



Full Length Article

Investigation of the effect of DC electric field on a small ethanol diffusion flame

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HIGHLIGHTS

- Effects of DC electric field on a small ethanol diffusion flame are analyzed.
- A numerical study has been performed to elucidate the experimental observations.
- Applied electric field increases flow velocity, promotes the fuel/oxidizer mixing.
- Applied electric field enhances combustion resulting in higher flame temperature.

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ABSTRACT

A small ethanol diffusion flame exhibited interesting characteristics under a DC electric field. A numerical study has been performed to elucidate the experimental observations. The flow velocity, chemical reaction rate, species mass fraction distribution, flame deformation and temperature of the flame in the applied DC electric field were considered. The results show that the applied electric field changes the flame characteristics mainly due to the body forces acting on charged particles in the electric field. The charged particles are accelerated in the applied electric field, resulting in the flow velocity increase. The effects on the species distribution are also discussed. It was found that the applied electric field promotes the fuel/oxidizer mixing, thereby enhancing the combustion process and leading to higher flame temperature. Flame becomes shorter with applied electric field and its deformation is related to the electric field strength. The study showed that it is feasible to use an applied DC electric field to control combustion and flame in small-scale.

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1. Introduction

The characteristics of micro- and meso-scale combustors with different configurations or under different external conditions have attracted lots of attention, e.g. [1–5]. The effects of an electric field applied on flame have been studied recently, e.g. [6–9]. Researchers found that the electric field can improve the stability of combustion. The external electric field was also used as a means for flame control, such as taking flame as an electrically active component based on voltage-current characteristics in the circuit [10]. The effects of electric fields on flame included stabilizing the combustion, increasing flame speed, reducing the soot production and emission, changing the flame temperature and shape [11–15]. These findings also imply that the efficiency of practical

non-premixed combustion systems could be improved by applying an electric field [16]. It has been identified that there are three major effects produced by the electric field on the flame, including the thermal effect, the ionic wind effect and the electrical-chemical effect [17].

Experimental and numerical methods have been used to study the effects of electric fields on flame behavior. Meng et al. [18] found that the flame propagation and combustion properties were significantly affected by the DC electric fields and the flame shape would become a prolate spheroid by the electric body force in the electric field. Imamura et al. [19,20] investigated the flame deformation of ethanol droplets in different vertical electric fields experimentally and the relation between the applied voltage and electrode distance was observed. Kim et al. [21] considered the stabilization characteristics of liftoff and blowoff in nonpremixed laminar jet flames in a coflow for propane fuel by applying AC and DC electric fields experimentally. van den Boom et al. [22] studied the influence of a DC electric field on the laminar burning

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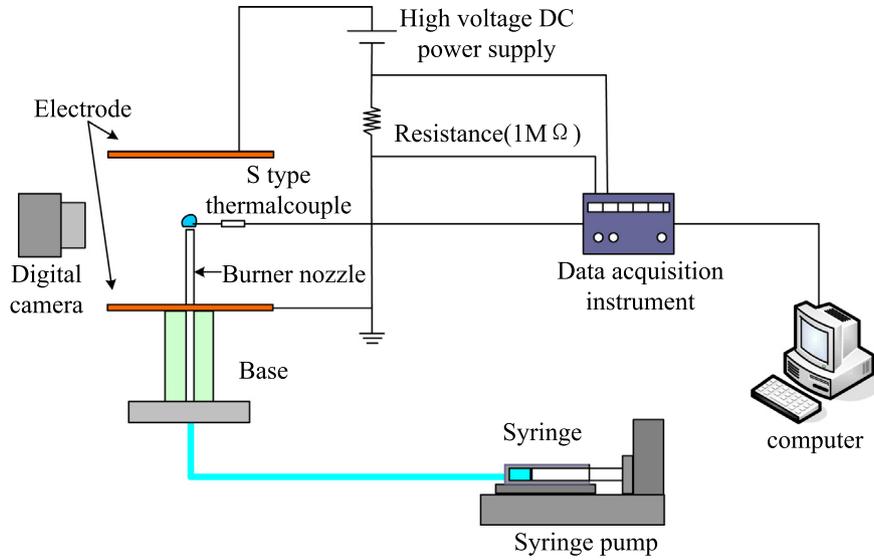


Fig. 1. Experimental system.

