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Key Points:

- Results of fluorescence emissions from air agree with laboratory observations
- Distribution of electrons in the full energy range for two TGF mechanisms
- Model-based quantifications of optical emissions generated during TGF production

Correspondence to:

W. Xu, wxx5015@psu.edu

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Optical emissions associated with terrestrial gamma ray flashes

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Wei Xu¹, Sebastien Celestin², and Victor P. Pasko¹

¹Communications and Space Sciences Laboratory, Department of Electrical Engineering, Pennsylvania State University, University Park, Pennsylvania, USA, ²Laboratory of Physics and Chemistry of the Environment and Space, University of Orleans, CNRS, Orleans, France

Abstract Terrestrial gamma ray flashes (TGFs) are high-energy photon bursts produced by high-energy electrons originating in the Earth's atmosphere through bremsstrahlung processes. In this paper, we present modeling studies on optical emissions resulting from the excitation of air molecules produced by the large population of electrons involved in TGF events based on two possible production mechanisms: relativistic runaway electron avalanches (RREAs) and acceleration of thermal runaway electrons produced by high-potential intracloud lightning leaders. Numerical models developed in this study are first validated through the calculation of fluorescence emissions from air excited by energetic electrons and comparison with available laboratory observations. Detailed discussion of the role of excitation and ionization collisions on the formation of the electron energy distribution is presented. Moreover, using Monte Carlo simulations, we show that electron energy distributions established from the two TGF production mechanisms considered here are inherently different over the full energy range. The strong energy dependence of the capability of electrons to generate excited states responsible for optical emissions from neutral and ionized nitrogen molecules leads to intrinsic differences in optical emissions produced by different mechanisms of TGF production. We also show that TGFs are most likely accompanied by detectable levels of optical emissions and that the distinct optical features are of significant interest for constraining and validating current TGF production models.

1. Introduction

Brief and intense photon bursts that originate from the Earth's atmosphere, known as terrestrial gamma ray flashes (TGFs), were first discovered in 1994 by *Fishman et al.* [1994] using the Burst and Transient Source Experiment detector aboard the Compton Gamma-Ray Observatory. After their discovery, TGFs have been detected by the Reuven Ramaty High Energy Solar Spectroscopic Imager (RHESSI) [*Smith et al.*, 2005], the Fermi Gamma ray Space Telescope [*Briggs et al.*, 2010], and the Astrorivelatore Gamma a Immagini Leggero satellite [*Marisaldi et al.*, 2010]. Spaceborne measurements indicate that TGFs typically consist of single or multiple pulses, last from a few tens of microseconds to a few milliseconds [e.g., *Fishman et al.*, 2011], and exhibit energy spectra that may extend up to 100 MeV [e.g., *Tavani et al.*, 2011].

In addition to satellite observations, radio signals radiated by TGF-associated lightning flashes have been used to infer the conditions that are required for the production of this high-energy phenomenon. Observational evidence shows that TGFs are closely associated with the initial development stages of normal polarity intracloud lightning that transports negative charge upward (+IC) [e.g., *Stanley et al.*, 2006; *Shao et al.*, 2010; *Lu et al.*, 2010]. Moreover, *Marshall et al.* [2013] have performed dedicated analyses on the initial breakdown (IB) stage of 10 IC flashes that emitted radio signals similar to TGF-producing ones and speculated that avalanches of relativistic runaway electrons caused IB pulses.

Two main mechanisms have been proposed and developed for interpreting TGF observations. The first mechanism is that of relativistic runaway electron avalanches (RREAs), which involves acceleration and multiplication of relativistic seed electrons (e.g., cosmic ray secondary electrons) under the application of large spatial scale electric fields within thunderstorms [*Gurevich et al.*, 1992; *Dwyer and Smith*, 2005]. Since natural background radiation or extensive air showers have been shown to provide insufficient number of seed electrons for the RREA theory to explain the TGF fluences observed from satellites, relativistic feedback mechanisms of positrons and X-rays have been invoked [*Dwyer*, 2008]. The other proposed mechanism is based on production of thermal runaway electrons [*Gurevich*, 1961] by the negative corona flash stages of stepping lightning leaders [*Moss et al.*, 2006; *Celestin and Pasko*, 2011]. Modeling studies by *Xu et al.*



Figure 1. Illustration of optical emissions produced by two TGF production mechanisms: RREAs in the large-scale homogeneous electric field within thunderstorms and the acceleration of thermal runaway electrons under the highly inhomogeneous electric field produced around the tip region of +IC lightning leaders during the negative corona flash stage.

[2012] and *Celestin et al.* [2012] support this mechanism by demonstrating that further acceleration of these thermal runaway electrons in the electric field produced near the tip region of long unbranched +IC lightning leaders leads to results consistent with satellite measurements of TGF spectra. The two production mechanisms, including the associated optical emissions and the fan-shaped streamer zone, in front of the lightning leader, are illustrated in Figure 1.

Although photon energy spectra established from both mechanisms show good agreement with satellite measurements [e.g., *Dwyer and Smith*, 2005; *Xu et al.*, 2012; *Celestin et al.*, 2012], there are notable differences between these two mechanisms. Lightning flashes do not play a critical role in the RREA theory and the high-energy parts of the electron distribution and corresponding photon spectrum are weakly dependent on the magnitude of the homogeneous electric field driving the RREAs [e.g., *Babich et al.*, 2004; *Dwyer and Smith*, 2005], whereas the energy spectrum of gamma rays emitted during the acceleration of thermal runaway electrons in the highly inhomogeneous electric fields around lightning leader tips is influenced by the physical properties and spatial electric field configurations associated with lightning stepped leaders [*Xu et al.*, 2012; *Celestin et al.*, 2012].

In spite of numerous experimental and theoretical efforts, little observational evidence is available about the TGF source. One of the most important unknowns is whether and which measurable optical emissions are generated during the production of TGFs. Optical emissions usually provide insightful knowledge about the energetics of electrons and the driving electric field in atmospheric discharges [e.g., Celestin and Pasko, 2010a] and are, therefore, extensively used in the studies of thunderstorm-related Transient Luminous Events (TLEs) at high altitudes in the Earth's atmosphere [e.g., Kuo et al., 2005, 2009; Liu et al., 2006; Adachi et al., 2006]. Recently, Østgaard et al. [2013] have reported optical emissions, detected for the first time, by the Lightning Imaging Sensor on the Tropical Rainfall Measuring Mission, from TGF-associated IC flashes. It is also worth mentioning that, based on high-speed camera observations, it has been suggested in Stolzenburg et al. [2013] that the impulsive breakdown associated with initial leaders during IB stages of cloud-to-ground (CG) and IC flashes can generate considerable amount of visible light. Furthermore, Dwyer et al. [2013] quantified optical emissions produced by relativistic feedback discharges based on the RREA mechanism. These studies motivate the present work that aims at the development of an experimentally testable framework that would allow to distinguish between different production mechanisms of TGF events by quantifying the optical emissions generated by the large population of high- and low-energy electrons.

To achieve this goal, we first introduce a full energy range relativistic Monte Carlo model to study the dynamics of electrons and the characteristic electron energy distributions in both TGF production mechanisms (section 2.1) and another numerical model for quantifying the associated optical emissions

(sections 2.2 and 2.3). In section 3.1, we validate our models by comparing results of fluorescence emissions excited by energetic electrons with available laboratory observations. Then, we present simulation results of electron energy distributions in the full energy range obtained in the cases of homogeneous electric fields (section 3.2.1) and a strongly inhomogeneous electric field produced by high-potential lightning leaders (section 3.2.2). We quantify optical emissions that can be possibly generated during production of TGFs in section 3.3. We further analyze in section 4.1 the concept of fluorescence efficiency in the context of TGFs and discuss the scaling of optical emissions with air density/altitude in the Earth's atmosphere in section 4.2. Finally, we discuss the implications of these calculations for understanding the mechanisms behind TGFs in section 4.3.

