A computational study of positive streamers interacting with dielectrics

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Abstract

We use numerical simulations to study the dynamics of surface discharges, which are common in high-voltage engineering. We simulate positive streamer discharges that propagate towards a dielectric surface, attach to it, and then propagate over the surface. The simulations are performed in air with a two-dimensional plasma fluid model, in which a flat dielectric is placed between two plate electrodes. Electrostatic attraction is the main mechanism that causes streamers to grow towards the dielectric. Due to the net charge in the streamer head, the dielectric gets polarized, and the electric field between the streamer and the dielectric is increased. Compared to streamers in bulk gas, surface streamers have a smaller radius, a higher electric field, a higher electron density, and higher propagation velocity. A higher applied voltage leads to faster inception and faster propagation of the surface discharge. A higher dielectric permittivity leads to more rapid attachment of the streamer to the surface and a thinner surface streamer. Secondary emission coefficients are shown to play a modest role, which is due to relatively strong photoionization in air. In the simulations, a high electric field is present between the positive streamers and the dielectric surface. We show that the magnitude and decay of this field are affected by the positive ion mobility.

Keywords: surface discharge, positive streamer, plasma fluid model, secondary emission, photoemission, dielectric

(Some figures may appear in colour only in the online journal)

1. Introduction

Electric discharges in electronic devices and HV (high-voltage) equipment often occur along dielectric materials. In the regions of HV stress around an insulator, electron avalanches and streamer discharges can develop. These partial discharges may eventually result in surface flashover of the insulator, i.e., electric breakdown. In [1] it was found that around atmospheric pressure, surface flashover voltages were 10\%–50\% lower than flashover voltages in pure gas gaps. A dielectric present in the vicinity of the electrodes not only modifies the fields between the electrodes, but also serves as a possible source or sink of electrons during the breakdown process. Studying the interaction between dielectrics and streamer discharges is therefore important to understand surface flashover.

Early studies of surface discharges focused on the measurement of flashover voltage [2, 3]. In the past few decades, the use of high-speed cameras has revealed more details about the early stages of surface discharges. In several experiments, streamer discharges were observed to have an affinity to propagate along dielectric surfaces rather than through the background gas only [4, 5]. This affinity for a dielectric surface was reported to depend on the discharge gap geometry [6], gas composition, pressure [7], and dielectric properties [8, 9].
To gain more insight into the physics of surface discharges, different types of numerical simulations have been performed, see e.g. [7, 10–15]. Studies on the interaction between plasmas and dielectrics have often been performed at lower pressure and in noble gases, where the discharge mechanisms are relatively well understood [16, 17]. Several authors have also studied surface discharges in atmospheric air. An incomplete list is given below.

Jørgenson et al [10] investigated the role of photoemission in the surface breakdown process. With Monte Carlo simulations, they concluded that photoemission plays a role at low field values near the breakdown threshold. Celestin et al [18] studied dielectric barrier discharges in air both experimentally and computationally, and highlighted the importance of surface charge. Jánksý et al [11] presented simulations of an air plasma discharge at atmospheric pressure, initiated by a needle anode set inside a dielectric capillary tube. Meyer et al [19] studied surface streamers with a plasma fluid model in a 2D geometry. Agreement with empirical estimates for streamer lengths was found, and it was observed that the surface charge quickly reaches so-called ‘saturation charge’ conditions. Babaeva et al [12, 20] performed a computational investigation of nanosecond pulsed surface discharges of positive and negative polarity. A hybrid fluid-Monte Carlo model was used to more accurately capture secondary electron emission caused by positive ions and photons. Sima et al [15] presented 2D axisymmetric fluid simulations of discharges spreading radially over a dielectric surface in a N₂/O₂ mixture. Furthermore, several computational studies of plasma–liquid interaction and plasma–tissue interaction have been performed at atmospheric pressure, see for example [21, 22]. In such studies, the liquid or skin is often modelled as a dielectric, sometimes with a finite conductivity.

The studies mentioned above have greatly improved our understanding of surface discharges in air. Here, this past work is extended in several ways. We consider a different geometry, namely a flat dielectric placed between parallel-plate electrodes. Streamers in this geometry can propagate along the dielectric or through the gas, in contrast to surface dielectric barrier discharges (SDBD), in which electrodes are completely separated by a dielectric [12, 23, 24]. This allows us to investigate the whole streamer–dielectric interaction, including discharge inception, attachment to the dielectric and propagation over the surface. Moreover, this geometry resembles some actual HV insulation applications [9, 25], e.g., insulators inserted between HV and ground electrodes in gas-insulated switchgear. Our focus here is on positive surface streamers, which can be computationally expensive to simulate. We have therefore developed an efficient fluid model with adaptive mesh refinement. For simplicity and efficiency, 2D simulations are used here, as full 3D simulations would still be very costly.

The content of the paper is as follows. The simulation model is described in section 2. In section 3.1, we focus on the attraction of streamers to dielectrics, and we look at the differences between surface and gas-phase streamers. Afterwards, several discharge parameters are varied, to study their effect on the streamer’s inception time, propagation velocity and morphology:

- The applied voltage in section 3.2.
- The permittivity (ε) of the dielectric material in section 3.3.
- The secondary electron emission coefficients (for positive ions and photons) in section 3.4.
- The positive ion mobility in section 3.5.

2. Simulation model

The 2D fluid model used in this paper is based on Afivo-streamer [26, 27], which is an open-source plasma fluid code for streamer discharges that features adaptive mesh refinement (AMR), geometric multigrid methods for Poisson’s equation, and OpenMP parallelism. For a recent comparison of six streamer simulation codes, including Afivo-streamer, see [28]. We have made several modifications to be able to simulate surface streamers:

- Electrons, ions and photons can be absorbed by dielectric surfaces.
- Surface densities and fluxes are stored separately from their equivalents in the gas.
- The electric field computation takes the surface charge into account.
- A new Monte Carlo photoemission module was implemented, and the photoionization routines were adjusted to account for the dielectric.

