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FLAME INHIBITION RESEARCH

QUARTERLY PROGRESS REPORT

1 November 1961 - 31 January 1962

THIS RESEARCH PROGRAM ON FLAME PROCESSES AND CHEMICAL KINETICS IS DIRECTED TOWARD A SCIENTIFIC UNDERSTANDING OF THE BASIC MECHANISMS OF FLAME INHIBITION. IT IS SUPPORTED BY THE U.S. ARMY ENGINEER RESEARCH AND DEVELOPMENT LABORATORIES, FORT BELVOIR, VIRGINIA, THROUGH CONTRACT NO. d 7386 WITH THE BUREAU OF NAVAL WEAPONS, DEPARTMENT OF THE NAVY.

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QUARTERLY PROGRESS REPORT

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SUMMARY OF PROGRESS 1 November, 1961 - 31 January, 1962

- (1) To circumvent the corrosion problems encountered in previous metal burner systems a new spherical flame apparatus has been designed and constructed using exclusively glass, teflon and ceramic materials. The sampling probe has been modified to allow temperature measurements to be made by the pneumatic probe method thus allowing the direct association of temperature and composition measurements.
- (2) The problem of handling and precisely metering small amounts of corrosive materials has been solved by the construction of a new type of pressure transducer whose critical parts are constructed of teflon and glass.
- (3) The scavenger probe studies of atom concentrations have continued. A newly designed apparatus has been constructed and tested in studies of oxygen atom concentration in a spherical methane flame. The results were satisfactory in reproduceability. Quantitative comparison of these results with equilibrium calculations were frustrated by serious enthalpy losses through conduction to the flask walls.

TABLE OF CONTENTS

List of Figures	iv
List of Tables.	v
I. BACKGROUND MATERIAL	1
Report Organization	1
Program Objectives	1
Historical Summary	1
II. PROGRAM OUTLINE	2
Flame Structure Experimental Studies	3
Simple Reaction Studies	5
Data Analysis	5
Interpretation.	6
Recommendations on Flame Inhibition	7
III. SUMMARY OF PREVIOUS WORK	8
IV. REPORT OF PROGRESS (1 November, 1961 - 31 January, 1962).	11
Corrosion Resistant Burner System	11
Pressure Transducer	14
The Concentrations of Atomic and Free Radical Species in Flames	17

LIST OF FIGURES

<u>Figure</u>	<u>Page</u>
II-1 Flow of Information in the Program to Study Flame Inhibition	2
II-2 Over-All Program to Study Flame Inhibition	2
II-3 Characteristic Profiles for a Methane- Oxygen Flame	3
III-1 Effect of HBr on the Characteristic Profiles of a Stoichiometric Methane Flame	8
III-2 The Effect of HBr on Intermediate Species in a Stoichiometric Methane Flame	8
III-3 Effect of HBr on the Rate of Methane Disappearance in a Stoichiometric Flame	9
III-4 Diffusion Coefficients of H ₂ -Ar as a Function of Temperature	9
III-5a Schematic of Scavenger Probe	10
5b Effect of Added HBr on H Atom Concentration	10
5c Test of Oxygen Atom Scavenger Probe	10
III-6a Effect of Added HBr on Flame Radical Concentrations	10
6b Effect of Added HBr on Inhibitor Species Concentrations	10
IV-1 Schematic Diagram of Spherical Flame Inhibition Apparatus	12
IV-2a Schematic Diagram of Corrosion Resistant Critical Orifice Flowmeter	15
2b Picture of Corrosion Resistant Transducer	15
2c Drawing of Transducer Base Plate	16
IV-3a Oxygen Atom Concentration in a Spherical Methane Flame	18
3b Composition Profile of Major Species	18

LIST OF TABLES

<u>Table</u>		<u>Page</u>
II-1	Methane-Oxygen Flame Reaction Kinetic Constants	5
III-1	Test of Oxygen Atom Scavenger Probe	10

FLAME INHIBITION RESEARCH

I. BACKGROUND MATERIAL

REPORT ORGANIZATION

This report is divided into four sections: I. Background Material, II. Program Outline, III. Summary of the program to the beginning of the reported quarter, and IV. Quarterly Progress. Only the fourth section contains new material (although Section III is updated quarterly). It is hoped that the background material will aid the unfamiliar reader in following the program without extensive references. Those who are familiar with the work may find it profitable to turn directly to the fourth section (Page 11).

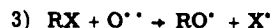
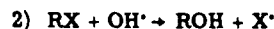
PROGRAM OBJECTIVE

The objective of this program is to obtain a scientific understanding of the mechanism of chemical inhibition of flames. Although this research is primarily restricted to the fundamental aspects of the problem, there are several potentially practical by-products. A quantitative understanding of flame inhibition should allow the prediction of the relative effectiveness of inhibitors on a flame system from their known chemical kinetics. Thus, a rational initial choice of inhibitor should be possible without extensive empirical studies. (This factor may become of increasing importance as "exotic" fuels become common.) It would also seem possible to set an absolute upper limit for the effect of an inhibitor, so that their efficiencies can be evaluated.

HISTORICAL SUMMARY

Over the past two decades, a number of studies have been made on flame inhibition of hydrocarbon-oxygen flames by halogen-containing compounds (Refs. 1 and 2). The experimental evidence suggests that the effectiveness of these materials is due to an effect on the chemical reactions in such flames. It has been established that the inhibition is a function of the concentration of the halogen, and that the order of effectiveness of the halogens is inverse to the reactivity of the free atoms, i.e., $F < Cl < Br < I$. Most practical extinguishers contain bromine compounds. Iodine compounds are usually unstable and expensive.

It is generally accepted that the mechanism for such flame inhibition is the exchange reactions between the inhibitor and the free radicals that are responsible for the primary propagation reactions in flames. Such reactions lead to the formation of stable molecules and free halogen atoms, which are much less reactive than the flame radicals. In hydrocarbon-oxygen flames the important radicals are hydrogen atoms, oxygen atoms, hydroxyl radicals, and possibly methyl radicals. Other radicals exist in flames, but these are probably less important to the flame propagation. On this hypothesis one would expect that the following reactions would be important in flame inhibition:



Here, R represents a hydrogen or hydrocarbon radical, and X represents a halogen. Dots indicate unpaired electrons. Reactions of this type are well known (Ref. 3), but much of the available data is too fragmentary to allow extrapolation of the kinetic information from the low-temperature regime of most laboratory studies to radical reactions at the high temperatures encountered in flames.

1. R. Friedman and J. R. Levy, "Survey of Fundamental Knowledge of Mechanisms by Action of Flame Extinguishing Agents," WADC Tech. Report 56-568, Jan. 1957, Supplement I, Sept. 1958, Supplement II, April 1959.
2. W. A. Rosser, H. Wise, and J. Miller, "Mechanism of Combustion Inhibition by Compounds Containing Halogen", Seventh Symposium (International) on Combustion, pp. 175-82, Butterworths Scientific Publications, London (1959).
3. E. W. R. Steacie, Atomic and Free Radical Reactions, Reinhold Publ. Co., New York (1954) (2 volumes).

FLAME INHIBITION RESEARCH

II. PROGRAM OUTLINE

The research program will utilize studies of the detailed microstructure of flames (Ref. 4) with and without added inhibitor, together with such other chemical and physical studies as seem necessary to elucidate the detailed mechanisms of chemical inhibition of flames. The relation between these studies can be visualized in Fig. II-1. Following this figure the report will be divided into five major sections (flame structure studies, simple reaction studies, analysis, interpretation, and recommendations of flame inhibition), and eight minor sections (temperature profiles, composition profiles, aerodynamic profiles, and atomic and free-radical composition profiles). This is a long-range program of modest scale whose completion is expected to take from three to five years. Therefore, although work is planned in all of these categories, current work will usually occupy only a few of these sections.

