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FUNDAMENTALS OF ARCJET THRUSTER THERMOPHYSICS

prepared by

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Abstract

This final technical report summarizes the research carried out by our group concerned with arcjet plasma propulsion. Section I describes the non-equilibrium plasma model and a detailed formulation for kinetic non-equilibrium. Section II describes experimental measurements in which measurements of electron number density, electron temperature, ion speed ratio and gas velocity are carried out by advanced Langmuir probes. Section III outlines our research with a modified 1-2 kW class arcjet, where surface-mounted Langmuir probes are used to investigate the constrictor and nozzle current attachment regions.

Overview

A general, two-dimensional, steady, non-LTE (local thermodynamic equilibrium) model is presented for nitrogen/hydrogen arcjet thrusters. Both thermal and chemical nonequilibrium processes are incorporated to describe a multi-temperature, multi-species N₂/H₂ plasma. The current and anode temperature distributions are fully coupled to the plasma flowfield with realistic boundary conditions. Thermal nonequilibrium, chemical equilibrium results are presented for simulated hydrazine. Results for both thermal and chemical nonequilibrium hydrazine and hydrogen arcjets show significant thermal nonequilibrium near the anode, and high frozen flow losses. It is concluded that nonequilibrium chemistry models are necessary to capture accurately the interior distributions of the plasma flowfield.

With recent advances made in low power (1-2 kW) arcjet numerical modeling, validation of numerical models through experimentation has become increasingly important. We describe multiple electrostatic probe techniques for characterizing flow conditions at the exit plane of a 1 kW-class hydrazine thruster. An improved electrostatic quadruple probe technique, which is also modified to account for plasma gradients over the finite probe volume and multiple ion species, provides measurements of ne, Te and ui/cm at the thruster exit. Quadruple probe results are also presented for measurements at several centerline locations immediately downstream of the thruster exit plane. A spatially-resolved time-of-flight electrostatic probe technique is employed for measurements of radial profiles of the plasma axial velocity ui. The quadruple probe results (Te, ne, ui/cm), coupled with independent measurements of plasma velocity, are used to estimate ion temperature Ti=Tg and the extent of thermal non-equilibrium at the thruster exit. Exit plane experimental data for ui, Te and Tg. The model underpredicts Te and a direct comparison of ne results is precluded by large axial gradients at the thruster exit.

A 1.4 kW arcjet thruster has been designed for internal probing of the nozzle. The arcjet employs a single-piece nozzle body, facilitating modeling of the anode heat transfer. Flush-mounted planar single Langmuir probes are used for internal diagnostics of the boundary layer flow inside the arcjet nozzle. Two nozzle probes and one constrictor probe are implemented to obtain n_e, T_e, current density and sheath voltage data in the arcjet for $11 \le P/m \le 20$ and flow rates of 60 and 85 mg/sec. Comparison between the experimental results and a computational model is made. Based on the data obtained with a nozzle probe the sheath voltage is predominantly negative, so that an electric field exists near the anode which retards the electrons. The results indicate a high degree of ohmic heating at the sheath edge, especially for positive sheath voltages.

The primary interest of the Air Force for arcjet propulsion is with storable propellants. The research reported here emphasizes gaseous mixtures of N_2 and H_2 for the simulation of N_2H_4 and H_2 arcjets. Because the arcjet is a low Reynolds number device (Re<1000), viscous behavior dominates energy transfer processes in the thermal and viscous nozzle boundary layers, which couples with multi-dimensional heat conduction in and from the massive conducting anode wall. Therefore, energy transfer must be described by coupled Navier-Stokes conservation relations which depend in a complex way on the fluid properties. Previously, no single computational model could be said to describe accurately the entire physics of arcjet flows.

The energy transport from the plasma located in the equilibrium constrictor region of the arcjet is a strong function of the number density and distribution of electrons from ionized species. The degree of non-equilibrium can significantly alter the expected electron number density. The diffusion of species, particularly electrons, is likewise important. The electric current distribution, not known, is strongly coupled to the n_e and T_e distribution in the vicinity of the relatively cold constrictor and anode walls. Thermal and chemical non-equilibrium plays a dominant role, resulting in spatial variation in the current attachment and anode heating. The proper description of this aspect of arcjet nozzle flow has proven particularly difficult to achieve in numerical models of the arcjet.

I. A Plasmadynamics Model for Nonequilibrium Processes in N2/H2 Arcjets

(AIAA Paper No. 95-1961)

Abstract

A general, two-dimensional, steady, non-LTE (local thermodynamic equilibrium) model is presented for nitrogen/hydrogen arcjet thrusters. Both thermal and chemical nonequilibrium processes are incorporated to describe a multi-temperature, multi-species N_2/H_2 plasma. The current and anode temperature distributions are fully coupled to the plasma flowfield with realistic boundary conditions. Thermal nonequilibrium, chemical equilibrium results are presented for simulated hydrazine. Results for both thermal and chemical nonequilibrium hydrazine and hydrogen arcjets show significant thermal nonequilibrium near the anode, and high frozen flow losses. It is concluded that nonequilibrium chemistry models are necessary to capture accurately the interior distributions of the plasma flowfield.

Nomenclature

- c_i mean thermal speed (m/s)
- C coulomb charge
- D_i effective diffusion coefficient (m²/s)

elst elastic energy transfer (W/m^3)

- g_0 gravitational acceleration at sea level (m/s²)
- h Planck's constant (J-s)
- h_i enthalpy of species i(J/kg)
- H heavy species total particle energy (J)
- I integrated total current (A)

I_{sp} specific impulse (s)

j current density (A/m^2)

| k _b | Boltzmann's constant (J/K) |
|------------------|--|
| k _c | Coulomb constant (Nm ² /C ²) |
| k f | Forward rate coefficient $(m^{6}/s \text{ or } m^{3}/s)$ |
| k _r | Reverse rate coefficient $(m^{6}/s \text{ or } m^{3}/s)$ |
| Le | Electron inelastic loss rate (W/m ³) |
| L _h | Heavy species inelastic loss rate (W/m ³) |
| mi | mass of species i (kg) |
| m _{ij} | reduced mass (kg) |
| ṁ | mass flow rate (kg/s) |
| Μ | Mach number |
| ni | number density of ith species (m ⁻³) |
| р | pressure (N/m ²) |
| P_{Ω} | integrated ohmic power deposition (W) |
| Q _{ij} | collision cross section (m ²) |
| Ta | anode temperature (K) |
| Te | electron temperature (K) |
| Tg | heavy species temperature (K) |
| Т | thrust (N) |
| u | axial velocity (m/s) |
| v | radial velocity (m/s) |
| v | velocity vector (m/s) |
| v | voltage (volts) |
| w | azimuthal swirl velocity (m/s) |
| x,r,θ | cylindrical coordinates |
| xi | mole fraction of species i |
| Уi | mass fraction of species i |
| x _{con} | constrictor length (mm) |

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 x_{mol} mixture parameter in $x_{mol}N_2 + H_2$

- Z_i thermodynamic partition functions
- α_i catalytic efficiency of species i
- ϵ_d dissociation energy (J)
- ϵ_i ionization energy (J)
- ε anode emissivity
- λ_a anode thermal conductivity (W/m-K)
- λ_e electron thermal conductivity (W/m-K)
- λ_h heavy species thermal conductivity (W/m-K)
- Λ_e Spitzer logarithm term
- μ gas viscosity (kg/m-s)
- v collision frequency (s⁻¹)
- θ_{vH2} H₂ vibrational temperature (K)
- θ_{vN2} N₂ vibrational temperature (K)
- ρ mass density (kg/m³)
- σ electrical conductivity (mhos/m)
- τ stress tensor (N/m²)
- $\overline{\Omega}_{ii}$ collision integral (m²)

I. Introduction

Electrothermal arcjets offer a significant advantage in specific impulse and cost over conventional satellite propulsion systems. Low power 1-2 kW systems have been flight qualified and are now used for north-south station keeping, while higher power 20-30 kW designs are in advanced development for orbit transfer applications.¹

The arcjet is shown schematically in Fig. 1. The fluid dynamics and energy transfer within the arcjet thruster are nonlinear and strongly coupled. The propellant is injected upstream of the constrictor, typically with an azimuthal (or 'swirl') velocity component. The energy conversion mechanism for an arcjet is an arc discharge, produced by a voltage difference between the thruster nozzle, which is the anode, and a conical cathode on the upstream side of the constrictor. Typical operating voltages for 1 kW class arcjets are 100 V DC at a current of I = 10 A. The arc current distribution depends on several factors in addition to the geometry. The distribution is coupled to the propellant mass flow rate, composition, thermal properties, electrical conductivity, and gas-dynamic properties.

Both chemical and thermal processes in the arcjet are described as nonequilibrium. Arc current is converted to electron thermal energy through ohmic dissipation. The electrons transfer thermal energy to heavy species in the arc plasma through collisions. This energy is then converted to thrust as the fluid accelerates through the nozzle. In regions of low pressure and/or elevated ohmic heating, collisional coupling between electrons and heavy species may not establish equal gas and electron temperatures. Additionally, flow velocities are large so that fluid residence times in the nozzle are of the order of a few μ s; much of the energy invested in dissociation and ionization is frozen. Pressures drop from roughly 1 atm at the constrictor to millitorr conditions at the exit; radial diffusion of electrons from the arc core is important in determining the arc structure.

The thrust and specific impulse produced by an arcjet are determined by several factors including the power transferred to the propellant, the extent of both kinetic and chemical

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nonequilibrium, the propellant gas mixture, and the arcjet nozzle geometry. Other factors which contribute to the overall performance include the thermal loading of the anode,² and the voltage sheaths at the electrodes. A comprehensive model of an arcjet thruster is highly complex, including plasmadynamic, fluid dynamic, radiative, and surface and volumetric heat transfer phenomena.

Most arcjet design strategies to date have been empirical, with design improvements based on experimental observations. The need for a more complete description of the complex physical processes has resulted in a variety of numerical efforts ranging from simplified one-dimensional models³⁻⁸ to more comprehensive two-dimensional descriptions of the gas dynamics and energy transfer processes.⁹⁻²² Comprehensive arcjet models are required to (1) interpret experimental results, (2) understand the physical processes in regions of the thruster where diagnostic techniques are challenging, (3) improve arcjet performance and (4) scale the devices to higher or lower power levels.

II. Nonequilibrium Model

An axisymmetric, steady, laminar, continuum flow, two-temperature kinetic nonequilibrium model is formulated for a direct current arcjet with flow swirl, a variable nozzle geometry, and a variable mixture ratio of nitrogen and hydrogen. Flow swirl is modeled by including an azimuthal momentum equation. Two methods are used to determine species populations. If chemical equilibrium is assumed, the species populations are computed from a generalized law of mass action. For chemical nonequilibrium simulations, populations are determined from species continuity equations.

The model predicts thrust, specific impulse, and internal fields for pressure p, mass density ρ , seven species densities n_i , and anode T_a , electron T_e and heavy species T_g temperatures. Additionally, the model predicts the current density distribution **j**, voltage potential V, and velocity components u, v, and w.

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The details of the model are presented in six sections below as: (1) the fluid dynamic equations, (2) the energy equations, (3) the electromagnetic equations, (4) the species number density equations, (5) transport coefficients, and (6) boundary conditions. The arcjet geometry is described as (x,r,θ) where x is the axial coordinate, r is the radial coordinate, and θ is the azimuthal coordinate.

1. Fluid Dynamic Equations

The viscous fluid dynamic equations are summarized below:

Axial momentum:

$$\frac{\partial(\rho u u)}{\partial x} + \frac{1}{r} \frac{\partial(r\rho u v)}{\partial r} = \frac{4}{3} \frac{\partial}{\partial x} \left(\mu \frac{\partial u}{\partial x} \right) + \frac{1}{r} \frac{\partial}{\partial r} \left(\mu r \frac{\partial u}{\partial r} \right) - \frac{2}{3} \frac{\partial}{\partial x} \left(\mu \frac{\partial v}{\partial r} \right) - \frac{2}{3r} \frac{\partial(\mu v)}{\partial x} + \frac{1}{r} \frac{\partial}{\partial r} \left(\mu r \frac{\partial v}{\partial x} \right) - \frac{\partial p}{\partial x}$$
(1)

Radial momentum:

$$\frac{\partial(\rho uv)}{\partial x} + \frac{1}{r} \frac{\partial(r\rho vv)}{\partial r} = \frac{\partial}{\partial x} \left(\mu \frac{\partial v}{\partial x} \right) + \frac{4}{3r} \frac{\partial}{\partial r} \left(\mu r \frac{\partial v}{\partial r} \right) + \frac{\partial}{\partial x} \left(\mu \frac{\partial u}{\partial r} \right) + \frac{2\mu}{3r} \frac{\partial u}{\partial x} - \frac{2v}{3r} \frac{\partial \mu}{\partial r} - \frac{4}{3} \frac{\mu v}{r^2} - \frac{2}{3r} \frac{\partial}{\partial r} \left(\mu r \frac{\partial u}{\partial x} \right) - \frac{\partial p}{\partial r} + \frac{\rho w^2}{r}$$
(2)

Azimuthal momentum
$$(\partial p/\partial \theta = 0)$$
:

$$\frac{\partial(\rho u(wr))}{\partial x} + \frac{1}{r} \frac{\partial}{\partial r} ((r\rho v + 2\mu)(wr)) = \frac{\partial}{\partial x} \left(\mu \frac{\partial(wr)}{\partial x} \right) + \frac{1}{r} \frac{\partial}{\partial r} \left(\mu r \frac{\partial(wr)}{\partial r} \right)$$
(3)

Axisymmetric continuity:

$$\frac{\partial}{\partial x}(\rho u) + \frac{1}{r\partial r}(r\rho v) = 0$$
(4)

2. Energy Equations

The assumption of kinetic nonequilibrium gives separate energy equations for the electrons and heavy species. For the chemical equilibrium model, the electron energy equation is:

$$\nabla \cdot \left(\left(n_e \mathbf{v} - \frac{\mathbf{j}}{e} \right) C_{pe} T_e \right) = \nabla \cdot \left(\lambda_e \nabla T_e \right) + \nabla \cdot \left(\frac{1}{m_e} \rho C_{pe} T_e D_e \nabla y_e \right) + \frac{\mathbf{j} \cdot \mathbf{j}}{\sigma} - \text{elst - radiation}$$
(5)

where 'elst' represents energy lost through elastic collisions with the heavy species,²³ and 'radiation' denotes optically thin radiation loss due to continuum bremsstrahlung.²⁴ The elastic energy transfer source term is calculated as the sum of electron-molecule, -atom, and -ion contributions as:

elst =
$$3k_b(T_e - T_g)n_e m_e \sum_{i \neq e} \frac{\overline{v}_{ei}}{m_i}$$
 (6)

where the average collision frequencies \overline{v}_{ei} between electrons and heavy species are calculated using the mean electron thermal speed and collision cross sections Q_{es} as:²⁴

$$\overline{\nu}_{ei} = \sqrt{\frac{8k_b T_e}{\pi m_e}} n_i Q_{ei}$$
⁽⁷⁾

where the electron-ion cross sections are:

$$Q_{e-ion} = \frac{\pi e^4 k_c^2 \ln \Lambda_e}{2(k_b T_e)^2}$$
(8)

Electron-molecule collision frequencies are multiplied by an energy loss factor δ to account for inelastic losses to the internal energy modes.^{8, 25} A constant loss factor of δ = 3000, chosen for the simulations presented here, is discussed below.

The chemical equilibrium energy equation for heavy species (molecules, atoms, and ions) is:

$$\nabla \cdot (\mathbf{n}_{h} \mathbf{v} \mathbf{H}) = \nabla \cdot (\lambda_{h} \nabla \mathbf{T}_{g}) + \sum_{i \neq e} \frac{1}{m_{i}} \nabla \cdot (\rho(\mathbf{C}_{pi} \mathbf{T}_{g} + \mathbf{h}_{i}^{0}) \mathbf{D}_{i} \nabla \mathbf{y}_{i}) + \text{elst} + \nabla \cdot (\mathbf{v} \cdot \tau)$$
(9)

where

$$\mathbf{H} = \frac{1}{n_{h}} \sum_{i \neq e} \left(n_{i} \left(C_{pi} T_{g} + h_{i}^{0} + \frac{m_{i} \mathbf{v} \cdot \mathbf{v}}{2} \right) \right)$$
(10)

The ionization and dissociation energies of molecules and atoms are arbitrarily included in the heavy species energy equation as reference enthalpies h_i^0 . The reference enthalpies and specific heats are summarized in Table 1.

For chemical nonequilibrium simulations, a finite rate chemistry model is used to determine inelastic energy exchange rates between the particle classes. This model is summarized in Table 2, which lists the individual reaction rates and energy exchanges assumed for each chemical process.²⁶⁻²⁹ The species continuity equations are multiplied by the reference enthalpies associated with each chemical reaction, and the terms appearing in the energy equations as the convection and diffusion of ionization and dissociation potentials are replaced with energy transfer rates. The electron energy equation for chemical nonequilibrium is then

$$\nabla \cdot \left(\left(n_e \mathbf{v} - \frac{\mathbf{j}}{e} \right) (\mathbf{C}_{pe} \mathbf{T}_e) \right) = \nabla \cdot \left(\lambda_e \nabla \mathbf{T}_e \right) + \nabla \cdot \left(\frac{1}{m_e} \rho \mathbf{C}_{pe} \mathbf{T}_e \mathbf{D}_e \nabla \mathbf{y}_e \right)$$

$$+ \frac{\mathbf{j} \cdot \mathbf{j}}{\sigma} - \text{elst} - \text{radiation} - \sum L_e$$
(11)

where individual energy loss rates L_e are summarized in Table 2.