These changes are described in more detail below.

With our 2D model, we effective simulate planar surface discharges. This leads to some differences compared to a full 3D description. First, the electric fields and charge densities in 3D are typically higher, as the streamer heads have a stronger curvature. Second, it is often observed that both surface and gas streamers are present in experiments [8, 29]. We do not observe these two components in our 2D model, but have seen them in preliminary 3D simulations that are still under development.

2.1. Fluid model

The fluid model used here is of the drift-diffusion-reaction type with the local field approximation [30]. The model keeps track of the electron density nₑ, the positive ion density nᵢ⁺ and the negative ion density nᵢ⁻, which involve in time as

\[
\frac{\partial n_e}{\partial t} = -\nabla \cdot \mathbf{\Gamma}_e + S_i - S_a + S_{pi} + S_{sc},
\]

(1)

\[
\mathbf{\Gamma}_e = -n_e \mu_e \mathbf{E} - D_e \nabla n_e,
\]

\[
\frac{\partial n_{i+}}{\partial t} = -\nabla \cdot \mathbf{\Gamma}_{i+} + S_i + S_{pi},
\]

(2)

\[
\mathbf{\Gamma}_{i+} = n_{i+} \mu_{i+} \mathbf{E},
\]

\[
\frac{\partial n_{i-}}{\partial t} = -\nabla \cdot \mathbf{\Gamma}_{i-} + S_i,
\]

(3)

\[
\mathbf{\Gamma}_{i-} = -n_{i-} \mu_{i-} \mathbf{E},
\]
Here, fluxes are indicated by a $\Gamma$, $\mu_e$ is the electron mobility, $D_e$ the electron diffusion coefficient, $E$ the electric field, and $\mu^+_i$ and $\mu^-_i$ the positive/negative ion mobilities. Furthermore, several source terms are present. The electron impact ionization and electron attachment terms are given by $S_i = \alpha e \mu_e |E|n_e$ and $S_a = \eta e \mu_e |E|n_e$, respectively, where $\alpha$ and $\eta$ are the ionization and attachment coefficients. The production of photoelectrons from photoionization is included with the term $S_{pi}$. Secondary electron emission due to the impact of both ions and photons is accounted for by the term $S_{se}$, defined below.

The local field approximation is used, so that $\mu_e$, $D_e$, $\alpha$ and $\eta$ are functions of the local electric field strength. Electron transport and reaction coefficients for air (1 bar, 300 K) were generated with Monte Carlo particle swarm simulations (see e.g. [31]), using Phelps’ cross sections [32]. The positive ion mobility $\mu^+_i = 3 \times 10^{-4} \text{ m}^2 \text{ V}^{-1} \text{ s}^{-1}$ is here considered to be constant, but in section 3.5 it is varied to investigate its effect on surface discharges. For simplicity, the negative ion mobility is set to zero ($\mu^-_i = 0$) throughout the paper.

We assume that electrons and ions attach to the surface when they flow onto a dielectric. They do not move or react on the surface, but secondary electron emission from the surface is taken into account. For the impact of positive ions, a SEE (secondary electron emission) coefficient $\gamma_i$ is used. When a photon hits a dielectric surface, we assume that the photon is absorbed. A SEE photoemission coefficient $\gamma_{pe}$ is used to determine the photoemission flux, see section 2.3. The effect of these SEE coefficients is studied in section 3.5, elsewhere they are set to zero. The secondary electron emission source term $S_{se}$ in equation (1) is non-zero only in cells adjacent to the dielectric surface. In these cells, it is given by

$$S_{se} = -\nabla \cdot (\tilde{\Gamma}_{pe} - \gamma_i \tilde{\Gamma}^+_i),$$

(4)

where $\tilde{\Gamma}^+_i$ is flux of positive ions onto the surface, and $\tilde{\Gamma}_{pe}$ is the photoemission flux coming from the surface. By definition, both these fluxes are non-zero only at the surface.

Secondary emission leaves behind positive surface charge on the dielectric. Therefore, the surface charge density $\sigma_s$ changes in time as

$$\partial_t \sigma_s = -e(\tilde{\Gamma}_e + \tilde{\Gamma}^-_i) + e(1 + \gamma_i)\tilde{\Gamma}^+_i + e\tilde{\Gamma}_{pe},$$

(5)

where $e$ is the elementary charge and the other terms correspond to the fluxes onto the dielectric surface: $\tilde{\Gamma}_e$ for electrons, $\tilde{\Gamma}^-_i$ for negative ions, $\tilde{\Gamma}^+_i$ for positive ions, and $\tilde{\Gamma}_{pe}$ for photons. We study positive streamers, which means that electrons generally move away from dielectrics. Therefore, an accurate description of the electron flux towards the surface [33] is not required here.

2.2. Electric field

The electric field $E$ is calculated by first solving Poisson’s equation for the electric potential $\phi$:

$$\nabla \cdot (\varepsilon \nabla \phi) = -(\rho + \delta_s \sigma_s),$$

(6)

where $\varepsilon$ is the dielectric permittivity, $\rho$ is the volume charge density, and $\delta_s$ maps the surface charge $\sigma_s$ on the gas–dielectric interface to the grid cells adjacent to the dielectric. Afterwards, the electric field is computed as

$$E = -\nabla \phi.$$  

(7)

At the dielectric interface, we ensure that the normal component of the electric field satisfies the classic jump condition

$$\varepsilon_2 E_2 - \varepsilon_1 E_1 = \sigma_s,$$  

(8)

where $\varepsilon_1$ and $\varepsilon_2$ denote the permittivities on both sides of the interface, and $E_1$ and $E_2$ the electric field components normal to the interface.