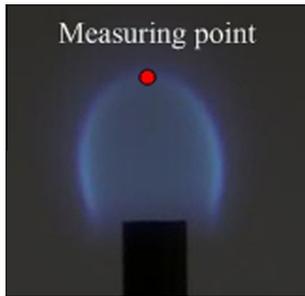


Fig. 2. Flame temperature measuring point.

3. Model description

3.1. Geometric and mathematical model

Fig. 3 shows a schematic of the axisymmetric model of the small-scale combustor with two plate electrodes. The model scale depended on the actual experimental system. In the experiments,

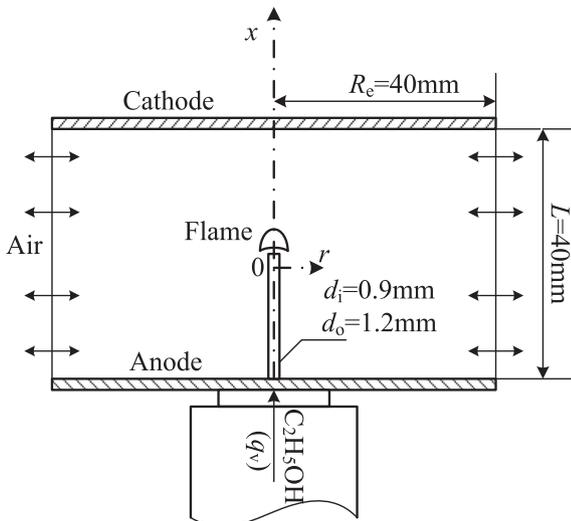


Fig. 3. The configuration studied.

electrode spacing had been adjusted to 40 mm. The upper electrode was connected to the high voltage of the power supply as the anode and the upper electrode was connected to the low voltage as the cathode. In consideration of the domain symmetry, a half of the cross-section of the domain was selected in the axisymmetric model to reduce the computational cost.

Since the main focus of the study was on the steady mean flow field, only steady and axisymmetric numerical simulations were considered. The mixture flow and combustion process followed conservation equations including the continuity, momentum, energy and species conservations for the governing equations.

In steady state, the governing equations of axisymmetric cylindrical coordinates are presented as follows:

Continuity:

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

where ρ is the density of flow, u is the x direction of flow velocity, and v is the r direction of flow velocity.

Momentum:

x direction:

$$\begin{aligned} \frac{\partial(\rho uu)}{\partial x} + \frac{1}{r} \frac{\partial(r\rho v v)}{\partial r} = \\ - \frac{\partial p}{\partial x} + 2 \frac{\partial}{\partial x} \left(\mu \frac{\partial u}{\partial x} \right) + \frac{1}{r} \frac{\partial}{\partial r} \left(r\mu \frac{\partial u}{\partial r} \right) + \frac{1}{r} \frac{\partial}{\partial r} \left(r\mu \frac{\partial v}{\partial x} \right) + F_x \end{aligned} \quad (2)$$

r direction:

$$\begin{aligned} \frac{\partial(\rho u v)}{\partial x} + \frac{1}{r} \frac{\partial(r\rho v v)}{\partial r} = \\ - \frac{\partial p}{\partial r} + \frac{\partial}{\partial x} \left(\mu \frac{\partial v}{\partial x} \right) + \frac{\partial}{\partial x} \left(\mu \frac{\partial u}{\partial r} \right) + \frac{2}{r} \frac{\partial}{\partial r} \left(r\mu \frac{\partial v}{\partial r} \right) + \frac{2\mu v}{r^2} + F_r \end{aligned} \quad (3)$$

where μ is the dynamic viscosity, p is the pressure on the flow element, and F_x , F_r are the body force on the element.

The source term of momentum equation in the axial direction is given as:

$$F_x = F_e = Ee(n_+ - n_-) = Een_c \quad (4)$$

where e is the electron charge, n_+ is the positive charge density, n_- is the negative charge density, and n_c is the net charge density.

The model was simplified and the net charge density was considered. The number of charged particles was estimated according

to the literature (10^9 – 10^{12} cm $^{-3}$) [17–19]. According to Eq. (4), the electric field force was estimated as

$$F = E n_c = 1600 \text{ N/m}^3, \quad E = 100 \text{ kV/m}, \quad n_c = 10^{11} \text{ cm}^{-3}$$

which is similar to the results in the literature (0–2000 N/m 3) [21].

