oil concentration is a factor of 100 below the lower ignition limits defined by this test program. It is felt that such a safety factor more than compensates for any variation in the lower ignition limit due to effective pipe length, wall temperature, and non-uniformity in oil contamination.

## I LITERATURE SURVEY

In recent years the phenomenon of compression ignition in pneumatic systems has been extensively studied. Such ignitions are induced in systems contaminated with sufficient quantities of a combustible organic material and exposed to rapid pressurization with an oxidizing gas.

Most of the work was concerned with high pressure air systems contaminated with lubricating oil from air compressors. Faeth and White<sup>(1)</sup> conducted a series of tests in which they investigated the combustion characteristics of 25 mg. samples of various lubricating oils placed at the dead end of a pipe exposed to rapid pressurization with air. They defined the combustion ranges as functions of various parameters including the rate of pressurization, air pressure, length of pipe, and the initial ambient system temperature. The results of these tests quantify the expected trends due to variation in the parameters. Of interest was the fact that the occurrence of combustion was greater, the shorter the length of pipe used down to a length of 1 ft. However, the change in combustion limit was only a mild function of the pipe length.

Wilson<sup>(2)</sup> and co-workers investigated the compression ignition characteristics of lubricating oils evenly coated on the internal surface of a 1 1/2" diameter test pipe. Using a 15 ft. length of pipe, they studied both auto-ignition and flame propagation using 2190 and Callulube lubricating oils. Wilson defined the lower combustible limit in the range of 1000 mg./ft<sup>2</sup> of oil. He also found that used degraded oil was much more reactive than new unused oils - attributing this to breakdown products in the oil. This conclusion was later verified by Zabetakis<sup>(3)</sup>. Both Wilson<sup>(2)</sup> and Perlee<sup>(4)</sup>, concerned with this type of ignition caused by degraded oil blow-by from air compressors, recommend proper cleaning and maintenance to prevent dangerous conditions from arising.

Faeth<sup>(1)</sup> did a theoretical analysis of the rapid compression process in an attempt to predict whether conditions for ignition would exist in a given system. Predictions based on his analysis compared well with his observed experimental data.

Dallinger and McGill<sup>(6)</sup>, concerned with high pressure air systems aboard submarines, surveyed work done in this field. They reported that an increase in pipe diameter increased the temperatures obtained during a rapid compression, thus lowering the concentration of oil necessary for ignition.

Russian interest in the problem of compression ignition with oxygen is indicated by the work of Gordeyev and coworkers<sup>(7)</sup>. They reported lower explosive limits of 26,000 mg./ft<sup>2</sup> of oil and lower combustion limits of 190 mg./ft<sup>2</sup>. However, they used oils with Russian designations, making comparison with 2190 difficult, and they failed to report pertinent parameters such as rates of pressurization and pipe diameters.

The dangers inherent in high pressure air systems, due to compression ignition, lead to questions about parallel phenomenon in high pressure oxygen systems. Baum<sup>(8)</sup> and his coworkers investigated 7500 psi oxygen systems, looking at effects of system temperature, vibration, shock, extended storage, contamination, and material compatibility. In regard to ignition, Baum<sup>(8)</sup> recommended "absolute" cleanliness from organics, and homogeneous hydrocarbon concentrations less than 50 ppm.

In 1961, Kehat<sup>(9)</sup> studied ignition and flame propagation at pressures up to 1500 psi using gaseous oxygen. Using a 1/4" test pipe evenly coated with oil, he used a spark and an electric match to induce ignition. Kehat found no significant danger of ignition at concentrations up to 1000 mg./ft<sup>2</sup> of oil. Also of interest was Kehat's technique of evenly coating a small diameter pipe, 1/2", with oil - namely of dissolving the oil in a solvent, carbon tetrachloride, applying the solution to the pipe, and then allowing the solvent to evaporate, leaving a thin film of oil behind.

A summary of the literature survey is presented in Section 4.2.2 and compared with the recommendations of this current test program.

## II TEST APPARATUS

### 2.1 General Description

The basic test apparatus is illustrated schematically in Figure 1.

The test apparatus basically consisted of a four foot section of 1/2 inch, schedule 80, monel pipe plugged at one end. The high pressure oxygen was supplied from a two tank system. One tank (15 ft<sup>3</sup> capacity) at 3800 psig was used to supply the second tank (9 ft<sup>3</sup>) at 3000 psig which, in turn, supplied the test pipe. The instrumentation and control equipment consisted of flame probes, pressure transducers, thermocouples, external strain gauges, a 5000 psi relief valve, a gas sampling bomb, a manually operated ball valve, a quick opening Marotte valve, a check valve and a manually operated vent valve. Photographs of the test pipe, in place for a run, are shown in Figures 2, 3, 4, and 5. A more detailed description of the components is found in Table I.

A four foot length of pipe was selected as the optimum length. This selection is a compromise between the minimum length recommended by Faeth and White and the physical instrumentation requirements necessary to measure flame propagation.

This test apparatus was modified slightly during the test program to accommodate a sparking device. This alteration consisted of removing the pressure transducer ( $P_1$ ) closest to the dead end and replacing it with the sparking device. This modification is shown in Figure 6.

In addition to the pipe just described, another test pipe was used during the final portion of testing with a minimum of instrumentation located adjacent to the dead end. This second pipe was also a 4 ft. length of 1/2" monel pipe, but it had only two instrumentation taps and a vent line. This test set up is shown schematically in Figure 7. Photographs of this test set up are shown in Figures 8 and 9. As can be seen in the figures a nichrome heating coil was added to the pipe to give the capability of operating at elevated initial temperatures.

TABLE I  
LIST OF APPARATUS AND COMPONENTS

<u>Component</u>	<u>Description</u>
1 3000 psig O <sub>2</sub> supply flask	9 ft <sup>3</sup> , MIL-C-1511A
1 3800 psig O <sub>2</sub> supply flask	15 ft <sup>3</sup> , MIL-C-28C9B
1 Solenoid valve	Marotta M.V. 173
1 Pressure gauge	0 to 10,000 psig scale
1 Check valve	3/4"
3 Nipples	3/4" pipe, NiCu
1 Pipe union	1/2", NiCu
1 Pipe cap	1/2", NiCu
1 Flame arrestor*	1/2" I.P.S. NiCu
1 Pipe tee	3/4" monel
1 Test pipe	4 ft. Sch. 80, 1/2" Monel MIL-T-1368 Ty. 1, Cond. 1
1 Relief valve	5,000 psi relieve pressure
1 Ball valve	3/4", manual
1 Vent valve	1/4", manual
1 Gas sampler bomb	See Section 2.2
4 Strain gauges	"
3 Pressure transducers	"
3 Temperature probes	"
4 Flame probes	"

\*The flame arrestor used in Runs A-1 through A-24, Runs D-1 through D-5, and F-36 was made of stainless steel.

For safety considerations, the test pipe was located in a concrete test pit and was surrounded by sand bags. The control and instrumentation read-outs were located in an adjacent test pit.

## 2.2 Instrumentation

Instrumentation on the basic test set-up consisted of four flame probes spaced 12 inches apart; three pressure transducers, 18 inches apart; four external strain gauges; and three thermocouples, 18 inches apart. The locations of this instrumentation can be seen in the figures presented in the previous section.

In addition to this basic instrumentation, a sparking device was used in an attempt to induce ignitions. The location of this sparking device can also be seen in the figures presented in the previous section. The outputs of the instrumentation probes were recorded on an oscillograph recorder with a paper speed of 64 inches/sec.

A close-up of a flame probe, a thermocouple, and the sparking device is shown in Figure 10; while a photograph of a pressure transducer, a strain gauge installation and the flame arrestor is shown in Figure 11.

A block diagram of the basic instrumentation is presented in Figure 12, while a schematic of the control circuitry is shown in Figure 13. A detailed list and description of the major instrumentation components is presented in Table II. The time sequence in which all the instrumentation and control circuitry operated is shown in Figure 14.

The instrumentation block diagram (Figure 12) shows the relationship of the control circuitry and the instrumentation probes to the recording device. The flame probes, the pressure transducers, and the strain gauges are energized by the circuitry; signals detected by them are then transmitted to the recorder. The thermocouples, using an ice water bath as a reference, generate voltages indicative of their temperatures and transmit them to the recorder. The sparking

TABLE II  
LIST OF INSTRUMENTATION EQUIPMENT

<u>Description</u>	<u>Manufacturer</u>
1. Chromel/alumel thermocouples (400 milliseconds response time)	Thermoelectric, Inc.
2. Chromel wire flame probes - porcelain base with 300 volt operating potential ( $< 1.7$ millisecond response time)	Electric Boat Division (not commercial)
3. Pressure transducers - 10,000 lbs/in <sup>2</sup> , air cooled type (.05 millisecond response time)	Norwood, Model 102
4. UV light beam oscillograph recorder - galvanometer (1.7 millisecond response time)	Consolidated Electrodynamics Corp. (C.E.C.), Model #124
5. 3 kc. carrier amplifiers for recorder (15 units)	C.E.C., Model #1-113B
6. Modulators for amplifiers (7 units) (2 milliseconds response time)	C.E.C., Model #15-605
7. Oscillator power supplies (2 units)	C.E.C., Model #2-105B
8. Variable AC Voltage Supply	Superior Electric
9. Displacement Transducer ( $< 1.7$ millisecond response time)	Underwater Explosion Research Division (USN)
10. Potentiometer and Switch (Auxilliary temperature measurements)	Leeds & Northrup
11. 120 OHM Strain Gauges	Baldwin-Lima-Hamilton



device discharges the electrical energy stored in its capacitors (Figure 13B) and this event is transmitted to the recorder. The potentiometer attached to the stem of the Marotta valve is energized by the circuitry and transmits the change in position of the valve to the recorder. The Marotta valve itself is activated by the control circuitry; while time delay relay 1 (TD-1) and (TD-2) are activated by the same circuitry and in turn control parts of this circuitry (Figure 13).

As seen in Figure 13A, the circuitry is activated by closing switch S-1 which energizes the indicating light plus the flasher and bell to warn personnel in the area that a test is in progress. A test is initiated by closing S<sub>3</sub> which activates relay K<sub>1</sub> and TD-1. Activation of relay K<sub>1</sub> closes contacts K<sub>1-1</sub> which keeps itself activated without regard to the position of S<sub>3</sub>. Activation of TD-1 closes the contacts TD1-1 which, in turn, activates TD-2, TD-3, and opens the Marotta valve. Activation of TD-2, after a preset period of time, shuts down the circuitry. Activation of TD-3, after a preset period of time, opens contacts TD3-1, closing the Marotta valve.

The circuitry of Figure 13B is activated by the closing of TD3-2, which has been closed by TD-3 of Figure 13A. With the contact TD3-2 closed, TD-5 is activated which in turn closes contacts TD5-1 activating K<sub>2</sub>. K<sub>2</sub> closes contact K<sub>2-1</sub> which discharges the spark. The 600  $\mu$ F capacitors are charged by closing switch S<sub>4</sub>. Contacts TD3-3 isolate this circuit while TD-3 is activated.

### III TEST PROCEDURE

#### 3.1 Preliminary Tests

Before proceeding with ignition tests, it was necessary to verify the proposed oil coating technique.

Preliminary tests were conducted to determine whether the oil coating technique selected would be satisfactory. This technique consisted of dissolving the oil in a low-boiling, non-combustible solvent Freon TF and then applying the resultant solution to the pipe. Upon evaporation of the Freon, a residual coating of oil was left on the pipe. In evaluating this technique, a sacrificial section of pipe was used. After the coating operation, the pipe was cut into small sections and the amount of oil present on each section was determined by chemical analysis. This analysis consisted of removing the oil from the pipe segments with a low boiling point solvent, then evaporating off the solvent. The amount of oil removed from each segment was then determined gravimetrically.

This same method of evaluation was used to determine the feasibility of other oil coating techniques as part of the verification. The other oil coating techniques evaluated were: a) applying the oil directly with a circular brush or swab, b) spraying the oil onto the pipe, c) partial removal of a heavy oil coating using a solvent.

#### 3.2 Test Plan

The test plan included the following areas of investigation:

- 1) The coating of the test pipe with various concentrations of 2190 oil at ambient temperatures and attempting a compression ignition with 3000 psi oxygen; the oil coatings were increased in small increments in concentrations ranging from 0.08 mg/ft<sup>2</sup> to 640 mg/ft<sup>2</sup>.
- 2) Placing droplets of 2190 oil near the dead end of the test pipe

and attempting a compression ignition with 3000 psi oxygen; the size of the oil droplets used ranged from 5 microliters ( $\mu\ell$ ) to 90  $\mu\ell$ . 3) Employing the combination of coating the test pipe with 2190 oil and placing a droplet of the oil near the dead end of the pipe, and attempting a compression ignition with 3000 psi oxygen; oil coating concentrations ranged from 12  $\text{mg}/\text{ft}^2$  to 88  $\text{mg}/\text{ft}^2$ , while the oil droplets ranged in size from 10  $\mu\ell$  to 100  $\mu\ell$ . 4) Coating the test pipe with various concentrations of 2190 oil or of Habcool cutting oil, using an electrical spark to induce ignition in a 3000 psi oxygen atmosphere; the 2190 oil concentrations ranged from 0.16  $\text{mg}/\text{ft}^2$  to 640  $\text{mg}/\text{ft}^2$ , while the Habcool concentrations ranged from 100  $\text{mg}/\text{ft}^2$  to 480  $\text{mg}/\text{ft}^2$ . 5) Coating a modified test pipe (minimum instrumentation) with various concentrations of 2190 oil and of Habcool cutting oil, and attempting a compression ignition with 3000 psi oxygen; the 2190 oil concentrations ranged from 30  $\text{mg}/\text{ft}^2$  to 480  $\text{mg}/\text{ft}^2$ , while the Habcool concentration ranged from 160  $\text{mg}/\text{ft}^2$  to 480  $\text{mg}/\text{ft}^2$ .

The 2190 (MIL-L-17331A) lubricating oil was selected to be used during this test program because most of the previous investigations of compression ignition and flame propagation were performed using this lubricating oil. Thus, in order to compare results with the earlier work, 2190 was used in this investigation. Habcool #318 cutting oil was selected as a typical organic cutting oil used in a machine shop with which piping might be contaminated during fabrication in actual practice.

### **3.3 Testing Procedure**

In making a test run the following procedure was used:

1. The test pipe was thoroughly cleaned by use of Freon (TF) rinses and a lint-free cloth.
2. With the instrumentation probes replaced by plugs, the pipe was filled with an oil/Freon solution of a known concentration.
3. The oil/Freon solution was emptied and the remaining Freon allowed to evaporate (15 to 20 minutes was allowed) leaving a thin, uniform oil film.
4. The instrumentation probes were replaced into their proper positions.
5. The test pipe was placed "on-line", putting the flame arrester into position within the union.
6. The calibration of the instrumentation was checked as well as the electrical control circuitry.
7. The position of all hand-operated valves was checked.
8. The test area was cleared of all personnel and the test was made.
9. After the run was completed, the electrical power was turned off and all valves secured.
10. A gas sample was taken for analysis and the test pipe was vented down to ambient pressure.
11. The test pipe was disconnected and was visually inspected.
12. The test pipe and instrumentation probes were thoroughly cleaned in preparation for the next run.

The gas sample at the conclusion of a test was analyzed for carbon dioxide, hydrogen, carbon monoxide, oxygen and nitrogen using gas chromatography.

No attempt was made to replace the air in the test pipe with pure  $O_2$  before the rapid pressurization. This initial air is compressed to within  $1/4$ " of the dead end during compression from 14.7 psi to 3000 psi and it was felt that this small amount of nitrogen, in comparison to the amount of oxygen present, would have a negligible effect on the results.