Details about the numerical implementation, which is compatible with adaptive mesh refinement, will be presented in a forthcoming paper.

2.3. Photoionization and photoemission

Positive streamer discharges need a source of free electrons ahead of them in order to propagate. Photons can generate such free electrons through photoionization in the gas or photoemission from a dielectric surface. In N\textsubscript{2}–O\textsubscript{2} mixtures, non-local photoionization can take place when an excited nitrogen molecule emits a UV photon in the 98 to 102.5 nm range, which has enough energy to ionize an oxygen molecule. Photoionization often plays an important role in electrical discharges, see e.g. [34, 35].

The role of photoemission in surface discharges is less well understood. Photons can be emitted from several excited states. The probability of photoemission not only depends on the photon energy, but also on the surface properties [10]. For simplicity, we consider only two types of photons in this paper: high-energy photons, which can generate photoionization and photoemission, and low-energy photons, which can only contribute to photoemission and are not absorbed in the gas. These processes are illustrated in figure 1.

Photoionization and photoemission are here modelled with a Monte Carlo (MC) method. We use the same Monte Carlo photoionization model as described in [36] and chapter 11 of
The idea is to approximate the photoionization source term $S_{\text{pi}}$ and the photon flux onto the dielectric $\Gamma_{\text{pe}}$ by randomly sampling discrete photons. For computational efficiency, these terms are updated every $\Delta t_r = 10\Delta t$, where $\Delta t$ is the time step used for solving equation (1).

First, the number of ionizing (high-energy) photons produced per grid cell during a time $\Delta t_r$ is determined. We use Zheleznyak’s model [38], in which the number of ionizing photons is proportional to the number of impact ionization events. The corresponding proportionality factor is set to $\xi p_h/(p + p_h)$, where $\xi = 0.05$ is a numerical factor, $p = 1$ bar is the gas pressure and $p_h = 40$ mbar is the collisional quenching pressure. We remark that $\xi$ should in principle depend on the electric field [38], but that it is here approximated by a constant, as in [36]. Per cell, a random number is drawn to determine how many photons are generated, see [37].

Since simulations are here performed in 2D, the discrete photons do not correspond to (single) physical photons. Instead, the total photon number $N_{\text{photons}}$ is fixed, so that the MC method always uses $10^5$ photons. The weight factor $w$ of these photons is given by

$$w = \int S, dV/N_{\text{photons}},$$

where

$$\int S, dV = \xi p_h/(p + p_h) \int S_0 dV,$$

is the volume-integrated production rate of ionizing photons. For simplicity, the number of produced low-energy photons is assumed to be equal to the number of high-energy photons. As the low-energy photons only contribute to photoemission, their effect can be controlled through the corresponding photoemission coefficient.

Second, an isotropically distributed direction is sampled for each photon. Afterwards, absorption lengths are determined. The absorption length of high-energy photons is sampled from the absorption function for air, see e.g. [36]. Low-energy photons are not absorbed by the gas, so their absorption length is set to a large value, making sure they always end up outside the computational domain.

Third, we determine which photons hit a dielectric surface, and where they do so. These photons are absorbed by the surface, where they contribute to the local photoemission flux $\Gamma_{\text{pe}}$. For the low-energy and high-energy photons, photoemission coefficients $\gamma_{\text{pel}}$ and $\gamma_{\text{peh}}$ are used, respectively. If a surface cell of area $\Delta A$ is hit by $n_i$ and $n_H$ low-energy and high-energy photons, then

$$\Gamma_{\text{pe}} = (\gamma_{\text{pel}} n_i + \gamma_{\text{peh}} n_H)w/\Delta A.$$  

(10)

The effect of photoemission is investigated in section 3.4; elsewhere in the paper photoemission is not taken into account (so that $\gamma_{\text{pel}} = \gamma_{\text{peh}} = 0$).

Fourth, the remaining high-energy photons that are absorbed in the gas contribute to photoionization source term $S_{\text{pi}}$. If $n_i$ photons are absorbed in a grid cell with volume $\Delta V$, then

$$S_{\text{pi}} = n_i w/\Delta V.$$  

(11)

Low-energy photons and high-energy photons that are absorbed outside the computational domain have no effect.

2.4. Computational domain and initial conditions

We use a parallel-plate electrode geometry with a flat dielectric in between, as shown in figure 2. The computational domain measures (40 mm)$^2$, and a dielectric is present on the left side with a width of 10 mm. The dielectric permittivity is set to $\varepsilon = 2$, but in section 3.3 it is varied to investigate its effect on surface discharges.

As a gas, artificial air (80% N$_2$ and 20% O$_2$) at 1 bar and 300 K is used. For the electric potential, Dirichlet boundary conditions are applied at the upper and lower boundaries, and Neumann zero boundary conditions on the left and right side. A voltage of 100 kV is applied. In section 3.2, this voltage is varied. The background densities of electrons and positive ions are set to $10^{10}$ m$^{-3}$ [39].

To start a discharge, the background field has to be locally enhanced. We do this by placing an ionized seed of about 2 mm long with a radius of about 0.4 mm. The electron and positive ion density are $5 \times 10^{19}$ m$^{-3}$ at the center, and they decay at distances above $d = 0.2$ mm with a so-called smoothstep profile: $1 - 3x^2 + 2x^3$ up to $x = 1$, where $x = (d - 0.2$ mm)/0.2 mm. When the electrons from a seed drift upwards, the electric field at the bottom of the seed is enhanced so that a positive streamer can form.

