

# INSTRUMENTATION DAYS ON GASEOUS DETECTORS

## Streaming phenomena in gases

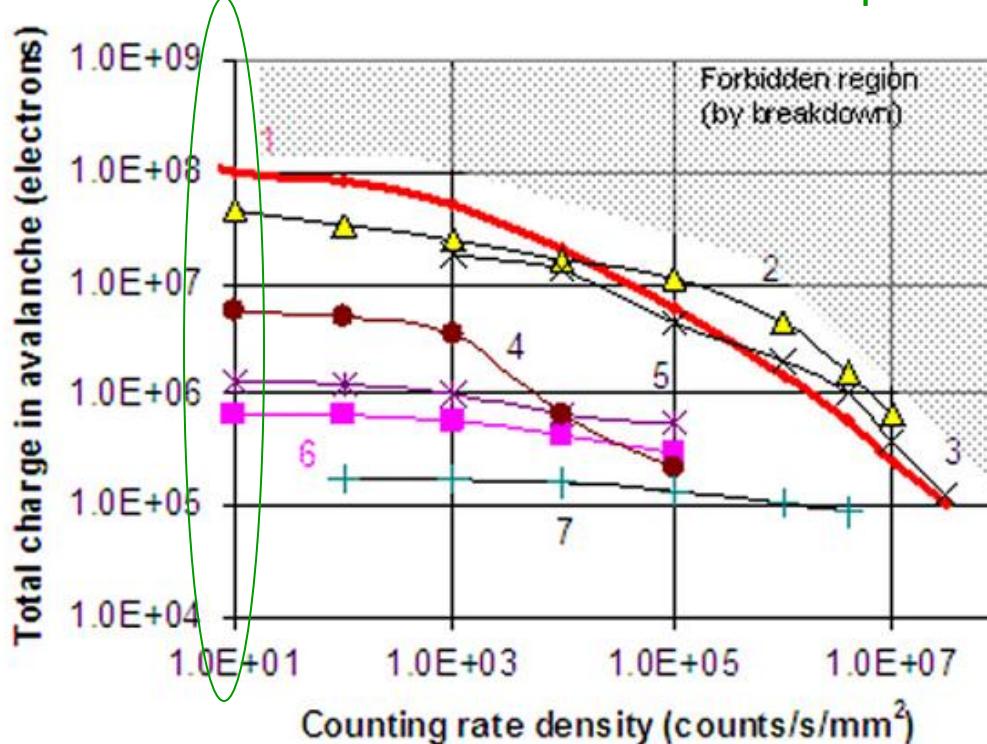
P.Fonte



RD51

# Fundamental detector limits

this work: low rate limit  $\Rightarrow$  space-charge limit

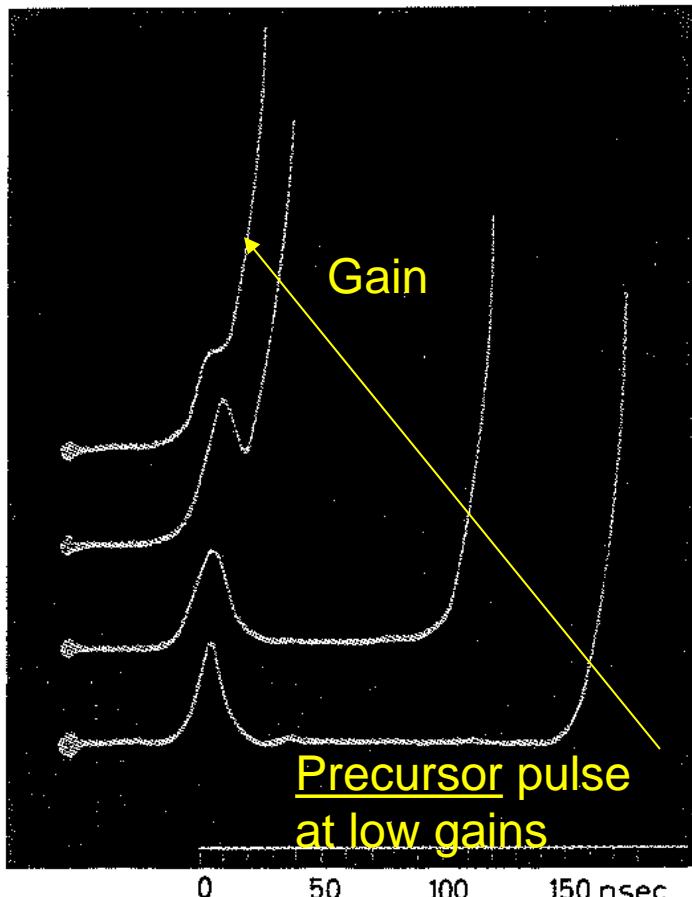


**Figure 4.** The maximum achievable gain, limited by breakdown, as a function of the x-ray flux for various detectors: (1) PPAC with 3 mm gap; (2) MICROMEGAS; (3) PPAC with 0.6 mm gap; (4) microstrip gas chamber with 1 mm strip pitch; (5) microstrip gas chamber with 0.2 mm strip pitch; (6) GEM; (7) microgap detectors with 0.2 mm strip pitch. Large counting rate densities require a reduction in the gas gain to prevent breakdown. See [52–55] and references therein for the original data and details. All data were converted to total avalanche charge.

# Fast breakdown - experimental evidence

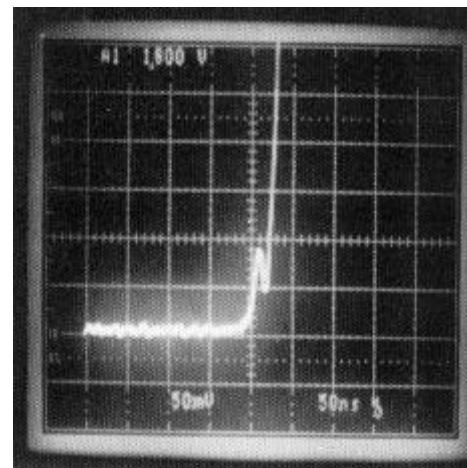
Very fast process featuring a “precursor” pulse

[RAE64]

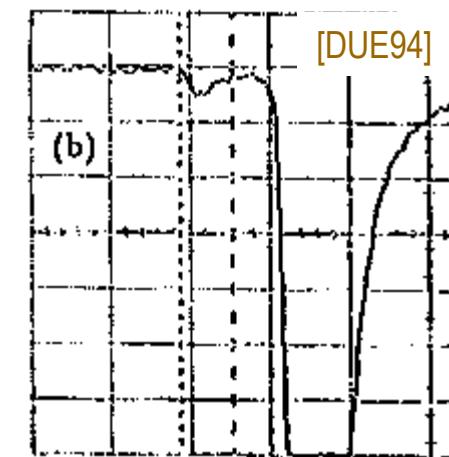


PPAC

[FON91]



RPC



A signature of low-gain cathode streamer-only breakdown

single-wire (SQS mode)

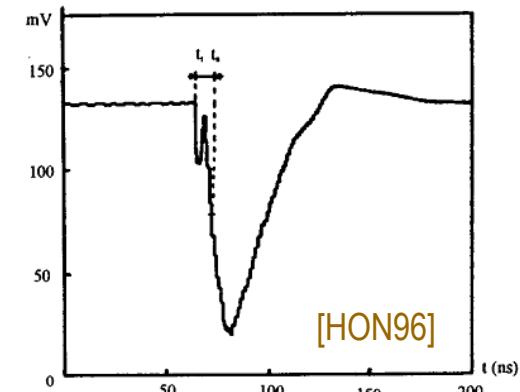


Fig. 1. The pulse shape of the SQS electrical signal  $V = 2.45$  kV. Methylal/(Methylal + Ar) = 16.6%.

Figure 5.14. Current oscillograms of static breakdown in methylal. Optical method.  $E/p = 64.4$ ,  $pd = 230$  Torr cm,  $d = 0.8$  cm,  $T_+ = 90$  nsec  $RC = 5$  nsec<sup>36</sup>

# Fast breakdown - experimental evidence

Cloud chamber observations (vapours, ~1cm gap)

High gain – anode and cathode streamers

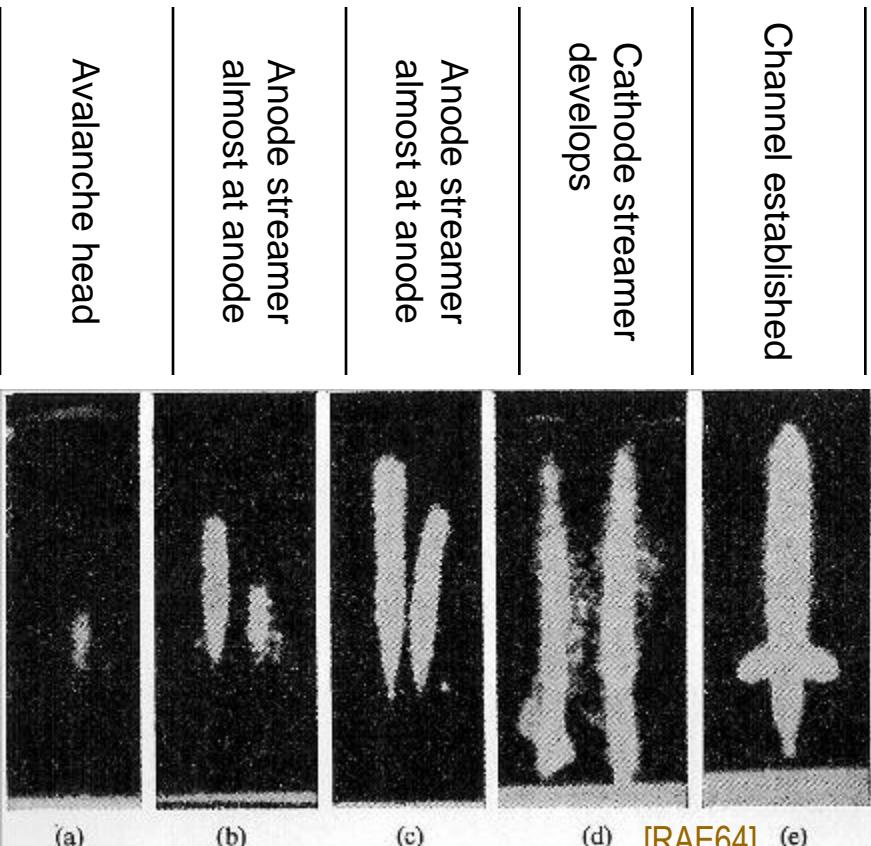


Figure 5.9. Development of one avalanche into a streamer, photographed in the cloud chamber (air, 270 Torr). The expansion ratio was reduced, so that in (a) only the head of the avalanche, as the region of the highest ion density, is visible as a track. If the voltage is slightly raised (at constant voltage pulse duration), an ‘anode directed’ streamer develops out of the avalanche head (b, c). Further increase of the voltage produces the development of the ‘cathode directed’ streamer, so that a plasma channel bridges the two electrodes. Therein occurs the spark. The same stages pass, if the voltage pulse height remains constant and the pulse duration is increased<sup>16</sup>.

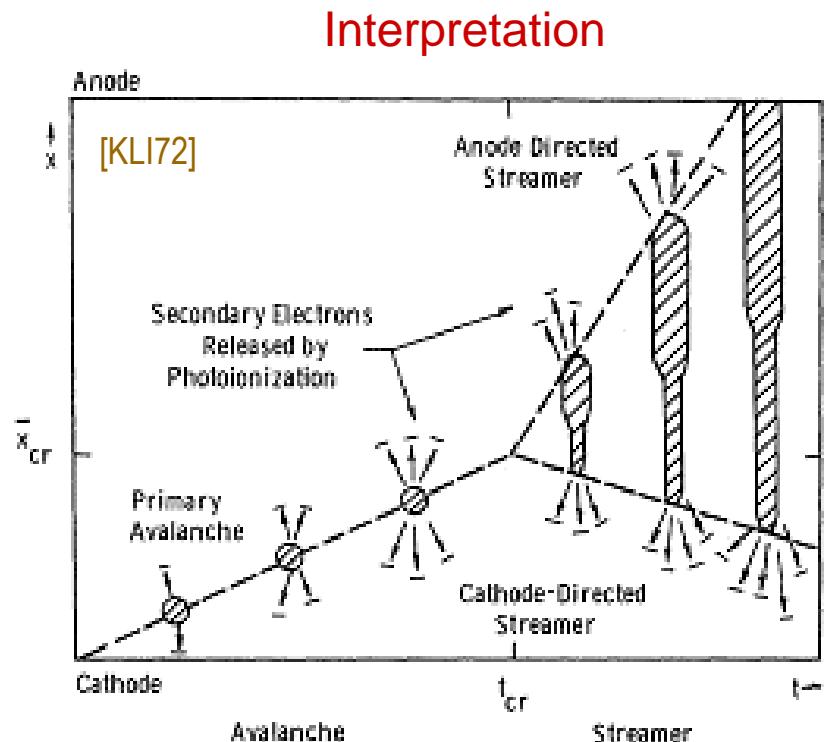


FIG. 6. Schematic representation of the qualitative description of streamer development given by Wagner. (Based on Figs. 22 and 27 of Ref. 11.) Anode- and cathode-directed streamer propagation begins at  $t_{critical}$  when the avalanche position equals  $x_{critical}$ .

# Fast breakdown - experimental evidence

Lower gain – only cathode streamer

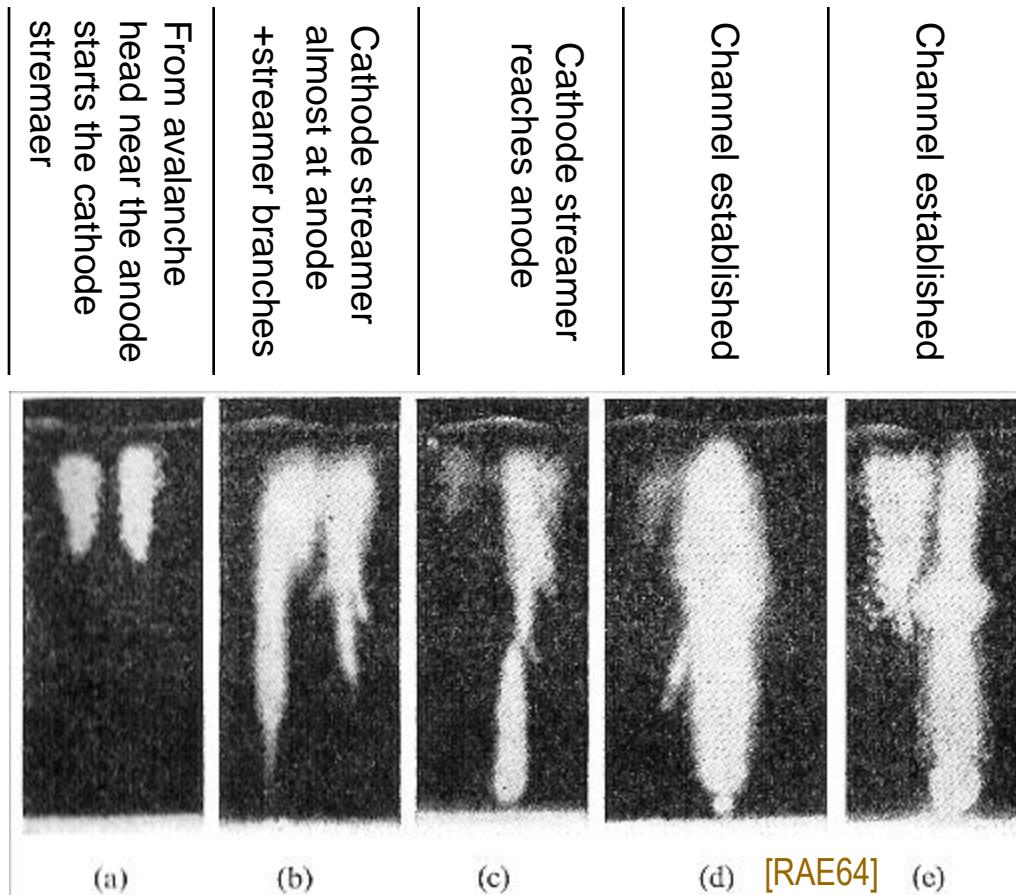
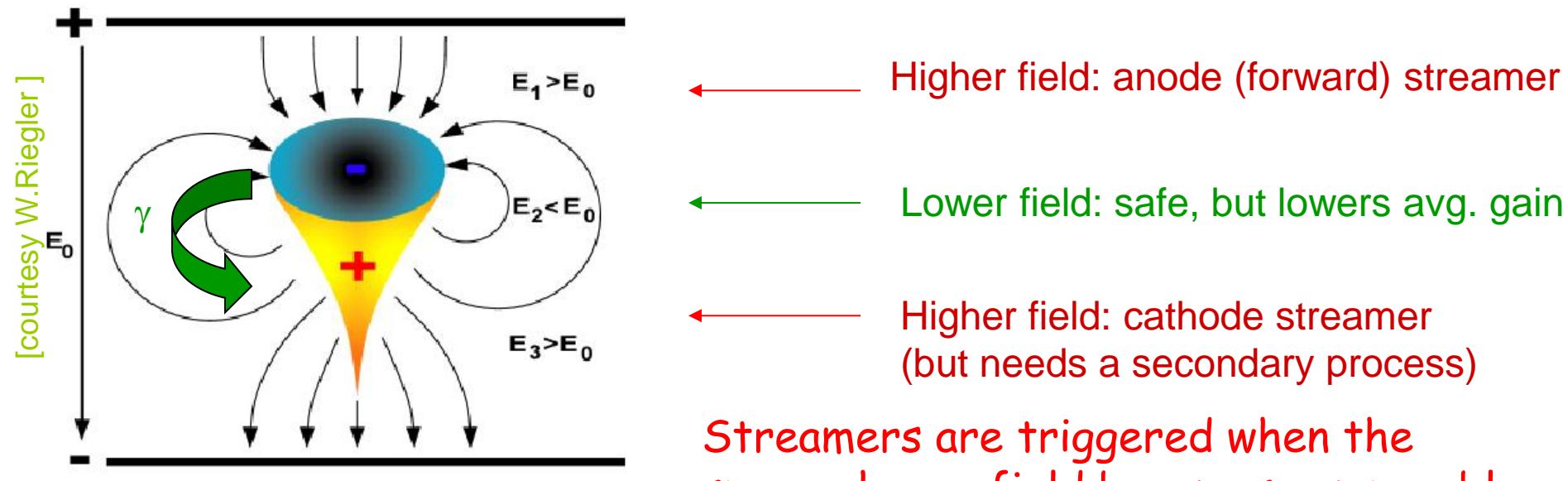


Figure 5.10. Development of the cathode directed streamer (with increasing pulse length). From the head of the avalanche (near the anode) (a) starts the cathode directed streamer ((b) (c)) till a plasma channel connects cathode and anode (d, e). These branched streamers resemble the discharge figures going out from a positive point, see Figure 5.12<sup>16</sup>

# Fast breakdown – accepted physical origin

(Meek and Raether's “streamer”/“Kanalaufbau” mechanism)

**Photon-mediated local feedback in a strong space-charge field**



Streamers are triggered when the space-charge field becomes comparable to the applied field:

a charge-dominated,  
geometry-dependent process.