2. Model Formulation

2.1. Monte Carlo Model for Electrons

The Monte Carlo model used in the present study to simulate the propagation and collisions of electrons in air is similar to that described in *Celestin and Pasko* [2011]. This model is three-dimensional (3-D) in the velocity space, 3-D in the configuration space, relativistic, and simulates electrons from sub-eV to GeV. The elastic scattering of high-energy (>500 eV) electrons by nitrogen and oxygen molecules is modeled using a shielded-Coulomb cross section, similar to that described in *Dwyer* [2007]. The singly differential cross sections of N₂ and O₂ are calculated over the full range of energy using the relativistic binaryencounter-bethe (RBEB) model [*Kim et al.*, 2000; *Celestin and Pasko*, 2010b]. In particular, this model enables orbital description of the differential cross section of ionization collisions for all target electrons. The energies of secondary electrons after ionization collisions are obtained through the knowledge of this differential cross section, and the scattering angles of primary and secondary electrons are determined from the relativistic equations of conservation of momentum and energy considering that the newly formed ion is static.

The driving electric field is a critical parameter in the present study as it directly controls the avalanche processes of runaway electrons. In the case of TGF production by thermal runaway electrons, the method of moments [*Balanis*, 1989, p. 670] is first used in order to calculate the electric field in the vicinity of a +IC lightning negative leader tip during the corona flash associated with the stepping of the leader (see *Celestin and Pasko* [2011], and references therein for discussion of the related phenomenology). The electric potential of the lightning leader tip with respect to the ambient potential can be approximated by $U_1 = E_0 I/2$ [*Bazelyan and Raizer*, 2000, p. 54], where E_0 is magnitude of ambient electric fields and *I* is the length of an unbranched leader channel. A representative electric potential that provides consistent results with RHESSI cumulative energy spectra has been suggested to be greater than 100 MV [*Xu et al.*, 2012]. For such high potential, the ambient large-scale thunderstorm electric field is assumed to be $E_0 = 5 \times 10^4$ V/m [e.g., *Marshall et al.*, 2001], the radius of the leader channel is chosen as 1 cm [*Rakov and Uman*, 2003, section 4.4.6, p. 134], and the IC lightning length is I = 4 km. Note that it has recently been shown in *Pasko* [2014] that, using electrostatic modeling, such a long unbranched leader can also explain magnitudes of the dipole moment changes and current moment changes believed to be associated with TGFs.

We study the dynamics of electrons in the full energy range in either large-scale homogeneous electric field producing RREAs or highly inhomogeneous electric field produced around the lightning leader tip region during negative corona flash stages. Most importantly, by keeping track of the acceleration of runaway electrons and the rapid generation of low-energy secondary electrons in both mechanisms, we focus on providing a rigorous description of the electron energy distribution in the full energy range.

Monte Carlo simulation encounters difficulty when modeled electrons cover many decades in the energy space, for instance, from sub-eV to GeV, for mainly two reasons. First, the copious amount of low-energy secondary electrons generated via ionization collisions of high-energy electrons severely increases the complexity of the simulation. Second, the simulation time that is required for the ensemble of electrons to achieve a statistical steady state is dramatically prolonged because of the high collision rate of low-energy electrons. In order to overcome these difficulties, the calculation of electron energy distribution is divided into two parts: a low-energy part (from sub-eV to 10 keV) and a high-energy part (from 10 keV to GeV). After steady state is reached in both parts, distribution function of electrons over the full range of energy is then obtained by normalizing the high-energy part calculation in order to connect to the low-energy part at 10 keV. This technique relies on adopting two characteristic time steps to maintain the collisional frequencies of high-energy and low-energy electrons separately. It is also supported by the similarity of

the low-energy parts of energy distributions for electron swarms initiated by either low-energy electrons or high-energy runaway electrons [*Colman et al.*, 2010]. This technique not only allows precise descriptions for both high- and low-energy electrons, but also greatly shortens the computation time. We have carefully validated this technique by comparing the results with full energy range simulations for representative test cases.

2.2. Optical Emissions Model

The main energy loss for runaway electrons occurs through ionization collisions with air molecules, resulting in low-energy secondary electrons. These primary and secondary electrons are able to generate excited species via impact excitation, and optical emissions can be generated by the radiative relaxation of these excited species. The model employed for evaluating optical emissions is similar to that documented by *Liu* and Pasko [2004]. Modeling optical emissions from the first positive band systems of N₂ ($B^3\Pi_g \rightarrow A^3\Sigma_u^+$, 1PN₂) is complex, particularly for cases at high air pressures with continuous excitation by electrons. The populating of N₂($B^3\Pi_g$) involves processes of direct impact excitation by electrons, cascading from higher electronic states of N₂, intersystem collisional transfer with adjacent states [e.g., *Morrill and Benesch*, 1996], and energy-pooling reactions [e.g., *Piper*, 1988]. Besides, the deexcitation of N₂($B^3\Pi_g$) by N₂ molecules is strongly vibrational-level-dependent [e.g., *Piper*, 1988; *Morrill and Benesch*, 1996]. Given that the focus of the present study is the fluorescence light produced by TGF sources at low altitudes where N₂($B^3\Pi_g$) is heavily quenched and 1PN₂ emissions are weak, in the present work, we mainly focus on optical emissions originating from the second positive band system of N₂ ($C^3\Pi_u \rightarrow B^3\Pi_g$, 2PN₂) and the first negative band system of N₂⁺ ($B^2\Sigma_u^+ \rightarrow X^2\Sigma_a^+$, 1NN₂⁺).

The intensity of optical emissions in Rayleighs is given by the following expression [*Chamberlain*, 1978, p. 213]:

$$I_{k} = 10^{-6} \int_{L'} A_{k} n_{k} dl$$
 (1)

where n_k [1/cm³] is the number density of excited species in state k, A_k is the radiation transition rate and the typical values for optical emissions from 2PN₂ and 1NN₂⁺ are, respectively, 2 × 10⁷ s⁻¹ and 1.4 × 10⁷ s⁻¹ [*Vallance Jones*, 1974, p. 119]. The integral is taken along L' (cm), representing the horizontal line of sight.

The quantity n_k is governed by processes including direct excitation by electrons, cascading from higher-energy electronic states, and collisional quenching by air molecules. The number density n_k can be calculated using the following relation [*Sipler and Biondi*, 1972]:

$$\frac{\partial n_k}{\partial t} = -\frac{n_k}{\tau_k} + \sum_m n_m A_m + v_k n_e \tag{2}$$

where $\tau_k = [A_k + \alpha_1 N_{N_2} + \alpha_2 N_{O_2}]^{-1}$ is the total lifetime of state k, α_1 and α_2 are the quenching rate coefficients due to collisions with N₂ and O₂ molecules, respectively, N_{N_2} and N_{O_2} are, respectively, the number densities of N₂ and O₂ molecules, the sum over the terms $n_m A_m$ represents cascading from higher-energy states, v_k is the excitation frequency for generating the upper excited states responsible for emissions from 2PN₂ and 1NN⁺₂, and n_e is the number density of electrons. For the sake of simplicity, steady state assumption is applied in order to find out n_k . This is valid given that the lifetimes of the upper excited states responsible for 2PN₂ and 1NN⁺₂ emissions are very short (less than several microseconds [*Vallance Jones*, 1974, p. 119]) when compared to the characteristic time scale of the TGF sources considered in this paper.