The number of possible inhibitors and the number of possible flame systems are both large, and the number of flame-inhibitor systems which may be of interest is even larger since it is the product of these two numbers. The situation can be represented by a very large "matrix" in which one index of the terms represents the flame system and the other represents the inhibitor system. (See Fig. II-2). Since it is impractical to study such a large number of systems, two possible approaches suggest themselves: The synthetic method, in which one would study the chemistry of inhibitors with radicals, and a separate study of the uninhibited flame systems. This is equivalent to studying the initial row and initial column in the "matrix". A satisfactory theory would allow the prediction of the effect of any inhibitor on any flame from this information. The other approach is the statistical technique, which is often used in agriculture. Studies would be made of a randomized set of these systems using one of the patterns of experiment design, such as the Latin square. By studying this fraction of the systems, a satisfactory theory should allow the derivation of information sufficient to predict the effect of any specific inhibitor on any specific flame system.

Both approaches assume that an adequate theory is developed. Therefore, a judicious combination of these two approaches seems in order. Early studies should be aimed at developing a theory, subsequent studies should be aimed at making a randomized statistical test of the efficacy of the theory, and final studies should be aimed at the accumulation of data. On this basis we propose to make a very detailed study

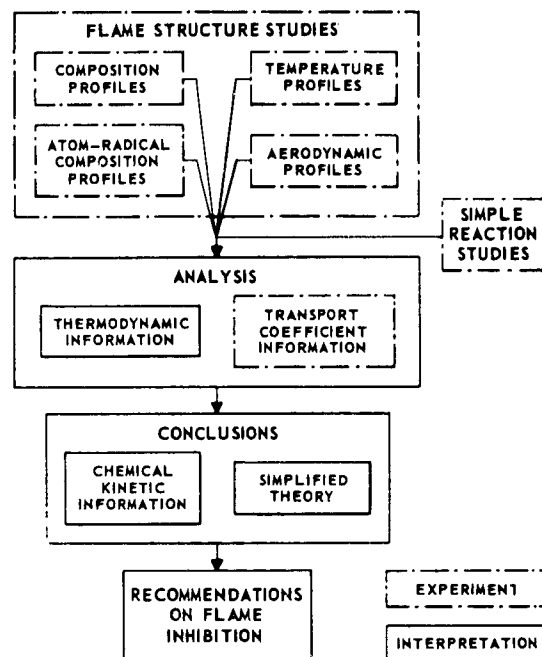


Figure II-1 FLOW OF INFORMATION IN THE PROGRAM TO STUDY FLAME INHIBITION

		FLAME SYSTEMS				
		H ₂ -O ₂	CO-O ₂	CH ₄ -O ₂	C ₂ H ₂ -O ₂	C ₂ H ₄ -O ₂
INHIBITOR-RADICAL SYSTEMS	HBr-DBr (H, O, OH)	TD	TD	TD	D	D
	Br ₂ (H, O, OH)	TD	TD	TD		
	HI (H, O, OH)	TD	TD	TD		
	HCl (H, O, OH)	TD	TD	TD		
	CF ₃ Cl (H, O, OH)	TD	TD	TD		
	CF ₃ Br (H, O, OH)	TD	TD	TD		
	CB ₄	TD	TD	TD		

TEST OF THEORY
INITIAL STUDY
DATA FOR APPLICATION

SYSTEM STUDY
SIMILAR TO FIG. 1

Initial Study, I - An intensive study of one inhibitor-flame system is directed toward the goal of a simplified theory of flame inhibition. The chosen system is methane-oxygen with HBr (and/or DBr) as an inhibitor.

Test of Theory, T - The theory from the initial study should then be tested on representative inhibitor-flame systems to confirm its generality.

Data for Application, D - If the theory seems satisfactory, then a store of data should be developed to permit its broad application. Such data is most readily obtained from simple systems—the flame systems alone and inhibitor-radical reaction systems. The required data include those of chemical kinetics, transport coefficients, and thermodynamic properties.

Figure II-2 OVER-ALL PROGRAM TO STUDY FLAME INHIBITION

4. R. M. Fristrom, and W. G. Berl, "Program for the Study of Flame Inhibition", proposal submitted by the Johns Hopkins University Applied Physics Laboratory, in response to USAEC Request No. 60-286, October 1960.

of the methane-oxygen flame with a simple inhibitor (probably HBr) to develop the theory. This will be followed by a randomized test on a restricted number of systems, and finally by a systematic study of parameters. The program is illustrated by Fig. II-2. It is discussed in more detail in Ref. 4.

FLAME STRUCTURE EXPERIMENTAL STUDIES

It is planned to use flame structure techniques in this research, on flames both with and without inhibitors. These techniques have been discussed in the literature (Refs. 5 and 6) and offer certain advantages for studying chemical reactions: 1) flames offer a steady-state for study, 2) there is no wall problem, 3) there are no mixing problems for studying fast reactions. The disadvantages are: 1) data analysis is complex and, 2) high precision is required of the data for obtaining kinetic information. These techniques nevertheless offer an excellent method for studying fast, high-temperature reactions and, in particular, offer the ideal technique for studying flame inhibition reactions in situ.

To characterize a flame system, it is necessary to specify at least $N + 1$ variables, where N is the number of molecular species (Ref. 5). The independent variable usually chosen is distance through the flame front, while the dependent variables are composition, temperature, velocity, or area ratio. Experimentally, it is common to classify the profiles in four categories according to the techniques required to measure them. These are: composition profiles, atomic and free-radical composition profiles, temperature profiles, and aerodynamic profiles. An example of such a set is given in Fig. II-3.

Composition Profiles

From the chemical standpoint, the most important information is the concentration profiles. The most satisfactory technique developed for making such studies is the microprobe sampling technique, which is followed by mass spectral analysis. The technique consists of withdrawing a sample of gas from the flame using a small, tapered quartz probe. The probe can be made small enough so that its effect on the flame is negligible, and the low pressure and rapid pressure drop in such a tapered probe quenches the sample so that reproducible meaningful samples can be obtained. Radical concentrations are not obtained by these techniques and are discussed in the next section. This technique is discussed in detail in the literature (Ref. 7).

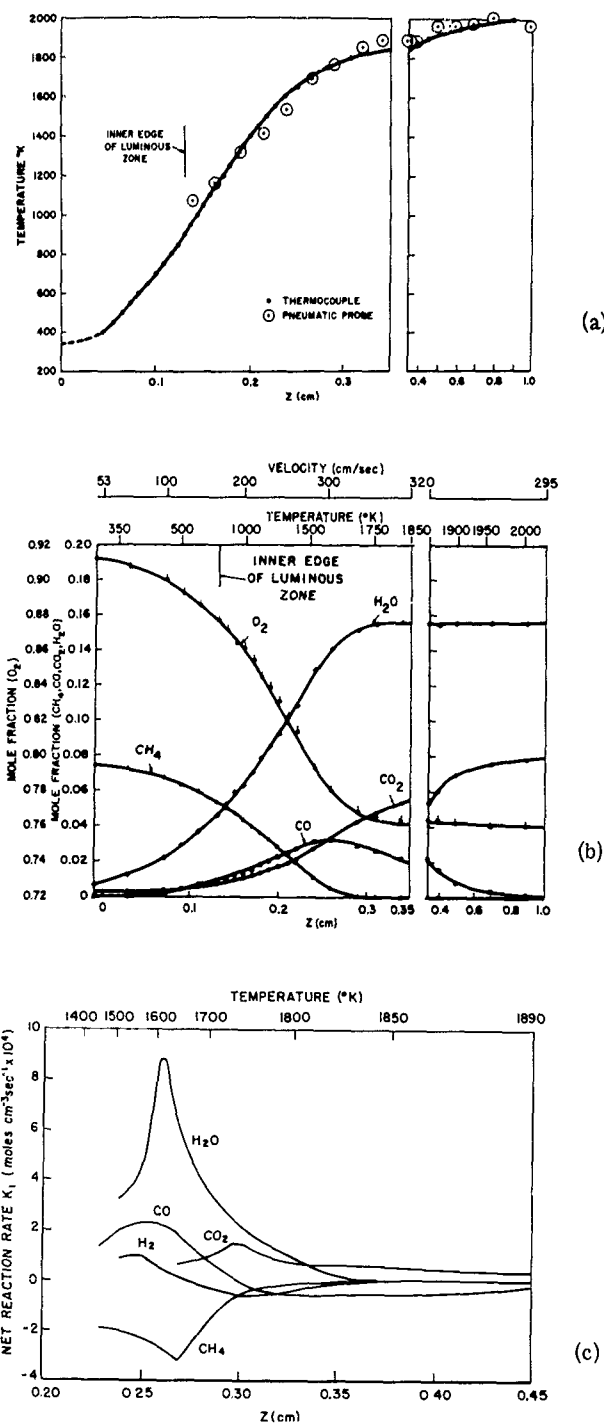


Figure II-3 CHARACTERISTIC PROFILES FOR A METHANE-OXYGEN FLAME

5. R. M. Fristrom, "Structure of Laminar Flames", Sixth Symposium (International on Combustion, pp. 96-110, Reinhold Publ. Co., New York (1957)).
6. R. M. Fristrom, C. Grunfelder, and S. Favon, "Methane-Oxygen Flame Structure I - Characteristic Profiles in a Low-Pressure, Laminar, Lean, Premixed Methane-Oxygen Flame", J. Phys. Chem. 64, 1386-92 (1960).