**Complex physical process, involving:**

- electron transport in variable fields
- electron multiplication in high fields
- space-charge distorted electric field
- emission of photons able to photoionize the gas at a certain distance (gas self-photoionization)

# Raether limit – parallel fields

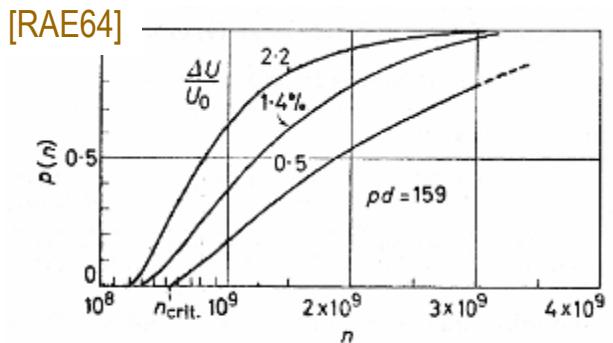
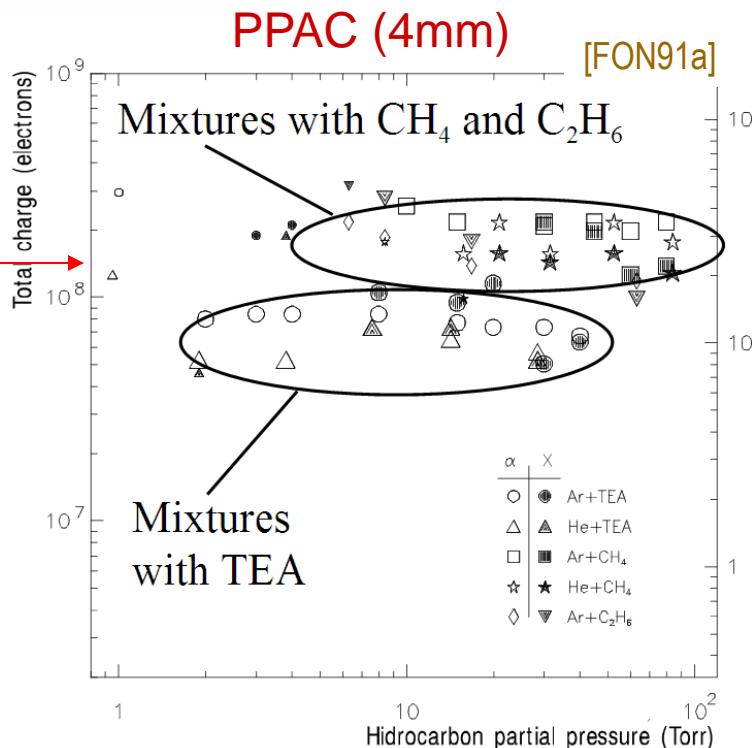
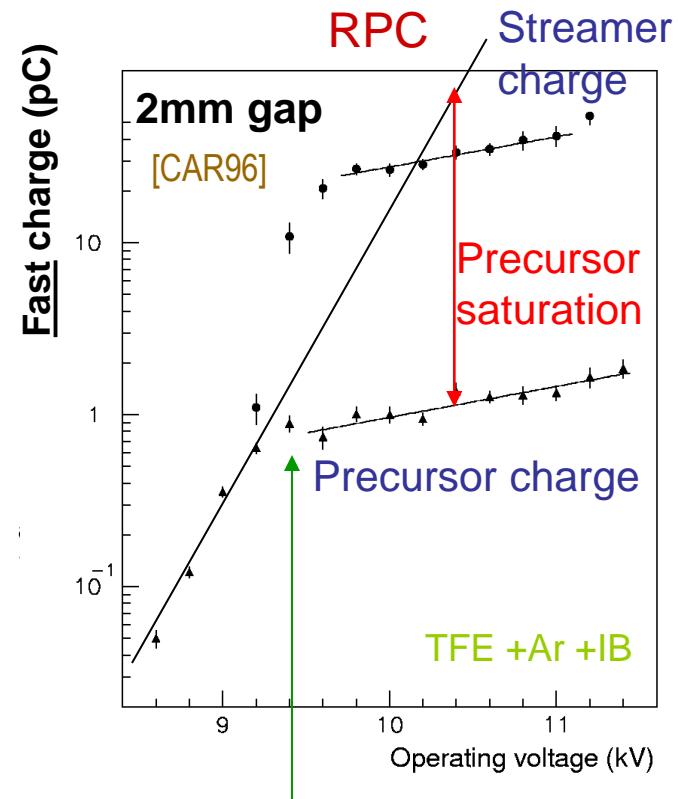


Figure 5.16. Probability  $p(n)$  that an avalanche produces a streamer as function of  $n$ ; ether,  $d = 0.8$  cm,  $p = 199$  Torr<sup>37</sup>

A charge-dominated process



The famous  
“Raether limit”  
of  $\sim 10^8$  electrons

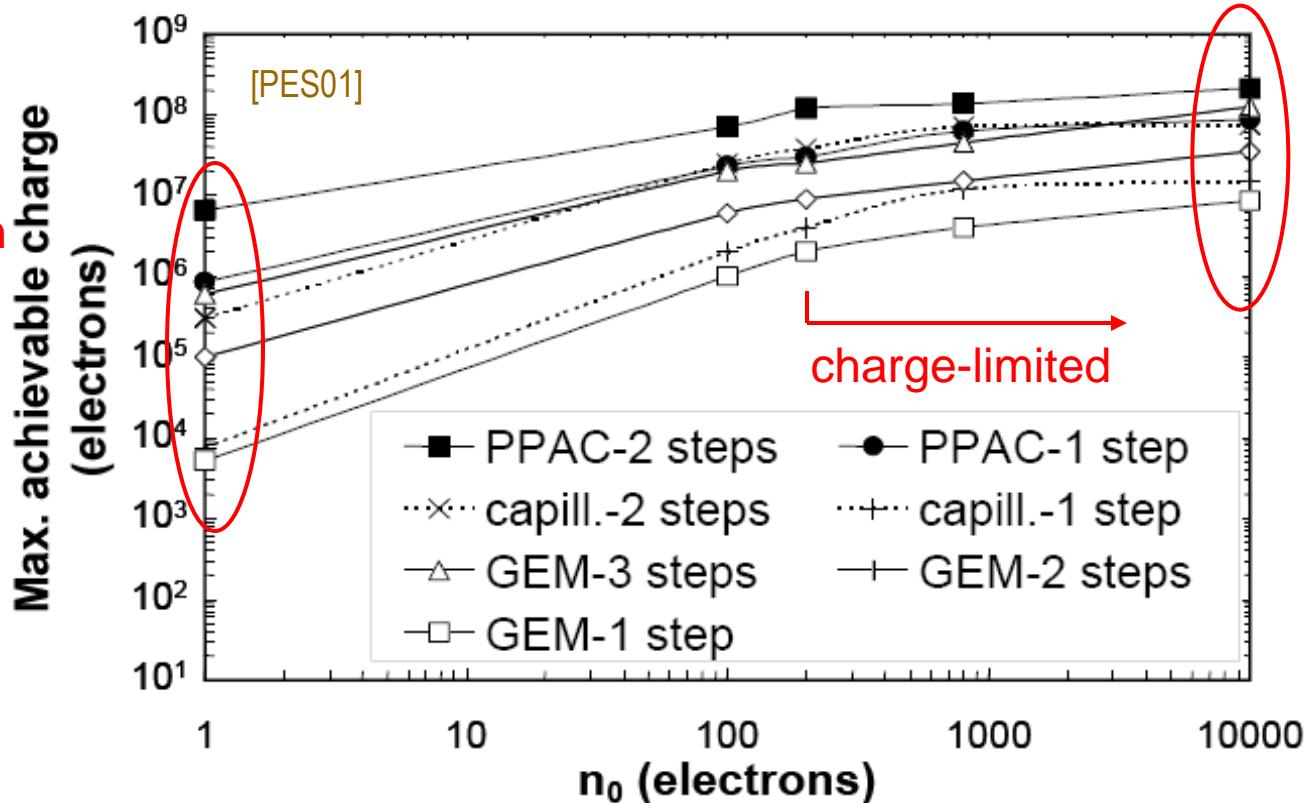


Avalanche gain saturation corresponds to the onset of streamers.  
No “limited proportionality” in parallel fields (except in  $SF_6$ ).

# Raether limit – micropattern detectors?

Geometry  
dependence  
+ multistep

Reduction at high gain  
Likely owing to:  
-avalanche statistics  
-Corona discharge



For  $n_0 > \sim 200$  electrons the Raether limit applies, but depends on geometry.  
For  $n_0 < \sim 200$  electrons other factors start to dominate, such as:  
avalanche gain fluctuation  
Corona discharge from sharp edges

# Hydrodynamic approach to streamer calculation

## Charge transport

$$\frac{\partial n_e(\vec{r},t)}{\partial t} = \underbrace{S}_{\text{other sources}} + \underbrace{(\alpha - \eta) |\vec{W}_e| n_e}_{\substack{\text{creation} \\ \text{multiplication} \\ \text{attachment}}} - \underbrace{\vec{\nabla} \cdot (\vec{W}_e n_e)}_{\substack{\text{transport} \\ \substack{\vec{W}_e \cdot \vec{\nabla} n_e + n_e \vec{\nabla} \cdot \vec{W}_e \\ \text{drift} \\ \text{pile-up}}}} + \underbrace{D_e \nabla^2 n_e}_{\text{diffusion}}$$

good reference: [DAV73]

electrons

$n(\vec{r},t)$  = charge density in space and time

$\vec{W}(E)$  = velocity of charges

$\vec{E}(\vec{r},t)$  = electric field: applied+space charge

$\alpha$ =first Townsend coefficient

$D$ =diffusion coefficient

Space-charge + applied field

$$\nabla^2 V = -\frac{e}{\epsilon_0} (n_{i+} - n_e - n_{i-})$$

Boundary conditions

initial densities:  $n_{e,i\pm}(\vec{r},0)$

behaviour of charges at the electrodes

Electrostatic B.C.

Slight drawback: no avalanche statistics

$$\frac{\partial n_{i+}(\vec{r},t)}{\partial t} = S + \alpha |\vec{W}_e| n_e$$

Ions, assuming stationary ions

$$\frac{\partial n_{i-}(\vec{r},t)}{\partial t} = \eta |\vec{W}_e| n_e$$

# Gas self-photoionization as a secondary process

It is possible that just transport accounts for the forward (anode) streamer but for the cathode streamer (growing backwards) something else is needed.

e.g. photoemission proportional to the electron multiplication

$$\frac{\partial n_f(\vec{r}, t)}{\partial t} = \delta |\vec{W}_e| n_e \quad \text{photon creation}$$

+ gas **self-photoionization** source term

$$S(\vec{r}, t) = \frac{Q}{\lambda} \int_{Volume} \frac{\partial n_f(\vec{r}', t)}{\partial t} \Omega(\vec{r} - \vec{r}') e^{|\vec{r}-\vec{r}'|/\lambda} d\vec{r}' \quad \text{distribute the photons around and ionize the gas}$$

$\delta$  = photon yield per electron

$\Omega$  = solid angle fraction from emission to absorption point

$Q$  = quantum efficiency

$\lambda$  = photon's mean free path

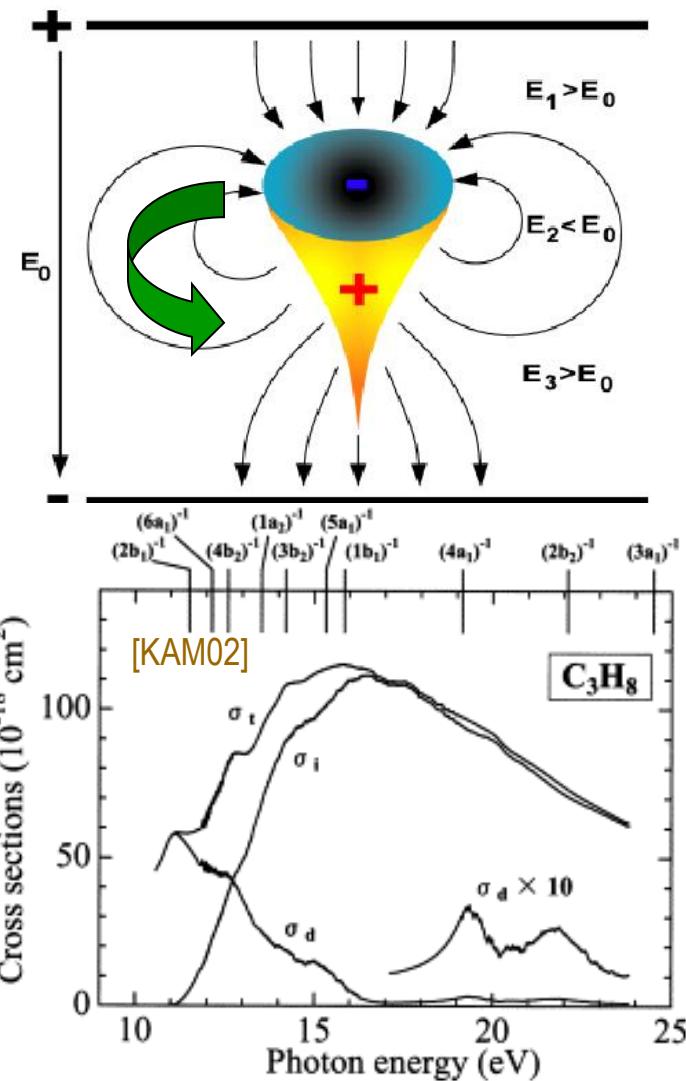
All this for each relevant emission wavelength...

# Gas self-photoionization as a secondary process?

- Streamer breakdown seems to be an universal phenomenon. (Generations of electrical engineers and detector physicists couldn't find a way to avoid it.)
- However, self-photoionization hardly allows an universal situation. Essentially every mixture requires its own theory!
  - Depends on the details of photoemission, photoabsorption and photoionization spectra and that must be considered for every component of a gas mixture;
  - Photoemission yields are essentially unknowable. Both hard to measure and hard to calculate. Yield depends on competing deexcitation processes: very mixture-dependent.
- In pure propane, for instance, at atm. pressure the absorption length at the edge of the photoionization band is only  $\sim 10\mu\text{m}$ . Hardly enough for closing a gain loop, even without considerations about yield.

IS THERE ANY OTHER WAY?

Closing the gain loop  
by self-photoionization?



# Diffusion as a streamer-supporting process

PHYSICAL REVIEW E

VOLUME 55, NUMBER 2

FEBRUARY 1997

## Propagation and structure of planar streamer fronts

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Université Paris VII, GPS Tour 23, 2 place Jussieu, 75251 Paris Cedex 05, France

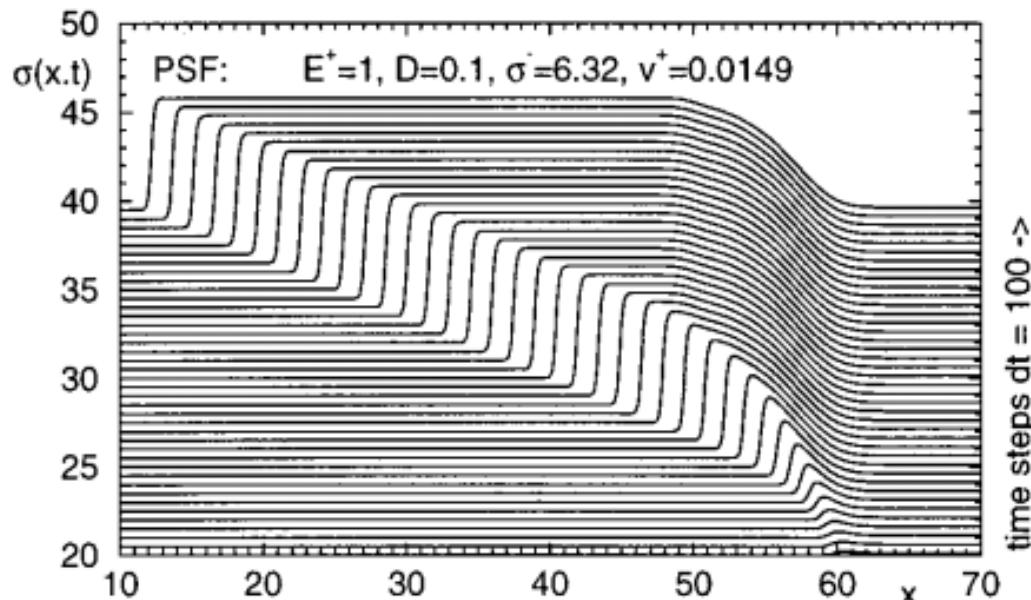


FIG. 10. Emergence of the uniformly translating PSF on the left for  $D = 0.1$ . Initial conditions identical with Fig. 9. The time range  $t = 4000\text{--}8000$  after an initial perturbation at  $t = 0$  and  $x_0 = 60$  is shown in time steps of  $\Delta t = 100$ . (Numerical grid size  $\Delta x = 0.01$  and  $\Delta \tau = 0.5$ .)

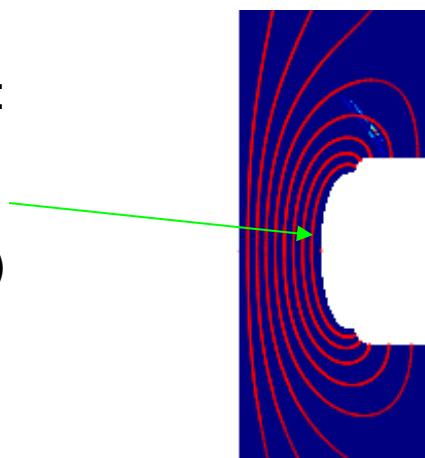
**Analytical and numerical proof that diffusion alone provides a sufficient mechanism for positive streamer front (PSF) propagation in some simplifying (but quite reasonable) conditions.**

# Simplified hydrodynamic model

$$\begin{cases} \frac{\partial n_e}{\partial t} = \alpha |\vec{W}_e| n_e - \vec{\nabla} \cdot (\vec{W}_e n_e) + D_e \nabla^2 n_e \\ \frac{\partial n_{i+}}{\partial t} = \alpha |\vec{W}_e| n_e \\ \nabla^2 V = -\frac{e}{\epsilon_0} (n_{i+} - n_e) \end{cases}$$

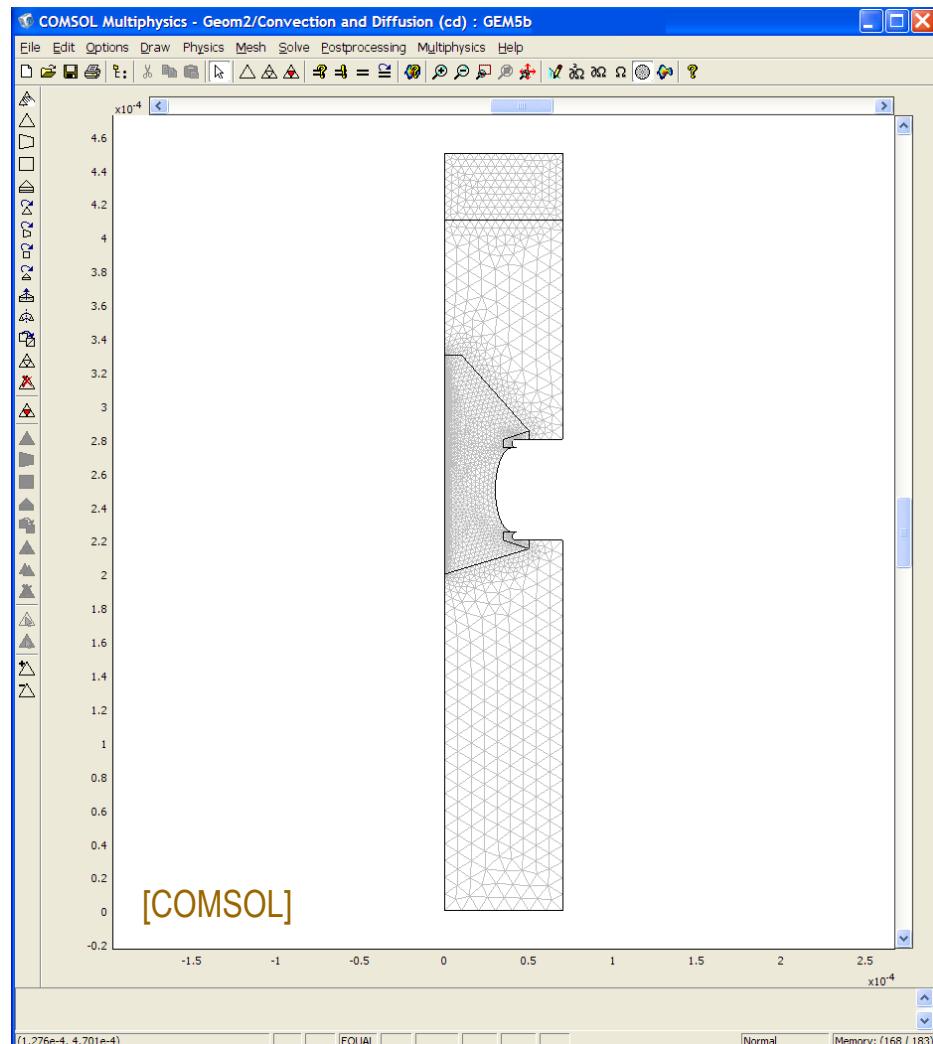
$n_{e,i+}(\vec{r},t)$  = charge density in space and time  
 $\vec{E}(\vec{r},t)$  = electric field =  $\vec{\nabla}V(\vec{r},t)$   
 $\vec{W}_e(\vec{E})$  = electron velocity  
 $\alpha(\vec{E})$  = first Townsend coefficient  
 $D_e(\vec{E})$  = electron diffusion coefficient

- Only electrons and positive ions
- No positive ion movement (in such short time span)
- No attachment
- **No photons**
- Assume axial symmetry (minimal condition for realism):  
2D calculation
- Applied field: boundary conditions on the potential
- Dielectrics: tangent (no charge flow into the surface)



