Flow actuation using radio frequency in partially ionized collisional plasmas

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We present a multidimensional theoretical model for a better understanding and the design of the dielectric barrier discharge-induced momentum exchange. Specifically, the formulation is used to predict surface discharge using two-dimensional asymmetric electrode configurations. Predictions for charge densities, the electric field, and gas velocity distributions are shown to mimic trends reported in the experimental literature. We also predict the electron charge accumulation on the dielectric surface self-limiting the discharge. © 2005 American Institute of Physics.

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Recent experimental focus on radio-frequency (rf)-based dielectric surface discharge at atmospheric pressures shows tremendous potential for boundary layer flow actuation which will be useful for aerospace and many other applications.1,2 These actuators can apply large electrodynamic (EHD) forces in a relatively precise and self-limiting manner, have rapid switch-on/off capabilities and require no moving parts. Typical electrode arrangements for these devices are similar to that shown in Fig. 1 where the grounded electrode is surrounded by an insulator and a voltage fluctuating with rf is applied to the electrode exposed to the gas. The electric forces are generated through electrodes embedded in the dielectric. As the applied voltage becomes sufficiently large for a given interelectrode distance d and pressure p, the dielectric surface adjacent to the rf electrode produces a barrier discharge weakly ionizing the surrounding gas. The experimental data suggest 1–20 kV peak-to-peak applied voltage with 2–50 kHz rf is suitable for these actuators operating at atmospheric pressure. The plasma at this pressure is highly collisional causing an efficient energy exchange between charged and neutral species. In this discharge, microfilaments of nanosecond duration with many current pulses in a half-cycle maintain the optical glow.

Due to a combination of electrodynamic and collisional processes, the proper mechanism of which is yet to be theoretically resolved, charged separated particles induce the gas particles to move. For accuracy and fidelity, it is essential that the force model be derived from first principles through a simulation of the elementary mechanisms that yield the discharge. Roy and Gaitonde3 demonstrated such a model for volume discharge between two dielectric coated electrodes using a finite element (FE)-based one-dimensional multifluid formulation for atmospheric conditions. The model was relatively basic but formed the framework of a first-principles-based methodology. Simulations showed reasonable comparison with reported rf discharges in partially ionized helium gas between two insulated electrodes.

Here, the multidimensional model for surface dielectric barrier discharge (DBD) at atmospheric pressures is presented as a first approximation. The formulation is consistent and calculates charge and neutral number densities, their momentum dynamics, electric field, and potential distribution in two spatial dimensions. For simplicity, inert helium is used as a carrier gas. The charge particle a = e, i distributions nα are considered non-Maxwellian obeying the continuity relation ∂nα/∂t + ∂nαVα/∂x = nαZ−rnαni. The electron temperature is nearly uniform at 1 eV = 11 600 K and the ions and neutrals are in LTE at 300 K. The working gas helium is maintained at bulk pressure p = 300 Torr and temperature T = 300 K. The current momentum follows the diffusion form (neglecting inertia), nαVαj = sgn(e)nαμaeVαj − Dα∂nα/∂xj, where Eα = −∂φj/∂xj is the electric field, and Vαj is the jth velocity component for the particle α. The potential distribution φ is calculated based on Poisson equation, e(∂2φ/∂x2)j = −q, where charge q = e(ni − nα). The electron diffusion is based on the Einstein relation, Dα = (Tα/εj)μα, where Tα is the energy in electron volts, ε is the elementary charge, μj is the permittivity, and μj = e/(meεα) is the mobility of an electron, where me ∼ 10−26 s is the electron-neutral collision frequency while μj ∼ 107 cm2/s. The ion diffusion Dj ∼ 500 cm2/s at 300 K, and the ion mobility μi is given as Ref. 5. The ionization rate z for helium gas used here is z/nα = (Vεiσei(Vth)j), the averaging is done over the velocities of the electrons whose energy is sufficient for ionization. The ionization process can be described as e + He → He++ e + e′, where e and e′ have different energy levels. The coefficient of recombination is given as r = (Vεiσei(Vth)) = 1.09 × 10−14 T−0.2 nα cm3/s. Here, Vth is the electron thermal velocity and σei is the

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electron-ion collision cross section. For simplicity, we ignored recombination effect on momentum at this stage.
Here, the electron plasma frequency, \( \omega_{pe} = 2\pi \sqrt{\left( \frac{n_i}{m_e} \right)} \), is much higher than the applied voltage frequency. Also, the momentum exchange terms due to electron-electron and ion-ion interaction are negligible in comparison with the electron-ion and ion-neutral momentum exchange as the relative drift between similar particles is small in comparison with the drift between electrons, ions, and neutrals.

It is important to calculate the neutral dynamics consistently so as to capture the momentum transfer process accurately. The conservation of neutral particles \( n_i \) of the carrier gas at atmospheric condition follows: 
\[
\frac{\partial n_i}{\partial t} + \nabla \cdot (n_i \mathbf{V}_f) = -n_i \mathbf{z} + m_n \mathbf{R}
\]

The dynamics of neutral particles are determined using the electron-neutral and ion-neutral collisional momentum exchange and ionization.