Energy:

$$\frac{\partial}{\partial x}(\rho u h_i) + \frac{1}{r} \frac{\partial}{\partial r}(r \rho v h_i) = \frac{\partial}{\partial x} \left(\frac{k}{c_p} \frac{\partial h_i}{\partial x} \right) + \frac{1}{r} \frac{\partial}{\partial r} \left(r \frac{k}{c_p} \frac{\partial h_i}{\partial r} \right) - \frac{\partial(h \mathbf{J}_i)}{\partial x} - \frac{1}{r} \frac{\partial(h \mathbf{J}_i)}{\partial r} + \frac{\partial(\tau u)}{\partial x} + \frac{1}{r} \frac{\partial(r \tau v)}{\partial r} + S_h \quad (5)$$

where k is effective heat transfer coefficient, h_i is the enthalpy of the species i , \mathbf{J}_i is the diffusive flux of the species i , c_p is the constant-pressure specific heat capacity of the mixture, the temperature, τ is the viscous dissipation stress, and S_h is the volumetric heat source term.

In our previous work [28], it was found that the external electric energy was very small compared with the actual burning thermal energy of ethanol in this study. This was also mentioned in Ref. [22]. So the external energy by the electric field was considered insignificant and ignored in the model.

Species:

$$\rho u \frac{\partial Y_i}{\partial x} + \rho v \frac{\partial Y_i}{\partial r} = - \frac{\partial \mathbf{J}_i}{\partial x} - \frac{1}{r} \frac{\partial(r \mathbf{J}_i)}{\partial r} + R_i + S_i \quad (6)$$

where Y_i is the mass fraction of species i , R_i is the net reaction rate, S_i is the additional generation rate caused by source terms.

In order to simplify the calculation, a one-step chemical reaction model was used.



The governing equations were solved using an implicit solver which is pressure based. The system was closed with appropriate boundary conditions on each side of the computational domain. For the small diffusion flame, the boundary conditions are consistent with the experimental condition, as shown in Table 1. Identical boundary conditions were employed for the condition with electric field and without electric field except for the electrode conditions. It was assumed that the purity of liquid ethanol was 99.7%, the temperature of the liquid fuel applied was 300 K and the ethanol was completely burnt. The model considered that the liquid ethanol had evaporated into gas near the nozzle outlet. This is consistent with the actual experimental observation.

3.2. Numerical simulation and validation

Taking the differences of the solid and the fluid zones into consideration, the computational domain was divided into the ceramic tube solid zone, the fluid and combustion zone. A systematic grid independence test was carried out. The final mesh chosen had a total number of elements of 159,980. Numerical simulations were performed for a reacting flow system with ethanol and air used as the fuel and oxidant respectively. In the simulation study, the oxygen mass fraction was taken as 23% (in ambient air). The ethanol was ignited above the burner nozzle outlet by assuming a temperature of 1000 K.

The accuracy of the present numerical model has been evaluated by comparing the measured and predicted flame shape and

Table 1
Boundary conditions.

Liquid ethanol inlet boundary condition	VELOCITY_INLET
Outlet boundary condition	PRESSURE_OUTLET
The wall of tube inside and outside	WALL
The electrodes condition	WALL

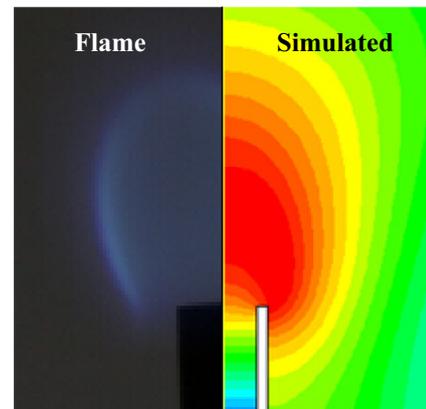


Fig. 4. Measured flame image and simulated temperature field without electric field ($q_v = 1.2$ ml/h).

Table 2
Flame temperature and errors with flow rate of 1.2 ml/h.

Temperature/K	No electric field	Electric field
Experimental data	1326	1388
Numerical data	1446	1472
Discrepancies	9.0%	6.1%

temperature. The small flame image captured by camera in the experiment and the flame temperature distribution of numerical calculation are shown in Fig. 4, respectively. Generally, in the high temperature region, the brightness of the flame will be high. So the flame image captured by camera and the calculated temperature field could indirectly reflect the flame shape. It was found that the flame shapes obtained from the two approaches were similar, which were both approximately spherical. Through the comparison of the flame temperature measured and calculated data, the difference between them was about 9%, and it is shown in Table 2. This was considered as acceptable. In addition, when the operating condition changed, the flame and temperature obtained by the measured results and calculated results both have the same trend of variation. It also implied the accuracy of the present numerical simulation.