The chemical nonequilibrium heavy species energy equation is:

$$\nabla \cdot (\mathbf{n}_{h}\mathbf{v}\mathbf{H}) = \nabla \cdot (\lambda_{h}\nabla \mathbf{T}_{g}) + \sum_{i \neq e} \frac{1}{m_{i}}\nabla \cdot (\rho \mathbf{C}_{pi}\mathbf{T}_{g} \ \mathbf{D}_{i}\nabla \mathbf{y}_{i})$$

+ elst + $\nabla \cdot (\mathbf{v} \cdot \boldsymbol{\tau}) - \sum \mathbf{L}_{h}$ (12)

where the total energy is now:

$$H = \frac{1}{n_{h}} \sum_{i \neq e} \left(n_{i} \left(C_{pi} T_{g} + \frac{m_{i} \mathbf{v} \cdot \mathbf{v}}{2} \right) \right)$$
(13)

The anode temperature distribution is determined from the energy equation for heat conduction in a solid. Neglecting ohmic dissipation in the anode gives

$$\nabla \cdot \left(\lambda_{a} \nabla T_{a}\right) = 0 \tag{14}$$

where $\lambda_a = \lambda_a(T_a)$.³⁰

3. Species Equations

For chemical equilibrium (CE) simulations, the species number densities are given as a function of the pressure p, the heavy species temperature T_g , and the electron temperature T_e . A seven-species plasma composition of N₂, H₂, N, H, N⁺, H⁺, and electrons is assumed. Dissociation and ionization equations are constructed from a generalized law of mass action as given by entropy maximization.³¹ This yields the following equations for the nitrogen and hydrogen species:

$$\frac{n_{\rm H}^2}{n_{\rm H_2}} = \exp\left(\frac{-\varepsilon_{\rm d,H_2}}{k_{\rm b}T_{\rm g}}\right) \left(\frac{\pi m_{\rm H} k_{\rm b} T_{\rm g}}{h^2}\right)^{3/2} \left(\frac{Z_{\rm H,ex}^2}{Z_{\rm H_2,ex} Z_{\rm H_2,rot} Z_{\rm H_2,vib}}\right)$$
(15a)

$$\frac{n_{N}^{2}}{n_{N_{2}}} = \exp\left(\frac{-\varepsilon_{d,N_{2}}}{k_{b}T_{g}}\right) \left(\frac{\pi m_{N}k_{b}T_{g}}{h^{2}}\right)^{3/2} \left(\frac{Z_{N,ex}^{2}}{Z_{N_{2},ex}Z_{N_{2},rot}Z_{N_{2},vib}}\right)$$
(15b)

$$\frac{n_e n_{H+}}{n_H} = 2 \exp\left(\frac{-\varepsilon_{i,H}}{k_b T_e}\right) \left(\frac{2\pi m_e k_b T_e}{h^2}\right)^{3/2} \left(\frac{Z_{H+,ex}}{Z_{H,ex}}\right)$$
(15c)

$$\frac{n_{e}n_{N+}}{n_{N}} = 2exp \left(\frac{\epsilon_{i,N}}{k_{b}T_{e}}\right) \left(\frac{2\pi m_{e}k_{b}T_{e}}{h^{2}}\right)^{3/2} \left(\frac{Z_{N+,ex}}{Z_{N,ex}}\right)$$
(15d)

In the above expressions, the excitation partition functions Z_{ex} are functions of T_e , while Z_{rot} and Z_{vib} are functions of T_g . Details for the calculation of the various partition functions can be found in References 24 and 31-34.

The plasma equation of state, quasi-neutrality, and the definition of the mixture parameter x_{mol} provide the additional relations required to solve for the seven unknown species. Note that the initial mixture composition is written as $x_{mol}N_2 + H_2$ so that pure hydrogen ($x_{mol} = 0$), simulated hydrazine ($x_{mol} = 1/2$) and simulated ammonia ($x_{mol} = 1/3$) can be easily investigated:

$$x_{mol} = \frac{2n_{N_2} + n_N + n_{N+}}{2n_{H_2} + n_H + n_{H+}}$$
(16)

$$\mathbf{n}_{\mathbf{e}} = \mathbf{n}_{\mathbf{H}+} + \mathbf{n}_{\mathbf{N}+} \tag{17}$$

$$p = n_e k_b T_e + (n_{N_2} + n_{H_2} + n_N + n_H + n_e) k_b T_g$$
(18)

For chemical nonequilibrium (CNE) simulations, Eqs. (15) and (16) are replaced by individual species continuity equations, which are written in terms of the species mass fractions $y_i = m_i n_i / \rho$:

$$\nabla \cdot (\rho \mathbf{v} \mathbf{y}_i) = \nabla \cdot (\rho \mathbf{D}_i \nabla \mathbf{y}_i) + \dot{\rho}_i \tag{19}$$

Mixture-averaged mass diffusion coefficients D_i are employed, rather than multi-component diffusion coefficients. With this method, Eq. (19) is solved for y_{N2} , y_H , y_N , y_{N+} , and y_{H+} . The electron population is then calculated from Eq. (17), and y_{H2} is determined from overall mass continuity, or equivalently $\Sigma y_i = 1$. The species productions rates, $\rho_i = m_i n_i$, are determined as a function of T_g and T_e from the finite-rate chemistry processes summarized in

Table 2. The references for the individual reaction rates are summarized in the right column of Table 2. For two-way reactions, the reverse reaction rates are extracted from the forward reaction rate and the 'equilibrium constants' given by Eqs. (15a-d).

4. Electromagnetic Equations

The calculation of the electrical current distribution is critical to understanding arc physics and poses a challenging problem in arcjet analysis. The current distribution is largely determined by the flow-field electrical conductivity σ , which is a strong function of the electron number density and temperature distribution. Models which assume local thermodynamic equilibrium (LTE) must artificially elevate σ in the cool boundary layer region to avoid excessive voltage drops near the anode.^{11,15} The artificial restriction that $T_e = T_g$ results in an underprediction in the ionization (and therefore electron number density and σ) in the boundary layer regions and a subsequent overprediction in power deposition. Hence, the 'conductivity floor' is employed elsewhere^{10,14} to provide agreement between the experimentally-observed and predicted power depositions.

Diffusion of electrons from the arc core towards the anode wall also elevates the electron number density and plays an important role in determining the boundary layer electrical conductivity. This has been demonstrated by Butler, Kull, and King,¹¹ who show that the incorporation of mass diffusion provides a reasonable current distribution without imposing artificial restrictions on σ or the current attachment region. However, the two-temperature solutions of Miller and Martinez-Sanchez¹⁷ and Keefer et al.¹⁶ demonstrate that significant kinetic nonequilibrium, with elevated T_e, exists outside of the arc core in the constrictor region. These results indicate that both diffusion and increased ionization due to elevated T_e are important factors which determine the boundary layer electrical conductivity.

The correct approach to the arc attachment problem is to model the anode as an equipotential surface and allow the current distribution to be a model output which is independent of artificial

restrictions. This is a self-consistent approach to solving for the current distribution and electric field simultaneously, for which the current distribution is properly coupled to the flow-field.

Assuming induced magnetic fields are negligible, and neglecting electron pressure gradient driving terms, Ohm's law and current conservation are used to solve for **j** and V:²⁵

$$-\nabla \mathbf{V} = \mathbf{j}/\boldsymbol{\sigma} \tag{20}$$

$$\nabla \cdot \mathbf{j} = 0 \tag{21}$$

5. Transport Coefficients

Transport coefficients for this model are calculated using mean free path mixture rules.³⁵ The viscosity is computed as:

$$\mu = \sum_{i=1}^{N} \frac{n_i}{\sum_{j=1}^{N} n_j M_{ij}} \mu_i$$
(22)

where M_{ij} is defined as

$$M_{ij} = \sqrt{\frac{2m_{ij}}{m_i}} \frac{\overline{\Omega}_{ij}^{(2,2)}}{\overline{\Omega}_{ii}^{(2,2)}}$$
(23)

and the pure species viscosity is

$$\mu_i = 1/2m_i n_i c_i l_i \tag{24}$$

where c_i is the mean thermal speed, l_i is the mean free path, and Ω_{ij} is the energy-averaged collision integral for interaction between species i and j.

The translational contribution to the heavy species thermal conductivity is calculated using an equation similar to (24) with the pure species conductivity given as:

$$\lambda_{i} = \frac{15}{8} k_{b} n_{i} c_{i} l_{i} \left(\frac{9\gamma_{i} - 5}{15\gamma_{i} - 15} \right)$$
(25)

The electron thermal conductivity is:

$$\lambda_{e} = \frac{\frac{15}{8} k_{b} n_{e}^{2} c_{e} l_{e} \left(\frac{9\gamma_{e} - 5}{15\gamma_{e} - 15} \right)}{\sum_{j=1}^{N} n_{j} M_{ij}}$$
(26)

and electrical conductivity is calculated from:

$$\sigma = \frac{n_e e^2}{m_e \sum_{i \neq e} n_i c_e \overline{\Omega}_{e-i}^{(1,1)}}$$
(27)

The effective diffusion coefficients D_i are calculated from:

$$D_{i} = \frac{1 - x_{i}}{\sum_{i \neq j} x_{j}/D_{ij}}$$
(28)

where the \mathbf{D}_{ij} are the binary mixture diffusion coefficients given as: 36

$$D_{ij} = \frac{3}{16n\Omega_{ij}^{(1,1)}} \left(\frac{2k_b T(m_i + m_j)}{\pi m_i m_j}\right)^{1/2}$$
(29)

The electron and ion diffusion coefficients are modified to account for the ambipolar electric field. Assuming $\frac{\nabla(n_e/\rho)}{n_e} \approx \frac{\nabla(n_{H+}/\rho)}{n_{H+}} \approx \frac{\nabla(n_{N+}/\rho)}{n_{N+}}$, the ambipolar diffusion coefficients for electrons, H⁺, and N⁺ are:

$$D_{a,e} = \frac{D_e(\mu_{H+}n_{H+} + \mu_{N+}n_{N+}) + \mu_e(D_{H+}n_{H+} + D_{N+}n_{N+})}{\mu_e n_e + \mu_{H+}n_{H+} + \mu_{N+}n_{N+}}$$
(30)

$$D_{a,H+} = \frac{D_{H+}(\mu_e n_e + \mu_{N+} n_{N+}) + \mu_{H+}(D_e n_e - D_{N+} n_{N+})}{\mu_e n_e + \mu_{H+} n_{H+} + \mu_{N+} n_{N+}}$$
(31)

$$D_{a,N+} = \frac{D_{N+}(\mu_e n_e + \mu_{H+} n_{H+}) + \mu_{N+}(D_e n_e - D_{H+} n_{H+})}{\mu_e n_e + \mu_{H+} n_{H+} + \mu_{N+} n_{N+}}$$
(32)

where the mobilities μ are calculated from $\mu_i = \frac{eD_i}{k_bT_i}$.

The above relations require the energy-averaged collision integrals for 28 species interactions. Collision integrals and collision cross sections are obtained from References 37-43. Temperature-dependent curve fits are used for the transport property and elastic transfer calculations.

6. Boundary Conditions

The physical boundary conditions for the continuity, momentum, energy, and electromagnetic equations are summarized in Table 3 and Fig. 1. At the inflow boundary 1A, the flow is axial and the total temperature T_t and total pressure p_t are specified. The total pressure is adjusted to obtain the desired mass flow rate. The u-velocities and the inlet static temperatures are then computed from the isentropic stagnation property relations. The effects of flow swirl are incorporated by specifying the azimuthal inlet w-velocities as a fraction of the u-velocities. At the solid cathode 1C and anode 1D surfaces, no-slip conditions are employed. Zero radial gradients are specified on the center-line 1E.

At the outflow boundary 1B, the static pressure is extrapolated from the interior of the flow. The outflow u-velocities are updated to satisfy continuity using velocity corrections as given by discretized forms of the momentum and continuity equations for the exit plane. This is consistent with the numerical solution method discussed in section III below. Note that a zero electron temperature gradient is employed at the walls 1C and 1D. At low temperatures, the electron thermal conductivity approaches zero so that this boundary condition will have little effect on the results.⁴⁴

The boundary conditions for the voltage are shown in Table 3. The anode is modeled as an equipotential surface, while the cathode voltage is equipotential with the voltage updated to obtain the specified total current I. Note that in this formulation the electrode sheath voltage drops V_s have been uncoupled from the bulk plasma. The axial gradient of the voltage is set to zero at the inflow and outflow boundaries so that current is forced to attach within the arcjet nozzle. Recent radial electric field and σ measurements along the exit plane, showing the existence of roughly 1 A/cm² current density, suggest that this boundary condition may need refinement.⁴⁵

The boundary conditions for the anode energy equation are also shown in Table 3. On the outer surfaces 2B and 2D, the local heat flux is given by radiation to the surroundings at a specified background temperature $T_{bg} \ll T_a$ and a constant emissivity of $\varepsilon = 0.31$. Along the anode surface 2C, the net heat flux is calculated as the sum of (1) the heat flux due to conduction and species diffusion from the flow-field, (2) the radiation loss through the exit plane to the surroundings, and (3) the sheath losses given by the product of the local current density and an assumed sheath voltage drop $V_s = 13$ V. Note that radiation exchange between the plasma and the anode is neglected in this formulation. The upstream anode surface 2A is either a fixed temperature or zero heat flux boundary.

The chemical nonequilibrium model requires boundary conditions for the species populations. At the anode surface, the diffusion velocities are equated to a fraction α_i of the species thermal velocities:²⁹

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$$D_{i}\frac{\partial y_{i}}{\partial \overline{n}} = \frac{\alpha_{i}}{4}\sqrt{\frac{8k_{b}T_{g}}{\pi m_{i}}}$$
(33)

The anode is assumed to be catalytic for recombination, so that $\alpha_{H+} = \alpha_{N+} = \alpha_H = \alpha_N = 1$. Assuming that molecules are reflected from the surface gives $\alpha_{H2} = \alpha_{N2} = 0$. The inflow is in chemical equilibrium up to the constrictor entrance, where the chemical nonequilibrium simulation begins. The mass fractions at the exit plane are extrapolated from the interior of the flow.

III. Numerical Solution Method

The solution method is structured around the global continuity and momentum equations. These are solved using a compressible form of the pressure-based PISO algorithm,⁴⁶ where the density variations are implicitly included in the pressure-correction procedure as described by Rhie.⁴⁷ Staggered velocity and scalar grids yield a well-connected pressure field. Upstream weighted densities are employed to insure numerical stability.⁴⁸ The equations are discretized over finite cell volumes, and power law differencing of convective and diffusive portions of the linearized coefficients preserves numerical stability in regions of the flow where convection dominates. The solution process is iterative, and values for the field variables from the latest iteration are used to linearize the source terms. Under-relaxation is employed so that updated solutions are fractionally added to the solution from the previous iteration. The governing equations are transformed into natural coordinates and solved on a uniform computational mesh.

Figure 2 illustrates the solution algorithm. The energy equations are solved to update gas, electron, and anode temperature fields. The species number densities, transport coefficients, and source terms are then updated using the latest available temperatures and pressures. The continuity and momentum equations are solved to update the pressure and velocity fields. Finally, the voltage and current density distributions are updated. Iterations are

continued until (1) the normalized energy equation residuals are within 1% of the electrical power input, (2) the difference between the inlet and exit plane mass flow rates is less than 1% of the total mass flow rate, and (3) the axial and radial momentum equation residuals are within 1% of their respective exit momentum fluxes.

IV. Results and Discussion

Overview of Previous Simulations

Before proceeding to the present results, a discussion of the effect of the electron-molecule inelastic loss factor δ on our initial voltage predictions is appropriate. For the NASA-Lewis 1 kW arcjet geometry, we found that the δ -factors quoted in Ref. 25 ($\delta \approx O(10-100)$) resulted in current attachment in the constrictor, with corresponding underpredictions of the operating voltage of V ≈ 40 V at I = 10 A. We also noted that increasing the mass flow rate had little or no effect on the attachment location. We electrically insulated the constrictor and observed little effect on the voltage prediction. Martinez-Sanchez recently re-examined the δ -factors for N₂, and values as high as $\delta \approx 4000$ were noted.⁸ We choose $\delta = 3000$ because (1) reasonable convective effects result, with the arc being swept further downstream as the mass flow rate is increased, as was originally postulated in Ref. 8, and (2) better agreement is achieved between the predicted and experimentally observed operating voltage, while maintaining the equipotential anode voltage boundary condition in the constrictor.

The constrictor has been electrically insulated by others to provide better agreement between measured and predicted voltage.^{16,17} These models simulated the German TT1 thruster, a higher power 10 kW arcjet with an \approx 5 mm long constrictor, and predicted that nearly all of the electrical power was deposited in the constrictor. The predictions for our geometry, where x_{con} = 0.25 mm, indicate that roughly 50% of the power is deposited downstream of the constrictor.

<u>Results</u>

We present three simulations for the geometry of a NASA-Lewis 1 kW arcjet. These are summarized in Table 4, where mass flow rate, voltage, current, thrust, and specific impulse are shown. The simulations are designated as either chemical equilibrium (CE) or chemical nonequilibrium (CNE). For each case, the upstream anode surface 2A (Fig. 1) temperature is constant at $T_a = 1000$ K, and the cathode 1C temperature varies linearly from $T_c = 1000$ K at the

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inlet to $T_c = 3000$ K at the cathode tip. The constrictor diameter is 0.63 mm. The cathode gap spacing, as measured axially from the converging portion of the anode surface, is $x_{cat} = 0.58$ mm for the hydrazine cases and $x_{cat} = 1.8$ mm for the hydrogen case. An anode voltage drop of $V_s = 13.0$ V is assumed, and the anode emissivity is $\varepsilon = 0.31$.¹² The inlet flow swirl velocity is 30% of the axial inflow velocity. The computational grids, shown in Fig. 3, are comprised of 62 axial by 17 radial nodes for the plasma, and 62 axial by 9 radial nodes for the anode domain. Simulation run times are several hours on a Convex C-240 mainframe computer.

