# The Photon-Assisted Cascaded Electron Multiplier Operation in CF<sub>4</sub> for Ion Backflow Suppression

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Abstract—The operation of the recently introduced Photon Assisted Cascaded Electron Multiplier (PACEM) in CF<sub>4</sub> is investigated. The PACEM uses the VUV scintillation produced in the electron avalanches of the first multiplier of the cascade to transfer the signal to the subsequent ones. The VUV scintillation induces the emission of a large number of photoelectrons from a CsI photocathode placed on the top-surface of the second multiplier. The photoelectrons are further multiplied in the subsequent stages of the cascade, resulting in efficient signal amplification. A mesh, set at a fixed voltage, is placed between the first and the second multipliers to block the charge transfer between them, thus suppressing all the ion backflow (IBF) heading to the first cascade element. The PACEM electrically isolates the first multiplier of the cascade and only the ions produced in the electron avalanches of the first element may flow back into the drift region. Operating in CF4, absolute IBFs as low as  $\sim 1$  ion per primary electron are achieved for drift fields of 0.1 kV/cm, while for fields of 0.5 kV/cm the absolute IBF is  $\sim 10$  ions/primary electron. This corresponds to an IBF of 10<sup>-4</sup> at gains of 10<sup>4</sup>, for Time Projection Chambers (TPC) operating conditions, and IBFs of  $\sim 10^{-5}$  at gains of  $10^6$  for Gaseous Photomultipliers (GPM) operating conditions.

## I. INTRODUCTION

THE Photon-Assisted Cascaded Electron Multiplier (PACEM) [1] was developed as an alternative for blocking avalanche-induced ions in cascaded gaseous electron multipliers. The avalanche originated in the first element propagates to the subsequent ones solely via its photons which, in turn, induce photoelectrons in the next multiplier of the cascade, while the charge transfer between the first and second cascade elements is blocked by a mesh electrode placed between them. This optical coupling is only effective in highly scintillating gases such as the heavy noble gases [2], [3], excluding applications in mixtures with quenching gases [2]. The PACEM principle and operational characteristics were demonstrated in xenon [1], [3]; by a proper choice of the gas, geometry and electric fields, the photon-assisted propagation can be made very efficient, while the charge transport between the first and the second element is completely blocked [3].

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Amongst the molecular gases, CF<sub>4</sub> is known to have a high scintillation output, either under bombardment by ionizing radiation [4] or produced in electron avalanches [5], having a significant component in the VUV region (peak around 170 nm) [4]. In addition, it is an important gas with many applications in high energy physics instrumentation and neutron detection. However, the scintillation produced in electron avalanches in CF<sub>4</sub> is lower than that achieved in xenon, which can be a drawback to the implementation of the PACEM concept in CF<sub>4</sub>. On the other hand, the collection efficiency of photoelectrons emitted by solid photocathodes in gaseous atmosphere is a few times higher in  $CF_4$  than in xenon [7], [8]. This effect is mainly due to the existence of vibrational excitation states in CF<sub>4</sub> that, at low electron impact energies, can compete efficiently with elastic scattering. As a result of the presence of this additional mechanism the electron energy may be reduced after just a few collisions to values that decrease the probability of returning to the photocathode [9]. This could compensate the lower scintillation output and result in a good performance of the PACEM in CF<sub>4</sub>.

In the present work, the operation of the PACEM in pure  $CF_4$  is investigated. The optical gain, i.e., the number of photoelectrons transferred through the second cascade element per primary electron, and the relative IBF, i.e., the number of ions flowing back to the drift region per primary electron, are studied as a function of applied voltages.

# II. ION BACKFLOW IN GASEOUS ELECTRON MULTIPLIERS

The positive ions produced in electron avalanches of gaseous detectors may present a serious drawback to the detector performance. These ions slowly flow back into the conversion/drift region of the detector and pose limitations to its operation. In particular, the trapping of the ions in gaseous detectors is of particular relevance for the development of Time Projection Chambers (TPC) and Gaseous Photomultipliers (GPM).

