ΝΟΤΙCΕ

THIS DOCUMENT HAS BEEN REPRODUCED FROM MICROFICHE. ALTHOUGH IT IS RECOGNIZED THAT CERTAIN PORTIONS ARE ILLEGIBLE, IT IS BEING RELEASED IN THE INTEREST OF MAKING AVAILABLE AS MUCH INFORMATION AS POSSIBLE

I

4.

A SEARCH FOR EXPERIMENTS TO EXPLOIT THE SPACE SHUTTLE ENVIRONMENT

VOLUME ONE

(NASA-CR-163030) A SEARCH FOR EXPERIMENTS N80-26337 TO EXPLOIT THE SPACE SHUTTLE ENVIRONMENT, VOLUME 1 Final Report (Relay Development Corp., Branford, Conn.) 70 p HC A04/MF A01 Unclas CSCL 22A G3/12 22353

Relay Development Corporation

BRANFORD, CONNECTICUT 06405



A SEARCH FOR EXPERIMENTS TO EXPLOIT THE SPACE SHUTTLE ENVIRONMENT

by

John B. Fenn

Final Report on Contract #955578 between Relay Development Corporation and NASA-JPL

"This work was performed for the Jet Propulsion Laboratory, California Institute of Technology sponsored by the National Aeronautics and Space Administration under Contract NAS7-100."

15 December 1979

RELAY DEVELOPMENT CORPORATION

226 Pleasant Point Road

Branford, Connectiout 06405

06405

203-488-5428

First of Two Volumes

Abstract

This study extends an earlier search for worthwhile experiments in pure and applied physics and chemistry which might take advantage of conditions achievable aboard the Space Shuttle. Of particular interest were the very large pumping speeds at high or ultra high vacuum, the highly non-equilibrium composition of the ambient atmosphere, and the relative absence of gravitational effects. Ideas and suggestions were solicited in the course of visits to 31 research establishments in Western Europe, India and Japan, conversations with over 90 scientists, and presentations at three international meetings. Intriguing possibilities emerged in five arenas: (1) Spectroscopy of the transition state in chemical reactions; (2) Flame structure and analysis; (3) Solid propellant combustion; (4) Analysis of atmospheric composition; (5) Turbulence Effects on Aerosol Coagulatior.

TABLE OF CONTENTS

¥

Volume I

	Summary1									
I.	Introduction3									
II.	Results 6									
III.	Conclusions and Recommendations 24									
IV.	References 25									
	Appendices 26									
	A. People Engaged in Direct Conversation 27									
	Field of Interest Key 31									
	B. Asilomar Conference on Dynamics of Molecular Collisions 32									
	C. llth International Symposium on Rarefied Gas Dynamics 41									
	D. 7th International Symposium on Molecular Beams 58									

Volume II

Places Visited	67
Appendix A - "From Sacred Cows to Satellites, India's Great Leap Upward"	86
Appendix B - "India's Hyderabad - An Old City With a New University"	93

Summary

This report comprises a geographic and substantive extension of an earlier attempt in 1976 to identify worthwhile experiments which might advantageously be carried cut aboard a Space Shuttle in orbit. The earlier investigation focussed on experiments with free jets which would take advantage of the very high pumping speed available in space. It was based on communication with scientists in America and Western Europe. The present author, who had participated in the earlier study, last year spent two months in India, three months in Japan and one month in Europe visiting laboratories and talking with research scientists and engineers. In addition, during the period since the first report was completed, he had made presentations at three international meetings in an attempt to stimulate interest in and suggestions for such experiments and others not involving free jets. This document reports on the results of these further attempts to identify worthwhile experiments. The harvest was rather lean but five interesting avenues of investigation were identified:

1. <u>Natural Lasers and Transition State Spectroscopy</u>. In the upper atmosphere the species OH is vibrationally very hot and rotationally very cold and thus invites attempts to obtain stimulated emission over very long path lengths. It might be rewarding to look very carefully at the infrared emission in the wings of excited OH. Because OH is formed by highly exothermic reaction of ozone and H atoms, any observed broadening could be interpreted in terms of the radiation occurring while the OH was still closely associated with the OOOH transition state complex.

2. <u>Very Low Pressure Flames</u>. It turns out that the volumetric flow rate of premixed fuel and oxidant required to maintain a stable flame is inversely proportional to the pressure. The lowest pressures which have been reached on earth are of order one torr where the mean free path is only a micron. To reach much lower pressures one

must have very high pumping speeds, such as might be achieved on the Space Shuttle. One inviting prospect is that flame front structure could be resolved in much more detail than is now possible. Another is that collisional quenching times could be increased to the same order as radiation life times, thus giving rise to possibly interesting radiation effects.

3. <u>Solid Propellant Combustion</u>. For reasons similar to those involved in gaseous flames, achieving stable combustion of solid propellants at very low pressure also requires very high pumping speeds. There are a number of interesting and unanswered questions about low pressure deflagration of such propellants which might be answered by experiments at high pumping speeds.

4. <u>Composition of Upper Atmosphere</u>. In spite of many measurement attempts there are large uncertainties in the composition of the upper atmosphere which stem from difficulties in avoiding perturbations of the composition by the measuring instrument. Possibilities for overcoming these problems include velocity discrimination in sampling and laser sensing techniques.

5. <u>Turbulence Effects on Aerosol Coagulation</u>. Agglomeration or coagulation of colloidal particles due to Brownian motion is reasonably well understood. Agglomeration of larger particles in gaseous systems may be strongly affected by velocity fluctuations due to turbulence. Such effects may have had a profound influence on the composition of planets and other bodies formed by condensation from primordial gas. Interesting agglomeration experiments may be possible on the Space Shuttle in the absence of gravitational settling in gaseous suspensions of large particles.

A SEARCH FOR EXPERIMENTS TO EXPLOIT THE SPACE SHUTTLE ENVIRONMENT

I. Introduction

In 1976 Relay Development Corporation (RDC) undertook a study program for NASA-JPL aimed at identifying and evaluating worthwhile experiments which might be advantageously carried out in an environment to which the Space Shuttle promises to provide access. In particular, RDC's assignment was to consider experiments in which supersonic free jets would play a vital role. Such free jets have been rewarding objects of study because they comprise a rich source of phenomena which occur under conditions readily achieved in such jets but which are not otherwise obtainable. The attraction of the Space Shuttle Environment (SSE) consists in its ability to provide virtually unlimited pumping speed at relatively low background pressures. Such large pumping speed would make possible much higher source Reynolds numbers at lower background pressures than would be economically feasible at the bottom of the ocean comprising Earth's atmosphere. Large Reynolds numbers would permit experiments at relatively large scale and/or at large pressure ratios for the expansion.

The RDC Study culminated in a report which summarized the results of an extensive literature search, conversations with nearly 100 scientists and analysis of a number of gedanken experiments.⁽¹⁾ The essential conclusions were:

- (1) No experiments emerged which in the sense of their scientific implications could be considered sufficiently critical to justify <u>ab initio</u> the investment of resources which would be required to carry them out aloft.
- (2) There were some interesting and worthwhile experiments which, though not "critical", the SSE could expedite or enhance but there could be no assurance that clever earthbound investigators might not find ways to achieve equivalent results at lower cost and in less time.

(3) The prospectively most interesting experiments were spectroscopic in the sense that they all involved the interaction of molecules with photons. The principal basis for their appeal stems from the ability of the SSE to provide long optical paths in nearly collisionless gas. These long paths mean that there can be relatively large numbers of molecules in the field of view of a detector for relatively long times. The absence of collisions means that the states of the molecules do not change. Thus, initially non-equilibrium conditions can be maintained.

Since the appearance of that RDC report there has been renewed consideration of the so-called Molecular Wake Shield or MWS. This concept involves the disposition of an impermeable surface of relatively large frontal area on the Shuttle itself or on some outboard vehicle. The idea is that this shield would sweep the molecules of ambient atmosphere out of its path so that in its immediate wake the effective background pressure or density would be substantially lower than that of the ambient atmosphere through which the Shuttle is flying. Subject to some unknowns, e.g., the contribution to be expected from outgassing of shield and vehicle surfaces, it has been estimated that the effective background pressure for a typical orbit altitude and velocity would be reduced from about 10^{-6} torr in the free stream to the vicinity of 10^{-12} torr^(2,3) in the wake of the shield. This ultra-high vacuum environment may make inviting some experiments which might not be feasible or interesting at the much higher background pressure presumed in the original study. Moreover, a corollary consequence of such a shield is the possibility of generating a collimated beam of molecules of ambient gas by means of a suitably shaped inlet aperture in the shield which would play the 'role of a skimmer in conventional nozzle beam systems. Oxygen atous are the dominant atmospheric species at probable oribiting altitudes. For example, they are present at concentrations of nearly $10^{10}/\text{cm}^3$ at an altitude of 200 kilometers. An orbiting velocity of 8 km/s would thus give rise to a beam of oxygen atoms with an intensity near 8 x $10^{15}/cm^2$ -s. Such a beam, which would have a translational energy of about 5 eV, cannot now be readily generated in earthbound laboratories and would make feasible many interesting scattering experiments.

In sum, the SSE in conjunction with the MWS opens up possibilities not contemplated in RDC's original study.

Another limitation of the original RDC study concerned the scientists polled for ideas, suggestions and comments. Though their number was appreciable, they represented a community which was restricted geographically to North America and Western Europe and scientifically to those who had some research experience with molecular beams and/or rarefied gas dynamics. Moreover, good ideas frequently have fairly long gestation periods and many of the scientists who were approached had no previous occasion to consider the possibility of doing experiments in space. Consequently, it seemed appropriate to undertake an extension of the original survey. One objective would be to include a broader segment of the scientific community. Another would be to renew encounters with some of those who have been accosted earlier to see whether some seeds sown the first time around might have since germinated.

As it happened, one of the authors of the original RDC report, Professor John B. Fenn, was to be on leave from Yale University during the spring term of 1979. He had been invited to spend two months at the Indian Institute of Science in Bangalore and at the Institute of Space and Aeronautical Sciences of the University of Tokyo. In addition, he planned to travel a month in Europe visiting laboratories and attending the International Symposium on Molecular Beams beginning on 28 May at Riva del Garda in Italy. This itinerary appeared to provide an excellent opportunity for him to extend the scope of the original survey by seeking ideas from a broader segment of the scientific community than had been included in the earlier report. Moreover, he would have an opportunity for second encounters with some of the people who had been previously exposed to the idea of experiments in the Space Shuttle Environment. Perhaps some seeds sown then might have fallen on fertile ground. Professor Fenn was willing to pursue these goals during his travels and was commissioned to do so. This

report summarizes his experiences and findings. Volume I includes a description of the possible experiments which emerged during the survey, a catalog of the scientists who were interviewed, and an account of the meetings at which formal presentations were made about the possibilities for Space Shuttle experiments. Volume II is essentially a travelogue describing the places which were visited. In the interests of informality and economy of words the writer will resort to the use of the vertical pronoun.

II. The Results

The word about possibilities for experiments on board the Space Shuttle was spread substantially farther and wider in this survey than during RDC's initial study. Even so, the harvest of ideas was in some sense still disappointing. As in our earlier census, most responses fell into one of the following categories; usually expressed in more euphemistic terms than I will employ.

- 1. This is all so new and different that I will have to think about it.
- 2. It is too far from my regular research beat and I can't see any connection which would make it worth my while to give it much attention.
- 3. Clearly a boondoggle. I could make better use of the money in my lab on the ground.
- 4. You Americans have enough money to contemplate such experiments. We have to devote our resources and attention to more immediate problems.
- 5. What a great idea! You could do this, and this, and this,..., all the "thises" being fairly thoughtless and naive.
- 6. Sounds like a great opportunity but I can't think of any way to take advantage of it.

Amidst all this chaff, there were a few grains of wheat, some proposed experiments which seemed interesting enough to report. They are set forth in the following paragraphs of this section.

6

1. Lasers Au Naturel and Transition State Spectroscopy. After nearly twenty-five years of molecular beam reactive scattering experiments and maybe a decade of spectroscopy and photochemistry by lasers, today's chemists know more about the microscopic details of chemical reactions than Arrhenius and Bodenstein could have dreamt of. In particular, these modern techniques have led to so-called "state-to-state" chemistry which contemplates complete experimental specification of velocities, internal energy levels and even the orientation of reactant and product species immediately before and after a reactive encounter. What still eludes the most intrepid investigators is any direct observation of that construct, the "transition state"--a configuration of species which marks at once the end of their identity as reactants and the beginning of their identity as products. Of extremely short lifetime and perhaps in some cases not even identifiable or definable, this ephemeral species has existed thus far mainly in the minds of investigators. Whatever characteristics have been ascribable to its reality are largely based on deductions as to plausible bridges between reactant states and product states. Before the beam and laser era, most experimental data related to the initial state of reactants and the final state of products. We now have increasingly direct information on the final state of reactants and the initial state of products so the bridges to be built are much shorter and therefore less speculative in their design.

The direct experimental identification and description of the transition state has been called by John Polanyi the "last frontier" of chemical kinetics. Of late there have been a number of suggestions by theorists that extremely high powered lasers might be able to probe the transition state structure, and indeed, take advantage of its asymmetric nature by bringing about laser promoted reactions between reactants which in their symmetric initial configurations are not capable of absorbing photons.(4,5) The calculated power requirements are

so large that many people are pessimistic about the outcome. Among the optimists is John Polanyi of the University of Toronto, and he sees the Space Shuttle, literally and figuratively, as a possible vehicle for experiments which might achieve the goal of direct observation of the transition state. After some discussion with me at the Riva meeting he wrote a letter summarizing his ideas. I know of no better way to present them than to reproduce his own words. What follows are essentially literal excerpts from his letter.

"The proposals for the skylab could be made to hinge on the observation of OH⁺ (vibrationally excited hydroxyl in its ground electronic state) and also Na* and Na** (electronically excited Na in the 3p and 4s states respectively). Since OH⁺ is emitting strongly from approximately 90 km and is hard to observe at ground level, I shall concentrate on that in setting forth the following two objectives:

(I) Look (for the first time) for evidence of what I termed in 1960 and 1961 publications, prior to the first laser, <u>a "natural iraser" comprised of the</u> <u>upper atmosphere</u>. The point was that vibrational temperature $T_{vib} \approx 10,000$ K and rotational temperature $T_{rot} \approx 230$ K. Hence hundreds of transitions in the P-branch of the fundamental ($\Delta v = 1$ at approximately 3 microns in the near infrared) have the potential for producing stimulated emission in view of the long path length.

The existence of "an inverse Frauenhofer effect" was postulated, i.e., the continuum emission from the sun should show narrow intervals of enhanced intensity when viewed through a long path-length of the upper atmosphere. (This, by the way, was the first time that $T_{vib}^{>>} T_{rot}^{-} > 0$ had been noted to be a sufficient criterion for what was termed "partial population inversion"--a concept that underlies the CO₂ and CO vibrational lasers).

(II) A far more venturesome proposal which, however, requires a similar line of experimentation, is to look for inframed emission in the wings of the OH^+ (v + v - 1) fundamental, or the OH^+ (v + v - 2, v + v - 3,...) overtones, extending into the photographic region. This would constitute observations of a type of collision-induced broadening due to strong (high energy) collisions that could only be due to the fact that OH^+ is produced chemically through $H + O_3 + [HOOO]^{\dagger} + HO^{\dagger} + O_2$ (as is highly likely to be the case) and that one is observing the OH radical before it has moved out of range of the other product, O_2 ; i.e., one would have (for the first time) spectroscopic data bearing on the forces operating in $[HOOO]^{\dagger} --$ <u>spectroscopy of the transition</u> state in a chemical reaction.

Such an observation has the best chance of success if one can observe a very long column of reacting $H + O_3$, as is the case in the upper atmosphere. The low pressure ensures that there is no local depletion of H through rapid reaction with O_3 , i.e., it solves the difficult problem of reagent mixing for fast reaction. Observation from space eliminates water absorption of radiation.

A similar experiment is possible for that part of Na* and Na** that is formed chemically in the upper atmosphere, naturally through reaction of $0 + \text{Na0}^+ + 0_2 + \text{Na*}$ (or Na**) or from the reaction of naturally-occurring atomic species with Na₂ injected into the surroundings from z supergonic orifice, thus $0 + \text{Na}_2 + \text{Na0} + \text{Na*}$.

The ratio of concentrations of transition state to product species is $10^{-10} - 10^{-11}$ for $0H^{\dagger}$, and $10^{-4} - 10^{-5}$ for Na*. This renders the experiment very difficult, but not impossible in view of the fact that photons can be counted against a dark background. In the case that Na₂ is injected, interposition of a chopper blade in the Na₂ flow will permit subtraction of any background from signal.

We very much want to try feasibility studies on Objective II in our laboratory and are ready to do so as soon as we get the signal to go ahead.

2. Flames At Very Low Pressures. Flames have fascinated man ever since he began to control and use fire, probably even before. Until Galileo introduced the controlled experiment and launched the era of modern science, fire as manifested by flame was regarded with awe as an embodiment of godly magic. The Greeks assigned it a fundamental role in the composition of the real world. Along with earth, air and water, it was considered one of the four elements from which all things were made. Flames are rather spectacular consequences of a complex interaction between physical and chemical processes. The combustion which gives rise to them is still the major source of the heat from which prime movers derive the work that powers industrial society. For these reasons, flames have been the objects of intense scientific study for nearly two centuries. Investigators have tried to understand their phenomenology in terms of their physics and chemistry. They have sought to relate their speeds, structure and stability to the thermodynamics and kinetics of chemical reactions and the physical transport of mass energy and momentum. In recent years there has been increasing interest in the role of reactants and products which are present in amounts too small to effect flame behavior but large enough to have major environmental consequences. Growing concern about the depletion of oil and gas resources has renewed interest in achieving even small increases in the efficiency of all kinds of combustion systems.