1. Hydrazine Simulations

The CE and CNE hydrazine simulations are systematically compared in Figs. 4 - 15. Contours of current density and anode temperature are indicated in Fig. 4 for CE hydrazine. At the cathode tip, the current density is $j \approx 40,000 \text{ A/cm}^2$. A peak anode attachment current density of $j \approx 18 \text{ A/cm}^2$ is noted at a location of $x \approx 11 \text{ mm}$, or roughly 5 mm downstream of the constrictor. The anode temperature increases from the upstream boundary value of $T_a = 1000 \text{ K}$ to $T_a \approx 1300 \text{ K}$ near the exit plane. The CNE hydrazine results are shown in Fig. 5. The peak anode current density of $j \approx 15 \text{ A/cm}^2$ occurs at $x \approx 8 \text{ mm}$, upstream from that for the CE case. The anode temperature is somewhat lower than for CE, with a value of $T_a \approx 1200 \text{ K}$ at the exit plane.

The marked difference in the current attachment predictions is more clearly illustrated in Fig. 6, where the anode current densities are indicated. For CNE, a more diffuse, bimodal distribution is noted, with the attachment closer to the constrictor. Consequently the voltage prediction, indicated in Table 4, is significantly lower at 87 V for CNE versus 113 V for the CE simulation. The anode T_e distributions are indicated in Fig. 7. A high degree of thermal nonequilibrium is noted, with maximum electron temperatures of $T_e \approx 8,000$ K and $T_e \approx 12,000$ K for the CE and CNE simulations respectively.

Heavy species and electron temperature contours in the constrictor region are indicated in Figs. 8-11. In Fig. 8, a maximum $T_g \approx 22,000$ K occurs near the center of the constrictor for

CE hydrazine. The central region of the arc is near thermal equilibrium, as indicated by a maximum $T_e \approx 23,500$ K shown in Fig. 9. The results for CNE hydrazine are similar, with maxima of $T_g \approx 20,000$ K and $T_e \approx 23,000$ K indicated in Figures 10 and 11.

Axial velocity contours are shown in Figs. 12 and 13. A maximum centerline velocity of $u \approx 8$ km/s occurs at x = 8 mm, and then decreases because of heating and viscous losses to $u \approx 6$ km/s at the exit for CE hydrazine. For CNE hydrazine, a maximum velocity of $u \approx 7$ km/s is predicted to occur at x = 9 mm, decreasing to $u \approx 6.5$ km/s at the exit.

Finite-rate chemistry must be included to capture frozen flow effects. In Figs. 14 and 15, exit plane predictions of electron number density n_e are presented. The predicted centerline density for CE hydrazine is $n_e = 1.6 \times 10^{11} \text{ cm}^{-3}$, while for CNE, the density is 3 orders of magnitude higher at $n_e = 1.6 \times 10^{14} \text{ cm}^{-3}$. Also shown are T_e and T_g profiles. More thermal nonequilibrium is indicated for the CE case with $T_e/T_g \approx 2.0$ at the centerline, while the CNE case is near equilibrium at $T_e/T_g \approx 1.1$.

2. Hydrogen Simulation

The results for the CNE hydrogen simulation are indicated in Figs. 16 - 20. Anode temperature and current density contours are shown in Fig. 16. The peak anode current density of $j = 16 \text{ A/cm}^2$ occurs at the attachment location of $x \approx 8 \text{ mm}$. Due to the higher specific power deposition, the thermal loading of the anode is somewhat more severe than for the hydrazine cases, with a predicted temperature near the exit plane of $T_a \approx 1400 \text{ K}$. Profiles of T_e and j at the anode surface are shown in Fig. 17. The results are qualitatively similar to the CNE hydrazine case, with high thermal nonequilibrium of $T_e \approx 12,000 \text{ K}$ predicted in the current attachment region.

Heavy species and electron temperature contours are shown in Figs. 18 and 19. In Fig. 18, a peak temperature of $T_g \approx 13,500$ K is noted. The electron temperature is similar in the central region of the flow with $T_e \approx 14,000$ K indicated in Fig. 19. At a location of x = 8 mm, T_e decreases in the radial direction reaching a minimum of $T_e \approx 3700$ K at r = 0.5 mm. The electron

temperature then increases, eventually reaching a value of $T_e \approx 12,000$ K at the anode surface. The electron temperature profile is qualitatively similar to the nonequilibrium predictions of Miller and Martinez-Sanchez¹⁷ and Keefer et al.¹⁶

Axial velocity contours are shown in Fig. 20. A maximum centerline value of $u \approx 15$ km/s occurs at x = 7 mm decreasing to $u \approx 12$ km/s at the exit plane.

V. Conclusions

A detailed, propellant-flexible, N_2/H_2 arcjet model is developed, and is sufficiently comprehensive to be used for arcjet design studies. A thermal nonequilibrium, chemical equilibrium model was first developed, and then generalized to chemical nonequilibrium by including finite rate chemistry and mass diffusion. The incorporation of nonequilibrium processes permits the current distribution to be coupled to the plasma flowfield with realistic boundary conditions and without artificial modifications to the electrical conductivity.

Chemical equilibrium and nonequilibrium results have been compared for hydrazine. High near-anode thermal nonequilibrium is indicated in both cases with $T_e/T_g \approx 8$ and $T_e/T_g \approx 12$ for CE and CNE predictions respectively. In the constrictor, the central region of the arc is near thermal equilibrium with $T_e \approx T_g \approx 20,000$ K. Mass diffusion and finite rate chemistry dramatically affect the arc structure. The incorporation of these chemical nonequilibrium processes reduces the arc length and lowers the predicted operating voltage. Additionally, higher frozen flow losses are noted. Predicted electron densities at the arcjet exit plane are three orders of magnitude higher with nonequilibrium chemistry.

Chemical nonequilibrium results have been presented for hydrogen. The results are qualitatively similar to those for hydrazine. However, lower arc temperatures are noted, with centerline $T_e \approx T_g \approx 14,000$ K in the constrictor.

The combined effects of viscous losses and supersonic heating were noted in all cases. Centerline axial velocities reach a maximum a few millimeters downstream of the constrictor, and then decrease to exit plane values of $u \approx 6$ km/s and $u \approx 12$ km/s for hydrazine and hydrogen respectively.

An anode temperature model was included. For CE hydrazine, maximum anode temperatures of $T_a \approx 1300$ K are predicted near the exit plane. For CNE hydrazine and hydrogen, maxima of $T_a \approx 1200$ and $T_a \approx 1400$ are predicted.

The model has generated much useful data; however, unresolved issues remain. The voltage predictions largely depend on the arc attachment location. We have found that parametrically increasing electron-molecule inelastic energy transfer increases the arc length. A more general physical model of electron-molecule inelastic transfer is required to describe accurately the near-anode transport properties. Additionally, a detailed electrical sheath model is needed to predict near-electrode voltage drops.

We postulate that the electron energy distribution may be non-maxwellian due to the high level of inelastic electron-molecule energy loss. We are considering a more complete description of the coupling between the electron energy and molecular internal energy modes. This may be required to describe the arc attachment and near-anode electrical conductivity more accurately.

The model must be compared with detailed experimental measurements of plasma properties for validation of both local and global performance predictions. Predictions from this model are compared with our experiments in Refs. 49 and 50.

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⁴⁹Bufton, S. A., Burton, R. L., and Krier, H., "Measured Plasma Properties at the Exit Plane of a 1 kW Arcjet," AIAA Paper 95-3066, 31st Joint Propulsion Conference, San Diego, July, 1995.

⁵⁰Tiliakos, N. T., Burton, R. L., and Krier, H., "Application of Internal Langmuir Diagnostics for Arcjets," AIAA Paper 95-2386, 31st Joint Propulsion Conference, San Diego, July, 1995.

| species | C _{pi} (J/K) | hi ⁰ (J/particle) |
|----------------|--|--|
| e | 5/2 kb | 0 |
| H ₂ | $k_{b}\left(7/2 + \frac{\theta_{vH2}}{T_{g}}\left(e^{\theta_{vH2}/T_{g}} - 1\right)^{-1}\right)$ | k _b /2 θ _{vH2} |
| N ₂ | $k_{b}\left(7/2 + \frac{\theta_{vN2}}{T_{g}}\left(e^{\theta_{vN2}/T_{g-1}}\right)^{-1}\right)$ | k _b /2 θ _{vN2} |
| H | 5/2 kb | 1/2 ɛd,H2 |
| N | 5/2 kb | 1/2 ε _{d,N2} |
| H+ | 5/2 kb | $1/2 \varepsilon_{d,H2} + \varepsilon_{i,H}$ |
| N ⁺ | 5/2 kb | $1/2 \varepsilon_{d,N2} + \varepsilon_{i,N}$ |

VIII. Figures and Tables

 Table 1: Species reference enthalpies and specific heats.

| Reaction* | Rate $(m^{6}/s \text{ or } m^{3}/s)$ | Loss rate (W/m ³) | Ref. |
|-----------------------------|---|--|------|
| $2H + M = H_2 + M$ | $k_{f1} = \frac{1.764 \times 10^{-42}}{T_g}$ | $L_{h} = \left(k_{r1}n_{H2} - k_{f1}n_{H}^{2}\right)n_{M}\varepsilon_{d,H2}$ | 26 |
| $H^+ + e + M = H + M$ | $k_{f2} = \frac{1.45 \times 10^{-33}}{T_g^{2.5}}$ | $L_{h} = (k_{r2}n_{H} - k_{f2}n_{H+}n_{e}) n_{M}\varepsilon_{i,H}$ | 26 |
| e + H ₂ > 2H + e | $k_{f3} = \langle \sigma v \rangle$ | $L_e = k_{f3}n_e n_{H2} \varepsilon_{d,H2}$ | 27 |
| $H^+ + e> H + hv$ | $k_{f4} = \frac{6.26 \times 10^{-17}}{T_e^{0.58}}$ | $\mathbf{L}_{\mathbf{e}} = \mathbf{k}_{f4} \mathbf{n}_{\mathbf{e}} \mathbf{n}_{\mathbf{H}+} \mathbf{\varepsilon}_{i,\mathbf{H}}$ | 26 |
| $H^+ + 2e = H + e$ | $k_{f5} = \frac{1.95 \times 10^{-20}}{T_e^{4.5}}$ | $L_{e} = (k_{16}n_{H} - k_{f6}n_{e}n_{H+})n_{e}\varepsilon_{i,H}$ | 26 |
| N ₂ + M> 2N + M | $k_{f7} = \frac{6.14 \times 10^{-9}}{T_g^{1.6}} e^{-113,200/T_g}$ | $L_{h} = k_{f7} n_{N2} n_{M} \epsilon_{d,N2}$ | 28 |
| M + 2N> N ₂ + M | $k_{f8} = \frac{8.10 \times 10^{-37}}{T_g^{1.39}} e^{1745/T_g}$ | $L_{\rm h} = - k_{\rm f8} n_{\rm N}^2 n_{\rm M} \varepsilon_{\rm d, N2}$ | 28 |
| e + N ₂ > 2N + e | $k_{f9} = \frac{4.98 \times 10^{-6}}{T_e^{1.6}} e^{-113,200/T_e}$ | $L_e = k_{f9}n_en_{N2}\varepsilon_{d,N2}$ | 29 |
| $N + e = N^+ + 2e$ | $k_{f10} = \frac{4.15 \times 10^3}{T_e^{3.82}} e^{-168,200/T_e}$ | $L_e = (k_{f10}n_N - k_{r10}n_en_{N+})n_e\varepsilon_{i,N}$ | 29 |

Table 2: Finite rate chemistry model. * ---> one way reactions, = two way reactions with the reverse rate evaluated using the chemical equilibrium constants, M is any heavy species third body.




| Flow-Field Domain | | | | | | | | |
|-------------------|---|--|--|--|---|--|--|--|
| variable | 1A | 1B | <i>1C</i> | | 1D | IE | | |
| u | from stagnation properties | from momentum | | = 0 | = 0 | $\frac{\partial \mathbf{u}}{\partial \mathbf{r}} = 0$ | | |
| v | = 0 | extrapolated | | = 0 | = 0 | = 0 | | |
| w | specified | extrapolated | | = 0 | = 0 | = 0 | | |
| Te | $\frac{\partial T_e}{\partial x} = 0$ | extrapolated | $\frac{\partial T_e}{\partial r} = 0$ | | $\frac{\partial T_e}{\partial r} = 0$ | $\frac{\partial T_e}{\partial r} = 0$ | | |
| Тg | from M and T _t | extrapolated | specified | | $T_g = T_a$ | $\frac{\partial T_g}{\partial r} = 0$ | | |
| Р | extrapolated | extrapolated | $\frac{\partial P}{\partial \overline{n}} = 0$ | | $\frac{\partial \mathbf{P}}{\partial \overline{\mathbf{n}}} = 0 \qquad \qquad \frac{\partial \mathbf{P}}{\partial \overline{\mathbf{n}}} = 0$ | | | |
| v | $\frac{\partial V}{\partial x} = 0$ | $\frac{\partial V}{\partial x} = 0$ | V = V _{cat} | | $V = V_S$ | $\frac{\partial V}{\partial r} = 0$ | | |
| Anode Domain | | | | | | | | |
| variable | 2A | 2B | 20 | | С | 2D | | |
| Ta | T_a specified or $\frac{\partial T_a}{\partial x} = 0$ | $\dot{q} = \varepsilon \sigma \left(T_s^4 - T_{bg}^4 \right)$ | | $\dot{q} = \lambda_{h} \frac{\partial T_{g}}{\partial \overline{n}} + \mathbf{j} V_{s} + \sum_{i=1}^{N} \frac{1}{m_{i}} \rho \left(C_{pi} T_{i} + h_{i}^{0} \right) D_{i} \frac{\partial y_{i}}{\partial \overline{n}} \\ - \varepsilon \sin 20^{\circ} \sigma \left(T_{s}^{4} - T_{bg}^{4} \right)$ | | $\dot{q} = \varepsilon \sigma \left(T_s^4 - T_{bg}^4 \right)$ | | |

Table 3: Boundary conditions for the fluid dynamic, energy, and electromagnetic equations. Energy equation boundary conditions are shown for both the fluid and anode domains. For radiative boundary conditions, T_s is the local anode surface temperature.

| CE/CNE | Propellant | m (mg/s) | I (Amps) | Voltage | I _{sp} (s) | Thrust (N) |
|--------|-------------------------------|----------|----------|---------|---------------------|---------------|
| CE | N ₂ H ₄ | 52.9 | 10 | 113 | 441 | 0.229 |
| CNE | N ₂ H ₄ | 50.2 | 10 | 87 | 439 | 0.216 |
| CNE | H ₂ | 13.2 | 10 | 138 | 934 | 0.121 |

 Table 4: Summary of predictions.
 CE denotes chemical equilibrium and CNE denotes chemical non-equilibrium simulations.



Figure 2: Numerical solution algorithm. The solution method is iterative, employing successive under-relaxation of the indicated field variables.



Figure 3: Computational grids. The grid is tuned axially for high density in the constrictor and radially for high density near the electrode surfaces.



Figure 4: Anode temperature and current density for hydrazine; chemical equilibrium (CE) simulation. The maximum anode current density is $j \approx 18$ A/cm² at $x \approx 11$ mm.



Figure 5: Anode temperature and current density for hydrazine; chemical nonequilibrium (CNE) simulation. The maximum anode current density is $j \approx 15 \text{ A/cm}^2$ at $x \approx 8 \text{ mm}$.



Figure 6: Current density at the anode surface for hydrazine. Results are shown for chemical equilibrium (CE) and chemical nonequilibrium (CNE) simulations.



Figure 7: Electron temperature at the anode surface for hydrazine. Results are shown for chemical equilibrium (CE) and chemical nonequilibrium (CNE) simulations.



Figure 8: Heavy species temperature for hydrazine; chemical equilibrium (CE) simulation. The maximum is $T_g \approx 22,000$ K at the constrictor centerline. The exit plane is located at x = 18.5 mm.



Figure 9: Electron temperature for hydrazine; chemical equilibrium (CE) simulation. The maximum is $T_e \approx 23,500$ K at the constrictor centerline. The exit plane is located at x=18.5 mm.



Figure 10: Heavy species temperature for hydrazine; chemical nonequilibrium (CNE) simulation. The maximum is $T_g \approx 20,000$ K at the constrictor centerline. The exit plane is located at x = 18.5 mm.



Figure 11: Electron temperature for hydrazine; chemical nonequilibrium (CNE) simulation. Maximum T_e is $\approx 23,000$ K at the constrictor centerline. The exit plane is located at x=18.5 mm.



Figure 12: Axial velocity contours for hydrazine; chemical equilibrium (CE) simulation. The maximum is $u \approx 8$ km/s at $x \approx 8$ mm. The centerline velocity decreases to $u \approx 6$ km/s at the exit plane.



Figure 13: Axial velocity contours for hydrazine; chemical nonequilibrium (CNE) simulation. The maximum is $u \approx 7$ km/s at $x \approx 9$ mm. The centerline velocity decreases to $u \approx 6.3$ km/s at the exit plane.



Figure 14: Radial distributions of electron density, electron temperature, and heavy species temperature at the exit plane for hydrazine; chemical equilibrium (CE) simulation.



Figure 15: Radial distributions of electron density, electron temperature, and heavy species temperature at the exit plane for hydrazine; chemical nonequilibrium (CNE) simulation.







Figure 17: Electron temperature and current density at the anode surface for hydrogen; chemical nonequilibrium (CNE) simulation.



Figure 18: Heavy species temperature for hydrogen; chemical nonequilibrium (CNE) simulation. The maximum is $T_g \approx 13,500$ K at the centerline near the cathode tip. The exit plane is located at x = 18.5 mm.



Figure 19: Electron temperature for hydrogen; chemical nonequilibrium (CNE) simulation. The maximum is $T_e \approx 14,000$ K at the centerline near the cathode tip. High near-anode thermal nonequilibrium is indicated downstream of the constrictor, with $T_e \approx 12,000$ K at the anode surface. The exit plane is located at x = 18.5 mm.