In TPCs, the presence and accumulation of ions in the drift region can disturb the electric field, resulting in dynamic track distortions [10]. This effect is most important in high-multiplicity experiments, e.g., relativistic heavy-ion physics applications [11]. GPMs equipped with bi-alkali photocathodes for visible light detection are even more sensitive to the ion back flow; ions impinging the photocathode have a high probability of inducing secondary emission of electrons, creating a mechanism of positive feedback that limits the gain [12]. In addition, ions reaching the photocathode are also a cause for faster aging, limiting the operation of the GPM for extended periods of time.

The fraction of ions that reach the drift region relative to the total number of electrons collected at the anode of the detector

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Fig. 1. Schematic view of the PACEM detector in a MHSP/GEM configuration.

is defined as the Ion Back Flow (IBF). A goal for IBF with values of  $10^{-4}$  for gains of  $10^4$  and of  $10^{-5}$  for gains of  $10^6$  has been established for TPCs [13] and GPMs [12], respectively. Short overviews of different methods and results obtained for IBF reduction are presented in [12] and [14]. So far, the lowest achieved values for IBF,  $1.5 \times 10^{-4}$  for a detector gain of  $10^4$ at a drift field of 0.2 kV/cm and  $3 \times 10^{-4}$  for a detector gain of  $10^5$  at a drift field of 0.5 kV/cm, were obtained in DC mode by electrostatic deviation and capture of the ions using a Micro Hole and Strip Plate (MHSP0) cascaded with Gas Electron Multipliers (GEM) [16], [17]. With this technique, a large amount of the total number of ions produced in the detector is trapped, and only a small fraction flows back to the drift region. The drawback of this method is that the absolute number of ions that reach the drift region increases when the gain of the detector increases, as more ions are produced.

A different approach to the IBF blocking was introduced with the Photon Assisted Cascaded Electron Multiplier (PACEM) detector [1], [3]. In this detector, only a fraction of the ions produced in the first cascade element will flow back to the drift volume. The ions produced in the last elements of the detector are completely blocked from flowing in to the drift region and the IBF is independent of the detector total gain. The operation of the PACEM in 1 bar of xenon has reached IBF values of  $\sim 2 \times 10^{-4}$  for a detector gain of  $10^4$  and a drift field of 0.1 kV/cm and  $10^{-5}$  for a detector gain of  $10^6$  and a drift field of 0.5 kV/cm [3].

## III. EXPERIMENTAL METHOD AND SETUP

The PACEM used in this work is presented in Fig. 1, in a MHSP/GEM configuration, and is the one used in [3]. The MHSP and the GEM share the same production technique and are both made of a micro-perforated 50  $\mu$ m Kapton<sup>®</sup> foil with a 28 × 28 mm<sup>2</sup> active area, covered on both sides with a 5  $\mu$ m copper layer, gold coated. The main difference between these two structures is the strip pattern present on the bottom face of



Fig. 2. Top (upper view) and bottom (lower view) side of the MHSP.

the MHSP. These strips are of two different and independent kinds: 100  $\mu$ m wide cathode strips, perforated with 70  $\mu$ m diameter biconical holes along the centre, and thinner, 20  $\mu$ m wide anode strips etched between the cathode strips, Fig. 2. With this arrangement the MHSP presents two independent electron multiplication regions: inside the holes, which is established by the voltage difference between the top electrode and the cathode strips of the MHSP, V<sub>CT</sub>, and in the region around the anode strips, which is established by the voltage difference between cathode and anode strips, V<sub>AC</sub>. Compared to the GEM, the MHSP allows for an extra multiplication stage, resulting in higher scintillation and charge gains [1], [18]. In addition, depending on the electric field between the MHSP and the wire mesh, only ~20% of the anode-avalanche ions may drift back through the MHSP holes [19].

The scintillation produced in the avalanches of the MHSP induces the extraction of photoelectrons from the reflective CsI photocathode (2500 Å thickness) deposited on the top of the GEM. The stainless steel grid (80  $\mu$ m diameter wires, 900  $\mu$ m spacing), G<sub>1</sub>, is connected to ground potential, blocking all the charge transfer between the MHSP and the GEM. The photoelectrons emitted by the CsI photocathode are transferred into the GEM holes, where they undergo multiplication. The avalanche electrons extracted from the holes are collected in the GEM bottom electrode. The reflective CsI Photocathode has very low sensitivity to charged-particles background, as discussed in [20].