4. Results and discussion

4.1. Velocity field

The flame of liquid ethanol in our small-scale combustor can only maintain stability within a certain range of fuel flow rates. When the fuel flow rate is too low or too high, the small flame cannot be ignited or will become oscillatory. In this study, the flame in a steady state was examined. The fuel flow rate of 1.2 ml/h where the flame could maintain stability was selected to perform the studies. Results about flow field of the flame were obtained. Fig. 5 shows the flow velocity vector diagrams near the flame without electric field and under positive electric field. Fig. 6 shows the comparison of flow field variations without and with the applied electric field. It was found that the velocity magnitude increased and the velocity changed more intensely when the electric field was applied.

This is mainly due to the influence of ionic wind [23]. There is a large number of positively and negatively charged particles being produced by chemi-ionization in the hydrocarbon flame reaction zone [30]. The positive electrode is below the flame and the negative electrode is above. These charged particles would affect the combustion progress when the DC electric field was applied. When

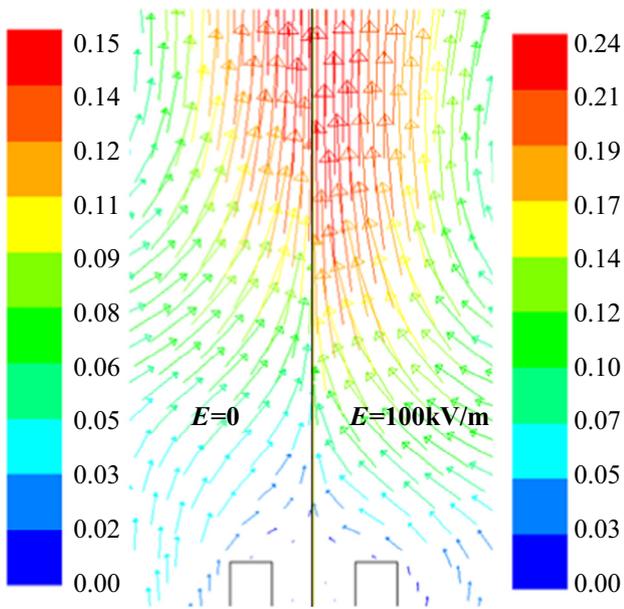


Fig. 5. Calculated velocity field for the diffusion flame. Left: no electric field, right: $E = 100 \text{ kV/m}$ ($q_v = 1.2 \text{ ml/h}$).

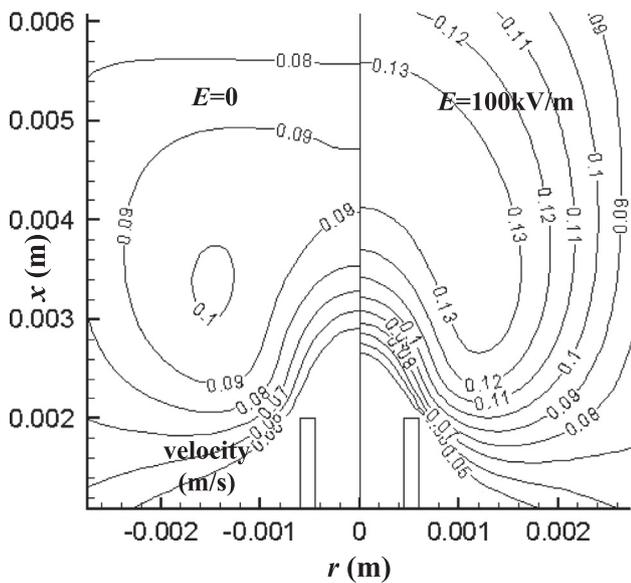


Fig. 6. Calculated velocity contour. Left: no electric field, right: $E = 100 \text{ kV/m}$ ($q_v = 1.2 \text{ ml/h}$).

the electric field was applied, the positively and negatively charged particles would move toward the opposite polarity electrode by the force of electric field. Charged ions were accelerated in the electric field and transferred their momentum to the neutral molecules by colliding with them. With the relatively small mean free path, the ions were accelerated by the electric field after each collision [31], which produced a large number of neutral molecules moving toward the electrodes. The net effect of this process is a significant body force produced by the electric field or the so-called “ionic wind”. In the same region, the flow velocity near the flame changed more intensely under the electric field compared with the case without the electric field. It suggested that the velocity gradient is larger, which also indicated the existence of the electric field force as a source for the momentum [21].