The excitation frequency, describing contributions to n_k from direct electronic excitation by electrons, is consistently calculated using Monte Carlo simulation results. The quantity v_k , defined as the number of molecular species *j* in the excited state *k* produced in unit time per one electron, is given by

$$v_k = N_j \int \sigma_k(\varepsilon) \mathbf{v}(\varepsilon) f(\varepsilon) d\varepsilon$$
(3)

where N_j is the number density of unexcited species j, $\sigma_k(\epsilon)$ is the cross section for the excitation of state k, which depends on the kinetic energy of the incident electron ϵ , $v(\epsilon)$ is the velocity of the colliding electron with an energy ϵ defined as $v(\epsilon) = c\sqrt{1 - [1 + \epsilon/(m_e c^2)]^{-2}}$, where c is the speed of light and m_e is the electron rest mass, and $f(\epsilon)$ is the electron energy distribution function, which is normalized so that $\int_0^{+\infty} f(\epsilon) d\epsilon = 1$. Electron impact excitation cross sections used for determining the excitation frequency of generating the upper excited states of $2PN_2$ are taken from BOLSIG+ database [*Hagelaar and Pitchford*, 2005]. These cross sections are provided between the threshold energy and 1 keV. However, for the energy distributions considered here, their values for energies >100 eV are small and their effects are negligible (see Figure 5b). Concerning $1NN_2^+$, it is considered that an $N_2^+(B^2\Sigma_u^+)$ ion is produced if one of the two $(2\sigma_u)$ electrons from the nitrogen molecule is removed [e.g., *Van Zyl and Pendleton*, 1995]. Thus, an accurate representation for the ionization cross section of $(2\sigma_u)$ electrons from the threshold energy to hundred MeV is needed. The RBEB model, due to its orbital-based description for the differential ionization cross section, is therefore well suited for the calculation of the excitation frequency of $N_2^+(B^2\Sigma_u^+)$ leading to $1NN_2^+$ emissions. Indeed, the branching ratio fraction for producing $N_2^+(B^2\Sigma_u^+)$ during ionization reactions of N_2 molecules by 100 eV electrons calculated using the RBEB model yields 0.163 and agrees well with the value 0.145 \pm 0.019 suggested by *Van Zyl and Pendleton* [1995].

Collisional quenching processes are critical at the source altitudes of TGFs [e.g., *Dwyer and Smith*, 2005; *Xu* et al., 2012], which are far below the quenching altitudes of both $2PN_2$ and $1NN_2^+$ [e.g., *Liu et al.*, 2006]. In the present study, we consider that $N_2(C^3\Pi_u)$ can be quenched by collisions with N_2 and O_2 molecules with rate coefficients of 10^{-11} cm³/s [*Kossyi et al.*, 1992] and 3×10^{-10} cm³/s [*Vallance Jones*, 1974, p. 119], respectively. The primary quenchers of $N_2^+(B^2\Sigma_u^+)$ are N_2 , with a rate coefficient of 4.53×10^{-10} cm³/s [e.g., *Mitchell*, 1970; *Kuo et al.*, 2005; *Pancheshnyi et al.*, 1998], and O_2 , with a rate coefficient of 7.36×10^{-10} cm³/s [e.g., *Mitchell*, 1970; *Kuo et al.*, 2005; *Pancheshnyi et al.*, 1998]. It is important to note that optical emissions evaluated using the modeling approach adopted in this paper significantly rely on the choice of quenching scheme and corresponding rate coefficients.

As in *Dwyer et al.* [2013], aurora spectra of $2PN_2$ and $1NN_2^+$ presented in *Vallance Jones* [1974] are used as an approximate spectral distribution for fluorescence photons to investigate the wavelength dependence of optical features. We note that our present calculations do not take into account the effects of radiative transfer between the source of emission and the observer.

2.3. Characteristic Radial Dimensions

The characteristics of optical emissions that can be possibly generated during TGF production, including intensities and intensity ratios between optical band systems, mostly depend on the energy distribution and number density of electrons. In this work, except for the comparison with laboratory observations in section 3.1, we assume a uniform spherically symmetric spatial distribution for source electrons involved in TGFs. The number density of electrons is then obtained using the full energy distribution and an assumption that 10¹⁷ electrons with energy above 1 MeV are generated during TGFs, that is believed to be required for explaining TGF fluences and spectra [*Dwyer and Smith*, 2005]. Although this assumption is not accurate enough for capturing the morphological characteristics of TGF-associated optical emissions, it does not have a significant impact on quantifying the total intensities of optical emissions. The characteristic radial dimensions representing processes of RREAs, acceleration and multiplication of thermal runaway electrons in the electric field produced by the 100 MV lightning leader, and the streamer zone of the 100 MV lightning leader are discussed below, schematically depicted in Figure 1 and summarized in Table 2.

In RREAs, propagation over multiple runaway avalanche lengths is required for relativistic electrons to gain energies on the order of tens of MeV and to explain TGF spectra, corresponding to an acceleration distance of approximately 1 km for typical electric fields in thunderclouds [e.g., *Dwyer et al.*, 2012]. In contrast, the electric field produced by a 100 MV lightning leader is highly inhomogeneous, intense close to the leader tip, and weak far from it [e.g., *Celestin et al.*, 2012]. This electric field is able to accelerate thermal runaway electrons over relatively short distances and a representative acceleration distance is found to be approximately 50 m for a 100 MV lightning leader.

Additionally, in this work, the intensity of optical emissions radiated from the streamer zone of the 100 MV lightning leader is also estimated using the streamer modeling results presented in *Liu et al.* [2008] and the knowledge about the number of streamers and the size of the streamer zone. Specifically, the number of streamers present in a streamer zone is $N_s = Q_s/q_s$ [e.g., *Bazelyan and Raizer*, 2000; *Celestin and Pasko*, 2011], where Q_s is the total electric charge contained in the streamer zone, and q_s is the average charge carried by a streamer, typically on the order of 1 nC [*Bazelyan and Raizer*, 2000, p. 69–71]. As pointed out by *Celestin and Pasko* [2011], Q_s is quadratically dependent on the electric potential difference formed by



Figure 2. Simulation results of fluorescence radiation from air at ground pressure ($N_0 \simeq 2.688 \times 10^{25} \text{ m}^{-3}$) excited by a continuous beam of 50 keV electrons. Fluorescence emissions exhibit a conical shape. The diameter of this beam, best represented by the strongest emissions from 2PN₂, is approximately 4 cm.

lightning leaders with respect to ambient potential (U_{l}) and can be calculated as follows:

$$Q_{\rm s} = \frac{\pi \varepsilon_0 U_{\rm l}^2}{2E_c^2} \tag{4}$$

where E_s^- is the electric field in the steamer zone of negative leaders and its value is taken as 12.5 kV/cm [*Babaeva and Naidis*, 1997, Figure 7]. From these relations, we estimate that, for long unbranched lightning leaders with electric potential of 100 MV, the total number of streamers constituting the streamer zone should be on the order of 10⁸. As for the characteristic size of the

streamer zone, it is proportional to $U_1: R_{sz}^- = U_1/2E_s^-$. Therefore, the radius of the streamer zone in front of a 100 MV lightning leader is approximately 40 m at ground level.

