Study of electron energy distribution function and transport parameters for CF₄, Ar gases mixture discharge by using the solution of Boltzmann equation-*Part II*

Baha T. Chiad, Hamid K. Radam, Mohammed J. Jader*

Department of Physics, College of Science, University of Baghdad, Jadiriya, Baghdad, Iraq * Department of Physics, College of Science, University of Babylon, Babylon, Iraq

Abstract The Boltzmann transport equation is solved by using two- terms approximation for pure gases and mixtures. This method of solution	Keywords
is used to calculate the electron energy distribution function and electric transport parameters were evaluated in the range of E/N varying from $1 \times 10^{-17} V \ cm^2 \le E \ / N \le 5 \times 10^{-15} V \ cm^2$.	
The electron energy distribution function of CF_4 gas is nearly Maxwellian at (1,2)Td, and when E/N increase the distribution function is non Maxwellian. Also, the mixtures are have different energy values depending on transport energy between electron and molecule through the collisions. Behavior of electrons transport parameters is nearly from the experimental results in references. The drift velocity of electron in carbon tetraflouride is large compared with other gases and mixtures. The mean electron energy to mixture is increasing at Argon ratios increased.	Article info Received: Mar. 2010 Accepted: Apr. 2010 Published: Dec. 2010

دراسة دالة توزيع طاقة الإلكترون ومعلمات الأنتقال لتفريغ مزيج غازات Ar,CF₄ بواسطة استخدام حل معادلة بولتزمان *II*

بهاء طعمة جياد, حامد كريم ردام, محمد جواد جادر* جامعة بغداد-كلية العلوم-قسم الفيزياء *قسم الفيزياء, كلية العلوم, جامعةبابل/ بابل-العراق

الخلاصه

معادلة بولتزمان حلت بواسطة استخدام التقريب ذي الحدين للغازات النقية والمزيجات. هذه الطريقة استخدم لحساب دالة توزيع طاقة الإلكترون ومعلمات الانتقال الالكتروني حسبت في مدى متغير من E/N $E/N = 5 \times 10^{-15} V \ cm^2$ $M^2 = 10^{-17} V \ cm^2 \le E \ N \le 5 \times 10^{-15} V \ cm^2$ أظهرت تلك النتائج أن دالة توزيع طاقة الإلكترون لغاز CF_4 قريب من التوزيع الماكسويلى عند (1.2) تاوسند، بعد

اظهرت تلك النتائج أن ذالة توزيع طافة الإلكترون لغاز CF₄ قريب من التوزيع الماكسويلي عند (1،2) تاوسند، بعد ذلك عندما تزداد E/N التوزيع يصبح غير ماكسويلي، كذلك المزيحات تمتلك قيم طاقه مختلفه معتمده على الطاقة المنتقلة بين الإلكترون و ألجزيئه . سلوك معلمات الانتقال قريب من النتائج التجريبية في المصادر . سرعة انجراف الإلكترون في غاز رباعي فلوريد الكاربون كانت كبيره مقارنة مع بقية الغازات والمزيجات. متوسط طاقة الإلكترون يزداد عند زيادة نسبة غاز الاركون.

Introduction

electron -velocity distribution The function plays a central role in defining the plasma. physical properties of The statistical behavior of electrons is governed by the electron energy distribution function (EEDF). Transport properties of electron are directly dependent on EEDF.

Theoretically EEDF can be obtained by solving Boltzmann equation. The EEDF is determined by the balance between the gain of the electrons from acceleration electric field and losses of electrons through the collisions [1]. The electron transport parameters of pure gas and gas mixtures have been studied for a wide range of applied electric field. These parameters such as the drift velocity, mobility, diffusion coefficient, ionization coefficient and mean electron energy, which are defined in collision cross section and EEDF is represented the backbone of the electron swarm behavior of pure gas and gas mixtures in discharge of plasma [2].