Radical-Atom Concentration Profiles

Logically, atom-radical concentration flame studies are part of the concentration profile determinations. However, since their study requires special techniques, they are considered separately.

Radicals and atoms are not susceptible to conventional sampling techniques because of their extreme reactivity which transforms them into stable species before analysis is possible. In most flame structure studies (Refs. 5 and 7) these concentrations are considered (usually correctly) to be negligible compared with the concentrations of the stable species, and are therefore neglected in the analysis. This reasonable approximation is, of course, only a stopgap measure to obtain information while satisfactory methods are developed for determining atom and radical concentrations. None of the techniques thus far developed (Refs. 8 through 10) possess the generality and spatial resolution necessary for the quantitative interpretation which we wish to make. Therefore, we are developing a new technique for studying radical concentration profiles under a companion flame structure study supported by ARPA (Task R). This is the scavenger probe method, which combines the techniques of probe sampling with those of chemical scavenging. A sample is taken through a probe, and the radical-containing gases are mixed rapidly with a large excess of a suitable "scavenger" gas which reacts with the radical to produce a characteristic product. Scavenger studies are well known in radiochemistry and photochemistry (Ref. 11), and mass spectrometer studies (Ref. 12) have indicated that radicals can be sampled successfully through orifices.

An absolute measurement of concentration can be made if 1) there is a one-to-one correspondence between the reaction product and the precursor radical and, 2) the analytical system (mass spectrometer) can be calibrated for the stable reaction product. This is a significant advantage since it is notoriously difficult to obtain absolute radical concentrations. Spatial resolution will be limited by the probe diameter, and sensitivity should be as high as the analytical system allows for stable species. As this technique is developed it will be used in these studies.

Temperature Profiles

Temperature profiles offer an important technique for characterizing flame fronts. Three methods have been used at this Laboratory: thermocouple traverses, aerodynamic measurements, and pneumatic probe measurements. Their data are in essential agreement (Ref. 6), although of varying precision.

The most precise method is that with thermocouple traverses in which temperatures are measured with fine (0.0005") Pt-Pt 10% Rh thermocouples coated with silica. Profiles with a precision of 10°K and excellent spatial reproducibility are obtained.

Temperature can be determined by using aerodynamic measurements of velocity and area ratio, deriving the point-by-point density through the continuity relations, and then applying the perfect gas law. The results are not of high precision (2-4%), but within their limits, they provide a reliable measure of the translational temperature of the gas.

Temperatures can also be measured with a pneumatic probe (Ref. 13). This device is a quartz probe similar to those used in the composition studies. If gas flows under choking conditions through two orifices in series, and the temperature is known at the second orifice, temperature at the inlet orifice of such a probe can be derived from the relation between ambient pressure and the pressure between the two orifices. This technique provides a moderately precise (1-2%) and reliable method for determining temperature and has the added advantage that it can be directly associated with a composition, since it can also be used for composition sampling.

Aerodynamic Profiles

Flames can be characterized aerodynamically by two profiles - velocity and area ratio. As mentioned under temperature measurements, these can be combined to derive a density profile. The area ratio is simply the geometry of the stream tubes of flow through the flame front. In the general case it is necessary to make this measurement, since stream tube expansion through

7. R. M. Fristrom, "Experimental Determination of Local Concentrations in Flames", APL/JHU CM-978.
8. C. G. James and T. M. Sugden, "Photometric Investigations of Alkali Metals in Hydrogen Flame Gases I - A General Survey of the Use of Resonance Radiation in the Measurement of Atomic Concentration", *Proc. Roy. Soc. A*-227, p. 312 (1955).
9. W. E. Kaskan, "Hydroxyl Concentrations in Rich Hydrogen-Air Flames Held on Porous Burners", *Comb. and Flame* 2, 229-43 (1958).
10. T. Grever and H. G. Wagner, "Die Reaktionszone von Flammen", *Zeit. fur Phys. Chem. (Neue Folge)* 20, 371-74 (1959).
11. R. H. Schuler, "Scavenger Methods for Free Radical Detection in Hydrocarbon Radiolysis", *J. Chem. Phys.* 62, 37-41 (1958).
12. S. Foner and R. Hudson, "The Detection of Atoms and Free Radicals in Flames by Mass Spectrometric Techniques", *J. Chem. Phys.* 21, 1374-82 (1953).

the flame is appreciable. With a symmetrical flame, such as the spherical flame, it is unnecessary to make this measurement, since it can be obtained directly from the geometry of the position determinations. Direct aerodynamic measurements are made by introducing MgO particles as a tracer and photographically visualizing their paths through the flame front, using a Zr flashbulb for the stream tube measurements and a pulsed electronic flashlamp for the direct velocity measurements (Ref. 14).

SIMPLE REACTION STUDIES

Flame systems have relatively complex chemistry (see Table I) which makes the identification of the dominant reactions difficult. For this reason it is desirable to supplement the flame studies with studies of the individual reactions with simplified chemistry. A second reason for doing so is that these studies will be made at lower temperature, thus enabling more precise estimates to be made of the activation energy.

The activation energy of most flame reactions lies below 10 kilocalories per mole, so that at flame temperatures the rate is a very insensitive function of the activation energy. On the other hand for this very reason, flame studies allow an accurate determination of the steric factor and may ultimately allow a determination of its temperature dependence - a problem which has plagued chemical kineticists for many years.

The techniques we propose to use are the conventional discharge tube-mixing techniques, with the addition of direct measurement of radical concentrations using the scavenger probe technique being developed. This work has been summarized by Steacie (Ref. 3) to 1954. More recent work has been carried out by a number of authors (Ref. 15).

DATA ANALYSIS

To make quantitative interpretation of flame structure information it is necessary to analyze the data. An analysis consists of calculating the actual flux profiles and rate of species and heat production profiles by quantitatively accounting for the effects of molecular diffusion and thermal

Table I
Methane-Oxygen Flame Reaction Kinetic Constants

Reaction	Activation Energy, E (kcal mole ⁻¹)	Frequency Factor, A (10 ¹⁴ cm ³ mole ⁻¹ sec ⁻¹)
CH ₄ + OH → CH ₃ + H ₂ O	8.5	2
CH ₄ + O → CH ₃ + OH	8	2*
CH ₃ + O ₂ → OH + H ₂ CO	0	1
H ₂ CO + OH → HCO + H ₂ O	0	2
HCO + OH → CO + H ₂ O	0	2
CO + OH → CO ₂ + H	7	0.3
H + H ₂ O → OH + H ₂	25	1
OH + H ₂ → H ₂ O + H	18	5
O + H ₂ → OH + H	10	1
OH + OH → H ₂ O + O	10	1
O + O + M → O ₂ + M	0	-
H + O ₂ → OH + O	18	5

* The value given is an estimate, normal for this type of reaction.

conductivity. This step is necessary to avoid false impressions of the rate processes involved (Fig. II-3), since in many flames the effects of diffusion and thermal conduction are large.

This analysis is based on the flame equations consisting of conservation laws and the differential equations of transport and chemical kinetics. They have been formulated in generality by Hirschfelder (Ref. 16) and adapted for flame structure analysis by Westenberg (Refs. 17 and 18).