3. Results & discussion

In section 3.1, the initial seed is placed at different distances from the dielectric to study the streamer–dielectric interaction. We also point out the main differences between surface and gas streamers. Next, we systematically study the effect of several parameters on the surface discharges: the applied voltage (section 3.2), the dielectric permittivity (section 3.3), the secondary electron emission coefficients (section 3.4), and finally the ion mobility (section 3.5). The parameters investigated and their values in each section are shown in table 1.

3.1. Streamer–dielectric interaction

Previous experiments have revealed that dielectrics attract positive streamers, see e.g. [4, 5]. This attraction is also present in
Table 1. Investigated parameters and their values in each section. Here $d$ is the distance between seed center and the dielectric; $U$ the applied voltage; $\varepsilon_r$, the relative permittivity of the dielectric; $\gamma_i$, the ion-induced secondary electron emission coefficient; $\gamma_{pe}$, the photoemission coefficient and $\mu_i^+$, the positive ion mobility.

| Section | $d$ (mm) | $U$ (kV) | $\varepsilon_r$ | $\gamma_i$ | $\gamma_{pe}$ | $\mu_i^+$ (m$^2$ V$^{-1}$ s$^{-1}$) |
|---------|----------|----------|----------------|------------|---------------|----------------------------------|
| 3.1     | (0.5, 1, 2, 5) | 100      | 2              | 0          | 0             | $3 \times 10^{-4}$               |
| 3.2     | 0.5      | (92, 100, 112) | 2              | 0          | 0             | $3 \times 10^{-4}$               |
| 3.3     | 0.5      | 100      | (2, 3, 5)      | 0          | 0             | $3 \times 10^{-4}$               |
| 3.4.1   | 0.4      | 100      | 2              | (0, 0.5)   | 0             | $3 \times 10^{-4}$               |
| 3.4.2   | (0.5, 1) | 100      | 2              | 0          | (0, 0.5)      | $3 \times 10^{-4}$               |
| 3.5     | 0.5      | 100      | 2              | 0          | 0             | $(0, 1, 5, 10) \times 10^{-4}$   |

Figure 3. The streamer development process between 10 ns and 18 ns for seed placed at 1 mm from the dielectric. White equipotential lines spaced by 2 kV are shown in part of the domain.

Figure 4. The electron density for streamers starting from different locations. For panels (a)–(d), the initial seed was placed at 0.5 mm, 1 mm, 2 mm and 5 mm from the dielectric. For comparison, a streamer in bulk gas is shown in panel (e). Results are shown at 20 ns, except for panel (a), which has the fastest propagation. White equipotential lines spaced by 2 kV are shown in part of the domain.

our numerical model. Figure 3 shows the evolution of the electron density for an initial ionized seed placed 1 mm away from the dielectric. It can be seen that the streamer starts to grow in air and then gradually develops towards the dielectric. After connecting with the dielectric, the streamer propagates down over its surface. The evolution shown here is in qualitative agreement with the discharge photographs in [4].

To study the streamer–dielectric attraction, we have placed the initial ionized seed at different distances from the dielectric. Figures 4(a)–(d) show the electron density for seeds placed at 0.5 mm, 1 mm, 2 mm and 5 mm from the dielectric. For comparison, the electron density for a streamer in bulk gas is also shown in figure 4(e). In this simulation the dielectric was removed, so that the whole computational domain contained gas, and the initial seed was placed at $x = 20$ mm. Figure 4(a) shows the electron density at 15 ns, for the other cases, which develop more slowly, results at 20 ns are shown. The closer the streamer is located to the dielectric, the stronger the attraction to the dielectric becomes. It can also be seen that a nearby dielectric increases the streamer’s velocity in the gas, and that streamers here propagate faster on the surface than in the gas.

3.1.1. Attraction to the dielectric. As photoemission is disabled here (see table 1), the attraction of the streamer to the dielectric is purely electrostatic. The net charge in the streamer head polarizes the dielectric, which increases the electric field between the streamer and the dielectric. This effect is illustrated in figure 5, which shows the horizontal electric field ($E_x$) around the streamer heads. With a dielectric present, $|E_x|$ increases on the dielectric side, and it is reduced on the other side. The closer the streamer is to a dielectric, the stronger this effect becomes. Eventually, the streamer will turn into a surface streamer. As shown in figure 4, such a surface streamer is thinner and quite asymmetric compared to a gas streamer.

In past research, the attraction of streamers to dielectrics could often be explained by the enhanced static field between pointed electrodes and dielectrics, see e.g. [4, 6]. In contrast, the attraction to the dielectric is here due to the space charge from the streamer itself, since in our plate-to-plate geometry, the electric field has no horizontal component before a discharge is present. Assuming that streamers propagate approximately along electric field lines [40], we can therefore state that streamers lead themselves to the dielectric: their space charge modifies the background field and causes a horizontal field component that attracts them to the dielectric.

Surface charge on the dielectric could also play a role in attracting streamer discharges, by modifying the background electric field. However, there is negligible surface charge here, as secondary electron emission is disabled in this section and electrons move away from the dielectric. Positive ions do move towards the dielectric, but there are initially few of them near the dielectric, and they drift with a relatively low mobility, so that they hardly contribute to the surface charge.
3.1.2. Effect on streamer velocity. Several experimental studies have found that surface discharges are faster than bulk gas streamers [5, 8, 41]. Their increased velocity was attributed to increased ionization rates near the dielectric, accumulated negative charge and electron emission from the dielectric surface. Electron emission from the dielectric is not included here (it is in section 3.4), but we still find that the surface streamers are significantly faster. Figure 6 shows the streamer velocity and its maximal electric field for the case \( d = 1 \) mm. Note that both the velocity and the maximal electric field increase when the surface streamer forms, at around \( y = 35 \) mm. Even though the higher field is mostly in the horizontal direction, see figure 5, it still contributes to a faster vertical growth.