## IV TEST RESULTS

### 4.1 Preliminary Tests

The oil coating tests showed that the selected technique of dissolving the oil in a non-combustible solvent, Freon 12, gave the best results. The oil was evenly coated onto the surface of the pipe and the concentration of oil on the pipe was proportional to the concentration of oil in the Freon. This relationship is shown in Table III. It was also found to be the best technique from an operational point of view - being less prone to technician error and having good reproducibility.

TABLE III  
OIL COATING CONCENTRATIONS  
(1/2" Monel Pipe)

<u>Oil Concentration in Freon - PPM</u>	<u>Oil Coating on Pipe - mg/ft<sup>2</sup></u>	<u>Oil Film Thickness - microns</u>
5	0.0125	$1.2 \times 10^{-3}$
10	0.025	$2.4 \times 10^{-3}$
100	0.25	$2.4 \times 10^{-2}$
1000	2.5	0.24
10,000	24.6	2.4
50,000	123.0	12.0
100,000	246.0	24.0
200,000	492.0	48.0
300,000	738.0	72.0

### 4.2 Ignition Test Results

The results of the test program are presented in the following sections. The significant data are presented in Section 4.2.1 and Appendix A, while an interpretation and discussion of these data is found in Section 4.2.2.

#### 4.2.1 Data

The parameters measured during each of the test runs are presented in Appendix A. The parameters include: oil concentration, ambient temperature, maximum pressure, rate of pressure rise, incident temperature, flame probe saturation, and strain gauge indication. The temperatures presented are the maximum temperatures recorded. Due to the lag and response time of the thermocouples, the actual temperatures experienced are much higher.

It should be noted that at various points throughout the test program blank runs were made - i.e. runs in which no oil was put into the test pipe. These runs were made to determine the time histories of the various parameters due only to the rapid pressurization of the test pipe with oxygen, and to serve as a check on the instrumentation. The results of all the blank runs were identical, therefore, the results of only one of these runs is presented.

#### 4.2.2 Correlation and Interpretation of Results

The first test run with oil contamination resulted in a violent explosion causing extensive damage to the test apparatus. This damage is shown in Figures 15-20. A comparison can be made with the undamaged test pipe shown in previous sections.

As can be seen in the figures a two inch section of the test pipe, at the inlet end, was disintegrated along with part of the couplings; one of the pressure probes was blown out of its fitting; several of the electrical connectors to the instrumentation probes were burned; and the copper "O" ring seals on the two remaining pressure transducers were blown out. In addition, the flame arrestor was disintegrated and the check valve welded closed. It should be noted that for this run the coupling at the inlet end of the test pipe and the flame arrestor were both made of stainless steel.

After the explosion, it was felt that the level of oil contamination,  $160 \text{ mg/ft}^2$ , was well into the explosive range and that the testing should be conducted in a much lower range of oil concentration. Subsequent test runs showed, however, that levels of oil contamination higher than  $160 \text{ mg/ft}^2$  can withstand rapid pressurization by pure oxygen without resulting in an explosion. These subsequent tests employed monel and nickel-copper components only, while the initial run at  $160 \text{ mg/ft}^2$ , which resulted in an explosion, had a stainless

steel flame arrestor. In an attempt to duplicate the explosive run, a stainless steel flame arrestor was put into the system (Run F-3). In this run, though the test pipe did not rupture, there was an ignition and flame propagation, as opposed to no indications of ignition or flame propagation under the same conditions using a monel flame arrestor (Run F-18).

No completely satisfactory explanation of the explosion is available, but it is significant that no indications of ignition were obtained with the monel flame arrestor, while very definite indications were observed with the stainless steel flame arrestor.

As a result of the explosion in the first run, subsequent tests were conducted (Runs A2-A24, F1-F28, F35) by evenly coating the test pipe with 2190 oil with concentrations of 0.08 mg./ft<sup>2</sup> up to 160 mg./ft<sup>2</sup> without any indication of ignition. At oil concentrations between 173 mg./ft<sup>2</sup> and 266 mg./ft<sup>2</sup> there were indications of ignition and flame propagation on some instrumentation probes but not on others. At oil concentrations above 266 mg./ft<sup>2</sup>, there were definite indications of ignition and flame propagation. These results can be seen graphically in Figure 21.

On the supposition that it was possible to have sufficient oil in the test pipe to propagate a flame while not having a high enough concentration to cause an auto-ignition, a series of runs were conducted in which it was attempted to induce an ignition. Thus runs (Runs D1-D14, AD1-AD10) were made with small drops of oil placed near the dead end of the test pipe (thus giving a high local concentration of oil). Oil droplets ranging in size from 5 $\mu$ l to 100 $\mu$ l were used alone and in combination with a uniform oil film ranging in concentrations from 12 mg./ft<sup>2</sup> to 88 mg./ft<sup>2</sup>. In Run D-12, the gas analysis after the run revealed the presence of CO<sub>2</sub>, indicating combustion of the oil. However, the other instrumentation did not confirm this combustion, and the results could not be duplicated in subsequent tests at similar or more severe conditions (Runs AD-8 and AD-10).



In the test runs with oil droplets of 70  $\mu$ l or greater,  $H_2$  was found in the gas after the test run. It is believed that this  $H_2$  is a product of the thermal decomposition of the oil. However, oil droplets were used only in an attempt to induce ignitions. This they failed to do, so the appearance of  $H_2$ , though interesting, was not investigated further. However, two things should be noted in this regard: 1) the occurrence of oil droplets, 70  $\mu$ l or greater, in a clean oxygen system is highly unlikely and 2) the  $H_2$  appeared in concentrations well below its lower ignition limit.

Having failed to induce ignition using small droplets of oil, an attempt was made to induce ignition by means of an electrical spark. A series of runs were made using the sparking device (Runs F1-F28, F35, F36). In those runs in which ignitions were observed, they were of the compression ignition type, and involved oil concentrations above 160 mg./ft<sup>2</sup>. The spark did not initiate any ignitions nor affect flame propagation in any way.

Test runs made in which the pipe was evenly coated with Habcool at 70°F (Runs F29-F34) indicated that the no ignition range was below about 110 mg./ft<sup>2</sup>, while the ignition range was somewhere above 250 mg./ft<sup>2</sup>. Comparison of these results with those obtained for 2190 (Figure 21) show Habcool to be somewhat more reactive.

In an attempt to attain higher temperatures from the rapid compression with pure oxygen, a series of tests were run using a pipe with a minimum of instrumentation located adjacent to the dead end. Because of the minimum amount of instrumentation, it was more difficult to get accurate indications of ignition and flame propagation. However, from the instrumentation available, the following results were obtained:

- A. Using 2190 oil at 70°F, tests were run at concentrations of 80 mg./ft<sup>2</sup> to 480 mg./ft<sup>2</sup> (Runs C1-C6). No ignition was obtained.

below  $160 \text{ mg/ft}^2$ , while the ignition range was found to be above  $200 \text{ mg/ft}^2$ . These results approximate the results shown in Figure 21.

- B. Using 2190 oil heated to  $125^\circ\text{F}$ , tests were run at concentrations ranging from  $160 \text{ mg/ft}^2$  to  $480 \text{ mg/ft}^2$  (Runs C7-C11). Indications of ignition were seen at the lower concentration of  $160 \text{ mg/ft}^2$ , indicating that the level of oil contamination necessary for ignition and flame propagation are somewhat lower at  $125^\circ\text{F}$  than at  $70^\circ\text{F}$ .
- C. Using Habcool cutting oil at  $70^\circ\text{F}$ , tests were run at concentrations ranging from  $160 \text{ mg/ft}^2$  to  $480 \text{ mg/ft}^2$  (Runs C12-C16). Definite ignitions were obtained above  $200 \text{ mg/ft}^2$ , while the non-ignition range is somewhere below  $160 \text{ mg/ft}^2$ . The non-ignition and ignition ranges are about the same as those found with the regular test pipe.
- D. Using Habcool oil at  $125^\circ\text{F}$ , tests were run at concentrations ranging from  $160 \text{ mg/ft}^2$  to  $480 \text{ mg/ft}^2$  (Runs C17-C21). The results from these tests were not distinguishable from those at  $70^\circ\text{F}$ .

A more quantitative correlation can be made by plotting the resultant %  $\text{CO}_2$ , the maximum recorded temperature, and maximum pressure obtained as functions of level of oil concentration. Such plots are shown in Figures 22, 23 and 24. It can be seen that these results match the non-ignition, transient, and ignition ranges earlier defined more qualitatively.

The carbon dioxide found was considerably less than would be dictated with 100% combustion of the oil. This is, however, consistent with the fact that carbon deposits and residual oil were found in the test pipe after ignitions, indicating considerably less than complete combustion.

It is possible to compare these results with those obtained by earlier investigators. Kehat reports little danger of ignition in oxygen

systems up to 1500 psig at concentrations up to 1000 mg/ft<sup>2</sup>. This compares to our lower ignition limit of 160 mg/ft<sup>2</sup> at 3000 psi. Kehat used a pure compound, n-hexadecane, for his work, while our work was performed with a commercial lubricating oil, 2190. Comparison of the properties of n-hexadecane and 2190 - e.g., molecular weights, auto-ignition temperature - and considering the differences in the oxygen pressures used reveal our results to be in reasonable agreement with Kehat.

It is also possible to compare our results with the results of Wilson et al. Wilson reports the lower ignition limit in high pressure air systems at 3000 psig of 2190 oil to be 1000 mg/ft<sup>2</sup>. By extrapolating these results to pure oxygen at 3000 psig, a lower ignition limit would be approximately 200 mg./ft<sup>2</sup>. This is in good agreement with the current results.

A summary comparison of the current results and those obtained by earlier investigators is presented in the following chart:

<u>Investigator</u>	<u>Test Gas</u>	<u>Test Apparatus</u>	<u>Oil Used</u>	<u>Lower Ignition Limit</u>
Kehat	O <sub>2</sub> , 1500 psi	1/2" pipe	C <sub>12</sub> H <sub>26</sub>	1000 mg/ft <sup>2</sup>
Wilson	Air, 3000 psi	1-1/2" pipe	2190	1000 mg/ft <sup>2</sup>
Gordeyev	O <sub>2</sub>	-	-	190 mg/ft <sup>2</sup>
Presti	O <sub>2</sub> , 3000 psi	1/2" pipe	2190	160 mg/ft <sup>2</sup>
Baum	O <sub>2</sub> , 1500 psi	(50 ppm homogeneous mixture - recommended lower limit)		

## V CONCLUSIONS AND RECOMMENDATIONS

From the test results obtained the following conclusions can be drawn for 3000 psi oxygen systems:

1. Compression ignition and flame propagation will not occur below oil concentrations of  $160 \text{ mg/ft}^2$  at oil temperatures below  $125^\circ\text{F}$  in a  $1/2"$ , schedule 80, monel pipe.
2. Compression ignitions will not occur with oil droplets or puddles smaller than  $100 \mu$  in size.
3. Habcool cutting oil was found to be somewhat more reactive than 2190.
4. Present oxygen clean standards (as specified by Shipyard Standard Practice 1.9 Rev. F, Oxygen and Nitrogen Systems, paragraph 3.2.3.1d) which require oil film concentrations of  $0.013 \text{ mg/ft}^2$  or less, are much too rigorous and can be relaxed. The oil film concentration of  $0.013 \text{ mg/ft}^2$  corresponds to obtaining a concentration of 5 ppm in a Freon rinse, while  $160 \text{ mg/ft}^2$  corresponds to 65,000 ppm in Freon.

It is recommended that the allowable oil film concentration specified by the above Shipyard Standard Practice be set at 500 ppm oil concentration in a freon rinse. This concentration allows a safety factor of 100 over the values at which oil ignitions begin to occur.

It is felt that such a safety factor more than adequately allows for any variation in the lower ignition limit due to variations in effective pipe lengths below 4 feet which might be found in an actual pressurized oxygen system.

5. To obtain pressure rises greater than 25% requires oil concentrations greater than 500 ng./ft<sup>2</sup>. This can be seen from Figure 24.

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## APPENDICES



ASTM D 153  
Oxygen Pressure Vessel Test

DATE	RUN NO.	CONC. (mg/ft <sup>2</sup> )	AMBI- ENT TEMP. (°F)	MAXIMUM PRESSURE IN X TIME (psig in x msec)			RATE OF PRESSURE RISE INSIDE PIPE (psig/sec)	MAXIMUM INSIDE P. (°F in x msec)	
				P <sub>1</sub>	P <sub>2</sub>	P <sub>3</sub>		P <sub>1</sub>	P <sub>2</sub>
	BLANK	0	70	3000 in 100	3000 in 100	3000 in 100	26000	200 in 100	100 in 100
10/6	A-1	160	70	>3000 in 90	>3000 in 90	>3000 in 90	20900	>2000 in 100	>2000 in 100
10/21	A-2	0.08	70	3000 in 100	3000 in 100	3000 in 100	23000	200 in 100	100 in 100
10/24	A-3	0.16	70					SAME AS RUN A-2	
10/24	A-4	0.32	70					SAME AS RUN A-2	
10/25	A-5	0.64	70					SAME AS RUN A-2	
10/26	A-6	1.60	70					SAME AS RUN A-2	
10/26	A-7	2.13	70					SAME AS RUN A-2	
10/26	A-8	2.67	70					SAME AS RUN A-2	
10/27	A-9	3.20	70					SAME AS RUN A-2	
10/27	A-10	4.00	70					SAME AS RUN A-2	
10/27	A-11	5.25	70					SAME AS RUN A-2	
10/28	A-12	7.82	70					SAME AS RUN A-2	
10/28	A-13	10.3	70					SAME AS RUN A-2	
10/28	A-14	12.8	70					SAME AS RUN A-2	
10/31	A-15	20.0	70					SAME AS RUN A-2	
10/31	A-16	24.6	70					SAME AS RUN A-2	
11/1	A-17	35.6	70					SAME AS RUN A-2	
11/2	A-18	45.7	70					SAME AS RUN A-2	
11/2	A-19	55.2	70					SAME AS RUN A-2	
11/2	A-20	64.0	70					SAME AS RUN A-2	
11/14	A-21	72.3	70	2900 in 139	2700 in 139	2700 in 139	28000	225 in 100	150 in 100
11/15	A-22	80.0	70	2960 in 141	2800 in 141	2800 in 141	28000	240 in 100	170 in 100

A

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[illegible]

DATE	RUN NO.	CONC. (mg/ft <sup>2</sup> )	AMBI- ENT TEMP. (°F)	MAXIMUM PRESSURE IN X TIME (psig in x msec)			RATE OF PRESSURE RISE INSIDE PIPE (psig/sec)	MAXIMUM DISCHARGE (cm <sup>3</sup> in x sec)	
				P <sub>1</sub>	P <sub>2</sub>	P <sub>3</sub>		T <sub>1</sub>	T <sub>2</sub>
		<b>[2190]</b>							
11/15	A23	87.2	70	3000 in 143	2800 in 143	2760 in 143	27000	240 in 153	165 in 105
11/15	A-24	94.1 (μl)	70	2940 in 141	2800 in 141	2740 in 141	23000	225 in 160	160 in 100
11/3	D-1	5	70	3000 in 133	2800 in 133	2760 in 133	28500	250 in 165	175 in 100
11/4	D-2	10	70					SAME AS RUN A-2	
11/4	D-3	15	70					SAME AS RUN A-2	
11/7	D-4	20	70					SAME AS RUN A-2	
11/7	D-5	5	70					SAME AS RUN A-2	
11/18	D-6	30	70	2900 in 105	2800 in 105	2720 in 105	37000	210 in 135	160 in 100
11/18	D-7	40	70	2900 in 104	2800 in 109	2760 in 109	35000	225 in 180	160 in 100
11/21	D-8	50	70	2950 in 101	2840 in 101	2760 in 101	33500	235 in 195	150 in 100
11/21	D-9	60	70	2950 in 104	2880 in 104	2760 in 104	36500	215 in 175	150 in 100
11/22	D-10	70	70	2920 in 105	2840 in 105	2720 in 105	36500	215 in 155	150 in 100
11/22	D-11	80	70	2940 in 105	2720 in 105	2720 in 105	38000	225 in 190	160 in 100
11/23	D-12	90	70	3100 in 97	2760 in 97	2650 in 97	42000	245 in 285	200 in 165
11/8	D-13	10	125					SAME AS RUN A-2	
11/9	D-14	10 (Habeool)	125					SAME AS RUN A-2	
11/9	AD-1	12.3+ 10 μl	125					SAME AS RUN A-2	
11/10	AD-2	24.6+ 10 μl	125					SAME AS RUN A-2	
11/10	AD-3	35.6+ 10 μl	125					SAME AS RUN A-2	