3. Results

3.1. Comparison With Laboratory Observations

In order to perform a validation of the present numerical models, fluorescence emissions produced by energetic electrons are first calculated in the framework of Monte Carlo simulation and compared to laboratory observations of *Davidson and O'Neil* [1964] and *Nagano et al.* [2004]. One of the advantages that is afforded by the current Monte Carlo model is that it is capable of accurately describing and recording the spatial and temporal information for all the collisions taking place in the system. Owing to this advantage, an analog of photograph of fluorescence from air generated by a continuous beam of 50 keV electrons at ground pressure can be derived from first principles and is presented in Figure 2. As clearly shown in this figure, these fluorescence emissions exhibit a conical shape, following the spatial distribution of electrons, and the diameter of this beam, best represented by the strongest emissions from 2PN₂, is approximately



Figure 3. Steady state energy distribution of a continuous beam of 50 keV electrons in air at 600 torr. Also shown is the dynamic friction force of electrons at ground pressure due to collisions with air molecules computed using the set of cross sections adopted in Monte Carlo simulation (red dashed line). The dominant energy loss processes in regions I, II, and III, are vibrational and rotational excitations, electronic excitation, and ionization collisions, respectively. The circle around 14.1 eV represents the enhancement in the dynamic friction force, mainly arising from energy losses to the electronic excitation of $N_2(C^3\Pi_u)$, that is also reflected in the electron energy distribution.

4 cm. The morphological features of these optical emissions, including the conical shape and the size of the illuminated region, are in excellent agreement with experimental observations [*Davidson and O'Neil*, 1964].

In the absence of external electric fields, the energy distribution, especially in the low-energy range (up to a few tens of eV), is characterized by the dynamic friction force of electrons. This point can be readily illustrated by a direct comparison of the dynamic friction force of electrons resulting from collisions with air molecules at ground pressure and the steady state electron energy distribution produced by the continuous beam of 50 keV electrons, as shown in Figure 3. This energy distribution is calculated by adopting a convolution technique. In particular, we first model the temporal evolution of the deceleration process undergone by an ensemble of 50 keV electrons due to collisions with air molecules in the case that they are injected instantaneously into air at 600 torr.



Figure 4. Comparison of fluorescence efficiency from air at 600 torr for major emission bands of $2PN_2$ and $1NN_2^+$, in the wavelength range between 300 nm and 430 nm, excited by (a) 50 keV electrons between experimental measurements of *Davidson and O'Neil* [1964] and present modeling results; (b) 850 keV electrons between experimental measurements of *Nagano et al.* [2004] and present modeling results.

Namely, we obtain the impulse response of the system. The steady state energy distribution of the continuous 50 keV electron beam is then calculated by convolving over the temporal evolution of this electron pulse. We have verified that directly modeling a continuous beam of 50 keV electrons in the simulation leads to identical results, while being a lot more time consuming. From Figure 3, we see that the depletion in the electron energy distribution around 2.5 eV precisely pinpoints the first peak of the dynamic friction force, defined by strong vibrational excitations of nitrogen molecules, while the peak of electron energy distribution around 4 eV coincides with a trough in the dynamic friction force. Furthermore, enhancement around 14.1 eV in the dynamic friction force (marked by a circle in Figure 3), that mainly comes from energy losses to the electronic excitation of $N_2(C^3\Pi_u)$, is reflected in the energy distribution.

In the region I marked in Figure 3, where the electron energy is below ~4 eV, vibrational and rotational excitations are the dominant energy loss processes. In particular, vibrational excitations facilitate fast removal of electrons from this energy region and accordingly cause the precipitous decrease in the energy distribution from 1 eV to ~2.5 eV. Region II is delineated by electrons with energy between ~4 eV and ~28 eV. Electronic excitation is the principal channel of energy loss in this region and the cross sections of electronic excitation collisions for generating the upper excited states of $2PN_2$ peak in this region. In region III, electrons possess energy in excess of the binding energy of electrons in N₂ and O₂ molecules and mostly convert their kinetic energy into the generation of secondary electrons, namely, ionization collisions.

Fluorescence efficiency is defined in the studies of extensive air showers [*Bunner*, 1967] as the fraction of the total energy deposited during the passage of electrons in a gas that is eventually transferred into fluorescence photons. For the comparison with *Nagano et al.* [2004], fluorescence efficiency is calculated by dynamically recording the number of fluorescence photons generated and the amount of energy deposited by 850 keV electrons in Monte Carlo simulations. However, in contrast with the thin target technique used in *Nagano et al.* [2004], *Davidson and O'Neil* [1964] employed a thick target technique to measure the fluorescence efficiency of 50 keV electrons by injecting a continuous electron beam into the gas chamber. In order to compare with the experimental measurements in *Davidson and O'Neil* [1964], we first calculate the production of fluorescence photons in the wavelength range between 300 and 430 nm and the energy deposition into air molecules by an ensemble of 50 keV electrons in the case that they are injected instantaneously into air at 600 torr. Fluorescence emissions corresponding to the steady state of continuous injection are then obtained by convolving the fluorescence pulse produced by the instantaneous injection of 50 keV electrons and assuming the same injection current as that in *Davidson and O'Neil* [1964]. Fluorescence efficiency is further determined as the ratio of the energy released in fluorescence photons to the energy deposited by the incident electron beam.

The comparison with wavelength-resolved measurements [*Davidson and O'Neil*, 1964; *Nagano et al.*, 2004] is shown in Figure 4 and documented in Table 1. Our modeling results are generally in very good agreement with laboratory measurements of *Nagano et al.* [2004] for both 2PN₂ and 1NN₂⁺ band systems.

Table 1. Comparison of Fluorescence Efficiency ($\times 10^{-5}$) From Air at 600 Torr for Major Emission Bands of 2PN₂ and 1NN⁺₂, in the Wavelength Range Between 300 nm and 430 nm, Excited by Energetic Electrons Between Experimental Measurements of *Davidson and O'Neil* [1964]^a (50 keV electrons), *Nagano et al.* [2004] (850 keV electrons), and Present Modeling Results

	Davidson and O'Neil [1964]	Nagano et al. [2004]		
Wavelength (Å)	50 keV	850 keV	50 keV	850 keV
3371, 2PN ₂ (0–0)	2.10	2.23-2.45	3.306	2.703
3537, 2PN ₂ (1–2)	0.32	0.237-0.331	0.695	0.568
3577, 2PN ₂ (0–1)	1.50	1.56-1.90	2.076	1.697
3756, 2PN ₂ (1–3)	0.30	0.415-0.563	0.582	0.476
3805, 2PN ₂ (0-2)	0.52	0.480-0.680	0.783	0.640
3914, 1NN ₂ +(0–0)	0.70	0.558-0.638	0.951	0.791
3943, 2PN ₂ (2–5)	0.05	0.058-0.188	0.131	0.107
3998, 2PN ₂ (1–4)	0.18	0.212-0.286	0.333	0.272
4059, 2PN ₂ (0-3)	0.18	0.209-0.243	0.230	0.188
4278, 1NN ₂ ⁺ (0-1)	0.27	0.181-0.259	0.265	0.220

^aThe systematic error for the measurements of *Davidson and O'Neil* [1964] is $\pm 15\%$.