# Numerical approach: finite elements

Used the commercial program *COMSOL Multiphysics*



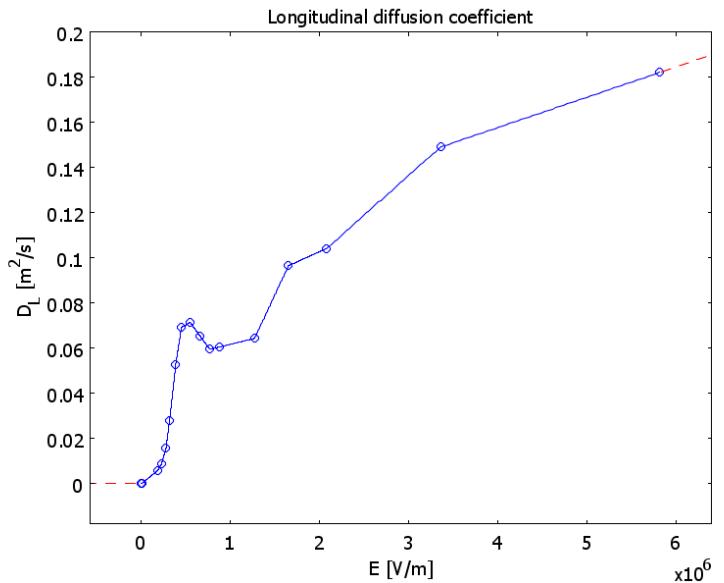
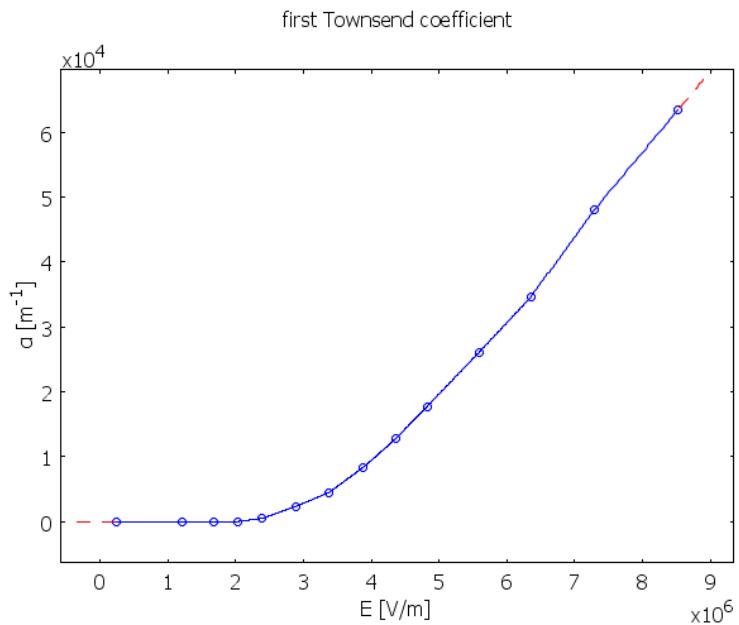
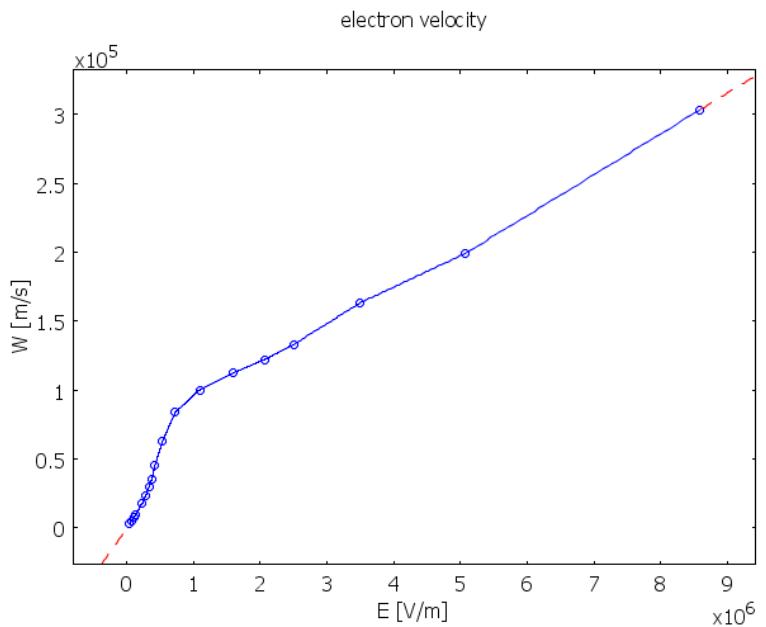
Solves a coupled set of a basic differential equation,

$$\left\{ \begin{array}{ll} e_a \frac{\partial^2 u}{\partial t^2} + d_a \frac{\partial u}{\partial t} + \nabla \cdot (-c \nabla u - \alpha u + \gamma) + \beta \cdot \nabla u + \alpha u = f & \text{in } \Omega \\ \mathbf{n} \cdot (c \nabla u + \alpha u - \gamma) + q u = g - h^T \boldsymbol{\mu} & \text{on } \partial\Omega \\ h u = r & \text{on } \partial\Omega \end{array} \right.$$

with arbitrary coefficients (any function of any variable) in a mesh (finite elements).

Covers most cases needed for applied physics.

# Gas: pure CO<sub>2</sub> @ atm. pressure



[HER02]

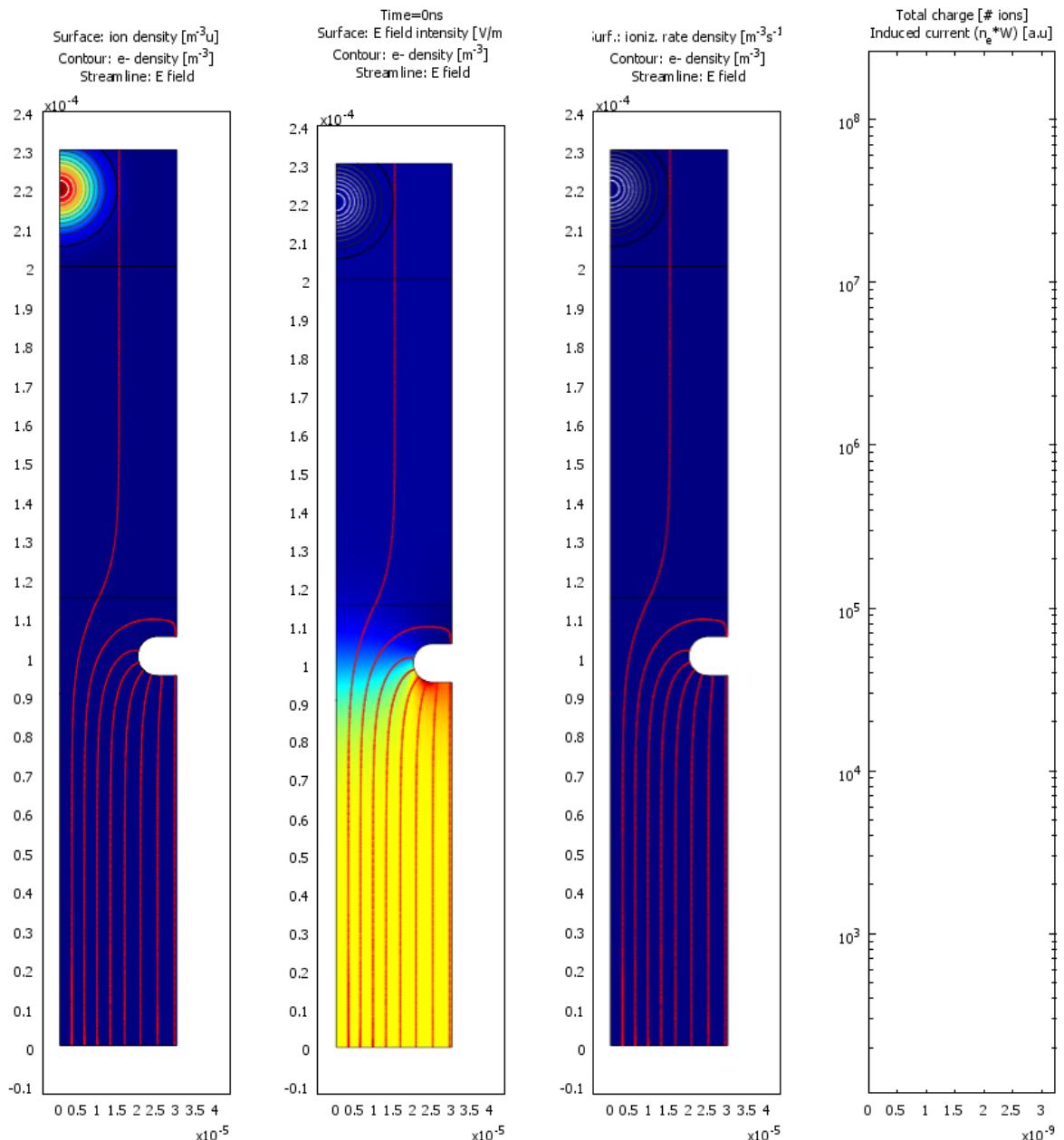
# MICROMEGAS

hole: 40 µm  
gap: 100 µm  
 $N_0=100 e^-$

Total induced current

$$\propto \int_{Volume} |\vec{j}| dV = \int_{Volume} n_e |\vec{W}_e| dV$$

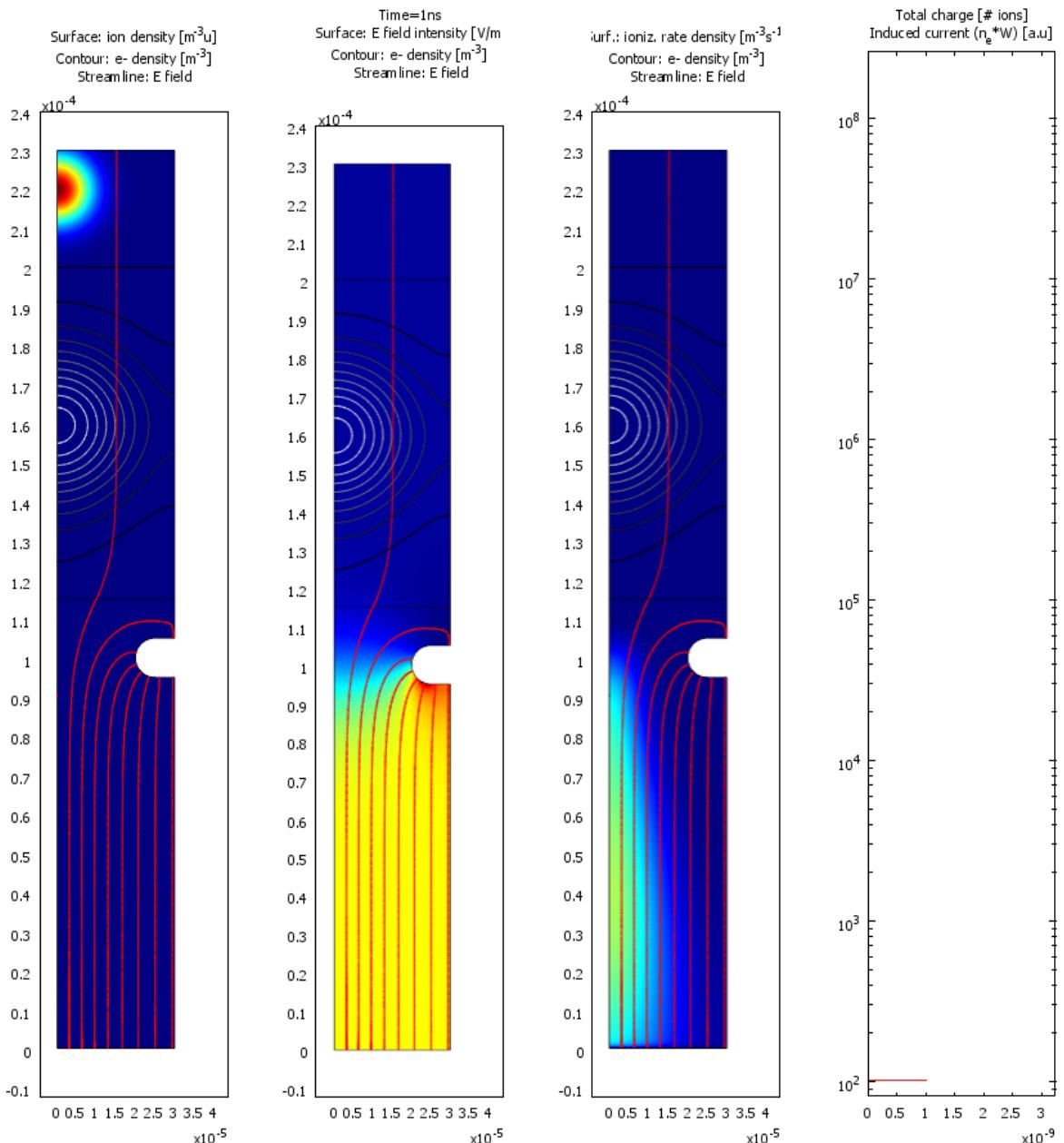
The details depend on which electrodes we are actually collecting this current.



# MICROMEGAS

hole: 40 µm  
gap: 100 µm  
 $N_0=100 e^-$

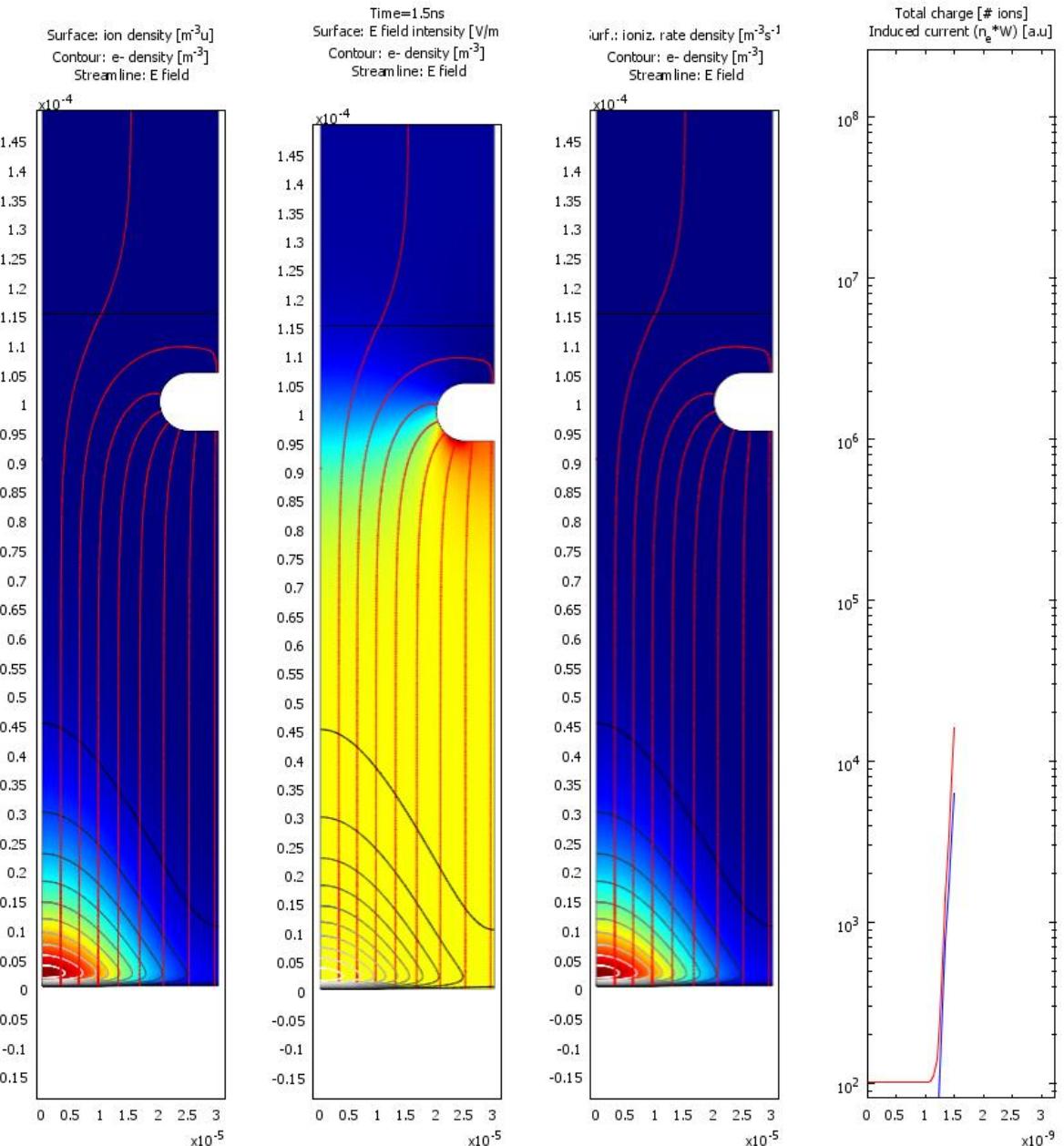
Start of multiplication  
while most of the electrons  
are still in the drift region



# MICROMEGAS

hole: 40 µm  
gap: 100 µm  
 $N_0=100 e^-$

Proportional gain

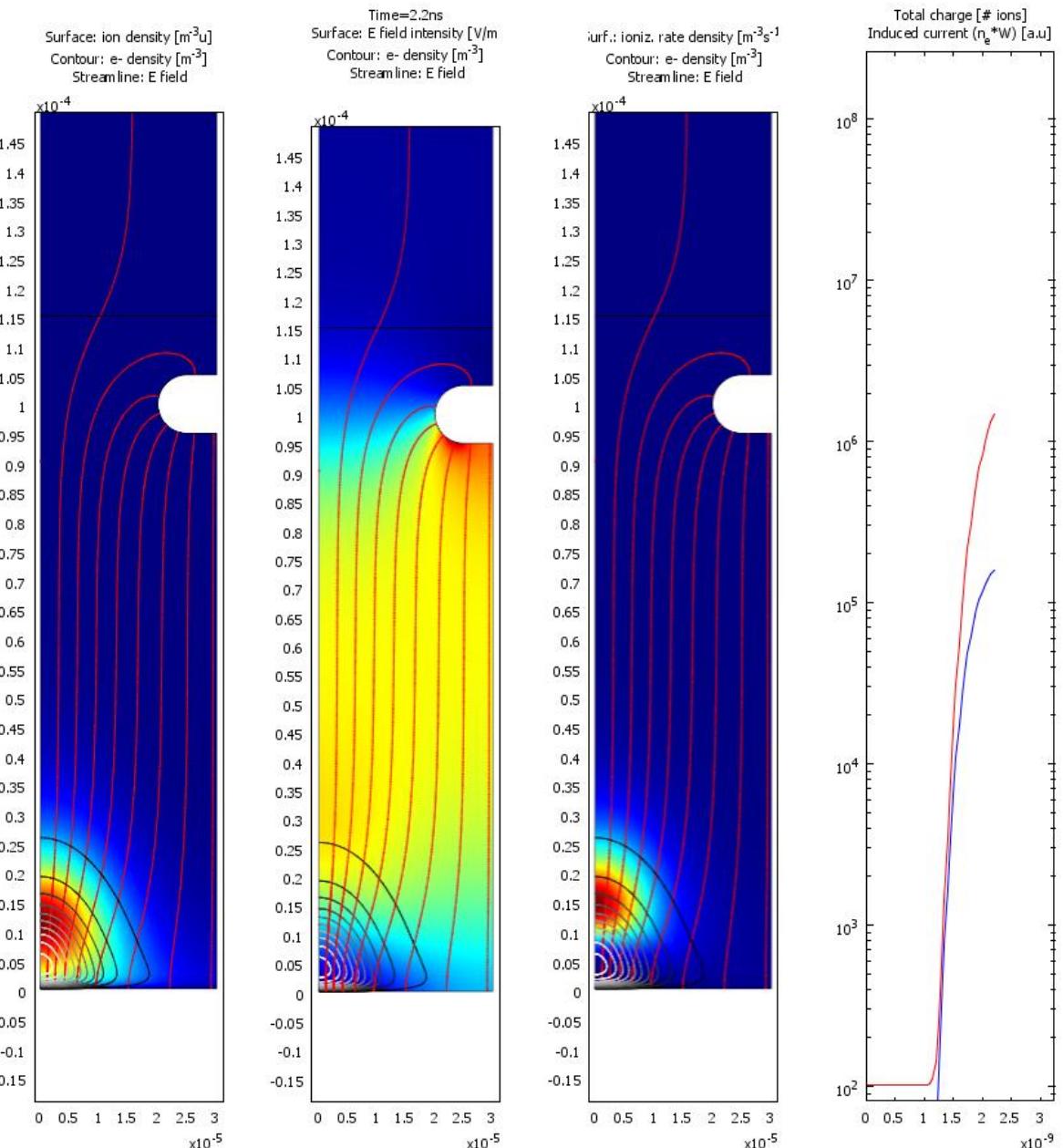


# MICROMEGAS

hole: 40 µm  
gap: 100 µm  
 $N_0=100 e^-$

Onset of space-charge:  
maximum multiplication  
region moves away from  
anode owing to space-  
charge.