For the measurements presented on this paper only the photoelectron current emitted by the reflective photocathode was recorded and there was no further photoelectron multiplication at the GEM holes. The grid  $G_1$  was polarized while the top GEM electrode was at ground potential resulting in a extraction field of 1.0 kV/cm. The detector was filled with  $CF_4$  up to the pressure of 1 bar and operated in sealed mode. Prior to the filling, the detector was pumped down to a pressure of  $10^{-6}$  mbar. During the operation of the detector, the  $CF_4$  was continuously purified using non-evaporable getters, SAES St707. All the electrodes of the detector were independently biased. The present studies were performed operating the PACEM in current mode.

The primary electron current,  $I_{PC0}$ , is due to the photoelectron emission from the semitransparent photocathode placed above a 10 mm long drift region as shown in Fig. 1. This photocathode is composed by a 250 Å CsI layer deposited on a 5 mm thick quartz window, pre-coated with a 100 Å Al layer to ensure electrical contact trough the whole surface of the photocathode. An Hg(Ar) UV lamp is used to produced the primary electrons that are extracted from the photocathode into the gas by the electric field in the drift region. Since the semitransparent photocathode is grounded, the electric field is determined by the voltage of the MHSP top electrode. The currents on the semitransparent and reflective photocathode were measured using Keithley 610C electrometers.

We define the optical gain as the ratio between the number of photoelectrons extracted from the reflective photocathode due to the gas scintillation and the number of primary electrons emitted by the semitransparent photocathode. As the current measured at the reflective photocathode,  $I_{\rm RP}$ , includes the contribution from both photoelectrons induced by the gas scintillation and photoelectrons induced by the direct exposure of the reflective photocathode to the UV photons emitted by the Hg lamp,  $I_{\rm RP0}$ , the optical gain is given by

Optical Gain = 
$$\frac{I_{\rm RP} - I_{\rm RP_0}}{I_{PC_0}}$$
. (1)

Both  $I_{RP0}$  and  $I_{PC0}$  are measured for null electric fields in the MHSP, i.e.,  $V_{CT} = V_{AC} = 0$  V, and in the GEM, i.e.,  $\Delta V_{GEM} = 0$  V, and extraction field of 1.0 kV/cm in the region between the reflective photocathode and the wire mesh G<sub>1</sub>. This value was found to ensure good extraction efficiency in the gas, above 90% [7].

The optical gain depends, amongst other factors, on the scintillation produced in the gas, on the photocathode quantum efficiency, on the solid angle subtended by the photocathode and on the photoelectron collection efficiency.

We define the absolute IBF as the number of ions flowing back to the drift region per primary electron, IBF/pe. In our setup, the absolute IBF is obtained by measuring the current at the semitransparent photocathode,  $I_{\rm PC}$ . This current includes both the primary photoelectron current emitted by the photocathode and the current due to the positive ions produced in the MHSP avalanches that flow back to the drift region,  $I_{\rm ions}$ . Therefore

$$\frac{\text{IBF}}{pe} = \frac{I_{\text{ions}}}{I_{\text{PC}_0}} = \frac{I_{\text{PC}} - I_{\text{PC}_0}}{I_{PC_0}}.$$
(2)

The IBF is obtained dividing the absolute IBF by the total gain of the cascade multiplier.

#### IV. EXPERIMENTAL RESULTS AND DISCUSSION

We have investigated the optical gain as a function of voltage difference between cathode and anode strips of the MHSP,  $V_{AC}$ , for different values of voltage difference applied to the MHSP



Fig. 3. PACEM optical gain in CF\_4 as a function of  $\rm V_{AC}$  for different  $\rm V_{CT}$  voltages and for  $\rm E_{Drift}$  = 0.1 kV/cm.