Solution of transport Boltzmann equation for the electron –velocity distribution function $f(\vec{r}, \vec{v}, t)$ in spherical harmonics is given by [3].

$$f\left(\vec{r},\vec{v},t\right) = \sum_{\ell=0}^{\infty} f_{\ell}\left(\vec{r},\vec{v},t\right) p_{\ell}(\cos\theta) \quad (1)$$

where $f_{\ell}(\vec{r}, \vec{v}, t)$ represent coefficient of expansion and $p_{\ell}(\cos\theta)$ is Legendre Polynomials. By using two-terms approximation, the equation form become :

$$f_{0}(\vec{r}, \vec{v}, t) = f_{0}(\vec{r}, \vec{v}, t) + f_{1}(\vec{r}, \vec{v}, t) \cos \theta$$

$$, \qquad (2)$$

$$f_{0}(\vec{v}) = f_{0}(\vec{v}) + \frac{\vec{v}}{v}f_{1}(\vec{v}) \qquad (3)$$

first term $f_0(\vec{v})$, is isotropic The distribution in velocity space. This occurs in elastic collisions, where the electrons could be expected to suffer large directional changes of velocity but relatively small energy losses because of mass difference between electrons and molecules. The second term $\frac{\vec{v}}{v}f_1(\vec{v})$, the distribution is anisotropic because of large energy losses of electron in inelastic $f1{\mathord{\leftrightarrow}} f2$ collisions,

The electron energy distribution function in CF_4 gas discharge was numerically calculated for E/N=15-300Td using Boltzmann kinetic equation in 2006 by Masek [4]. The momentum transfer and vibration cross-section were derived from available experimental swarm parameters while the cross section for inelastic electron-molecule collisional processes were taken from the literature. The two terms spherical harmonic expansion of the distribution function was used in the calculations.

Niktovic and Potrovic in 2008 [5] are obtained results by using simple solution for Boltzmann equation in mixture of CF_4 with its radicals for ratios of the electric field to the gas number density E/N from (1Td – 1000 Td). The analysis of non – conservative collisions revealed a range of E/N where electron attachment introduced by radicals significantly changes electron kinetics obtained for pure CF_4 gas.

The Aim of this work is to study theoretically the electron energy distribution function and electron transport parameters in DC electric discharge processes for CF₄, Argon gases and its mixtures to different ratios from solution of Boltzmann equation by using two terms approximation.

Theory

The fundamental equation governing the electron distribution function is the Boltzmann equation. For spatially uniform gas in the presence of steady electric field. The Boltzmann equation for electrons in collisions state is given by [6]:

$$\frac{\partial f}{\partial t} + \mathbf{v} \quad \frac{\partial f}{\partial r} - \frac{e\vec{E}}{m} \frac{\partial f}{\partial \mathbf{v}} = \left(\frac{df}{dt}\right)_c \quad \dots (4)$$

where $\left(\frac{df}{dt}\right)_{c}$ represents the rate of change of $f\left(\vec{r}, \vec{v}, t\right)$ due to collision, thus, when collisions are considered. The total derivative $\frac{df}{dt}$ represents all particles moving in the phase space, where, the partial derivative $\frac{\partial f}{\partial t}$ represents the change in the number of electrons at a given point in the phase space. Now one considers the case of spatially uniform gas, also, $f\left(\vec{r}, \vec{v}, t\right)$ function of velocity only, where $\nabla_{r} f = 0$ and thus

equation (4) becomes:

$$\frac{\partial f}{\partial t} - \frac{e \vec{E}}{m} \nabla_{v} f = \left(\frac{d f}{d t}\right)_{c}$$
(5)

In weakly ionized gases, the most collisions occurrence between electrons and neutral atoms, the collisions between electron –electron and electron – ion are rare.

Consider now only neutral atoms collision. The contribution of collision of each kind to the change the distribution function can be quite simply added together. One will divide collisions into elastic and inelastic collisions :

$$\left(\frac{df}{dt}\right)_{c} = \left(\frac{df}{dt}\right)_{elastic} + \left(\frac{df}{dt}\right)_{inelastic}$$
(6)

The first part of equation (6) represents the change of function due to elastic collisions and the second part of equation represents the change of function because the inelastic collisions. The class of inelastic collisions do not include excitation process of atoms or molecules but also the creation of new electrons as a result of ionization.