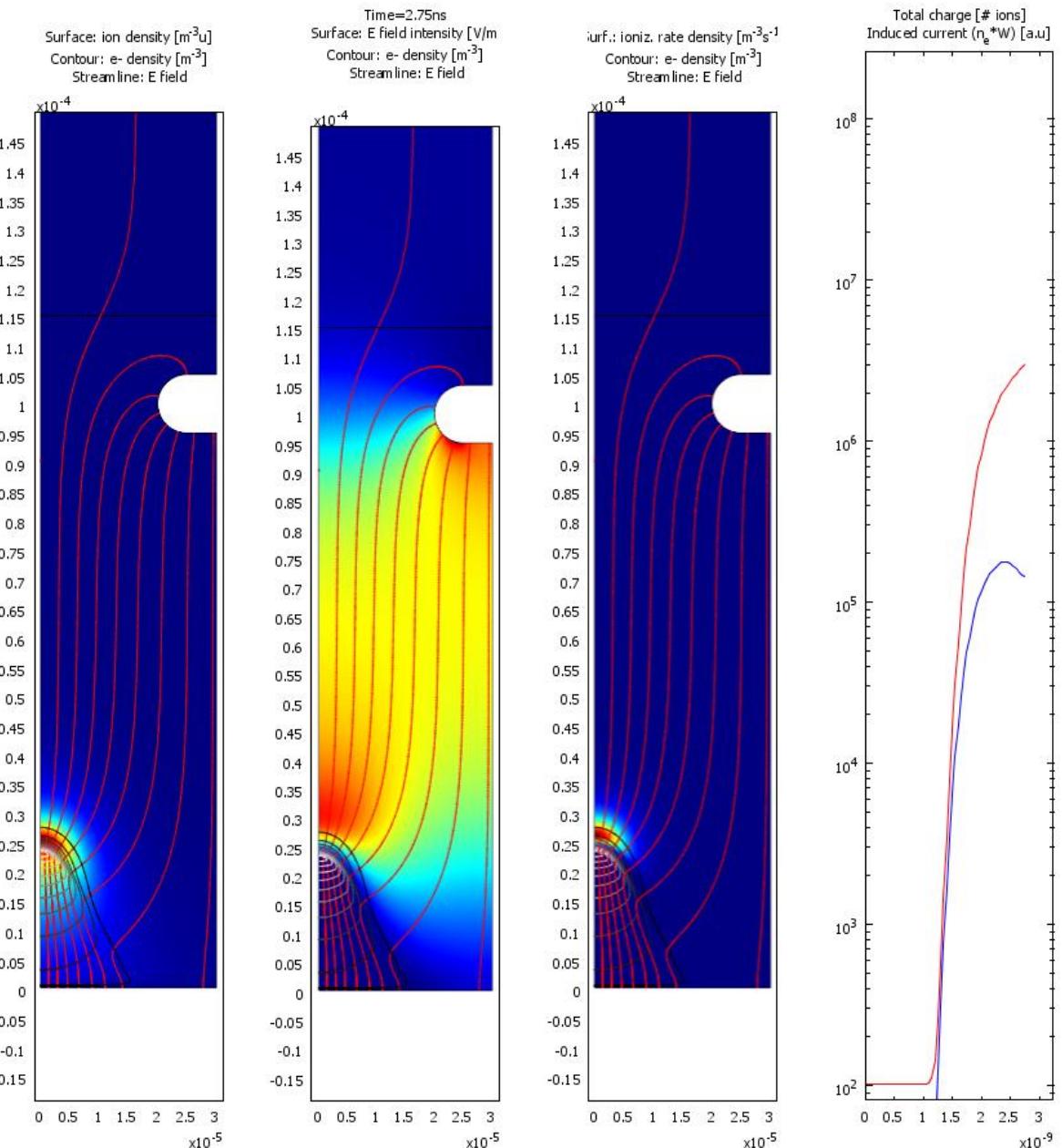
This is not a streamer yet.  
Depends on the inflow of  
electrons from the drift for  
growth.



# MICROMEGAS

hole: 40 µm  
gap: 100 µm  
 $N_0=100 e^-$

The induced current actually decreases because the multiplication of the inflow of electrons is not enough to compensate for the electron losses to the anode.

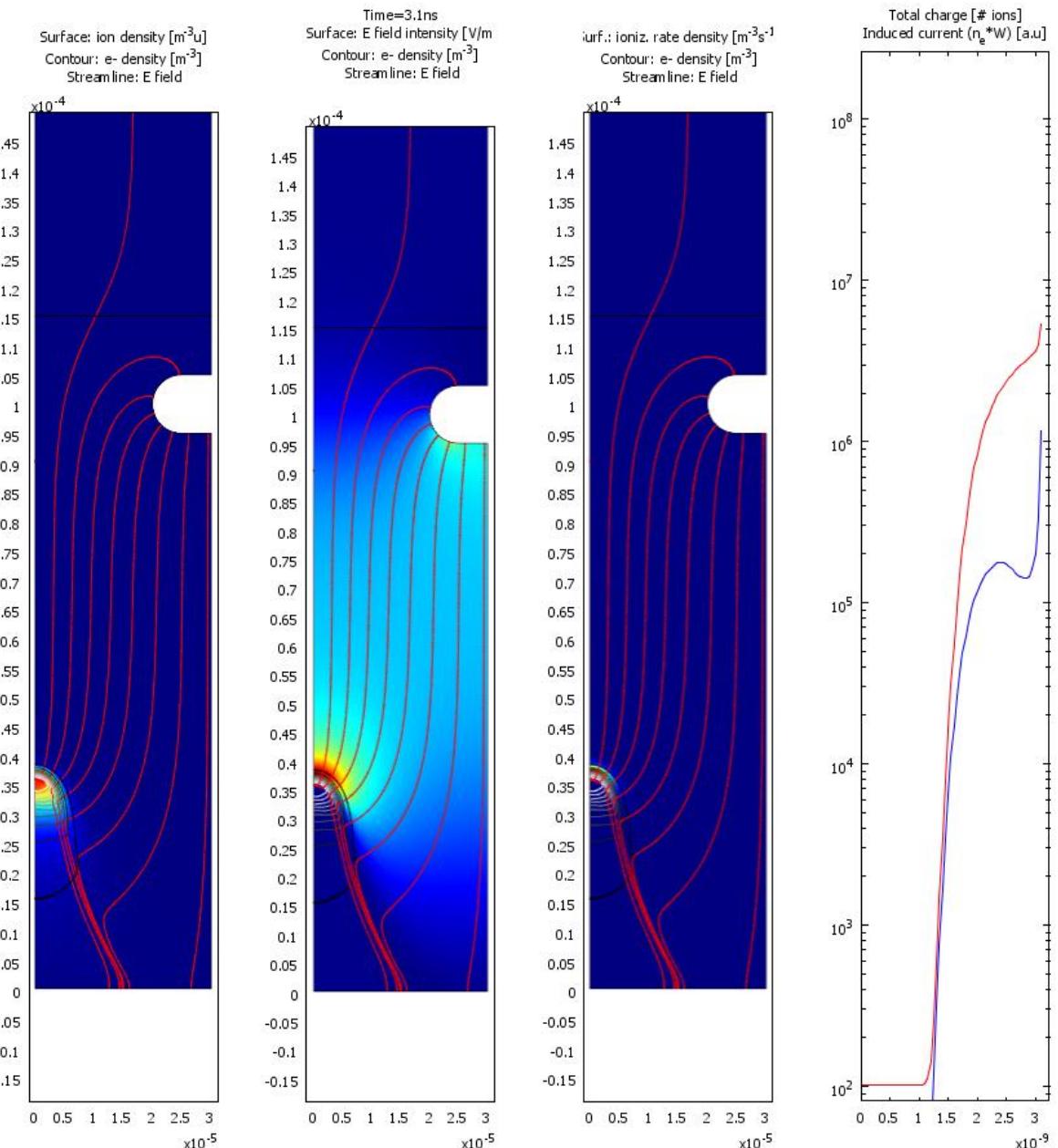


# MICROMEGAS

hole: 40 µm  
gap: 100 µm  
 $N_0=100 e^-$

A streamer structure is formed on the tip of a plasma conductive “needle”.

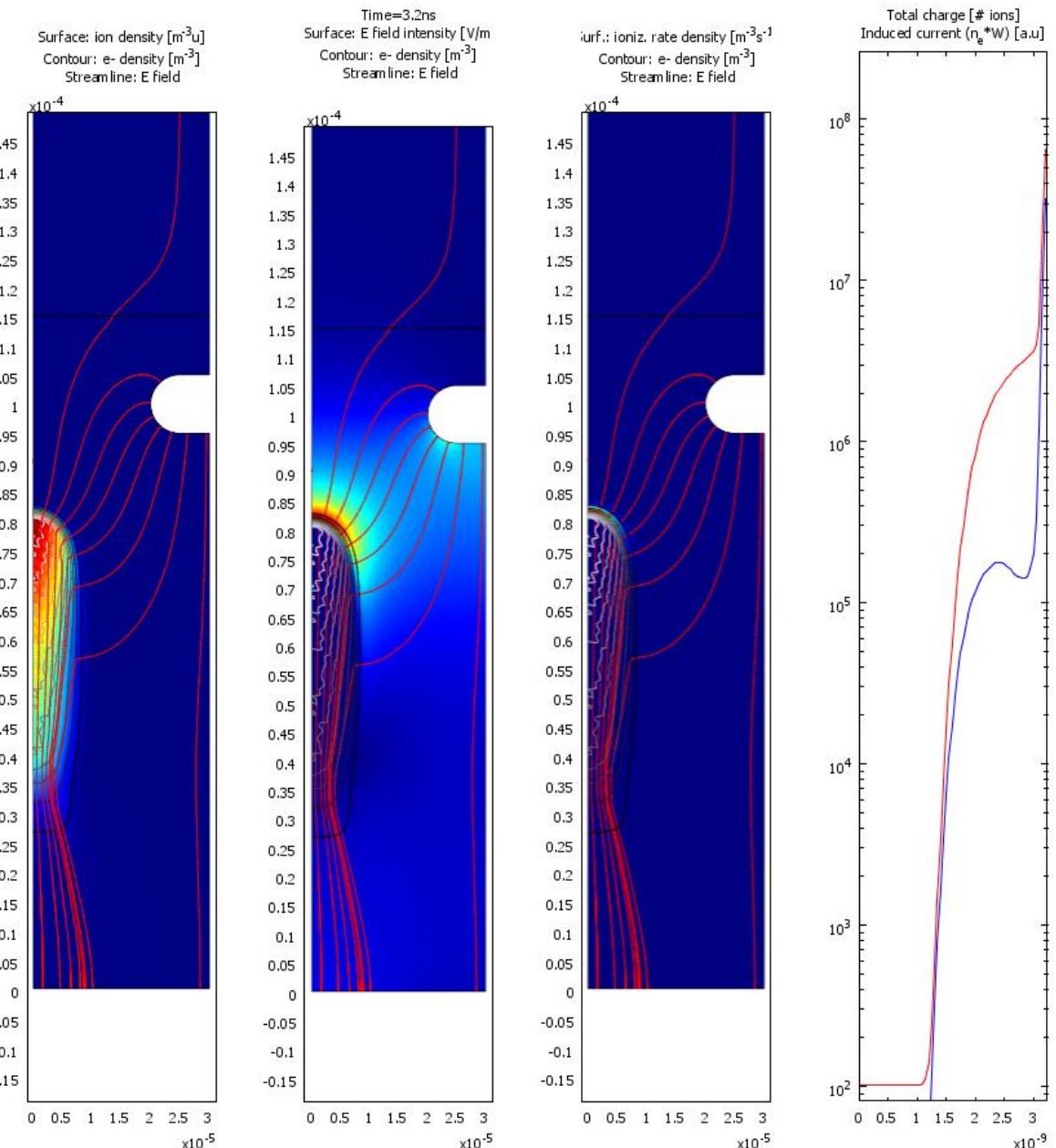
The current increases and the structure known as “the precursor” is formed.



# MICROMEGAS

hole: 40 µm  
gap: 100 µm  
 $N_0=100 e^-$

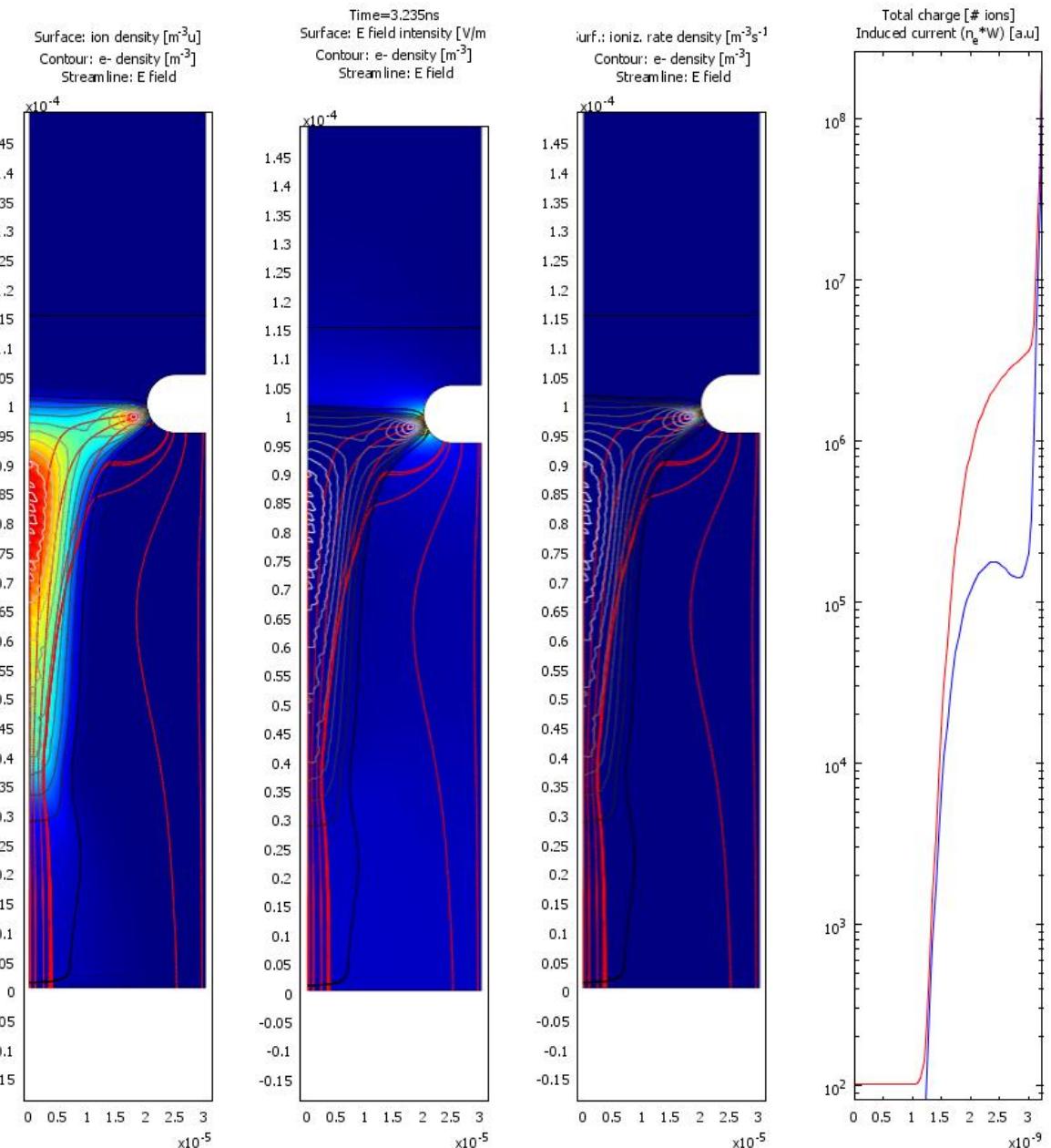
The streamer is self-sustained and propagates towards the cathode at very high speed.



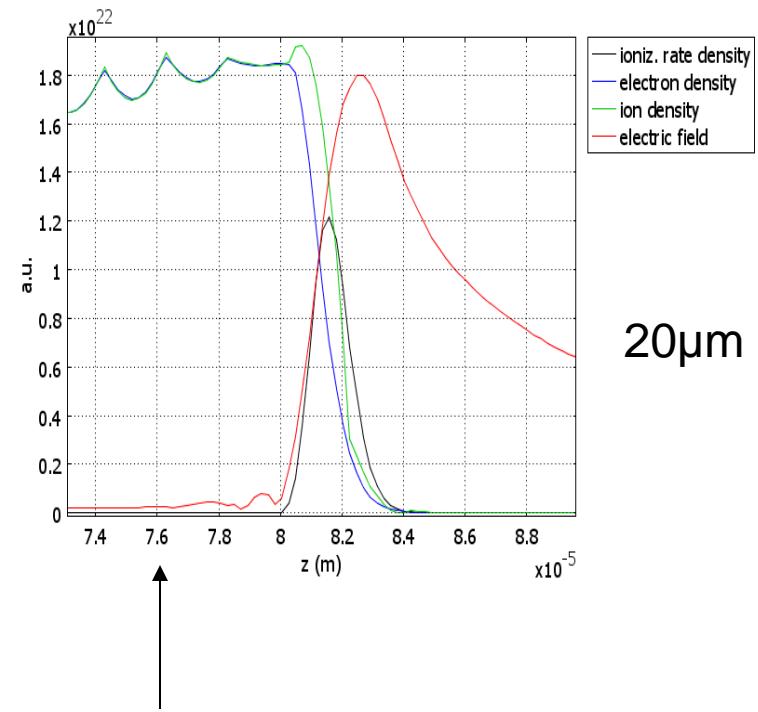
# MICROMEGAS

hole: 40 µm  
gap: 100 µm  
 $N_0=100 e^-$

The streamer branches sideways towards the mesh, bridging anode and cathode with a conductive plasma path.  
Further discharge stages will follow, eventually ending with a spark.

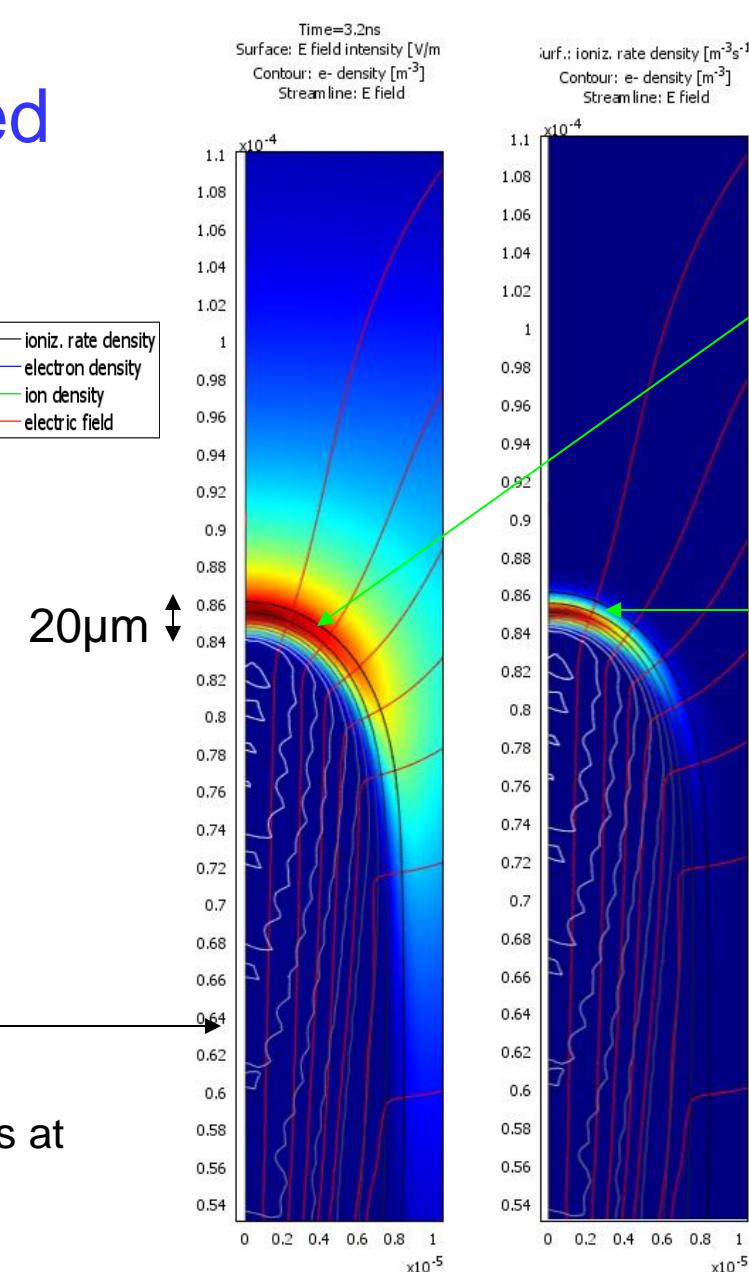


# Structure of the diffusion-assisted streamer



Very low field region  
↓

the anode potential appears at  
the tip of the streamer



Owing to the strong density gradient some electrons actually diffuse against the E field into the high gain region on the tip of the streamer.