The averaged velocity \( \mathbf{V}_f \) of the fluid of density \( \rho \) following \( \frac{\partial \rho}{\partial t} + \nabla \cdot (\rho \mathbf{V}_f) = 0 \) is based on the Navier–Stokes equations with EHD body force: 
\[
\frac{\partial \mathbf{V}_f}{\partial t} + \nabla \cdot (\mathbf{V}_f \mathbf{V}_f) = -\nabla p + \frac{1}{\rho} \nabla \cdot \mathbf{q} E \cdot \mathbf{V}_f + \nabla \cdot \mathbf{V}_f
\]

The electric-field lines clearly represent a directional bias to the right. The most ions accelerate toward the right along these lines leaving only a few to move in the opposite direction. The net result is an order of magnitude higher gas velocity downstream of the right edge of the exposed anode with a two orders of magnitude smaller negative velocity near the left edge. These results stress that the asymmetry enhances the electric field and thus influences flow direction.

The multiscale ionized gas (MIG) flow code was used to solve the charge and gas hydrodynamic equations. MIG implements a powerful high-fidelity FE procedure adapted from fluid dynamics to overcome the stiffness of the equations generated by multispecies charge separation phenomena. FE techniques are especially suitable for their adaptability to arbitrary multidimensional geometries and boundary conditions. Here, a two-dimensional (2D) bilinear FE formulation is employed along with implicit time marching to solve the continuity and momentum equations for all species and gas. The solution process consists of two steps—first solve for all species (including neutrals) to calculate the charge separation and electric field up to a periodic asymptote with a time step of up to 100 times the dielectric relaxation time scale, then use those as body force to predict the fluid velocity and continuity evolving in time with the flow time scale. The Galerkin weak statement associated with a variational integral underlines the development of this numerical algorithm. The solution of the Newton iteration for the resulting matrices is convergent at any given time step when the maximum value of the residual relative \( L_2 \) norm for each of the state variable becomes smaller than a convergence criterion of \( 10^{-4} \).

The computational domain \((x, y, z): 0.01 \) consists of a lower half \((y: 0, 0.005) \) insulator with dielectric constant

\[
\epsilon_{r,i} = 1.0055 \epsilon_0
\]

FIG. 2. Instantaneous solution for asymmetric configuration: (a) Electric-field lines, and (b) streamwise component of the electric field.

FIG. 3. Contours and line plots show negative charge accumulation on the downstream dielectric surface of the rf electrode: (a) Line plots of charge distribution at various time and spatial locations, and (b) 2D contours of \((n_i, n_e)\) at peak voltage.
important effect on flow control. The small time scale of the discharge allows minimal charge transport and energy dissipation, thus less gas heating.

The 2D charge solutions plotted in Fig. 3(b) show the nonuniform nature of the distribution along the edge of the dielectric. A residual of electrons remains deposited on the dielectric surface downstream of the exposed powered electrode for the entire duration of the cycle and causes a net electric force in the direction from the electrode to the downstream surface. This is why the net near wall jet velocity is unidirectional. The peak streamwise electron current for this solution is predicted at the two corners of the exposed electrode with a maximum magnitude (~3 mA) around \( \pi \) radian and the minimum current (~0.2 mA) is computed for the \( \pi/2 \) radian. This periodic behavior of the electron current explains the charge build up on the insulator surface opposing the applied voltage just beyond the peak as the applied voltage starts decreasing, thus bounding the discharge. This is also confirmed by the experimental data (see, for example, Fig. 2 of Ref. 12) where the current magnitude peaks at about \( \pi \) radian.

Figure 4(a) plots the computed force due to charge separation and the electric field and shows a local maximum about the exposed electrode with nearly ten times force near the right edge as compared to the left edge. The force dissipates quickly beyond the electrode. The time evolution of the force components due to charge separation is plotted in Fig. 4(b) for the positive half of the cycle at a point 1 mm downstream of the exposed electrode and 0.1 mm above the dielectric. The figure shows a steady rise and fall of the force following with the applied frequency. The negative half of the cycle is not shown as the magnitude of force is order of magnitude smaller in that segment. As the interelectrode gap \( d \) reduces the force increases, however the characteristic remains the same. The velocity field based on a quiescent flow initial condition shows streamwise component of the computed gas velocity in Fig. 5(a) and depicts a strong wall jet downstream of the rf electrode away from the dielectric surface. The highest momentum transfer between the gas and the charge particles occur slightly downstream of the peak force location. The local vertical line plots at six streamwise locations describe how the flow velocity increases responding to the increase in axial electric field (Fig. 5(b)) and experimental results show very similar feature. The peak jet velocity is nearly 1.7 times the freestream velocity.

In conclusion, a 2D FE-based formulation of plasma-fluid interactions is given for modeling the dielectric barrier discharge for flow control purposes. The partially ionized plasma is modeled using the multicomponent fluid equation for helium gas. The computed results are similar to the experimental data showing the exposed electrode is situated upstream of the peak location of the electric field and thus the velocity. The 2D predictions mimic the self-limiting discharge mechanism due to the charge accumulation on the insulator surface and the lowest electron current just beyond the peak applied voltage. The results indicate that the surface charge on the dielectric is the key for unidirectional momentum transfer for asymmetric electrode configuration. A better selection of electrochemical data along with computation of electron temperature gradient is necessary for accuracy of the solution. More investigation is needed to ascertain the optimum electrical parameters and applied voltage shape (e.g., saw tooth, square wave, etc). In the near future, the model should be extended to air with negative ions and additional mechanisms in the source terms.

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