One important and powerful variable in the experimental study of flames is pressure. In early experiments pressures were almost always atmospheric or higher, partly because such pressures were easy to achieve in the laboratory and partly because practical systems operated in this range. With the advent of high altitude flight internal combustion engines were required to perform at subatmospheric pressures. In the laboratory experimenters began to appreciate and exploit the fact that reaction zones in flames could be greatly extended and magnified by lowering the pressure and therefore the density of the reacting

gases. In flames of premixed gases at one atmosphere much of the action occurs in a region somewhat less than a few tenths of a millimeter in thickness. At 25 torr this flame zone expands to a centimeter or more, depending upon the composition of the entering gas. This increase in thickness greatly improves the resolution of structure which can be achieved by various kinds of probes.

On a very elementary plane of flame phenomenology the question was often asked whether there was a limiting pressure below which a flame front could not propagate into unburned gas. In an attempt to answer this question and at the same time to increase the resolution of their spectroscopic probing of flame front structure, Gaydon and Wolfhard in 1950 carried out a then heroic effort to produce flames at very low pressures. (6,7)They found that there was in effect a characteristic Reynolds number below which stable flames could not propagate. In terms of more directly measurable quantities Reynolds number at a particular flame speed is proportional to the product of pressure and burner diameter. Thus, what Gaydon and Wolfhard found was that as long as this product was maintained at a characteristic value for a particular composition, they could lower the pressure as much as they wanted to as long as they increased the burner diameter correspondingly. They also found that for many fuel-oxidant combinations the burning velocity did not change appreciably with pressure. Thus, as the pressure was lowered the volumetric flow increased and they reached the limit of their available pumping speed at pressures of about one torr with a burner diameter of 10 cm. The pumping speed which they had available was not specified. They simply noted that they were using the largest available industrial vacuum pumps. With these low pressure flames and spectroscopic techniques, Gaydon and Wolfhard probed the species distribution in various flames. More recently, Wolfhard in collaboration with Vanpee, Hinck and Seamans examined the nature of OH radiation in flames at pressures down to about 10 torr

on a burner 10 cm in diameter.⁽⁸⁾ Their system was exhausted by a Roots pump with a speed of 300 liters/sec. Even more recently Vanpee has made observations on flames down to about one torr in a system whose pumping speed is about 2400 liters/sec.⁽⁹⁾ None of these investigators apparently made any particular attempt to cool the burned gases before they arrived at the pump. Appropriate cooling together with judicious use of cryopumping when the exhaust gases are readily condensible could have greatly enhanced their effective pumping speeds. Even so, if one wanted to burn flames at pressures substantially lower than one torr it seems clear that it would be difficult and expensive to provide sufficient pumping speed because the required speed goes up with the inverse square of the pressure. To remove the gases from a flame at a pressure of one millitorr would require 10⁶ times as much pumping speed as would be required at one torr!

Is there any good reason to go to such low pressures? The answer to this natural question is not entirely obvious. A number of reasons can be cited. How good they are is a matter of conjecture. We can give such conjecture a starting point by noting that an age old question in combustion circles has been whether there is really a low pressure limit to flame propagation. To the extent that simple extrapolation of past experience can be relied upon, the answer seems to be that one can maintain flames at as low a pressure as one wants provided only that one makes the diameter of the burner sufficiently large. Testing of this empirical rule to pressures well below one torr would be perhaps interesting but certainly not of overwhelming importance per se. However, upon more careful reflection some intriguing possibilities emerge. In the first place, it is to be recalled that some of the elementary exothermic reactions in many flames almost certainly involve atom and radical recombination which are thought to require three body encounters. At one torr these events are relatively scarce compared to

binary collisions but still possibly numerous enough to allow ordinary flame reactions to proceed. Indeed, Wolfhard noted that in flames of carbon monoxide or hydrogen with oxygen there was a decrease in burning velocity with decreasing pressure which did not occur with hydrocarbons as fuels. He tentatively attributes this decrease in speed to the decrease in three body collisions but observed that another effect having to do with radiation could be involved.⁽¹⁰⁾

Flames clearly have fairly large populations of radiating species. Indeed, those that radiate in the visible portion of the spectrum are what make flames visible and attractive both esthetically and scientifically. The radiation lifetime of many of these species, especially those radiating in the infrared is much longer than the time between collisions at pressures as low as one torr. For example, radiation lifetimes for the various modes in CO, are all greater than three milliseconds. At one torr the time between collisions at flame temperatures is of the order of a microsecond. Thus, at that pressure the average CO_2 molecule undergoes several thousand collisions during its radiation lifetime, probably enough to insure thermalization of its internal energy. At substantially lower pressures, however, the possibility of radiation before thermalization becomes enhanced giving rise to a couple of provocative questions. Does the shift in mode of de-excitation for internally excited species make a difference in the structure and propagation characteristics of the flame? Is it possible that going to low pressures will greatly suppress collisional de-excitation so that population inversions capable of lasing might be achieved? It is to be remembered that in many exothermic reactions there is growing evidence that product molecules are formed in vibrationally or electronically excited states. It is this phenomenon which makes so-called chemical lasers possible. A possible advantage of flame systems for such lasers is that the reactants can be premixed because the relatively high activation

energy of the initial flame reactions prevents the premixed reactants from reacting until they enter the flame front which is a region of very rapid rise in temperature and/or the concentration of free radicals and chain carriers which cause the reaction to occur very quickly. The other requirement for achieving and maintaining the inversion necessary for lasing is that collisional deactivation of the excited species must not occur too rapidly. It is this latter requirement that low pressures could make possible in flame systems.

Of course, even if effective high power chemical lasers could be developed around flame systems by using the high pumping speed available in space, it might represent something of a pyrrhic victory because there might not be much use for them up there. Nevertheless, exploration of the possibility by spectroscopic studies of very low pressure flames would seem to offer the prospects of some interesting science.

3. <u>A Variation on the Low Pressure Flame Theme</u>. Flames generated by the combustion of solid propellants as used in rocket motors also offer opportunities for study at pressures low enough to require very large pumping speeds. These propellants are of two general classes. So-called colloidal or double base propellants are based on dispersions of nitrocellulose in nitroglycerine, i.e., very similar to the smokeless powder or cordite used in guns. More popular in rocket motors are the so-called composite propellants which comprise a granular oxidizer, usually ammonium perchlorate, embedded in a matrix of resin boinder which also is the fuel. These formulations are molded into large monolithic "grains," usually cylinders with an axial cavity whose size and shape are determined by the burning characteristics of the propellant and the ballistic requirements of the motor. In designing the grain for a particular motor and mission it is extremely important to know and if possible to control the burning rate of the propellant and its dependence upon pressure. Moreover,

there are a number of concomitant phenomena such as combustion instabilities with which are associated violent pressure oscillations which must be taken into account in the design of a motor. For these reasons there has been a great deal of experimental and theoretical study of solid propellant combustion. Even so, the theory of solid propellant burning is in a much more primitive state than the theory of flames in premixed gaseous systems. The difficulties are due in part to the much greater chemical complexity of solid propellant systems and in part to the complications stemming from the phase change as the solid reactants become gaseous products. Much of the experimental study of solid propellant combustion aimed at developing a theoretical understanding of the process is done with relatively small samples of the propellant, usually in the form of rectangular or cylindrical prisms called "strands". These strands are placed in a bomb in which the ambient pressure can be controlled independently of the combustion behavior. All strand surfaces but the uppermost one are coated with an inhibitor. The exposed surface is ignited and the deflagration rate, i.e., the linear regression rate of the burning surface into the substrate propellant, is monitored. Of primary interest is the dependence of this rate on the ambient gas pressure and the temperature and composition of the strand. It is well known that for any particular strand composition. and size, i.e., cross sectional area of the burning surface, there is a critical pressure below which a flame cannot be sustained. This so-called Low Pressure Deflagration Limit or LPDL is of some practical interest because of problems associated with extinguishing and re-igniting a solid propellant rocket motor at high altitudes. Of more fundamental interest is its use as a test of models for the combustion process. Any self respecting model should be able to predict and characterize the LPDL phenomenon.

Some years ago Roy Cookson and I carried out some experiments on LPDL in which we measured the pressure at which the flame was extinguished for a range

of strand sizes, i.e., areas of burning surface.⁽¹¹⁾ We found that the LPDL varied directly with the hydraulic radius of the strand, i.e., the ratio of perimeter to cross sectional area. Consequently, we were able to make a straight line extrapolation to a zero value of this hydraulic radius which would correspond to an infinite burning area. Somewhat surprisingly we found that at this limit the LPDL was finite. That is to say, the flame would go out at a finite pressure even for a burning surface of infinite area. Most theories indicate that the LPDL is due simply to a heat loss by radiation from the surface and the hot gases. Our results suggest that some other mechanism may be involved because at zero pressure and infinite area of burning surface the flame front would be infinitely thick. It is not entirely clear that radiative heat loss could be significant under these circumstances. It would seem likely that as the pressure decreases the accompanying decrease in the number of three body collisions and the increase in the ratio of radiation life time to time between collisions might well be expected to play an important role in solid propellant flames as well as in the premixed gaseous flames discussed in the previous section.

I hasten to point out that the pumping speed which was available was only about 10 liters/sec and was just sufficient to cope with a strand about 2.5 cm in diameter for which the LPDL was 40 torr. It is a very long extrapolation from 2.5 cm to infinity! Because the deflagration rate is inversely proportional to pressure and the gas density decreases directly with pressure, the pumping speed required per unit area of burning surface is roughly constant. But the burning area increases with the square of strand diameter. Consequently, the required pumping speed also goes up with the square of diameter. Thus, to go from 2.5 cm to 100 cm would require a pumping speed of something like 16 thousand liters a second at a pressure somewhere between 5 and 25 torr. That would seem to require a very large and expensive rig in which to dump very dirty hot gas.

By going to much larger strand sizes than Cookson and I were able to accommodate in our rig, one could test our extrapolation out to much larger sizes. In addition, of course, lowering the pressure would spread out the flame zone and make possible much more effective probing by spectroscopy and sampling probes. Thus, the SSE might lend itself to some interesting studies of solid propellant combustion.

4. On the Composition of the Space Shuttle Environment. One of the items on which there was unanimous agreement of Professor James Mayer's ad hoc Committee on Experiments in Space was the need to characterize as completely as possible the composition of the gas through which the Space Shuttle would be flying. In addition, for many experiments, it would be important to know the composition of the gas actually "in" the laboratory or the immediate environs of the space craft. The obvious way to determine chemical composition of low density gas is with a mass spectrometer and it would seem to be almost a routine problem in the case of "in lab" atmosphere. For the free stream gas, however, there are substantial sampling problems. Contamination by outgassing from the vehicle is a serious problem. Moreover, because many of the free stream species are atomic and can undergo reaction upon collision with a surface, it becomes very difficult to be sure that the measured ion currents are truly characteristic of free stream gas and do not include contributions from species which have collided with a surface somewhere on the vehicle or even in the instrument. For these reasons there is substantial uncertainty as to the true composition of the outer atmosphere in spite of the many measurements which have been made with instruments on board sounding rockets and balloons. A further reason for pursuing the problem of composition measurement is that the composition is not uniform in time or space. Consequently, one would like a lot of data taken at various times and places in order to obtain information on the scale and intensity of the fluctuations in composition in order to learn more about the structure of the atmosphere.

17

Constraint States of the second states of the secon

In the course of my visits in Europe I learned about two projects relating to mass spectrometry measurements on board the shuttle. Although both of them are probably well known to JPL-NASA I will mention them briefly here for the sake of completeness in reporting my discussions. The German Space Group have access to some space on one of the early flights which one of its component companies (Dornier or Messerschmidt, I believe) committed itself to "rent" or "buy" when NASA accepted bids a few years ago. They have also obtained a spare of the gold plated mass spectrometer which von Zahn had designed for the Mars mission. They are now in the process of designing and building the auxilliary gear which will adapt this instrument for use in their "rented" space. For reasons which are not clear to me, this adaptation involves a major electronics development which is to cost several hundred thousand marks. Everyone seems to recognize that the sampling problems will not be solved in this experiment. They claim justification on the basis that they will get some hands on experience in carrying out an experiment in space. In particular, they hope to find out what kinds of problems they will have to solve before they can contemplate obtaining truly reliable mass spectrometric analyses of the free stream atmosphere.

A more advanced project has been proposed to NASA by Berndt Feuerbacher of the Space Science Department of the European Space Agency (ESTEC) at Noordwijk in the Netherlands. In a proposal responsive to NASA AO-OSS-2-78 he outlines a method for discriminating between free stream gas and any distortions due to outgassing or surface interactions in mass spectrometric analysis. This discrimination is achieved by forming a molecular beam using two skimmers aligned with the velocity vector of the shuttle and analyzing the beam with an integrated time-of-flight quadrupole mass spectrometer. Because the vehicle flight velocity is much higher than the thermal velocity of most species, contaminants from outgassing or reflection from surfaces will have markedly different velocities

(both speed and direction) from the free stream species. Thus, fairly crude time-of-flight velocity analysis obtained by chopping the incoming beam can readily distinguish between free stream and contaminant species.

In my view this approach seems sound and I hope that NASA decides to support the proposed program. It should be noted that the proposed apparatus will also lend itself to experiments in which the incoming beam species can be deliberately scattered from various kinds of surfaces in order to study gas surface interactions. It has often been pointed out that the Shuttle can generate a reasonably intense highly collimated beam of 0 atoms with an energy of about 5 eV and a speed ratio of about 9 or 10. Such beams cannot now be readily produced in earthbound laboratories and would permit many kinds of interesting gas-gas and gas-surface scattering experiments. In particular gas-gas scattering experiments with hydrocarbons and other fuel molecules might provide much needed information on the chemistry of the flame combustion reactions in which 0 atoms are presumed to play an important role. Because these possibilities have been much discussed, I will not spend time and space in this report ruminating over them anew. I mention them at all by way of indicating that the Feuerbacher proposal would not be an isolated experiment but could be an important starting point for an extended series of interesting and worthwhile experiments.

There is one drawback to the use of mass spectrometers. They cannot readily provide information about the internal energy states of the species in the sample. In the lower atmosphere essentially all species are in the ground state. Because of its low density and exposure to the full intensity of solar radiation, the upper atmospheric gas can have substantial concentrations of internally excited species. Whether one wishes to carry out experiments in the SSE or whether one is concerned primarily with atmospheric structure and processes, it would be most valuable to know about the internal states of upper atmospheric species as well as their relative abundance. The availability of tunable lasers promises to make

possible the acquisition of data on internal energy states. Two laser methods which have been fairly extensively investigated and which seem applicable to the SSE conditions are laser induced fluorescence and laser multiphoton ionization. These are both very sensitive but, of course, require lasers tunable in the absorption range of the species of interest. The former works better when the fluorescence is in the near visible or shorter wavelength range of the spectrum because the photons are higher energy and more readily detectable and because the radiation lifetimes are usually short so that emission will occur while the excited species are still in the field of view of the detector. The latter requires very high photon densities and consequently samples a very small volume of gas because the laser beam must be focused to a small area. If tunable lasers become available in the uv portion of the spectrum, single photons will become energetic enough to cause ionization and the requirement for photon beam intensity will be relaxed and the sampling volume will be correspondingly increased.

There is a third possibility for using tunable lasers to identify internal energy states which has recently been developed by Professor G. Scoles and his colleagues at the University of Waterloo and which seems worthy of note here. Scoles has been a long time devotee of the use of super conducting bolometers as molecular beam detectors. Bolometers respond to the total energy content of the incident molecular beam. Thus, he finds that he can cross a dc molecular beam with a laser beam which is chopped and obtain an ac signal from his detector at the frequency of the laser beam chopping. That is, when a beam molecule absorbs a photon, it deposits the photon energy on the detector along with any other internal or translational energy which it possesses at the time of the absorption. These other energies are dc so that by appropriate adjustments of chopping frequency and phase his detector output indicates only the absorbed photons. He tunes the laser over a frequency range and obtains signal corre-

sponding to absorption when the laser frequency matches the absorption frequency of the molecule. In this way he has been able to do absorption spectroscopy (12.13)at the very low densities characteristic of molecular beams. Of course. the flight time of the excited molecule between the laser beam and the bolometer must be short relative to the radiation lifetime. This situation is quite characteristic of the infrared region of the spectrum so that his method becomes an admirable supplement to detection based laser induced fluorescence where long radiation lifetimes are a great handicap. Scoles estimates that with a bolometer sensitivity of 10^{-13} watts/hz^{1/2} he can detect fluxes of 3 x 10^6 molecules/sec photon absorption in the near to middle infrared. If one could achieve an equivalent sensitivity for a detector surface area of one cm² and if the shuttle flight velocity is 8 km/s, then one could detect about 40 excited molecules/cc of sampled gas. Even if the photon absorption cross sections are very small, the effective sensitivity would be extremely high. Of course, this method like all laser methods depends upon the availability of tunable lasers over the spectral range of interest, but the tunable range is increasing steadily. Already, with color center lasers and other solid state devices there is a very wide range of infrared already within reach.

Quite clearly, the advantages of the Scoles detection method lends itself to many of the kinds of spectroscopic experiments which were discussed in the original RDC report. It further enhances the prospects of many kinds of scattering experiments which might be done with beams extranced from the flux of atmospheric gas by the spacecraft. For these reasons I sincerely hope that Scoles will receive the support necessary to develop further this most ingenious application of bolometer detectors.

5. Microgravity and Coagulation in Turbulent Aerosols. The opportunities for experiments to take advantage of the microgravity condition in the SSE have been recognized from the beginning, especially in the field of fluid mechanics. Indeed in the European plans for space lab there is a whole module devoted to fluid mechanics experiments. One area of interest is in the effects due to surface tension which can become very important when gravitational forces become negligible. There are prospective practical implications of these effects because they can be very influential in the process of drawing single crystals from melts, for example. The surface tension of such melts is very high and in the presence of temperature gradients it can cause strong convection currents, i.e., the socalled Magangoni effect. At the DFVLR-AVA Institute at Goettingen I ran across some very interesting research on this problem. Dr. Ch.-H. Chun has developed a technique for studying surface tension effects in an earth bound system. He minimizes distortions due to gravity by going to very small masses (i.e., volumes) of liquid so that the surface tension forces are large relative to the gravitational forces. He suspends a drop of liquid between the ends of two vertical copper rods 3 mm in diameter. Each rod can be rotated in either direction over a range of speeds and the temperature of each rod can be independently controlled. He induces Marangoni convection by providing a downward temperature gradient through the liquid, i.e., making the top rod slightly warmer than the bottom rod. Dispersed in the fluid is particulate material which is illuminated by a vertical sheet of light passed through the vertical axis of the drop so that only the particles in the plane of the sheet are illuminated. With an enlarging lens and a TV camera he projects the illuminated plane on a screen which very clearly displays the flow streamlines. He finds he can markedly affect the flow patterns if he rotates one rod with respect to the other. In a counter rotating mode at an appropriate differential velocity the Marangoni flow cell is completely inhibited. These experiments

are being done in anticipation of studies to be performed in the Fluid Dynamics Module. Thus, they are already tuned in to the Shuttle Program and need no further elaboration here. I report them simply because they were to me beautiful and interesting.