4.2. Mass fraction distribution

Obtained by numerical simulations, Figs. 7 and 8 show the ethanol and CO_2 mass fraction distributions, respectively. The effects of applied electric field on the species distribution can be observed. As seen from the figures, when the DC electric field was applied, the concentration of each species has a tendency of inward contraction. The increasing flow velocity near the flame due to the addition of electric field enhances the mixing of species. The applied electric field could increase the flame propagation velocity [18] and flame temperature by increasing the ion number density and redistributing the ion concentration, which enhances the combustion. The fuel rapidly spread to the air and met the oxidant and then the reaction took place. The fuel and oxidant burnt faster and were quickly consumed. The intensity of combustion process increased, and species distribution had an inward contraction.

A similar phenomenon can be seen from the calculated mass fraction profiles of O_2 and CO_2 on the axis as shown in Fig. 9. The maximum value of the mass fraction of CO_2 under positive electric field was a little higher than that without the electric field and the point of maximum value moved upstream. The mass fraction of O_2 also moved upstream and became more sharply decreasing close to the flame. This observation agreed with the experimental results, which showed the reduction of flame height with the increasing electric field strength. These results showed that the flame had a tendency of contraction, which also implied the intensification of the reaction.

4.3. Chemical reaction rate

From the numerical results, some changes in chemical reaction were observed when the electric field was applied. Fig. 10 shows the chemical reaction rate in the flame. It can be found that when the electric field was applied, the reaction rate in the flame increased and the scope of the reaction expanded slightly. While the surface of chemical reactions has a tendency of contraction, which is consistent with the mass fraction discussed before. It also suggests that the flame size decreases. It can be inferred from the above mentioned that the velocity near to flame zone was increased, which enhanced the fuel/oxygen mixing. The charged particles received an acceleration by electric field, which made

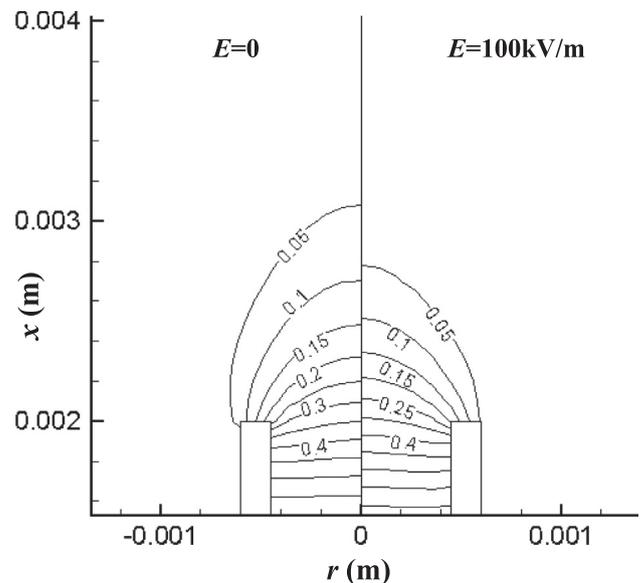


Fig. 7. Calculated ethanol mass fraction. Left: no electric field, right: $E = 100 \text{ kV/m}$ ($q_v = 1.2 \text{ ml/h}$).