Fig. 4. PACEM optical gain in CF4 as a function of  $\rm V_{AC}$  voltage for  $\rm V_{CT}$  voltage of 500 V and different drift electric fields.

holes,  $V_{CT}$ , Fig. 3, and for different drift electric fields, Fig. 4. The results of Figs. 3 and 4, at low  $V_{AC}$  values, show that a GEM as the first element of the PACEM will not lead to optical gains high enough for an effective operation of the PACEM, presenting optical gains well below 1. Values similar to those achieved with the PACEM operating in GEM mode ( $V_{AC} = 0$  V) were obtained by the authors of [2], using a GEM coupled to a CsI photocathode in a Kr atmosphere. The additional charge and scintillation gain achieved in the anode strips of the MHSP results in a fast increase in optical gain, which may reach values as high as 100.

Compared to the same conditions of operation in xenon, the optical gains obtained in  $CF_4$  are lower, denoting the lower scintillation output achieved in  $CF_4$ . Nevertheless, values above 10 are achieved for  $V_{CT}$  and  $V_{AC}$  above 450 V, and maximum values achieved are comparable of those obtained when operating in pure xenon. The results of Fig. 4 also show that the optical gain is almost independent on the drift electric field.

To ensure that no primary charge is lost during the first multiplication stage, a relatively large gain on this stage is required (>10). On the PACEM detector this takes place at the MHSP holes, where, for the operational conditions presented on this paper, the gain is above 70 [21]. In addition, during previous work done with the PACEM detector operating at pulse mode in xenon, no energy resolution degradation from the first to the



Fig. 5. Absolute IBF, ions reaching the semitransparent photocathode per primary electron, as a function of the PACEM optical gain for  $V_{\rm CT}=500$  V and different drift electric fields (a) and for drift field of 0.1 kV/cm and different  $V_{\rm CT}$  voltages (b).

second stage was observed [1], confirming that the gain on the first multiplication stage is large enough to assure full detection efficiency of the primary charge.

The absolute IBF was investigated as a function of optical gain for drift electric fields of 0.5 and 0.1 kV/cm (GPMs and TPCs operating conditions, respectively), Fig. 5. For GPM operating conditions,  $E_{Drift} = 0.5$  kV/cm, 24 ions per primary electron were measured to reach the photocathode at an optical gain of 12. The operation of the PACEM with a triple-GEM coupled to a reflective photocathode [7], [22] as the second stage of the cascade multiplier may achieve a total gain of  $10^6$  with an IBF value of  $2.4 \times 10^{-5}$ . The same consideration can be made for the TPC operating conditions ( $E_{Drift} = 0.1 \text{ kV/cm}$ ) for which we have measured 2.5 ions per primary electron at an optical gain of 13. This value results in an IBF of  $2.5 \times 10^{-4}$  for total gains of  $10^4$ . These figures are very close to the goal established for the operation conditions of these devices. Comparing to the PACEM operation in xenon, the operation in  $CF_4$  leads to the similar levels of ion blocking capability.

#### V. CONCLUSION

We have demonstrated the capability of the PACEM operating in CF<sub>4</sub>. The obtained results show that CF<sub>4</sub> presents good scintillation and photoelectron extraction capabilities enabling to achieve optical gains as high as 100. The ion blocking capability of the PACEM in CF<sub>4</sub> allows the reduction of the number of ions flowing back to the drift region per primary electron to 2.5 and 24 in TPC conditions ( $E_{Drift} = 0.1$  kV/cm) and GPM conditions ( $E_{Drift} = 0.5$  kV/cm), respectively. This corresponds to IBF values of  $2.5 \times 10^{-4}$  and  $2.4 \times 10^{-5}$  at optical gains of about 12 for drift electric fields of 0.1 and 0.5 kV/cm and total cascade gains of  $10^4$  and  $10^6$ , respectively. The obtained results are similar to those achieved for the PACEM operation in xenon, denoting that the lower scintillation output achieved in CF<sub>4</sub> is compensated by the higher transmission efficiency of photoelectrons in CF<sub>4</sub>.

Future work in this area will involve the study of the pulse mode operation of the PACEM in  $CF_4$  and the study of the PACEM in other gases with high scintillation yields.

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