More appropriate to the purpose of this report are some experiments being carried out by Professor Hiroshi Sato at the Institute of Space and Aeronautical Studies at the University of Tokyo. He is addressing himself to the problem of how the pullence may affect the growth of droplets in a gas undergoing condensation. He mixes a stream of moist air with a stream of colder air and produces a stream of air with small droplets in it, i.e., a stream of fog. He then passes that stream vertically through grids of various sizes in order to induce varying degrees of turbulence. At various distances above the grid he probes the foggy stream with a hot wire in an anemometer configuration. The wire cools and gives a signal via resistance change when it is struck by a droplet. The size of the signal pulse is proportional to the size of the droplet. The frequency of pulses indicates the number density of droplets. Thus he obtains a measure of growth rate as a function of time by knowing the velocity of flow and comparing probe signal patterns at various heights in the stream. The apparatus has not yet been perfected but meaningful results are beginning to accumulate.

Professor Sato is convinced that an understanding of droplet agglomeration and growth by turbulence has important and far reaching implications. He seems particularly intrigued by the effects this phenomena may have had upon the composition of the planets during the condensation and accretion from initial primordial gas. He even thinks that it may have something to do with the fact that all planets rotate in the same direction. Of course, the coagulation of solid colloidal particles in liquids and the effects of stirring on the coagulation process have been studied for a long time. Smoluchowski of kinetic theory fame even did some theoretical anal-

ysis of the process many years ago. There seems to have been very little if any work done on gaseous dispersions of liquids. The diminution of gravitational effects which could be achieved in the SSE is obvious. Experiments like those which Sato is trying can deal only with particles small enough so that drag forces will keep them suspended *i*⁻ the gas. It is not entirely clear just how such experiments might be designed and carried out on the shuttle but they certainly would seem to deserve some careful consideration. Professor Sato himself has a great interest in pursuing this kind of work and assured me that he would be happy to cooperate in any project which might be undertaken in anticipation of performing such experiments.

III. Conclusions and Recommendations

I talked to many people from many disciplines. The response was somewhat less than overwhelming but I did glean a few ideas which seem to merit further consideration and analysis. Moreover, I was able to identify some capable investigators who are not now associated with a space program but who are interested in pursuing possibilities for extending their studies to take advantage of the Space Shuttle Environment:

- 1. Professor John C. Polanyi of the Department of Chemistry at the University of Toronto is keenly interested in high resolution spectroscopy of species native to or introduced into the upper reaches of the atmosphere.
- 2. Professor Marcel Vanpee of the Department of Chemical Engineering at the University of Massachusetts would be very interested in a spectroscopic study of ephemeral but kinetically important species in low pressure flames and tube reactors taking advantage of the high speed pumping that SJE offers.
- 3. Professor Giacinto Scoles of the Department of Chemistry at the University of Waterloo in Waterloo, Ontario is eager to develop his laser-bolometer detector system for use in the analysis of atmospheric species and the determination of their internal energy levels.

4. Professor Hiroshi Sato of the Institute of Space and Aeronautical Sciences wants very much to cooperate in any project which might extend his study of aerosol coagulation by turbulence, possibly taking advantage of the low gravity forces in orbiting laboratories.

If NASA gets to the point where it can seriously entertain proposals for research projects toward any of these goals, it would do well to get in touch with these investigators.

IV. References

- J. B. Fenn, S. B. Ryali and M. P. Sinha, Final Report on JPL Contract, #954327, Relay Development Corp., (1976).
- 2. R. A. Outlaw and F. J. Brock, J. Vac. Sci. & Tech. <u>14</u>, 169 (1977).
- L. T. Melfi, R. A. Outlaw, J. E. Hueser and F. J. Brock, J. Vac. Sci. and Tech., <u>13</u>, 698 (1976).
- 4. T. F. George, I. H. Zimmerman, P. L. DeVries, J. M. Yuan, K. S. Lam, J. C. Bellum, H. W. Lee, M. S. Slutsky and J. T. Liu in <u>Chemical and</u> <u>Biochemical Applications of Lasers</u>, Ed. C. B. Moore (Academic Press, New York), Vol. IV, 1976.
- 5. A. E. Orel and W. H. Miller, Chem. Phys. Letters <u>57</u>, 362 (1978).
- 6. H. G. Wolfhard, Z. Tech. Phys. 24, 206 (1943).
- 7. A. G. Gaydon and H. G. Wolfhard, Fuel 29, 15 (1950).
- 8. E. C. Hinck, T. S. Seamans, M. Vanpee and H. G. Wolfhard, 10th Intl. Symposium Combustion, Proceedings, Williams and Wilkinson (1965).
- 9. Private Communication.
- 10. Private Communication.
- 11. R. A. Cookson and J. B. Fenn, AIAA J., 8, 864 (1970).
- 12. T. E. Gough, R. Miller and G. Scoles, Appl. Phys. Lett <u>30</u>, 338 (1977).
- 13. T. E. Gough, R. Miller and G. Scoles, J. Chem. Phys. <u>69</u>, 1588 (1978).
- 14. Ch.-H. Chun, "Marangoni Convection in Floating Zone Under Reduced Gravity Effect", Proceedings 2nd European Conference of Crystal Growth, Sept. 1979.

APPENDICES

"Spreading the Word"

This section attempts to provide some dimension to the extent of my evangelistic activities. Section A comprises a list of all those with whom I have had direct conversations in the past year on possible SSE experiments. In addition to these "person-to-person" encounters, I also described the opportunity and the advantages of the spacecraft environment for experiments to audiences of various sizes in the course of lectures and seminars presented during my travels. On three occasions I made organized presentations on the subject to relatively large groups. At the 1978 Conference on Dynamics of Molecular Collisions at Asi lomar, California in June 1978 and at the Eleventh International Symposium on Karefied Gas Dynamics at Cannes in July 1978, I made oral presentations with slides and transparencies. These meetings were prior to the term of the contract for which this document is the final report, but they were subsequent to the report on the original RDC study. Therefore, to indicate the extent of exposure to the idea of Space Shuttle experiments, Section B is an attendance list for the Asilomar Conference and Section C is a list of participants at the Cannes meeting.

The third occasion for a selected audience was the International Symposium on Molecular Beams which was at Riva del Garda, Italy in June 1979. At that meeting the message was presented in a Poster Session which remained on display throughout the week of the symposium. During scheduled Poster Sessions I was on hand for discussion. Unfortunately, I do not have an attendance list for that meeting but I am including a program in Section D. Because most scientists have to present papers if they are to qualify for travel support from their sponsors, this program provides a fairly representative census of those in attendance. Needless to say it also includes authors who stayed home, but maybe they got the word by hearsay from their colleagues.

APPENDIX A

People Engaged in Direct Conversation

(Numbers in parentheses refer to key for Fields of Interest at end of II-A)

Akamatsu, Professor Teruaki Dept. of Mechanical Engr. Kyoto University Yoshida Honmachi, Sakyo-ku, Kyoto

Anderson, Professor J. B.(4,11,23,10,70) Dept. of Chemistry Pennsylvania State Univ. University Park, Pa. 16802

Andres, Professor R. P.(4,11,14,70,71) Dept. of Chemical Engr. Princeton Univ. Princeton, NJ

Balasubramanian, Professor D.(22,23,37,47) Dept. of Chemistry Univ. of Hyderabad Hyderabad, India

Becker, Professor E. W. (4,11,69,70) Institut f. Kernverfahrenstechnik Kernforschungzentrum D-7500, Karlsruhe, W. Germany

Bernstein, Professor R. B.(6,11,21,23) Dept. of Chemistry Columbia University New York, NY 10027

Bose, Professor T. K.(24,26) Aeronautical Engr. Dept. Indian Inst. of Technology Madras, India

Brooks, Professor Philip R.(6,11,21,23) Chemistry Dept. Rice University Houston, TX 77001

Brusdeylins, Dr. G. (4,11,12,13) Max Planck Inst. f. Strömingsforsch. Postfach 867, Böttingerstrasse 4/8 3400 Göttingen, W. Germany Campargue, Dr. Roger(4,11,70) C.E.N. Saclay, SCPH Lab. Jets Moleculaires BP 2, 91190, Gif-sur-Yvette, France

Cassuto, Dr. A. (5,17,18,19,34) CNRS Laboratoire M. Letort Rt. de Vandoeuvre 54600 Villers, Nancy, France

Chandrasekahr, Professor (35,45) Liquid Crystal Group Raman Inst. Bangalore, India

Chun, Dr. C. H. DFVLR-AVA Bunsenstrasse 10 3400 Göttingen, W. Germany

Chupka, Professor William(6,23,37,69) Dept. of Chemistry Yale Univ. New Haven, CT 06520

Cross, Professor R. J., Jr.(6,12,11,23) Dept. of Chem. Yale Univ. New Haven, CT 06520

Deshpande, Professor S.M.(24,26,27) Aeronautical Engr. Indian Inst. of Science Bangalore, India

deVries, Dr. A. E. (6,11,12,21) FOM Inst. for Atomic & Molec. Physics Kruislaan 407 Amsterdam-Oost, The Netherlands

Ding, Dr. A. (11,12,37,68) Hahn-Meitner Inst. f. Kernforschung Berlin GMBH D-100 Berlin 39, W. Germany

A. .-3

Ehrfeld, Dr. W. (4,11,69,70) Institut f. Kernverfahrenstechnik Kernforschungszentrum D-7500, Karlsruhe, W. Germany

Faubel, Dr. M. (6,11,12,23) Max Planck Inst. f. Strömungsforsch. Postfach 867, Böttingerstrasse 4/8 3400 Göttingen, W. Germany

Feuerbacher, Dr. Berndt (11,18,19,69) Space Science Dept. European Space Agency, ESTEC Noordwijk, The Netherlands

Franck, Professor Ulrich (22,16,22,37) Inst. for Physical Chemistry Univ. of Karlsruhe Karlsruhe, W. Germany

Fujiwara, Professor Toshi (2,24,20) Dept. Aeronautical Engr. Nagoya University Furo-cho, Chikusa-ku, Nagoya

Fukuda, Professor Kuniya (20,23,37,54) Dept. of Engr. Physics Kyoto University Yoshida Honmachi, Sakyo-ku, Kyoto

Gentry, Professor W. R. (6,12,11,20,21) Chemistry Dept. Univ. of Minnesota Minneapolis, MN 55455

Govinda-Raju, Professor G. R.(55,56,57,59) High Voltage Engr. Indian Inst. of Science Bangalore, India

Gowariker, Dr. V. R. (25,26) Chemicals and Materials Group Vikram Sarabhai Space Center Trivandrum, India

Greene, Professor E. F. (2,6,11,18) Dept. of Chemistry Brown University Providence, RI 02912

Grover, Dr. James R. (11,23,72) Dept. of Chemistry Brookhaven National Lab Upton, NY 11973

Gspann, Dr. J. (4,11,70,71) Institut f. Kernverfahrenstechnik Kernforschungszentrum D-7500, Karlsruhe, W. Germany Hagena, Dr. Otto (11,14,70,71) Inst. f. Kernverfahrenstechnik Kernforschungszentrum D-7500, Karlsruhe, W. Germany

Herschbach, Professor D. R. (5,6,11,20,21,2) Dept. of Chemistry Harvard University Cambridge, MA 02138

Hishinuma, Professor Naoshi (11,68) Inst. of Physics College of General Education, Univ. of Tokyo Komaba, Meguro-ky, Tokyo

Hulpke, Dr. G. (11,18,19) Max Planck Inst. f. Strömungsforsch. Postfach 867, Böttingerstr. 4/8 3400 Göttingen, W. Germany

Jain, Professor S. R. (29,30,31) Aeronautical Engr. Indian Inst. of Science Bangalore, India

Katayama, Professor Mikio (23,37,21) Dept. of Pure & Applied Science College of Gen. Education., Univ. of Tokyo Komaba, Meguro-ku, Tokyo

Kaufman, Professor F. (5,6,23,37,49,66) Dept. of Chemistry Univ. of Pittsburgh Pittsburgh, Pa. 15261

Kitagawa, Professor Taiji (5,11,12,37) Faculty of Pharmaceutical Sciences University of Medical and Pharm. Sci. Toyama

Kobayashi, Professor Yasunori (3,4) Inst. of Applied Physics Univ. of Tsukuba Sakura, Niihari, Ibaraki 300-31, Japan

Kodera, Professor K. (Emeritus) Dept. of Chemistry Kyoto University Yoshida Honmachi, Sakyo-ku, Kyoto

Kolb, Dr. C. E. (5,6,11,12,23,37) Aerodyne Research Inc. Crosby Dr. Bedford, MA 01730

Kondow, Dr. Tamotsu (22,23,65,66,67) Dept. of Chemistry Univ. of Tokyo Hongo, Bunkyo-ku, Tokyo

Kotake, Professor Susumu (11,14,32) Inst. of Space & Aeronautical Sci. Univ. of Tokyo Komaba, Meguro-ku, Tokyo

Kuchitsu, Professor Kozo (22,23,65,66,67) Murakami, Dr. M. (10,15,32,64) Dept. of Chemistry Univ. of Tokyo Hongo, Bunkyo-ku, Tokyo

Kumar, Professor R. (41, 42, 43)Chemical Engr. Dept. Indian Inst. of Science Bangalore, India

Koppenwallner, Dr. G. (4,1,18)DFVLR-AVA Bunsenstrasse 10 3400 Göttingen, W. Germany

Kupperman, Professor Aron (5,6,11,65) Dept. of Chem. California Inst. of Technology Pasadena, CA 91125

Kuratani, Professor Kenji (24,25,26,20) Inst. of Space & Aeronautical Sci. Univ. of Tokyo Komaba, Meguro-ku, Tokyo

Lai, Professor Hon-Ming (23,46,58) Dept. of Physics Chinese University of Hong Kong Shatin, N.T., Hong Kong

Lee, Professor Y. T. (5,6,11,21,69) MMRD - 70A - 4414Lawrence Berkeley Labs Berkeley, CA 94720

Legge, Dr. H. (4,11,15) DFVLR-AVA Bunsenstrasse 10 3400 Göttingen, W. Germany

Leung, Professor Alfred F. (35,37) Dept. of Physics The Chinese Univ. of Hong Kong Shatin, N.T., Hong Kong

Lang, Dr. H. (4,9,12,15) Max Planck Inst. f. Strömungsforsch. Postfach 867, Böttingerstr. 4/8 3400 Göttingen, W. Germany

Miller, Professor D. R. (4,6,11,13,18) Dept. of Applied Mech. & Engr. Sci. University of California San Diego, Ca.

Inst. of Space & Aeronautical Sci. Univ. of Tokyo Komaba, Meguro-ku, Tokyo

Oguchi, Professor Hakuro (2,3,4,9,10) Inst. of Space & Aeronautical Sciences Univ. of Tokyo Komaba, Meguro-ku, Tokyo

Onji, Dr. A. (2,4,11,18) National Aerospace Laboratory Mitaka, Tokyo

Oshima, Professor Koichi (1,2,3,4,9,32) Inst. of Space & Aeronautical Sciences Univ. of Tokyo Komaba, Meguro-ku, Tokyo

Nair, Dr. K. Narayanan (52,53,58) Space Physics Division Vikram Sarabhai Space Center Trivandrum

Narasimha, Professor R. (1,2,3,4,28) Aeronautical Engr. Indian Inst. of Science Bangalore, India

Polanyi, Professor J. C. 262 Lash Miller Chem. Lab. Univ. of Toronto Toronto, Ontario, M5S 1A1

Prasad, Professor K. K. (2,3,4,44) Mechanical Engr. Dept. Indian Inst. of Science Bangalore, India

Raghunandan, Dr. B. N. (24,26,32) Aeronautical Engr. Indian Inst. of Science Bangalore, India

Rao, Professor C. N. R. (34,36,37,38) Solid State and Structural Chem. Indian Inst. of Science Bangalore, India

Rao, Professor K. H. (36,39,37,40) Solid State & Structural Chemistry Indian Inst. of Science Bangalore, India

Rao, Dr. P. V. M. (50,51,52,59) Space Physics Div. Vikram Sarabhai Space Center Trivandrum, India

Rao, Dr. U. R. (60,61,62) Director ISRO Satellite Center Peenya, Bangalore, India

Rebrov, Professor A. K. (4,13,14) Inst. of Thermophysics Novosibirsk-90 USSR

Reddy, Dr. C. A. (49,50,51) Space Physics Division Vikram Sarabhai Space Center Trivandrum, India

Reuss, Professor J. (6,11,14,20,71) Fys. Lab. Kath. Univ. Toernooiveld, Driehuizerweg 200 Nijmegen, The Netherlands

Riley, Professor S. J. Chemistry Dept. Yale Univ. New Haven, CT 06520

Rothe, Professor E. W. (5,6,11,21,67) RIES, College of Engineering Wayne State University Detroit, MI 48202

Sano, Professor Taeko (3,11,14) Inst. of Industrial Science Tokai University Kita-kaname, Hiratsuka, Japan

Sato, Professor Hiroshi (1,3,27,28) Inst. of Space & Aeronautical Sciences Univ. of Tokyo Komaba, Meguro-ku, Tokyo

Schumacher, Professor Ernst (6,11,20,23) Inst. for Physical Chemistry Univ. of Bern Bern, Switzerland

Scoles, Professor G. (11,18,20) Dept. of Chemistry Univ. of Waterloo Waterloo, Ontario Professor A. K. Srikante (1,3,4) Dept. of Aeronautical Engineering Indian Institute of Science Madras, India

Takayanagi, Professor Kazuo (9,12,13,54) Inst. of Space & Aeronautical Sciences Univ. of Tokyo Komaba, Meguro-ku, Tokyo

Teshima, Dr. Koji (2,11,18) Dept. of Aeronautical Engr. Kyoto University Yoshida Honmachi, Sakyo-ku, Kyoto

Toennies, Professor J. P. (2,4,11,12,23) Max Planck Inst. f. Strömungsforschung Postfach 867, Böttingerstrasse 4/8 3400 Göttingen, W. Germany

Tsuchita, Professor Soji (12,13,21,22) Dept. of Pure & Applied Sciences College of General Education, Univ. of Tokyo Komaba, Meguro-ku, Tokyo

Vanpee, Professor Marcel (24,25,37) Dept. of Chemical Engr. Univ. of Massachusetts Amherst, Mass.