As can be seen in figure 4, the electron density inside a surface streamer (\( \approx 10^{20} \) to \( 10^{21} \) m\(^{-3} \)) is higher than in a gas streamer (\( \approx 10^{19} \) m\(^{-3} \)) when the discharge conditions are otherwise similar. This was also observed in (for example) [23]. We remark that besides the presence of a dielectric, the electron density inside a streamer discharge also depends on other factors, such as the applied voltage, the streamer radius, the gas composition, the amount of photoionization etc. Also note that we use a Cartesian 2D model, in which there is less field enhancement around the streamer heads than in full 3D, reducing the electron density in both surface and gas streamers.

There seem to be several related effects that lead to the increased surface streamer velocity. The strong electric field between a surface streamers and a dielectric pulls surface streamers towards the dielectric. This reduces their radius (as shown in figure 4), and results in an asymmetric streamer head shape, which also leads to stronger electric field enhancement. The result is that the ionization rate is increased, that the streamer has a higher degree of ionization, and that it propagates faster. This behavior is quite distinct from gas streamers, which typically propagate faster when they have a larger radius [42].

3.1.3. Cathode sheath. As shown in figure 7, the surface streamer ‘hovers’ over the dielectric surface without fully connecting to it. This phenomenon was also observed in simulations of dielectric barrier discharges [12, 43–45], and it only occurs for positive streamers. The reason is that positive streamers grow from incoming electron avalanches, but such avalanches require sufficient distance before they reach ionization levels comparable to the discharge. Positive streamers can therefore not immediately connect to the dielectric surface. Due to the net charge in the streamer head, a very high electric field is present in the narrow gap between streamer and dielectric. The effect of secondary electron emission on these dynamics is studied in section 3.4, and the role of the positive ion mobility is investigated in section 3.5.

We remark that the maximum electric field of a positive surface streamer can rapidly rise to very high values, as shown in figures 6(a) and 7. These high-field areas typically contain a low electron density, but they still pose a problem for plasma fluid simulations. In the present fluid simulations, the transport and reaction coefficients (e.g., the ionization coefficient or the electron mobility) are functions of the local electric field.
Figure 6. The maximum electric field and streamer velocity versus the vertical position of the streamer head.

Figure 7. The electron density and the electric field around the positive streamer head at 20 ns, for an initial seed at 1 mm from the dielectric. Note the gap between the streamer and the dielectric.

Figure 8. The maximum electric field versus time for streamers in background electric fields of 2.3 kVmm$^{-1}$, 2.5 kVmm$^{-1}$ and 2.8 kVmm$^{-1}$ between. The indicated stages are (I) initial stage, (II) gas propagation, (III) transition towards a surface streamer, (IV) stable surface propagation.

field strength. These coefficients are tabulated up to a certain maximum electric field, which is here 35 kVmm$^{-1}$; for higher fields, we use the tabulated value at 35 kVmm$^{-1}$. More generally, the validity of the local field approximation is questionable when there are such high electric fields (and corresponding strong gradients). For future studies in such ultra-high electric fields, particle-in-cell simulations could therefore be more suitable, as was also observed in [20]. Finally, we remark that a potential physical limitation for this maximum electric field is field emission of electrons from the surface.

3.2. Effect of applied voltage

To study the effect of the applied voltage on surface discharges, we have performed simulations for applied voltages of 92 kV, 100 kV and 112 kV, which correspond to background electric fields of 2.3 kVmm$^{-1}$, 2.5 kVmm$^{-1}$ and 2.8 kVmm$^{-1}$, respectively. In all cases, the initial seed was located at 0.5 mm from the dielectric, and the evolution up to 15 ns was simulated. Figure 8 shows the maximum electric field versus time, and Figure 9 shows the electron density for three cases at 7 ns and 9 ns. Both figures reveal the following stages in the streamer’s development:

(I) The inception stage, in which the maximum electric field is from 0 to about 9 kVmm$^{-1}$ in our setup and the streamer is hardly propagating, as shown in figure 9(a).

(II) The gas-propagation stage, in which streamers propagate in the gas with a maximum electric field below 12.5 kVmm$^{-1}$. This stage is visible in figures 9(a) and (b).

(III) The transition stage from a gas streamer to a surface streamer, in which the maximum electric field increases sharply. The streamer also loses its rounded head shape, as shown in figures 9(b) and (c).

(IV) The surface propagation stage. The growth of the maximum electric field is slowing down, and the streamer propagates along the dielectric, as shown in figure 9(c).

Figure 8 shows that when the voltage is changed, the streamers still exhibit similar behavior in these four stages. The main difference is that the inception stage becomes shorter. For background fields of 2.3 kVmm$^{-1}$, 2.5 kVmm$^{-1}$ and 2.8 kVmm$^{-1}$, the inception stages last 8.5 ns, 5.5 ns and 3.75 ns, respectively. The second stage also becomes slightly shorter for a higher applied voltage.
Figure 9. The electron density for streamers in different background electric fields (2.3 kV/mm, 2.5 kV/mm, and 2.8 kV/mm) at 7 ns and 9 ns.

Figure 10. Streamer velocities versus their vertical location for different background electric fields.

Figure 11. The streamers’ maximal electric fields versus time for dielectrics with relative permittivities $\varepsilon_r$ of 2, 3 and 5.

Figure 12. The streamers’ electron density for dielectrics with relative permittivities $\varepsilon_r$ of 2, 3 and 5. Results are shown at different times, at the moment when the streamer heads are at $y = 36$ mm.

3.3. Effect of dielectric permittivity

The relative permittivity $\varepsilon$ of dielectric materials varies over a wide range. To study how $\varepsilon$ affects surface discharges, we have performed simulations with $\varepsilon$ set to 2, 3 and 5. As before, the initial seed was placed at 0.5 mm from the dielectric, and simulations were performed up to 15 ns.