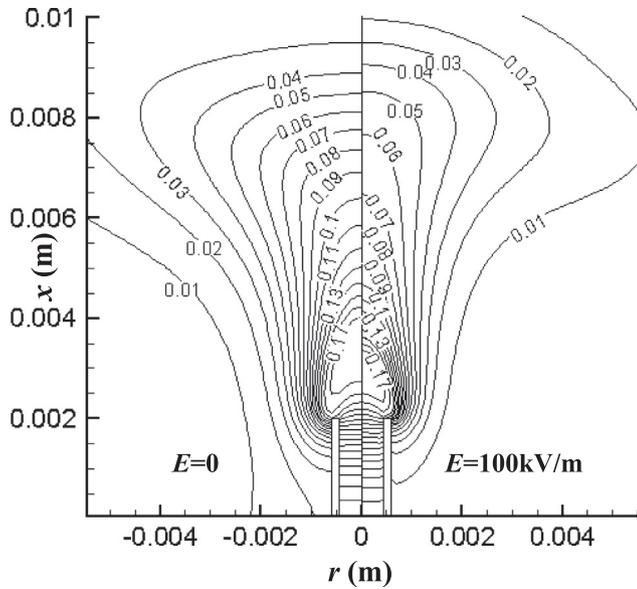


Fig. 8. Calculated CO₂ mass fraction. Left: no electric field, right: $E = 100\text{ kV/m}$ ($q_v = 1.2\text{ ml/h}$).

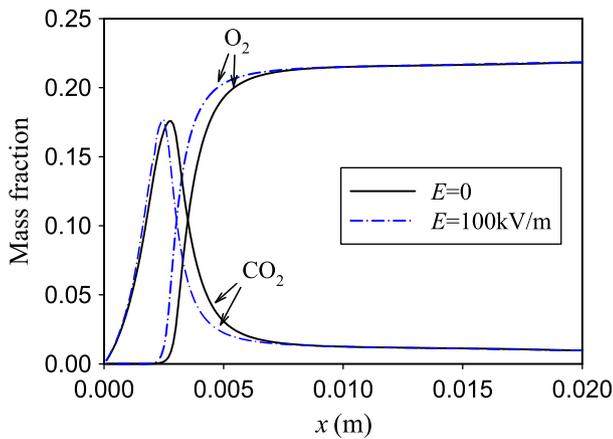


Fig. 9. Calculated mass fraction profiles of O₂ and CO₂ on the axis ($q_v = 1.2\text{ ml/h}$).

charged particles mix quickly with oxygen in the flame front [10] and the reaction took place in a wider space. Thus, the reaction rate increased resulting in the fuel burning faster and the inward contraction of the flame front to the fuel side. It means that the DC electric field can enhance the combustion process and in turn lead to a higher flame temperature.

4.4. Flame deformation in electric field

One important characteristic of laminar diffusion flame is the flame shape or structure. Fig. 11 shows the calculated temperature distribution of flame without electric field condition (left) and that under positive electric field (right) which also reflects the flame shape. Fuel flow rate was 1.2 ml/h , and the applied voltage was 4.0 kV ($E = 100\text{ kV/m}$). It shows that the flame became shorter and smaller after the electric field was applied. The flame height decreased by 50% with an applied electric field.

The maximum temperature with electric field appeared to be 1472 K which was slightly above (by 4.7%) the temperature with no electric field (1446 K). Owing to these aerodynamic effects associated with the electrical field, the entire flow field is affected, and in particular the flame shape, which is well known to be very sen-

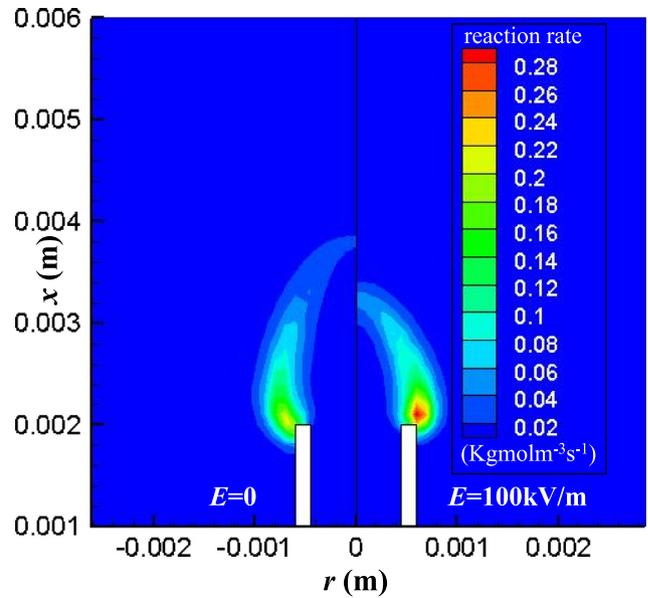


Fig. 10. Chemical reaction rate. Left: no electric field, right: $E = 100\text{ kV/m}$ ($q_v = 1.2\text{ ml/h}$).