Vasagam, Dr. R. M. (32,33) Director, Ariane Passenger Payload Exp. ISRO Satellite Center Peenya, Bangalore, India

Venugopalan, Dr. S. (37,45) Liquid Crystal Group Raman Institute Bangalore, India

Wharton, Professor Lennard (6,11,18,20,23) James Franck Inst., Univ. of Chicago 5640 S. Ellis Ave. Chicago, IL 60637

Wolfhard, Dr. H. G. (23,24,25,37) Inst. for Defense Analysis 400 Army Navy Drive Arlington, Va.

Yamashita, Dr. M. (11,14,20) Inst. of Space and Aeronautical Sciences Univ. of Tokyo Komaba, Meguro-ku, Tokyo

Yamazaki, Dr. Satoshi (2,5,13,11) Toyota Central Res. & Development Labs. Hisakata, Tempaku-ku, Nagoya, 408

Yoshikawa, Professor Kenneth K. (4,9,10) Inst. of Applied Physics Univ. of Tsukuba Sakura, Niihari, Ibaraki 300-31, Japan

Zare, Professor R. N. (6,11,18,20,21,37,70) Dept. of Chemistry Stanford University Stanford, CA 94305

Field of Interest Key

- 1. Fluid Mechanics
- 2. Shock Tubes
- 3. Gas Dynamics
- 4. Rarefied Gas Dynamics
- 5. Chemical Kinetics
- 6. Molecular Dynamics
- Thermodynamics
 Statistical Mechanics
- 9. Kinetic Theory
- 10. Monte Carlo Methods
- 11. Molecular Beam Experiments
- 12. Molecular Energy Transfer
- 13. Relaxation Rutes
- 14. Nucleation-Condensation
- 15. Vaporization Studies
- 16. Physical Organic Chemistry
- 17. Heterogeneous Catalysis
- 18. Molecule Surface Scattering
- 19. Surface Physics
- 20. Laser Spectroscopy
- 21. Laser Induced Chemistry
- 22. Physical Chemistry
- 23. Chemical Physics
- 24. Combustion
- 25. Liquid Rocket Propellants
- 26. Solid Rocket Propellants
- 27. Propulsion Dynamics
- 28. Turbulence
- 29. Astronautics
- 30. Orbital Mechanics
- 31. Satellite Trajectory Analysis
- 32. Hear Transfer
- 33. Molecular Structure
- Surface Chemistry
 Solid State Physics

- 36. Solid State Chemistry
- 37. Spectroscopy
- 38. Structural Chemistry
- 39. Amorphous Materials
- 40. Solution Chemistry
- 41. Chemical Engineering

 - 42. Chemical Process Dynamics & Control43. Chemical Reaction Engineering & Reactor Design
- 44. Gas Dynamic Lasers
 - 45. Liquid Crystals
 - 46. Theoretical Physics
 - 47. Biochemistry
 - 48. Biophysics
 - 49. Meterology
 - 50. Ionospherics
 - 51. Radio Wave Propagation
 - 52. Geomagnetics
 - 53. Ultrasonic Acoustics
 - 54. Plasma Physics
 - 55. Power Transmission
 - 56. High Field Phenomena
 - 57. Dielectric Materials
 - 58. Properties of Materials
 - 59. Electrostatics 60. Astro Physics
 - 61. UV Astronomy
 - 62. X-Ray Astronomy
 - 63. Electronics
 - 64. Heat Pipes

- 65. Low Energy Electron Scattering
- 66. Photochemistry
- 67. Excited Atom Chemistry
- 68. Charge Exchange
- 69. Mass Spectrometry
- 70. Isotope Enrichment
- 71. Molecular Clusters
- 72. Synchroton Radiation Spectroscopy

APRIAIIA - B

Participants 1978 Conference on Dynamics of Molecular Collisions Asilomar, California June 26-30, 1978

Dr. O. William Adams Dept. of Energy Washington, DC 20545

Professor Millard H. Alexander Dept. of Chemistry University of Maryland College Park, Maryland 20742

Prof. James B. Anderson 152 Davey Laboratory Dept. of Chemistry Pennsylvania State Univ. University Park, PA 16802

Dr. Roger W. Anderson Natural Sciences II University of California Santa Cruz, CA 95064

Brian Kitfield Annis Oak Ridge National Lab. P. O. Box X Oak Ridge, Tenn. 37830

Prof. Vincenzo Aquilanti Dipartimento di Chimica Universita 'degli Studi 06100 Perugia (ITALY)

Prof. Daniel J. Auerbach Dept. of Chemistry Johns Hopkins University Baltimore, MD 21218 Dr. Michael Baer Soreq Nuclear Research Center Yavne, ISRAEL

Prof. Thomas L. Bailey Dept. of Physics University of Florida Gainesville, Florida 32601

Dr. Andre D. Bandrauk 103 Lewis Hall University of California Berkeley, CA 94720

Dr. John R. Barker SRI International Menlo Park, CA 94025

Mr. Chris Becker MMRD - 70A-4429 Lawrence Berkeley Lab. Berkeley, CA 94720

Prof. R. B. Bernstein Dept. of Chemistry Columbia University New York, NY 10027

Dr. Normand C. Blais MS-732 Los Alamos Scientific Lab. Los Alamos, NM 87545

Dr. Stephen M. Bobbio Chemistry Dept. University of Georgia Athens, Georgia 30602 Dr. Dietrich Brandt University of Toronto Chemistry Dept. 80 St. George St. Toronto, CANADA

Prof. Philip R. Brooks Chemistry Dept. Rice University Houston, TX 77001

Dr. John H. Brophy Room 2-045, Chemistry Dept. M.I.T. Cambridge, Mass. 02139

Nancy J. Brown E & E Division Lawrence Berkeley Laboratory Berkeley, CA 94720

Dr. Brunetto Brunetti Dipartimento Di Chimica Dell'Universita-Via Elce Di Sotto, 06100 Perugia (ITALY)

Mr. Richard Buss MMRD - 70A-4425 Lawrence Berkeley Laboratory Berkeley, CA 94720 Dr. Piergiorgio Casavecchia MMRD - 70A-4429 Lawrence Berkeley Laboratory Berkeley, CA 94720

Miss Sylvia Ceyer MMRD - 70A-4418 Lawrence Berkeley Laboratory Berkeley, CA 94720

Prof. Sally Chapman Dept. of Chemistry Barnard College Columbia University New York, NY 10027

Dr. David C. Clary IBM Research Laboratory K34/281, 5600 Cottle Road San Jose, CA 95193

Dr. Michael J. Coggiola SRI International Menlo Park, CA 94025

Dr. Ronald B. Cohen Aerospace Corporation P. O. Box 92957 Los Angeles, CA 90009

Dr. J.N.L. Connor Dept. of Chemistry University of Manchester Manchester, M13 9PL, ENGLAND

Dr. Robert D. Coombe Science Center Rockwell International 1049 Camino Dos Rios Thousand Oaks, CA 91360 Dr. William H. Cramer Chemistry Division National Science Foundation 1800 G St. NW Washington, DC 20550

Prof. R. James Cross, Jr. Dept. of Chemistry Yale University New Haven, Conn. 06520

Dr. John B. Cross CNC-2, MS-732 Los Alamos Scientific Lab. Los Alamos, NM 87545

Prof. Paul J. Dagdigian Dept. of Chemistry The Johns Hopkins University Baltimore, Maryland 21218

Dr. Sheldon Datz Oak Ridge National Laboratory P. O. Box X Oak Ridge, Tenn. 37830 Dr. Adalbert Ding Hahn-Meitner-Institut fur Kernforschung Berlin GmbH D-1000 Berlin 39, GERMANY

Dr. Thomas H. Dunning Argonne National Laboratory Chemistry Division Argonne, Illinois 60439

Dr. Rudolf M. Duren Max-Planck-Institut fur Stromungsforschung Bottingerstr.4-8 D3400 Gottingen, WEST GERMAN

Dr. Marie Durup Lab. de Resonance electron. et ionique, Univ. de Paris 91405 Orsay, FRANCE Dr. Michael B. Faist Chemistry Dept. Brookhaven National Laboratory Upton, NY 11973

Prof. James M. Farrar Dept. of Chemistry Univ. of Rochester Rochester, NY 14627

Prof. John B. Fenn Mason Laboratory Yale University New Haven, Conn. 06405

Dr. Andrew Freedman Havemeyer Hall Columbia University New York, NY 10027

Prof. Jean H. Futrell Dept. of Chemistry University of Utah Salt Lake City, Utah 84112 Dr. Bruce C. Garrett Dept. of Chemistry Univ. of Minnesota Minneapolis, MN 55455

Prof. W. R. Gentry Chemistry Dept. University of Minnesota Minneapolis, MN 55455

Dr. Dieter Gerlich MMRD - 70A-4405 Lawrence Berkeley Laboratory Berkeley, CA 94720

Dr. Anna Giardini-Guidoni C.N.E.N. Divisione N. Attivita C.P. 65 - 00044 Frascati (Rome), ITALY

Prof. Clayton F. Giese Tate Laboratory of Physics University of Minnesota Minneapolis, MN 55455

Dr. Robert G. Gilbert Dept. of Theoretical Chem. University of Sydney N.S.W. 2006, AUSTRALIA

Dr. Keith T. Gillen Molecular Physics Laboratory SRI International Menlo Park, CA 94025

Dr. Graham P. Glass Chemistry Dept. Rice University Houston, TX 77001 Dr. David M. Golden SRI International Menlo Park, CA 94025

Prof. Robert J. Gordon Dept. of Chemistry University of Illinois Chicago, Illinois 60680

Dr. Thomas R. Govers Lab. de Resonance Electronique et Ionique Univ. Paris-SUD 91405 Orsay, FRANCE

Prof. E. F. Greene Chemistry Dept. Brown University Providence, RI 02912

Dr. James R. Grover Dept. of Chemistry Brookhaven National Lab. Upton, New York 11973

Dr. Hellmut Haberland Fakultat fur Physik der Universitat Freiburg Hermann-Herder-Strasse 3 7800 Freiburg, WEST GERMANY

Prof. William L. Hase 35 Chemistry Wayne State University Detroit, MI 48202

Dr. Raymond F. Heidner The Aerospace Corp. Bldg. 130 - Room 145 El Segundo, CA 90245

Dr. Irving Philip Herman 155 Monte Cresta Ave. #203 Oakland, CA 94611

Prof. Dudley Herschbach Dept of Chemistry Harvard University Cambridge, Mass. 02138

Prof. Peter Hess Dept. of Chemistry University of California Berkeley, CA 94720

Prof. Peter M. Hierl Chemistry Department Kansas University Lawrence, Kansas 66045

Dr. Tomohiko Hirooka MMRD - 70A-4418 Lawrence Berkeley Laboratory Berkeley, CA 94720 Prof. William M. Jackson Dept. of Chemistry Howard University Washington, DC 20059

Prof. F. Kaufman Department of Chemistry University of Pittsburgh Pittsburgh, PA / 526/

Prof. James L. Kinsey Room 6-229 M.I.T. Cambridge, MA 02139

Prof. Fritz S. Klein Isotope Dept. Weizmann Institute Rehovot, ISRAEL

Prof. Kumasaburo Kodera 31-2 Shimogamokitazonocho Sakyoku, Kyoto 606 JAPAN Dr. Herbert F. Krause Oak Ridge National Lab. 4500 W, F-17 Oak Ridge, TN 37830

Prof. John R. Krenos Dept. of Chemistry Douglass College Rutgers University New Brunswick, NJ 08903

Prof. Aron Kuppermann Dept. of Chemistry (127-72) California Inst. of Technology Pasadena, CA 91125

Dr. George H. Kwei Mail Stop 732 Los Alamos Scientific Lab. Los Alamos, NM 87545

Dr. Neil Lang Lawrence Livermore Laboratory Livermore, CA 94550

Prof. Y. T. Lee MMRD - 70A-4414 Lawrence Berkeley Laboratory Berkeley, CA 94720

Prof. Stephen R. Leone JILA, Nat. Bureau of Standards University of Colorado Boulder, Colorado 80309

Dr. William Lester NRCC Lawrence Berkeley Laboratory Berkeley, CA 94720

Prof. Jacob J. Leventhal Dept. of Physics University of Missouri St. Louis, MO 63121

Dr. M. C. Lin Code 6110 Naval Research Laboratory Washington, DC 20375

Dr. Hans J. Loesch Universitat Bielefeld Fakultat fur Fhysik Universitatsstrasse D-4800 Bielefeld 1, W GERMANY

Prof. Dr. J. Los FOM-Inst for Atomic & Molecular Physics Kruislaan 407, Amsterdam THE NETHERLANDS Dr. Alan C. Luntz IBM Research Laboratory 5600 Cottle Road San Jose, CA 95193

Dr. Chang-Chi Mei Dept. of Chemistry University of Calif. Berkeley, CA 94720

Prof. C. Bradley Moore Dept. of Chemistry University of Calif. Berkeley, CA 94720

Dr. John Moseley Molecular Physics-106B SRI Menlo Park, CA 94025

Dr. James T. Muckerman Chemistry Dept. Brookhaven Nat. Laboratory Upton, NY 11973

Prof. M. Menzinger Lash Miller Chemical Laborator: 80 St. George Street Toronto, CANADA, M5S 1A1

Univ. of Notre Dame Notre Dame, Indiana 46556

Miss Elizabeth R. Manzares

Radiation Laboratory

Prof. R. A. Marcus 174 Noyes Lab. University of Illinois Urbana, Illinois 61801

Dr. Raymond P. Mariella Jr. Allied Chemical Co., MRC P. O. Box 1021R Morristown, NJ 07960

Prof. Richard M. Martin Dept. of Chemistry University of California Santa Barbara, CA 93106 Prof. J. Douglas McDonald 366 Noyes Laboratory University of Illinois Urbana, Illinois 61801

Dr. Charles A. McDowell Dept. of Chemistry University of British Columbia 2075 Wesbrook Place Vancouver, BC, CANADA

Prof. David McFadden Dept. of Chemistry Boston College Chestnut Hill, MA 02167 Dr. Jean B. Ozenne 3192 Bryant St. Palo Alto, CA 94306 Dr. James R. Peterson Molecular Physics Lab-106B SRI International Menlo Park, CA 94025

Prof. Norman C. Peterson Polytechnic Institute 333 Jay Street Brooklyn, NY 11201

Prof. Leon F. Phillips Department of Chemistry University of Canterbury Christchurch, NEW ZEALAND

Prof. J. C. Polanyi 262 Lash Miller Chem. Lab. University of Toronto Toronto, Ontario, CANADA

Dr. Eli Pollak Dept. of Chemistry Columbia University New York City, NY 10027

Dr. Lise Lotte Poulsen Institute for Chemistry University of Copenhagen Copenhagen, DK-2200 N, DENMARK

Dr. Jack M. Preses Division of Basic Energy Sci. U.S. Dept. of Energy Washington, DC 20545

Prof. David E. Pritchard Room 26-231 M.I.T. Cambridge, Mass 02139

Dr. Robert K. Nesbet IBM Research Laboratory San Jose, CA 95193

Dr. Roy H. Neynaber IRT Corporation 7650 Convoy Court San Diego, CA 92111

Dr. Cheuk-Yiu Ng Dept. of Chemistry 'Iowa State University Ames, IA 50011

Dr. Nikitin Institute of Chemical Physics Academy of Sciences Moscow, V-334, U.S.S.R. Dr. Eric K. Parks Bldg. 200 Argonne National Laboratory Argonne, Illinois 60439

Dr. Christopher A. Parr Univ. Texas at Dallas BE 2.6, P.O. Box 688 Richardson, TX 75080

Prof. John M. Parson Chemistry Dept, OSU 140 W. 18th Avenue Columbus, OH 43210

Prof. Dr. Hans Pauly Max-Planck-Institute Bottingerstr 4-8 D3400 Gottingen, WEST GERMANY Dr. Alfred T. Pritt Rockwell International P. O. Box 1085 Thousand Oaks, CA 91360

Dr. J. Gary Pruett Dept. of Chemistry University of Pennsylvania Philadelphia, PA 19104

Dr. Martin Quack Institut fur Physikalische Chemie, der Universitat Tammanstrasse 6 3400 Gottingen, WEST GERMANY Dr. Lionel M. Raff Dept. of Chemistry Oklahoma State University Stillwater, Oklahoma 74074

Prof. Gene P. Reck 35 Chemistry Wayne State University Detroit, MI 48202

Prof. Stephen J. Riley Chemistry Department Yale University New Haven, Conn. 06520

Prof. John Ross Room 6-123 M.I.T. Cambridge, Mass 02139

Prof. Erhard W. Rothe RIES, College of Engineering Wayne State University Detroit, MI 48202 Prof. George C. Schatz Dept. of Chemistry Northwestern University Evanston, IL 60201

Prof. Christoph Schlier Fakultat fur Physik der Universitat 78 Freiburg, WEST GERMANY

Mr. Peter A. Schulz MMRD - 70A-4418 Lawrence Berkeley Laboratory Berkeley, CA 94720