The maximum electric field versus time for the three permittivities is shown in Figure 11. The main difference we observe is that the second and third stages are shorter for a higher permittivity. A higher $\varepsilon$ means the dielectric polarizes more strongly, which leads to a stronger attraction of streamers to the dielectric. Streamers therefore start the surface propagation stage earlier, and their maximum electric field increases more rapidly. Note that their maximum electric field is also higher during the surface propagation stage. In summary, we can conclude that a higher permittivity leads to a faster transition into a surface streamer, and a higher maximum field during the surface propagation stage.

Figure 12 shows the electron density for these three cases when all the streamers are at $y = 36$ mm. It can clearly be seen that the streamers attach more rapidly to the dielectric with a higher permittivity. Notice also that the surface streamer’s radius is smaller with a higher dielectric permittivity, a result of the stronger electrostatic attraction.

Figure 13 shows the velocity versus the streamer’s vertical position for the different $\varepsilon_r$. The permittivity has only a small effect on the streamer’s velocity, in agreement with the experimental observations of [6]. In contrast, a negative correlation between the permittivity and the streamer velocity was found in [9]. The discrepancy could come from the different geometry that was used, in which multiple surface and gas streamers propagated next to a cylindrical dielectric.

3.4. Effect of secondary electron emission from dielectrics

Electron emission from dielectrics may influence streamer velocities [46] and affect the high electric field in the dielectric–plasma gap [47]. In this section, we study how secondary electron emission affects surface streamers in our computational geometry. Both ion-induced secondary emission (ISEE) and photo-emission are considered.
3.4.1. Ion-induced secondary electron emission. The ion-induced secondary electron emission (ISEE) yield $\gamma_1$ can vary over a wide range [48–50]. Here we consider two cases, $\gamma_1 = 0.5$ and $\gamma_1 = 0$ (i.e., no secondary emission). In this section, the initial ionized seed’s center is placed 0.4 mm away from the dielectric, so that streamers start directly next to the dielectric. Figure 14 shows the electron density and electric field distribution at 15 ns for both ISEE coefficients. It can be seen that ISEE here has little effect on the streamer length and the electric field. The electron density in the streamer–dielectric gap is slightly higher behind the head for the streamer with $\gamma_1 = 0.5$, but this has negligible influence on the electric field in the gap. We can conclude that the ISEE yield hardly affects the streamer head and its propagation (the latter is mainly determined by the former). The reason for this is that positive ions have a much lower mobility than electrons. Most positive ions are generated close to the streamer head, and they will not reach the dielectric in time to affect the rapidly propagating streamer head. Most ISEE electrons are therefore released after the streamer has passed by.

3.4.2. Photoemission. The photo-emission coefficient $\gamma_{pe}$ for typical dielectric materials varies between $10^{-4}$ to $10^{-1}$ for photon energies of 5–20 eV [51, 52]. This yield can be higher if the material contains stains or defects, or when it is negatively charged [10, 48]. The measurement of $\gamma_{pe}$ of dielectrics in air is often quite challenging [53]. We here use several values for $\gamma_{pe}$ to demonstrate how photoemission affects positive streamers.

As discussed in section 2.3, we consider low-energy and high-energy photons, with the main distinction that high-energy photons can be absorbed in the gas. The following four cases are considered for the photoemission coefficients $\gamma_{pel}$ and $\gamma_{peh}$ for low-energy and high-energy photons:

(a) $\gamma_{pel} = 0$, $\gamma_{peh} = 0$.
(b) $\gamma_{pel} = 0.5$, $\gamma_{peh} = 0$.
(c) $\gamma_{pel} = 0$, $\gamma_{peh} = 0.5$.
(d) $\gamma_{pel} = 0.5$, $\gamma_{peh} = 0.5$.

In the simulations, streamers start near the dielectric, with the seed placed at either 0.5 mm or 1 mm from the dielectric.

As shown in figure 15, photoemission by low-energy photons helps to start a discharge near a dielectric. At 5.5 ns, the $\gamma_{pel} = 0.5$ cases show the streamer already bending towards the dielectric with a sharp tip, due to photoemission. However, we remark that the inception time for these four cases is the same when the seeds are placed at 1 mm away from the dielectric. The secondary electrons from the dielectric then need more time to reach the streamer, and the streamers have already started due to the photoionization they generate. We conclude that photoemission can be important for discharges close to dielectrics and for discharges in gases with less photoionization than air.

Figure 16(a) shows the electron density distribution for the above four cases at 11 ns. The streamers with $\gamma_{pel} = 0.5$ are longer than the other two, since they start earlier. Photoemission also causes them to attach to the dielectric more rapidly. Another difference is that the narrow gap between streamer and dielectric is smaller with more photoemission. This happens because photoemission provides seed electrons in the gap, which allows the streamer to get closer to the dielectric. To see this more clearly, the electron density distributions at $y = 36.4$ mm [the dashed line in figure 16(a)] are shown for these four cases in figure 16(b). Without photoemission, the electron density has a wider profile with a lower maximum, and it is located farther from the dielectric. When photoemission is included, the effect of the low-energy photons (i.e., $\gamma_{pel} = 0.5$) is most important here.

Figure 17 shows the streamer velocities versus their vertical position for all four cases. The photoemission coefficients have only a small effect on the velocity, which is a little higher with $\gamma_{pel} = 0.5$. We think this is somewhat surprising. A possible explanation is that photoemission mostly leads to growth towards the dielectric, whereas photoionization in the gas contributes most of the free electrons that cause growth parallel to the dielectric. Another effect in the simulations presented here is that high-energy photons contribute less to a streamer’s growth very close to a dielectric. There are two reasons for this. First, these photons are absorbed at shorter distances if they hit a dielectric. Second, their photoemission coefficient is here

Figure 13. Streamer velocities versus their vertical location for different relative dielectric permittivities.