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1. ...  
2. ... 55,000 rpm/min

S	T <sub>1</sub>	T <sub>2</sub>	MAXIMUM VISIBLE TEMP. (°F)	GAS TEMPERATURE INDICATION IN X TESTS (°F/in in X msec)				FLAME PROBE INDICATION TIME (msec)				PER- CENT CO <sub>2</sub>
				S <sub>1</sub>	S <sub>2</sub>	S <sub>3</sub>	S <sub>4</sub>	F <sub>1</sub>	F <sub>2</sub>	F <sub>3</sub>	F <sub>4</sub>	
1	103	100 in 20	70	150 in 103	150 in 103	150 in 103	150 in 103	-	-	-	-	0
2	100	100 in 21	70	150 in 101	150 in 101	150 in 101	150 in 101	-	-	-	-	0
3	100	100 in 20	70	150 in 103	150 in 103	150 in 103	150 in 103	-	-	-	-	0
4	100	100 in 20	70	150 in 105	150 in 105	150 in 105	150 in 105	-	-	-	-	0
5	100	100 in 10	70	150 in 109	150 in 109	150 in 109	150 in 109	-	-	-	-	0
6	100	100 in 20	70	150 in 101	150 in 101	150 in 101	150 in 101	-	-	-	-	0
7	100	100 in 10	70	150 in 104	150 in 104	150 in 104	150 in 104	-	-	-	-	0
8	100	100 in 20	70	150 in 105	150 in 105	150 in 105	150 in 105	-	-	-	-	0
9	95	95 in 10	70	150 in 105	150 in 105	150 in 105	150 in 105	-	-	-	-	0
10	100	100 in 100	70	150 in 97	150 in 97	150 in 97	150 in 97	-	-	-	-	6.58
			125									1.1% <sub>H2</sub>
			125									
			125									
			125									
			125									

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SECTION 1 - TESTS

Oxygen Concentration in Air

DATE	RUN NO.	CONC. (mg/ft <sup>2</sup> )	AMBI- ENT TEMP. (°F)	MAXIMUM PRESSURE IN X TUBE (psig in x msec)			RATE OF PRESSURE RISE INSIDE PIPE (psig/sec)	MAXIMUM TEMPERATURE	
				P <sub>1</sub>	P <sub>2</sub>	P <sub>3</sub>		T <sub>1</sub>	T <sub>2</sub>
		[2190]							
11/10	AD-1	45.7+ 10 µl	125						SAME AS RUN A-2
11/14	AD-5	64.0+ 10 µl	125						SAME AS RUN A-2
11/28	AD-6	55.2+ 70 µl	70				12000		SAME AS RUN A-2
11/29	AD-7	64.0+ 80 µl	70				16000		SAME AS RUN A-2
11/29	AD-8	72.2+ 90 µl	70				18000		SAME AS RUN A-2
11/30	AD-9	80.0+ 100 µl	70	2920 in 109	2840 in 109	2720 in 109	17000	280 in 300	100 in 100
11/30	AD-10	87.2+ 100 µl	70	2880 in 106	2760 in 106	2720 in 106	16000	280 in 350	115 in 100
3/30	F-1	0.16	70				38000		SAME AS RUN A-2
3/30	F-2	1.60	70				38000		SAME AS RUN A-2
3/31	F-3	7.81	70				38000		SAME AS RUN A-2
3/31	F-4	15.2	70				38000		SAME AS RUN A-2
3/31	F-5	24.6	70				38000		SAME AS RUN A-2
4/3	F-6	35.6	70				38000		SAME AS RUN A-2
4/3	F-7	45.8	70				38000		SAME AS RUN A-2
4/3	F-8	55.2	70				38000		SAME AS RUN A-2
4/4	F-9	64.0	70				38000		SAME AS RUN A-2
4/4	F-10	72.3	70				38000		SAME AS RUN A-2
4/4	F-11	80.0	70				38000		SAME AS RUN A-2
4/4	F-12	87.2	70				38000		SAME AS RUN A-2

A

INJECTION RATE: 100

Flow rate: 100,000 gal/sec

INJECTION RATE		MATERIAL OUTSIDE TEMP. (°F)	DEPART GASES & INJECTION IN X SECS (min/hr. in X msec)				FLAME PROBE EXTINGUISHING TIME (msec)				PER- CENT CO <sub>2</sub>
S	T		S <sub>1</sub>	S <sub>2</sub>	S <sub>3</sub>	S <sub>4</sub>	P <sub>1</sub>	P <sub>2</sub>	P <sub>3</sub>	P <sub>4</sub>	
		125									0
		125									0
		70									0
		70									1.4% <sub>H<sub>2</sub></sub>
		70									1.3% <sub>H<sub>2</sub></sub>
		70									1.0% <sub>H<sub>2</sub></sub>
5-1-102	105 in 25	70	150 in 109	150 in 109	150 in 109	150 in 109	20	20	20	20	0
5-1-100	95 in 20	90	155 in 106	150 in 106	150 in 106	155 in 106	-	-	-	-	0

B

DATE	RUN NO.	CONC. (mg/ft <sup>2</sup> )	AMBI- ENT TEMP. (°F)	MAXIMUM PRESSURE IN X TIME (psig in x msec)			RATE OF PRESSURE RISE INSIDE PIPE (psig/sec)	MAXIMUM PRESSURE	
				P <sub>1</sub>	P <sub>2</sub>	P <sub>3</sub>		T <sub>1</sub>	T <sub>2</sub>
4/5	F-13	107	70				38000	SAME AS RUN A-2	
6/6	F-14	120	70				38000	SAME AS RUN A-2	
6	F-15	133	70				38000	SAME AS RUN A-2	
4/7	F-16	147	70	3000 in 100	3000 in 100	3000 in 100	38000	200 in 100	140 in 100
7	F-17	147	70				38000	SAME AS RUN A-2	
4/7	F-18	160	70				38000	SAME AS RUN A-2	
7/10	F-19	133+ 104	70	3000 in 100	3000 in 100	3000 in 100	38000	200 in 100	140 in 100
10	F-20	173	70		2930 in 121	2680 in 121	34500	207 in 582	230 in 582
12	F-21	173	70			2800 in 100	35000	203 in 300	100 in 300
7/12	F-22	266	70		3000 in 102	3000 in 102	38000	205 in 502	210 in 502
13	F-23	266	70		2910 in 103	2680 in 108	40250	306 in 500	236 in 500
4/14	F-24	373	70		2880 in 83	2660 in 83	40250	345 in 300	623 in 300
18	F-25	480	70		2960 in 90	2680 in 98	43000	290 in 550	411 in 550
19	F-26	640	70		4000 in 79	3720 in 79	44000	670 in 332	775 in 332
7/20	F-27	640	70		4320 in 87	3880 in 89	40000	1034 in 140	723 in 140
24	F-28	640	70		4320 in 91	4120 in 92	38000		429 in 140
6/15	F-35	640	70		3760 in 88	3680 in 217	66500	1194 in 207	832 in 137
16	F-36	160	70	2940 in 140	2760 in 140		28500	245 in 277	410 in 280
HARCOOL									
26	F-29	160	70		3320 in 190	3040 in 190	35000	363 in 244	190 in 244
27	F-30	160	70		2800 in 121	2520 in 121	32000	272 in 390	190 in 390
28	F-31	320	70		3000 in 125	2800 in 125	34500	420 in 400	201 in 400
28	F-32	320	70		3120 in 113	2960 in 202	35400	447 in 510	1 in 510
28	F-33	480	70		4680 in 36	6520 in 35	42000	2170 in 103	87 in 221
3	F-34	107	70		3000 in 110	2950 in 110	37000	200 in 110	110 in 110

DATE: 12-10-1964

NO.	INLET TEMP	T <sub>3</sub>	THERMOCOUPLE TEMP. (°F)	FLAME PROBE TEMPERATURE (°F) (MIL/IN IN X MSEC)				FLAME PROBE EXTINGUISHING TIME (msec)				PER- CENT CO <sub>2</sub>
				S <sub>1</sub>	S <sub>2</sub>	S <sub>3</sub>	S <sub>4</sub>	F <sub>1</sub>	F <sub>2</sub>	F <sub>3</sub>	F <sub>4</sub>	
100	120 in 100	70	160	160 in 100	180 in 100	160 in 100	160 in 100	90	-	-	-	0
110	120 in 100	70	165	170	180 in 100	160 in 100	160 in 100	100	100	100	100	<0.10
120	200 in 609	70	170	180 in 100	180 in 100	180 in 100	180 in 100	-	-	161	161	0
130	215 in 20	70	165	160	160 in 100	160 in 100	160 in 100	-	-	-	-	0
140	273 in 1202	70	150	160 in 100	160 in 100	160 in 100	160 in 100	100	100	100	100	0
150	300 in 678	70	170	175 in 100	180 in 100	180 in 100	180 in 100	100	100	100	100	<0.10
160	433 in 600	70	170	186	220 in 116	180 in 100	180 in 100	100	100	100	100	<0.10
170	484 in 1000	70	150	179	185 in 100	180 in 100	180 in 100	100	100	100	100	0.45
180	650 in 272	70	190	180	285 in 246	275 in 246	280 in 246	100	100	117	140	2.00
190	745 in 370	70	290	186	275 in 73	260 in 73	260 in 370	100	89	87	172	2.50
200	934 in 310	70	295	192	290 in 250	210 in 92	200 in 92	100	93	241	254	-
210	800 in 280	70	355	219	285 in 218	235 in 218	230 in 217	100	2	147	214	3.40
220	160 in 420	70	105	140	150 in 140	140 in 140	140 in 140	100	258	351	470	0.16
230	217 in 144	70	200	112	170 in 186	180 in 186	195 in 186	97	1000	91	74	0
240	127 in 20	70	150	130	140 in 130	140 in 130	140 in 130	-	-	-	-	0
250	261 in 690	70	180	125	150 in 125	160 in 125	160 in 125	72	-	325	500	0
260	180 in 850	70	170	100	165 in 133	160 in 200	155 in 123	69	200	320	480	0.57
270	1270 in 45	70	465	35	300 in 35	1380 in 35	260 in 36	34	33	36	36	1.95
280	115 in 10	70	30	105	150 in 105	140 in 105	150 in 105	2	-	-	-	0

B



Oxygen Pressure (psig)

DATE	RUN NO.	CONC. (mg/ft <sup>2</sup> )	AMBI-ENT TEMP. (°F)	MAXIMUM PRESSURE IN X TIME (psig in x msec)			RATE OF PRESSURE RISE INSIDE PIES (psig/sec)	MAXIMUM PRESSURE (psig)	
				P <sub>1</sub>	P <sub>2</sub>	P <sub>3</sub>		T <sub>1</sub>	T <sub>2</sub>
		2190							
5/8	C-1	80.0	70	3160 in 108	-	-	38600		
5/9	C-2	160	70	3000 in 102	-	-	38500		
5/9	C-3	240	70	3160 in 97	-	-	42500		
5/9	C-4	320	70	2960 in 104	-	-	40000		
5/9	C-5	400	70	3050 in 118	-	-	40000		
5/9	C-6	480	70	3240 in 96	-	-	45000		
5/10	C-7	160	125	3050 in 100	-	-	42500	80	100
5/10	C-8	240	125	3200 in 97	-	-	44500	80	100
5/10	C-9	320	125	3070 in 94	-	-	40000	80	100
5/10	C-10	400	125	3100 in 90	-	-	45000	80	100
5/11	C-11	480	125	3190 in 72	-	-	40000	80	100
		HABCOOL							
5/11	C-12	160	70	2910 in 88	-	-	45000		
5/11	C-13	240	70	2880 in 87	-	-	47500		
5/11	C-14	320	70	2960 in 76	-	-	52000		
5/12	C-15	400	70	2910 in 73	-	-	43500		
5/12	C-16	480	70	2910 in 70	-	-	53500		
5/12	C-17	160	125	2800 in 81	-	-	48000	80	100
5/12	C-18	240	125	2890 in 49	-	-	51000	90	110
5/12	C-19	320	125	2810 in 80	-	-	52000	85	100
5/12	C-20	400	125	3000 in 76	-	-	38000	90	110
5/12	C-21	480	125	2900 in 67	-	-	50000	90	110

A

1000 1000 1000

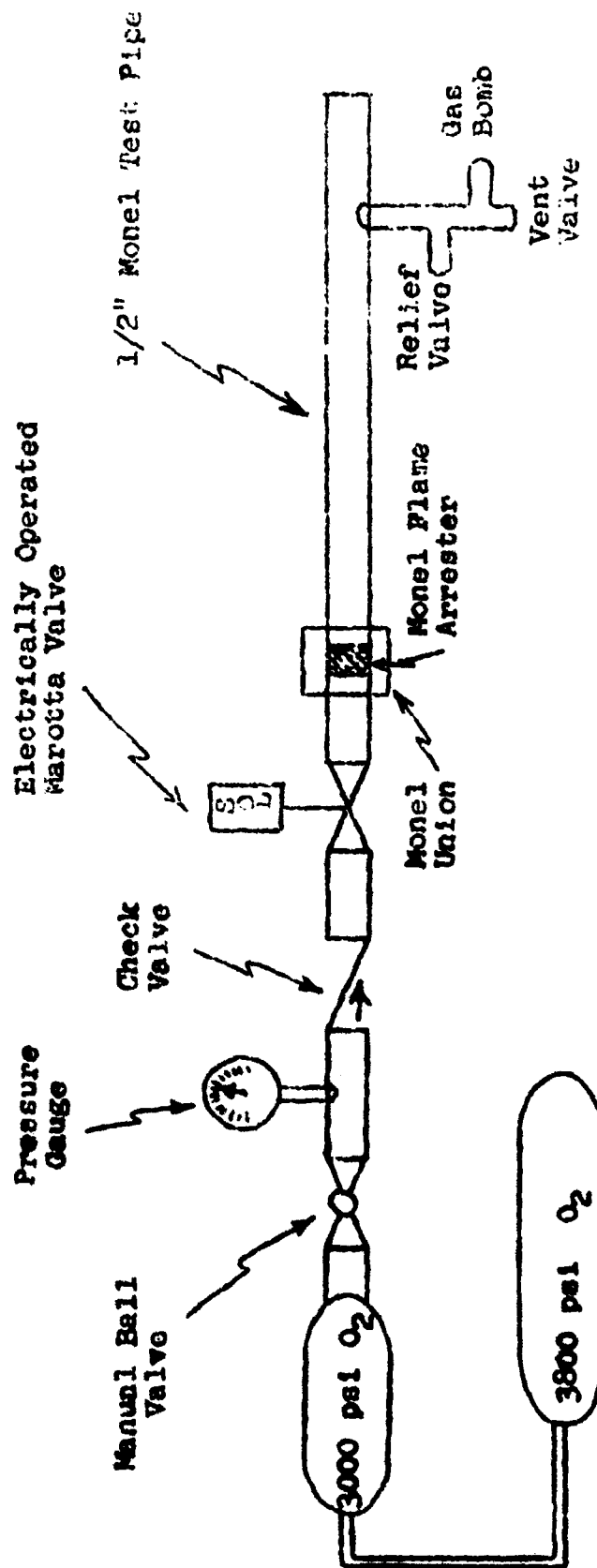
ber 30 1000 1000

SIDE VENTILATION				FLAME PROBE INDICATION TIME (msec)				PER-CENT CO <sub>2</sub>
TIME (msec)				TIME (msec)				
1	2	3	4	5	6	7	8	9
100	100	100	100	100	100	100	100	0
100	100	100	100	100	100	100	100	0
100	100	100	100	100	100	100	100	0.62
100	100	100	100	100	100	100	100	0.32
100	100	100	100	100	100	100	100	0.68
100	100	100	100	100	100	100	100	1.97
100	100	100	100	100	100	100	100	0.23
100	100	100	100	100	100	100	100	0.23
100	100	100	100	100	100	100	100	0.45
100	100	100	100	100	100	100	100	0.77
100	100	100	100	100	100	100	100	2.25
100	100	100	100	100	100	100	100	0.12
100	100	100	100	100	100	100	100	0.50
100	100	100	100	100	100	100	100	1.32
100	100	100	100	100	100	100	100	0.46
100	100	100	100	100	100	100	100	1.85
100	100	100	100	100	100	100	100	0
100	100	100	100	100	100	100	100	0.23
100	100	100	100	100	100	100	100	0.38
100	100	100	100	100	100	100	100	1.09
100	100	100	100	100	100	100	100	1.58