Prof. G. Scoles Facolta di Scienze Universita di Trento Povo(Trento) 1TALY

Prof. H. K. Shin Dept. of Chemistry Univ. of Nevada Reno, Nevada 89557

Dr. Kosuke Shobatake MMRD - 70A-4435 Lawrence Berkeley Laboratory Berkeley, CA 94720

Dr. Joel A. Silver Aerodyne Research Inc. Crosby Drive Bedford, Mass. 01730

Prof. Peter A. Siska Dept. of Chemistry Univ. of Pittsburgh Pittsburgh, PA 15260 Prof. James G. Skofronick Dept. of Physics Florida State University Tallahassee, Florida 32306

Dr. James J. Sloan National Research Council Division of Chemistry 100 Sussex Drive Ottawa, CANADA

Dr. Thompson M. Sloane Physical Chemistry Dept. General Motors Research Lab. Warren, Michigan 48090

Dr. Ian W.M. Smith Dept. of Physical Chemistry Univ. Chemical Laboratories Lensfield Road Cambridge, ENGLAND

Mr. Randal K. Sparks MMRD - 70A-4435 Lawrence Berkeley Laboratory Berkeley, CA 94720

Prof. Leonard D. Spicer Chemistry Dept. University of Utah Salt Lake City, UT 84112

Dr. Richard C. Stern L-468 Lawrence Livermore Laboratory Livermore, CA 94550

Dr. Steven Stolte University of Nymegen Physics Dept. Toernooiveld, Nymegen HOLLAND Dr. Michael Stuke Max-Planck-Institut Postfach 968 3400 Gottingen, W. GERMANY

Prof. William C. Stwalley Dept. of Chemistry University of Iowa Iowa City, IA 52242 Prof. Dr. Juergen Troe Inst. fur Physik. Chemie Universitaet Goettingen D-3400 Goettingen, W GERMANY

Prof. Donald G. Truhlar Dept. of Chemistry University of Minnesota Minneapolis, Minn. 55455

Dr. Shen Y. Tang IRT Corporation P. O. Box 80817 San Diego, CA 92138

Dr. William L. Taylor Monsanto Research Corp. P. O. Box 32 Miamisburg, OH 45342

Dr. Peter W. Tiedemann MMRD - 70A-4418 Lawrence Berkeley Laboratory Berkeley, CA 94720 Prof. Trina Valencich 1370 Palms Blvd. Venince, CA 90291

Dr. James J. Valentini Dept. of Chemistry Harvard University Cambridge, Mass. 02138 Dr. Jochen Wanner Projektgruppe fur Laserforschung der Max-Planck-Gesellschaft, D-8046 Garching/Munich, WEST GERMANY

Dr. Sol Wexler Argonne National Laboratory 9700 S. Cass Ave. Argonne, Illinois 60439 Dr. Charles Edward Young CHM 200-A101 Argonne National Laboratory Argonne, Illinois 60439

Prof. Lennard Wharton James Franck Inst. University of Chicago 5640 S. Ellis Ave. Chicago, IL 60637

Dr. John Wiesenfeld Dept. of Chemistry Cornell University Ithaca, NY 14853

Prof. John Winn Dept. of Chemistry University of California Berkeley, CA 94720

Prof. Daniel H. Winicur Chemistry Dept. Univ. of Notre Dame Notre Dame, Indiana 46556

Dr. Francis J. Wodarczyk Rockwell International Science Center, P.O. Box 1085 Thousand Oaks, CA 91360 Prof. Richard N. Zare Dept. of Chemistry Stanford University Stanford, CA 94305

Dr. Reinhard Zellner Inst. fur Physik. Chemie Universitat Gottingen 3400 Gottinge, W. GERMANY AII-15

APPENDIX C

Attendance List 11th International Symposium on Rarefied Gas

Dynamics, Cannes, July 3-8, 1978

I.C. A.S. ABBASS DEPT. OF PHYSICS - COLLEGE OF SCIENCE UNIVERSITY OF BAGHDAD - ADHAMIYA BAGHDAD IRAG

T. ADE INST. SPACE AND AERON, SCI.; TOKYO UNIV. Komađa;meguro-ku Tokyo Japan

PROF, G. ADOMEIL INSTIT, F. ALLG, MECHAMIK , TECHN, MOSCHULE TEMPIERGRABEN 51 AACHEN RFA

DR. M. ALEXANDRE C.E.N. SACLAY; D.G.I./SEPCP B.F. N• 2 91190 GIF SUR YVETTE FRANCE

"PROFESSOR B.V. ALEXEJEV MOSCOW AVIATION INSTITUTE MOSCOW A-80 VOLOKOLAMSCOJE HIGHWAY 4 USSR

PR. J.A. ANDERSON DEPT, OF CHEMISTRY 152 DAVEY LAB.-PENNSYLV. STATE UNIV. UNIVERSITY PARK, PENNSYLVANIA 16802 USA

PROFESSOR R.P. ANDRES DEP. OF CHEMICAL ENGINEERING PRINCETON UNIVERSITY PRINCETON, NEW JERSEY 08540 USA

DR. J. ANDRE UNIV. DE PROVENCE - U.E.R. DE PHYSIQUE CENTRE DE ST-JEROME - SERVICE 232 13397 HARGEILLE CEDEX 4 FRANCE PROFESSOR J.B. ANDERSON DEPT, OF CHEMISTRY PENNSYLVANIA STATE UNIV. UNIVERSITY PARK, PA. 16802 USA

DR. K. AOKI DEPT. OF AERON. ENGIN. ;KYOTO UNIV. 7 MATSU-MACHI,MORIGUCHI OSAKA 570 JAPAN

DR. G. ARMAND C.E.N. SACLAY; BAT. 62 BP N+2 91190 GIF SUR YVETTE FRANCE

MADAHE HUETZ-AUBERT CNRS GR 14 ECAM 92290 - CHATENAY MALABRY FRANCE

~

PR. S.P. BAKANOV DEPT, SURF, PHENOM.- INST, PHYS, CHEM, USSR ACAD, OF SC. LENINSKI PROSPECT, 31 MOSCOW, B-71 - USSR

DR. A.G. BASHKIROV USSR NATIONAL COMMITTEE ON THEORETICAL & APPLIED MECHANICS VERNADSKOVO PROSPECT, 101 117526 MOSCOW B-526 - USSR

PROFESSOR E.W. BECKER KERNFORSCHUNGSZENTRUM 7500 KARLSRUHE POSTFACH 3640 RFA

PROFESSOR J.J.M. BEENAKKER RIJKSUNIVERSITEIT LEIDEN - HUYGENS LAB. WASSENAARSEWEG 78 LEIDEN 2405 THE NETHERLANDS

PROFESSOR N. BELLONO POLIT. DI TORINO,1ST. DI HECC. RAZ. Corso duca degli Abruzzi, 24 10129 Torino Italia

PROFESSEUR J.J. BERNARD UNIV. P. ET H. CURJE; LAB. AEROTHERMIQUE 4 TER, ROUTE DES GARDES 92190 MEUDON FRANCE

PROFESSOR S.B. BERNDT Royal Inst. of Technology ; K.T.H. S-10044 Stockholm 70 Sweden

DR. H.C.W. BEYERINCK PHYSICS DEPARTMENT EINDHOVEN UNIV. OF TECHNOLOGY EINDHOVEN - P.O. BOX 513 THE NETHERLANDS

PROF. DR. ALFRED E. BEYLICH TECHNISCHE HOCHSCHULE 5100 AACHEN

RFA

DR. G.K. BIENKOWSKI GAS DYNAHICS LABORATORY FORRESTAL CAMPUS PRINCETON UNIV. PRINCETON, NEW JERSEY 08540 USA

PR, DR. K. BIER INST. FUER TECH. THERM. & KALT. DER UNIV. KARLSRUHE 75 KARLSRUHE 1 RFA

DR. N.S. BIRAJDAR DEPT. OF MATHEMATICS YESHAWANT COLLEGE NANDED (431602) INDIA PROFESSOR G.A. DIRD DEPARIMENT OF AERONAUTICS THE UNIVERSITY OF SYDNEY SYDNEY, N.S.W. 2006 AUSTRALIA

DR. P. BLEY INST. FUER KERNVERFAHRENSTECHNIK DER UNIV. UND DES KERNFORSCHUNGSZENTRUMS D-7500 KARLSRUHE RFA

PROFESSOR G. BOATO FAC. DI SCIENZA MATEMATICHE UNIVERSITA DI GENOVA Genova Italy

DR. R. BOETTGER HAMBURG UNIV.; MPJ FUR KERNPHYS. POSTFACH 103980 69 HEIDELBERG 1 R.F.A.

DR. VICINIO BOFFI Lab. DI Ingegneria Nucleare 16, VIA dei Colli Bologna 40137 Italy

DR. PETEK BRANKO Ljubljana UNIV.; Physics Dept.

61000 LJUBLJNA Yugoslavia

•.

DR. B. BRIGOLI C.I.S.E. P.O.B. 3986 20100 Milano Italy

BR, GIAN PAOLO BRIVIO IST. DI FISICA DELL'UNIV, GRUPPO SOLIBI VIA CELORIA, 16, MILANO ITALY

. .

PROFESSOR C.L. BRUNDIN OXFORD UNIVERSITY DEPT. OF ENG. SC.-PARKS ROAD OXFORD OX1 3PJ ENGLAND

PROFESSEUR R. BRUN UNIV. DE PROVENCE; LAB. DYNAM. THERHOPH. FLUIDES CENTRE ST-JEROME 13397 MARSEILLE CEDEX 4 FRANCE

DR. G. BRUSDEYLINS MAX-PLANCK-INST. FUER STROMUNGS-FORSCHUNG - BOTTINGER STR. 6-8 D 3400 GOETTINGEN RFA

ŧ

DR. MANFRED BECKER D.F.V.L.R. INSTITUT FUER ANGEWANDTE GASDYNAMIK 5000 KOLN 90 - POSTFACH 90 60 58 RFA

PROFESSOR E.A. BRUN MEMBRE DE L INSTITÚT 8-10 PLACE DU COMMERCE 75015 PARIS FRANCE

PROFESSOR H. CABANNES UNIVERSITE PARIS V3 MECANIQUE THEORIQUE - TOUR 66 4, PLACE JUSSIEU 75005 - PARIS, FRANCE

DR. R. CAMPARGUE CEN SACLAY;SCPH LAB. JETS HOLECULAIRES BP 2 91190 GIF SUR YVETTE FRANCE

DR. D.H. CAMPBELL UNIV. OF SOUTHERN CALIFORNIA LOS ANGELES, CALIFORNIA 20007 USA

· 43

-

.

DR. C.D. CANTRELL LOS ALAMOS SCIENTIFIC LADORATORY LOS ALAMOS NEW MEXICO 87545 USA

DR. M. CATHALA SNIAS B.P. N•52 06322 Cannes La Rocca France

DR. T.D. NC CAY ARD, INC. AEUC DIVISION A SVERDRUP CORP. COMPANY ARNOLD AIR FORCE STATION, TENNESSEE 37389 - USA

PROFESSOR CARLO CERCIGNANI ISTITUTO DI HATEMATICA 20133 MILANO PIAZZA LEONARDO DA VINCI, 32 ITALY

MADAME SYLVIE CHABAUTY CNRS - INFORMASCIENCE CENTRE DE DOC. SCIENT. & TECH. 26, RUE BOYER 75971 PARIS CEDEX 20 - FRANCE

DR. ASHOK U. CHATWANI INST. FUER THERM. & FLUIDDYNAMIK RUHR-UNIVERSITAET BOCHUM BOCHUM RFA

DR. K.Y. CHIEN NAVAL SURFACE WEAPONS CENTER Silver Spring Maryland 20910 USA

A.A. CHIKHAOUI LAB. DYN. THERMOPHYS. FLUIDES CENTRE DE ST. JEROME 13397 MARSEILLE CEDEX 4 FRANCE

DR: P. CODA C.N.E.N. S.P.ANGUILLARESE KH 1+300 Roma Italy

DR. G.T. COLEMAN Royal Aircraft Establ.; Aerodynamics Dept.

FARNBOROUGH; HAMPSHIRE G.B.

ì

DR. R. COLE BOLOGNA UNIVERS.;LAB. NUCLEAR ENGIN. VIA DEI COLLI.16 40136 BOLOGNA ITALY

PROFESSOR GLORGE COMSA I.F. GRENZFLAECHENFORSCHUNG UNI VAKUUNPKYSIK POSTFACH 1913, 5170 JUELICH 1 RFA

PROFESSOR NOEL CORNGOLD AFPLIED SCIENCE DEPT. CAL. INSTITUTE OF TECHNO. PASADENA, CALIF, 91125 USA

RR. C. CREEMERS FYSICO-CHEMISCH LABORATORIUM VAN DE K.U. LEUVEN CELESTIJNENLAAN 200 G B - 3030 HEVERLEE - BELGIUM

DR. J. CUVELLIER C.E.N. SACLAY; PHYS, ATUHIQUE B.P.N+2 91190 GIF SUR YVETTE FRANCE

DR. A.E. DABIRI MECHANICAL ENGIN, DEPT, ARYA MEHR UNIV, OF TECHNOLOGY P.O. BOX 3406 - TEHRAN IRAN

طالع

DIPL. PHYS. C. DANKERT DFVLR, AERO. VERBUCHSANSTALT I. F. DYN. VEPBUENNTER GASE PUNSENGTR. 10, 3400 GUETTINGEN RFA

DR. J. DARROZES ENSTA 32, BD VICTOR 75015 PARIS FRANCE

•

DR. R. DAVID J.G.V. DER KERNFORSCH, JUELICH KFA JUELICH/JGV 5170 JUELICH, POSTFACH 1713 R.F.A.

HADAME C. DEGEN CHRS - INFORMASCIENCE CENTRE DE DOC. SCIENT. & TECHN. 26, RUE BOYER 75971 PARIS CEDEX 20 - FRANCE

DR, S.M. DESHPANDE · DEPARTMENT OF AERON. ENG. INDIAN INSTITUTE OF SCIENCE BANGALORE INDIA

DR. F. M. DEVIENNE LAB. PHYSIQUE MOL. DES HAUTES ENERGIES BP N•2 04530 PEYHEINADE FRANCE

DR. I. DIMICOLI C.E.N. SACLAY; D.R.A./SIRMA BP N•2 91190 GIF SUR YVETTE FRANCE

PROFESSOR J. DORNING LANRENCE LIVERNORE LABORATORY UNIVERSITY OF CALIFORNIA P.O. ROX 808 - LIVERNORE, CA 94950 USA

.

DR. C. DREVET C.E.A. 3, CLOS PEROULT 91200 ATHIS-MONS FRANCE

DR. D. DREYFUSS M.I.T. R.M. 37-438 77 MASSACHUSETTS AV., CAMBRIDGE MA 02139 U.S.A.

DR. ANIONI DROBNIK Politechnika Lodzka Instytut Fizyki UL. Wolczanska 219 93-005 Lodz - Poland

DR. B. BUBORGEL C.E.N. LIMEIL BP N•27 94190 VILLENEUVE ST GEORGES FRANCE

DR. RICHARD EDWARDS DEPARTMENT OF AEROSPACE ENGINEERING UNIVERSITY OF SOUTHERN CALIFORNIA LOS ANGELES, CALIFORNIA 90007 USA

DR. W. EHRFELD KERNFORSCHUNGSZENTRUM KARLSRUHE GHBH INSTITUT FUER KERNVERFAHPENSTECHNIK POSTFACH 3640 - D 7500 KARLSRUHE 1 RFA

DR. T. ENGEL INST. FUER PHYS. CHEMIE UNIVERSITAET MUENCHEN SOPHIENSTR. 11, D-8000 MUENCHEN 2 RFA

DR. HIROSHI ENJOJI THERMONUCLEAR FUSION LABORATORY INST. OF PHYS. & CHEM, RESEARCH 2-1, HIROSAWA, WAKO-SHI, SAITAMA 351 JAPAN DR. X. DE L EPREVIER D.R.E.T. 26 BD VICTOR 75976 PARIS ARMEES FRANCE

DR. EROFEEV COMP. CENTER OF ACAD, OF SC. OF USSR VAVILOVA 40 MOSCOW 117333 USSR

DR. M. EYB Dornier System GmrH Postfach 1340 7970 Friedriechshafen RFA

DR. U. EHRFELD KERNFORSCHUNGSZENTRUM 7500 KARLSRUHE Postfach 3640 RFA

DR. ALAN MC FALL DIV. 5735 - SANDIA LABORATORIES ROX 5800 Albuquerque NM 87115 USA

DR. JEAN FARGES GROUPE DES AGREGATS MOLECULAIRES BAT. 210 - UNIV. PARIS-SUD 91405 ORSAY CEDEX FRANCE

PROFESSOR A. FAVRE DIRECTOR, I.M.S.T. 12, AVENUE DU GENERAL LECLERC 13003 MARSEILLE FRANCE

PROFESSOR JOHN B. FENN Yale University - Mason Laboratory 9 Hillhouse Avenue New Haven, Conn, 06520 USA

PROFESSOR M. FIEBIG RUHR UNIV., BOCHUH, INSTITUT FUR THERHO-FLUID DYNAMIK RUHR UNIVERSITAT GOCHUM 463 BOCHUM - RFA

PROFESSOR S, S, FISHER UNIVERSITY OF VIRGINIA ENGINEERING AND APPLIED SCIENCE CHARLOTTESVILLE, VA 22901 USA

PROFESSOR W. FISZDON INST. OF FUND. TECH. RES. DEPT. OF FLUID DYNAMICS SWIETOKRZYSKA 21, OO-049 WARSZAWA, POLAND

PROFESSOR MARC FITAIRE LAB. DE PHYSIQUE DES PLASMAS BATIMENT 212 91405 ORSAY CEDEX FRANCE

DR. D. FRADKIN LOS ALAMOS SC. LAB. P.O. BOX 1663 H.S. 565 LOS ALAMOS NEW MEXICO 87545 U.S.A.