Figure 20: Axial velocity contours for hydrogen; chemical nonequilibrium (CNE) simulation. The maximum is $u \approx 15$ km/s at $x \approx 8$ mm. The centerline velocity decreases to $u \approx 12$ km/s at the exit plane.

II. Measured Plasma Properties at the Exit Plane of a 1 kW Arcjet

(AIAA Paper No. 95-3066)

Abstract

With recent advances made in low power (1-2 kW) arcjet numerical modeling, validation of numerical models through experimentation has become increasingly important. We describe multiple electrostatic probe techniques for characterizing flow conditions at the exit plane of a 1 kW-class hydrazine thruster. An improved electrostatic quadruple probe technique, which is also modified to account for plasma gradients over the finite probe volume and multiple ion species, provides measurements of n_e, T_e and u_i/c_m at the thruster exit. Quadruple probe results are also presented for measurements at several centerline locations immediately downstream of the thruster exit plane. A spatially-resolved time-of-flight electrostatic probe technique is employed for measurements of radial profiles of the plasma axial velocity u_i. The quadruple probe results (T_e, n_e, u_i/c_m), coupled with independent measurements of plasma velocity, are used to estimate ion temperature T_i=T_g and the extent of thermal non-equilibrium at the thruster exit. Exit plane experimental data for u_i, T_e and T_g are compared with computational arcjet model predictions, showing agreement for u_i and T_g. The model underpredicts T_e and a direct comparison of n_e results is precluded by large axial gradients at the thruster exit.

Nomenclature

| The second geometric barrage area in i | Α | Electrode | geometric | surface | area | $[m^2]$ |
|--|---|-----------|-----------|---------|------|---------|
|--|---|-----------|-----------|---------|------|---------|

- c_m Most probable thermal speed [m/s]
- e Electron charge [coulombs]
- f Ion composition parameter $(=n_{\rm H}+/n_{\rm e})$

I Probe electrode current [A]

- Iarc Arcjet operating current [A]
- j_i Ion saturation current density [A/m²]

- k Boltzmann constant [J/K]
- m Particle mass [kg]
- m Propellant flow rate [mg/s]

n Particle density [cm⁻³]

- P Thruster power [kW]
- Pk Designation for electrode "k"
- Q Collision cross section [m²]
- rp Electrode radius [mm]
- s Electrode clearance [mm]
- T Temperature [K, eV]
- ui Ion flow velocity [m/s]
- v Particle thermal speed [m/s]
- V Probe electrode voltage [Volts]
- V_d Electrode relative bias voltage [Volts]
- V_f Floating electrode voltage [Volts]
- V_p Plasma potential [Volts]
- Varc Arcjet operating voltage [Volts]
- x Axial distance from thruster exit [mm]
- x_{H⁺,N⁺} Wake effect collection area parameters
- δ Flow divergence angle [Degrees]
- ε_d Dissociation energy [eV]
- **Γ** Mathematical Gamma function
- θ Probe angle [Degrees]
- κ Correction to n_e from N⁺ ions
- μ Equal to $(m_{H+}/m_{N+})^{1/2}$
- τ_L Electrode "end effect" parameter
- λ_D Debye length [m]

 λ_{mfp} Collision mean free path [m]

σ Standard deviation [mm]

 ϕ Equal to e/kT_e , $[V^{-1}]$

Subscripts

e,i,n Electron, ion, neutral species

g Gas

1,2,3,4 Electrode designation

I. Introduction and Background

Although research in the area of electrothermal thrusters is relatively mature, only recently has the arcjet become applied to commercial satellites.¹ With these thrusters beginning to find a niche in industry, accurate numerical models become increasingly important as design tools to help in the understanding of fundamental arcjet physics. As such, adequate measurements of fundamental plasma properties must be available for the purpose of validating numerical efforts. It is the purpose of this research to: 1) provide fundamental data for validation of H_2/N_2 computational arcjet models; 2) add to the existing base of fundamental data in the very near-field plume region; and 3) demonstrate the feasibility of electrostatic probing techniques as useful diagnostics in the small geometries associated with the 1 kW arcjet. In this paper we present electrostatic time-of-flight and quadruple probes results for the exit plane of a 1 kW hydrazine arcjet. Where applicable, comparisons are made with experimental data and computational model predictions performed by our group.

Electrostatic probes have found much use in experimentally determining plasma parameters in space and laboratory plasmas. Several authors²⁻⁶ have employed classical Langmuir⁷ single probes in the characterization of low power

arcjet plumes. While single probes are appealing due to their relatively simple construction, interpreting the probe V-I characteristic can be difficult. A viable alternative is the triple electrostatic probe,⁸ which allows simultaneous measurement of T_e and n_e without the necessity of a voltage sweep and a corresponding probe V-I characteristic. The utility of triple probes for electric propulsion devices has been demonstrated in MPD thruster plumes⁹⁻¹² and in a 1 kW hydrogen arcjet plume.¹³ Additionally, a general review¹⁴ of electrostatic probe techniques and their uses in electric propulsion devices has recently been presented.

Under certain conditions, the crossed-probe technique¹⁵⁻¹⁸ can be used to measure u_i/c_m (=plasma velocity/ion most probable thermal speed) in flowing plasmas. The quadruple electrostatic probe¹⁹ combines the crossed-probe and triple probe techniques for simultaneous measurements T_e , n_e , and u_i/c_m . The utility of the quadruple probe has been demonstrated in an MPD thruster^{19,20} and, more recently, in the very near-field plume of a 1 kW hydrazine arcjet.²¹

II. Quadruple Probe Technique

Figures 1a. and 1b. show an electrical schematic and potential plot of the quadruple probe, consisting of three cylindrical electrodes (P₁, P₂, P₃) which are aligned with the plasma flow vector and one electrode (P₄) which is perpendicular to the plasma flow. P₃ and P₄ are biased at constant voltages V_{d3} and V_{d4} relative to P₁, and P₂ assumes the floating potential V_f of the local plasma (I₂=0). The circuit comprised of electrodes 1, 3 and 4 is electrically floating such that ion current collected at P₃ and P₄ is balanced by the electron current collected at P₁, thereby determining the potential difference V_{d2} [=V_{d2}(T_e)]. The three aligned electrodes 1-3 are similar to the triple probe and yield simultaneous

measurements of n_e and T_e. The addition of the perpendicular electrode P₄ allows the measurement of the parameter u_i/c_m , determined from the ratio I₄/I₃ for crossed electrostatic electrodes with V_{d3} = V_{d4}. For the quadruple probe, the measured quantities are the electrode currents I₃ and I₄ and the potential difference between electrodes 1 and 2, V_{d2}. Currents I₃ and I₄ are determined by measuring voltage drops across resistors (R₃ = R₄ = 100 Ω). The electrode bias voltages V_{d3} and V_{d4} are provided by batteries, with V_{d3}=V_{d4}=12V.

It has been shown⁹ that probe contamination can have a marked effect on T_e measurements through drastic changes in the measured quantity V_{d2} . In the present study, the probe was cleaned thoroughly before each data acquisition session using ion bombardment cleaning.¹³ This was accomplished by biasing all four electrodes 24 V below facility ground to attract ions to the probe. The electrodes were then placed in the arcjet plasma stream for several ~ 1 s intervals, so that the electrodes would glow and "burn off" the contaminant layer. Quadruple probe sweeps acquired before cleaning always yielded electron temperatures that were higher than the post-cleaning values by a factor of ~ 2.

Probe Construction

A schematic of the quadruple probe used in this study is shown in Fig. 2. The probe is comprised of 4 individual tungsten wire electrodes of 0.25 mm diameter with an exposed length of 2.5 mm for electrodes 1-3 and 2.0 mm for electrode 4. The location of the perpendicular electrode has been modified from that of previous probes,²¹ to coincide with the axial location of the probe center. Each wire is mounted in round single-bore alumina (Al₂O₃) tubing, which is in turn mounted in, and supported by, thin-wall stainless steel tubing. This assembly is mounted in a larger stainless steel tube to facilitate connecting the probe tip to

the probe positioning mechanism discussed below. Inside this larger tubing is a four-bore alumina tube that electrically insulates the four probe leads. A high temperature ceramic-based adhesive²² is used to bond each of the probe electrodes and components in place. A small quantity of adhesive is placed at the junction between P₄ and its single bore alumina tube to electrically insulate part of the electrode, thereby reducing the error associated with determining its geometric surface area (Fig. 2.).

The center-to-center separation between electrodes 1 and 2 is 1.4 mm. This spacing, along with electrode lengths of 2.5 mm, allows an unaligned quadruple probe to handle flow divergence angles $\delta \sim 30^{\circ}$ without "shadowing" of one electrode by the other. In an effort to minimize the effect of the probe geometry on the plasma flow, a gradual transition between the electrodes and the probe support has been implemented. The radius of the single-bore alumina electrode supports is ~ 0.4 mm, which minimizes perturbations of the plasma flow at the junction between the electrodes and their support. The exposed length of the single-bore alumina is ~ 5 mm. Ceramic adhesive is used to fill the void, and ease the transition, between the electrodes and the stainless steel probe support tubing.

Probe Length Scales

For the conditions expected at the exit plane probe location ($T_e \sim 0.6 \text{ eV}$, $n_e \sim 4 \times 10^{12} \text{ cm}^{-3}$), the Debye length is 2.9×10^{-4} cm and the ratio of electrode radius to Debye length $r_p/\lambda_D=45$, so that the ion sheath surrounding the electrodes can be classified as thin with minimal error. For probe measurements 5 mm downstream, this ratio increases to $r_p/\lambda_D \sim 55$. The clearance (s) between electrodes 1 and 3 (Fig. 2) is s~1 mm so that $s/\lambda_D > 250$ and the sheaths on adjacent electrodes do not interact.

To avoid strong sensitivity of the ion current to small misalignments between the flow vector and the electrode axis, the so-called end effect parameter²³ τ_L [=(L/u_i λ_D)(kT_e/m_i)^{1/2})] should be greater than 50. For our plasma conditions, this parameter is conservatively estimated at τ_L >250. The electrodes should also be long enough that the Bohm sheath ion current collected along the cylinder length (I_b) is much larger than the current collected at the probe end due to the convection of charged particles into the probe tip (I_t). A first order analysis of the ratio I_b/I_t for the conditions at the exit plane showed I_b/I_t > 30, so that the error introduced to the collected ion current by neglecting the cylinder tip is less than 4%.

The radius r_p of the tungsten electrodes (Fig. 2) was chosen as a compromise between the thin sheath assumption $(r_p/\lambda_D >>1)$ and the assumption of free molecular flow over the probe electrodes $(\lambda_{mfp}/r_p >>1)$. For the seven species plasma expected at the exit plane (H₂, N₂, H, N, H⁺, N⁺, e), all relevant collision mean free paths were calculated. The following assumptions are made in calculating mean free paths: $T_e=7000$ K, $T_g=2500$ K, $n_e = 4 \times 10^{12}$ cm⁻³, and $n_{neutral}$ = 3 × 10¹⁵ cm⁻³ with 10% dissociation at the thruster exit. The following (worst case) mean free paths were calculated: $\lambda_{n-n}/r_p > 20$, $\lambda_{i-n}/r_p > 5$, $\lambda_{i-i}/r_p > 30$, λ_{i-e}/r_p =20, $\lambda_{e-n}/r_p > 20$, and $\lambda_{e-e}/r_p = 30$. Clearly, the probe electrodes are operating in the collisionless regime. Additionally, $\lambda_{mfp}/\lambda_D > 200$ for all collisions, with an overwhelming majority of the collisions having $\lambda_{mfp}/\lambda_D > 1000$, indicating that the collisionless sheath assumption is appropriate.

Quadruple Probe Theory

Quadruple probe analysis assumes: 1) The ion sheath surrounding the tungsten wires is thin compared to the wire radius $(r_p \gg \lambda_D)$; 2) Each probe wire and sheath are collisionless $(\lambda_{mfp} \gg r_p \gg \lambda_D)$; 3) The electron energy distribution is Maxwellian; 4) Velocity slip between plasma species is negligible; and 5) A wake region of low ion and electron density exists downstream of the perpendicular electrode P4.

Derivation of quadruple probe response is similar to that of the triple probe⁸ and has been presented elsewhere.¹⁹⁻²¹ In our previous work²¹ we presented some of the revisions necessary to allow the application of triple and quadruple probe theory to multicomponent (N-H) plasmas. In the present study, the probe is applied to a simulated hydrazine plume, with multiple species of ions present, whereas most of the previous applications of triple and quadruple probes have been for plasmas with one dominant species of positive ion. Single Langmuir probes have been used in nitrogen and hydrogen-based arcjet plumes,²⁻⁵ but their use is not contingent on knowledge of the species.

In the present study, we present a significant modification of the quadruple probe theory. Rather than utilizing an analytic expression for the current collected at electrode P4, I4 is related to the current collected at P3 through the ratio of the measured quantities I3 and I4. The significance of this modification is that it completely removes errors in T_e and n_e measurements that were due to assumptions regarding effective current collection area of P4. For a quadruple probe with electrode geometric surface areas A3 (=A1=A2) and A4 and electrode biasing such that Vd3=Vd4, the following expres-sion can be derived:

$$1 = \frac{1 + \exp(\phi V_{d3}) - 2\exp[\phi(V_{d3} - V_{d2})]}{(I_4/I_3) \{\exp[\phi(V_{d3} - V_{d2})] - 1\}}$$
(1)

Since V_{d3} is prescribed and V_{d2} is experimentally determined, Eq. (1) determines $T_e = e/k\phi$ through iteration.

The ion saturation current density ji is related to measured quantities by

$$j_{i} = \frac{(I_{3}/A_{3})(1 + I_{4}/I_{3})}{\exp(\phi V_{d2}) - 1}$$
(2)

Eq. (2) is used to determine n_e from I₃, I₄ and V_{d2} by relating j_i to the electron density through the Bohm sheath analysis. For a plasma with more than one species of collected ion, the Bohm analysis must be modified to account for the collection of each ion species via the Bohm sheath. For the region of the hydrazine plume of interest, H⁺ and N⁺ are the dominant ions species, and the Bohm expression becomes:²¹

$$h_i = e (kT_e)^{1/2} exp(-\frac{1}{2}) \left(\frac{n_{H^+}}{(m_{H^+})^{1/2}} + \frac{n_{N^+}}{(m_{N^+})^{1/2}} \right)$$
 (3)

For $n_e=n_H^+ + n_N^+$, Eqs.(2) and (3) yield

$$n_{e} = \frac{\kappa \left(\frac{I_{3}}{A_{3}}\right) \left(1 + \frac{I_{4}}{I_{3}}\right) \exp\left(\frac{1}{2}\right) (m_{H^{+}})^{1/2}}{e(kT_{e})^{1/2} \left[\exp(\phi V_{d2}) - 1\right]}$$
(4)

In Eq. (4), κ represents the effect of multiple ion species on the probe response to electron density:

$$\kappa = \frac{1 + \left(\frac{n_N^+}{n_H^+}\right)}{1 + \mu\left(\frac{n_N^+}{n_H^+}\right)}$$
(5)

where $\mu = (m_H + /m_N +)^{1/2}$. For a plume contain-ing only H⁺, $n_N + = 0.0$ and κ reduces to unity, such that Eq. (4) reduces to single component (H⁺) form.

Experimental observations^{24,25} show that low nitrogen ion densities are found near the exit plane in ammonia and hydrazine arcjets. It is suggested that this is a result of charge exchange collisions in the expanding nitrogen-hydrogen propellant. It can be shown (Fig. 3) that equilibrium number density ratios n_N/n_H and n_N^+/n_H^+ are equal for T < 7500 K at p = 1 atm. For this temperature range, $n_N/n_H < 0.4$, due to the preferential dissociation of H₂ ($\epsilon_d = 4.48 \text{ eV}$) over N₂ ($\epsilon_d = 9.61 \text{ eV}$). The arcjet exit plane N-H plasma is highly dissociated and partially ionized, with $T_e > T_g$, and the estimated range of gas temperature is $T_g < 6000$ K. Ionization by electron collision is negligible, and nitrogen rapidly undergoes charge-exchange collisions with hydrogen, the cross-section for which is on the same order of magnitude as N-H momentum transfer collisions. Thus: H + N⁺ <---> H⁺ + N, and in the absence of other ionization processes, the two charge-exchange reactions balance in steady state. Writing these volumetric collision rates in the form nQv, with relative thermal speeds vH,N \approx vH, and cross sections QH-N+ \approx QN-H+ gives:

$$\frac{\mathbf{n}\mathbf{N}^{+}}{\mathbf{n}\mathbf{H}^{+}} = \frac{\mathbf{n}\mathbf{N}}{\mathbf{n}\mathbf{H}} \tag{6}$$

which directly implies a low nitrogen ion density in the exit plane (Fig. 3). Thus, the nitrogen ion density is effectively determined by the gas temperature T_g and not the electron temperature. For $T_g < 6000$ K, $n_N^+/n_H^+ < 0.1$ and $\kappa < 1.07$, indicating a ~7% increase of n_e due to the presence of nitrogen ion probe current.