Above a certain gain threshold this structure is self-sustained.

A thin active region forms upstream of the ion density maximum, moving the maximum upwards at high speed.

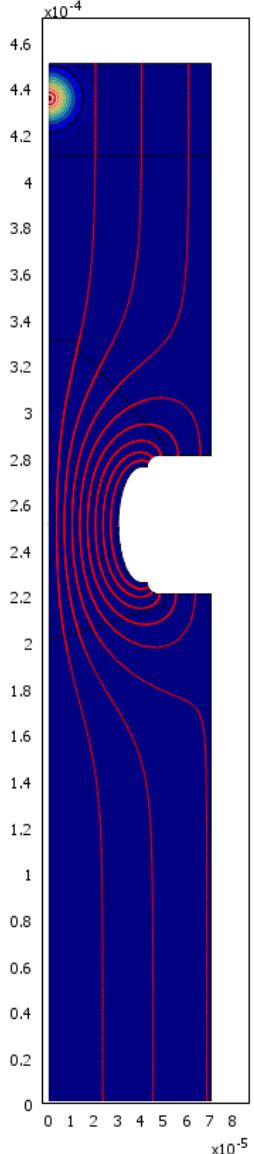
The movement of the streamer is the movement of the density maximum, not a physical velocity of an object.

# GEM

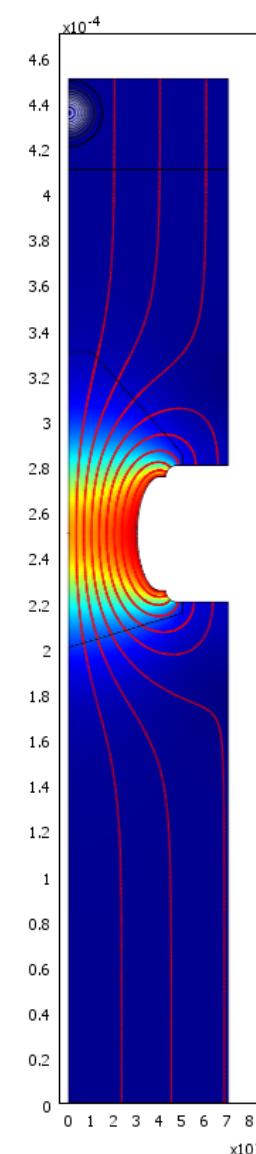
hole: 60 µm  
gap: 100 µm

$N_0=100$  e<sup>-</sup>  
 $V=1250$ V

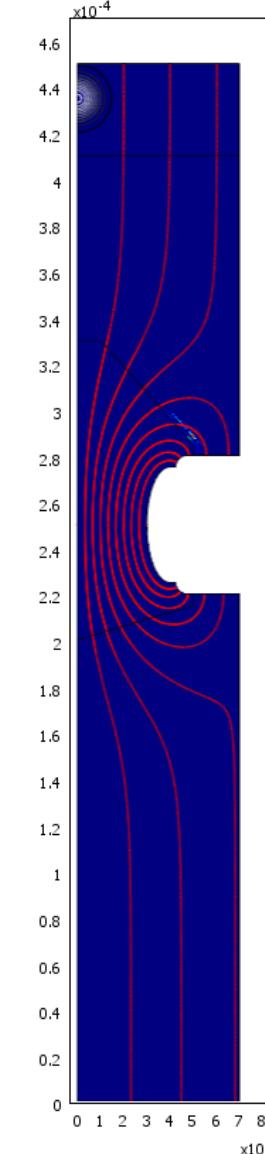
Surface: ion density [ $\text{m}^{-3}\text{u}$ ]  
Contour: e- density [ $\text{m}^{-3}$ ]  
Streamline: E field



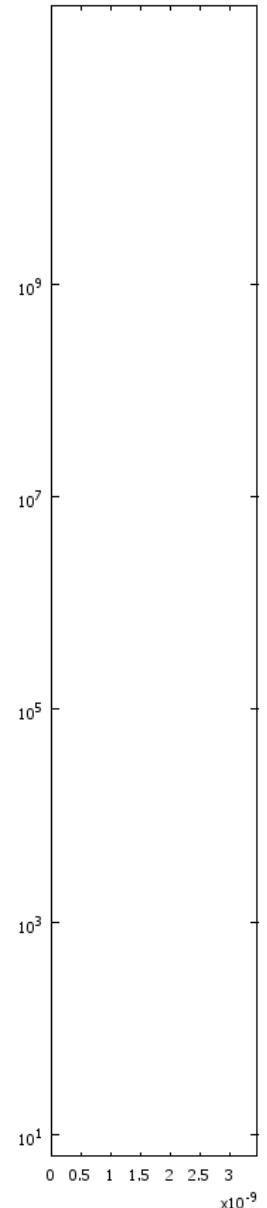
Time=0ns  
Surface: E field intensity [V/m]  
Contour: e- density [ $\text{m}^{-3}$ ]  
Streamline: E field



Surf.: ioniz. rate density [ $\text{m}^{-3}\text{s}^{-1}$ ]  
Contour: e- density [ $\text{m}^{-3}$ ]  
Streamline: E field



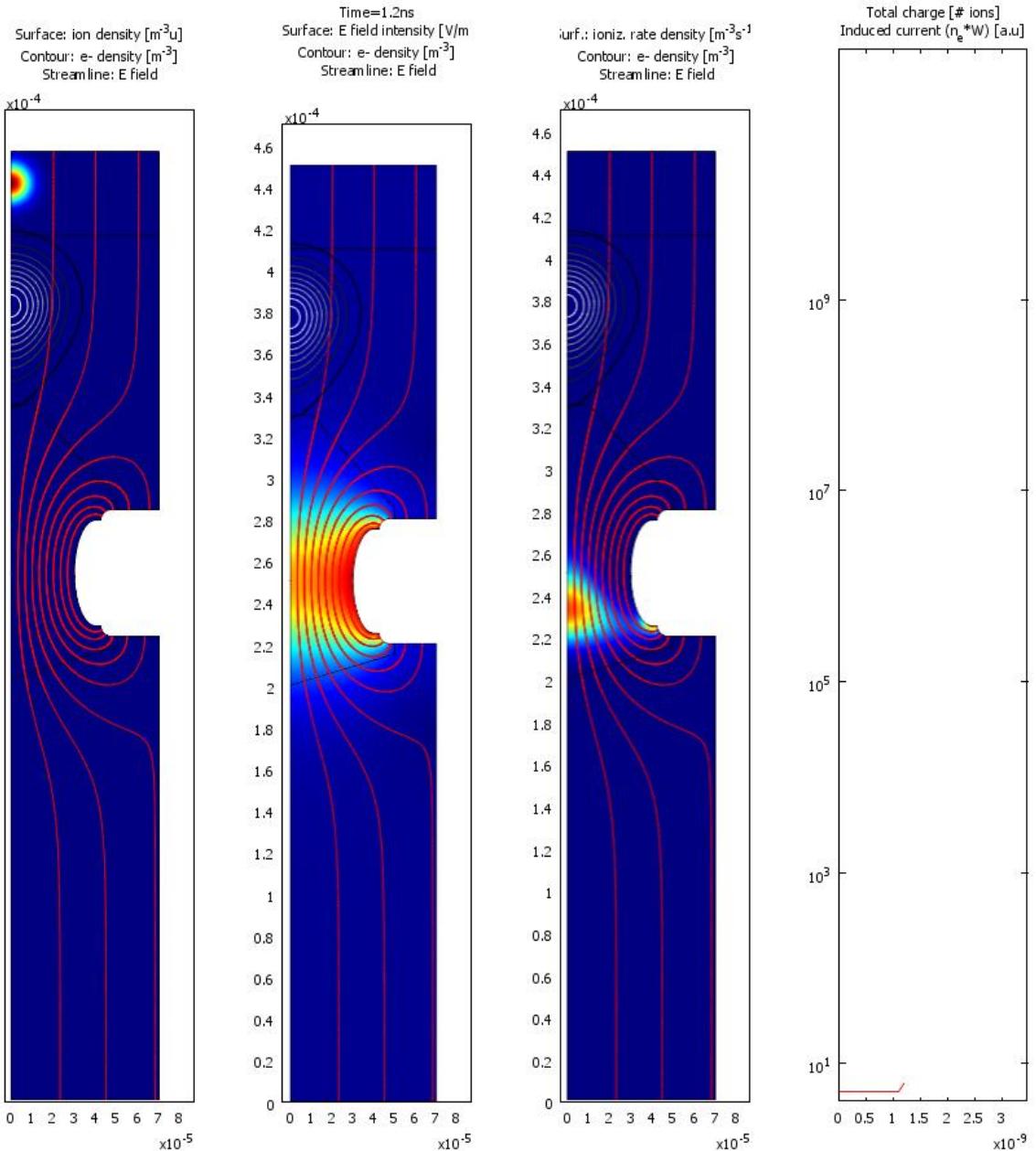
Total charge [# ions]  
Induced current ( $n_e \cdot W$ ) [a.u]



# GEM

hole: 60 µm  
gap: 100 µm

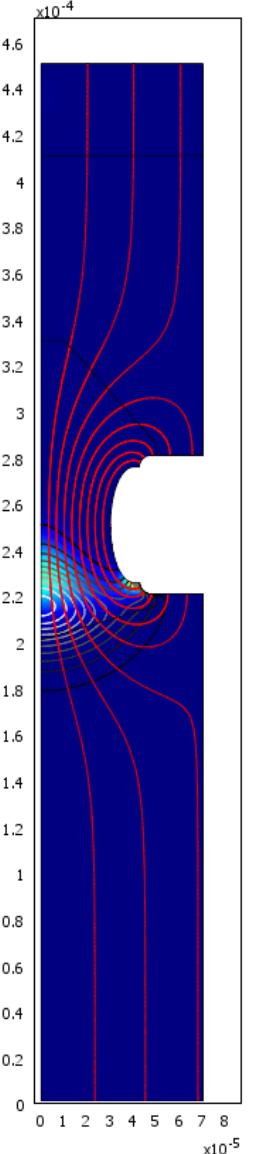
$N_0=100$  e<sup>-</sup>  
 $V=1250V$



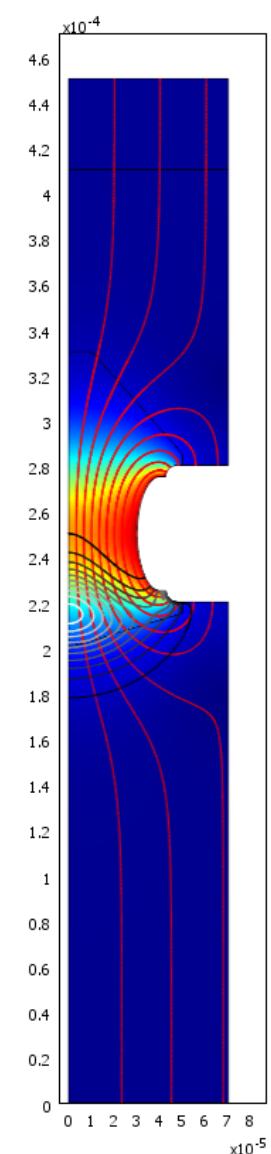
# GEM

hole: 60 µm  
 gap: 100 µm  
 $N_0=100 e^-$   
 $V=1250V$

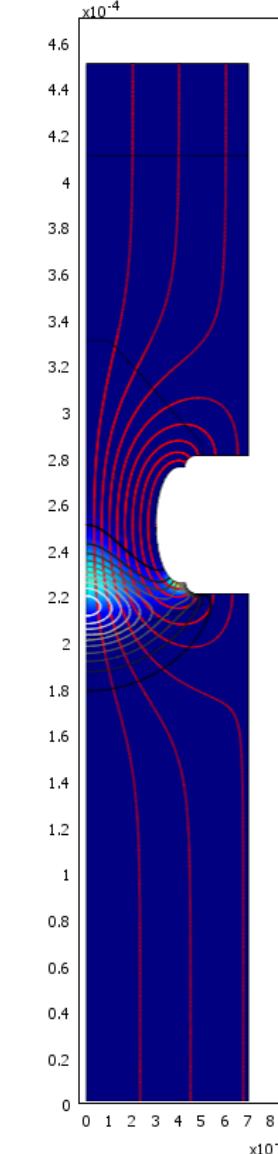
Surface: ion density [ $m^{-3} u$ ]  
 Contour: e- density [ $m^{-3}$ ]  
 Streamline: E field



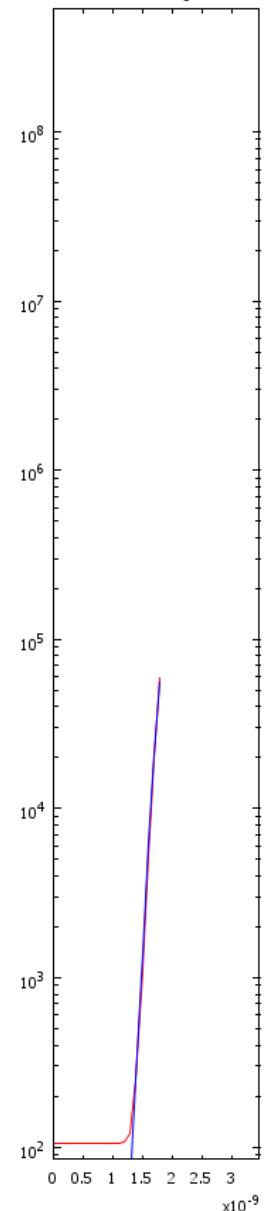
Time=1.8ns  
 Surface: E field intensity [V/m]  
 Contour: e- density [ $m^{-3}$ ]  
 Streamline: E field



Surf.: ioniz. rate density [ $m^{-3}s^{-1}$ ]  
 Contour: e- density [ $m^{-3}$ ]  
 Streamline: E field



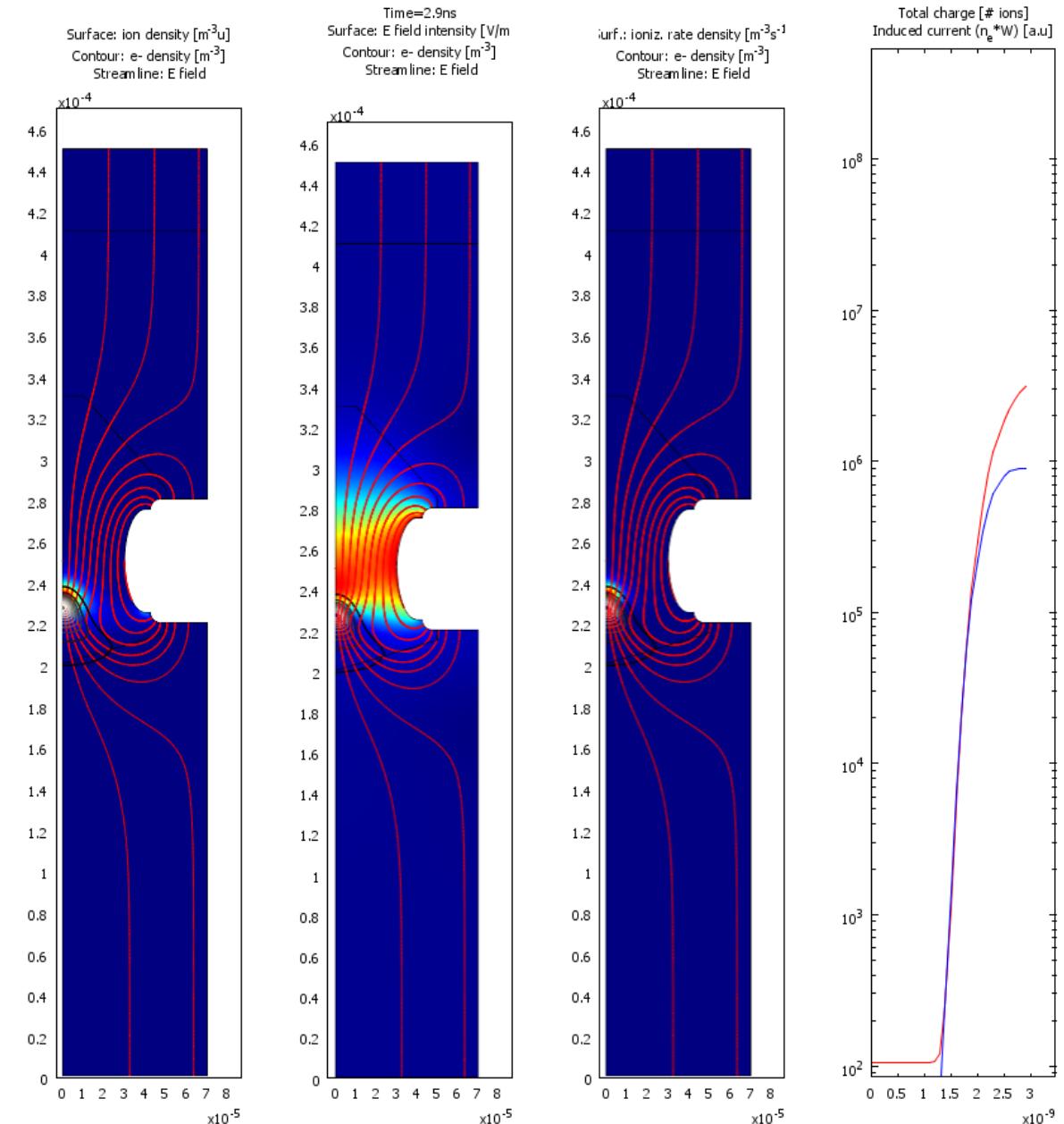
Total charge [# ions]  
 Induced current ( $n_e \cdot W$ ) [a.u.]





# GEM

hole: 60 µm  
 gap: 100 µm  
 $N_0=100 e^-$   
 $V=1250V$

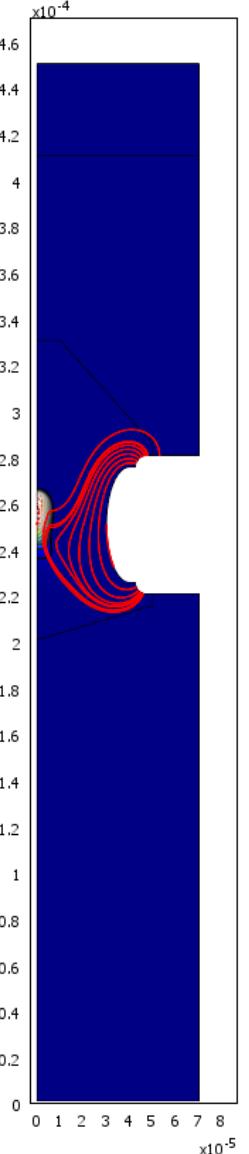


# GEM

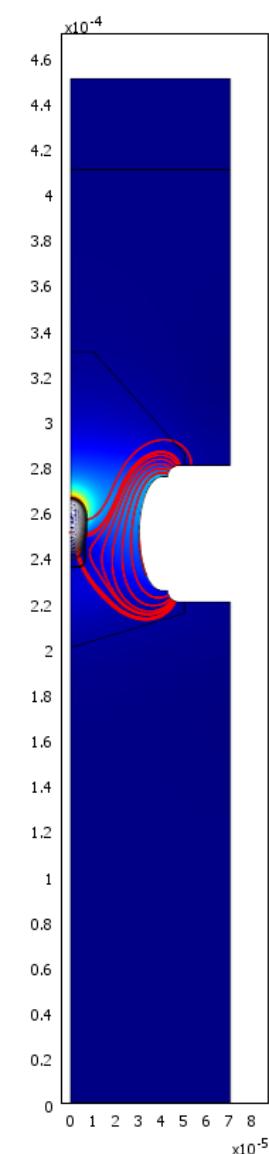
hole: 60  $\mu\text{m}$   
gap: 100  $\mu\text{m}$

$N_0=100 \text{ e}^-$   
 $V=1250\text{V}$

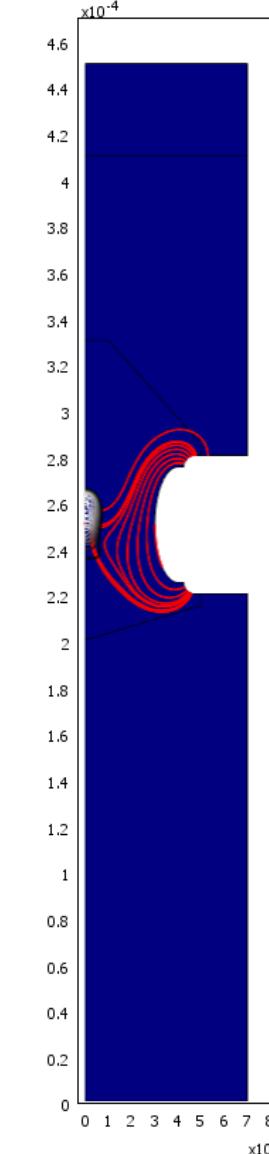
Surface: ion density [ $\text{m}^{-3}\text{u}$ ]  
Contour: e- density [ $\text{m}^{-3}$ ]  
Streamline: E field