PROFESSOR J.B. FRENCH INST. FOR AEROS.STUDIES-UNIV. OF TORONTO 4925 DUFFERIN ST. DOWNSVIEW, ONTARIO M3H ST6 - CANADA

PROFESSOR DR. A. FROHN INSTITUT FUER THERHODYNAMIK DER LUFT-UND RAUMIAHRT UNIV. STUTIGART 7 - STUTIGART B0 (VAIHINGEN) RFA

DR. INZ, PIDTR GAJFWSKI INSTYTUT TECHNIKI LOTNICZEJ I MECHANIKI STOSOWANEJ NOWOWJEJSKA 24 00-665 WARSZAWA – POLAND DR. ROPERT J. CALLAGHER 8321, FLUID NECHANICS DIVISION SANDIA LABORATORIES LIVERMORE, CALIFORNIA 94550 USA

PRIV. DOZ. DR. ING. B. GAMPERT UNIVERSITAET ESSEN - GESANTHOCHSCHULE FACHDEREICH 12 - MASCHINENTECHNIK 4300 ESSEN 1 - POSTFACH 6843 RFA

DR. R. GATIGNOL LABORATOIRE DE MECANIQUE THEORIQUE UNIVERSITE P. & M. CURIE TOUR 66 - 4 PLACE JUSSIEU 75230 CEDEX 05 PARIS - FRANCE

DR. P. GAUCHEREL LAB. D'AEROTHERMIQUE DU CNRS 4 TER ROUTE DES GARDES 92190 MEUDON FRANCE

IR. M. A. GAVEAU CEN SACLAY,SCPH LAD. JETS MOLECULAIRES BP 2 91190 GIF SUR YVETTE FRANCE

DR. G. GEORGANTOPOULOS PATRAS UNIVERSITY; NECHANICS DEPT.

PATRAS GREECE

DR. D.P. GIDDENS SCHOOL OF AERO, ENG. GEORGIA INST. OF TECH. ATLANTA, GA. 30332 GEORGIA - USA

DR. GERSTENKORN LAR, AYNE COTTON - CNRS II CAMPUS D ORSAY 91905 ORSAY CEDEX FRANCE

nan kanan dara ya 199 Tani Mara Matika Ingila

DR. VON HALLE UNION CARBIDE CORP. OAK RIDGE GAS DIF. PLANT K-25 - P.O. BOX P OAK RIDGE, TENNESSEE 37830 USA

DR. B.B. HAMEL THERMAL & FLUID SCI, GR. DREXEL UNI, 32ND AND CHESTNUT STREET PHILADELPHIA, PENNA 19104 USA

DR. P.J. HARBOUR Culhan Laboratory Abingdon Oxon, OX 14 3 BB

ENGLAND

DR. J.K. HARVEY IMPER. COL. OF SC. & TECHNOLOGY DEPT. OF AERONAUTICS PRINCE CONSORT ROAD, LONDON SW7 2BY - ENGLAND

DR. L.J.F. HERMANS HUYGENS LABORATORIUM RIJKSUNIVERSITEIT LEIDEN LEIDEN THE NETHERLANDS

DR. G. HERMAN AMSTERDAM UNIVERSITY; NATUURKUNDIG LAD. VALCKENIERSTRAAT 45 AMSTERDAM THE NETHERLANDS

DR. A. HOAREAU UNIV. CLAUDE BERNARD LYON I 43 BD. DU 11 NOV. 1918 69621 VILLEURBANNE FRANCE

PROFESSOR HEINRICH HORA DEPT. OF THEURETICAL PHYS. THE UNIV. OF NEW SOUTH WALES KENSIGTON NSW 2033 - P.O. BOX 1 AUSTRALIA PROFESSOR Y, K, HSU BANGOR COMMUN. COLL, UNIV, OF MAINE AT ORONO ROOM 111 BANGOR HALL BANGOR MAINE 04401 USA

PROF. DR. H. HULSMAN DEPT. NATUURKUNDE, UNIV. INSTELLING ANTWERPEN B-2610 WILRIJK BELGIUM

PROFESSOR FRANKLIN C. HURLBUT DEPT, OF MECH. ENGINEERING UNIVERSITY OF CALIFORNIA BERKELEY, CALIFORNIA 94720 USA

DR. A. E. LE JEUNE CEN SACLAY; DGI/SEPCP BP N+2 91190 GIF SUR YVETTE FRANCE

DR. E.A. JOHNSON DEPARTMENT OF PHYSICS UNIV. OF SURREY, GUILDFORD SURREY, GU2 5XH ENGLAND

DR. NIKOS KAFOUSIAS Patras University; Dept Mechanics Patras

GREECE

PROFESSOR K. KARAMCHETI DEPT. OF AERO. & ASTRONAUTICS STANFORD UNIVERSITY STANFORD, CALIFORNIA 94305 USA

PROFESSOR GERALD R. KARR UNIVERSITY OF ALABAMA P.O. BOX 1247 HUNTSVILLE, ALABAMA 35807 USA

DR RUBERT G, GILDERT DEPT. OF THEOR. CHEMISTRY UNIVERSITY OF SYDNEY N.S.W. 2006 AUSTRALIA . .

DR, A, GINZBERG HEBREW UNIVERSITY EINSTEIN ST. 25/A HAIFA ISRAEL

DR, GIUSEPPE GOBDINI MASSACHUSETTS INST. OF TECHNOLOGY DEPT. OF AERONAUTICS & ASTRONAUTICS CAMBRIDGE, MASSACHUSETTS 02137 USA

DR. G. GODEFROY C.E.N. LIMEIL BP N•27 94190 VILLENEUVE ST GEORGES FRANCE

DR. L. GOTTESDIENER LABORATOIRE D'AERCTHERMIQUE DU CNRS 4 TER ROUTE DES GARDES 92190 MEUDON FRANCE

PROFESSOR C. GOUDAS PATRAS UNIVERSITY; MECHANICS DEPT.

PATRAS GREECE

4 **ا**

PROFESSOR T.E. GOUGH GUELPH-WATERLOO CENTRE FOR GRAUUATE WORK IN CHEMISTRY UNIV. OF WATERLOO, ONTARIO N2L 3G1 - CANADA

PROFESSOR R. GRICE THEORETICAL CHEMISTRY DEPARTMENT CAMBRIDGE UNIVERSITY CAMBRIDGE CB 2 1EW ENGLAND PROFESSOR A.E. GROSSER DEPARTMENT OF CHCHISTRY MC GILL UNIV., 801 SHERBROOKE STREET WEST NONTREAL H 3A 2K6 - CANADA

DR. R. DE GRYSE Gent State University; Solid State LAB. Si Krijgslaan 271 9000 gent Belgium

DR. J. GSPANN I.F. KERNVERFAHRENSTECHNIK KERFORSCHUNGSZENTRUM 7500 KARLSRUHE - POSTFACH 3640 RFA

DR, F, GUENTHER Siemens ag 8000 Huenchen

RFA

DR. F. GUNTHER INST. FUER TECH. CHEMIE DER TECHN. UNIVERSITAET HANNOVER Callinstr. 3 3000 Hannover - RFA

DR. M. GILBERT CEN SACLAY ; SCPH BP N•2 91190 GIF SUR YVETTE FRANCE

DR. H. HABERLAND FAK, FUER PHYSIK DER UNIVERSITAET 79 FREIBURG I. BR. HERMANN-HERBER-STRASSE **3** RFA

PROF, DR. O.F. HAGENA I.F. KERNVERFAHRENSTECHNIK KERNFORSCHUNGSZENTRUM 7500 KARLSRUHE, PF, 36%0 RFA

DR, M.D. KARTALEV INST. OF HATH. AND MECHAN. P.O, BOX 373 Sofia 1000 Bulgaria

DR. H. KIESEWETTER Ruth Aachen Abtlg Physiologie Melatenerstr, 211 51 Aachen RFA

PROFESSOR F. KLEIN WEIZHANN INST. OF SCIENCE; ISOTOPE RES, DEPT,

• •

REHOVOT Israel

.

.

PROFESSOR H.F. P. KNAAP Leiden Univ.; Huygens Lab. Wassenaarseweg 78 2300 Ra Leiden The Netherlands

PROFESSOR E, KNUTH 5531 BOELTER HALL UCLA LOS ANGELES, CALIFORNIA 90024 USA

DR. A. KOCIC INST, OF NUCL, SC. BORIS KIDRIC 11001 BEDGRAD P.O. BOX 522 YUGOSLAVIA

PROFESSOR M.N. KOGAN COMP. CENTER OF ACAD, OF SC. OF USSR VAVILOVA 40, Moscow 117333 USSR

DR. G. KOPPENWALLNER DFLVR - INST. DYN, VERD. GASE 3400 GOETTINGEN RUNSENSTRASSE 10 RFA

٤

DR.J. KORVING LEIDEN UNIV.; HUYGENS LAB. WASSENAARSEWEG 78 2300 RA LEIDEN THE NETHERLANDS

DR. Y.A. KOSHMAROV Muscow aviation institute Moscow

USSR

DR. S. KOSOWSKI Inst. of Fund. Technology Res, UL. Pereca 2 m 1214 00-849 Warsaw Poland

DR. K. KOURA National Aerospace LAB. 1830 Jindaiji-Machi Chofu Tokyo Japan

DR. MIROSLAV KRANYS DEPARTEMENT DE PHYSIQUE UNIVERSITE DE MONTREAL MONTREAL - CASE POSTALE 6128 CANADA

DR. G. KRIEG Kernforsch.zentr. Karlsruhe Postfach 3640 75 Karlsruhe R.F.A.

DR. J.G. KUHRY C.R.NUCL. DE STRASPOURG 23 RUE DU LOESS 67037 STRASBOURG CEDEX FRANCE

,

DR. A. KULICKI INST. OF HEAT ENGINEERING WARSAW TECHN. UNIVERSITY UT. NOWOMIEJSKA 25 00-665 WARSAW - POLAND

DR. JOSEF KUNC Warsaw Technical University Nowowiejska 25 00-665 Warsaw Poland

PROFESSOR IVAN KUSCER UNIVERSITAET KAISERSLAUTERN 6750 KAISERSLAUTERN POSTFACH 3049 RFA

PROFESSOR S.S. KUTATELADZE CORR, MEMMER OF THE USSR AC, OF SC. DIRECTOR - INST, OF THERHOPHYSICS NOVOSIDIRSK-90, 630 090 USSR

DR. K. KYNEFORS - Goteborg Univ., Phys. Chem. Dept. Fack; S-402 20 Goteborg

SWEDEN

- ÷

DR. F. LABERNEDE C.E.N. LIMEIL BP N•27 94190 VILLENEUVE ST GEORGES FRANCE

DR. WOLTERBEEK LAMPERT TWENTE UNIV. OF TECHNOLOGY AFD. TN., T.H.TWENTE POSTBUS 217 ENSCHEDE THE NETHERLANDS

DR. J.A.C. LAMBREGTS LIMBURG UNIV.; BIOMEDL.CENT. MED.FACULTY BEELDSNIJDERSDREEF 101 6216 EA MAASTRICHT THE NETHERLANDS

DR. M. LAMPIS IST, DI MATEM. DEL POLIT. DI MILANO PIAZZA LEONARDO DA VINCI, 32 20133 MILANO ITALY DR. LANG MAX-PLANCK-INST. FUER STROMUNGS-FORSCHUNG - BUTTINGER STR. 6-8 B 3400 GOETTINGEN RFA

DR. J. LAPUJOULADE C.E.N. SACLAY; PHYS. ATOM. BAT.62 DP N+2 91190 GIF-SUR-YVETTE FRANCE

DR. A.R. LARSON UNIV. OF CALIF.,LAS. SCIENT. LAB. P.O. BOX 1663 LOS ALAMOS NEW MEXICO 07545 USA

DR. S. LEACH CNRS-LAB. DE PHOTOPHYSIQUE Batiment 213 Faculte des sciences 91405 ORSAY - FRANCE

DR. A. LEBEHOT CEN SACLAY ; SCPH LAB. JETS MOLECULAIRES RP 2 91190 GIF SUR YVETTE FRANCE

PROFESSOR J.H. DE LEEUW UNIVERSITY OF TORONTO 30 VALLONCLIFFE POAD TOWNHILL, ONTARIO CANADA

PROFESSOR Y.T. LEE LANNENCE BERKELEY LABORATORY UNIV, OF CALIFORNIA - BERKELEY CALIFORNIA 54720 USA

DR. D. LEGUILLON LAB. DE MECAN. THEOR. UNIVERSITE P. & M. CURIE TOUR 66 - 4 PLACE JUSSIEU 75230 CEDEX 05 PARIS - FRANCE DR. Y. LEJAY C.E.N. SACLAY; PHYS, ATOM. BAT.62 BP N=2 91190 GIF SUR YVETTE FRANCE

DR. J.C. LEMONNIER CEN SACLAY;SCPH LAB, JETS MOLECULAIRES BP 2 91190 GIF SUR YVETTE FRANCE

MONSIEUR JC. LENGRAND CNRS-LAB, D'AEROTHERMIQUE 4 TER, ROUTE DES GARDES 92190 MEUDON FRANCE

PROFESSOR V.B. LEONAS SPACE RESEARCH INST. USSR ACAD. SCI. 68, PROFSOYUZNAJA ST. 117810 GSP 312, MOSCOW USSR

PROFESSOR L.L. LEVENSON DEPARTMENT OF PHYSICS UNIV. OF MISSOURI-ROLLA ROLLA, MISSOURI 65401, USA

 \sim

PROFESSOR CLARK H. LEWIS VIRGINIA POLYT. INST. & STATE-UNIV. AEROS & OCEAN ENGR. BLACKSBURG VIRGINIA 24061 - USA

DR. C.P. LI APPLIED MECHANICS DEPARTMENT LOCKHEED ELECTRONICS CO., INC. HOUSTON, TEXAS 77058 USA

PROFESSOR J. LOS . FOM INST. FOR ATOM.& MOL. PHYS. KRUISLAAN 407 Amsterdam-oost The Netherlands DR. P, LOUVET CEN SACLAY; SCPH BP N+2 91190 GIF SUR YVETTE FRANCE

DR. D.Y.S. LOU DEPT. OF MECH, & AEROS. ENGIN. UNIVERSITY OF DELAWARE NEWARK, DE 19711 USA

DR. A, LUBBERT INST. FUER TECH. CHEMIE DER TECHN. UNIVERSITAET HANNOVER Callinstr. 46, Eingang 3 -D 3000 Hannover - RFA

DR. J.W.L. LEWIS Ard, Inc. AEDC Division A Sverdrup Corp. Company Arnold Air Force Station Tennessee 37389 - USA

PROFESSOR R.J. HADIS DEPT, OF PHYSICAL CHEMISTRY UNIVERSITY OF HUNICH MUNICH RFA

DR. N.K. MAJUMDAR Saha Inst. of Nuclear Physics Calcuita

INDIA

PROFESSOR J.R. MANSON CENS - SERVICE DE PHYS. ATOMIQUE INTERACTIONS GAZ-SOLIDES B.P. NO. 2 91190 GIF-SUR-YVETTE - FRANCE

DR. C. MANUS CEN SACLAY ; DPH.GZ PA BP N•2 91190 GIF SUR YVETTE FRANCE

DR. D. MARETTE CEN SACLAY;SCPH LAB. JETS NOLECULAIRES BP 2 91190 GIF SUR YVETTE FRANCE

DR. K. H. MARTENSSON A.B. ATOMENERGI

61101 NYKOPING Sweden

Ì

DR. J.P. MARTIN CNRS - ECOLE CENTRALE (050014) GRANDE VOIE DES VIGNES 92270 CHATENAY MALABRY FRANCE

DR. NATAKEYAMA MASAYUKI Tokyo Met. College Aeron. Engen. 8-53-1 minahisenju Arkawa-ku Tokyo Japan

PROF. D. MASSIGNON CEN SACLAY; SCPH &P N•2 71190 GIF SUR YVETTE FRANCE

.

DR. P. MENGER EINDHOVEN UNIV. OF TECHNOLOGY PO BOX 513 EINDHOVEN THE NETHERLANDS

MR. S.C. METCALF HEAD, HIGH SPEED FLOWS SECT. AER. DEPT. ROYAL AIRCRAFT ESTABLISHMENT FARNBORDUGH/HANTS ENGLAND

PROFESSOR HISASHI MIKANI PEBEARCH LAP. OF NUCL. REACTOR TOKYO INSTITUTE OF TECHNOLOGY O-OKAYAMA, NEGURO-KU TOKYO - JAPAN DR. D.R. HILLER UNIV. OF CALIFORNIA, SAN DIEGO DEPT. OF APPL. MECHAN.& ENG. SC.B-010 LA JOLLA, CALIFORNIA 92093 USA

DR. N.K. MITRA INSTITUT FUER THERMO-UND FLUIDDYNAMIK 463 BOCHUM-OUERENBURG POSTFACH 2148 RFA

PROF, V. MOLINARI BOLOGNA UNIV.; LAB, NUCL. ENG. VIA DEI COLLI 16 BOLOGNA ITALY

DR. R. MONACO ISTITUTO DI MECCANICA RATIONALE-POLITECH, TORINO CORSO DUCA DEGLI ABRUZZI 24 TURIN ITALY

DR. G, MONNOM LAB. PHYS. MOL. DES HAUTES ENERGIES RP N•2 04530 PEYMEINADE FRANCE

DR. G. HONNON LAD. PHYS. MOL. DES HAUTES ENERGIES BP N•2 06530 PEYMEINADE FRANCE

DR, J.P. HORERA CEN F.A.R.; DPH/PFC BP N+6 72260 FONTENAY AUX ROSES FRANCE

DR. A. MORTELETTE AEROSPATIALE-CANNES BP 52 06322 CANNES LA BOUCA FRANCE

and the second second second

'1

PROFESSOR E.P. MUNTZ UNIV. OF SOUTH. CALIFORNIA DEPT. OF ACRO. ENG. UNIV. PARK LOS ANGELES, CALIFORNIA 90007 USA

PROFESSOR W.H. MILLER DEPT. OF CHCH. LAWRENCE BERKELEY LAB. UNIVERSITY OF CALIFORNIA BERKELEY, CA 94720 USA

PROFESSOR ROODAM NARASIMHA DEPARTMENT OF AERONAUTICAL ENG. INDIAN INSTITUTE OF SCIENCE BANGALORE INDIA

PROF. H. NEUERT HAMBURG UNIV.; INST. EXPER. PHYSICS 36 JUNGRUNSTR. 2000 HAMBURG R.F.A.