Thermodynamic Information

The thermodynamic functions of interest are enthalpies, and heat capacities. This information is generally available for almost all of the species

13. C. Gunfelder, Jr., "Instruments for the Measurement of Local Flame Temperatures in High-Velocity Streams", APL/JHU CM-768 (1953).
14. R. M. Fristrom, W. H. Avery, R. Prescott, and A. Mattuck, "Flame Zone Studies by the Particle-Track Technique I - Apparatus and Technique", J. Chem. Phys. 22 106-9 (1954).
15. Symposium on Some Fundamental Aspects of Atomic Reactions, Can. J. of Chem., Vol. 38, October (1960), pp. 1643-2032.
16. J. O. Hirschfelder, C. F. Curtiss, and R. B. Bird, Molecular Theory of Gases and Liquids, John Wiley and Sons, New York, pp. 756-83, (1954).
17. R. M. Fristrom and A. A. Westenberg, "Flame Zone Studies IV - Microstructure and Material Transport in a Laminar Propane-Air Flame Front", Comb. and Flame 1, 217-8 (1957).

found in flame inhibition studies over the temperature range of interest. Several standard reference sources are available (such as Ref. 18) which provide data of sufficient accuracy for our purposes.

Transport Coefficient Studies

Transport coefficient information is necessary for the quantitative interpretation of flame structure data. The required data are the binary diffusion coefficients for the species being studied and the thermal conductivities of the mixtures. The multicomponent diffusion coefficients necessary for the interpretation of such systems are a complex function of all of the binary diffusion coefficients (Ref. 19). To avoid this problem it is usual to choose systems in which one species can be considered dominant; in this case, the true multicomponent diffusion coefficients can be accurately approximated by a set of binary diffusion coefficient of the individual species with the dominant species. The general case, with no single dominant species, could also be handled, but only at the expense of an order of magnitude increase in the necessary diffusion coefficient information (in the general multicomponent case, it is necessary to know the diffusion coefficients of all of the possible pairs of species) (Ref. 20).

The principal problem is to obtain precise, reliable data at sufficiently high temperatures to be useful in the flame studies. A new technique has been developed at APL which allows such measurements (Ref. 21). Diffusion coefficient data are available for the methane-oxygen system (Ref. 22).

The problem of thermal conductivities of mixtures is in a less satisfactory state, but fortunately this information is unnecessary for kinetic studies. An analogous "point source" technique is being developed at APL under Task R for making such measurements (Ref. 23).

INTERPRETATION

The interpretation of this information on flame structure consists of deriving chemical kinetic information and information on flame processes (Refs. 24 through 26). The analysis

provides data on rates and fluxes of enthalpy and the various molecular species. Such a description is unambiguous and complete, but if the information is to be useful on other systems it is necessary to deduce the detailed reaction scheme and derive the appropriate kinetic constants, activation energies and steric factors. This step is largely an art, since in a real system all possible reactions occur to some extent. What is desired is to provide a description of the dominant reactions. For this reason there may be several interpretations with varying levels of sophistication and complexity. For flame inhibition work what is usually desired is the simplest possible interpretation consistent with a quantitative description.

Chemical Kinetic Information

Once a mechanism is assigned, the derivation of chemical kinetic data from flame structure and simple reaction rate information is straightforward. The choice of mechanism is largely a subjective process. We will not discuss the process other than to point out that a useful beginning is often a systematic collection of the possible reactions. The data which we have available from flame structure and simple reaction studies are: temperature, concentrations, and rates. The constants which are of interest are: the activation energy E , the steric factor P , and the temperature dependence of the steric factor n . In most kinetic studies, and probably in our initial work, the precision is not sufficient to allow separation of the temperature dependence factor from the exponential dependence of the activation energy. Flame studies, however, do offer information in a temperature regime which may ultimately allow such a separation to be made. If this is possible, it will represent an important addition to kinetic information. The relation of the exponential and power dependence of the reaction rate on temperature is best visualized by expanding the Arrhenius exponential factor as a series:

$$R_i = R_j = X_i X_j Z P T^n e^{-E/RT} \\ = X_i X_j \frac{Z_0}{T_0^{\frac{n-1}{2}}} P \left\{ T^{n-\frac{1}{2}} - \frac{E}{R} T^{n-3/2} + \frac{E^2}{2! R^2} T^{n-5/2} + \dots \right\}$$

18. A. A. Westenberg and R. M. Fristrom, "Methane-Oxygen Flame Structure II - Conservation of Matter and Energy in the One-Tenth Atmosphere Flame", *J. Phys. Chem.* **64**, 1393-8 (1960)
19. F. D. Rossini, et. al, Selected Values of Physical and Thermodynamic Properties of Hydrocarbons and Related Compounds, Carnegie Press, Pittsburgh, Pa. (1953).
20. R. E. Walker, N. deHaas, and A. A. Westenberg, "Measurements of Multicomponent Diffusion Coefficients for the CO₂ -He-N₂ System Using the Point-Source Technique", *J. Chem. Phys.* **32**, 1314-16 (1960).
21. R. E. Walker and A. A. Westenberg, "Molecular Diffusion Studies in Gases at High Temperature I - The 'Point Source' Technique", *J. Chem. Phys.* **29**, 1139-46 (1958)
22. R. E. Walker and A. A. Westenberg, "Molecular Diffusion Studies in Gases at High Temperature IV - Results and Interpretation of the CO₂ -O₂, CH₄ -O₂, and H₂O-O₂ Systems", *J. Chem. Phys.* **32**, 436-42 (1960).

Since the species are interrelated, a number of cross checks should be possible on the derived kinetic constants, so that there will be a reasonable assurance of the essential correctness of the derived mechanism.

Simplified Theory

The goal of these studies is to provide a description of flame inhibition which is relatively simple and yet still will yield quantitative predictions. This may not always be possible, but it appears that in the case of the methane-oxygen flame a start can be made. The goal of such a simplified theory would be to enable quantitative predictions to be made of the effect of inhibitor on flame reactions (and hence burning velocity) with a precision of 20 per cent. In a sense this simply represents the limit of interpretations, since all mechanisms represent abstractions of the true system.

RECOMMENDATIONS OF FLAME INHIBITION

As a result of this work it is hoped that it will be possible to make some practical recommendations on flame inhibition. It is hardly possible to predict in advance what such recommendations will be, but it seems reasonable to expect that they will include recommendations of specific inhibitors for specific flame systems.

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23. R. E. Walker, N. deHaas, and A. A. Westenberg, "New Method of Measuring Gas Thermal Conductivity", Phys. of Fluids **3**, 482 (1960).
 24. R. M. Fristrom, A. A. Westenberg, and W. H. Avery, "Etude du Mechanisme de la Reaction Propane-Air par l'Analyse de Profils de Fronts de Flamme", de l'Institut Francais du Petrole et Annales de Combustibles Liquides, Vol. XIII, No. 4, pp. 544-553 (1958)
 25. A. A. Westenberg and R. M. Fristrom, "Methane-Oxygen Flame Structure IV - Chemical Kinetic Considerations", J. Phys. Chem. to be published Feb. 1961.
 26. R. M. Fristrom and A. A. Westenberg, "Experimental Chemical Kinetics from Methane-Oxygen Laminar Flame Structure", Presented at the Eighth Symposium (International) on Combustion, Sept. 1960.

FLAME INHIBITION RESEARCH

III. SUMMARY OF PREVIOUS WORK

SUMMARY OF WORK PRIOR TO 1 NOV. 1960

The work prior to the initiation of this contract (on 1 November 1960) consisted of a survey of the possibilities of applying flame structure data to flame inhibition studies, and a suggested program for such studies. This work was summarized in a proposal submitted to the U. S. Army Engineering Research and Development Laboratory in October 1960 (Ref. 4, on page 2).

SUMMARY OF PROGRESS, 1 NOV. 1960 - 31 JAN. 1961

(1) A flat flame burner was set up for flame inhibition studies. The previously studied methane flame (CH_4 - 0.078, O_2 - 0.92; pressure 0.05 atm) was successfully stabilized with added inhibitor (HBr) up to concentrations of 0.0035 mole fraction.