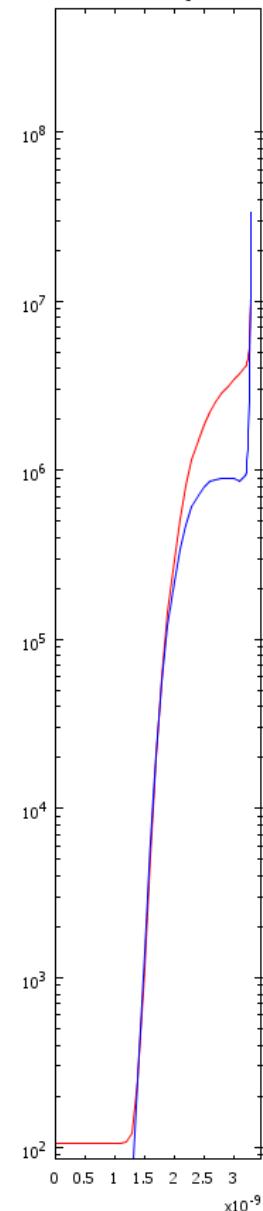
Time=3.3ns  
Surface: E field intensity [V/m]  
Contour: e- density [ $\text{m}^{-3}$ ]  
Streamline: E field



Surf.: ioniz. rate density [ $\text{m}^{-3}\text{s}^{-1}$ ]  
Contour: e- density [ $\text{m}^{-3}$ ]  
Streamline: E field

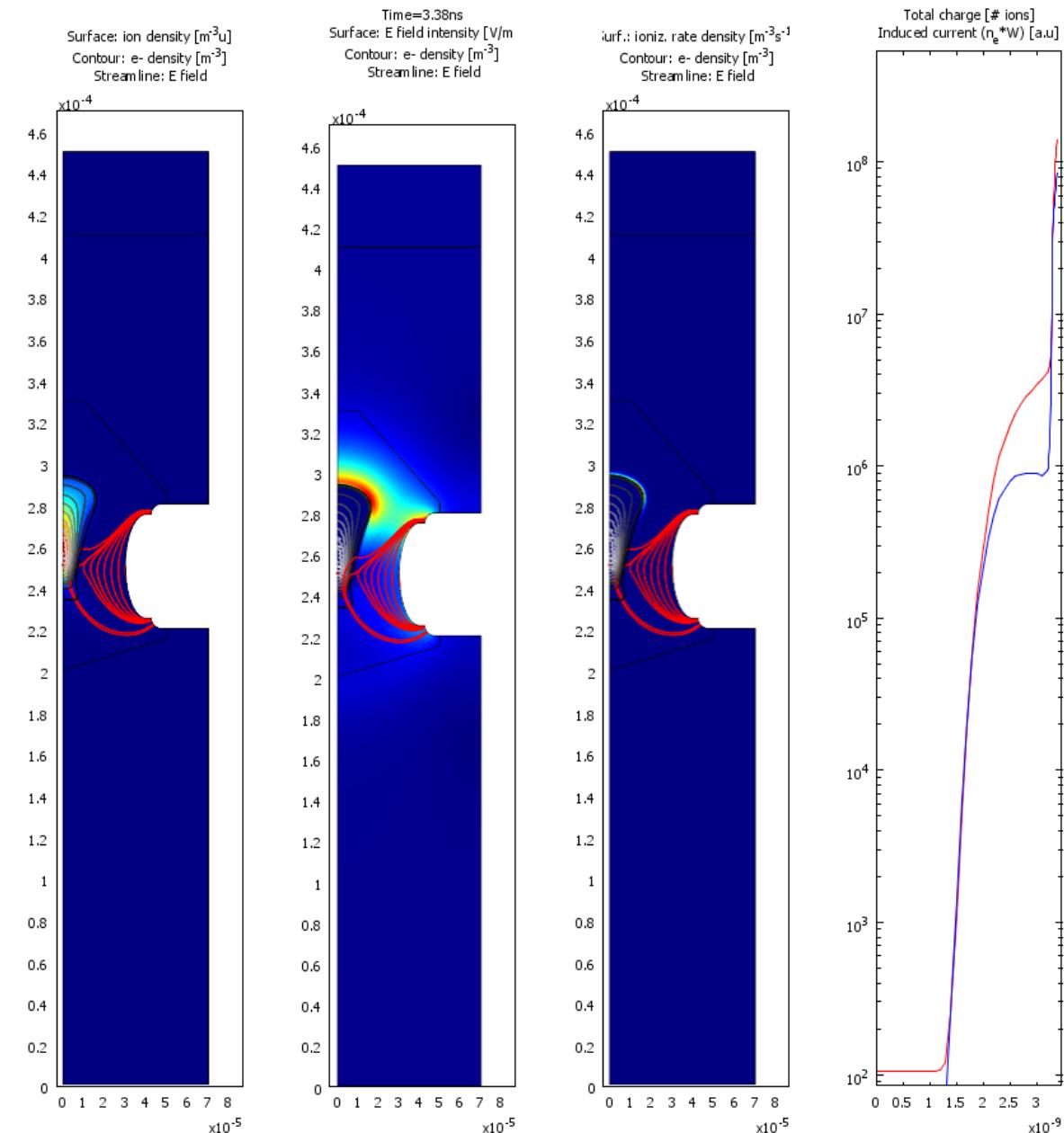


Total charge [# ions]  
Induced current ( $n_e \cdot W$ ) [a.u.]



# GEM

hole: 60 µm  
 gap: 100 µm  
 $N_0=100 e^-$   
 $V=1250V$

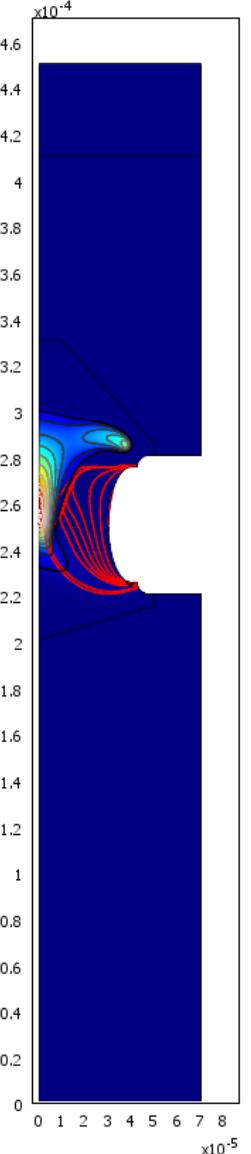


# GEM

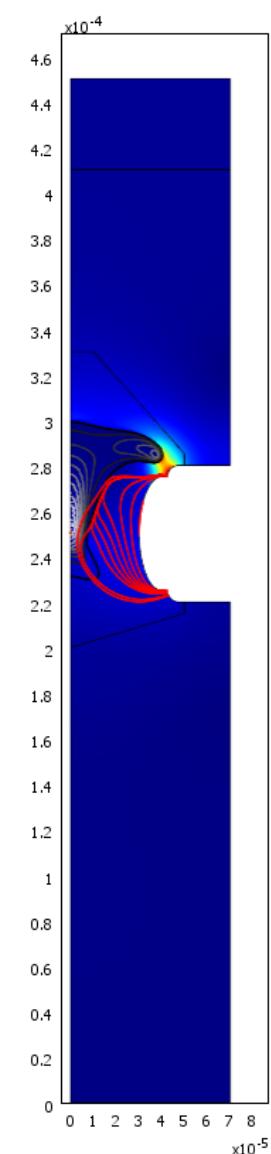
hole: 60 µm  
gap: 100 µm

$N_0=100 e^-$   
 $V=1250V$

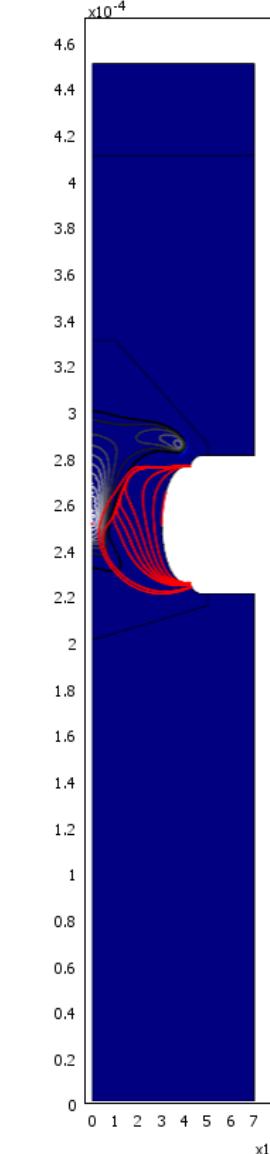
Surface: ion density [ $m^{-3} u$ ]  
Contour: e- density [ $m^{-3}$ ]  
Streamline: E field



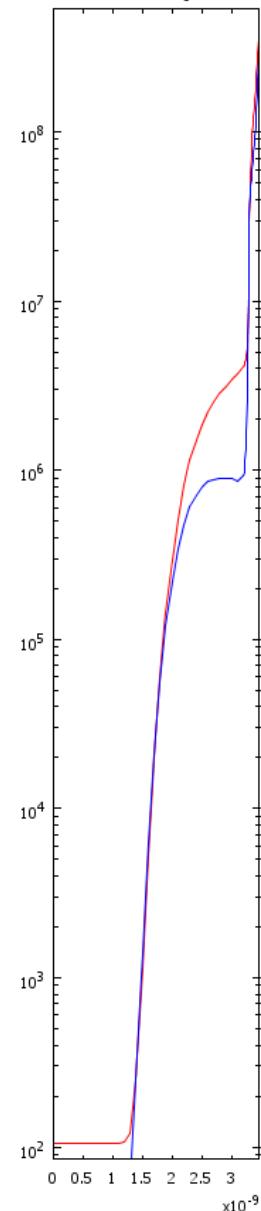
Time=3.46ns  
Surface: E field intensity [V/m]  
Contour: e- density [ $m^{-3}$ ]  
Stream line: E field



Surf.: ioniz. rate density [ $m^{-3}s^{-1}$ ]  
Contour: e- density [ $m^{-3}$ ]  
Streamline: E field



Total charge [# ions]  
Induced current ( $n_e * W$ ) [a.u.]