Crossed Electrostatic Probes

The use of crossed electrostatic probes for measuring plasma velocity^{15,17} and the implementation of the crossed probe technique into the quadruple probe have been presented elsewhere.^{19,21} For a cylindrical electrode oriented at an angle θ relative to the plasma flow, the collected ion current is a function of the quantity (u_i/c_m)sin θ , where c_m = (2kT_i/m_i)^{1/2}. For the case of two equally biased probes, with one probe aligned with the flow vector and the other one normal to it, the single species collected ion current ratio for the thin sheaths case is:¹⁷

$$\frac{I_4}{I_3} = \frac{2 A_4}{\sqrt{\pi} A_3} \exp\left[-\left(\frac{u_i}{c_m}\right)^2\right] \sum_{n=0}^{\infty} \left[\frac{\left(\frac{u_i}{c_m}\right)^n}{n!}\right]^2 \Gamma(n+\frac{3}{2})$$
(7)

In order to apply the crossed probe technique to the multicomponent (H₂-N₂) plasmas of interest here, significant revisions must be made to Eq. (7). In the original derivation of the crossed probe technique, a Maxwellian velocity distribution is assumed for the single component ion collection.¹⁵ For the present study, this expression has been rederived using a two-component form of the Maxwellian velocity distribution. The derivation is similar to the single component case, and results in a contribution to the ratio I4/I3 from each of the collected ion species. For the two species of ions assumed present, the ion composition is defined by $f = n_H + /n_e$ and $(1-f) = n_N + /n_e$ ($n_e = n_H + n_N +$). The resulting expression for the measured current ratio is

$$\frac{I_{4}}{I_{3}} = C \left\{ f(x_{H^{+}}) \exp\left[-\left(\frac{u_{i}}{c_{m,H^{+}}}\right)^{2} \right] \sum_{n=0}^{\infty} \left[\frac{\left(\frac{u_{i}}{c_{m,H^{+}}}\right)^{n}}{n!} \right]^{2} \Gamma(n + \frac{3}{2}) + \mu(x_{N^{+}})(1 - f) \exp\left[-\left(\frac{u_{i}}{\mu c_{m,H^{+}}}\right)^{2} \right] \sum_{n=0}^{\infty} \left[\frac{\left(\frac{u_{i}}{\mu c_{m,H^{+}}}\right)^{n}}{n!} \right]^{2} \Gamma(n + \frac{3}{2}) \right\}$$
(8)

where the constant C is given by

$$C = \frac{2 A_4}{\sqrt{\pi} A_3 [f + (1-f)\mu]}$$
(9)

In Eq. (8), the first term in the brackets { } is the H⁺ contribution and the second term is the N⁺ contribution. The most probable thermal speed for N⁺ ($c_{m,N+}$) has been written in terms of $c_{m,H+}$ (since $c_{m,N+=} \mu c_{m,H+}$). Note that each of the terms is weighted by the relative density of its corresponding ion [f, (1-f)].

Because a wake is formed behind P4, a portion of the electrode area A4 does not collect ions. The parameters x_{H+} and x_{N+} are the fractions of the electrode area A4 available for H⁺ and N⁺ collection, respectively. Because the extent of the wake effect is determined by the relative magnitudes of the directed ion velocity u_i and thermal speed of the ions, x_{H+} and x_{N+} are not necessarily equal. For Tg=T_i=2500 K and a directed velocity u_i ~ 6500 m/s, u_i/c_{m,H+} ~ 1 and u_i/c_{m,N+} ~ 4. Several references discuss the wake effect for high speed flows.¹⁷⁻¹⁹ It has been postulated and experimentally verified for an argon plasma at u_i/c_m ~ 1, that the wake effect causes the effective collection area to be half of the geometric electrode area.¹⁷ Since u_i/c_{m,H+} is expected to be ~ 1, x_{H+} is assumed to be 1/2 so that only the front half of P4 collects ions. For the nitrogen ions, the wake effect reduction in collection area is expected to be more pronounced. Since the directed velocity u_i is several times c_{m,N+}, N⁺ is assumed to be collected by the projected area¹⁹ of the perpendicular electrode only, such that x_N +=1/ π .

Given f, Eq. (8) becomes an expression for $u_i/c_{m,H^+}$ versus the measured current ratio I₄/I₃. Equation (8) is plotted in Fig. 4 for $0 \le f \le 1$, $x_{H^+=1/2}$ and $x_{N^+=1/\pi}$. Pure H⁺ collection is represented by f=1.0, 2H⁺ + N⁺ collection by f=0.67, and pure N⁺ collection by f=0.0.

If an independent measure of the directed plasma velocity u_i is known, Fig. 4 can be used to determine $c_{m,H+} [=(2kT_i/m_{H+})^{1/2}]$ and, hence, the gas temperature $T_g=T_i$. We describe below u_i measurements which lead to an estimate of T_i .

Effect of Radial Gradients on Probe Response

Quadruple probe measurements at the thruster exit plane show smooth symmetric profiles for n_e and the electrode currents I₃ and I₄, as shown below. However, V_{d2} is known to have a very asymmetric profile,²¹ which is interpreted as a result of the steep radial gradient in n_e off-axis, and to a lesser extent, the radial gradient in T_e. These gradients are perpendicular to the quadruple probe electrodes and the arcjet thrust axis. In the presence of plasma radial gradients, the individual electrode current equations²⁰ are rewritten in terms of the plasma parameters at each electrode, and are rearranged to solve explicitly for V_{d2} and I₃ in terms of radial profiles of T_e and n_e :

$$V_2 = V_2 \left(T_{e2}, \kappa \right) \tag{10}$$

$$V_1 = V_1 (T_{e1}, T_{e3}, n_{e1}, n_{e3}, I_4/I_3, \kappa)$$
(11)

$$V_{d2} = V_2 - V_1$$
 (12)

$$I_3=I_3 (T_{e3}, n_{e3}, \kappa)$$
 (13)

The subscripts on the parameters T_e and n_e in Eqs. (10)-(13) denote the local value of those parameters at each of the electrodes (1-3) of the quadruple probe. The ratio I₄/I₃ is a measured quantity and κ is a function of the plasma composition discussed above. Since the centerline experimental data are at zero gradient conditions, these data can be used with Eqs. (1) and (4) to determine the centerline values of T_e and n_e.

Using Eqs. (10)-(13), Fig. 5 displays V_{d2} vs. radial position for the quadruple probe of Fig. 2, for flat and Gaussian n_e and T_e profiles. For Gaussian n_e and constant T_e , V_{d2} increases monotonically across the thruster. For a Gaussian T_e distribution broader than n_e , the V_{d2} profile shows more asymmetry. The constant T_e case can be solved analytically for V_{d2} , giving:

$$V_{d2} = \frac{kT_e}{e} \ln\left[\left(1 + \frac{I_4}{I_3}\right) \times \left(\frac{n_{e3}}{n_{e1}}\right) + 1\right]$$
(14)

In this equation the effect of the density gradient can be clearly seen. As the swept probe starts outside the plume and enters it, $n_{e3}/n_{e1} < 1$. At the axis $n_{e1} = n_{e3}$, and as the probe leaves the plume, $n_{e3}/n_{e1} > 1$. The result [Eq. (14)] is a monotonically increasing V_{d2} across the thruster face.

Given arbitrary profiles of $T_e(r)$ and $n_e(r)$, we see that $V_{d2}(r)$ and $I_3(r)$ can be predicted. The inverse problem, of extracting n_e and T_e profiles from measured V_{d2} and I_3 , can also be solved. Since V_{d2} [Eq. (12)] is a strong function of T_e , and I_3 [Eq. (13)] is a strong function of n_e , there is sufficient uncoupling that an iterative routine is used to find the unique $T_e(r)$ and $n_e(r)$ profiles which reproduce the measured V_{d2} and I_3 profiles. Starting with centerline values and assumed Gaussian profiles, the routine marches the probe position across the thruster face, while generating a new T_e profile. This updated profile is used with Eq. (13) to update $n_e(r)$, and the process is repeated until both profiles converge. Typically five numerical iterations are required to reach convergence.

III. Time-of-Flight Velocimetry

Since in addition to the quantities n_e and T_e a quadruple probe measurement yields the quantity $u_i/(2kT_i/m_H+)^{1/2}$, knowledge of the heavy particle velocity u_i can be used to determine the heavy particle temperature $T_g = T_i$. We have designed and implemented a time-of-flight velocimetry technique for measuring spatially resolved axial velocities in arcjet plumes. This technique is derived from a previous Current Modulation Velocimetry (CMV) method.^{13,26} In our method, a short duration current deficit is superimposed on the arcjet current, the effect of which is monitored by an electrostatic time-of-flight (T-O-F) probe as it convects axially in the thruster plume.

Time-of-Flight Probe

The CMV technique can be made spatially resolved with electrostatic probes, thereby eliminating the need to Abel-invert the data as required by line-of-sight optical methods. Figure 6 shows the electrostatic time-of-flight sensing probe used in this study. It consists of two 0.75 mm diameter tungsten wires which are electrically insulated along most of their length by two 1.6 mm o.d. alumina tubes. The exposed length of each tungsten electrode is 0.25 mm and the axial separation between the two is 5 mm (\pm 2%). The elevation of the probe is such that the arcjet thrust axis is located between the sensing electrodes. The electrodes are biased 24 V below facility ground (arcjet anode) to monitor the local electron saturation current density [~ n_e, (T_e)^{1/2}] at each measurement location. Variations in probe electrodes, with the temporal separation of these signals, combined with the probe geometry, indicating the flow velocity.

Although this probe is physically intrusive, the probe electrodes (0.75 mm dia.) are in the near-free molecular flow regime and are thus expected to have little effect on the flow. Additionally, any plasma perturbations due to the electrodes (shocks, collisional effects, etc.) are expected to have a similar result at each electrode, thereby having minimal effect on the convective time delay.

Arcjet Current Deficit Circuitry

The arcjet power processing unit (PPU) produces current with a high frequency ripple (~ 15-20 kHz, \pm 8% about the mean) which causes "natural" fluctuations in n_e and T_e. These inherent fluctuations are difficult to utilize with the T-O-F probe due to their long period (~ 55 µs), compared with the expected

convective time delay of the T-O-F probe (~ 1 μ s). Instead, fluctuations are artificially introduced to the arcjet plasma flow by superimposing a short duration (~ a few μ s) current deficit pulse on the arcjet operating current.

The circuit used to modify the arc current is similar to that used in previous CMV studies²⁶ and is shown in Fig. 7. The current pulse is initiated by closing a switch to a fast R-C circuit in parallel with the arcjet, causing a sudden decrease in arc current while the capacitor charges. This short duration (~ 4 μ s) current deficit results in a "tagging" of the plasma in the arc-heating region. Switch closure is accomplished with an IRF-350 field-effect transistor (FET), which is closed by the gate pulse shown in Fig. 7. Originally, the R-C circuit element was replaced by a single resistor (20-30 Ω), but ringing on the edges of the resulting square current pulse obscured the desired T-O-F probe signals.

Spatially resolved axial velocities are measured by triggering the arcjet current deficit pulse at several different T-O-F probe locations. Since the probe is swept radially through the plume during the experiment, it is necessary to trigger the current deficit pulse repeatably at each desired radial location. Probe location is measured by monitoring the voltage of a precision potentiometer (Fig. 7) which has been calibrated against the arcjet anode for probe position. The position (voltage) of the desired measurement location is set at the voltage comparator circuit, so that a trigger signal is issued to the HP 214A pulse generator when the probe reaches the desired radial location. In turn, the pulse generator supplies a square pulse to "close" the FET and initiate the arcjet current deficit. Because 1) the gate pulse must be positive (relative to the FET voltage), and 2) the arcjet cathode (and FET) operates at ~ -100 V (anode grounded), the FET Driver Circuit (Fig. 7) is a necessary buffer between the FET and the HP pulse generator. Since the

probe is swept relatively slowly (~ 20 cm/s), the probe location is effectively "frozen" during the ~ 3-4 μ s elapsed time between the initial trigger and the monitoring of the tagged plasma at the T-O-F electrodes.

IV. Experimental Apparatus

Electrostatic probe experiments were carried out at the exit plane of a 1 kW NASA arcjet thruster²⁷ operating at a flow rate of 50 mg/s of $2H_2 + N_2$ to simulate fully decomposed hydrazine. Each component of the propellant was individually metered by Unit mass flow controllers and mixed in the propellant line. During steady state operation, the thruster operated at 10.0 A and 112 V (P/m = 22.4 MJ/kg). Arc current was measured with a Hall effect current transducer. The thruster exhausted into a 1 m diameter x 1.5 m long vacuum tank²¹ at ~200 mTorr during arcjet operation. Anode temperatures were monitored with an optical pyrometer, reaching ~ 1200 K during steady state operation.

The quadruple probe and T-O-F probe are accurately positioned and repeatably swept through the thruster plume with the probe mount system shown in Fig. 8. The complete assembly is fastened to a step motor (Fig. 8) mounted on a linear translation carriage.²¹ The motor and probe assembly carriage is capable of \pm 5 cm linear translation perpendicular to the arcjet axis, allowing the probes to be manually swept through the plume at ~ 20 cm/s. Carriage and probe position are determined to 0.25 mm by a precision 10 k Ω linear potentiometer coupled to the carriage manual drive shaft, and calibrated against the outer diameter of the anode. The probe support arm is coupled directly to the motor shaft with the probe tip on the shaft axis, allowing probe angle to be varied (in 0.9° increments) without changing the probe tip location.

Probe elevation is repeatably located to within 0.5 mm by aligning the probe center with a line etched on the arcjet anode. The separation between the probe tips and the thruster exit plane is reliably set to within 0.1 mm with a spark gap gauge. The arcjet mount is known to deflect toward the probe ~0.7 mm while under vacuum. Any error associated with the arcjet exit plane position because of this is systematic and does not affect the relative spacing of subsequent axial measurement positions.

V. Experimental Results and Analysis

Experimental quadruple and T-O-F probe results are presented for the very near-field plume of the arcjet. To quantify the extent of axial gradients in T_e and n_e , quadruple probe measurements are made at several probe tip locations from 1-5 mm from the exit plane. Radial profiles with probe tips located 1 mm from the exit are presented for both the quadruple and T-O-F probes.

Quadruple Probe Results

For the quadruple probe, T_e and n_e are determined by measurements of V_{d2} , I₃ and I₄. Since these parameters are expected to be influenced by the inherent current ripple associated with the arcjet PPU, a discussion of the effect of the ripple on the probe is warranted. Figure 9 shows the centerline ion current density j₃ (= I₃/A₃) measured at electrode 3 and the arc current I_{arc} versus time. The current measured at electrode 3 is in phase with the arc current, except for a delay of ~ 3-4 µs associated with the time required for the plasma to flow from the arc-heating region near the cathode to the quadruple probe. Accounting for differences in average velocity and measurement location, this delay is consistent with previous triple probe results for the hydrogen arcjet.¹³

The relationship between centerline measurements of Vd2 and j3 is shown in Fig. 10. While j3 closely follows the phase of the PPU current ripple, Vd2 is 180° out of phase with j3. Since j3 ~ $n_e(T_e)^{1/2}$ and V_{d2} increases monotonically with T_e [Eq. (1)], Figs. 9 and 10 indicate that, except for the convective time delay, ne increases and T_e decreases with increases in I_{arc} during a typical PPU ripple cycle. Specifically, Eqs. (1) and (4) indicate that the \pm 10% current ripple causes an associated \pm 18% ripple in n_e (in phase) and a \pm 5% ripple in T_e (180° out of phase) about their means. For the quadruple probe centered at x=2.25 mm from the exit plane, the mean centerline values are T_e ~0.6 eV and n_e ~3.6 x $10^{12}~\rm cm^{-3}.$ The T_e results, although out of phase with Iarc, are in phase with the arc voltage Varc (which has a \pm 3% ripple) due to the negative impedance characteristic of the arcjet. Thus the T_e behavior is consistent with the variation of the E-field, which heats the electrons. These results are different from previous hydrogen arcjet data,¹³ which indicate that n_e decreases and T_e increases with increasing I_{arc} . It is unclear at this time if this is an indication of a fundamental difference in the operating physics of the hydrazine and hydrogen arcjets.

Because of the large variation in the measured quantities V_{d2} , I_3 and I_4 with the PPU current, the mean values of these parameters are used to determine the mean values of T_e , n_e and $u_i/c_{m,H+}$ through Eqs. (1), (4) and (8). For radial profile data, a moving average smoothing routine is applied to the raw data so that the effects of the PPU ripple are eliminated.

Te and ne Axial Profiles

Previous n_e data indicate that axial gradients over the length of the quadruple probe may be large.^{4,28,29} To quantify the effects of gradients in T_e and n_e over the 2.5 mm quadruple probe length, centerline data were acquired for

several probe axial locations. Five probe tip locations from 1 to 5 mm from the thruster exit, in 1 mm increments, were utilized. Results for the measured current density j3 for five centerline axial locations are shown in Fig. 11. The geometry of the three aligned electrodes is also shown, with its placement corresponding to the probe location for the first axial location. A 50 % decrease in j3 is indicated over the 5 mm region investigated, which represents the combined effect of changes in T_e and n_e, since j3 ~ n_e(T_e)^{1/2}.

Figure 11, along with the corresponding V_{d2} data and Eqs. (1) and (4), yields the axial centerline profiles of T_e and n_e shown in Fig. 12. As a check of the quadruple probe and to insure that the presence of the perpendicular electrode does not influence the T_e and n_e data, independent measurements were made with a triple probe at the first axial location (x= 2.25 mm). The quadruple and triple probe T_e and n_e data were consistent within 5 %, well within the experimental error associated with the experiment. This is not a surprising result, since in the limit I₄/I₃ --> 0, Eqs. (1) and (4) reduce to the triple probe equations for T_e and n_e .

The electron temperature (Fig. 12) varies from 6700 to 3600 K (~0.6 - 0.3 eV) between 2.25< x <6.25 mm from the thruster exit. These results are roughly consistent with previous far-field plume results of T_e ~0.2-0.3 eV for 30<x<150 mm and T_e ~ 0.1-0.2 eV for x=320 mm.^{3,4} Further, the present data seem to corroborate previous estimates²⁸ of T_e \ge 0.5 eV in the arcjet nozzle based on arguments that T_e is on the order of the atomic excitation temperature.