B

# OIL CONTAMINATION IN OXYGEN SYSTEMS DIAGRAM OF TEST APPARATUS

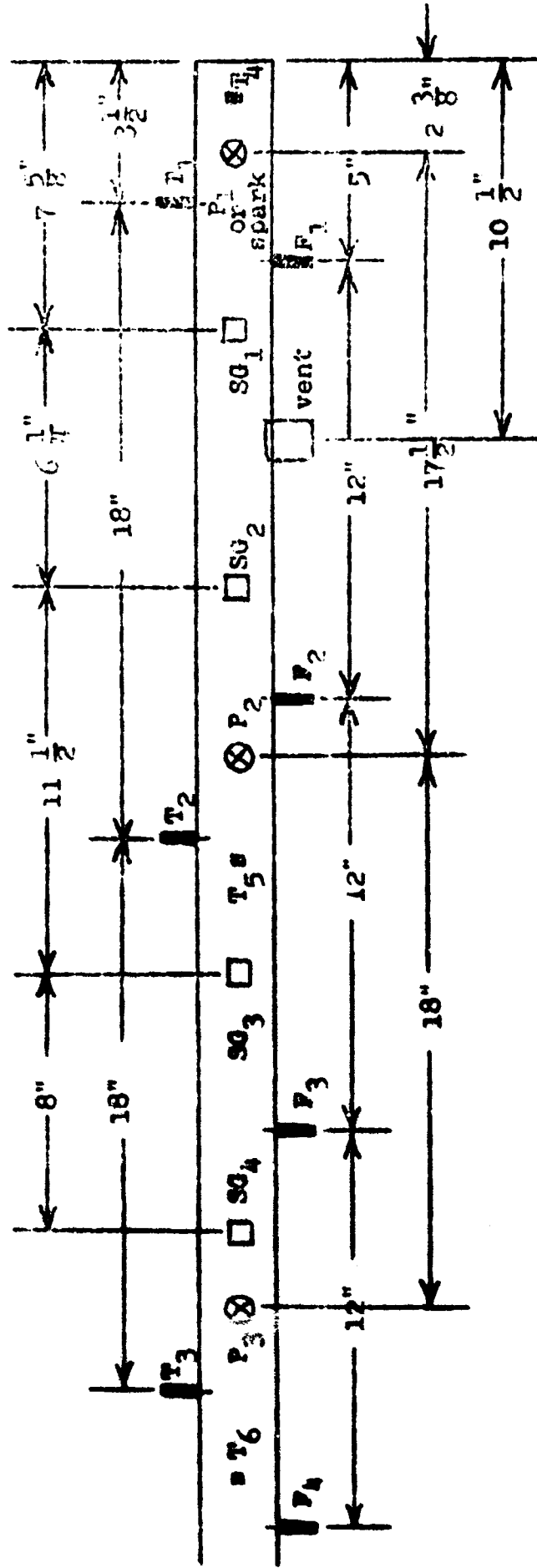
FIGURE 1



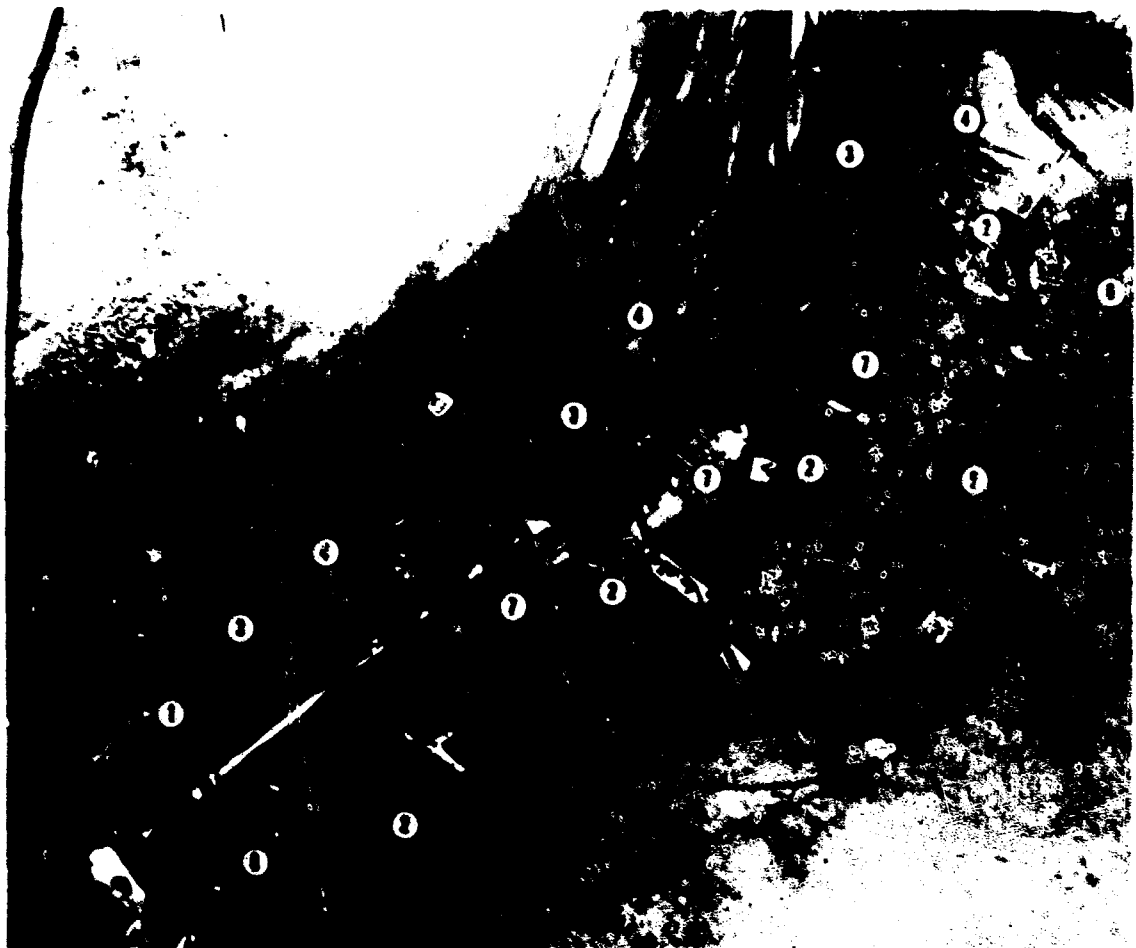
# WIND PIPE LAYOUT OF INSTRUMENTATION (TOP VIEW)

FIGURE 1A

- T - Thermocouple
- P - Flame Probe
- P - Pressure Transducer
- SG - Strain Gauge

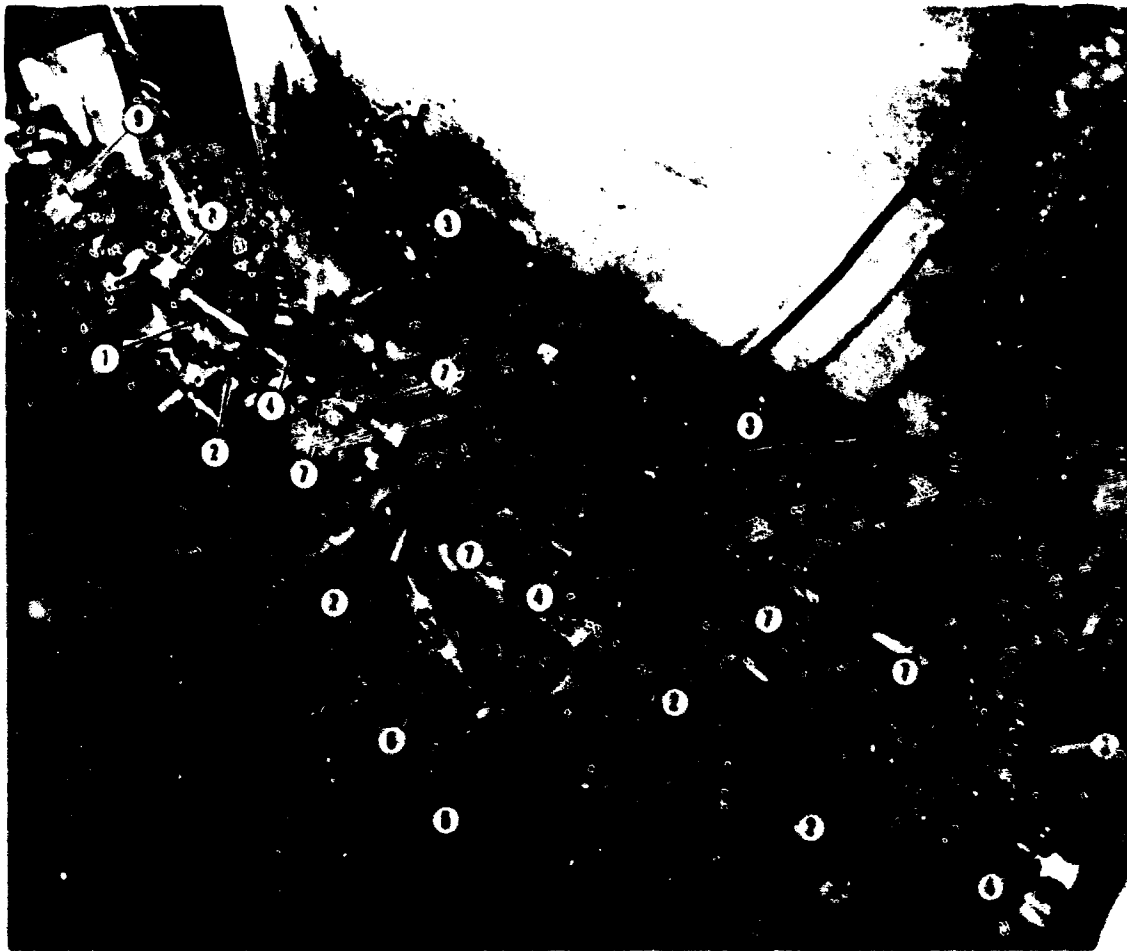


Pipe Length: 48"



**TEST PIPE  
FIGURE 2**

- |                         |  |
|-------------------------|--|
| 1) 1/2" MONEL PIPE      | 5) RELIEF VALVE                        |
| 2) FLAME PROBES         | 6) GAS SAMPLE BOMB                     |
| 3) TEMPERATURE PROBES   | 7) STRAIN GAUGES                       |
| 4) PRESSURE TRANSDUCERS | 8) MONEL UNION (FLAME ARRESTER INSIDE) |



**TEST PIPE**  
**FIGURE 3**

- |                         |                    |
|-------------------------|--------------------|
| 1) 1 1/2" MONEL PIPE    | 5) RELIEF VALVE    |
| 2) FLAME PROBES         | 6) GAS SAMPLE BOMB |
| 3) TEMPERATURE PROBES   | 7) STRAIN GAUGES   |
| 4) PRESSURE TRANSDUCERS | 8) MC NEIL UNION   |
|                         | 9) MAROTTA VALVE   |



OXYGEN SUPPLY PIPING  
FIGURE 4

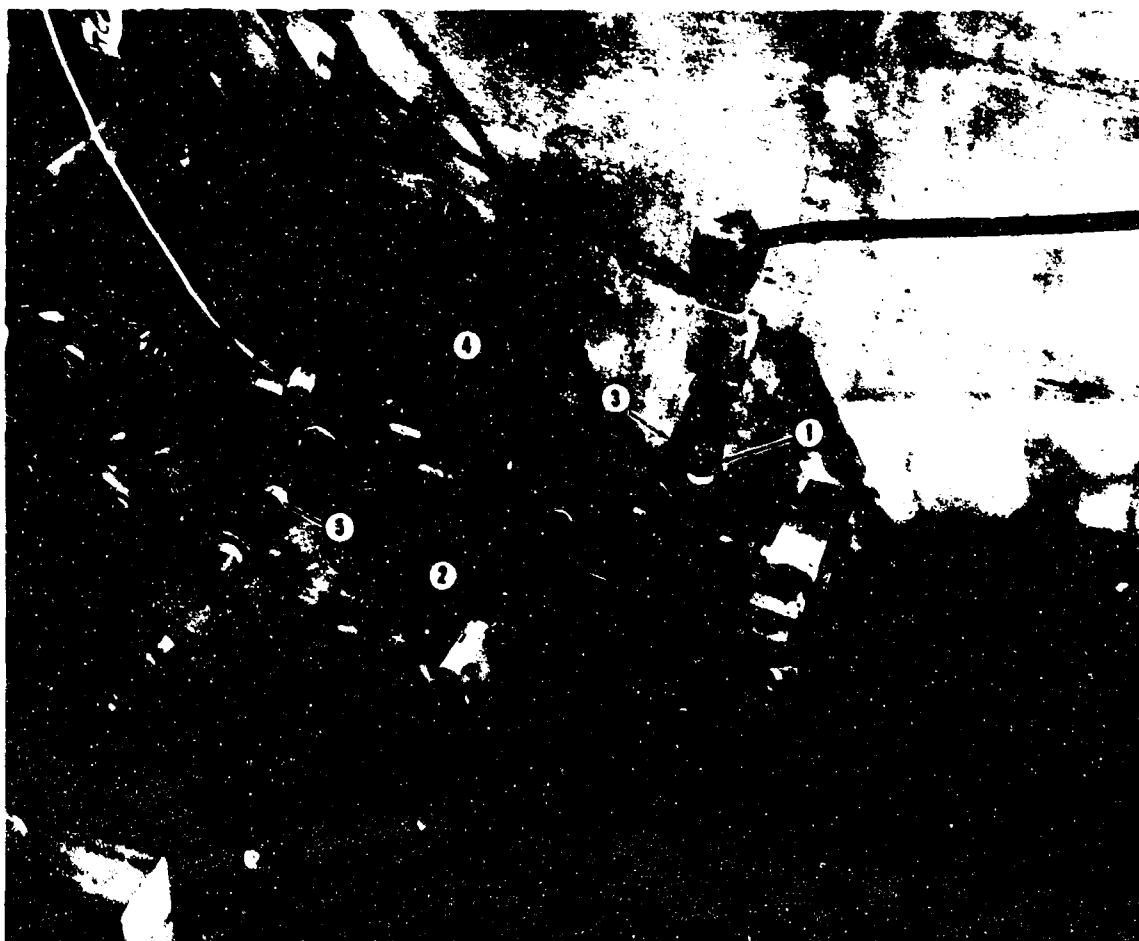
- 9) MAROTTA VALVE
- 10) CHECK VALVE
- 11) PRESSURE GAUGE
- 12) HAND OPERATED BALL VALVE