This agreement also validates our use of aurora spectra in order to quantify the wavelength dependence of optical emissions produced by energetic electrons. In addition, we note that the mean energy loss of 850 keV electrons derived from Monte Carlo simulation is approximately 1.74 keV/cm, close to the theoretical value defined by the dynamic friction force at this pressure. The slight difference observed for the (0–0) transition of $1NN_2^+$ between calculated and experimental results, at 391 nm, is likely due to three-body deactivation processes as pointed out in *Mitchell* [1970] and *Nagano et al.* [2004], which are able to weaken the efficiency by approximately 20%. As for the fluorescence efficiency produced by continuous injection of 50 keV electrons, our modeling results also agree with experimental measurements of *Davidson and O'Neil* [1964] for $1NN_2^+$, while the agreement is not as good for $2PN_2$ and we note that, for some bands of $2PN_2$ (i.e., $2PN_2(1-2)$), as large as a factor of 2 differences are observed.

3.2. Distribution of Electrons in the Full Energy Range 3.2.1. Relativistic Runaway Electron Avalanches

Figure 5a shows the distribution of electrons in the full energy range produced by the acceleration and multiplication of relativistic runaway electrons in a large-scale homogeneous reduced electric field of E/N=70 Td (~18.8 kV/cm at $N\simeq 2.688 \times 10^{25}$ m⁻³, 1 Td = 10^{-21} Vm²) in air at ground pressure. This energy



Figure 5. (a) Distribution of electrons in the full energy range produced by the acceleration and multiplication of relativistic runaway electrons in a large-scale homogeneous electric field of 70 Td (18.8 kV/cm). The results are obtained in air at ground level ($N_0 \simeq 2.688 \times 10^{25}$ m⁻³). Also presented are BOLSIG+ calculations [*Hagelaar and Pitchford*, 2005] and modeling results presented in *Colman et al.* [2010, Figure 11]. The inset is a zoom-in view of the energy range below 20 eV. (b) Cumulative sum of the frequency for generating $N_2(C^3\Pi_u)$ and $N_2^+(B^2\Sigma_u^+)$ excited by the large population of electrons in RREA processes. Fifty percent of the upper excited states of N_2 responsible for 2PN₂ emissions is produced by electrons with energy below 12.75 eV, while 50% of the upper excited states of N_2^+ responsible for 1NN₂⁺ emissions is produced by electrons with energy below 70.8 keV.



Figure 6. Distribution of electrons in the full energy range produced by the acceleration and multiplication of (a) relativistic runaway electrons in homogeneous electric fields: 4.3 kV/cm and 18.8 kV/cm and (b) relativistic runaway electrons in a homogeneous electric field of 12.5 kV/cm and thermal runaway electrons in the highly inhomogeneous electric field near the tip region of a 100 MV lightning leader during the negative corona flash. The energy distributions are normalized so that the integration over electron energy from 1 MeV to 100 MeV yields unity. The results are obtained in air at ground level atmospheric density. The diamonds in Figure 6a show Monte Carlo simulation results obtained from *Dwyer and Babich* [2011, Figure 3], representing the energy distribution of electrons in RREAs driven by the electric field of 4.3 kV/cm.

distribution is obtained after approximately 11 avalanche lengths, or *e*-folding times, and has already reached steady state. In spite of the noise in the energy distribution, it is clearly shown that over 6 orders of magnitude, the low-energy part (<100 eV) is consistent with recently published full energy range model results [*Colman et al.*, 2010] and BOLSIG+ calculations [*Hagelaar and Pitchford*, 2005]. We also note that the ratio between the number of electrons with energy below and above 1 MeV is found to be 9365, in agreement with the analytical calculation documented in *Dwyer and Babich* [2011].

Steady state electron energy distributions in RREAs, for the cases of homogeneous electric fields with magnitudes of 4.3 kV/cm and 12.5 kV/cm, are presented in Figures 6a and 6b, respectively. The energy distributions are normalized so that the integration from 1 MeV to 100 MeV yields unity, following the assumption that the same number of electrons with energies above 1 MeV is produced in the source of TGFs. Also presented in Figure 6a are Monte Carlo simulation results (diamonds) documented in *Dwyer and Babich* [2011, Figure 3], representing the energy distribution of electrons in RREAs driven by an electric field of 4.3 kV/cm. For this electric field, the high-energy part of electron energy distribution shows excellent agreement between present work and results reported in *Dwyer and Babich* [2011, Figure 3]. We therefore estimate that our results on the electron energy distribution are validated over more than 15 orders of magnitude of the dynamic range. To our knowledge, such a result, over the full energy range of interest here, had never been obtained.

Unlike the case in Figure 3 (see discussion in section 3.1), energy distribution for electrons propagating under external electric fields, like in RREAs, is governed by the competition between the energy gained from electric fields and energy losses in collisions with neutral gas molecules, i.e., the dynamic friction force. The avalanche of relativistic runaway electrons is characterized by abundant generation of large quantities of secondary electrons over the full energy range. Secondary electrons with energies below ~2 eV are mostly trapped in this region because the applied electric field is insufficient to energize them to penetrate the vibrational barrier at ~2 eV (see Figure 3). Secondary electrons in the "valley" of the dynamic friction force (between ~3 eV and ~15 eV) may be subject to an electric force that is capable of overcoming their energy losses in collisions with air molecules and therefore be accelerated to higher energies, depending on the strength of the driving electric field and the momentum of electrons. A driving electric field with a higher strength can surmount a larger dynamic friction force, corresponding to higher electron energies in the sharp increase between ~4 eV and ~123 eV (see Figure 3). Therefore, one sees from Figure 6a that, under a stronger electric field, more electrons are sustained in the energy range between ~2 eV and ~15 eV. Also, in the high-energy part, the energy cutoff corresponding to the distribution obtained in 18.8 kV/cm is higher than that in 4.3 kV/cm.

For the purpose of understanding the energy dependence of the capability of electrons in generating the upper excited states responsible for optical emissions from $2PN_2$ and $1NN_2^+$, the excitation frequencies

Table 2. Intensity of Optical Emissions from $2PN_2$ (Column 3) and $1NN_2^+$ (Column 4) in Rayleighs and Intensity Ratio Between $2PN_2$ and $1NN_2^+$ (Column 5) in the Visible Range With Wavelengths Between 390 nm and 700 nm for Different Acceleration Processes (Column 1) With Different Characteristic Sizes (Column 2) Calculated at Ground Level

	Radius (m)	2PN ₂ (R)	1NN ₂ ⁺ (R)	$\frac{2PN_2}{1NN_2^+}$
RREA (4.3 kV/cm)	1000	$8.99 imes 10^{8}$	1.22×10^{9}	0.74
RREA (12.5 kV/cm)	1000	1.70 × 10 ⁹	1.55 × 10 ⁹	1.10
RREA (18.8 kV/cm)	1000	6.63 × 10 ⁹	1.31 × 10 ⁹	5.06
Thermal runaway electrons	50	8.28×10^{11}	5.23×10^{11}	1.58
Streamer zone	40	6.83×10^{10}	6.75 × 10 ⁸	101.19

are calculated using equation (3) and the energy distribution obtained in the present work and shown in Figure 5a. Figure 5b presents the cumulative sum of the frequency for generating N₂($C^3\Pi_u$) and N₂⁺($B^2\Sigma_u^+$) in RREAs driven by an electric field of 70 Td. It is found that 50% of the upper excited states of N₂ responsible for 2PN₂ emissions is produced by electrons with energy below 12.75 eV while 50% of the upper excited states of N₂⁺ responsible for 1NN₂⁺ emissions is produced by electrons with energy below 70.8 keV. This feature can be attributed to the fundamental difference in cross sections. The excitation cross section for generating N₂⁺($B^2\Sigma_u^+$) rises rapidly above the threshold, attains a maximum value around 114 eV, drops off slowly to around 1 MeV, and then reaches a relativistic plateau above 1 MeV. Consequently, even though they are in minority, relativistic electrons efficiently ionize nitrogen molecules, resulting in comparable amount of N₂⁺($B^2\Sigma_u^+$) when compared to that produced by low-energy electrons. In contrast, the cross section of 2PN₂ falls off much faster after the peak and becomes negligible in the region above 1 keV, where ionization and elastic collisions dominate. As a result, optical emissions from 2PN₂ are mainly produced by low-energy electrons. We note that since the electron impact excitation cross section for generating N₂($B^3\Pi_g$) has a similar behavior as that of 2PN₂, optical emissions from 1PN₂ should also mainly come from low-energy electrons.