After the compensation (two- terms approximation), we obtain two equations:

$$\frac{\partial f_0}{\partial t} - \frac{e\vec{E}}{m} \frac{1}{3v^2} \frac{\partial}{\partial v} \left(v^2 f_1\right) = \left(\frac{df_0}{dt}\right)_c$$
(7)

$$\frac{\partial f_{1}}{\partial t} - \frac{e \vec{E}}{m} \frac{\partial f_{0}}{\partial v} = \left(\frac{d f_{1}}{d t}\right)_{c} \quad (8)$$

When the electron collision frequency for momentum- transfer is much larger than the electron collision frequency for excitation, the necessary condition for the two terms expansion for distribution function to be valid , the momentumtransfer collisions play the major role in reducing the asymmetry in the distribution function .the function change in elastic collisions state is given by [7] :

$$\left(\frac{df_{1}}{dt}\right)_{c} = -\upsilon_{e} \left(\mathbf{v}\right) f_{1} \left(\vec{\mathbf{v}}\right)$$
(9)

 v_e is the electron momentum- transfer collision frequency where

$$\mathcal{D}_{e} (\mathbf{v}) = N \mathcal{Q}_{m} (\mathbf{v}) \mathbf{v}$$

$$\left(\frac{df_{\perp}}{dt}\right)_{c} = -N \mathcal{Q}_{m} (\mathbf{v}) \mathbf{v} f_{\perp} (\vec{\mathbf{v}})$$
, (10)

V represented electron velocity and $Q_m(v)$ is momentum transfer cross section.

From the solution of equation (8), we will find $f_1(v)$ value, where all quantities are assumed to independent of time, is given by:

$$f_{\perp}\left(\vec{\mathbf{v}}\right) = \frac{e\vec{E}}{\mathbf{v}\,m\,N\,\mathcal{Q}_{m}\left(\mathbf{v}\right)}\frac{\partial f_{0}}{\partial \mathbf{v}} \quad (11)$$

The change of function in elastic collisions is given by:

$$\frac{\partial f_0}{\partial t} - \frac{2e}{mv} \frac{\partial}{\partial \varepsilon} \left[\frac{\vec{E}^2 \varepsilon}{3NQ_m} \frac{\partial f_0}{\partial \varepsilon} + \frac{2m}{M} NQ_m \varepsilon^2 f_0 + \frac{2mK_B T}{Me} NQ_m \varepsilon^2 \frac{\partial f_0}{\partial \varepsilon} \right] = \left(\frac{df_0}{dt} \right)_{\mathcal{O}}$$

(12) The right hand side of equation (12) represented the combined effect on the distribution function of elastic and inelastic electron –neutral collisions only, assuming that the electron-electron collisions and electron-ion are negligible.

The change of the function
$$\left(\frac{df}{dt}\right)_c$$
 in

inelastic collisions is given by:

$$\left(\frac{df}{dt}\right)_{LOSS} = \sum_{j} N_{0} \left[f_{0} \left(\varepsilon + \varepsilon_{j}\right) Q_{j} \left(\varepsilon + \varepsilon_{j}\right) \left(\varepsilon + \varepsilon_{j}\right) - f_{0} \left(\varepsilon\right) Q_{j} \left(\varepsilon\right) \varepsilon \right] 2e / mv$$

$$. (13)$$

The energy gain from electric field in superelastic collisions is given by [6].

$$\left(\frac{df}{dt}\right)_{GAIN} = \sum_{j} N_{j} \left[f_{0}\left(\varepsilon - \varepsilon_{j}\right)Q_{j}\left(\varepsilon - \varepsilon_{j}\right)\left(\varepsilon - \varepsilon_{j}\right) - f_{0}\left(\varepsilon\right)Q_{j}\left(\varepsilon\right)\varepsilon\right] 2e / m_{T}$$