OXYGEN SUPPLY PIPING  
FIGURE 5

- 10) CHECK VALVE
- 11) PRESSURE GAUGE
- 12) HAND OPERATED BALL VALVE
- 13) 3000 PSI OXYGEN SUPPLY TANK



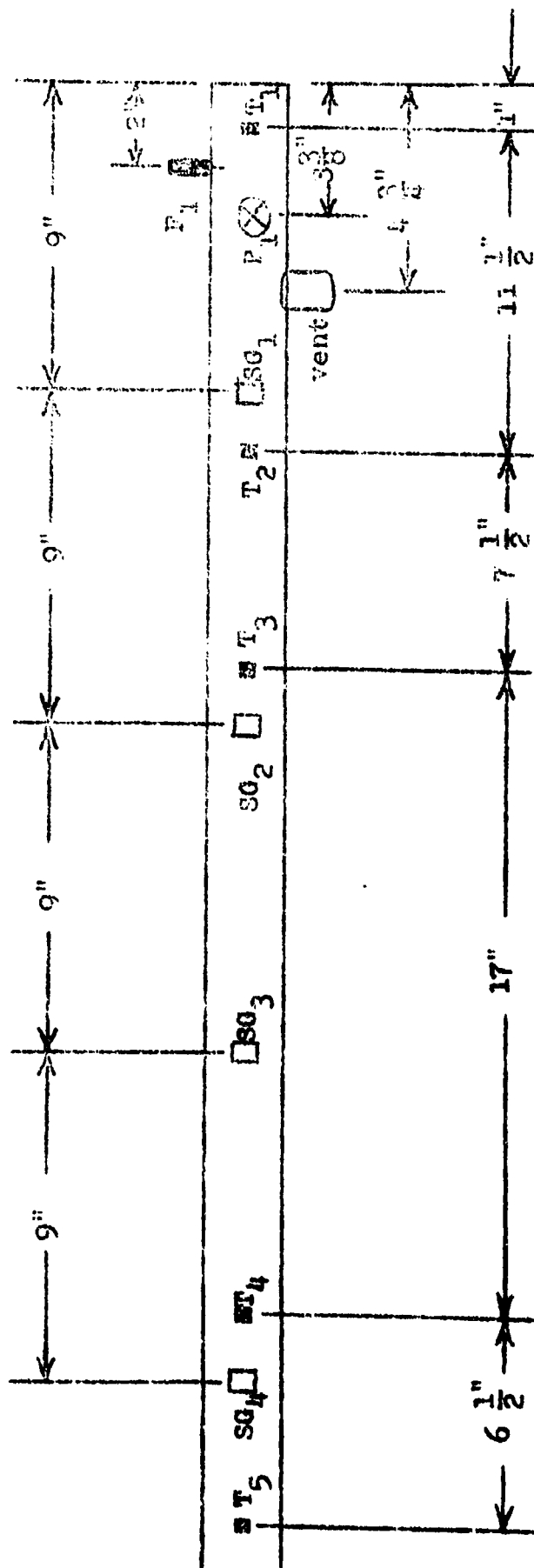


TEST PIPE WITH SPARKING DEVICE  
FIGURE 6

- 1) SPARKING DEVICE
- 2) FLAME PROBE
- 3) TEMPERATURE PROBE
- 4) STRAIN GAUGE
- 5) VENT LINE

FIGURE 7

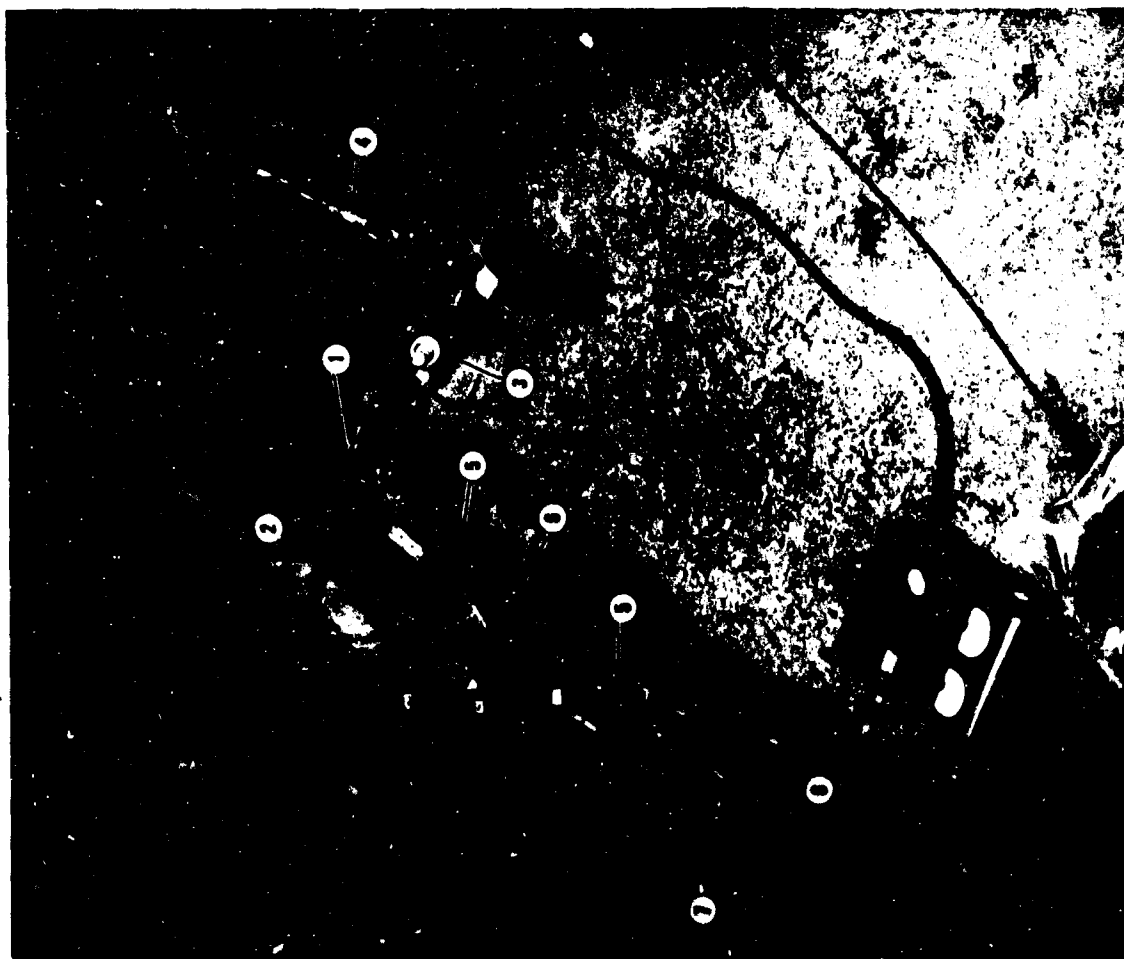
- T - Thermocouple
- F - Flame Probe
- P - Pressure Transducer
- SG - Strain Gauge



Pipe Length: 45 1/2"

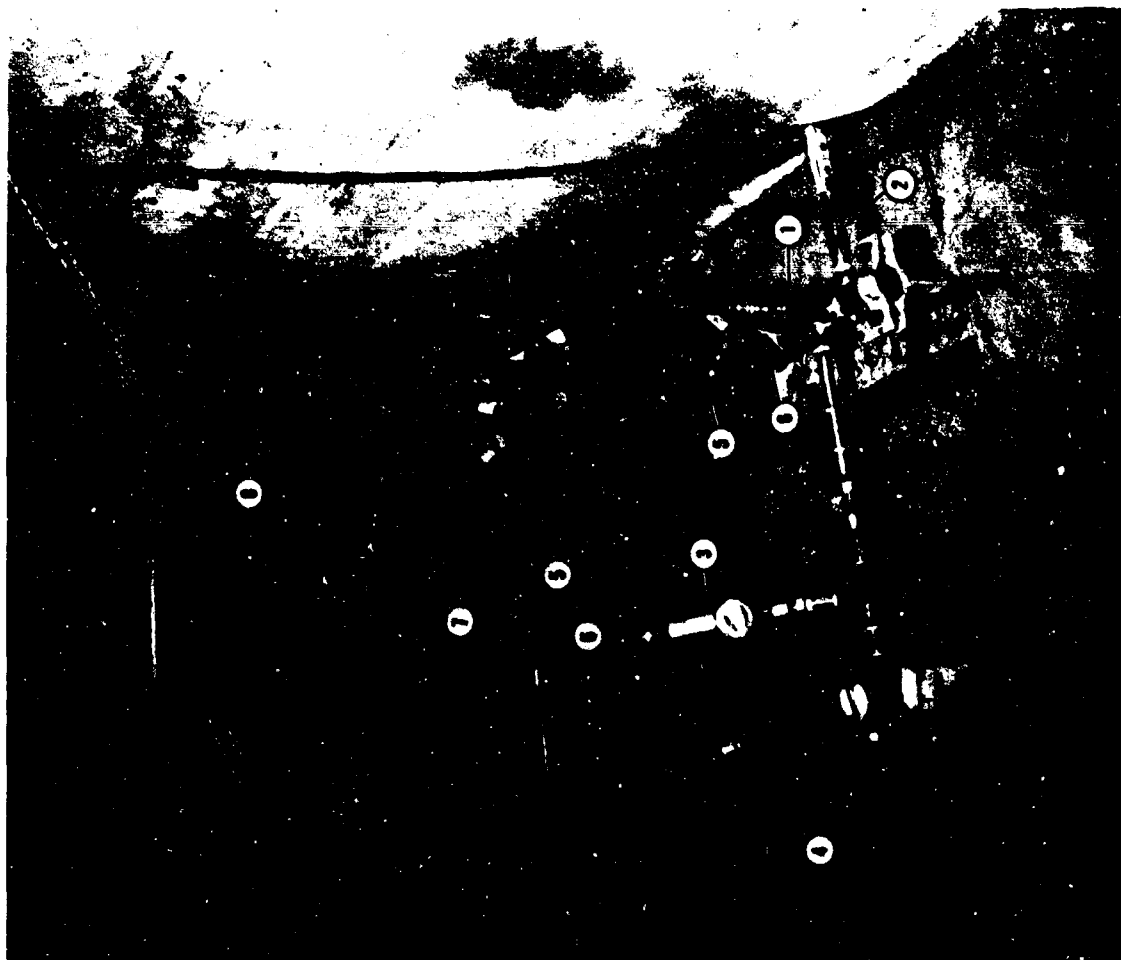
MODIFIED TEST PIPE  
FIGURE 8

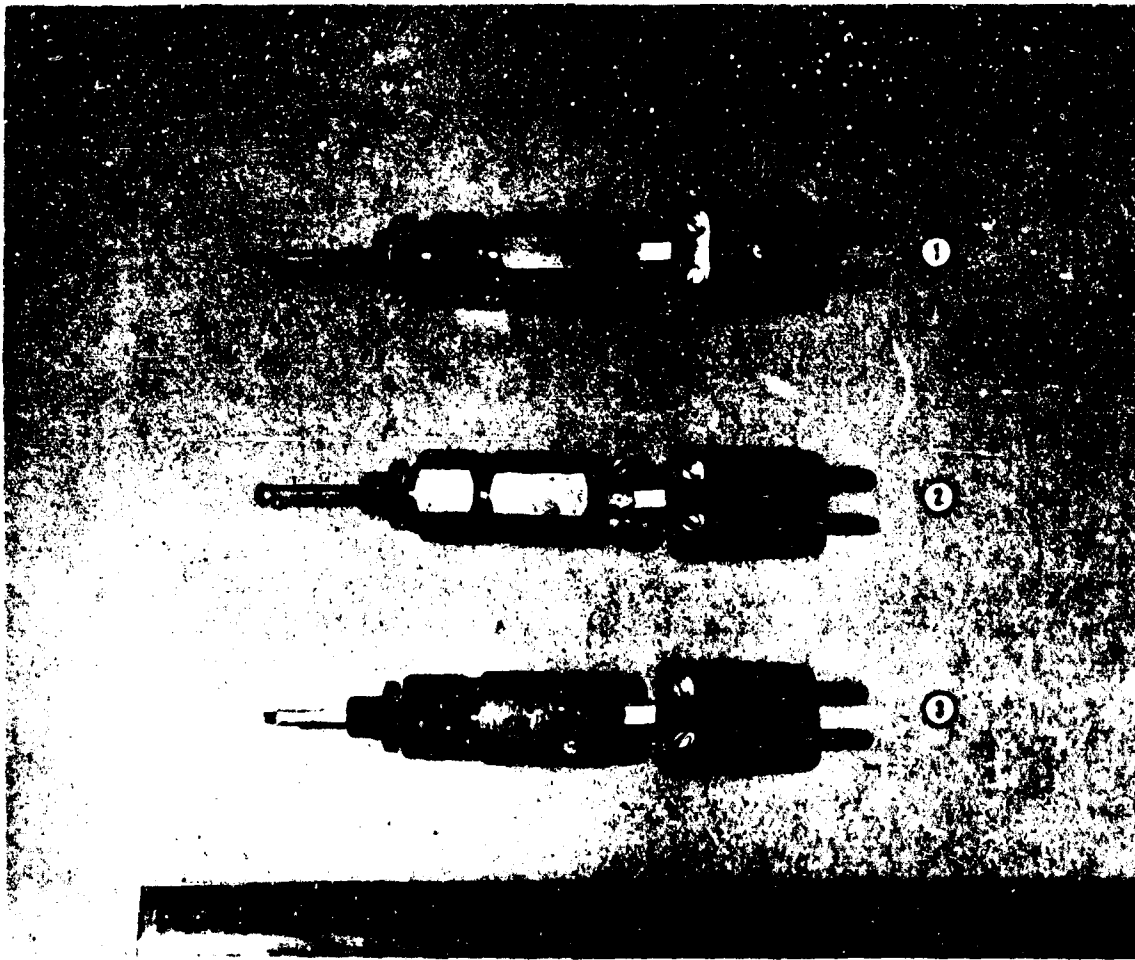
- 1) PRESSURE TRANSDUCER
- 2) FLAME PROBE
- 3) RELIEF VALVE
- 4) GAS SAMPLING BOMB
- 5) STRAIN GAUGES
- 6) HEATING WIRE
- 7) MONEL UNION