3.2.2. Acceleration of Thermal Runaway Electrons in Lightning Leader Field

Figure 6b shows the full energy distribution for the ensemble of electrons accelerated in the highly inhomogeneous electric field around the tip region of a 100 MV lightning leader during the negative corona flash stage. The electron energy distributions are normalized so that the integration from 1 MeV to 100 MeV yields unity. Due to the differences in geometry and magnitude between the homogeneous and inhomogeneous electric fields, a deviation from the RREA energy distribution is observed: a much smaller amount of electrons is generated and accumulated in the low-energy range and the high-energy tail does not follow the typical energy cutoff observed in RREAs.

3.3. Optical Emissions

Table 2 shows modeling results of optical emissions in the visible range with wavelengths between 390 nm and 700 nm from the following processes at ground level: RREAs, acceleration of thermal runaway electrons in the electric field produced by the 100 MV lightning leader, and the streamer zone of the 100 MV lightning leader. The columns 3 and 4 represent the maximum intensities of optical emissions from 2PN₂ and 1NN⁺₂, respectively, during the listed processes. Column 5 shows the intensity ratio of 2PN₂ to 1NN⁺₂ associated with the listed processes. Interestingly, both TGF production mechanisms produce considerable amount of optical emissions, with magnitudes comparable to those from the streamer zone associated with the 100 MV lightning leader. The intensities of optical emissions from the streamer zone of the 100 MV lightning leader are, respectively, 6.83×10^{10} R and 6.75×10^8 R for 2PN₂ and 1NN⁺₂. The intensity ratio of 2PN₂ to 1NN⁺₂ is ~101.19 as indicated in *Liu et al.* [2008].

The optical emissions from $2PN_2$ (column 3 of Table 2) associated with RREAs intensify with increases in driving electric fields. For instance, the intensities of optical emissions from $2PN_2$ corresponding to RREAs under electric fields of 4.3 kV/cm, 12.5 kV/cm, and 18.8 kV/cm are, respectively, 8.99×10^8 R, 1.70×10^9 R, and 6.63×10^9 R. However, optical emissions from $1NN_2^+$ associated with RREAs do not indicate the same tendency. The intensities for 4.3 kV/cm, 12.5 kV/cm, and 18.8 kV/cm are, respectively, 1.22×10^9 R, 1.55×10^9 R, and 1.31×10^9 R. To understand how these characteristic tendencies in optical output stem from the differences in electron energy distributions, it is helpful to look at the comparison presented in Figure 6a. As the homogeneous electric field increases from 4.3 kV/cm to 18.8 kV/cm, a larger population of

electrons is generated and sustained in the energy region from ~11 eV (the threshold energy of producing $N_2(C^3\Pi_u)$) to ~15 eV, which plays a critical role in generating optical emissions from 2PN₂ (see Figure 5b and discussion in section 3.2.1). Therefore, a stronger driving electric field should demonstrate more intense optical emissions from 2PN₂ during RREAs as evident in Table 2. Contrary to 2PN₂, optical emissions from $1NN_2^+$ can be as efficiently produced by high-energy electrons as by low-energy ones (see Figure 5b and discussion in section 3.2.1). Considering that the energy distribution of electrons accelerating in 18.8 kV/cm only slightly differs from that of 4.3 kV/cm in the energy region above ~20 eV (the threshold energy of producing $N_2^+(B^2\Sigma_u^+)$), optical emissions from $1NN_2^+$ should not change considerably, as also evident in Table 2.

The two TGF production mechanisms exhibit distinct optical features. First, when compared to RREA processes, optical emissions associated with the acceleration of thermal runaway electrons in lightning leader fields are more intense, with 8.28×10^{11} R from $2PN_2$, and 5.23×10^{11} R from $1NN_2^+$. This is primarily because the production of TGFs by thermal runaway electrons is naturally confined to a relatively compact region in the vicinity of negative leader tip [e.g., Xu et al., 2012; Celestin et al., 2012]. Second, the intensity ratios of 2PN₂ to 1NN⁺₂ are different between the two production mechanisms. For example, the intensity ratios between 2PN₂ and 1NN₂⁺ are, respectively, 0.74, 1.10, and 5.06 for RREAs driven by homogeneous electric fields of 4.3 kV/cm, 12.5 kV/cm, and 18.8 kV/cm while this ratio is 1.58 for thermal runaway electrons produced and accelerated in the vicinity of the tip of a 100 MV lightning leader. This feature is also a reflection of the inherent differences in electron energy distributions. In the case of RREAs, as discussed in the previous paragraph, a stronger driving electric field enables more intense optical emissions from $2PN_2$ while producing similar levels of emissions from 1NN₂⁺. For this reason, one observes in Table 2 that the intensity ratio between 2PN₂ and 1NN⁺₂ in RREAs becomes larger with the increase in driving electric fields. Nevertheless, as illustrated in Figure 6b, the electron energy distributions are similar in the energy regions critical for optical emissions from $2PN_2$ and $1NN_2^+$ between the processes of thermal runaway electrons accelerating in a 100 MV lightning leader field and RREAs under an electric field of 12.5 kV/cm. Thus, the intensity ratios of 2PN₂ to 1NN⁺₂ are close between these two processes as presented in Table 2.

4. Discussion

4.1. Fluorescence Efficiency

For the sake of comparison with previously published results [*Dwyer et al.*, 2013], we have calculated the fluorescence efficiency without collisional quenching corresponding to the steady state of RREAs for $2PN_2$ and $1NN_2^+$ using:

$$\epsilon_{k} = \frac{\nu_{k} \sum_{\lambda} E_{\lambda} I_{k}(\lambda)}{\int F_{d}(\varepsilon) v(\varepsilon) f(\varepsilon) d\varepsilon}$$
(5)

where E_{λ} is the energy of fluorescence photons with a wavelength of λ defined as $E_{\lambda} = hc/\lambda$, where *h* is the Planck's constant and *c* is the speed of light in free space, $I_k(\lambda)$ is the intensity distribution of optical emissions from either 2PN₂ or 1NN₂⁺ obtained from *Vallance Jones* [1974] and is normalized so that $\sum_{\lambda} I_k(\lambda) = 1$, and $F_d(\varepsilon)$ is the dynamic friction force due to collisions with air molecules that electrons with an energy of ε experience at ground level (see Figure 3). The summation in the numerator is taken over wavelengths of major emission bands from 2PN₂ or 1NN₂⁺.

It is found that the fluorescence efficiencies of $2PN_2$ for RREAs driven by electric fields of 4.3 kV/cm, 12.5 kV/cm, and 18.8 kV/cm, in the visible range of 390–700 nm without collisional quenching, are 0.0146%, 0.00287%, and 0.0208%, respectively. Fluorescence efficiencies of $1NN_2^+$, in the same wavelength range without collisional quenching, are 0.213%, 0.0283%, and 0.0443% for RREAs driven by electric fields of 4.3 kV/cm, 12.5 kV/cm, and 18.8 kV/cm, respectively. Based on these calculations, fluorescence efficiency in RREAs can vary notably for driving electric fields with different strengths.