(2) Using the uninhibited flame, two flame composition profiles were run on the new time-of-flight mass spectrometer. A third run was made on an inhibited flame to test the sensitivity of the instrument for HBr. From these test runs it was concluded that the instrument requires the addition of a trap current regulator to allow quantitative work and a forepump trap to shorten the HBr pump-out time. These items have been ordered, and the inhibitor studies have been temporarily transferred to the CEC magnetic-deflection spectrometer.

(3) A computing machine routine for the automatic reduction of flame structure data has been developed.

(4) A report on the flame inhibition program was presented 15 November 1960 at the meeting on "Inhibition of Ignition and Flames with Chemicals." This meeting was sponsored by the Committee on Fire Research of the National Academy of Sciences, National Research Council.

SUMMARY OF PROGRESS, 1 FEB. - 30 APR. 1961

(1) A spherical flame apparatus was set up for inhibition studies on the system (CH_4 - 0.09, O_2 - 0.179, Ar - 0.727, pressure 0.05 atm). Composition profiles were measured for both the inhibited and uninhibited flames with the same total mass flow. In addition to measuring the concentrations of the species which were stable to sampling, an attempt was made to measure hydrogen-atom concentration profiles using a new device, "the scavenger sampling probe." Qual-

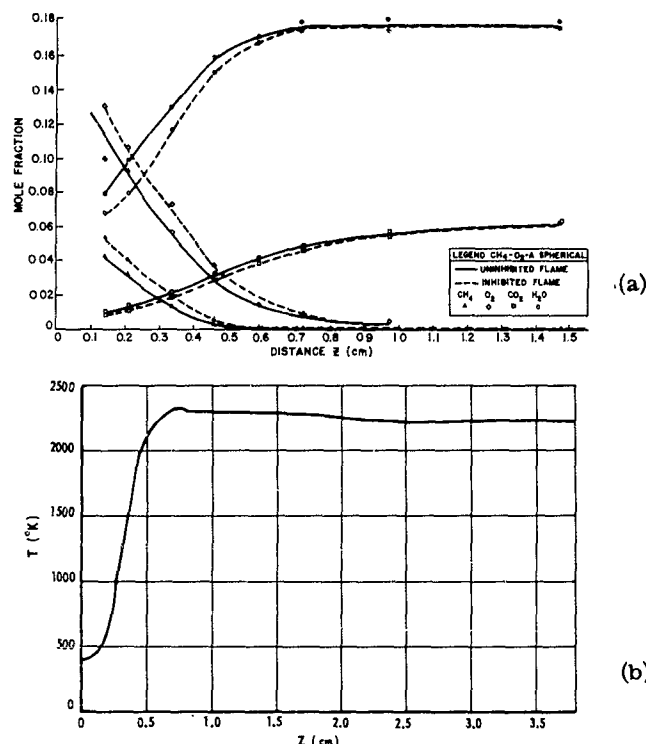


Figure III-1 THE EFFECT OF ADDED HBr ON THE CHARACTERISTIC PROFILES OF A STOICHIOMETRIC METHANE FLAME. COMPOSITION OF THE MAJOR SPECIES (mole fractions) AND TEMPERATURE IS PLOTTED AGAINST DISTANCE, Z (cm.).

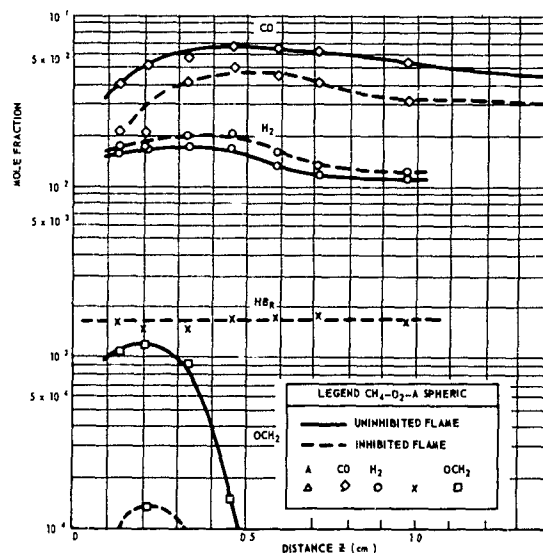


Figure III-2 THE EFFECT OF ADDED HBr ON THE CONCENTRATIONS OF INTERMEDIATE SPECIES IN A STOICHIOMETRIC METHANE FLAME.

itative consideration of these results yields the following information:

- (a) The addition of this amount of inhibitor ($X(\text{HBr}) = 0.0015$) appreciably lowers the rate of reaction of the major species (CH_4 , O_2 , and CO).
- (b) Peak concentrations of the intermediate species (OCH_2 , CO , and H_2) are changed and shifted relative to the major species. When data on diffusion and temperature become available, a quantitative interpretation may yield H , OH , and CH_3 radical concentration data.
- (c) The apparent concentration of H atoms was depressed almost five-fold by the addition of HBr even in the equilibrium region.
- (d) HBr was the only bromine-containing compound detected ($X(\text{Br}_2) < 10^{-4}$).

These latter two results are probably artifacts of the techniques. They may, however, yield useful information and provide a guide for subsequent work.

(2) A series of computer calculations were made of the adiabatic temperatures and compositions of the equilibrium burned gases of some CH_4 - O_2 - Ar mixtures and of the effect of adding small amounts of HBr . A routine developed by the U. S. Naval Ordnance Test Station was used on the APL computer to determine these flame properties. One of the most interesting points developed was that the HBr is almost completely dissociated and that bromine atoms are the commonest inhibitor species in the burned gases.

SUMMARY OF PROGRESS, 1 MAY - 31 JULY 1961

(1) A set of composition and temperature profiles was determined for a flat methane-oxygen flame (CH_4 - 0.078, O_2 - 0.92, $P = 0.05$ atm) with and without added HBr . Corrosion was so severe that the burner assembly became unusable. A new teflon-glass-ceramic system including traps to protect the pumps has been designed and is being fabricated.

(2) A program to measure the high-temperature diffusion coefficients for the analysis of the inhibited flames has been started. The apparatus has been activated and tested at 300°K and 500°K on the well-known $\text{He}-\text{N}_2$ system. Measurements have been started on the $\text{Ar}-\text{H}_2$ system.

(3) Scavenger probe studies of radical concentrations in flames have been continued. The use of chlorinated hydrocarbons as scavenger for H atoms has been shown to be proportional to H , but the scavenging efficiency was found to be only 10%. Studies of oxygen-atom concentration using NO_2 as a scavenger have been made.

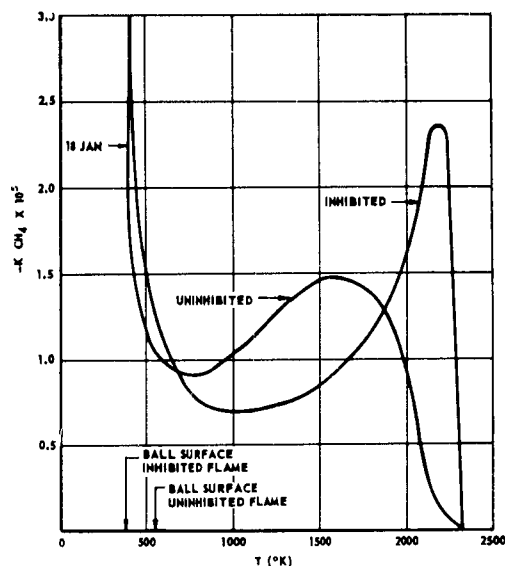


Figure III-3 THE EFFECT OF HBr ON THE RATE OF METHANE DISAPPEARANCE IN A STOICHIOMETRIC FLAME (CH_4 - 0.09 O_2 - 0.179 Ar - 0.727; $P = 0.05$ atm.). K_{CH_4} (MOLES PER CUBIC CM. PER SEC $\times 10^5$) VERSUS TEMPERATURE ($^\circ\text{K}$)
NOTE THAT AN APPRECIABLE PART OF THE REACTION APPEARS TO BE OCCURRING AT OR NEAR THE SURFACE OF THE BALL. THE PRINCIPLE EFFECT OF THE ADDED HBr APPEARS TO BE A SHIFT OF THE REACTION TO THE HIGH TEMPERATURE REGION.