# GEM

## lateral (ring)

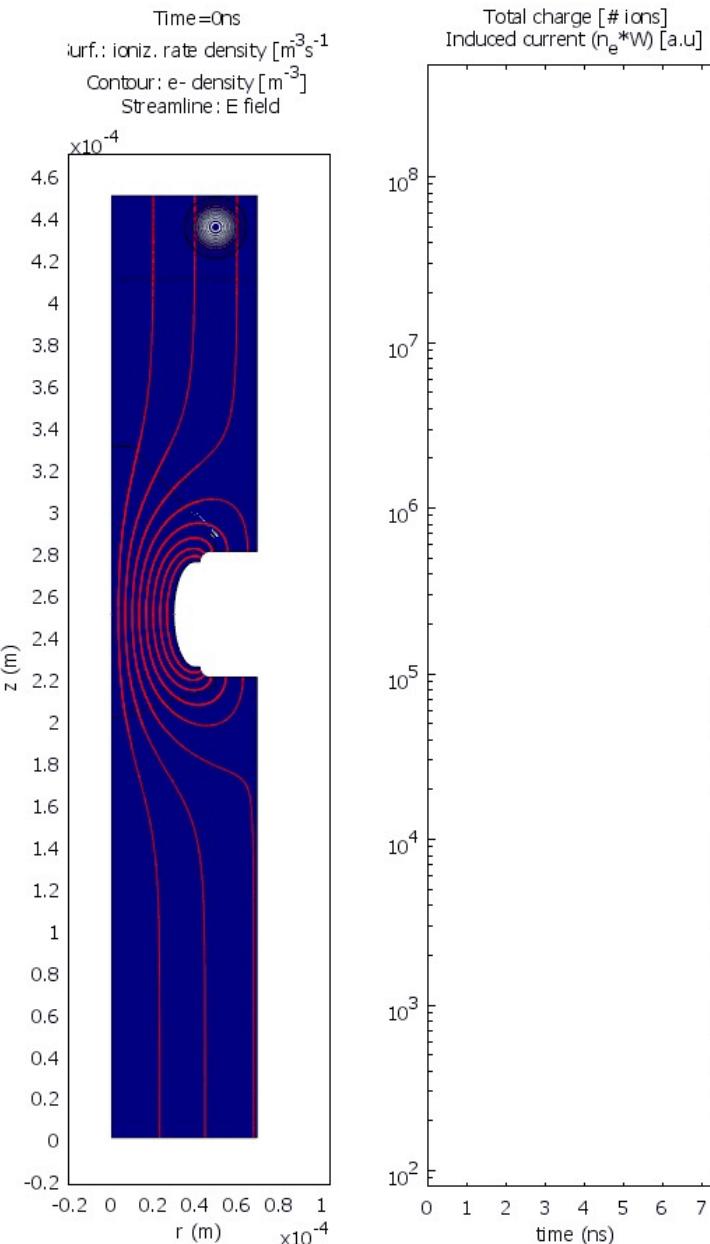
### avalanche

hole: 60 µm

gap: 100 µm

$N_0=100$  e<sup>-</sup>

V=1250V



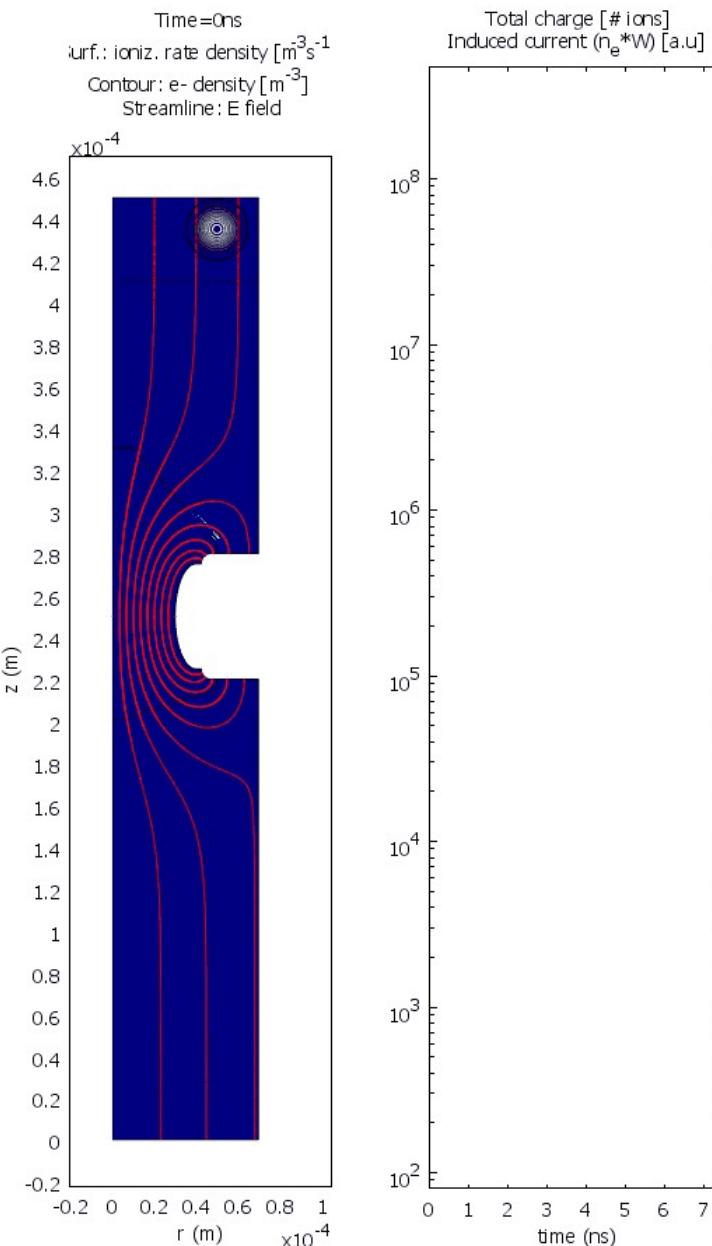
# GEM lateral (ring) avalanche

hole: 60 µm

gap: 100 µm

$N_0=100$  e<sup>-</sup>

V=1250V



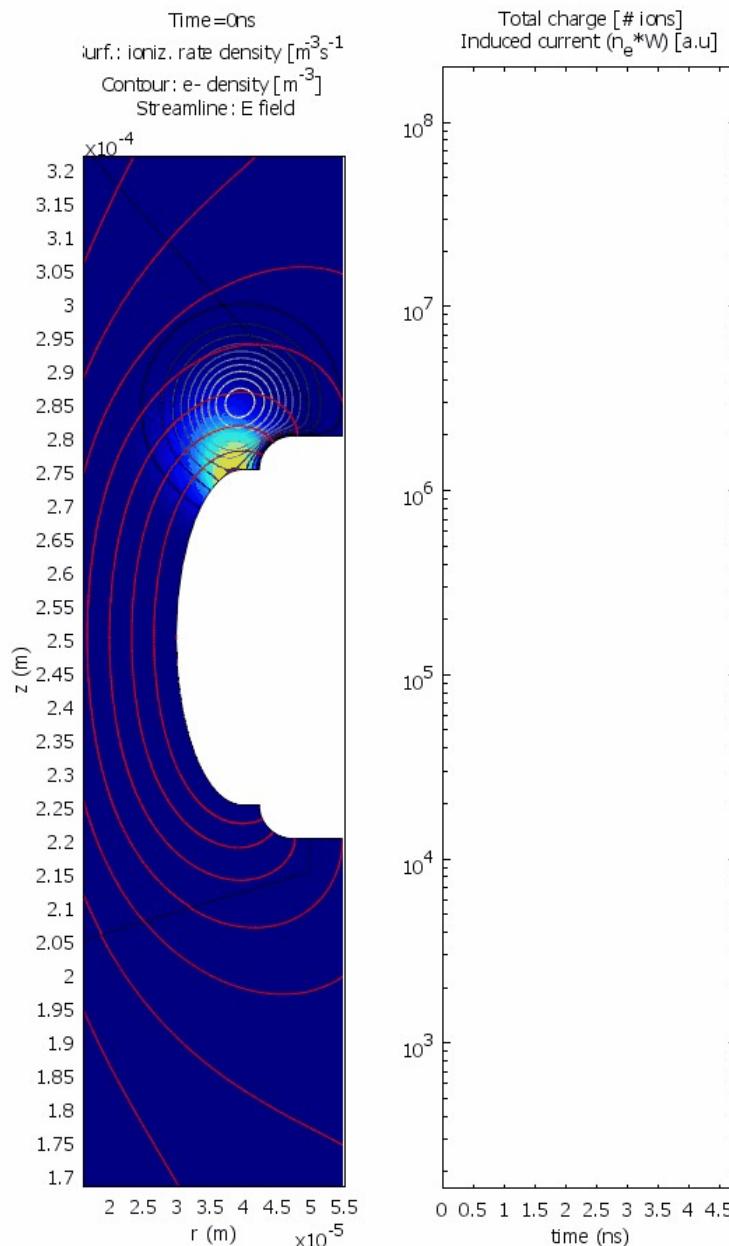
# GEM surface avalanche

hole: 60 µm

gap: 100 µm

$N_0=100$  e<sup>-</sup>

V=1150V



# GEM surface avalanche

hole: 60 µm

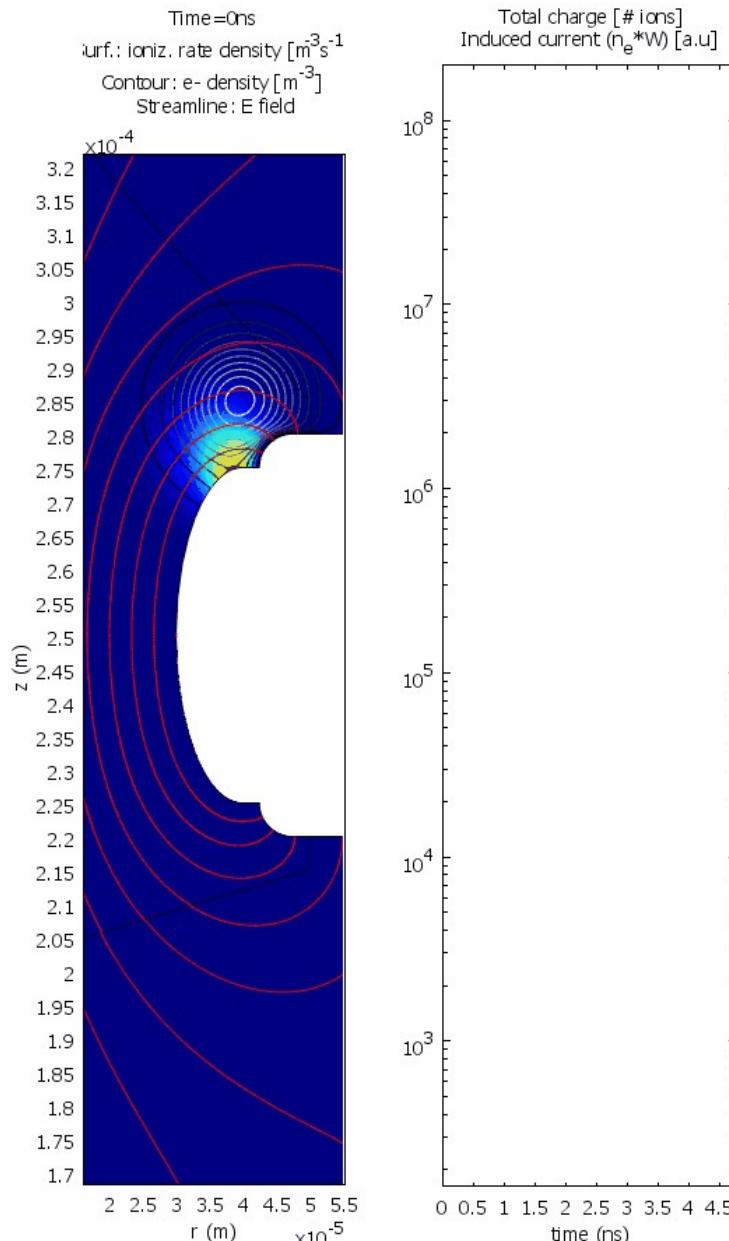
gap: 100 µm

$N_0=100$  e<sup>-</sup>

V=1150V

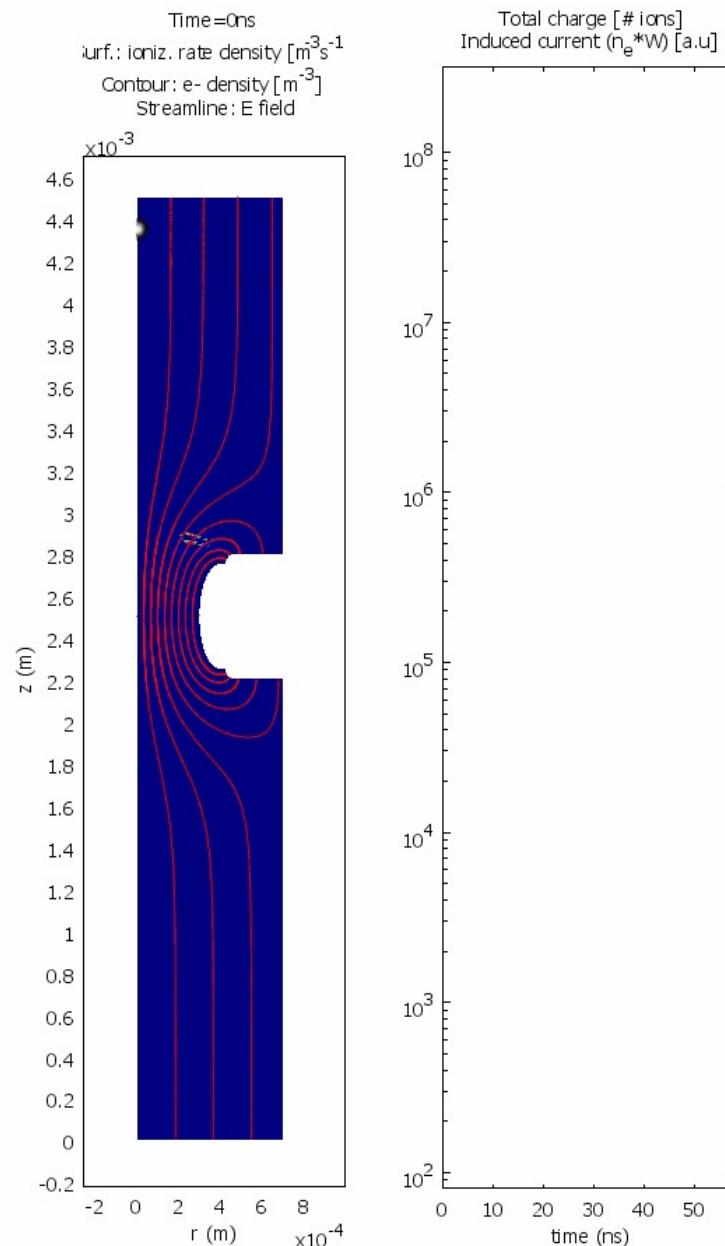
This happens 100V below the streamer limit in the space, limiting the practical GEM gain.

Solved by multisteping.



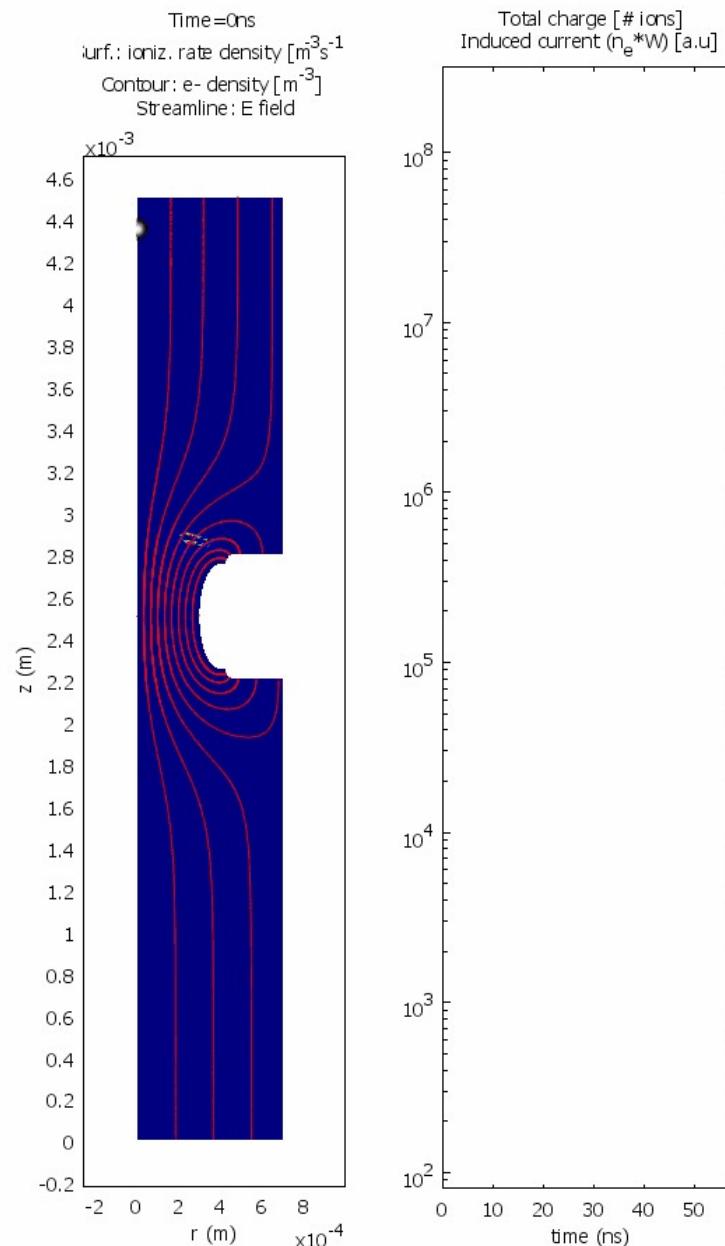
# THGEM (GEMx10)

hole: 600 µm  
gap: 1 mm  
 $N_0=100$  e<sup>-</sup>  
 $V=4600V$



# THGEM (GEMx10)

hole: 600 µm  
gap: 1 mm  
 $N_0=100$  e<sup>-</sup>  
 $V=4600V$

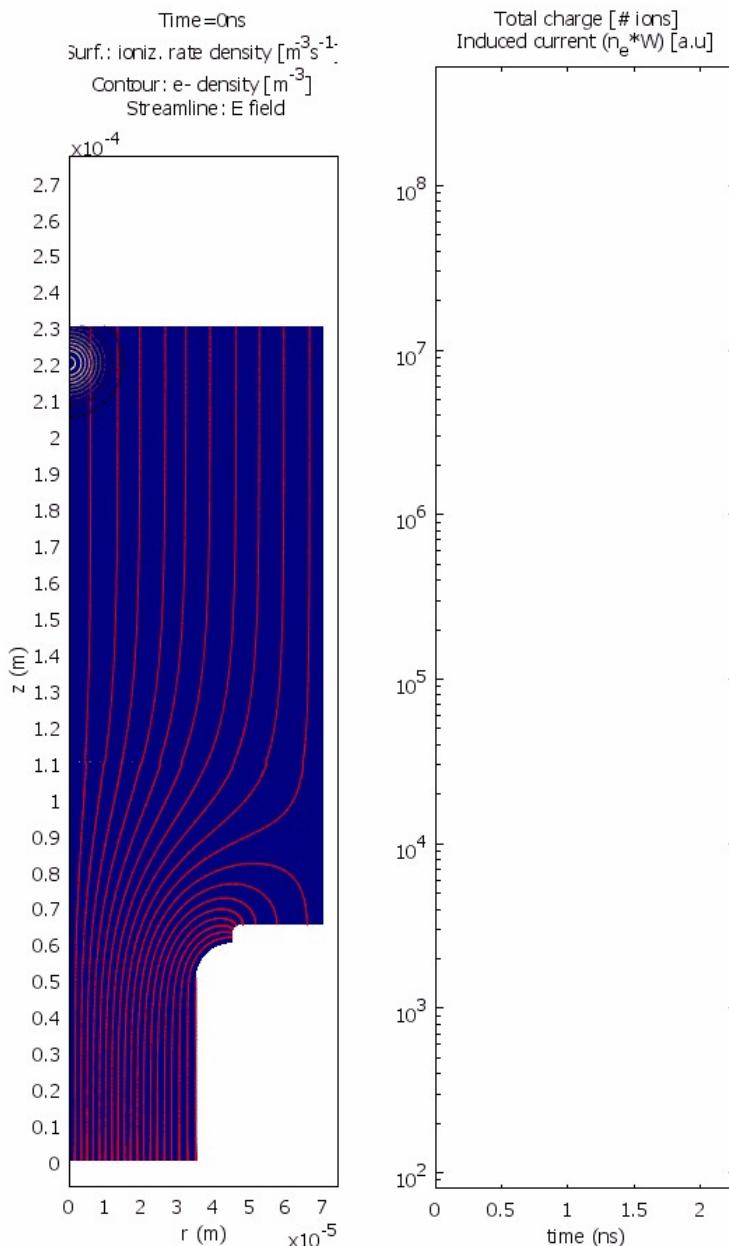


# CAT

diameter: 70  $\mu\text{m}$

height: 650  $\mu\text{m}$

$N_0=100 \text{ e}^-$

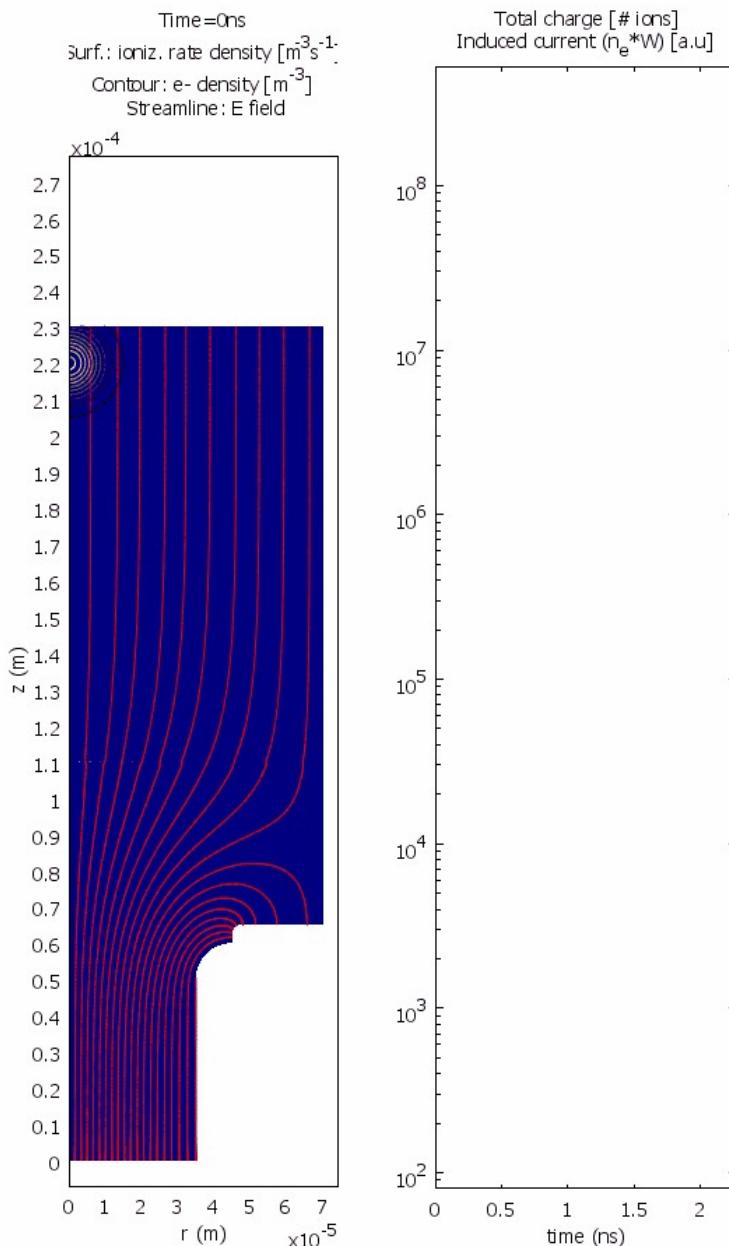


# CAT

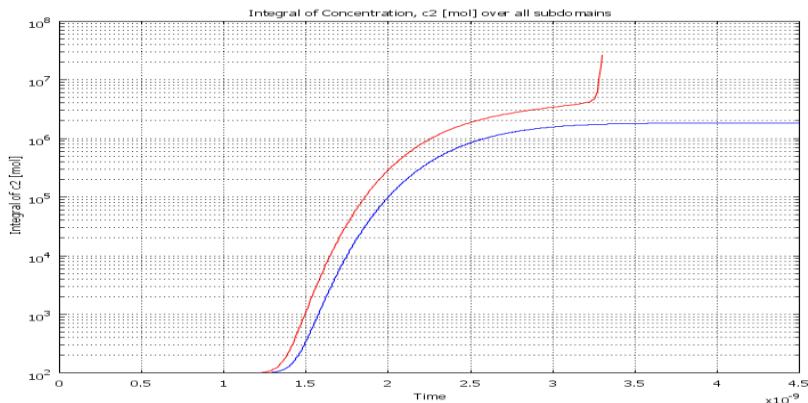
diameter: 70  $\mu\text{m}$

height: 650  $\mu\text{m}$

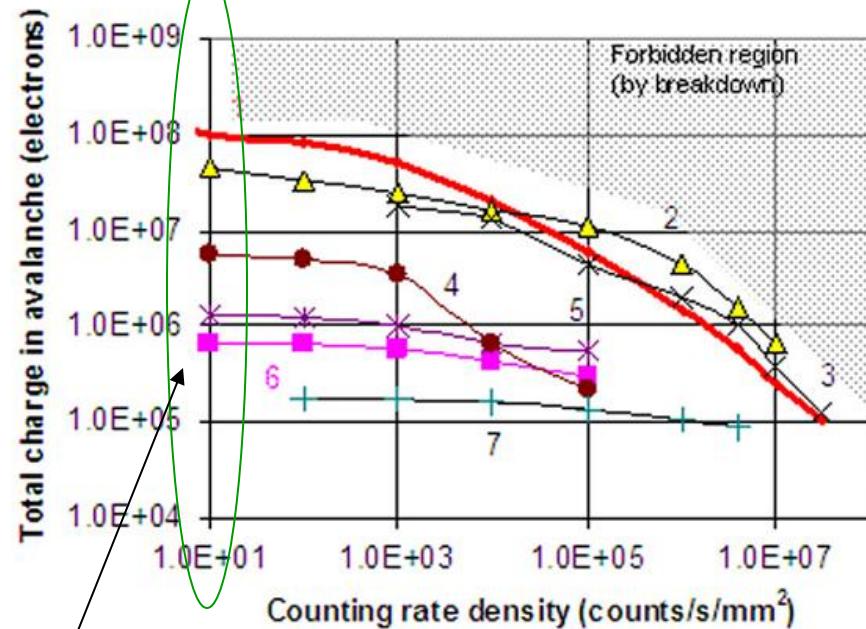
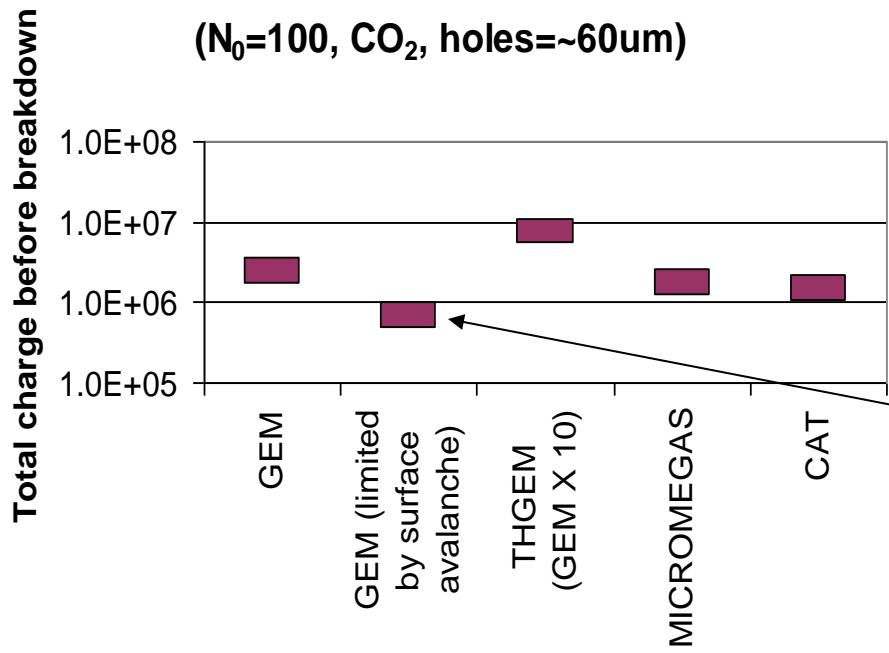
$N_0=100 \text{ e}^-$



# Determination of the space-charge limit



**BREAKDOWN LIMIT**  
 $(N_0=100, \text{CO}_2, \text{holes}=\sim 60\mu\text{m})$



**Figure 4.** The maximum achievable gain, limited by breakdown, as a function of the x-ray flux for various detectors: (1) PPAC with 3mm gap; (2) MICROMEGAS; (3) PPAC with 0.6 mm gap; (4) microstrip gas chamber with 1 mm strip pitch; (5) microstrip gas chamber with 0.2 mm strip pitch; (6) GEM; (7) microgap detectors with 0.2 mm strip pitch. Large counting rate densities require a

~ OK for GEM  
Much lower for MM.  
(maybe not the same exact geometry,  
gas, etc)

# Streamer-resistant detectors?

single-wire (SQS mode)

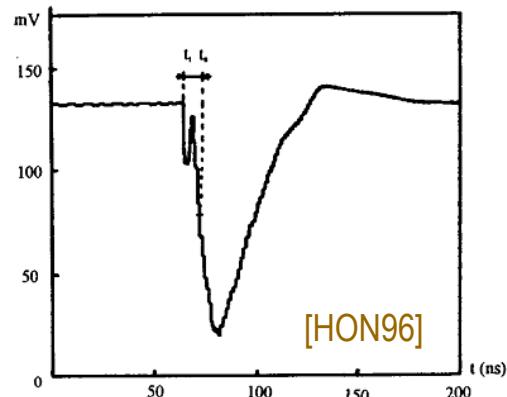
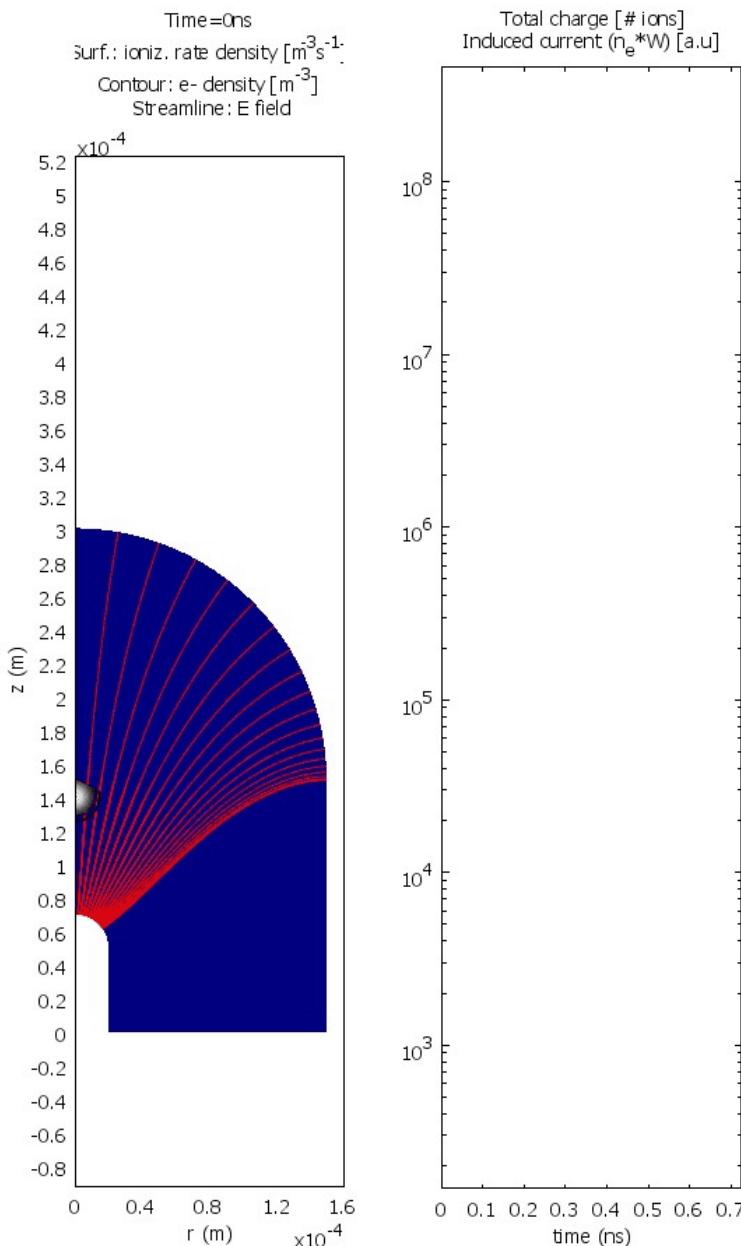


Fig. 1. The pulse shape of the SQS electrical signal  $V = 2.45$  kV.  
 $\text{Methylal}/(\text{Methylal} + \text{Ar}) = 16.6\%$ .