DR. MICHIO NISHIDA DEPT, OF AERON, ENGIN, Kyoto University Kyoto Japan

DR. W. OBERT KERNFORSCHUNGSZENTRUM KARLSRUHE GMBH INST. FUER KERNFAHRENSTECHNIK POSTFACH 36 40 - D 7500 KARLSRUHE RFA

DR. M.J. OFFERHAUS FOM-INST. VOOR ATOOM-EN MOLEC. KRUISLAAN 407 AMSTERDAM THE NETHERLANDS

PROFESSOR HAKURO DGUCHI I. OF SPACE & AERONAUTICAL SCIENCE UNIVERSITY OF TOKYO KOMADA, MEGURO-KU TOKYO, JAPAN PROFESSOR D.C. PACK UNIV.OF STRAT.-DEPT. OF MATHEMATICS LIVINGSTONE TOWER 26 RICHMOND STREET GLASGOW G1 1XH, SCOTLAND

BR. G. C. PANDE UNIV. PATRAS ; DEPT. MECHANICS PATRAS

GREECE

.

PROFESSOR H. PAULY MPI F. STROEMUNGSFORSCHUNG 3400 GOETTINGEN BOETTINGERSTR. 6/8 RFA

DR. T. PLATKOWSKI WAPSAW UNIV.; DEPT. MATH. & MECHANICS WARSAW P.KI. N. WARSAW POLAND

DR. P. PLURIEN CEN SACLAY ; DGI BP N•2 91190 GIF SUR YVETTE FRANCE

DR. B.T. PORODNOV URALS POLYTECHNICAL INSTITUTE PHYSICAL & TECHN. FACULTY 620002, SVERDLOVSK K-2 USSR

DR. J. LEITH POTTER VON KARMAN GAS DYNAMICS FAC. ARO, INC. ARNOLD AFG TENN. 37389 USA

MME. C. POUX SOPEMEA 18 RUE EDOUARD BELIN 31029 TOULOUSE FRANCE

PROFESSOR I. PRIGOGINE PRIX NOBEL DE CHIMIE UNIVERSITE LIBRE DE BRUXELLES BRUXELLES BELGIUM

UR. STEFAN RADEV INST. OF MATHEMATICS & MECHANICS ACADEMY OF SCIENCES KV. GEO MILEV, UL. 36, BL. 8 SOFIA 1113 - BULGARIA

DR. S.M. RAHACHANDRA HINDUSTAN AERONAUTICS LIMITED POST BOX 1789 Bangalore 560017 India

DR. A.K. RAY 2012, HOLYBROOK CRESCENT OTTAWA KIJYY6 ONTARIO CANADA

FROFESSOR A.K. REBROV INSTITUTE OF THERMOPHYSICS SIBERIAN BRANCH OF THE USSR AC. SCI. NOVOSIPIRSK-90 USSR

DR. R. REICHENBACH EUROPEAN RESCAPCH OFFICE; US ARMY 223 OLD MARYLEBONE ROAD LONDON NW1 5TH U.K.

PROFESSOR J, REUSS FYS.¹ LAB. KATH. UNIV. TOERNOOIVELD, DRIEHUIZERWEG 200 NIJMUGEN, THE NETHERLANDS

DR. P. RIGNY CEN SACLAY; DGI/SEPCP FP N+2 91190 GIF SUR YVETTE FRANCE PROF, Y. RIJOV MOSCOW AVIATION INSTITUTE Volokolamskojc Road 4 Moscow A-80 USSR

MR. MILTON ROGERS 6307 PEARCE AVE. Baltinore Maryland USA

DIPL, ING, F. ROMANI LEHRSTUHL FUER ALLG. MECHAN. AN DER RHEIN, WESTF. TECHN. HOCHSCHULE 5100 AACHEN RFA

PROFESSEUR A. RO "GARD CEN SACLAY; SCPH BP N•2 91190 GIF SUR YVETTE FRANCE

DR. G. PITTION-ROSSILLON CEN LIMEIL BP N•2 94190 VILLENEUVE ST GEORGES FRANCE

DR. J.C. ROUSTAN LAB. PHYS. MOL. DES HAUTES ENERGIES BP N•2 04530 PEYHEINADE FRANCE

DR. B. ROWE LAB. D'AEROTHERMIQUE DU CNRS 4 TER ROUTE DES GARDES 92190 HEUDON FRANCE

PROFESSOR OLEG RYZHOV COMPUTING CENTER USSR ACAD, OF SCI. 40 VAVILOV ST. - 117333 MOSCOW USSR

DR. W.C. SCHIEVE CENTER FOR STAT. MECH. & THERM. THE UNIV. OF TEXAS AT AUSTIN, AUSTIN, TEXAS 78712 USA

DR. E. SCHNID Kernforsch.zentr. Karlsruhe abt. : ikvt Postfach 3640 D 75 Karlsruhe RFA

PROFESSOR G. SCOLES LIBERA UNIV. DEGLI STUDI DI TRENTO FACOLTA DI SCIENZE 38050 POVO (TRENTO) ITALY

LT.COL. D. W. SEEGMILLER US AIR FORCE EUROP. OFFICE SCIENT. RES. & DEVEL, EOARD, 223 OLD MARYLEBONE RD. LONDON NW1 UK

DR. M. SEIDL DORNIER SYSTEM GMBH 7990 FRIEDRICHSHAFEN POSTFACH 1360 RFA

~

DR. F. SEILER INST. F. STROM. & STROMUNGSMASCH. UNIVERSITAET KARLSRUHE D-7500 KARLSRUHE RFA

DR. P.J. SEVERIN PHILIPS RESEARCH LABORATORIES EINDHOVEN

THE NETHERLANDS

DR..E.M. SHAKHOV COMPUTING CENTER USSR ACADENY OF SCIENCES MOSCON USSR DR. R.G. SHARAFUTDINOV INSTITUTE OF THERMOPHYSICS SIBERIAN BRANCH OF THE USSR AC. SC. NOVOSIBIRSK-90 USSR

PROFESSOR FREDERICK SHERMAN DEPT. OF MECHANICAL ENGINEERING UNIVERSITY OF CALIFORNIA BERKELEY CALIFORNIA 94720 TH01 USA

DR. STEVEN J. SIBENER LAWRENCE BERKELEY LABORATORY UNIVERSITY OF CALIFORNIA BERKELEY, CALIFORNIA 94720 USA

DR. D. SIEGELMAN AVCO SYSTEMS DIV. 201 Lowell St. Wilmington, MA 01887 USA

.

DR. R.E. SMALLEY DEPT. OF CHEMISTRY - RICE UNIVERSITY P.G. BOX 1892 HOUSTON, TX 77001 USA

BR. T, SOGA DEPARTMENT OF AERON. ENG. Nagoya University Chikusa-ku Nagoya 464 Japan

PROFESSOR YOSHIO SONE DEPARTEMENT OF AERONAUTICAL ENG, KYOTO UNIVERSITY KYOTO 606 JAPAN

DR. V.H. SOUNDALGEKAR DEPT. OF MATHEMATICS INDIAN INST. OF TECH. POWAI EOMBAY -400075 INDIA PAGE IS

OF POOR QUALITY

PROFESSOR GILBERT D. STEIN GASDYNAMICS LABORATORY DEPT. OF NECH. ENG. & AST. SC. Northwestern University Evanston, Illinois 60201 - USA

\$

ACAD. V.V. STRUMINSKII DEPT. OF MECH. OF HETEROG. MEDIA USSR ACAD. OF SCIENCES 125040 MOSCOW A-40 LENINGRADSKII PROSP. 7 - USSR

PROFESSOR L. TALBOT UNIVERSITY OF CALIFORNIA MECHANICAL ENGINEERING BERKELEY, CALIFORNIA 94720, TM01 USA

DR. S.P. TANG TRW - DEFENSE & SPACE SYST. GRP ONE SPACE PARK R1-1196 Redondo Beach, California 90273 USA

PROFESSOR C.M. TCHEN 2 STORSKOVSVEJ 4682 TUREBY

DENMARK

٩

PROFESSOR J.G. THEOBALD FAC. DES SCIENCES LAB. SPECT. HERT. & D'ELECTRONIQUE F-25030 BESANCON CEDEX FRANCE

PROFESSOR L.B. THOMAS PHYSICAL CHENICAL LABORATORY UNIVERSITY OF MISSOURI COLUMBIA, MISSOURI 55201 USA

PROFESSOR DR. J.P. TOENNIES MAX-PLANCK-INST. F. STRUEMUNGSFORSCH. D 3400 GOETTINGEN POSTFACH 847, BOETTINGERSTRASSE 4/8 RFA DR. G. TORCHET GROUPE DES AGREGATS MOLECUL. UNIVERSITE PARIS SUD 91405 ORSAY FRANCE

DR. K.J. TOURYAN FLUID & THERMAL SCIENCES DEPT, 1260 SANDIA LAHORATORIES, P.O. BOX 5800 ALBUQUERQUE, HEW MEXICO 87115 USA

PROFESSOR L. TRILLING AERONAUTICS AND ASTRONAUTICS MASSACHUSETTS INST, OF TECHNOLOGY CAMERIDGE, MASSACHUSETTS 02139 USA

PROFESSOR J. VALENSI LABORATOIRE DE MECANIQUE DES FLUIDES 1, RUE HONORAT. 13000 - MARSEILLE FRANCE

DR. Y. VENDENBOOMGAERDE CEN LIMEIL BP N•27 94190 VILLENEUVE ST GEORGES FRANCE

PROFESSOR DR. N.F. VERSTER TECHN, HOGESCHOOL TE EINDHOVEN INSULINDELAAN 2 EINDHOVEN THE NETHERLANDS

DR. H. VESTNER INSTITUT F. THEORETICAL PHYSIK DER UNIVERSITAET ERLANGEN – NUERNBERG 852 ERLANGEN, GLUECKSTRASSE 6 RFA

PROFESSOR Y. WACHMAN MASSACHUSETTS INST. OF TECHN. DEPT. OF AEKO. & ASTRONAUTICS CAMERIDUE, HASSACHUSETTS 02139 USA

•

PROFESSOR L. WALDMANN INST. F. THEORETISCHE PHYSIK UNIV. ERLANGEN-NUERNBERG 8520 ERLANGEN GLUECKSTRASSE 6 RFA

DR. Z. WALENTA FLUID MECH. DEPT.; INST. FUNDTL. TECHNOL. RES.;AC SWIETOKRZYSKA 21 00-049 WARSAW POLAND

DR. G.H. WEGDAM UNIVERSITEIT VAN AMSTERDAM LABORATORIUM VOOR FYSICHE CHEMIE NIEUWE PRINSENGRACHT 126 AMSTERDAN - THE NETHERLANDS

PROFESSOR J.F. WENDT VON KARMAN INST. FOR FLUID DYNAMICS CHAUSSEE DE WATERLOO 72 B - 1640 RHODE-ST-GENESE BELGIUM

PROFESSOR LENNARD WHARTON THE J. FRANCK INST., UNIV. OF CHICAGO 5640 ELLIS AVE. CHICAGO, ILLINOIS 60637 USA

PROFESSOR DR. M.M.R. WILLIAMS NUCL. ENGIN. QUEEN MARY COLLEGE LONDON UNIVERSITY LONDON ENGLAND

DR. KLAUS WINKELMANN MAX-PLANCK-INST. FUER STROMUNSFORSCH. 3400 GOETTINGEN BOETTINGERSTRASSE 6/8 RFA

DR. A. NISHIEWSKI WARSAW TECHNICAL UNIVERSITY UL.ZWYCIEZCOW 13 M.7 03-936 WARSAW POLAND

يكرز

DR. H. WITEK INT. FUR MOLPHYS.-FREIE UNIV. BERLIN BOLTZMANNSTRASSE 20 1000 BERLIN 33 RFA

PROFESSOR CURT WITTIG UNIVERSITY OF SOUTHERN CALIFORNIA DEPARTMENT OF ELECTRICAL ENGINEERING LOS ANGELES, CALIFORNIA 90007 USA

PR. G. WORTBERG Lehrstuhl fuer Allg. Mech. Technischen Hochschule Aachen 5100 Aachen RFA

PROFESSOR DR: W. WUEST DFVLR-INST. DYN. VERDUENNTER GASE 3400 GOETTINGEN. BUNSENSTRASSE 10 RFA

DR. KYOJI YAMAMOTO DEPT. OF MECHANICAL ENGINEERING SCHOOL OF ENGINEERING, OKAYAMA UNIV. OKAYAMA JAPAN

PROFESSOR SHEE-MANG YEN UNIVERSITY OF ILLINOIS AT URBANA-CHAMPAIGN URBANA, ILLINOIS \$1801 USA

۰.

DR. KENNETH K. YOSHIKAWA NASA-AMES RESEARCH CENTER N 230-3 MOFFETT FIELD CALIFORNIA 94035 USA

PROFESSOR TOR YTREHUS NTH-INSTITUTT FOR MEKANIKK NORGES TEKNISKE HOGSKOLE 7034 TRONDHEIM-NTH NORWAY

Dr. Bernard Zappoli Societe Europ de Propulsion 27207 Vernon, B.P. 802 France

,5.7

MCNDAY - INELASTIC SCATTERING

- 8.30 * K. Bergmann (Kaiserslautern) LASER SPECTROSCOPY AND INELASTIC COLLISIONS
- 9.00 ** U. Buck (MPI Göttingen) MOLECULAR SCATTERING FROM NON-SPHERICAL POTENTIALS
- 10.00 * D. Beck, U. Ross, W.Schepper (Bielefeld) ON THE BULGE EFFECT OF MOLECULAR SCATTERING
- 10.30 COFFEE BREAK

MINISYMPOSIUM ON ELASTIC SCATTERING

- 11.00 * J.J.H. v.d.Biesen, E.H.v.Veen, F.A.Stokvis, C.J.N.v.d. Meijdenberg (Leiden) MEASUREMENT OF THE GLORY STRUCTURE IN THE TOTAL CROSS SECT. OF AR-KR, KR-KR, KR-XE AND XE-XE
- 11.30 U. Buck, G:Maneke, J.Schleusener (MPI Göttingen), R.A. Aziz (Waterloo) G.Scoles (Waterloo - Trento) U.Valbusa (Waterloo)

ON THE QUESTION OF THE WELL DEPTH OF THE HEAY INTERATOMIC POTENTIAL

- 11.45 F. Torello, M.G. Dondi (Genova) A NEW H-H SPHERICAL POT. FROM HIGH RES. MOL.BEAM EXPERI-MENTS
- 12.00 B. Brunetti, G. Liuti, E.Luzzatti, F.Pirani, F.Vecchiocat tivi (Perugia) ABSOLUTE TOTAL CROSS SECT. FOR $0 - 0_2, 0_2 - 0_2, 0 - N_2, 0_2 - N_2$, O - N2, 02 - N2 COLLISIONS
- 12.15 R. Düren; H.O.Hoppe, H.Tischer (MPI Göttingen) OBSERVATIONS OF THE "ORBITING" PHENOMENON AS A BACKWARD PEAK IN THE DIFFERENTIAL CROSS SECTION
- 12.30 U. Schwalm, J.P. Toennies (MPI Göttingen) ORBITING RESONANCES IN THE SCATT. OF H ATOMS FROM MOLECU LES
- 12.45 H.C.W. Beijerinck, P.M.A. v.d.Kam, N.F. Verster (Eindhoven) THE SMALL ANGLE DIFF. CROSS-SECTION FOR AN INVERSE POWER POTENTIAL.