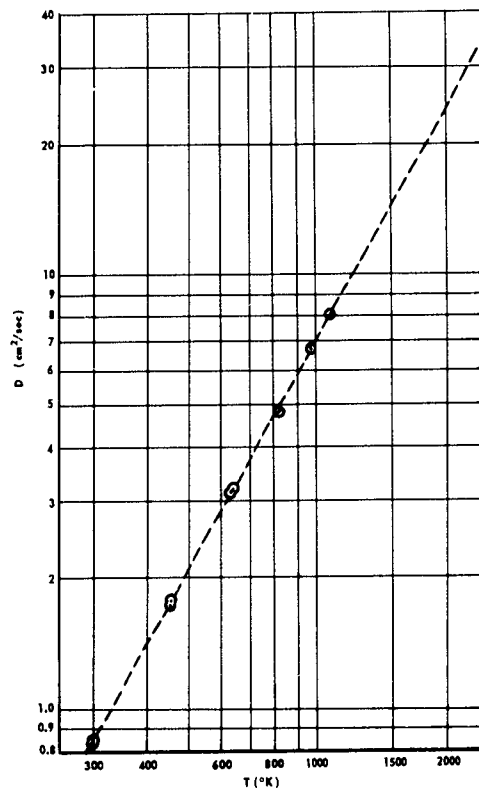


Figure III-4 DIFFUSION COEFFICIENT OF THE SYSTEM H_2 (trace) - Ar AS A FUNCTION OF TEMPERATURE

The efficiency of scavenging appears to be high and may approach 100%.

SUMMARY OF PROGRESS 1 AUG. - 31 OCT. 1961

(1) A temperature profile was measured on the spherical stoichiometric methane-oxygen-argon flame system whose composition structure was reported in TG 331-2. (See pp 9)

(2) The diffusion coefficient of the system A-H₂ (tract) was measured at atmospheric pressure over the range 300 - 1070°K. (See pp 12)

(3) An analysis was made of the effect of added HBr on the fluxes and rates on the stoichiometric methane flame. (See pp 14). This indicated that a significant fraction of the reaction occurred on the surface of the ball.

Table III-I

Test of Oxygen-Atom Scavenger Probe*

System							
CH ₄	O ₂	Ar	T ₀ (°K)	P (atm)	Expt'l.	Calculated**	
0.078	0.92	0.002	313	4.1	0.0019	0.0020	
0.10	0.18	0.72	313	3.8	0.0030	0.00325	

* Using the reaction $O + NO_2 \rightarrow NO + O_2$ for scavenging in flame microprobe-sampling studies (see Figure III-5a below), the O-atom concentrations were found in the equilibrium burned gas region.

** Values interpolated from calculation made using $T_0 = 400^\circ K$.

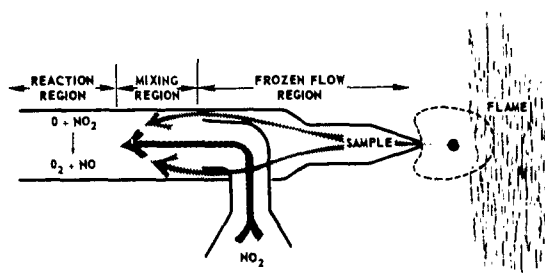


Figure III-5a SCHEMATIC DIAGRAM OF SCAVENGER PROBE USED IN MEASURING OXYGEN ATOM CONCENTRATIONS

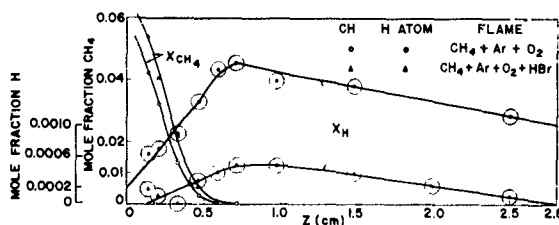


Figure III-5b EFFECT OF ADDED HBr ON COMPOSITION PROFILES OF METHANE AND HYDROGEN ATOMS IN A METHANE FLAME

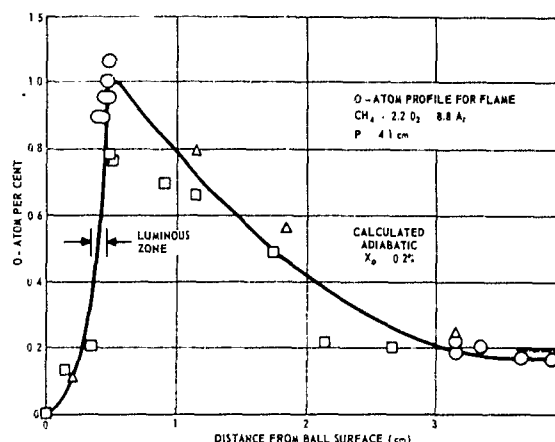


Figure III-5c TEST OF OXYGEN-ATOM SCAVENGER PROBE

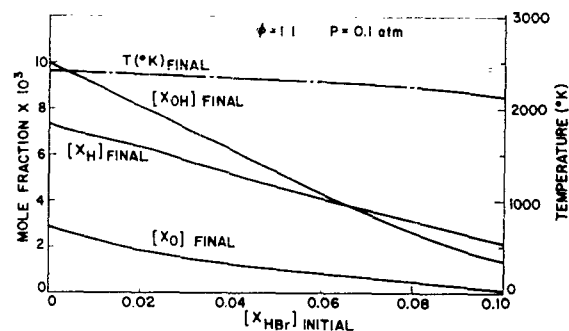


Figure III-6a EFFECT OF ADDED HBr ON FLAME RADICAL CONCENTRATIONS (H, O, OH)

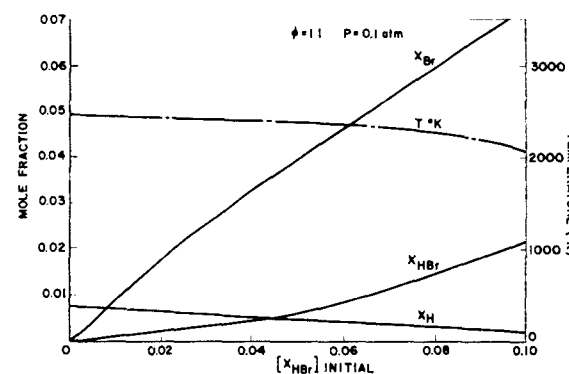


Figure III-6b EFFECT OF HBr CONCENTRATION ON INHIBITOR SPECIES CONCENTRATIONS (HBr, Br, Br₂)

FLAME INHIBITION RESEARCH

IV. REPORT OF PROGRESS

1 NOVEMBER 1961 - 31 JANUARY 1962

CORROSION RESISTANT BURNER SYSTEM

W. E. Wilson, J. T. O'Donovan, and R. M. Fristrom

Previous inhibition studies of flame structure (see pp 8-10) were unsatisfactory because of corrosion problems. Bromine atoms formed in the flame attacked stainless steel parts of the burner forming ferric bromide which distilled throughout the system contaminating it. In addition it was found that corrosion of the hydrobromic acid orifice inlet resulted in irreproducible flow rates.

To avoid these problems a new spherical burner system (see Fig. IV-1) has been designed and constructed using exclusively glass, teflon and ceramic parts. All gaskets, "O" rings and entry ports are of teflon, and in addition a potassium hydroxide trap is used to protect the pumps and pump oils.

The system also will incorporate a critical orifice flow metering system featuring an all teflon pressure transducer for handling corrosive inhibitors.