(14) Now, the sum of function change in two state (ground state and excitated state), after the equations compensation (12), (13) and (14), we obtain on work equation [7]:

$$\left(\frac{m\varepsilon}{2\epsilon}\right)^{l_{2}} \frac{\widetilde{g}_{0}}{\partial} = \frac{E^{2}}{3} \frac{\partial}{\partial \varepsilon} \left(\frac{\varepsilon}{NQ_{n}} \frac{\widetilde{g}_{0}}{\partial \varepsilon}\right) + \frac{2m\partial}{M\partial \varepsilon} \left(\varepsilon^{2}NQ_{n}f_{0}\right) + \frac{2mk_{B}T}{M\epsilon} \frac{\partial}{\partial \varepsilon} \left(\varepsilon^{2}NQ_{m} \frac{\widetilde{g}_{0}}{\partial \varepsilon}\right) + \sum_{j} \left[\left(\varepsilon + \varepsilon_{j}\right)f_{0}\left(\varepsilon + \varepsilon_{j}\right)N_{0}Q_{j}\left(\varepsilon + \varepsilon_{j}\right) - \varepsilon f_{0}\left(\varepsilon\right)N_{0}Q_{j}\left(\varepsilon\right)\right] + \sum_{j} \left[\left(\varepsilon - \varepsilon_{j}\right)f_{0}\left(\varepsilon - \varepsilon_{j}\right)N_{j}Q_{-j}\left(\varepsilon - \varepsilon_{j}\right) - \varepsilon f_{0}\left(\varepsilon\right)N_{j}Q_{-j}\left(\varepsilon\right)\right]$$

$$(15)$$

In right hand side of equation (15), the first part accounts for the gain of energy result from electric field. The second part of equation accounts for gain in energy (KT) and elastic losses by 2m/M. Also, in third part of the equation, few energy losses occur because of elastic scattering of electron due to electron collision with molecule. These, processes all occurs in ground state of molecules. The first term of excitation state, accounts energy losses of electrons result the inelastic collisions. The last part of equation (15), is the energy gain of electrons given by the electric field electrons to compensation the energy loss of electrons because the collisions.

When the left hand side of equation (15), equal zero, the function $f(\varepsilon)$ becomes isotropic.

Equation (15) is used in pure gases state, if the gas is mixture of different species. The momentum transfer cross section, excitation cross section and super elastic for molecules of species (k) in inelastic processes are represented by Q_m^k , Q_j^k , Q_{-j}^k respectively. Also, the electron collision frequency in elastic collisions state for molecules of species k (mixtures) become: v_e^k (v) = $N_K Q_m^K$ (v) v (16)

where N_{K} is the number of molecules of species k.

Transport coefficients

The drift velocity is nonlinear function with electric field, and the mobility depends on strength field. at sufficiently low E/N, where an electron loses all equal to the gain from the electric field at one elastic collision , the drift velocity is proportional to E/N.

The relation between drift velocity and distribution function of electron energy is given by [8]:

$$\mathbf{v}_{d} = -\frac{E}{3} \left(\frac{2e}{m}\right)^{1/2} \int_{0}^{\infty} \frac{u}{N Q_{m}} \left(u\right) \frac{df_{0}}{du} du$$
(17)

The mobility is defined as the proportionally coefficient between the drift velocity of a charged particle and electric field. The mobility of electrons is:

$$\mu_e = \frac{e}{m \upsilon_m} = \frac{\mathbf{v}_d}{E} \tag{18}$$

Where v_m represent the electron momentum- transfer collision frequency. The electron mobility decreased with E/N increase, this occur energy loss result of electron through the collisions between electrons and neutral molecules.

From the relation between the drift velocity and mobility with EEDF, we can be calculate electron mobility equation [9]:

$$\mu_{e} = -\frac{1}{3} \left(\frac{2e}{m}\right)^{1/2} \int_{0}^{\infty} \frac{u}{\sum \delta_{s} \mathcal{Q}_{m}} \left(u\right)^{\frac{df_{0}}{du}} du$$
(19)

where δ_s represents fractional concentration of the s species $\left(\delta_s = \frac{N_s^{j}}{N_s}\right)$ and N_s^{j} is the number of molecules of

species s in the excited state j. when the density of charged particles is very low, the charges of opposite signs diffuse independently of each other, This

is known as free diffusion. In a weakly ionized gas discharge, the ambipolar diffusion coefficient is simplified by noting that $\mu_e \rightarrow \mu_i$ usually. The relation between diffusion coefficient and electron energy distribution function is given by [10]:

$$D_{T} = \frac{1}{3} \left(\frac{2}{m}\right)^{1/2} \int \frac{u^{1/2}}{N Q_{m}(u)} f(u) du$$
(20)