MINISYMPOSIUM ON NON SPHERICAL POTENTIALS

17.00 G. Rotzoll, A. Lübebrt (Hannover) FITTING OF TOTAL DIFFERENTIAL ATOM-MOLECULE CROSS SECT. WITH ANISOTROPIC POTENTIALS

* = Invited paper

- MONDAY -

- 17.15 H. Thuis (Nijmegen) ANISOTROPY OF NO-INERT GAS SYSTEMS
- 17.30 W.P. Kræmer (MPI München) QUANTUMCHEMICAL SCF AND CI CALCULATIONS OF THE ENERGY HY-PERSURFACE FOR THE SYSTEM HE-CO
- 17.45 A.Kuppermann, M.Kiel, J.T.Slankas, G.A.Parker (Caltech) ANISOTROPIC POTENTIALS FOR HELIUM BEAMS BY CROSSED MOLECU LAR BEAMS OF N₂, O₂, NO, CO AND CO₂
- 18.00 V.N. Khromov, V.B. Leonas (Moscow) ON THE USE OF SCATTERING DATA AS A TEST OF THE ACCURACY OF AB INITIO CALCULATIONS OF ANISOTROPIC POTENTIALS
- 18.15 W.R. Gentry, M.A. Hoffbauer, C.F.Giese (Minnesota) PULSED MOL. BEAM STUDIES OF STATE-RESOLVED ROT. EXCITATION
- 18.30 J. Andres, U.Buck, F.Huisken, J.Schleusener, F.Torello (MPI - Göttingen) STATE RESOLVED DIFFERENTIAL CROSS SECTIONS FOR 0 →2 ROTA-TIONAL TRANSITIONS IN D₂ + Ne COLLISIONS
- 18.45 J. Schaefer (MPI München) RECENT RESULTS OF He - H₂ (HD) CROSS SECT. FOR EL. AND INE LASTIC SCATTERING BELOW 1.5 eV CENTER OF MASS ENERGY
- 19.00 L. Monchick (Johns Hopkins), J.Schaefer (MPI München) AB INITIO CALCULATIONS OF THE TRANSPORT CROSS-SECTION OF H₂
- 19.15 CHAIRMAN'S REMARKS : G.P. TOENNIES (MPI Göttingen)

- T U E S D A Y -

REACTIVE SCATTERING

- 8.30 * R. Grice (Manchester) REACTIVE SCATTERING OF A SUPERSONIC OXYGEN AYOM BEAM
- 9.00 ** Y.T. Lee (Berkeley) RECENT ADVANCES IN STATE TO STATE REACTIVE SCATTERING
- 9.45 ****** J.C. Polanyi (Toronto) CHEMICAL DYNAMICS; THEORY VS. EXPERIMENT
- 10.30 COFFEE BREAK

MINISYMPOSIUM ON INELASTIC SCATTERING

- 11.00 S.B. Ryali (Connecticut), G.E.Kolb (Aerodyne Corp.), J.B. Fenn (Yale) VIBRATIONAL EXCITATION OF CO, BY COLLISIONAL T-V EXCHANGE 11.15 F.A. Gianturco, U.T.Lamanna, D.Ignazzi (Bari) VIB. EXCITATION OF SIMPLE MOLECULES BY COLLISION WITH PRO TONS : A THEORETICAL STUDY ON CO TARGETS 11.30 J.A.D. Stockdale, R.J. Warmack (Oak Ridge) COLLISIONAL IONIZATION OF Cs AND K BY O 11.45 M.J.P. Maneira, U.Weigmann, A.M.C. Moutinho, K.Lacmann (H. M.I. - Berlin) DOUBLE DIFF. CROSS SECT. FOR K FORMATION IN K + SnCl_COLL. 12.00 J. Krenos (Rutgers) CHEMICAL DYNAMICS OF ENERGY TRANSFER : FORMATION OF Ne IN COLLISIONS OF He (2'S) WITH GROUND-STATE Ne. 12.15 H. Haberland, P. Oesterlin (Freiburg) THE PUMPING PROCESS OF THE HENE-LASER, STUDIED IN DETAIL IN A CROSSED MOLECULAR BEAM EXPERIMENT 12.30 R. Düren; U. Krause, G. Moritz (MPI - Göttingen) ELECTRONIC EXCIT. IN COLLISIONS OF Na, K and Rb WITH Hg 12.45 CHAIRMAN'S REMARKS : V. AQUILANTI (Perugia) MINISYMPOSIUM ON REACTIVE SCATTERING 17.00 D. Brandt, J.C.Polanyi (Toronto) MICROSCOPIC BRANCHING IN REACTIONS H+FY (Y=C1, Br, I) \rightarrow HF (v', J') + Y17.15 J.R. Grover, D.E.Malloy, J.B.A. Mitchell (Brookhaven) APPLICATIONS OF RADIOACTIVE MOL. BEAMS : (1) THE CHEMI -STRY OF ASTATINE; (2) THE MEAS. OF ABSOLUTE DIFF. CROSS. SECT. K.K. Verma, W.C. Stwalley (Iowa) 17.30 THE PRODUCTION OF OH (A Σ^+) FROM COLL. OF SUPERTHERMAL H ATOMS WITH HOO AND O2 17.45 A.C. Luntz, P. Andresen (IBM - San José) DYNAMICS OF THE CHEMICAL REACTIONS OF O (3P) WITH SATURA TED HYDROCARBONS 18.00
 - .00 S. Chapman (Columbia) THEORETICAL STUDY OF Be + HF -----> BeF + H

manual contractions while contract of the second

- TUESDAY -

E. V	Vieta	zke,	M.Erdw	eg, j	.He	uschk	el, 1	Matu	ns, (3,
200	SKIII SCED	I LUU DEAM		TOC .	3.0	CTIID 3 (7		7.8 00 C		rana in
		DLAN			AD:	STRAC.	LTON .	AND 2		
TIU	N LIN	THE	REACTI	Or OF	BL	MTJH	-CH-3	L AND	(CH,	31/2
R. I	Dirso	herl	, H.U.	Lee	(St	uttga	rt)			
MOD	מידרים	D BE	M KTN	TOT	•	DEN (TT)		2020	$\cap \Sigma$	Cm

- 18.30 R. Dirscherl, H.U. Lee (Stuttgart) MOLECULAR BEAM KINETICS : REACTIVE SCATT. OF Sm WITH O₂ AND LASER FLUORESCENCE STUDY OF YDBr.
- 18.45 E.K. Parks, S. Wexler (Argonne) COLLISION-INDUCED DISSOC. OF THE CESIUM HALIDES BY RARE GAS ATOMS AND SF MOLECULES
- 19.00 W.L. Hase, R.J. Wolf (Wayne State) CHEMICAL DYNAMICS OF C₂H₅ DECOMPOSITION
- 19.15 CHAIRMAN'S REMARKS : R.B. BERNSTEIN (Columbia)

-WEDNESDAY-

FEAMS OF CLUSTERS

- 8.30 * E. Schumacher, W.H.Gerber, A.Herrmann, M.Hoffmann, S.Leutwyler, L.Wöste (Bern) SPECTROSCOPY OF METAL CLUSTERS IN MOLECULAR BEAMS
- 9.00 ****** D. Herschbach (Harvard) MOLECULAR CLUSTER BEAMS
- 10.00 * Dr. F.M.Devienne, M. Teisseire (Peymeinade) METAL AGGREGATES FORM. BY HIGH ENERGY MOL. BEAMS BOMBARDMENT
- 10.30 COFFEE BREAK

18.15

MINISYMPOSIUM ON CLUSTERS

- 11.00 K. Sattler, J.Mühlbach, A.Reyes Flotte, E.Reckna gel (Konstanz) A SOURCE FOR METAL ATOM AGGLOMERATES OF UNIF. MASS
- 11.15 S. Romano, E. Clementi (Donegani) MONTE CARLO SIMULATION OF SMALL ION-WATER CLUSTERS
- 11.30 G.L. Griffin, J.Kratsios, R.P.Andres (Princeton) PRODUCTION AND SIZE ANALYSIS OF MOL. BEAMS OF ME -TAL CLUSTERS
- 11.45 * R. Monot, E.R.Dietz, A.R. George, W.D.Knight (Berkeley) BEAM OF POTASSIUM CLUSTERS : STERN-GERLACH EXPERIMENT.

ىرىيى يەرىپىدىغان قۇللىغانىيە بىلىش يېلىغانىيە 1973 - ئەرىپىدىغان قۇللىغانىيە بىلىش يېلىغانىيە

AII-36

-WEDNCSDAY-

- 12.15 J Spann, H.Vollmar (Karlsruhe) MUTASTABLE CLUSTERS
- 12.30 M. Cavallini, G. Grillo (Assoreni) TWIN MOL. BEAM STUDY OF OXIDATION REACTIONS ON TRANS.ME-TALS
- 12.45 L. Holmlid, J.O. Olsson (Göteborg) SURFACE CATALYZED REACTIONS ON CARBON COVERED Pt(88W)

MINISYMPOSIUM ON ION AND MUONIUM CHEMISTRY

- 11.00 * D.M. Garner, D.G. Fleming (Vancouver) MUONIUM REACTION KINETICS : HYDROGEN ISOTOPE EFFECTS AND CHEMICAL DYNAMICS IN STUDIES OF Mu + X₂ AND Mu+ HX
- 11.30 J.N.L. Connor, A.Laganà (Manchester) QUASICLASSICAL DYNAMICS OF LIGHT HEAVY-HEAVY ATOM REAC-TIONS - THE REACTIONS Mu + F₂ AND Mu + Cl₂
- 11.45 H. Schmidt, M. Konrad, F. Linder (Kaiserslautern) CROSSED-BEAM MEASUREMENTS OF CHARGE TRANSFER REACTIONS FOR He + N AND He + O IN THE 1 eV COLLISION ENERGY RANGE.
- 12.00 I. Kusunoki, Ch. Ottinger (MPI Göttingen) CHEMILUMINESCENT ION-MOLECULE REACTIONS : ROT.-VIB. STA TE DISTRIBUTIONS OF CH (A)FROM C + H₂ (D₂) COLLI -SIONS.
- 12.15 W. Lindinger, E. Alge, H.Villinger, H.Störi (Innsbruck); D.L. Albritton, F.C.Fehsenfeld (Boulder) DEPENDENCE OF ION-MOLECULE REACT. ON THE ION VIB. ENERGY
- 12.30 * A. Ding, A.Redpath, U.Steinmetzger (H.M.I. Berlin) INFRARED CHEMILUMINESCENCE FROM ION MOLECULE REACTIONS

AII-37

- THURSDAY -

SURFACE SCATTERING

- 8.30 * F. Tommasini (Genova), U.Valbusa (Waterloo) ATOMIC HYDROGEN SCATTERING FROM CRYSTAL SURFACES
- 9.00 ****** V. Celli (Virginia) THE DYNAMICS UP GAS-SURFACE INTERACTIONS
- 9.45 ** L. Wharton (Chicago) SURFACE REACTIONS AND MOLECULAR BEAMS
- 10.30 COFFEE BREAK
- 11.00 POSTER SESSION (in alph. order of 1st author)

D.H.H. Al-Amiedy, D.E. Dugdale, D.C. Lainé (Keele) ELECTROSTATIC STATE SELECTION OF MOLECULES EMPLOYING A SINGLE WIRE HELIX.

E. Alge, H. Villinger, H.Helm, W.Lindinger (Innsbruck), D.L. Albritton, F.C. Fehsenfeld (Boulder) DRIFT TUBE MEAS. : THE LINK BEIWEEN THERMAL RATE CONST. AND CROSS SECT. OBTAINED IN BEAM EXPERIMENTS.

B. Andresen (Copenhagen), S.Hultberg, B.Jelencovic, L.Lilje by, S.Mannervick, E. Veje (Stokholm) DISSOCIATION AND EXCITATION OF SMALL MOLECULES UPON PASSA -GE THROUGH THIN FOILS

D. Bassi (Trento), M. Cavallini, G. Grillo (Assoreni) THE CATALYTIC OXIDATION OF CO ON PALLADIUM SURFACES

D. Bassi, A. Boschetti, S.Marchetti, M.Zen(Trento)T.E.Gough, R.E. Miller (Waterloo), G. Scoles (Waterloo & Trento) I.R. SPECTROSCOPY OF SUPERSONIC MOL. BEAMS WITH TUNABLE DIODE AND COLOR CENTRE LASERS

K.Bergmann, R.Engelhardt, U.Hefter, P.Hering (Kaiserslautern)

MOL. BEAM DIAGNOSTIC WITH INTERNAL STATE SELECTION : PER-PENDICULAR VEL. DISTRIBUTION IN A NA/NA SUPERSONIC BEAM

G. Brusdeylins, R.B.Doak (M.P.I. Göttingen) TOTAL INTEGRAL CROSS SECTIONS FOR He - CO

U.Buck, L.Mattera, D.Pust (M.P.I. Göttingen), D. Haaks (Wup pertal)

ULTRAVIOLET EMISSION IN Xe + Xe COLLISIONS NEAR THRESHOLD

J.M.Calo (Princeton) CRYOGENIC DEPOSITION AND DESORPTION STUDIES OF STRATOSPHE DICALLY DETENDED SDECTES

G. Caracciolo, T.H.Ellis, G.O.Este, G.Knight, G.Scoles, U. Valbusa (Waterloo)

SCATTERING EXPERIMENTS WITH H ATCMS. NON-SPHERICAL SY-STEMS : $H + C_2H_6$, $H + C_3H_8$, $H + CO_2$, $H + CO_2$.

M.A.A.Clyne, SJ.Davis, M.C.Heaven, I.S. McDermid (Queen Mary Coll.)

QUANTUM-RESOLVED DYNAMICS IN DIATOMIC HALOGENS AND INTER-HALOGENS, USING DOPPLER-LIMITED LASER EXCITATION.

C.Dankert, H.Legge (Göttingen) INVESTIGATION OF A BEAM SKIMMED AT SMALL KNUDSEN NUMBERS

A.Ding (H.M.I. Berlin) A SEMICLASSICAL INVERSION PROCEDURE FOR STUECKELBERG O -SCILLATIONS

E.N.Evlanov, Y.V. Lebedev, V.B. Leonas (Moscow) "COLD" BEAM OF HYDROGEN ATOMS

J.D. Ganière, R.Monot, R.Rechsteiner (Lausanne) BEAMS OF NA CLUSTERS : TIME OF FLIGHT MASS SPECTROMETRY

F.A. Gianturco, U.T.Lamanna, A.Attimonelli A SCATTERING-ORIENTED POTENTIAL ENERGY SURF. STUDY FOR VIB-ROT EXCITATIONS OF DIATOMICS BY H⁺.

H. Helm, K.Stephan, T.D.Märk (Innsbruck) MASS SPECTROMETRIC INVESTIGATION OR RARE GAS DIMER BEAMS

M.Hofmann, S.Leutwyler, E.Schumacher (Bern)W.Schulze (Berlin)

MATRIX SPECTROSCOPY OF NA AND K CLUSTERS FORMED IN A SU-PERSONIC MOLECULAR BEAM

A.P. Kalinin, V.N. Khromov, V.B.Leonas (Moscow), R.W. Wiynaendts van Resandt, J.Los (Amsterdam) DIFFERENTIAL SCATTERING OF Li IONS ON No MOLECULES

P.M.A. van der Kam, H.C.W. Beijerinck, N.F. Verster (Eindhoven)

MONTE CARLO CALCULATION OF ANGULAR RESOLUTION FUNCTIONS

M.E.Koch, W.C.Stwalley (Iowa) MULTIPHOTON IONIZATION OF Li2

M.J.P. Maneira, A.J.F. Praxedes, A.M.C.Moutinho (Lisboa) DIFF. CROSS. SECT. FOR ION PAIR FORM. IN K+CCl, COLLISIONS

D.Monzel, V.Ramakrishan (Garching) THERMAL ACCOMODATION OF HELIUM AND NEON ADSORBATE-COVERED TUNGSTEN SURFACES.

R.A.R. Porter, A.E.Grosser (McGill) CH, MDLECULAR BEAM SOURCE

مآند

P.G.A. Theuws, C.E.E.Pernot, H.C.W.Beijerinck, N.F.Verster (Eindhoven)

EXCITATION AS A DETECTION MECHANISM FOR GROUND STATE PARTICLES

P.G.A.Theuws, C.E.E.Pernot, H.C.W. Beijerinck, D.C.Schram, N.F. Verster (Eindhoven)

PRODUCTION OF METASTABLES IN A HOLLOW CATHODE ARC DISCHAR-GE

J. Verberne (Nijmegen) SPECTROSCOPY ON (H₂) 2 DIMERS

A.E.Zarvin, R.G. Sharafutdinov (Novosibirsk) DIRECT EXP. VERIFICATION OF SUPERSONIC MOL. BEAM FORMA -TION MODEL

MINISYMPOSIUM ON GAS-SURFACE INTERACTIONS

- 17.00 * M.J.Cardillo (Bell Labs.) THE DIFFRACTION OF HE ATCMS FROM SI SINGLE CRYSTALS
- 17.30 H.Conrad, G.Ertl, J.Küppers, S.W.Wang (München), K.Gérard, H. Haberland (Freiburg) PENNING IONIZATION ELECTRON SPECTROSCOPY OF CLEAN AND CO-COVERED METAL SURFACES
- 17.45 E. Ficocelli-Varracchio (Bari) FIELD-THEORETIC APPROACH TO SCATTERING OFF SOLID SURFACES
- 18.00 A.C. Levi (Genova and Trento) DEBYE-WALLER FACTOR AND INFLASTIC ATOM-SURFACE SCATTERING
- 18.15 A.Armand, J.Lapujoulade, Y.Lejai, N. Papanicolau (Saclay) THERMAL DEPENDENCE OF THE SPECULAR PEAK OF HELIUM SCATTE-RED FROM (100) COPPER
- 18.30 E. Semerad, E.M. Hörl (Seibersdorf) DEBYE-WALLER FACTOR IN Ne-BEAM SCATT. BY A LIF SURFACE
- 18.45 G. Boato, P. Cantini, C.Guidi, R.Colella (Genova) CHARGE DENSITY WAVES OBSERVED BY MOL. BEAM DIFFRACTION
- 19.00 L. Greiner, H.Hoinkes, H. Wilsch (Erlangen) CORRELATION OF SPEC. AND DIFF'. INTENSITY AT SELECTIVE AD-SORPTION EXP. INVESTIGATIONS WITH D ON LiF (001)

65

19.15 CHAIRMAN'S REMARKS : G. BOATO (Genova)

AII-40

- F R I D A Y =

BEAM-PHOTON INTERACTIONS

- 8.30 * W.R. Gentry, M.A. Hoffbauer, C. Giese (Minneapolis) PHOTODISSOCIATION OF v.d. WAALS DIMERS IN PULSED MOL.BEAMS
- 9.00 ** V.S. Letokhov (Moscow) MULTISTEP PHOTOIONIZATION OF MOL. BEAMS BY LASER LIGHT
- 9.45 ****** J. Durup (Orsay) PHOTODISSOCIATION
- 10.30 COFFEE BREAK

MINISYMPOSIUM ON PHOTOIONIZATION AND PHOTODISSOCIATION

- 11.00 * N.J.A. van Veen, M.S. de Vries, A.E. de Vries (F.O.M. Amsterdam) PHOTOFRAGMENTATION OF DIATOMIC HALOGEN COMPOUNDS
- 11.30 R.B. Bernstein (Columbia) LASER MULTIPHOTON IONIZ. AND FRAGMENT. OF MOL. BEAMS
- 11.48 R.K. Sparks, L.R. Carlson, K.Shobatake, M.L. Kowalczyk, Y.T. Lee (Berkeley) DYNAMICS OF PHOTODISSOCIATION OF 0,
- 12.06 S. Leutwyler, M. Hoffmann, E. Schumacher (Bern) SEQUENTIAL TWO-PHOTON-IONIZATION OF NAK AND K IN SUPER-SONIC MOLECULAR BEAMS
- 12.24 I.M. Beterov, Yu.V.Brzhazovskii, V.P.Chebotaev, A.K.Re brov, B.E. Semvachkin, A.A. Vostrikov INFLUENCE OF CO_LASER RADIATION ON SF_6 CONDENSATION AND MOLECULAR BEAM INTENSITY.
- 12.42 J.R. Grover, J.B.A. Mitchell (Brookhaven) A VERSATILE USER-ORIENTED ATOMIC AND MOL. BEAM APPARATUS FOR USE WITH THE NAT. SYNC. LIGHT SOURCE.