MODIFIED TEST PIPE  
FIGURE 9

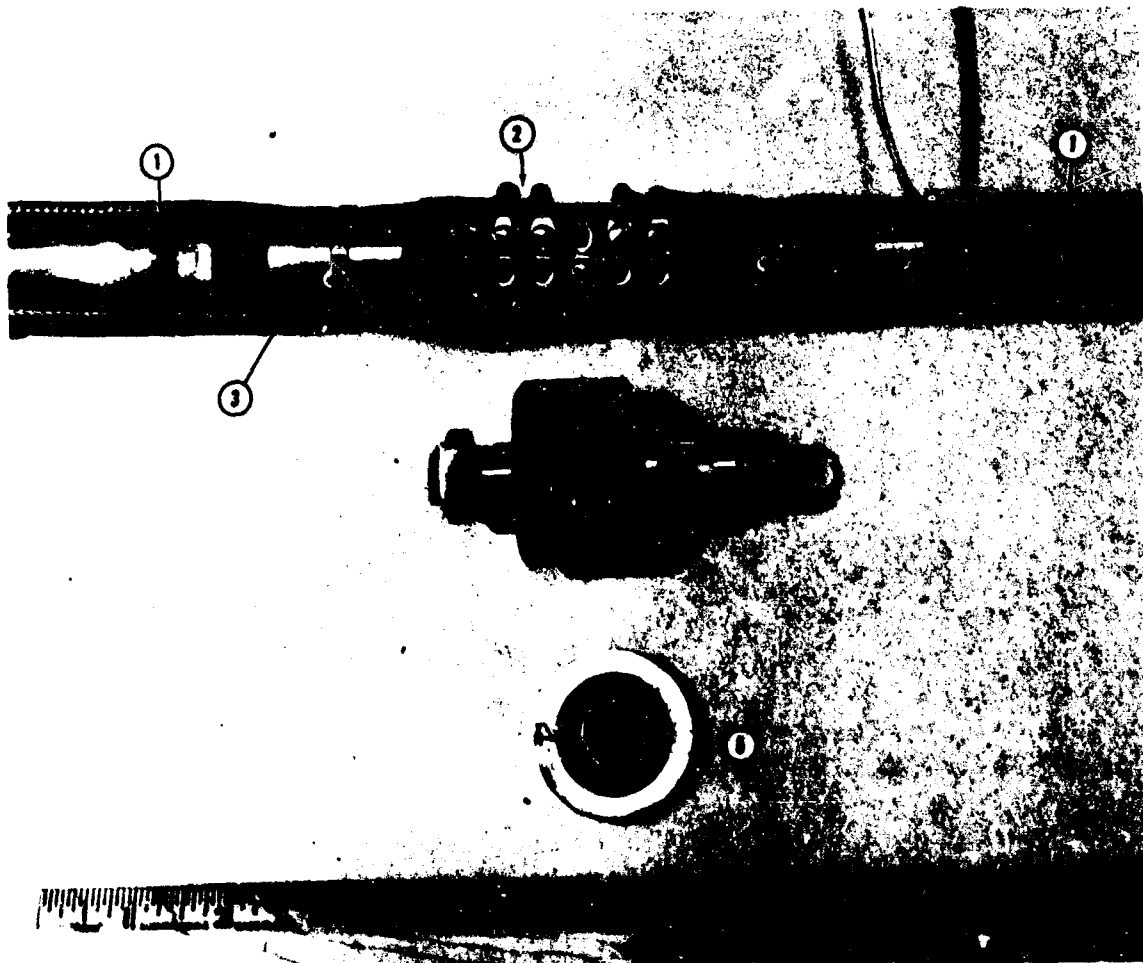
- 1) PRESSURE TRANSDUCER
- 2) FLAME PROBE
- 3) RELIEF VALVE
- 4) GAS SAMPLING BOMB
- 5) STRAIN GAUGES
- 6) HEATING WIRE
- 7) MONEL UNION
- 8) MAROTTA VALVE





INSTRUMENTATION PROBES  
FIGURE 10

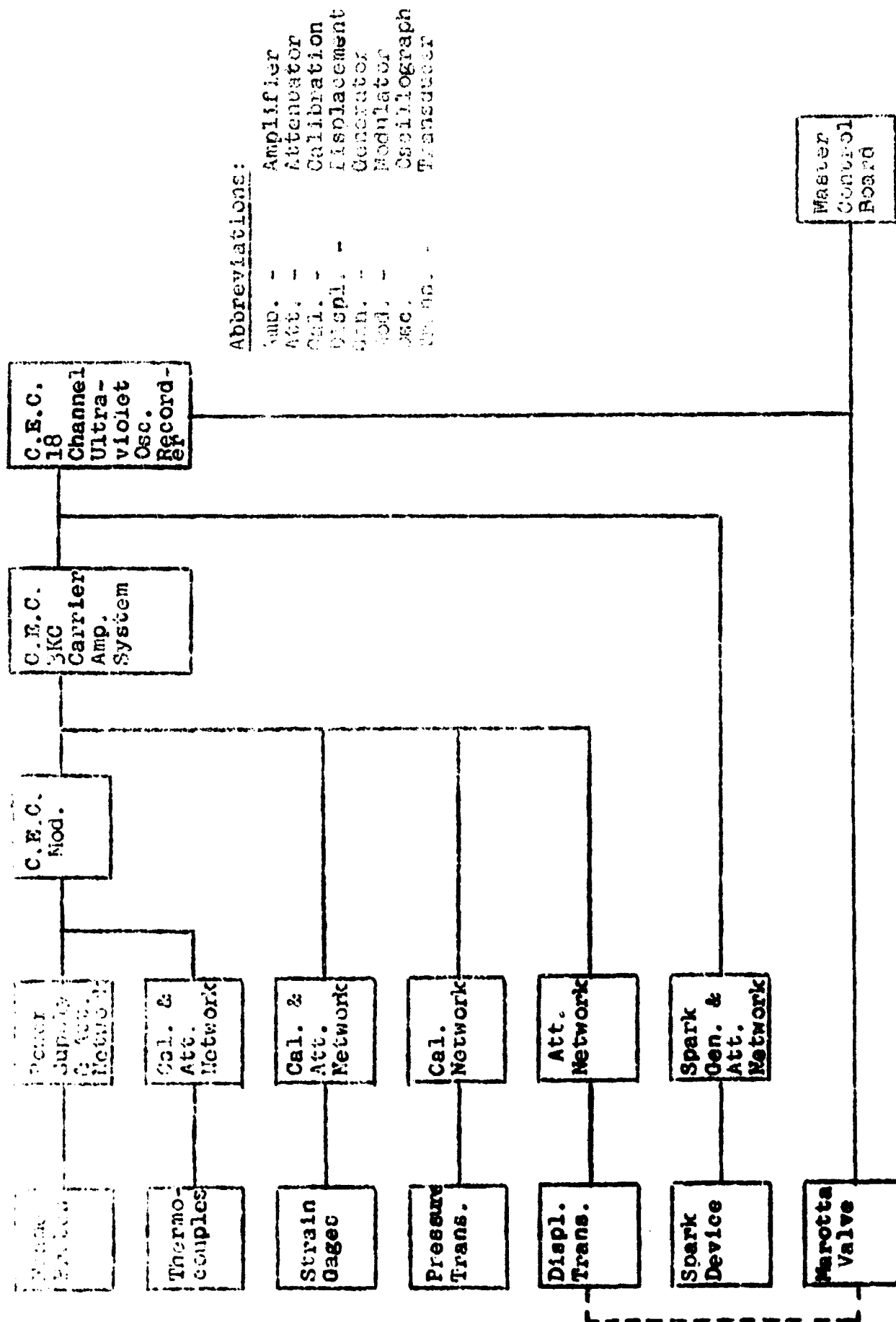
- 1) SPARKING DEVICE
- 2) THERMOCOUPLE
- 3) FLAME PROBE



**TEST PIPE COMPONENTS**  
**FIGURE 11**

- 1) STRAIN GAUGE
- 2) TERMINAL FOR STRAIN GAUGES
- 3) HEATING COIL
- 4) PRESSURE TRANSDUCER
- 5) FLAME ARRESTOR

OIL CONTAMINATION IN OXYGEN SYSTEMS  
INSTRUMENTATION BLOCK DIAGRAM - FIGURE 12



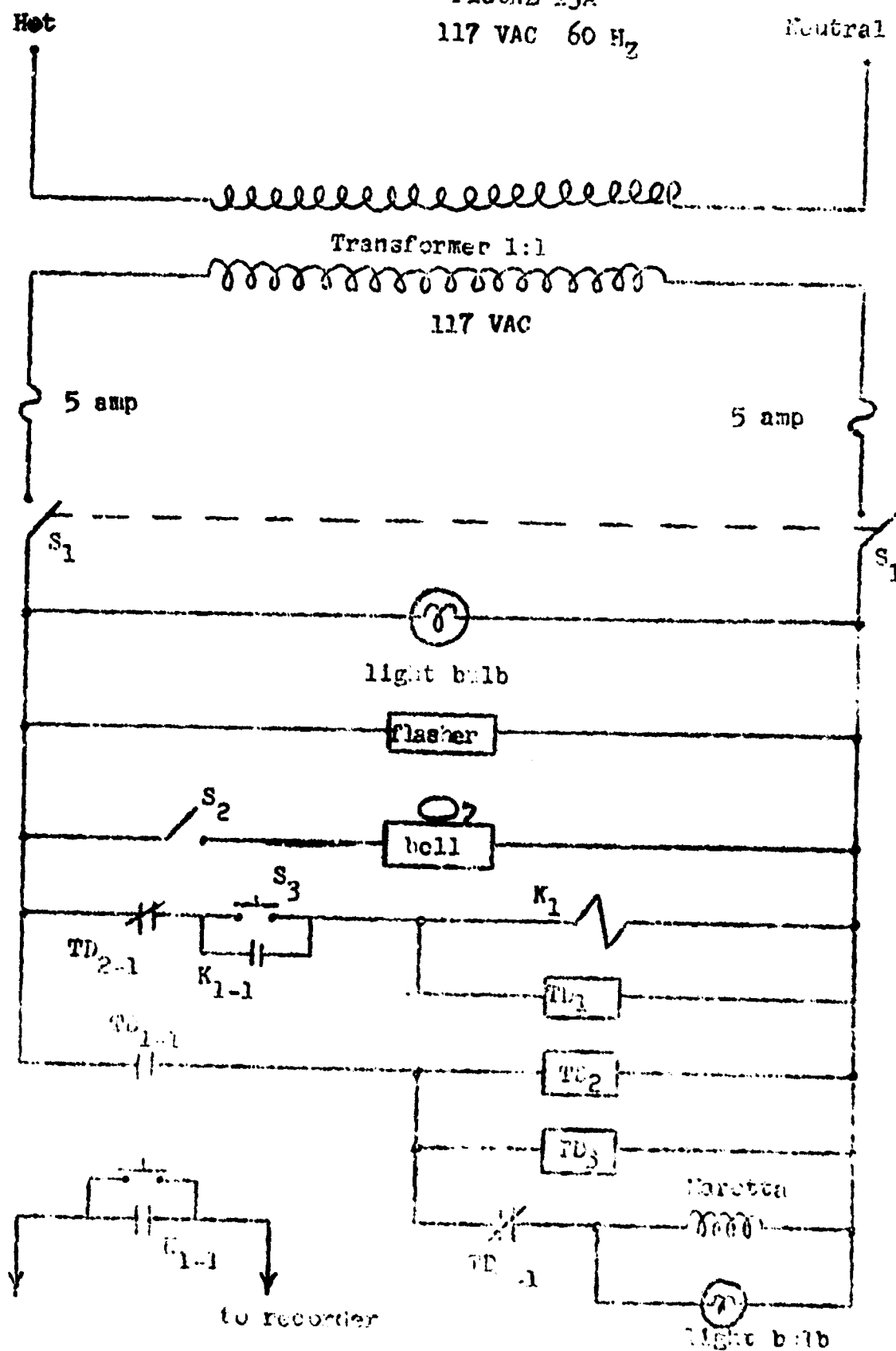
OIL CONTAMINATION IN OXYGEN SYSTEMS  
AUTOMATIC REMOTE CONTROL CIRCUIT