More intense driving electric fields do not necessarily facilitate the associated fluorescence efficiency, even if they result in more energy deposition. In RREAs, energy deposition and fluorescence emissions are caused by electrons in different energy regions. To illustrate this discussion, we have verified that, for the three electric fields studied in the present work, most of the energy deposited via collisions with air molecules is from the large ensemble of electrons in the energy region below ~4 eV while optical emissions from $2PN_2$ are mostly produced by electrons with energies between ~11 eV and 100 eV.

As the driving electric field changes from 4.3 kV/cm to 12.5 kV/cm, although secondary electrons in the energy region between ~11 eV and 100 eV are able to excite more $N_2(C^3\Pi_u)$ and release more energy to generate fluorescence photons from $2PN_2$, secondary electrons in the energy region below ~4 eV much more efficiently deposit their kinetic energy into air molecules through vibrational and rotational excitations. As a consequence, the fluorescence efficiency for $2PN_2$ decreases. However, when the driving electric field further increases to 18.8 kV/cm, a much larger population of secondary electrons can be accelerated to and maintained in the energy region that is critical for $2PN_2$ and, therefore, an enhancement in fluorescence efficiency is observed. Similarly, fluorescence efficiency for $1NN_2^+$ in RREAs is also highly dependent on the strength of the driving electric field.

Moreover, extensive air showers have also been studied through modeling of the deceleration of 1 MeV electrons [e.g., *Lafebre et al.*, 2009] in the atmosphere. Using the same model as discussed in the present paper, we have estimated the fluorescence efficiencies of 1 MeV electrons decelerating through air. It is interesting to note that the fluorescence efficiencies for $2PN_2$ and $1NN_2^+$, between 390 and 700 nm and without collisional quenching, are found to be 0.0405% and 0.554%, respectively, that are very close to the values reported and used in *Dwyer et al.* [2013]. We note that the calculations of *Dwyer et al.* [2013] were based on experimentally measured fluorescence efficiency for 0–0 bands of $2PN_2$ and $1NN_2^+$ [*Keilhauer et al.*, 2006].

Additionally, we have estimated the amount of fluorescence photons that can be generated per runaway electron per meter in the wavelength range between 300 and 430 nm corresponding to the steady state of RREAs. This wavelength range is chosen in order to cover major emission bands of $2PN_2$ and $1NN_2^+$. The values are approximately 6.2, 10.7, and 38.3 m⁻¹ for RREAs driven by electric fields of 4.3 kV/cm, 12.5 kV/cm, and 18.8 kV/cm at ground pressure, respectively. This shows that the stronger the sustaining electric fields, the more intense the associated fluorescence emissions. Note that this is consistent with the transition in electron energy distributions (i.e., enhancement in low-energy part with increasing applied electric field) presented in Figure 6a.

Fluorescence efficiency, in the study of air showers, describes a process in which primary electrons eventually deposit all their initial kinetic energy into air molecules of the Earth's atmosphere, which release it through emissions of fluorescence and collisional deexcitation. However, in TGFs, the acceleration of electrons and the subsequent deposition of energy into air molecules are supported by external electric fields. The amount of energy that is converted into fluorescence emissions mainly comes from that acquired from the driving external electric field. Therefore, we suggest that fluorescence efficiency in extensive air showers is not conceptually identical to that in TGFs. Also, the electron energy distribution obtained from the deceleration process in extensive air showers can be substantially different from that in TGFs. Due to the fact that the capability of electrons in generating fluorescence photons is strongly energy-dependent (see Figure 5b and discussion in section 3.2.1), fluorescence efficiency measured in extensive air showers is different from that in TGFs.

4.2. Scaling of Optical Emissions

Modeling results of optical emissions from the acceleration of thermal runaway electrons in lightning leader field and the associated streamer zone (Table 2) are obtained at ground level air density and can be applied to the source altitudes of TGFs using similarity scaling. As indicated in equation (1), the intensity of optical emissions is closely related to the number density of excited species n_k . For the two optical band systems considered in the present work, i.e., 2PN₂ and 1NN₂⁺, n_k is populated by electron impact excitations (the term $v_k n_e$ in equation (2)) and depopulated by spontaneous emissions (the term A_k in τ_k) and collisional quenching processes with air molecules (the terms $\alpha_1 N_{N_2}$ and $\alpha_2 N_{O_2}$ in τ_k). At or below the source altitudes of TGFs [e.g., *Dwyer and Smith*, 2005; *Xu et al.*, 2012], collisional quenching processes dominate over spontaneous emissions and, in the steady state, n_k is obtained by the balance between direct excitations and collisional quenching. Since both collisional quenching ($\alpha_1 N_{N_2} + \alpha_2 N_{O_2}$) and electron impact excitation frequency (v_k) scale as the number density of air molecules ($\sim N$), n_k scales as the number density of electrons n_e .

The total number of energetic electrons emitted from a lightning leader is invariant of air pressure [*Moss et al.*, 2006]. The characteristic radial dimension of the acceleration of thermal runaway electrons in the lightning leader field is defined as the region in which thermal runaway electrons accelerate and efficiently gain energy. This condition is fulfilled if the electric force exerted on thermal runaway electrons $q_e E$

exceeds the average friction force due to collisions with air molecules $q_e E_t$, where q_e is the electron elementary charge, E is the inhomogeneous electric field produced around lightning leader tips, and E_{+} is the relativistic runaway threshold field and its value at ground level atmospheric density is ~2.8 kV/cm [e.g., Dwyer et al., 2012]. As suggested by Celestin et al. [2012], the highly inhomogeneous electric field E produced by high-potential lightning leaders can be approximated by an A/r function, where $A = 0.04E_0/r$ is a fitting parameter and r is the distance from the leader tip. Since r varies inversely as E and E_t scales proportionally with the density of the neutral atmosphere, i.e., $E_t \sim N$, the characteristic radial dimension representing the acceleration of thermal runaway electrons in lightning leader field scales as $\sim 1/N$. Therefore, the number density of energetic electrons responsible for producing $N_2(C^3\Pi_u)$ or $N_2^+(B^2\Sigma_u^+)$ scales as $\sim N^3$. Furthermore, considering that the intensity of optical emissions in Rayleighs is obtained by integrating over the line of sight, which scales as $\sim 1/N$, and remembering that n_k scales in the same way as n_{e} , one sees that the cross-sectional intensity of optical emissions (number of photons per unit area) produced during the acceleration of thermal runaway electrons in lightning leader fields scales as $\sim N^2$. For example, the characteristic radius for the acceleration process of thermal runaway electrons in the 100 MV lightning leader case is 50 m at ground level. However, this quantity at an altitude of 12 km, corresponding to the source altitude of TGFs [e.g., Xu et al., 2012], would be \sim 211 m. The intensity of 2PN₂ is 8.28 \times 10¹¹ R at ground level and would be reduced to \sim 4.63 \times 10¹⁰ R at 12 km altitude.

As for the streamer zone associated with the 100 MV lightning leader, its radius R_{sz}^- is inversely proportional to E_s^- , which scales as $\sim N$. Thus, R_{sz^-} scales as $\sim 1/N$. For the same reason, the total amount of electric charge contained in the streamer zone Q_s also scales as $\sim 1/N$ (see equation (4)). Because the streamer electron density scales as $\sim N^2$ and the length scale of streamers scales as $\sim 1/N$ [e.g., *Liu and Pasko*, 2004], the average electric charge carried by a streamer q_s scales as $\sim 1/N$. Since both Q_s and q_s scale as $\sim 1/N$, the number of streamers constituting a streamer zone, which follows the relation $N_s = Q_s/q_s$, remains the same with respect to air pressure/altitude. In a single streamer, n_k scales as the streamer electron density, $\sim N^2$. As a result, by integrating over the line of sight, the total cross-sectional intensity of optical emissions from the streamer zone scales as $\sim N$.