Temperature measurements will be made by means of a pneumatic probe¹ which utilizes the sampling probe as its first orifice. This will allow direct association of the temperature with the composition measurements. The principle of these temperature measurements is that if two

1. See Reference 13 - Page 5.

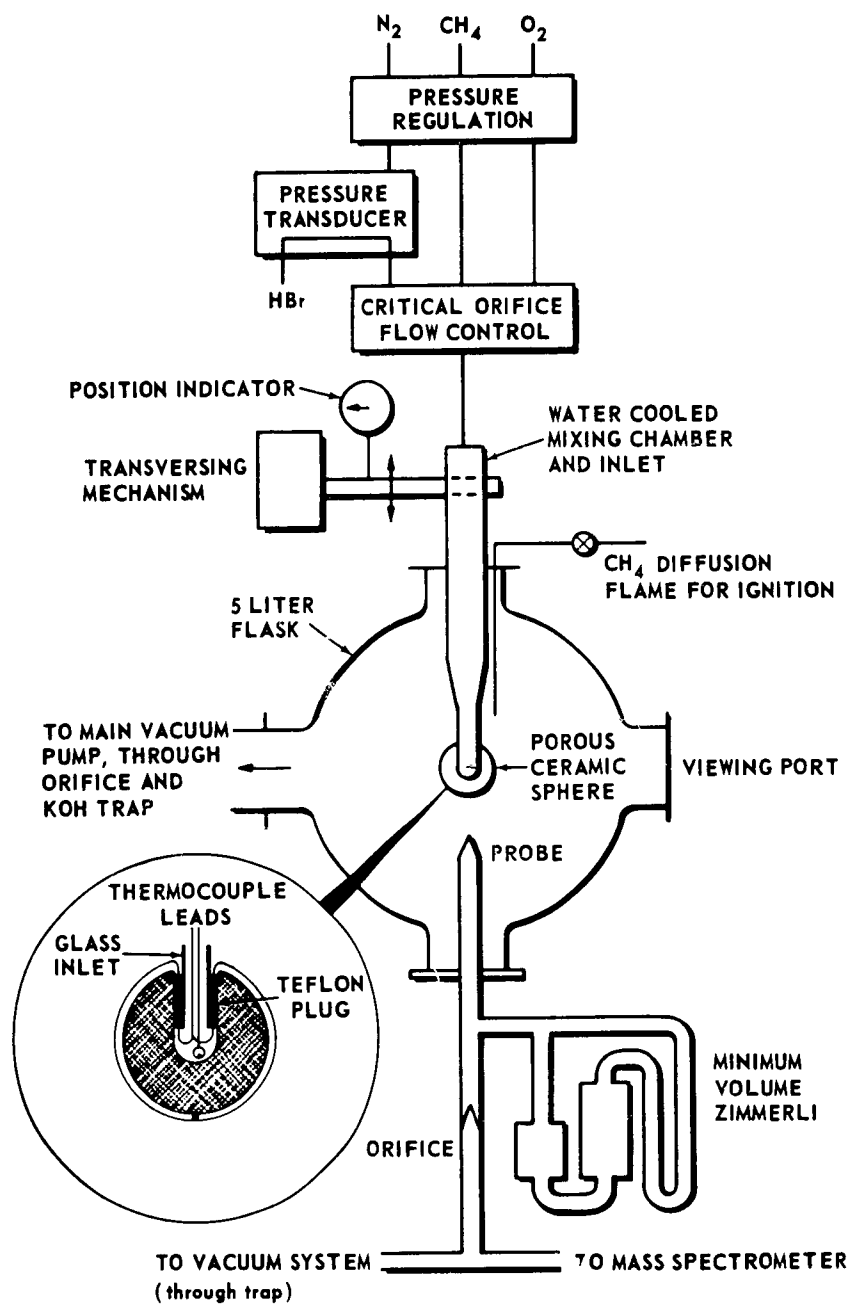


FIG. IV-1 SCHEMATIC DIAGRAM OF CORROSION RESISTANT
SPHERICAL FLAME INHIBITION APPARATUS

orifices are introduced in series both operating under choking continuum flow the pressure between the two orifices is a function of the temperature above the two orifices (it is assumed that specific heat and molecular weight are unchanged and any changes due to Reynolds number are accommodated by calibration). Successful operation of such a system using a microprobe inlet requires a minimum volume to avoid excessive delay in reaching pressure equilibrium. To accomplish this a minimum volume Zimmerli style mercury manometer has been designed and constructed.

PRESSURE TRANSDUCER

J. T. O'Donovan, W. E. Wilson, and R. M. Fristrom

One problem often encountered in flame inhibition studies is that of accurate flow control of a corrosive inhibitor. A standard flow system consists of a pressure regulator and a flow control device such as a critical orifice.¹ Since there are no corrosion resistant pressure regulators of sufficient accuracy available it was necessary to develop one for these studies.

The technique used was to pressurize the corrosive gas with a non-corrosive gas whose pressure could be regulated by standard methods. In one application of the pressure transducer (Fig. IV-2a), nitrogen, in the aluminum section, is used to transmit the desired pressure, through a slack teflon diaphragm to hydrogen bromide, in the glass section. As the hydrogen bromide flows out through the critical orifice (sapphire mounted in teflon) nitrogen flows in and the diaphragm moves to maintain a constant pressure.

A working model (Fig. IV-2b) has been constructed and tested. It appears to maintain a sufficiently stable pressure to give accurate flow control. The system has been checked on the mass spectrometer and has no internal or external leaks. It should permit the easy handling of corrosive gases.

It will also be used for expensive materials such as isotopic species. Since it may be heated it will also be suitable for metering solids or liquids which can be volatilized below 500°F.

1. J. W. Andersen and R. Friedman, Rev. Sci. Inst. 20, 61 (1949).

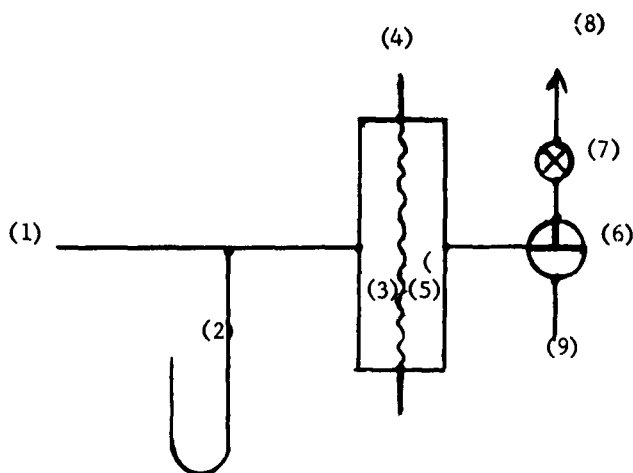


FIG. IV-2a SCHEMATIC DIAGRAM OF CORROSION RESISTANT
CRITICAL ORIFICE FLOWMETER

- (1) Adjustable constant pressure source of non-corrosive gas.
- (2) Manometer for determining pressure.
- (3) Transducer Base Plate (see Fig. 2b).
- (4) Slack teflon diaphragm (0.003").
- (5) Corrosion resistant reservoir. This is the cover of a large dessicator (Pyrex No. 3120) With appropriate modification any size could be used.
- (6) Three way 6mm. stopcock.
- (7) Critical orifice (saphire watchmakers jewel mounted in teflon plug).
- (8) Regulated corrosive gas flow.
- (9) Reservoir of corrosive gas for filling transducer.