FIGURE 13A

117 VAC 60 Hz

Neutral

Ground

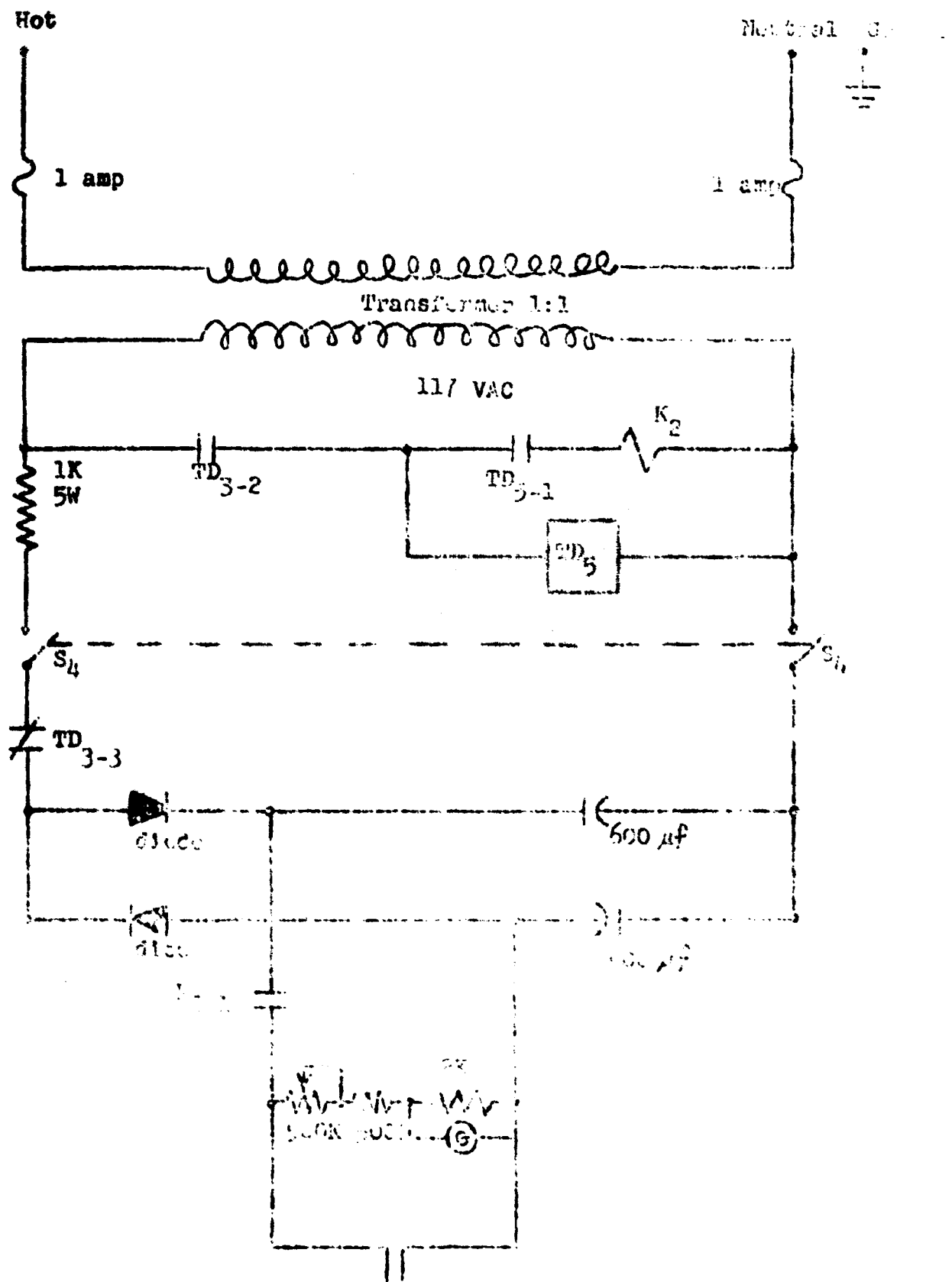




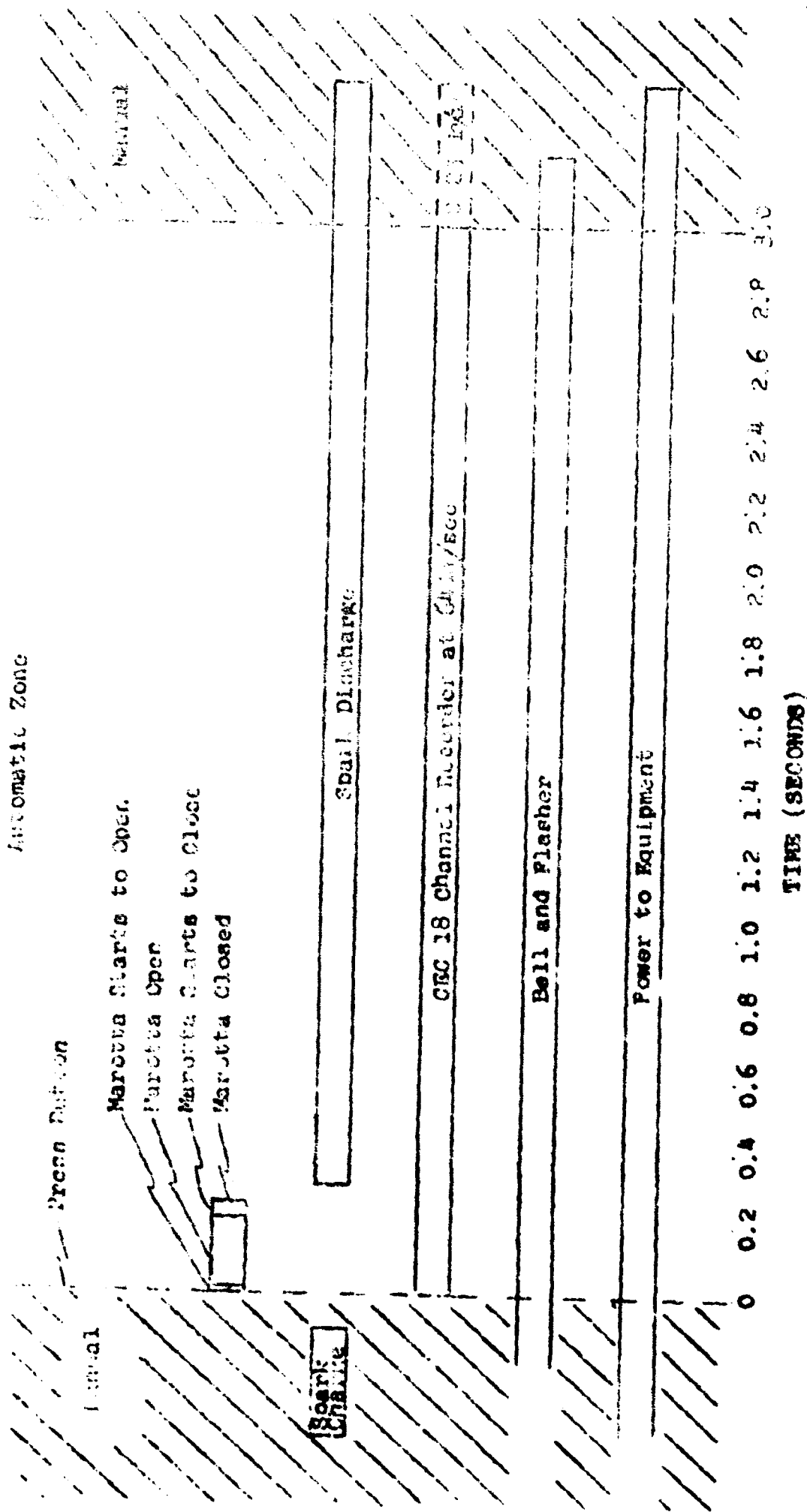
## SPARK GENERATOR CIRCUIT

**FIGURE 13B**

117 VAC 60 H<sub>Z</sub>



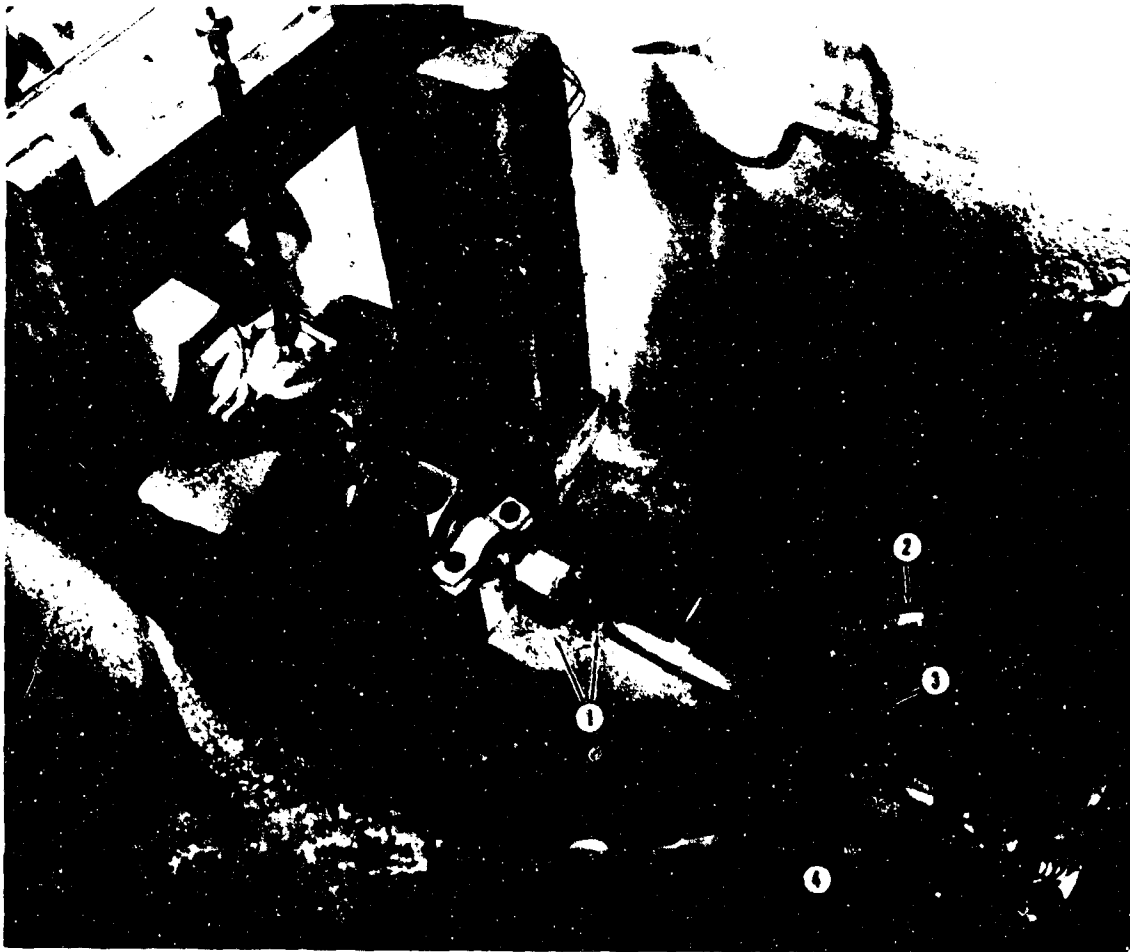
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TEST PIPE AFTER EXPLOSION  
FIGURE 15

- 1) BURNED UNION AND TEST PIPE FITTING
- 2) CHARRED, BENT FLAME PROBE
- 3) BENT THERMOCOUPLE CONNECTOR
- 4) MELTED, EMPTY PRESSURE TRANSDUCER BOSS
- 5) DISPLACED STRAIN GAUGE TERMINALS
- 6) DISPLACED COPPER O-RING ON PRESSURE TRANSDUCER
- 7) RELIEF VALVE



TEST PIPE AFTER EXPLOSION  
FIGURE 16

- 1) MELTED UNION AND TEST PIPE FITTING
- 2) BROKEN THERMOCOUPLE
- 3) MELTED, EMPTY PRESSURE BOSS
- 4) DISPLACED STRAIN GAUGE TERMINAL



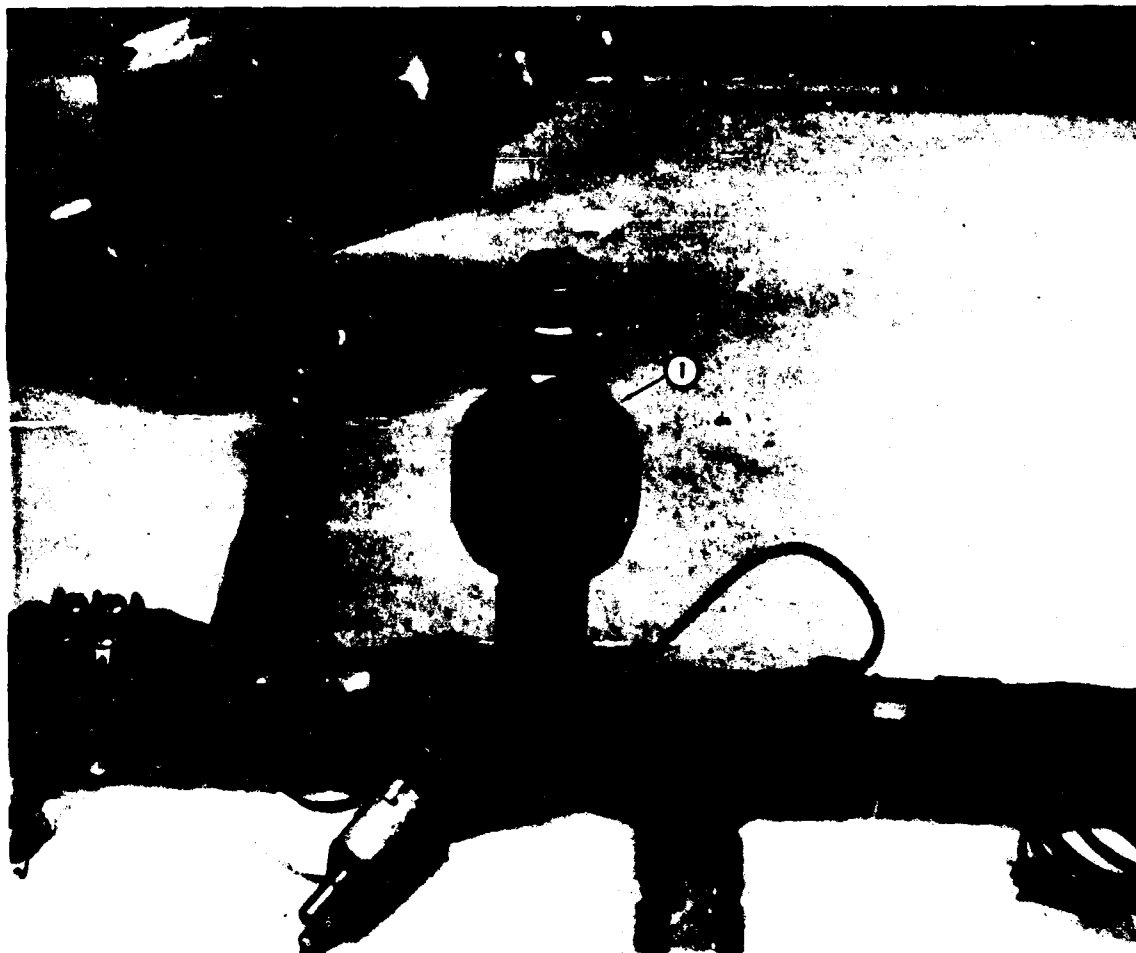
TEST PIPE AFTER EXPLOSION  
FIGURE 17

- 1) MELTED UNION
- 2) MELTED TEST PIPE FITTING FRAGMENTS
- 3) TEST PIPE



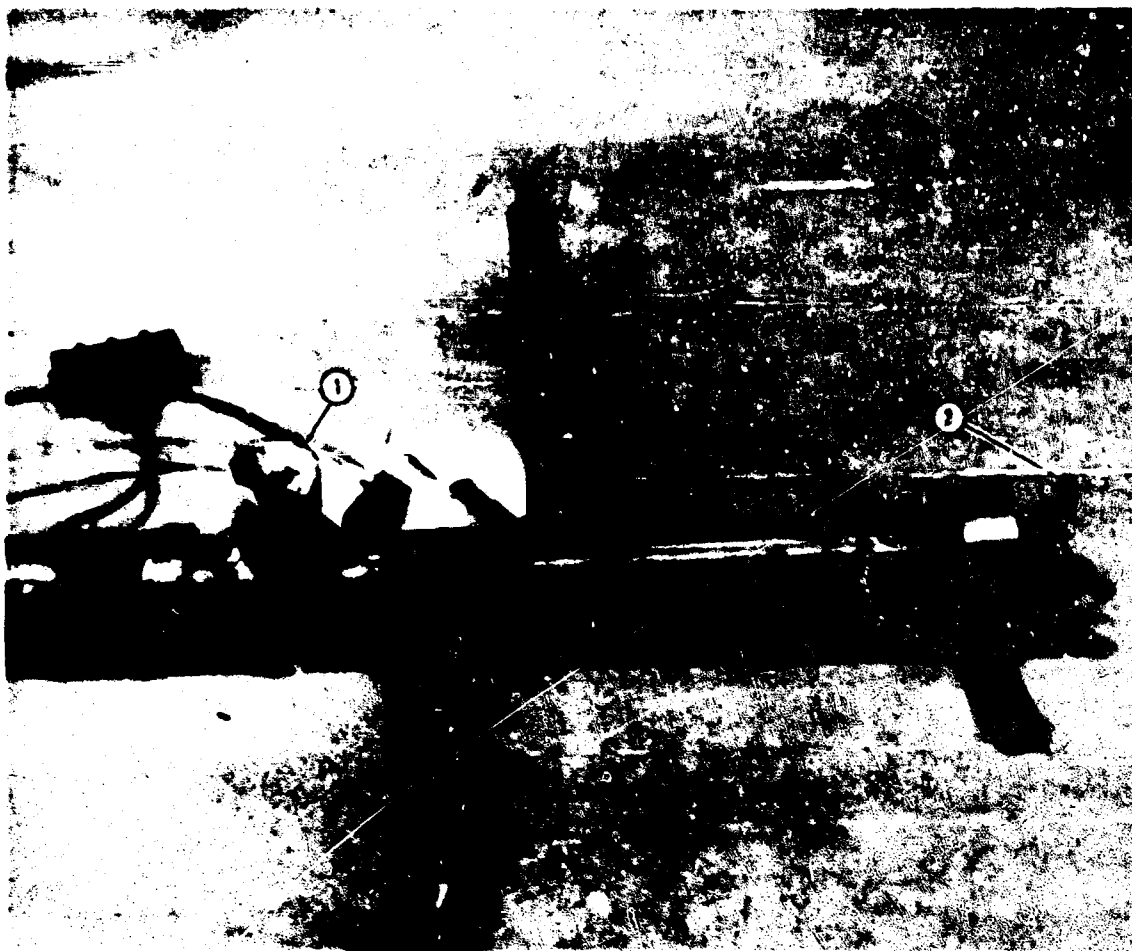
PRESSURE TRANSDUCER AFTER EXPLOSION  
FIGURE 18