 Table (1): the number densities values and mass density of pure gases

Gas	Mass density (10 ⁻³ gm /cm ³)	Number of density(10 ¹⁹ atom/ cm ³)
CF ₄	3.85	2.640161 [13]
Ar	1.783	2.68772 [14]

An electron colliding with a neutral atom can produce a negative ion in a process termed electron attachment. Dissociative and non dissociative electron attachment reactions depend strongly on the electron energy

The primary ionization coefficient is a basic parameter in discharge physics and is defined as the number of ionizing collisions made by an electron in moving 1cm in the direction of the applied $\frac{\alpha}{N}$ electric field. The coefficient is used in describing the behavior of a swarm of electrons traveling through a gas [11].the ionization coefficient is calculate from the relation:

$$\frac{\alpha}{N} = \frac{1}{v_d} \left(\frac{2e}{m}\right)^{1/2} \sum_k \int_{uik} \frac{N_K}{N} Q_{ik} (u) u f_0 (u) du$$
(21)

is called ionization coefficient. N_{K} represent the number of atoms of type k

method of calculation

The work deals with the theoretical studies about electron energy distribution function and electron transport parameters in DC electric discharge processes for CF_4 , Ar gas and its mixtures with different ratios at low temperature of the plasma.

The search dependent upon solve the equation of Boltzmann for collisions by two-terms approximation method in gas discharge processes at regular uniform electric field by using NOMAD computer program in Fortran language.

The electron energy distribution function and transport parameters was calculated for range (1 -500) Td with NOMAD program by input data for this program such as collision cross section, the number density of gases, number and types of gases, concentration of gas, distribution density, E/N represent the ratio between electric field intensity to the number density of gas, molecular weight, gas temperature, electron density, as well as the program contain other condition must be taken into account this work like, electron energy distribution function dependent of collision cross sections.

The number density of gas was calculated theoretically depended on weight density and molecular weight, the relation between these rewrite [12]:

$$N = \frac{\rho N_A}{\omega}$$
(22)

Table (2): The number of density for

Gas Mixtures	Mass	Number of	
	density	density	
	(10	$(10^{19} \text{atom}/$	
	³ gm	cm^{3})	
	$/cm^{3})$		
90%CF ₄ +10%Ar	3.650	1.718	
75%CF ₄ +25%Ar	3.339	1.572	
50%CF ₄ +50%Ar	2.820	1.328	
25%CF ₄ +75%Ar	2.742	1.293	
10%CF ₄ +90%Ar	1.990	0.937	
mixtures calculated in this work.			

where ρ represent weight density, ω is molecular weight, N_A represent Avogadro constant. The table (1) shown that the number density of gases and mass density in pure gases (CF₄, Ar) at room temperature.

Results and Discussion

Carbon tetrafluoride - Argon mixtures

The transport Boltzmann equation was solved for CF_4 –Ar mixtures of different ratios. The electron swarm parameters in different mixtures have been analyzed for

range of E/N values $(1 \times 10^{-17}V \ cm^2 \le E \ / N \le 5 \times 10^{-15}V \ cm^2)$. The different ratios mixtures of CF₄-Ar gases are listed in Table (3-1).

The influence of different discharge parameters on the electron distribution function is shown in figures (1 to 3). The electron energy distribution function strongly affected by changing either the parameter E/N or gas mixtures. The electron energy distribution function is nearly Maxwellian at E/N = 1-2Td in most mixtures but the electron energy distribution function is non Maxwellian at values of $E/N \ge 5Td$ because the electrons lose energy in inelastic collisions with atoms in the mixture.