Results for the electron density axial profile are also shown in Fig. 12. Over the 5 mm range of probe locations n_e drops by ~ 30 %, with the magnitude of dn_e/dx increasing slightly near the thruster exit. The n_e results of the present

study, along with those of other researchers, are summarized in Fig. 13. While the n_e gradient based on the present data is large near the exit $(dn_e/dx \sim -0.5 \times 10^{12} \text{ cm}^{-3}/\text{mm} \text{ at } x= 2-3 \text{ mm})$, it is small compared with the axial gradient indicated in the nozzle interior and at the exit plane. It thus appears likely that large axial variations in the n_e gradient (large d^2n_e/dx^2) are present in the thruster very near-field plume. The present n_e data appear consistent with previous results with the exception of previous data for x=0 and x=10 mm (Fig. 13).²⁹ However, these data represent upper limits placed on n_e based on Stark broadening of line-of-sight (non-Abel inverted) emission spectroscopy signals, and thus are consistent with our probe results.

Based on the axial variations of T_e and n_e reported above, it is apparent that the quadruple probe results represent plasma conditions averaged over the axial dimension (2.5 mm) of the probe. Additionally, the very large n_e gradients presumed to exist immediately downstream of the thruster exit require extra care in locating and reporting probe positions for both optical and intrusive measurement techniques.

Exit Plane Radial Profiles

Traces for the measured quadruple probe voltage V_{d2} and current I₃ are shown for a typical exit plane radial sweep in Fig. 14. These data have been smoothed so that the traces shown represent the mean signals (with the PPU ripple effects removed). As discussed above, the I₃ trace is symmetric about the thruster centerline, while the V_{d2} trace exhibits the asymmetry attributed to radial gradients in n_e and, to a lesser extent, T_e over the probe face.²¹ The modified quadruple probe theory [Eqs. (10)-(13)] is used to solve for the radial profiles T_e(r) and n_e(r) resulting from the radial quadruple probe data V_{d2} and I₃ (Fig. 14). This analysis indicates that the T_e profile is very flat near the centerline (r<3 mm) and nearly uniform at T_e ~ 0.6 eV. This uniform T_e, along with Eq. (13), is used to determine the radial n_e profile at x=2.25 mm from the thruster exit, shown in Fig. 15. Uncertainty in n_e values far from the centerline is larger due to misalignment between electrode 3 and the flow divergence angle δ .

T-O-F Velocity Probe Results

The effect of the time-of-flight current pulse on thruster operation is demonstrated in Fig. 16, which shows the arcjet current I_{arc} and voltage V_{arc} versus time during a typical pulse. The current drops suddenly as the FET switch closes (Fig. 7) and the 0.1 µF capacitor begins to charge. As the capacitor nears the end of the charge cycle (~ a few 1 µs R-C time constants), the current through the FET leg of the circuit decreases so that the arc current resumes the normal PPU ripple after a small positive ring. The width of the arcjet current deficit pulse is ~ 4 µs. The nature of the behavior of V_{arc} is not immediately clear from Fig. 16. However, the derivative of the arc current (dI_{arc}/dt) has the same qualitative behavior as V_{arc} , implying a possible inductive effect resulting from the arcjet geometry and/or from the current pulse circuitry.

The T-O-F probe response is shown in Fig. 17 for a typical centerline axial velocity measurement. The signal measured at the downstream electrode has been increased by a factor of 6 to account for the lower electron current (lower T_e and n_e) 5 mm downstream of the upstream electrode. Both the upstream and downstream electrodes show noise beginning at t ~ 2 µs that corresponds with the action of closing the FET switch. Approximately 3 µs later, the effect of the arcjet current pulse is monitored as a decrease in the electron current measured by the upstream electrode. After a time delay for the tagged flow to traverse the 5 mm

electrode separation, a similar decrease is noted at the downstream electrode. For both of the signals, the width of the decrease in the measured electron current is ~ $4 \mu s$, approximately the same as that of the current deficit pulse.

Axial velocities are inferred from the convective time delay demonstrated in Fig. 17 and the known separation (5.0 mm) between the two T-O-F electrodes. The time delays are derived from the probe response using the temporal separation between the minima (due to the decrease in electron current) measured at each electrode. This is accomplished by taking the derivative of each (smoothed) signal as shown in Fig. 18. The difference in the locations of the zero crossings is the time required for the tagged flow to travel the 5.0 mm electrode separation.

Results of T-O-F probe centerline velocity measurements are shown in Fig. 19 for data acquired during three individual periods after thruster ignition (cold start). The data show that the centerline axial velocity initially averages ~ 6.0 km/s during the period that the thruster is warming up. After approximately 5-7 minutes of run time, the axial velocity averages ~ 6.5 km/s. These data are consistent with our previous results²¹ which indicate that exit plane ne measurements reach steady state after ~ 4-8 minutes. During the warm-up period, the measured velocities fluctuate between 5.8 and 6.3 km/s. Similarly, the steady-state centerline axial velocities range from 6.3 to 6.7 km/s. These variations, although approximately within the experimental error, may be physical. Previous work utilizing CMV for a 1 kW hydrogen arcjet²⁶ has shown similar velocity fluctuations. In that work, it was determined that the phase of the PPU ripple was not responsible. Similarly, the arcjet current deficit pulse was always triggered at the same phase in the PPU current ripple (Fig. 16) for the data presented in this study. It is unclear what may be causing this phenomenon.

The T-O-F probe was also used for off-centerline measurements to generate an axial velocity profile at the thruster exit. For these data, an attempt was made to align the probe with the flow divergence angle δ . It was found that using a fixed angle ($\theta = 0^{\circ}$) probe produced a flat or even inverted profile with a minima on the centerline. An off axis T-O-F probe that is grossly misaligned with the flow divergence angle ($\theta << \delta$) will see a "faster" plasma at the downstream electrode than at the upstream electrode. Additionally, the electrode separation along the streamline is less than the geometric separation (5.0 mm) by the factor $\cos(\delta)$. Both of these factors artificially increase the measured axial velocities, with the former phenomenon having a greater effect than the latter. Although the plasma streamlines are not known a priori, the error associated with T-O-F probe misalignment for off axis measurements can be minimized by rotating the probe to align it with the expected flow divergence at a given location. The flow divergence has been shown⁴ to be quite large ~ 10 mm from the thruster exit, and is expected to be large at the exit as well. For T-O-F measurements at r = 0, 1, 2, 3and 4 mm, the probe angles used were 0° , 10.8° , 18.0° , 25.2° and 45.0° , respectively. It was assumed that velocities measured in this manner were along the probe angle and were converted to axial components.

Figure 20 shows the results of axial velocity measurements made at the thruster exit plane, with the upstream electrode located 1.0 mm from the exit. The profile is fairly symmetric and is compared with a computational model as discussed below.

Analysis of Crossed Probe Data

Since a measurement of the centerline value of the plasma velocity u_i has been made, quadruple probe data can be used to estimate the heavy particle kinetic temperature T_g=T_i. For the quadruple probe located at x=2.25 mm, the measured ratio of the ion currents collected by electrodes 3 and 4 is I₄/I₃ = 0.55 to 0.60. This current ratio, along with Fig. 4 for f=0.9 (κ =1.07), yields u_i/c_{m,H}+ = 0.95 to 1.05, which indicates that the most probable thermal speed for H⁺ (c_{m,H}+) is approximately equal to the directed ion velocity u_i. For the measured centerline velocity u_i=6.5 km/s, the heavy particle temperature [T_i=(c_{m,H}+)²m_H+/2] is T_g=T_i ~ 2500 K (0.2 eV). Therefore, with the previous results for T_e ~ 6700 K, the extent of thermal non-equilibrium on axis at the thruster exit is T_g/T_e ~ 0.4.

Since the perpendicular electrode P₄ in reality measures I₄ based on a distribution of u_i and T_i along its length, a discussion of the errors introduced by the probe size is in order. Because the quadruple probe theory for determining T_e and n_e uses the measured value of I₄/I₃, no error due to the presence of the perpendicular electrode is involved in determining T_e and n_e. This has been verified (above) with independent triple probe measurements. In interpreting the crossed probe data, however, the geometry of the probe becomes a factor. The current I₄ is likely slightly underrepresented due to changes in n_e and u_i along the electrode length. However, because u_i/c_m,H+ is proportional to 1/(T_i)^{1/2}, a T_i profile peaked at the centerline has the opposite effect on I₄. Assuming a 10% error in I₄, I₄/I₃= 1.1(0.57)~0.63. The ratio u_i/c_m,H+ becomes ~ 1.15, and T_i becomes 2000 K (0.17 eV).

Comparison of Computational Model and Data

A computational arcjet modeling $effort^{30-31}$ has been undertaken by our group concurrent with the exit plane probe studies discussed above. The model is
an axisymmetric, seven species hydrogen/nitrogen plasma code that utilizes a PISO algorithm to solve the computational domain up to the exit plane. Separate energy equations are formulated and solved for the electrons and the heavy species. The anode temperature distribution is included, and the plasma electrical conductivity is coupled with the plasma properties, which allows a self-consistent solution for the current distribution. The model is capable of both chemical equilibrium and non-equilibrium simulations. A complete discussion of the model and assumptions has been presented elsewhere.^{30,31}

Comparisons are made between experimental results and model predictions for a 1 kW-class arcjet operating on 50 mg/s of 2H₂+N₂ to simulate fullydecomposed hydrazine propellant. The arc current is 10.0 A for both model and experiment. The specific powers for the laboratory thruster (112 V, 10 A) and numerical model (94 V, 10A) are 22.4 and 18.2 MJ/kg, respectively. The model specific power is contingent on the empirical value of the anode fall potential used. For the model results discussed below, a chemical non-equilibrium simulation is employed.

The centerline exit plane n_e predicted by the model is $n_e=3 \times 10^{14}$ cm⁻³, which, given the largely varying axial n_e gradients discussed above, seems consistent with previous experimental data (Fig. 13). Because of these large gradients, a direct comparison between the quadruple probe exit plane data (probe tip = 1 mm from exit) and the computational model is difficult.

Centerline exit plane heavy particle temperature predictions agree favorably with experiment ($T_{g,model} = 2800 \text{ K}$, $T_{g,exp.} \sim 2000-2500 \text{ K}$), but T_e results differ by a factor of ~ 2 ($T_{e,model} = 3000 \text{ K}$, $T_{e,exp.} \sim 6600 \text{ K}$). Based on experimental results of

the present and previous studies, it thus appears that the numerical model may be underpredicting electron temperatures at the thruster exit plane. Note that model predictions of thrust, u_i, I_{sp} and T_g are unaffected, since the relative electron concentration is low ($n_e/n_n < 0.001$). Internal nozzle diagnostics³² performed by our group are expected to help resolve the discrepancy between experiment and numerical model predictions.

Fig. 20 shows a comparison between the predicted axial velocity profile and the results of the T-O-F probe surveys. Agreement is excellent and within experimental error in most locations. The numerically predicted profile is slightly more narrow and peaked than the measured profile. This is an expected result because the model flow is constrained in the nozzle (no-slip at wall), which differs from the free expansion that occurs in the plume in the vicinity of the T-O-F probe.

VI. Summary and Conclusions

Experimental results for two electrostatic probe diagnostic techniques are presented for the very near-field plume of a 1 kW hydrazine arcjet. An improved quadruple probe theory, showing excellent agreement with independent triple probe surveys, was used to measure centerline axial profiles of T_e and n_e for x=2.25-6.25 mm from the exit plane. While the gradient in n_e is slight over the region investigated, indications are that the density varies by 1-2 orders of magnitude within the first few millimeters downstream of the exit plane. For the axial position nearest the thruster exit, T_e=6600 K, n_e=3.6 x 10¹² cm⁻³ and T_g=T_i ~ 2000-2500 K. With the heavy particle temperature measurement, the extent of thermal non-equilibrium has been quantified at the thruster exit. A spatially resolved technique for measuring plasma axial velocities has also been developed

and implemented. The radial velocity profile was measured at the exit plane, with a peak centerline velocity of 6500 m/s. The above results were used to evaluate the predictions of a numerical H_2/N_2 arcjet model. Agreement was very favorable for both the exit plane centerline T_g and radial u_i profile. Based on previous and current data, it appears that the model may underpredict exit plane T_e . Internal nozzle diagnostics are expected to illuminate the causes of this discrepancy. Comparison of n_e predictions and experimental data could not be attempted due to the large gradients in n_e present at the exit plane. Clearly, there are significant advances to be made in understanding fundamental arcjet physics, through both experimentation and modeling efforts.

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Fig. 1a. Quadruple probe electrical schematic. Measured quantities are V_{d2} , I_3 and I_4 .







Fig. 2. Schematic of the quadruple probe tip swept through the arcjet plume.



Fig. 3. Ionic and atomic number density ratios for simulated N₂H₄ at p=1 atm.



Fig. 4. Velocity ratio $u_i/c_{m,H}$ + vs. I₄/I₃ for 0.0<f<1.0. For this study, f (= n_H +/ n_e) ~ 0.9, corresponding to K=1.07.



Fig. 5. Calculated quadruple probe response to axisymmetric Gaussian profiles of T_e and n_e with standard deviation σ (mm).



Fig. 6. Side view schematic of the time-of-flight electrostatic probe.



Fig. 7. Schematic of circuit used to trigger arcjet current deficit pulse for T-O-F velocimetry.



Fig. 8. Schematic of the probe mount. Probe axes can be rotated in 0.9° increments.



Fig. 9. Arcjet current and ion saturation current density showing the effects of the PPU ripple. The delay between I_{arc} and j₃ is ~ 3-4 μ s.



Fig. 10. V_{d2} and j₃ vs. time. Figs. 9 and 10 show that n_e increases and T_e decreases with increases in I_{arc} .



Fig. 11. Axial variation of the ion saturation current density. Horizontal error bars are 2.5 mm long, indicating the length of electrode 3.



Fig. 12. Axial variation of T_e and n_e . The n_e gradient is ~ -0.5 x 10^{12} cm⁻³/mm nearest the thruster. Experimental error in T_e is ± 15%.



Fig. 13. Comparison of centerline n_e results from the present work and previous studies.



Fig. 14. Smoothed profiles of $I_3(r)$ and $V_{d2}(r)$ measured by the quadruple probe.



Fig. 15. Measured n_e radial profile for x=2.25 mm from the thruster exit plane.



Fig. 16. Variation of I_{arc} and V_{arc} during the current deficit pulse. The pulse width is much smaller that the period of the PPU ripple.



Fig. 17. Variation of the T-O-F probe signals during the current deficit pulse. Probe electrode signal delay is ~ $0.77 \ \mu s$.



Fig. 18. Derivatives of the smoothed T-O-F probe signals depicting zerocrossings at signal minima locations.



Fig. 19. Measured centerline axial velocity vs. time for arcjet warm-up. The numerical model prediction is also shown.



Fig. 20. Experimental measurements and model predictions of the axial velocity profile at the thruster exit.

III. Application of Internal Langmuir Diagnostics to Arcjets

(AIAA Paper No. 95-2386)

Abstract

A 1.4 kW arcjet thruster has been designed for internal probing of the nozzle. The arcjet employs a single-piece nozzle body, facilitating modeling of the anode heat transfer. Flushmounted planar single Langmuir probes are used for internal diagnostics of the boundary layer flow inside the arcjet nozzle. Two nozzle probes and one constrictor probe are implemented to obtain n_e, T_e, current density and sheath voltage data in the arcjet for $11 \le P/m \le 20$ and flow rates of 60 and 85 mg/sec. Comparison between the experimental results and a computational model is made. Based on the data obtained with a nozzle probe the sheath voltage is predominantly negative, so that an electric field exists near the anode which retards the electrons. The results indicate a high degree of ohmic heating at the sheath edge, especially for positive sheath voltages.

Nomenclature

A_p Probe area [m²]

c Most probable speed [m/s]

cW Specific heat of tungsten $[J/(kg-^{\circ}K)]$

e Electronic charge [Coulomb]

E Total charge particle energy [J]

 E_{∞} Electric field in plasma [V/mm]

Eavg Average resistive electric field between electrodes [V/mm]

 I_D Total probe current [μA]

Ie,sat Electron saturation current [µA]

Iarc Arcjet applied current [A]

Ie, Ii Electron/ion current [µA]

I_{Sp} Specific impulse [sec]

 I_0 Current at zero volts [μA]

j Current density [A/cm²]

je,i Electron/ion thermal current density [A/cm²]

k Boltzmann constant [J/°K]

m Propellant mass flow rate [mg/sec]

me, i Electron, ion particle mass [kg]

ne Electron number density [m⁻³]

 $n_{e\infty}$ Electron density at sheath edge [m⁻³]

r_p Probe radius [μm]

 R_{S} Shunt resistor [Ω]

Te,i Electron /ion temperature [°K]

 ΔT_W Probe temperature rise [°K]

 V_{∞} Plasma potential [V]

V_f Floating potential [V]

V_p Probe biasing potential [V]

Greek Symbols

 $\Gamma(x)$ Flux of charge carriers $[m^2-sec]^{-1}$

κw Tungsten thermal conductivity [W/m-K]

 λ Mean free path [μ m]

 λ_D Debye length [µm]

Collision frequency [sec⁻¹]

 ρW Tungsten density [kg/m³]

 σ Electrical conductivity [ohm-m]⁻¹

 $\phi_{\rm S}$ Sheath voltage [V]

I. Introduction

The goal of this research is to develop an understanding of arc attachment and electrode heating processes at the anode of a low power hydrazine arcjet. The Langmuir probe is employed as a diagnostic for internal probing of the arcjet boundary layer. Single planar flush-mounted probes are situated in the constrictor and anode walls of a 1.4 kW arcjet thruster.

Ultimately, the high performance limit of an arcjet is determined by anode heating. The achievement of high levels of I_{sp} , e.g. above 600 seconds for a hydrazine arcjet, requires P/m (power/ propellant flow rate) exceeding 80 MJ/kg and results in levels of anode heating which can cause electrode failure. One of the major contributions to anode heating is the power dissipation in the anode sheath which is proportional to the sheath voltage. In this paper a diagnostic probe technique to provide measurements of that voltage is described.