Figure 14. The electron density and electric field at 15 ns with the ion-induced secondary electron emission coefficient $\gamma_1$ set to 0 (left) and 0.5 (right).
3.5. Effect of positive ion mobility

The positive ion mobility \( \mu_i^+ \) can affect surface streamers in two ways. First, a higher ion mobility increases the amount of ion-induced secondary electron emission (ISEE). However, since ISEE was found to play a negligible role in section 3.4.1, its role is not further studied here, and we set the ISEE yield to zero (i.e., \( \gamma_i = 0 \)).

A second effect is that a higher ion mobility increases the conductivity of the discharge, in particular in regions where the ion density is high compared to the electron density. For positive surface streamer discharges, such a region is present in the streamer–dielectric gap. This gap typically contains a high electric field, especially close to the streamer head, see section 3.1.3. Electrons rapidly drift away from the surface, whereas positive ions move from the high-density discharge region towards the surface, as illustrated in figure 18(a).

To investigate how the positive ion mobility (\( \mu_i^+ \)) affects the decay of the high electric field in the streamer–dielectric gap, we have performed simulations with positive ion mobilities of 0, \( 1 \times 10^{-4} \text{ m}^2 \text{ V}^{-1} \text{ s}^{-1} \), \( 5 \times 10^{-4} \text{ m}^2 \text{ V}^{-1} \text{ s}^{-1} \) and \( 1 \times 10^{-3} \text{ m}^2 \text{ V}^{-1} \text{ s}^{-1} \), using a seed placed 0.5 mm from the dielectric. For these simulations, we have recorded \( E_x \) in the middle of the gap at the point indicated in figure 18(a). The recorded fields are shown versus time in figure 18(b). The maximum electric field occurs when the streamer heads pass by the observation point indicated in figure 18(a). The decay of the peak in \( E_x \) is faster for higher \( \mu_i^+ \), which is
most clearly visible for the $\mu_1^+ = 5 \times 10^{-4} \text{ m}^2 \text{ V}^{-1} \text{ s}^{-1}$ and $\mu_1^+ = 1 \times 10^{-3} \text{ m}^2 \text{ V}^{-1} \text{ s}^{-1}$ cases. Note that the field also decays when the ions are immobile. This mainly happens because the amount of net space charge is lower behind the streamer head, but electron avalanches in the gap (due to e.g. photoionization) also contribute.

4. Conclusions

In this paper, we have studied positive surface streamers with numerical simulations. A 2D fluid model for surface discharges based on the Afivo-streamer code [26] was developed. The model includes a Monte Carlo procedure for secondary electron emission (from both high and low energy photons) and support for dielectric surfaces. These new features are compatible with the adaptive mesh refinement and the parallel multigrid solver provided by the underlying Afivo framework [27].

We have used the new model to investigate the interaction of positive streamers and dielectrics. We considered a parallel-plate geometry, with a flat dielectric between the two electrodes. Positive streamer discharges started from an ionized seed that was placed near the dielectric and the positive electrode. The effect of several parameters was investigated: the applied voltage, the dielectric permittivity, secondary electron emission caused by ions and photons, and the mobility of positive ions. Our main findings are summarized below:

(a) A narrow gap forms between positive streamers and dielectrics, as was also observed in earlier work [12, 44, 45]. A very high electric field can be present in this so-called ‘cathode sheath’.

(b) The attraction of positive streamers to the dielectric was found to be mostly electrostatic. In our geometry, this attraction was caused by the net charge in the streamer head, which polarized the dielectric, increasing the field between the streamer and the dielectric. A higher dielectric permittivity led to a more rapid attachment of the streamer to the dielectric.

(c) Compared to gas streamers, surface streamers had a smaller radius, a higher electric field, and a higher electron density. In our simulations, this gave surface streamers a higher propagation velocity than gas streamers.

(d) A higher applied voltage caused the positive surface discharges to start earlier, but they behaved qualitatively similar. A higher dielectric permittivity also accelerated the formation of surface streamers.

(e) Photoemission can accelerate streamer inception near dielectrics. However, photoemission hardly increases the velocity of surface streamers. A possible reason is that photoemission mostly leads to growth towards the surface, whereas photoionization contributes more to the growth parallel to the surface.

(f) The positive ion mobility affects the decay of the high electric field in the streamer–dielectric gap.

Availability of model and data

The source code and input files for the model used in this paper are available at https://gitlab.com/MD-CWI-NL/afivostreamer (git commit 08f1c828).