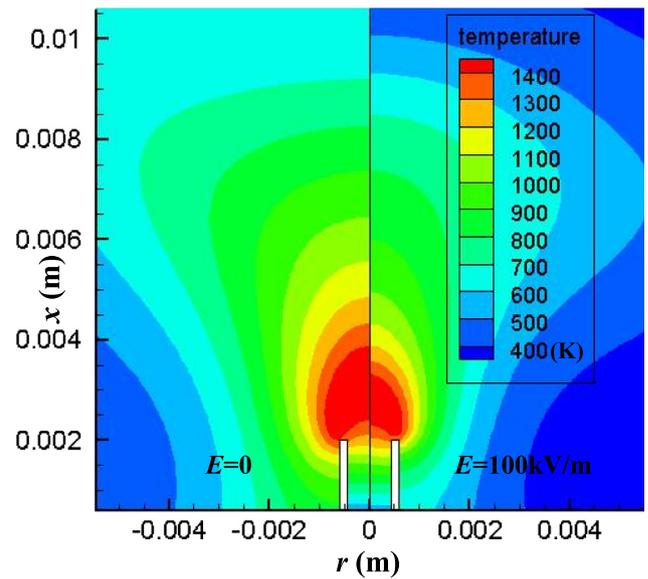


Fig. 11. Calculated temperature fields. Left: no electric field, right: $E = 100\text{ kV/m}$ ($q_v = 1.2\text{ ml/h}$).

sitive to the surrounding flow field [32], is influenced accordingly. Thus, the flame scale became smaller.

The aspect ratio was defined as the ratio of the flame height (H) to width (W)

$$\alpha = H/W \quad (8)$$

Fig. 12 shows the changes of flame deformation rate with the applied DC electric field strength obtained from the experimental results. The results show a satisfactory correlation with the best fit of the following equation with correlation coefficient $R = 0.9408$.

$$\alpha = \frac{0.8983 - 0.0047E}{1 - 0.002E} \quad (9)$$

It could be found that the flame deformation rate became smaller with the increasing electric field strength. When the electric field strength was stronger, the electric field force was greater and the flame became flatter.

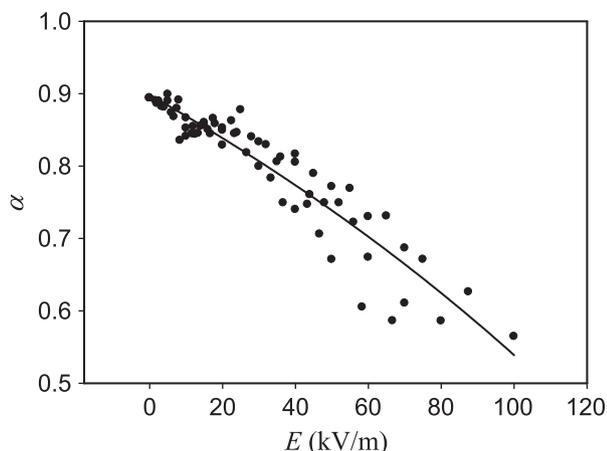


Fig. 12. The flame deformation rate with electric field strength collecting by experimental results ($q_v = 1.2$ ml/h).

The flame shape is affected by the surrounding flow field, reaction rate and diffusion. The height of the laminar diffusion flame is proportional to the fuel flow rate, and is inversely proportional to the diffusion coefficient D . As known from the Fick's Diffusion Law, D is nearly proportional to $T^{3/2}$. According to the results of flame temperatures and discussions above, the flame temperature with applied electric field is higher than that without electric field. So the diffusivity increased and the flame became shorter under the effect of DC electric field. The electric field increased the flow velocity by the electric force and promoted the reaction rate and the diffusion of the species which also made the flame length shorter.

5. Conclusions

The effects of DC electric field on the small ethanol diffusion flame were investigated. The flow velocity, chemical reaction rate, mass fraction distribution, flame temperature and deformation in the applied DC electric field were considered. The results show that the applied electric field changes the flame characteristics mainly due to the body forces acting on charged particles. The applied electric field accelerates the charged particles and they collide with the neutral particles and transfer the momentum, thus increasing the flow velocity. The effects on the species distribution and the flow field near the flame by the applied electric field promote the mixing of fuel and oxidant, which enhances the combustion process and leads to a higher flame temperature. Flame becomes shorter with applied electric field and its deformation is related to the electric field strength.

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