Langmuir probes have been used extensively in shock tubes and in hypersonic ionized flows¹⁻⁴ and have been a powerful tool of plasma diagnostics since the early 1920's.⁵ They are relatively simple in design and straightforward in implementation and can provide a convenient method of obtaining local plasma properties n_e , T_e , j and sheath voltage ϕ_s , near the anode surface. Such measurements will help validate numerical model predictions near the anode, and help determine whether these models correctly describe the physics governing arc attachment and anode heating.

An understanding of arc attachment is an important aspect of arcjet models. However, due to the complex near-anode physics and the lack of experimental data (such as n_e , T_e and σ) in this region, some models have used heuristic approaches toward arc attachment, e.g. an artificial conductivity "floor" near the wall for current attachment,^{6,7} or an insulated constrictor, forcing arc attachment in the supersonic region.⁸

Investigations of plasma conditions inside the nozzle and constrictor have been performed on low and medium power arcjets.⁹⁻¹¹ Zube *et al.*^{9,10} used drilled holes or a quartz window to obtain spectroscopic access. The radial profile of H₂ excitation temperature, electron density and atomic hydrogen density in the constrictor was measured. However, due to problems with line width determination near the walls only the data from the inner 60-70% of the constrictor diameter gave reliable results. Temperature restrictions on the window and its effect on energy transfer processes in the constrictor limit the utility of the approach.

Other internal diagnostics work includes that of Hargus *et al*, 12 who performed internal emission spectroscopy measurements in the nozzle expansion region of a 26 kW ammonia arcjet. Three optical access ports were equally spaced along the nozzle wall. Atomic and ionic excitation temperatures of H and NII were obtained as well as electron number densities.

Curran *et al.*¹³ studied arc energy deposition in a segmented anode 1-2 kW arcjet nozzle, the nature of arc attachment, and its effects on performance characteristics of the device. The current was found to attach diffusely to the nozzle wall, with more than 50% of the input power to the arcjet added in the diverging section of the nozzle. The current distribution was found to be dependent on the mass flow rate.

In addition to measuring plasma properties in the arcjet boundary layer, this work employs Langmuir probes in the nozzle to assist in determining azimuthal symmetry of the current distribution and in understanding the physics of arc attachment. Unlike work with segmented anodes, the Langmuir probe approach provides a minimum disturbance to heat flux and current attachment processes at the anode surface.

II. Experimental Approach

A 1.4 kW thruster was constructed with slightly larger dimensions than the standard NASA 1 kW thruster. The constrictor diameter was increased to 1.0 mm (from 0.63 mm) and the constrictor length to 1.5 mm (from 0.25 mm), with an area ratio of 225. By eliminating the anode insert and implementing a single-piece thruster body (Fig. 1), the arcjet can accommodate an array of Langmuir probes, as shown in Fig. 2.

Use of a monolithic body also facilitates modeling of the anode heat transfer. The arcjet is fabricated from a tungsten alloy (HD-18: 95% W, 3.5% Ni, 1.5% Cu) possessing better machining qualities than 2% thoriated tungsten. The cathode is 3.2 mm diameter at the rear of the arcjet, stepping up to 4.8 mm at the thruster head. The cathode gap is set at 1 mm. The converging cone half angle is 30°, while the diverging section is 20° half angle.

The initial experimental tests used three single probes inside the arcjet. These Langmuir probes are made of tungsten wire, surrounded by an alumina insulator, and inserted into a stainless steel tube. The tungsten wire and alumina tubing are held in place with high temperature (4000 °F) zirconia adhesive. Tungsten wire was selected due to its high melting point (3395 °C) and high work function (4.53 V), giving low electron emission from the probe. A calculation of the temperature rise of the probes due to a probe current I_p at a bias voltage V_p is given by:¹⁴

$$\Delta T_{\rm W} = \frac{2 I_{\rm p} V_{\rm p} t^{0.5}}{A_{\rm p} (\pi \rho_{\rm W} c_{\rm W} \kappa_{\rm W})} \tag{1}$$

where A_p is the probe collection area, t is time, and ρ_w , c_w , and κ_w are the probe material properties. Probe collection areas are 0.10 mm² for the nozzle probes and 0.022 mm² for the constrictor probe.

For a 15 volt probe bias at a typical 4 mA current for 50 msec, the maximum temperature rise ΔT_W of the constrictor probe is 30 °K and about 10 °K for the nozzle probes. Assuming the probes operate at the wall temperature, these calculated temperature rises indicate that probe heating is not a problem.

The constrictor contains one 0.18 mm diameter probe and the anode contains two 0.38 mm diameter probes (A and B) separated by 180° at an axial location x = 2.2 mm from the constrictor probe (Fig. 2).

Each probe is fastened to the grounded anode with a miniature tubing connector, and installed in the arcjet body, facilitating removal of the probes for inspection or replacement. The tungsten probe wires are connected to a subminiature thermocouple mount. Belden twisted pair shielding wires connect the mount to a vacuum connector, then to a circuit board, and terminate at the differential amplifier of a 10 MHz digital oscilloscope via BNC cables (Fig. 3). The thruster is seated on a mount inside a 1.5 m³ vacuum tank, with a background pressure of ~ 100 mTorr for flow rates of 50 mg/sec. The simulated hydrazine propellant flow rate can be varied between 50 and 90 mg/sec and is controlled by two Unit mass flow controllers. Plasma parameters, n_e, T_e and j are derived from the probe characteristic data, obtained by biasing the probe with a function generator and measuring the probe current.

The function generator supplies each probe with a ± 14.5 volt peak-to-peak sinusoidal 10 Hz pulse. This gated pulse is triggered externally by a 6 VDC signal. The function generator frequency is kept at 10 Hz to avoid unwanted voltage drops across the low inductance, 100 Ω current sensing resistors R_s. The oscilloscope collects 4096 samples at a rate of 32 kHz, twice the frequency of the 16 kHz arcjet power processing unit.

A 9.4 μ F capacitor is used across each shunt resistor (Fig. 3) to filter noise in the signals. The probe voltage is determined from the function generator output and is corrected for the potential drop across R_s during data reduction.

Prior to each experiment the probes are visually inspected for erosion and checked for continuity in the circuit. The arcjet is initially started at 60 mg/sec, taking about 10-12 minutes to achieve a steady state temperature of ~ 790 °C. Before the system is triggered to initiate data acquisition, the probes are cleaned by applying a 0.5 Hz, 14.5 volt square wave pulse, achieving probe cleaning by ion bombardment. Once the arcjet has achieved steady state operation the function generator is then externally triggered and all three probes are biased simultaneously. The data presented in this paper is for flow rates of 60 and 85 mg/sec, and $11 < P/m \le 20$.

Simple Probe Theory

The Langmuir probe is a metallic electrode inserted into a plasma, biased at positive and negative voltages V_p with respect to the plasma potential, V_{∞} . The current I_p collected by the probe is measured as a function of V_p , generating a probe characteristic (Fig. 4), providing information on the plasma properties.¹⁵

Region A (Fig. 4) is called the electron saturation region, where $V_p > V_{\infty}$ so that electrons are attracted to the probe and ions are repelled. Region A provides the magnitude of the electron saturation current, $I_{e,sat}$.

When the probe voltage equals the plasma potential, V_{∞} , there is no net voltage gradient around the probe and the charged particles migrate to the probe because of their random thermal velocities. In the transition region B, low energy electrons are repelled and ions are accelerated to the probe. If the slope d(lnIe)/dV_p is constant then the electron distribution is Maxwellian and the electron temperature is inversely proportional to the slope.

The equation governing the probe current collected in the transition region (B, Fig. 4), is derived by calculating the flux of charged particles to the probe, assuming a Maxwellian velocity distribution for the electrons and a collisionless thin sheath. The justification for the thin, collisionless probe sheath assumption is discussed later in the paper. The derivation for the probe current in the transition region (Appendix A), results in:

$$I_{p} = n_{e}eA_{p}\left(\frac{kT_{e}}{2\pi m_{e}}\right)^{1/2}exp\left[-\frac{e}{kT_{e}}(V_{p} - V_{\infty})\right]$$
(2)

Equation (2) is independent of probe shape, i.e. it holds for planar, cylindrical, or spherical probes. The electron temperature is found by taking the natural logarithm of Eq. (2) and differentiating with respect to the probe voltage to get:

$$\frac{d(\ln I_e)}{dV_p} = -\frac{e}{kT_e}$$
(3)

Once T_e is obtained from the transition region, the electron number density is found from the electron saturation current, $I_{e,sat}$, obtained from the "knee" in the curve between A and B, Fig 4. This point gives the plasma potential V_{∞} , and $I_{e,sat}$. The electron number density is found by setting $V_p=V_{\infty}$ in Eq. (2):

$$I_p = I_{e,sat} = en_e A_p \sqrt{\frac{kT_e}{2\pi m_e}}$$
(4)

giving:

$$n_{e} = (4.025 \times 10^{15}) \frac{I_{e,sat}}{A_{p}} \sqrt{\frac{1}{T_{e}}} \quad [m^{-3}]$$
(5)

(all units MKS)

When the probe potential equals the floating potential, V_f (Fig. 4) the flux of electrons equals the flux of ions to the probe so $I_p = 0$. In region C of Fig. 2, almost all electrons are repelled, resulting in an ion sheath. The ion saturation current is much less than the electron saturation current because $m_e \ll m_i$ and $T_e \gg T_i$.

III. Experimental Results

Probe Characteristic Data

Figure 5 shows a typical I_p vs V_p characteristic for the constrictor probe. Figure 6 shows a typical characteristic for nozzle probe B. Prior to data reduction the probe voltage and current data are smoothed using 5 point moving average smoothing.

Current Density

Current density j is determined from the characteristic by finding the current at $V_p = 0$ and dividing by the probe area. For the conditions in Figs. 5 and 6, P/m=13.8 MJ/kg, and m=60 mg/sec, j = 0.52 A/cm² for nozzle probe B and j = 0.06 A/cm² for the constrictor probe.

Data Analysis

The electron temperature is found from the slope of the transition region of the ln I_e versus V_p plot (Fig. 7). To obtain the electron current I_e from the total probe current I_p, the ion current I_i at all voltages less than V_{∞} is predicted and subtracted from I_p, using an analytic curve fit to the ion saturation current.

The theoretical form of the ion saturation current depends on the length scale regime, i.e. whether the sheath is thin or thick with respect to the probe radius, and collisional or collisionless with respect to the charged particle mean free paths. Calculations presented below in Table I show that both nozzle probes have borderline collisionless thin sheaths, while the constrictor probe has a thin collisional sheath. The data reduction process applies collisionless theory to all probes.

For the thin sheath case the Laframboise method is used.¹⁶ This approach utilizes a plot of nondimensional ion current versus nondimensional probe voltage (Appendix B). With an empirical fit to the Laframboise plot of nondimensional ion current vs probe voltage for various r_p/λ_D values, an expression for the theoretical ion current is found independent of T_e , for $V_p < V_{\infty}$:¹⁷

$$I_{i} = \frac{1}{4} en_{e} \sqrt{\frac{8e}{\pi m_{i}}} A_{p} (1.127 \sqrt{V_{\infty} - V_{p}})$$
(6)

The plasma potential V_{∞} is found from a graphical analysis of the probe characteristic. Ideally, the plasma potential is the probe voltage at the "knee" in the characteristic on a semilog plot of Ie vs V_p, indicating the end of the transition region and the start of the electron saturation regime.

To use Eq. (6), an estimate of n_e is required. This is done by solving Eq. (6) for n_e and choosing a point $(V_p, I_p=I_i)$ in the ion saturation region; e.g. (Fig. 6) $V_p = -14 \text{ V}$, $I_i = -193 \mu \text{A}$ and $V_{\infty} = 0.6 \text{ V}$. This data is substituted back into Eq. (6) for an estimate of n_e . Using this n_e and V_{∞} in Eq. (6), a theoretical fit to I_i is found for all $V_p < V_{\infty}$.

The resulting ion current $I_i(V_p)$ is subtracted from the total measured current to get I_e . Points $I_e < 0$ are discarded in order to obtain $ln(I_e)$ vs V_p . From this plot, the slope of the electron transition region is found by using a weighted least squares fit to the data. If this region is linear (Fig. 7), the electrons have a Maxwellian velocity distribution, and T_e can be found from Eq. (3). If the transition region is not linear then the T_e obtained from the slope is really an average energy $E_{avg} = 3/2kT_e$. Once T_e is known, the electron number density is found from Eq. (5).

Results Varying P/m

Data for two flow rates are presented, 60 and 85 mg/sec, for $8 \text{ A} \le I_{arc} \le 10.4 \text{ A}$. For the conditions used in these tests n_e and T_e data were unavailable from the constrictor probe because a probe characteristic with distinct saturation regions could not be obtained (Fig. 5).

Figures 8 and 9 are plots of electron temperature at the anode wall as a function of P/m. The data is from nozzle probes A and B, located about 2.2 mm downstream of the constrictor probe, in the supersonic portion of the arcjet nozzle. There appear to be no obvious trends with P/m , though the T_e data for probe A are consistently lower than that for probe B. On average, T_e ~ 16,600 °K ±15% for a flow rate of 60 mg/sec, while for 85 mg/sec the average T_e is about 15,000 °K ±15%.

Figures 10 and 11 present the electron number density measurements for both nozzle probes. For almost all values of P/m the n_e values from probe B are larger than that from probe A. For probe B and a flow rate of 60 mg/sec n_e varies between 1 x 10¹⁷ m⁻³ and 2 x 10¹⁸ m⁻³ for $14 \le$ P/m < 19. Similarly, for an 85 mg/sec flow rate n_e varies between $4x10^{16}$ m⁻³ and 1.5×10^{18} m⁻³ for probe B, for $11 \le$ P/m < 15.

Both the electron temperature and the electron number density data from nozzle probe A are lower than those from probe B. It was discovered, upon closer inspection, that probe B was flush with the wall, while probe A was recessed several probe diameters, reducing its data values compared to probe B, as shown in Figs. 8-11. This prevented simultaneous measurement of j at two azimuthal locations, a technique which will be used in the future to evaluate azimuthal symmetry of the current attachment.

Figures 12 and 13, for 60 and 85 mg/sec flow rates, show that for nozzle probe B the largest j value for 60 mg/sec is ~ 2.8 A/cm² for P/m = 17.9 MJ/kg. The current density has a maximum of about 3.7 A/cm², for a P/m of 12.5 MJ/kg and a flow rate of 85 mg/sec.

There is no obvious trend in the j data for 85 mg/sec. For 60 mg/sec, j appears to increase with P/\dot{m} . For all P/\dot{m} and both propellant flow rates, the j measured by the constrictor probe is generally significantly less than the j measured by nozzle probe B.

IV. Discussion

The plasma-wall interaction in the near-anode region of an arcjet presents a complex problem in understanding the physics of arc attachment. This is due to: (1) the non-neutral sheath layer;

(2) the effects of the wall condition, i.e. catalytic surface or emissive; and (3) non-equilibrium coupling of fluid dynamics and electrodynamics. Understanding the near-anode physics is important in determining the transition between diffuse and constricted modes of current transfer.¹⁸ In the diffuse mode the arc is dispersed over a large area, as opposed to the constricted mode where the arc is highly concentrated. Operation of an anode in a constricted mode leads to large ohmic heating and erosion, thus limiting the electrode lifetime.

Near the anode surface there are several coupled physical processes: convection, diffusion, conduction, and chemical kinetics. The region close to the anode can be divided into several sublayers where large temperature and velocity gradients, non-neutrality and chemical diffusion can play important roles in determining local plasma properties.¹⁹

To determine whether a thin collisionless sheath surrounds the probes the following length scales are evaluated, using experimental data: probe radius r_p , Debye length λ_D , and mean free paths (mfp) for e-H₂ and H⁺-H₂ collisions: λ_{e-H_2} , $\lambda_{H^+-H_2}$. Electron-H₂ collisions are important when electrons are collected by the probe, while ion-H₂ collisions are relevant for ion collection.

Inside the sheath region, (< λ_D from wall) charge neutrality is violated and electrostatic forces control the motion of electrons and ions. The sheath forms near an electrode so as to shield the main body of the plasma from the large electric fields that exist within a Debye length of the wall,²⁰ and maintain charge neutrality in the plasma.

Table I shows some typical values for the relevant length scales based on experimental data.

| | Constrictor | <u></u> |
|----------------------------|-------------|---------------------------------------|
| n_{e} (m ⁻³) | 1 | Nozzle 2 x 10 ¹⁷ |
| $T_{e}(\mathbf{K})$ | ~4,000 | 15,000 |
| ^r p (μm) | 85 | 190 |
| λ _D (μm) | 14 | 19 |
| $\lambda_{e-H_2}(\mu m)$ | 1 | 13 |
| $\lambda_{H}+H_{2}(\mu m)$ | 0.10 | 2 |

Table I. Relevant length scales based on experimental data.

From Table I, for the nozzle probes, $r_p/\lambda_D = 10$, so the sheath is thin. The electron mfp is on the order of the Debye length, and we use a collisionless approach for analyzing the electrons. The ion mfp is ~0.1 of the Debye length, so that the ion current is overstated by using the collisionless assumption.

The constrictor n_e and T_e (Table I) are estimated from the constrictor probe characteristic (Fig. 5) showing ~ 80 μ A of electron saturation current. The calculations show that the constrictor probe is collisional for both electrons and ions, but that the sheath is thin (r_p/ λ _D = 6).

Sheaths

A diagram of the potential distribution between the cathode and anode of an arcjet is shown in Fig. 14. The anode sheath voltage is shown negative with respect to the plasma potential V_{∞} , so that the sheath electric field is towards the anode, causing electrons moving toward the anode to be retarded. A negative sheath voltage can exist if there are large pressure and electron temperature gradients near the electrode, so that the current density, j, is less than the electron thermal current density, je.