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References

[1] Cookson A H 1970 Proc. Inst. Electr. Eng. 117 269
[2] Cookson A H 1981 IEE Proc. A Sci. Meas. Technol. 128 303–12
[3] Sudarshan T S and Dougali R A 1986 IEEE Trans. Electr. Insul. EI-21 727–46
[4] Sobota A, van Veldhuizen E M and Stoffels W W 2008 IEEE Trans. Plasma Sci. 36 912–3
[5] Trienekens D J M, Nijdam S and Ebert U 2014 IEEE Trans. Plasma Sci. 42 2400–1
[6] Sobota A, Lebouvier A, Kramer N J, van Veldhuizen E M, Stoffels W W, Manders F and Haverlag M 2009 J. Phys. D: Appl. Phys. 42 015211
[7] Dubinova A, Trienekens D, Ebert U, Nijdam S and Christen T 2016 Plasma Sources Sci. Technol. 25 055021
[8] Allen N L and Mikropoulos P N 1999 IEEE Trans. Dielectr. Electr. Insul. 6 357–62
[9] Meng X, Mei H, Chen C, Wang L, Guan Z and Zhou J 2015 IEEE Trans. Dielectr. Electr. Insul. 22 1193–203
[10] Jorgenson R E, Warne L K, Neuber A A, Kriile J, Dickens J and Krompholz H 2003 Effect of Dielectric Photo-emission on Surface Breakdown: An LDRD Report Tech. Rep. SAND2003-1731, 811483 US Department of Energy DOI:10.2172/811483
[11] Jánisky J, Tholin F, Bonaventura Z and Bourdon A 2010 J. Phys. D: Appl. Phys. 43 395201
[12] Babaeva N Y, Tereshonok D V and Naidis G V 2016 Plasma Sources Sci. Technol. 25 044008
[13] Georgiou G E, Papadakis A P, Morrow R and Metaxas A C 2005 J. Phys. D: Appl. Phys. 38 R303–28
[14] Sun G Y, Guo B H, Song B P, Su G Q, Mu H B and Zhang G-J 2018 Phys. Plasmas 25 063502
[15] Sima W, Liu C, Yang M, Shao Q, Xu H and Liu S 2016 Phys. Plasmas 23 063508
[16] Zhang G, Su G, Song B and Mu H 2018 IEEE Trans. Dielectr. Electr. Insul. 25 2321–39
[17] Sun A, Becker M M and Loffhagen D 2018 Plasma Sources Sci. Technol. 27 054002
[18] Celestin S, Bonaventura Z, Guaitella O, Rousseau A and Bourdon A 2009 Eur. Phys. J. Appl. Phys. 47 22810
[19] Meyer H K, Mauseth F, Marskar R, Pedersen A and Blassczyk A 2019 IEEE Trans. Dielectr. Electr. Insul. 26 1163–71
[20] Babaeva N Y 2015 Plasma Sources Sci. Technol. 24 034012
[21] Babaeva N Y and Kushner M J 2013 J. Phys. D: Appl. Phys. 46 025401
[22] Tian W and Kushner M J 2014 J. Phys. D: Appl. Phys. 47 165201
[23] Hua W and Fukagata K 2019 Phys. Plasmas 26 013514
[24] Brandenburg R 2017 Plasma Sources Sci. Technol. 26 055001
[25] Sima W, Liu C, Yang M, Xu H and Shao Q 2017 IEEE Trans. Plasma Sci. 45 906–12
[26] Teunissen J and Ebert U 2017 J. Phys. D: Appl. Phys. 50 474001
[27] Teunissen J and Ebert U 2018 Comput. Phys. Commun. 233 156–66
[28] Bagheri B et al 2018 Plasma Sources Sci. Technol. 27 095002
[29] Meng X, Mei H, Wang L, Guan Z and Zhou J 2017 IEEE Trans. Dielectr. Electr. Insul. 24 391–400
[30] Luque A and Ebert U 2012 J. Comput. Phys. 231 904–18
[31] Rabie M and Franck C 2016 Comput. Phys. Commun. 203 268–77
[32] Phelps A V and Pitchford L C 1985 Phys. Rev. A 31 2932–49
[33] Hagelaar G J M, de Hoog F J and Kroesen G M W 2000 Phys. Rev. E 62 1452–4
[34] Pancheshnyi S 2005 Plasma Sources Sci. Technol. 14 645–53
[35] Pancheshnyi S 2014 Plasma Sources Sci. Technol. 24 015023
[36] Bagheri B and Teunissen J 2019 Plasma Sources Sci. Technol. 28 045013
[37] Teunissen J 2015 PhD Thesis Department of Applied Physics Thesis https://research.tue.nl/en/publications/3-d-simulations-and-analysis-of-pulsed-discharges
[38] Zheleznyak M, Mnatsakanyan A and Sizykh S 1982 High Temp. 20 357–62
[39] Sun A, Teunissen J and Ebert U 2014 J. Phys. D: Appl. Phys. 47 445205
[40] Nijdam S, Takahashi E, Teunissen J and Ebert U 2014 New J. Phys. 16 103038
[41] Akyuz M, Gao L, Cooray V, Gustavsson T G, Gubanski S M and Larsson A 2001 IEEE Trans. Dielectr. Electr. Insul. 8 902–10
[42] Briels T M P, Kos J, Winands G J J, van Veldhuizen E M and Ebert U 2008 J. Phys. D: Appl. Phys. 41 234004
[43] Babaeva N Y and Kushner M J 2011 Plasma Sources Sci. Technol. 20 035017
[44] Stepanyan S A, Soloviev V R and Starikovskaia S M 2014 J. Phys. D: Appl. Phys. 47 485201
[45] Soloviev V and Krivtsov V 2017 J. Phys.: Conf. Ser. 902 10
[46] h Tan B, J Allen N and Rodrigo H 2007 IEEE Trans. Dielectr. Electr. Insul. 14 111–8
[47] Campanell M D and Umansky M V 2016 Phys. Rev. Lett. 116 085003
[48] Motoyama Y and Sato F 2006 IEEE Trans. Plasma Sci. 34 336–42
[49] Motoyama Y, Hirano Y, Ishii K, Murakami Y and Sato F 2004 J. Appl. Phys. 95 8419–24
[50] Tschiersch R, Nemschokmichal S, Bogaczyk M and Meichsner J 2017 J. Phys. D: Appl. Phys. 50 105207
[51] Fujihiara M and Inokuchi H 1992 Chem. Phys. Lett. 17 554–6
[52] Buzulutskov A, Breskin A and Chechik R 1997 J. Appl. Phys. 81 466–79
[53] Dubinova A A 2016 PhD Thesis TU Eindhoven Eindhoven et al https://research.tue.nl/en/publications/modeling-of-streamer-discharges-near-dielectrics