Fig. (1): The EEDF as a function of electron energy for several values of E/N in







Fig. (3): The EEDF as a function of electron energy for several values of E/N in 10% CF₄ + 90% Ar gases mixture



Fig.s(4) The EEDF as a function of energy for different ratios gases mixture of Mixtures where E/N value is constant

Figure (4) shows the change of distribution function for different ratios of CF₄-Ar gases mixture at constant value of $E / N = 5 \times 10^{-16} V \ cm^2$. The electron energy distribution function is minimum at low ratio of CF₄ and the probability of collisions is small and increase in electrons energy from the electric field. The energy

gain from electric field will be distributed on number of electrons, and therefore the



Fig. (5) The mean electron energy as a function E/N in different ratiosCF₄-Ar mixtures.

electron energy is higher in less collisions region.

The variation of the mean electron energy with E/N for different of the mixtures components is shown in figure (5). The mean electron energy of pure Argon is higher than other gases and mixtures (number of collisions is lower).

Figure (6) shows the characteristic energy for different ratios of mixtures, the higher value of thermal energy occurs in the ratio 10% $CF_4 + 90\%$ Ar compared with other ratios and it is lower than the value in pure CF_4 .



Fig. (6) The characteristic energy as of E/N in different ratios of gas mixtures (CF_4 -Ar).

The mobility of electron as a function of E/N is shown in figure (7), we denote at Argon gas the mobility is lower than that of CF_4 -Ar gases mixture because the electron energy in argon gas is higher than that with mixtures, and therefore the probability of collisions is larger in argon gas and the mobility in mixtures is very few at higher E/N value, and process is reversed in weak electric fields with low probability of collisions.



Figure(7): The mobility of electron as electron as a function of E/N in different ratios of mixtures (CF₄-Ar).



Figure(8): The ionization coefficient of a function of E/N in different ratios of mixtures (CF_4 -Ar).

Figure (8) shows the ionization coefficient as a function of E/N. This illustrates that at the low ratio of CF_4 mixture, the ionization coefficient is high, also, the ratios of negative ions becomes lower.

References

- [1] Y. Itikawa, " Molecular processes in plasma ", Tokyo, Japan (**2007**).
- H. Date and H. Sakai, "Boltzmann equation analysis of electron collision cross section and swarm parmeters for krypton", J. phys. D: Appl. Phys., Vol.<u>22</u>, P.1478-1481 (1989).
- [3] K. Smith and R. W. Thomson, "Computer Modeling of Gas", Plenum Press, New York (**1978**).
- [4] K. Masek, L. Laska, Inter science, Vol. <u>27</u>, P.15-21 (**2006**).
- [5] Z. D. Niktovic and V. D. Stojanovic," Modeling of high *E/N* in mixtures of CF4 and its radicals", Publ. Astron. obs, Belgrad,. No. <u>84</u>, 103-106 (**2008**).
- [6] T. Holstein," Energy distribution of electrons in high frequency gas discharges", Phys. Rev., Vol. <u>70</u>, No. 5-6 (1946).

- [7] S. R. Hussein, PhD thesis, "Solution of Boltzmann equation using two terms approximation for several SF₆&other gases mixture" University of Salahaddin / Erbil (2007).
- [8] W. L. Nighan, "Electron energy distribution and collision rats in electrically excited N₂, CO and CO₂," Phys., Rev., Vol. <u>2</u>, P. 1989-2000 (1970).
- [9] C. Truesdell, J. Chem. Phys, Vol. <u>37</u>, P. 2336 (1962).
- [10] T. Makabe and Z. petrovic, "Plasma Electronics: Application in Microelectronics Device Fabrication", Taylor and Francis Group, New York, (2006).
- [11] Y. P. Raizer, " Gas discharge physics ", Sepringer- Verlag (**1991**).
- [12] J. M. Meek and J. D. Craggs, "Electrical Break down of gases", Ch 1, p. 33, (John Wiley & Sons, 1978).
- [13] T. Nahi and A. Hamadi, " Atomic physics " Ch. 2, P. 105, (1980).
- [14] N.N. Greenwood and A. Earnshaw, "Chemistry of the element", 2nd, Eidition, Oxford (1997).
- [15] J. F. Ziegler, "Stopping cross section for energetic ions in all elements", Book, Vol. <u>5</u>, Pergamon Press Inc, (1980).