The voltage in the sheath is found by comparing the total current density arriving at the anode with the electron thermal current density near the anode. Assuming the probe sheath is collisionless, the sheath voltage is derived first by writing the absolute magnitude of current density j as:

$$\mathbf{j} = \mathbf{j}_{\mathbf{e}} + \mathbf{j}_{\mathbf{i}} \tag{7}$$

where je and ji are the electron and ion thermal current densities, respectively given as:

$$j_e = (1/4)en_ec_e; \quad j_i = (1/4)en_ic_i$$
 (8)

where c_e and c_i refer to the electron and ion most probable thermal speeds. At the sheath edge the electron density is given by the Boltzmann relation, so that $n_e = n_i = n_{e,\infty} \exp[e\phi_s/kT_e]$. Substituting this relation for n_e and n_i in Eq. (8) and using the results in Eq. (7) for j_e and j_i , an equation for the sheath voltage is found:

$$\phi_{s} = -\left(\frac{kT_{e}}{e}\right) \ln\left[\frac{en_{e\infty}c_{e} + en_{e\infty}c_{i}}{4 j}\right]$$
(9)

Typically, the ion thermal current density is a few percent of the electron thermal current density so that this term is dropped, resulting in Eq. (10):²¹

$$\phi_{s} = -\left(\frac{kT_{e}}{e}\right) \ln\left[\frac{en_{e\infty}c_{e}}{4j}\right]$$
(10)

The potential drop of the sheath can therefore be positive or negative, depending on the relative magnitudes of j and je, and can be determined experimentally given n_e , T_e and j near the anode, as extracted from a probe characteristic.

The sheath voltage ϕ_s from Eq. (10) is presented as a function of P/m in Figs. 15 and 16. It is observed that for almost all cases the sheath voltage is negative at the nozzle probe locations.

The experimental results for n_e, T_e, j and ϕ_s are now be compared with computational model results for the conditions 60 mg/sec flow rate, and I_{arc} = 8 and 10 A. The computational model is an axisymmetric, seven species H₂/N₂ plasma code, capable of both thermal and chemical nonequilibrium simulations.²² Comparing the experimental data with the numerical results for T_e in Fig. 8, it is seen that T_e from the model is slightly lower than T_e from the experiment. Closer agreement between the model and our experiment occurs at the lower P/m value of 14.0 MJ/kg for both T_e and n_e data (Table II).

| $I_{arc} = 8A$ | | | | |
|-----------------------------------|------------------------|------------------------|--|--|
| | Experiment | Model | | |
| P/m[MJ/kg] | 13.8 | 14.0 | | |
| n _e [m ⁻³] | 4.4 x 10 ¹⁷ | 8.4 x 10 ¹⁷ | | |
| T _e [K] | 13,600 | 10,300 | | |
| j [A/cm ²] | 0.52 | 11 | | |
| φ _s [V] | -1.0 | 1.46 | | |

Table II. Comparison of Nozzle Probe B data and Numerical Model. $\dot{m} = 60 \text{ mg/sec.}$

| $I_{arc} = 10A$ | | | |
|------------------------|------------------------|------------------------|--|
| | Experiment | Model | |
| P/m[MJ/kg] | 17.2 | 17.1 | |
| ne [m ⁻³] | 1.3 x 10 ¹⁷ | 1.0 x 10 ¹⁸ | |
| T _e [K] | 20,700 | 10,000 | |
| j [A/cm ²] | 0.4 | 13.8 | |
| φ _s [V] | -0.5 | 1.44 | |

A major difference between the experimental data and the model is in the current density, as the model shows current densities (Fig. 12) several times the measured values. The nozzle internal surface area is about 5.5 cm². For $I_{arc} = 10A$, this means that $j_{avg} \sim 1.8 \text{ A/cm}^2$, which has been observed with probe B, Figs. 12 and 13. Therefore, our Langmuir probe results are consistent with an even distribution of j in the nozzle, but based on the model, j is more narrowly attached, Fig. 17. This difference can be resolved by installing probes in different positions.

Figure 17 shows the model current density distribution for a flow rate of 60 mg/sec and two different arc currents. The model predicts that j is not uniform and reaches a maximum value of \sim 14 A/cm² downstream of the constrictor, but about 1.5 mm upstream of the Langmuir probe location. Therefore, it seems possible that the probes are not at the location of maximum current density.

From our experimental data for n_e , T_e , and j, a value for the sheath voltage has been calculated. For most of the experimental conditions ϕ_s is negative, i.e. the anode sheath electric field retards electron motion to the anode, Fig. 14. This implies less anode heating compared with the case $\phi_s > 0$.

For the experimental condition of 85 mg/sec flow rate and P/m = 14.6 MJ/kg, the sheath voltage is slightly positive, $\phi_s = 0.1$ V. The experimental probe characteristic for this positive sheath voltage is shown in Fig. 18 for probe B.

Comparing Figs. 6 and 18, the plasma potential $V_{\infty} = 0.6 \text{ V}$ for $\phi_S < 0$, and $V_{\infty} = -1.8 \text{ V}$ for $\phi_S > 0$. The entire Fig. 18 characteristic is shifted to the left for the $\phi_S > 0$ case.

The resistive electric field $E_{\infty} = j/\sigma$ can now be estimated from Figs. 6 and 18. The electrical conductivity is:²²

$$\sigma = \frac{n_e e^2}{m_e \sum_{i \neq e} n_i c_e \overline{\Omega}_{e-i}^{(1,1)}}$$
(11)

where $\Omega_{e-i}^{-(1,1)}$ is the energy-averaged collision integral. Based on the mean free path calculations in Section IV, the electron-H₂ collision frequency is dominate over ion-H₂ interactions, so that:

$$v_{c} \equiv \sum_{i \neq e} n_{i} c_{e} \overline{\Omega}_{e-i}^{(1,1)} = v_{e-H_{2}} + v_{e-N_{2}}$$
(12)

Using constant collision cross sections for Q_{e-H_2} and Q_{e-N_2} , combined with the experimental data of Figs. 6 and 18, a value of σ can now be estimated. Since j is measured, an estimate of $E_{\infty}=j/\sigma$ is also made. The results are summarized in Table III.

| - | 13.8 MJ/kg | 14.6 MJ/kg |
|-----------------------------------|----------------------|----------------------|
| n _e [m ⁻³] | 4.4×10^{17} | 6.8x10 ¹⁷ |
| T _e [°K] | 13,600 | 13,900 |
| j [A/cm ²] | 0.52 | 2.2 |
| σ [Ω–m] ⁻¹ | 0.15 | 0.23 |
| E∞[V/mm] | 35 | 95 |
| φ _s [V] | -1.0 | 0.1 |

Table III. Data for $\phi_S < 0$ and $\phi_S > 0$.

For $\phi_S < 0$, $E_{\infty} = 35$ V/mm and for $\phi_S > 0$, $E_{\infty} = 95$ V/mm. These values are larger than the average electric field in the arc of $E_{avg} \sim 100$ V/4mm arc length = 25 V/mm, indicating a high degree of ohmic heating at the sheath edge, especially for the case of positive sheath voltage.

V. Conclusion

We have presented preliminary experimental data from the anode boundary layer of an arcjet and have estimated the sheath voltage, the electrical conductivity and electric field values at the sheath edge, using single Langmuir probes. Though n_e and T_e data were not available from the constrictor probe, it was possible to obtain j data, which showed that under most conditions there was significantly more current downstream in the nozzle boundary layer than in the constrictor.

Based on the data obtained from a nozzle probe it appears that the sheath voltage is negative, so that an electric field exists near the anode which retards the electrons. There may be a critical applied current density that leads to a transition of negative to positive sheath voltage. What governs the change in sign of the sheath voltage is relevant to understanding the physics of arc attachment, since the voltage drop in the anode boundary layer and sheath determines the transition between diffuse and constricted modes of current transfer. Also, the sign of the sheath voltage can be indicative of a high degree of ohmic heating at the sheath edge, especially for $\phi_s > 0$.

Comparison between the experimental results and a computational model show approximate agreement for n_e and T_e . However, a discrepancy is found in the current density data. It is possible that the nozzle probes are not located in the region of maximum arc attachment; more probes will have to be placed in the anode wall to address the issue of arc attachment.

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VIII. Appendix A

Derivation of the Current in the Transition Region of the Probe Characteristic

The analysis used in this paper to obtain electron temperature and electron number density from the probe characteristic is based on the assumption of a thin collisionless sheath. The derivation of the electron current in the transition region follows that of Swift and Schar²³ and is presented here for convenience.

Assuming a collisionless thin sheath surrounds the probe, the flux density of charged particles reaching the probe is:²³

$$\Gamma = n(r_p)\overline{u(r_p)} = \int f(c) \, u \, d\gamma \tag{A1}$$

where $d\gamma$ represents the elementary volume in velocity space, f(c) is the velocity distribution function, n(r_p) is the total charged particle concentration at the probe's surface and u(r_p) represents the mean carrier velocity normal to the probe's surface. In the transition region the majority of the carriers are electrons. Therefore, the current to the probe is given by:

$$I_{p} = e\Gamma_{p}A_{p} \tag{A2}$$

To determine the probe current as a function of probe potential, Γ_p needs to be evaluated in terms of V_p. In Eq. (A1) f(c) is assumed to be a Maxwellian velocity distribution for the electrons. For a collisionless thin sheath, with no ionizing reactions, the total energy E, of the electrons is:²³

$$E = \frac{1}{2} m_e u^2 + eV(x)$$
 (A3)

where only the velocity component perpendicular to the probe surface is considered and V(x) is the potential relative to the unperturbed plasma at a point x from the probe's surface, i.e. $V(x) = V_p - V_{\infty}$. In the unperturbed plasma V(x) = 0 and E = (1/2)meu².

Substituting the Maxwellian distribution for f(c) into Eq. (A1).:

$$\Gamma(\mathbf{x}) = n_{e} \left(\frac{m_{e}}{2\pi k T_{e}}\right)^{3/2} \int_{u_{1}}^{\infty} \exp\left[\frac{-m_{e}}{2k T_{e}} u^{2}\right] u du \times$$
$$\int_{-\infty}^{\infty} \exp\left[\frac{-m_{e}}{2k T_{e}} v^{2}\right] dv \int_{-\infty}^{\infty} \exp\left[\frac{-m_{e}}{2k T_{e}} w^{2}\right] dw \quad (A4)$$

where u₁ is the required velocity for the electron to overcome the potential barrier V(x). Since we want the probe current as a function of probe potential, Eq. (A4) is recast in terms of E, the energy in the unperturbed plasma ($E = (1/2)m_eu^2$):

$$\Gamma(x) = n_e \left(\frac{m_e}{2\pi k T_e}\right)^{3/2} \int_{u_1}^{\infty} \exp\left[\frac{-m_e}{2k T_e} u^2\right] u du \times \int_{-\infty}^{\infty} \exp\left[\frac{-m_e}{2k T_e} v^2\right] dv \int_{-\infty}^{\infty} \exp\left[\frac{-m_e}{2k T_e} w^2\right] dw$$
(A5)

The first integration is recast in terms of E; therefore Eq. (A3) is rearranged to solve for u:

$$u = \pm \left\{ \frac{2}{m_e} [E - eV(x)] \right\}^{1/2}$$
 (A6)

Taking the differential of u and substituting Eq. (A6) into Eq. (A4) an equation for Γ is found in terms of E and V(x). Performing the integration:

$$\Gamma(\mathbf{x}) = \frac{n_e}{4} \left(\frac{8kT_e}{\pi m_e} \right)^{1/2} \exp\left[-\frac{eV(\mathbf{x})}{kT_e} \right]$$
(A7)

where $\Gamma(x)$ is the flux of electrons crossing the sheath edge. The flux density of electrons reaching the probe is found by setting $V(x) = V_p - V_{\infty}$. in Eq. (A7):

$$\Gamma_{\rm p}({\rm x}) = \frac{n_{\rm e}}{4} \left(\frac{8kT_{\rm e}}{\pi m_{\rm e}}\right)^{1/2} \exp\left[-\frac{e}{kT_{\rm e}} \left(V_{\rm p} - V_{\infty}\right)\right] \tag{A8}$$

Therefore, the electron current in the transition region of the probe characteristic is given by substituting Eq. (A8) into Eq. (A2):

$$I_{p} = en_{e}A_{p}\left(\frac{kT_{e}}{2\pi m_{e}}\right)^{1/2} exp\left[-\frac{e}{kT_{e}}\left(V_{p} - V_{\infty}\right)\right]$$
(A9)

Collisional Effects

Collisions generally reduce the current to the probe because of the necessity for particles to diffuse up to the probe instead of arriving there by free flight.²⁴ The collisionless equations presented above place an upper bound on the collected current.¹⁷ To calculate the reduction in the collected current, scattering collisions must be considered.

Collisional sheath effects are important if the electron-neutral or ion-neutral mean free paths are much less than the Debye length. If the probe sheath is collisional then the above probe theory requires modification. For example, the total energy of an electron in the sheath would not be given by Eq. (A3), but would have additional energy loss terms and/or gain terms due to ionization reactions or particle scattering. The species conservation and energy equations would have to be solved in addition to Poisson's equation in the sheath.

IX. Appendix B

Laframboise Method

The ion current, which must be known to derive I_e , is an empirical fit to a theoretical formulation of Laframboise¹⁶ for collisionless sheaths, and is presented here as a convenience to the reader.

The thin-sheath, transition to a thick sheath and the thick sheath regimes can all be treated by the Laframboise method. Lafram-boise solved Poisson's equation numerically, assuming a Maxwellian distribution of ions at T_i and electrons at T_e , for various values of r_p/λ_D .

The Laframboise approach utilizes a plot of j_i^* versus X_p^* for several values of r_p/λ_D . The parameter, j_i^* , is a dimensionless quantity that multiplies the probe collection area and accounts for the Bohm acceleration in the presheath region. X_p^* is a nondimensional parameter that indicates how far the probe is biased into ion saturation compared to T_e, given by:¹⁷

$$X_p^* = \frac{e(V_{\infty} - V_p)}{kT_e}$$
(B1)

The ion current is given by:¹⁷

$$I_i = \frac{1}{4} \operatorname{en}_e \sqrt{\frac{8kT_e}{\pi m_i}} A_p j_i^*$$
(B2)

To use Eq. (B1) both n_e and T_e are required. A simplification is made by an empirical fit to the Laframboise plot:

$$j_i^* = 1.127 (X_p^*)^{1/2}$$

With this empirical fit, an expression for the theoretical ion current is found independent of Te:

$$I_{i} = \frac{1}{4} en_{e} \sqrt{\frac{8e}{\pi m_{i}}} A_{p} (1.127 \sqrt{V_{\infty} - V_{p}})$$
(B3)

Equation (B3) can now be used as an analytical fit to the ion current data and subtracted from the total current to obtain I_e .



Fig. 1. The 1.4 kW thruster incorporates a single-piece thruster body to accommodate internal probing of the arcjet boundary layer.



Fig. 2. Anode design with planar Langmuir probes for internal n_e , T_e and current density measurements. Nozzle length is 19.2 mm; constrictor length is 1.5 mm. The nozzle probes are 2.2 mm downstream of the constrictor probe, at an area ratio of ~ 7. The constrictor probe is ~ 0.5 mm upstream of the constrictor exit.



Fig. 3. Experimental schematic for three single Langmuir probes.



Fig. 4. Schematic of a typical probe characteristic, showing the various regions of interest that provide information on the plasma properties.



Fig. 5. Experimental probe characteristic for the constrictor probe for conditions P/m =13.8 MJ/kg, $\dot{m} = 60$ mg/sec. V_p is measured with respect to the anode potential, which is zero. Probe collection area is 0.022 mm².



Fig. 6. Experimental probe characteristic for the anode wall probe B for conditions P/m =13.8 MJ/kg, $\dot{m} = 60$ mg/sec. Probe voltage is measured with respect to the anode potential, which is zero. The current density j is I₀/A_p. Probe collection area is 0.10 mm².



Fig. 7. A semilog plot of Ie versus V_p for P/m =13.8 MJ/kg, m = 60 mg/sec. The transition region, $-3.5 \le V < 0$, is linear over 2 decades, signifying a Maxwellian electron distribution. V_{∞} and Ie,sat are obtained from the "knee" in the characteristic.



Fig. 8. Electron temperature T_e versus P/m for 60 mg/sec. Probes A and B are located 2.2 mm downstream of the constrictor exit, and are separated by 180°. The values from a numerical model²² are also shown. Data from probe A is not valid, as discussed.


Fig. 9. Te versus P/m for 85 mg/sec. Data from probe A is not valid, as discussed.



Fig. 10. Electron number density n_e versus P/m for a fixed flow rate of 60 mg/sec. The values from a numerical model²² are also shown. Data from probe A is not valid, as discussed.



Fig. 11. Electron number density n_e versus P/m for 85 mg/sec. Data from probe A is not valid, as discussed.



Fig. 12. Current density at anode potential versus P/m for 60 mg/sec. The values from a numerical model²² are also shown.



Fig. 13. Current density at anode potential versus P/m for 85 mg/sec.



Fig. 14. Potential distribution diagram between the cathode and anode of an arcjet, and in the near-anode region showing a negative sheath voltage.



Fig. 15. Sheath voltage at anode potential versus P/m for 85 mg/sec.



Fig. 16. Sheath voltage at anode potential versus P/ \dot{m} for 60 mg/sec. Values from a numerical model²² are also shown.



Fig. 17. Current density distribution along the anode wall based on the numerical model²² for the 60 mg/sec and $I_{arc} = 8$ and 10 A. Constrictor entrance is located at 5.7 mm and the constrictor exit is at 7.2 mm.



Fig. 18. Probe B characteristic for $\phi_S = 0.11$ V at 85 mg/sec, P/m = 14.6 MJ/kg. The location of the "knee" makes $V_{\infty} < 0$. Compare this characteristic with Fig. 6, for $\phi_S = -1.0$ V.