4.3. Optical Emissions Associated With TGFs

Different driving electric fields can lead to different acceleration and multiplication processes undergone by runaway electrons, different energy distributions obtained, and thus different capabilities in generating optical emissions. In RREA mechanism, relativistic runaway electrons harvest energy from the homogeneous electric field and gradually build up a ~7 MeV high-energy cutoff while producing a large ensemble of low-energy electrons. Reciprocally, as demonstrated in *Celestin et al.* [2012], the intense inhomogeneous electric field naturally present in compact regions around negative lightning leader tips during stepping processes can accelerate thermal runaway electrons to high energy over much shorter distances, corresponding to much less low-energy electrons generated and a significantly different high-energy cutoff. A direct consequence of these differences is that the optical output associated with these two mechanisms is different. For example, typical magnitude of the large spatial scale electric field measured within thunderstorms is lower than 4.3 kV/cm [e.g., *Marshall et al.*, 2001], the optical output from RREA processes driven by such field would be much weaker than that resulting from the acceleration process of thermal runaway electrons in lightning leader fields (see Table 2). In addition, the intensity ratio between $2PN_2$ and $1NN_2^+$ would be 1 order of magnitude smaller than that of the acceleration and multiplication processes of thermal runaway electrons in lightning leader fields.

TGF events are most likely accompanied with detectable levels of optical emissions for both existing models as suggested by modeling results (see Table 2). These optical emissions would be dominantly blue, or purplish blue (as previously noted by *Dwyer et al.* [2013]). Moreover, using modeling results of optical emissions produced during the acceleration of thermal runaway electrons in lightning leader fields (see Table 2) and considering that this process typically lasts from a few tens of microseconds to a fraction of millisecond, we estimate that the total amount of optical energy radiated in the visible wavelength range in this scenario would be on the order of a few tens of joules. It is comparable to that theoretically produced during relativistic feedback discharges [*Dwyer et al.*, 2013] but much smaller than that of normal lightning [*Uman*, 2001]. Further comparisons between modeling results and ground-based or space-based observations would require the transport of optical photons through the atmosphere to be taken into account. This is especially important in view of the future space missions ASIM (ESA) and TARANIS (CNES)

that will detect TGFs from the nadir direction in association with photometric measurements. Furthermore, the temperature and humidity dependence of air fluorescence emissions have been recently observed [e.g., *Ave et al.*, 2008] and this effect should be accounted for to ensure accurate interpretation of optical observations.

The theory of TGFs produced by thermal runaway electrons initiated during negative corona flashes may introduce a chronological order between the optical emissions from molecules excited by thermal runaway electrons and their by-products, and those from the streamer zone. Indeed, one can expect that most thermal runaway electrons and their corresponding optical emissions would be produced during the early establishment of the streamer zone, the transient stage of negative corona flashes. According to the discussion in *Moss et al.* [2006], the duration of the negative corona flash can be estimated from the speed of streamers and the size of the eventual streamer zone. Considering that streamers, on average, have speeds on the order of ~ 5×10^6 m/s [e.g., *Stenbaek-Nielsen et al.*, 2013] and the typical size of the streamer zone associated with high-potential lightning leaders in +ICs is in the range 50–600 m [e.g., *Winn et al.*, 2011], the corresponding negative corona flashes should have a duration in the range 10–200 µs. However, streamer zones could last for much longer time, typically in the range from 100 µs to 1 ms in +ICs [e.g., *Marshall et al.*, 2013; *Lu et al.*, 2011]. The lightning leaders in –CGs can emit intense bursts of X-rays during the stepping process [e.g., *Dwyer et al.*, 2005]. It is possible that fluorescence light is also observable around the tip of these leaders.

Even if difficult to observe, optical emissions from molecules excited by thermal runaway electrons accelerating in lightning leader fields are temporally and spatially separated from those normally produced by lightning leaders. Optical emissions associated with the acceleration process of thermal runaway electrons in lightning leader fields would be produced right after the establishment of a new leader step, while the lightning leader would emit light during the connection of the new step. Additionally, since thermal runaway electrons gain energy from the electric field produced in the vicinity of lightning leader tips, fluorescence photons caused by thermal runaway electrons would be emitted from a compact region in front of lightning leaders. Moreover, considering that electrons involved in lightning discharges and those produced by thermal runaway electrons can follow significantly different energy distributions, it is conceivable that the intensity ratio of optical emissions from $2PN_2$ and $1NN_2^+$ can be considerably different between the two processes. Finally, the production of optical emissions by RREAs sustained by large-scale thunderstorm electric fields is not necessarily associated with stepping lightning leaders and could therefore be observed in the absence of light from lightning discharges.

5. Conclusions

Principal contributions of this paper can be summarized as follows:

- 1. Using a full energy relativistic Monte Carlo model, we have investigated the fluorescence emissions from air excited by energetic electrons at high pressures. Simulation results, including the morphological features of fluorescence emissions generated by a continuous beam of 50 keV electrons and the fluorescence efficiency for the 2PN₂ and 1NN₂⁺ band systems produced by 850 keV electrons, are consistent with laboratory observations [Davidson and O'Neil, 1964; Nagano et al., 2004]. The importance of the dynamic friction force in characterizing electron energy distribution has been emphasized.
- 2. We have studied the characteristics of electron energy distributions over the full energy range for two TGF production mechanisms: RREAs and production of thermal runaway electrons during the negative corona flash stages of lightning leaders. The simulated energy distribution of RREAs is in good agreement with results presented in e.g., *Dwyer and Babich* [2011] and *Dwyer et al.* [2012] in the high-energy region and BOLSIG+ calculation [*Hagelaar and Pitchford*, 2005] and simulation results in *Colman et al.* [2010] in the lower energy part. Furthermore, we have emphasized that the capability of RREA electrons in generating the upper excited states responsible for optical emissions from 2PN₂ and 1NN₂⁺ is strongly energy dependent. It has also been illustrated how RREAs and the acceleration of thermal runaway electrons in the electric field produced by high-potential lightning leader lead to inherently different electron energy distributions.
- 3. From the knowledge of the electron energy distribution, we have quantified the optical emissions that are possibly generated during the production of TGFs. Modeling results indicate that TGFs are most likely accompanied with detectable levels of optical emissions. However, optical emissions produced during

two TGF production mechanisms.

4. Calculations of fluorescence efficiencies corresponding to air showers and the steady state of RREAs have shown that, although broadly used in the study of air showers to identify the energy of primary particles,

> these fluorescence efficiencies are not appropriate for the study of optical emissions from TGF sources. 5. Optical emissions, appearing as the low-energy signature of the underlying electron acceleration mechanism, are a good probe to point to key ingredients in TGF production. The distinct optical features described in the present work can be used to constrain and validate existing TGF models. Moreover, since TGFs are closely associated with the initial development of +IC flashes [e.g., *Marshall et al.*, 2013], measurements of optical emissions also have important implications for studies of the initial breakdown stage of lightning flashes.

the acceleration of thermal runaway electrons in lightning leader fields are more intense than those associated with RREAs. The intensity ratio between 2PN₂ and 1NN⁺₂ is intrinsically different between these

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