FIG. IV-2b PICTURE OF CORROSION RESISTANT PRESSURE
TRANSDUCER

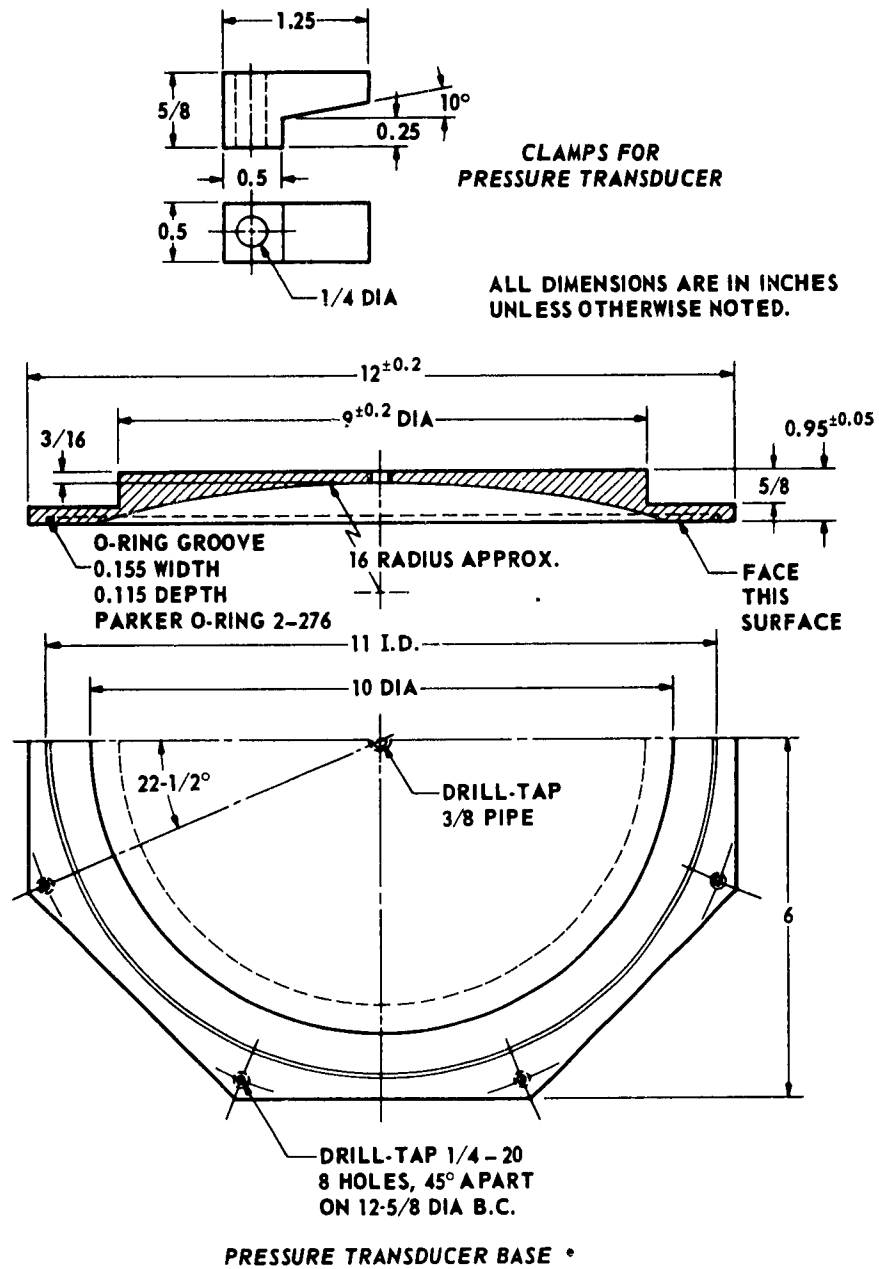


FIG. IV-2c DRAWING OF TRANSDUCER BASE PLATE

THE CONCENTRATIONS OF ATOMIC AND FREE RADICAL SPECIES IN FLAMES

R. M. Fristrom, and C. Grunfelder

Free radical and atomic species are important in flame reactions and therefore a method called "the scavenger probe" technique has been developed for measuring the concentration of these species.

Earlier studies (see pp 8-10) were promising and indicated a number of avenues for improvement. During the past quarter the apparatus was reconstructed to incorporate the following improvements: (1) The probe was watercooled over its entire length excluding only the inlet nozzle, (2) provision was made for cleaning the probe in situ, (3) the probe was oriented vertically so that captured dust particles would not lodge in the nozzle area, (4) provision was made for making blank runs using a scavenger by-pass inlet and a silver "spoiler" to destroy the atoms. (Metering of the scavenger gas was improved by incorporating a 1° taper stainless steel needle valve in the inlet line).

Operation of the system on the spherical flame burner was satisfactory. The results were reproducible within 2% from day to day (see Fig. IV-3) and were independent of NO_2 flow rate and sampling pressure level (from 50-300 microns). After cleaning, the system could be used for several days.

A study was made of the oxygen atom profile of a twentieth atmosphere methane-oxygen flame (see Fig. IV-3). The profile was reproducible, of the correct order of magnitude and of the expected shape. It was not possible to make a direct comparison of the results with

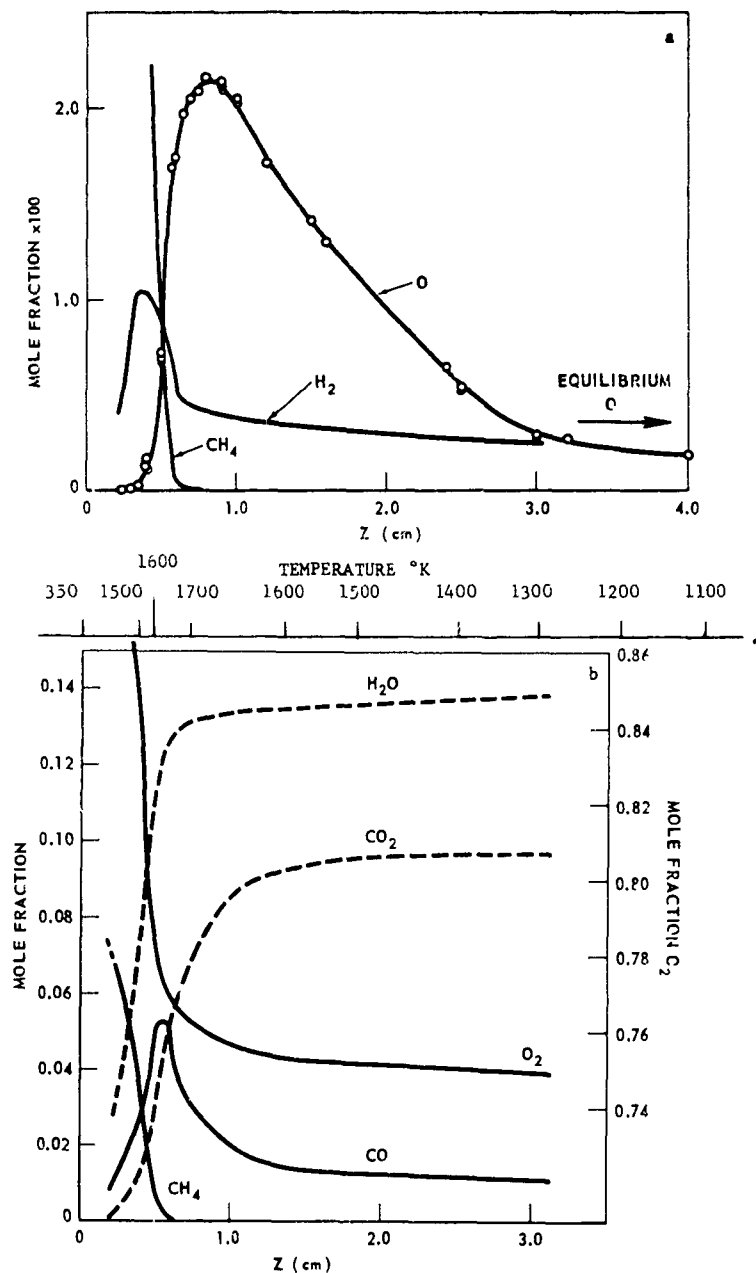


FIG. IV-3a OXYGEN ATOM CONCENTRATION IN A SPHERICAL
METHANE FLAME
-3b COMPOSITION PROFILES OF MAJOR SPECIES

equilibrium calculations because conduction losses lowered the flame temperature more rapidly than the relatively slow three-body recombination of oxygen could follow. Hence, equilibrium was not reached in the sampling region.

In spite of this problem it appears that the technique is sound since reproducible results were obtained under differing conditions with different apparatus. However, the final proof will have to await studies on a flame which is in true equilibrium or in an electric discharge where comparison can be made using other established techniques such as gas phase titration and Wrede gauges ¹.

1. K. R. Jennings, "The Production, Detection, and Estimation of Atoms in the Gaseous Phase", Quarterly Reviews 15, 237 (1961).

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