1) PRESSURE TRANSDUCER FITTING



PRESSURE TRANSDUCER AFTER EXPLOSION  
FIGURE 19

1) DISPLACED, BENT COPPER O-RING ON PRESSURE TRANSDUCER



INSTRUMENTATION BOSS AFTER EXPLOSION  
FIGURE 20

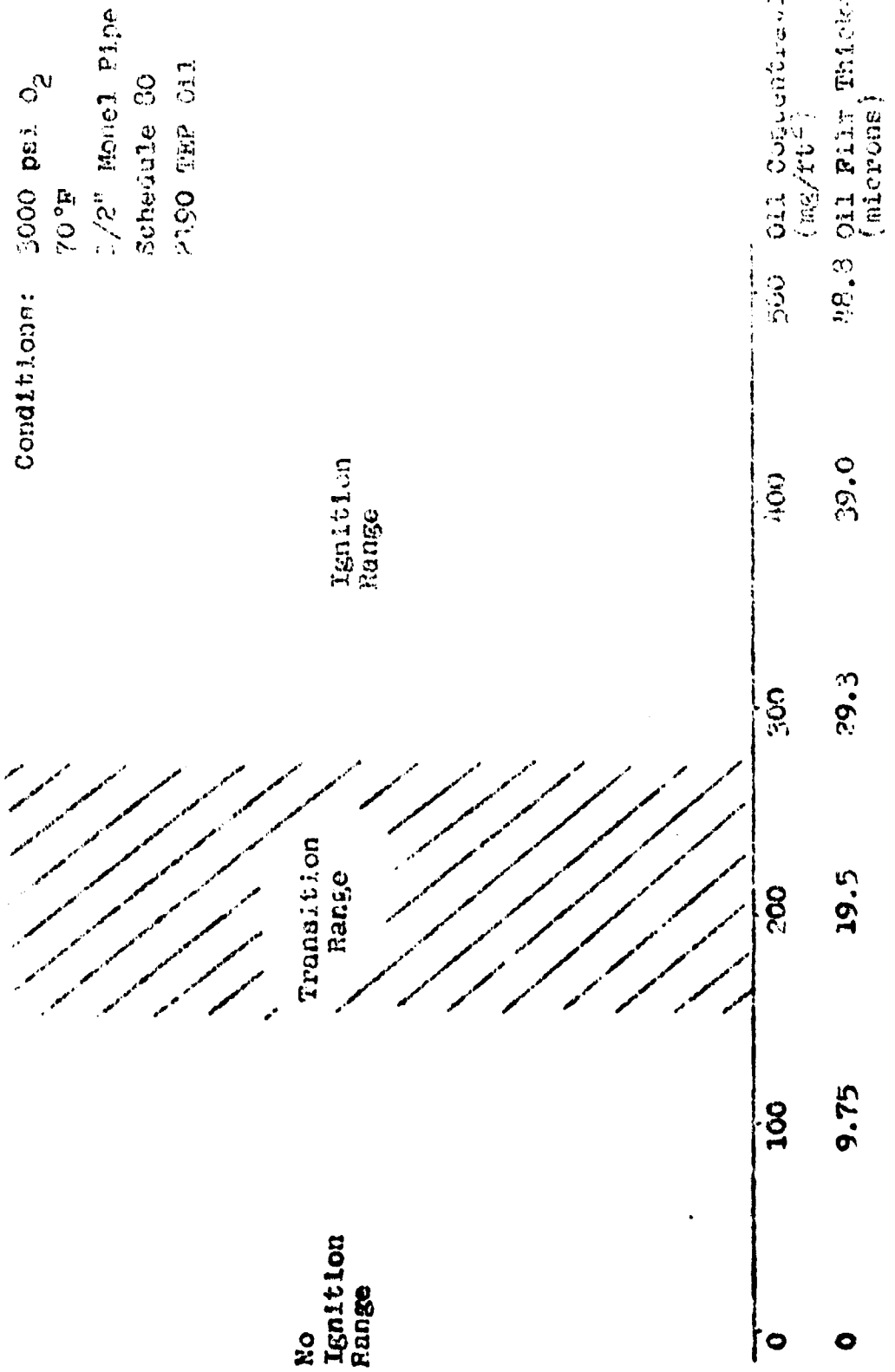
- 1) MELTED PRESSURE TRANSDUCER BOSS
- 2) MELTED TEST PIPE FITTING



# OIL CONTAMINATION IN OXYGEN SYSTEMS

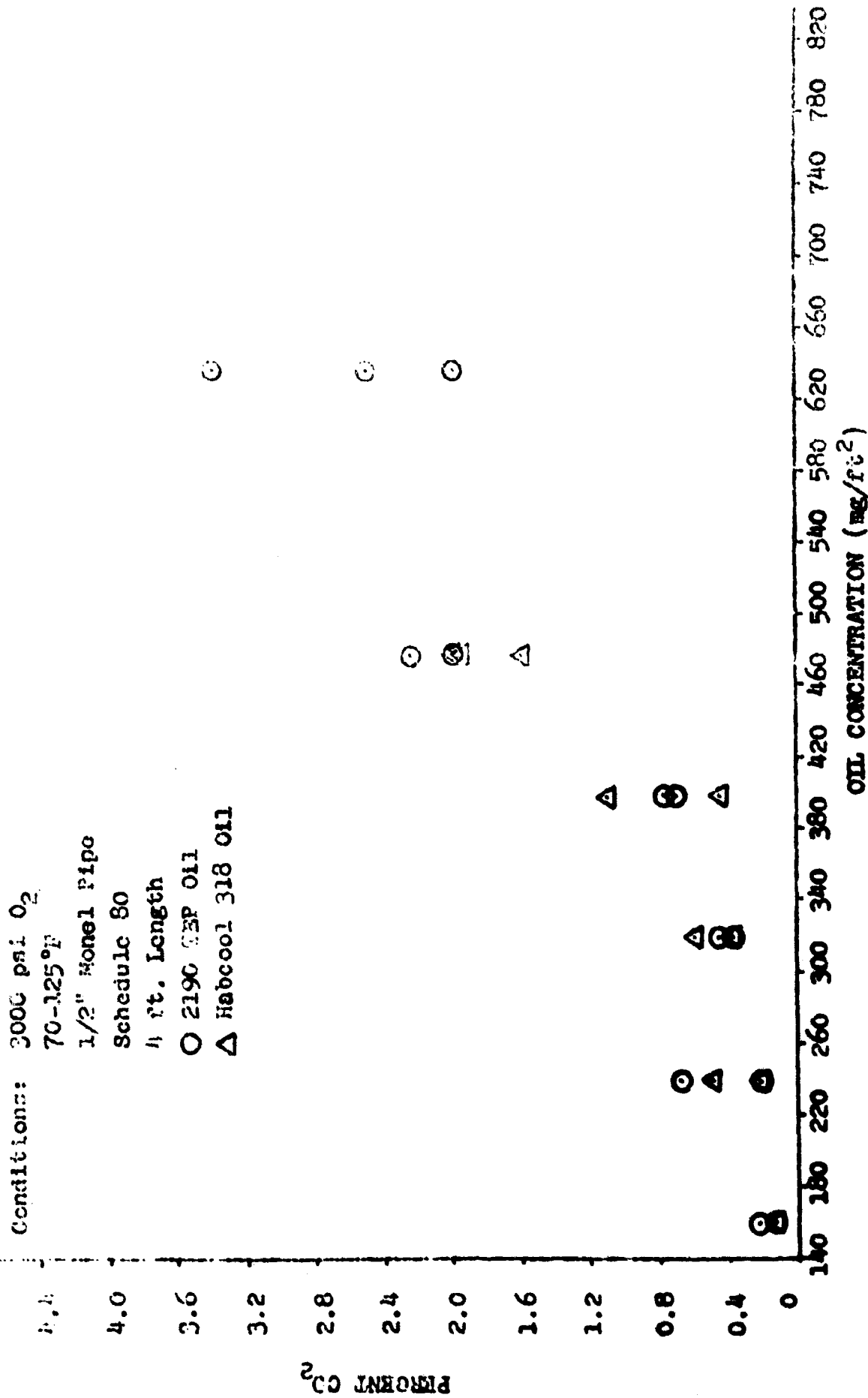
## IGNITION RANGES

FIGURE 21



PERCENT  $\text{CO}_2$  VS. OIL CONTAMINATION

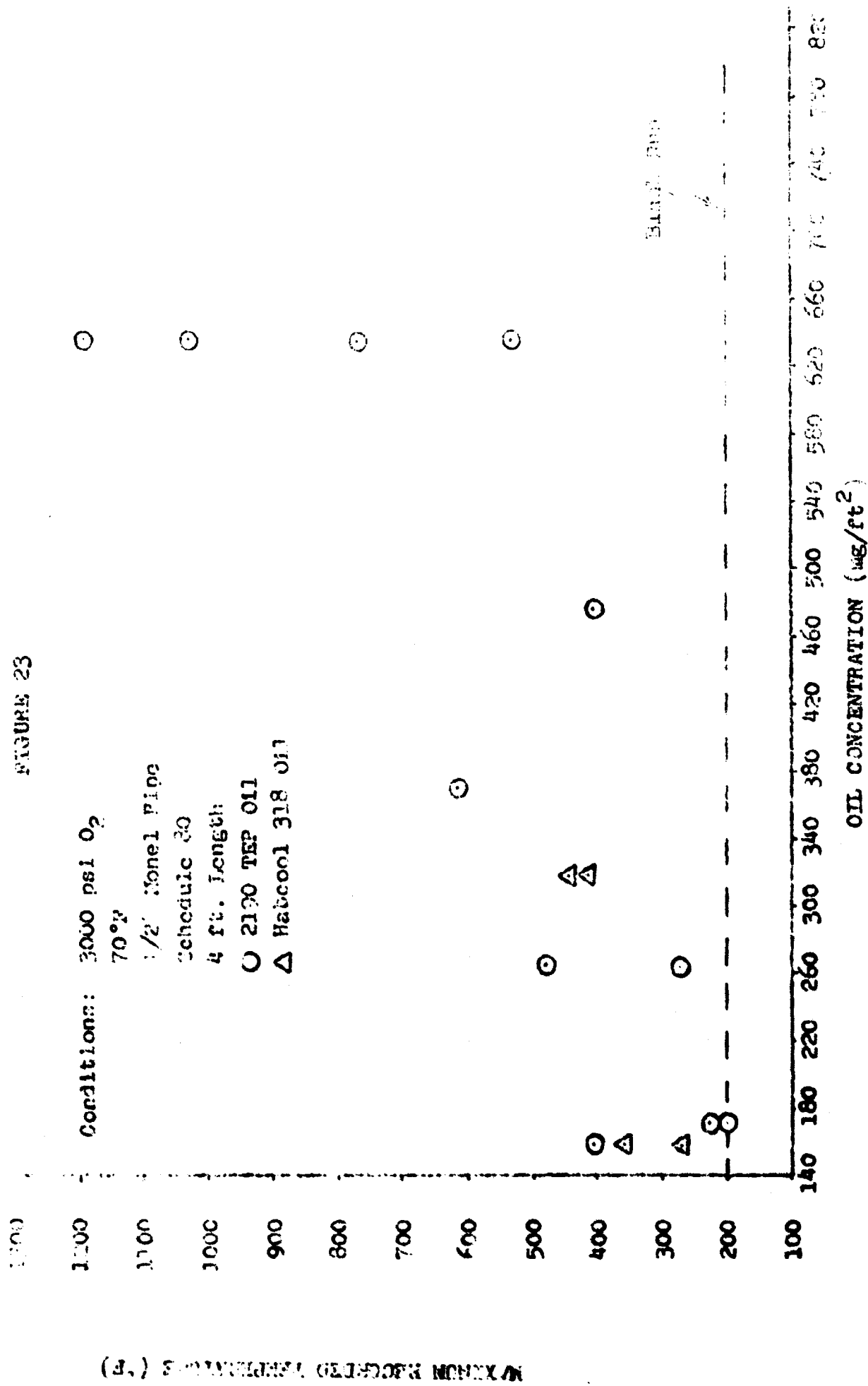
FIGURE 22



# OIL CONTAMINATION IN OXYGEN SYSTEMS

MAXIMUM PERMITTED TEMPERATURE VS. OIL CONCENTRATION

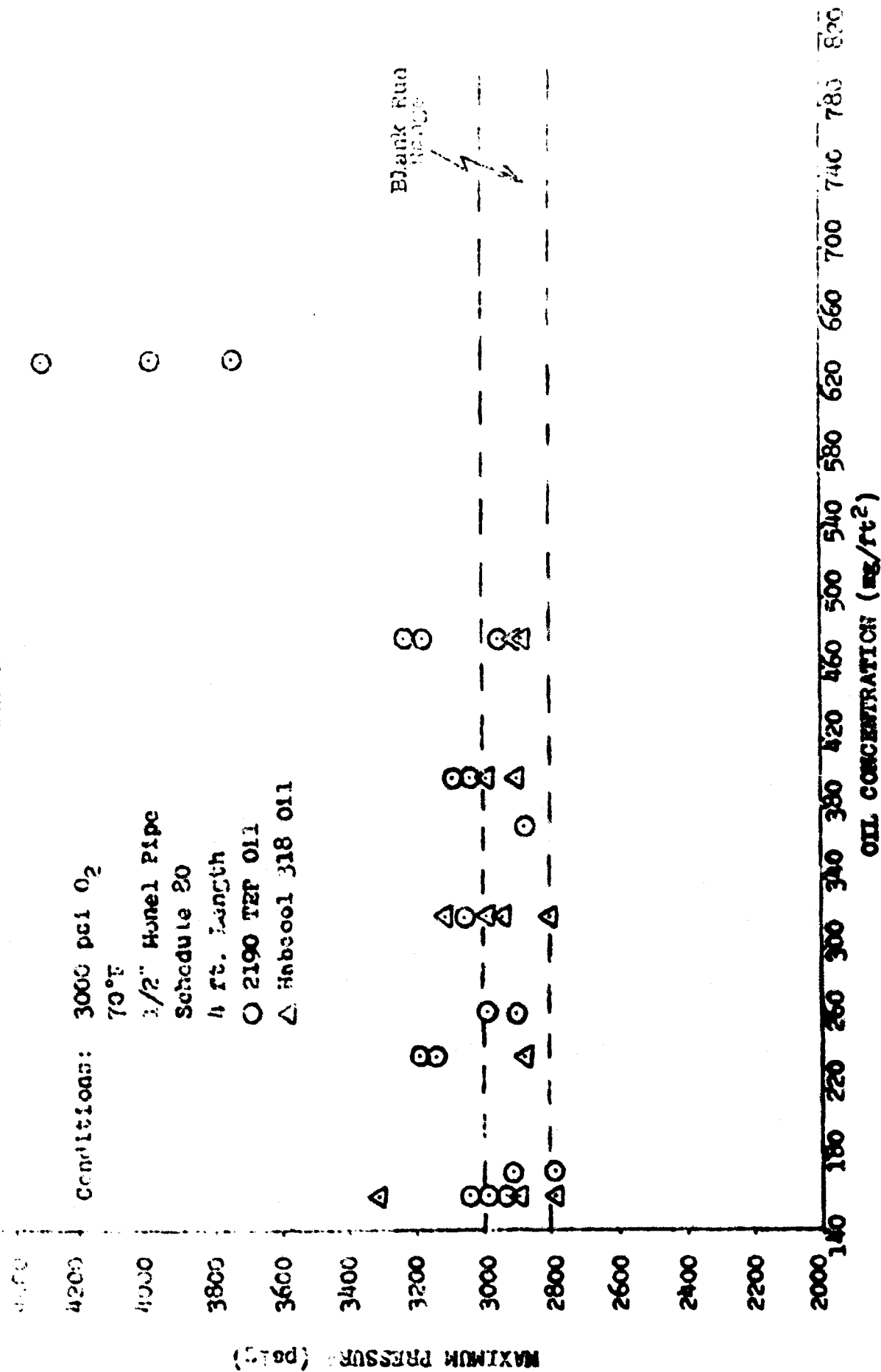
FIGURE 23



# Oil Oxidation in Oxygen Systems

Oil, 100% Sulfonated 100% Oil Concentration

FIGURE 24



UNCLASSIFIED

Security Classification

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4. DESCRIPTIVE NOTES (Type of report and inclusive dates) <b>Final Report (July 1967 - May 1971)</b>			
5. AUTHOR(S) (Last name, first name, initial) <b>Preotti, John R.</b> <b>DeLancey, Charles J., Jr.</b>			
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13. ABSTRACT <p>The ignition limits of 170 lubricating oil and Lubrol #31 cutting oil evenly coated on a 1/2" nominal pipe exposed to 3000 psi O<sub>2</sub> were determined. Little danger of ignition or flame propagation exists below oil concentrations of 160 mg/ft<sup>3</sup>; while concentrations above 160 mg/ft<sup>3</sup> will ignite in a pure O<sub>2</sub> atmosphere.</p>			

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