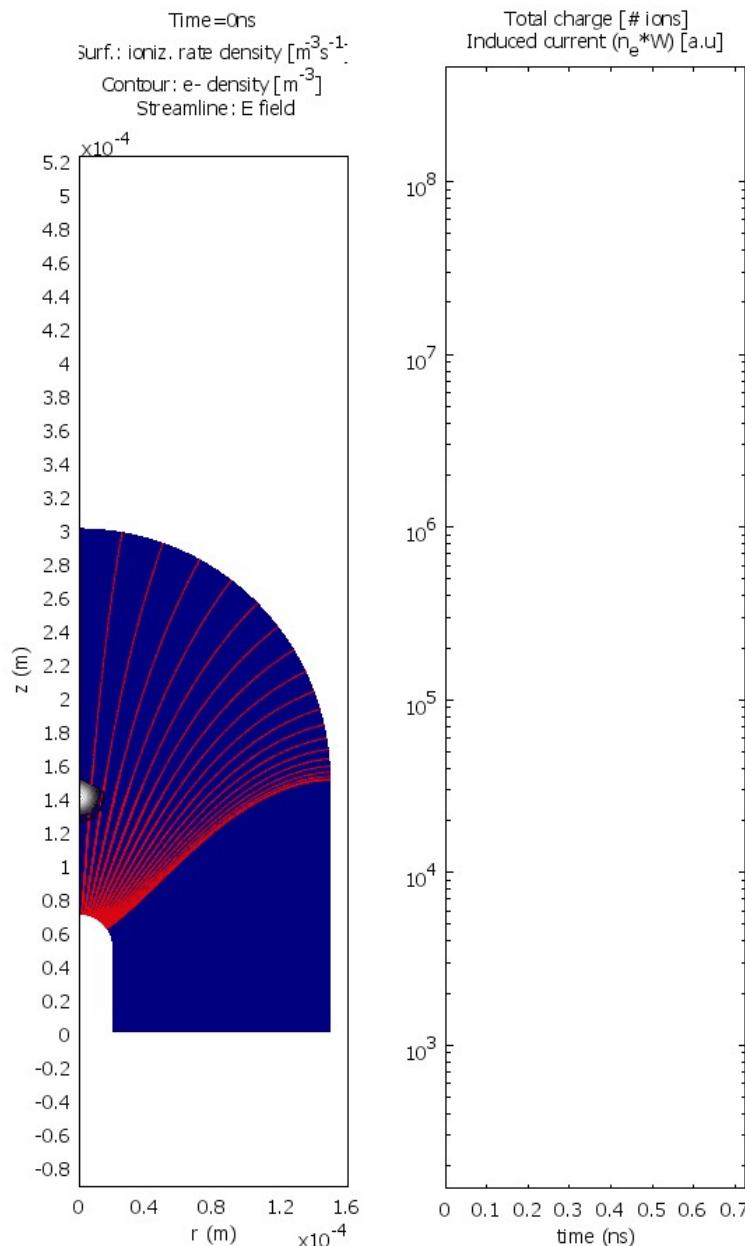
Could there be a self-quenching streamer (SQS) mode in MPGDs?

# SQS around needles

diameter: 40 µm  
 $N_0=100 e^-$



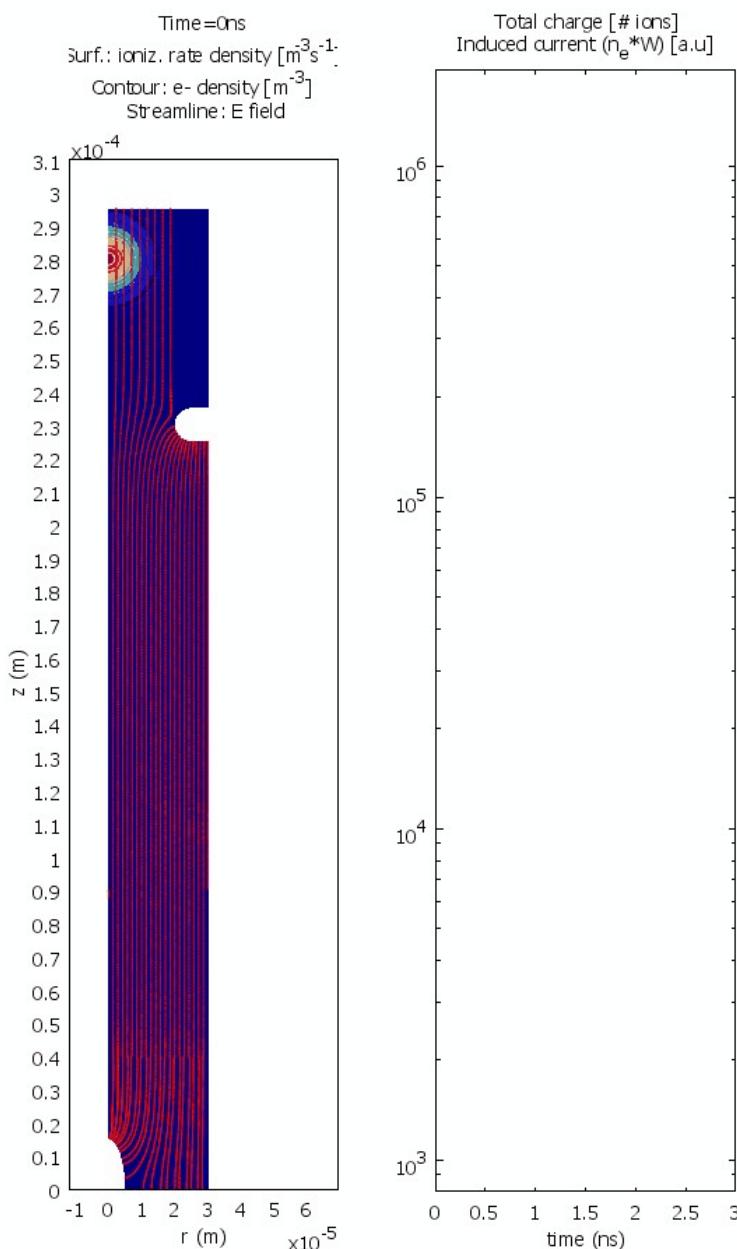
# SQS around needles



# Array of needles + InGRID

diameter: 10  $\mu\text{m}$   
 $N_0=1000 \text{ e}^-$

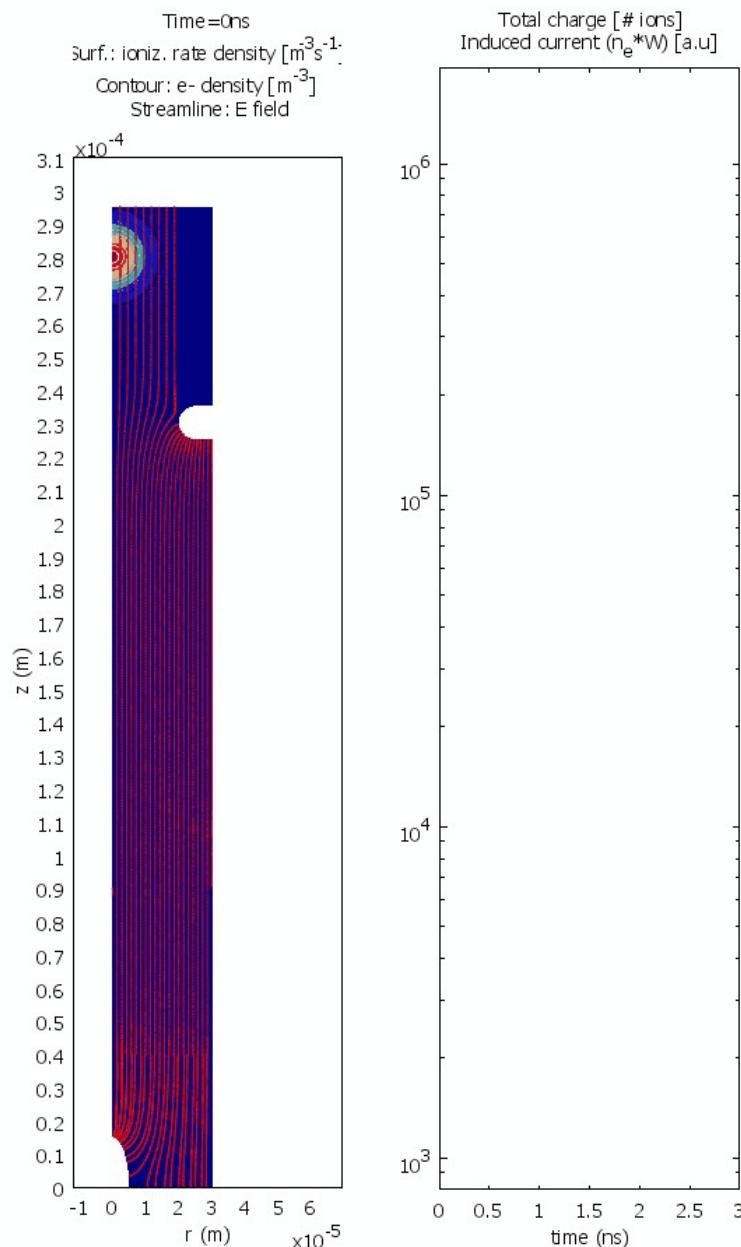
Or course:  
needle  $\Rightarrow$  low rate capability



# Array of needles + InGRID

diameter: 10  $\mu\text{m}$   
 $N_0=1000 \text{ e}^-$

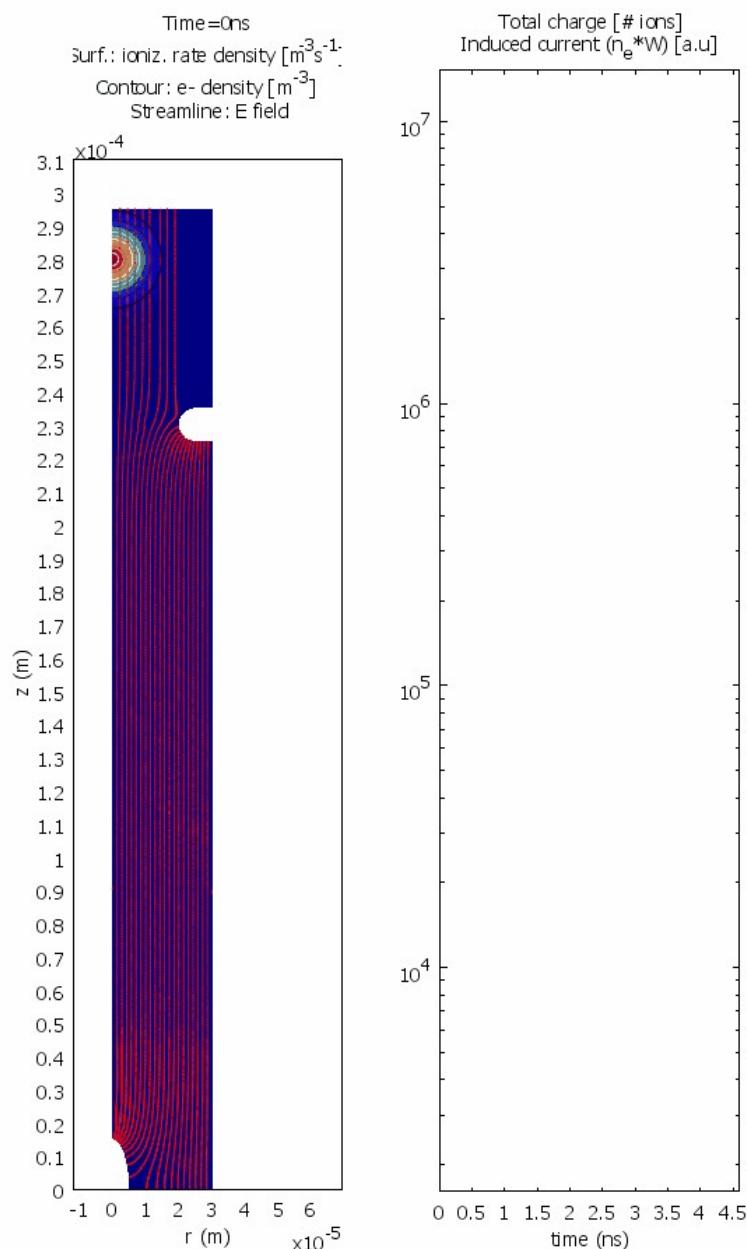
It seems there is indeed an  
SQS regime in the needle



# Array of needles + InGRID

diameter: 10  $\mu\text{m}$   
 $N_0=2000 \text{ e}^-$

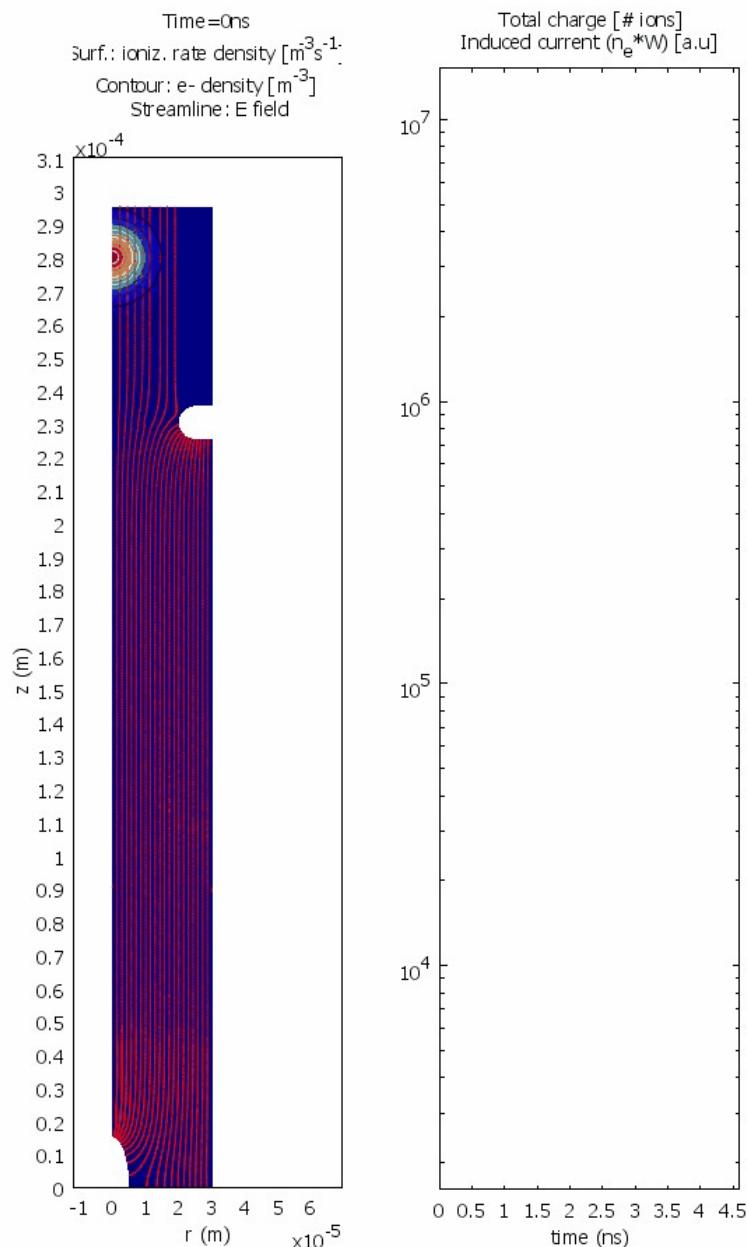
Larger charge



# Array of needles + InGRID

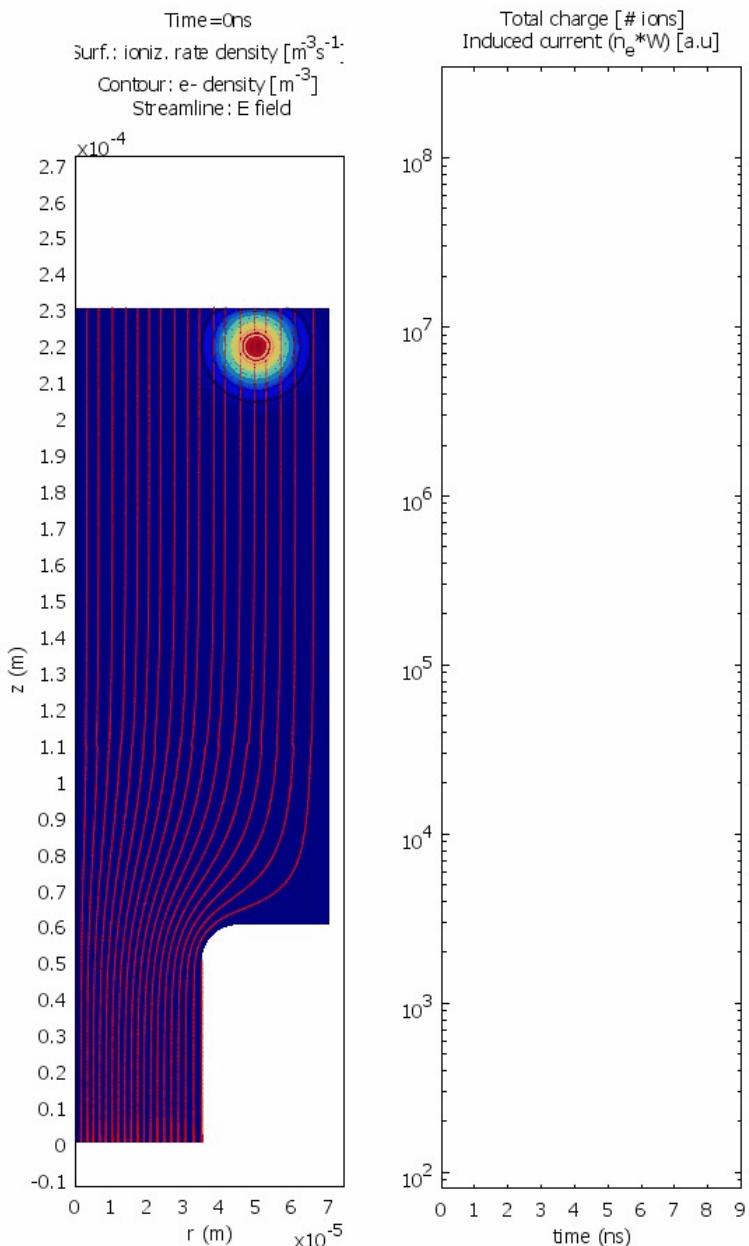
diameter: 10  $\mu\text{m}$   
 $N_0=2000 \text{ e}^-$

It seems there is indeed an SQS regime in the needle, but there is still too much parallel-field gain.



# Cathodeless CAT

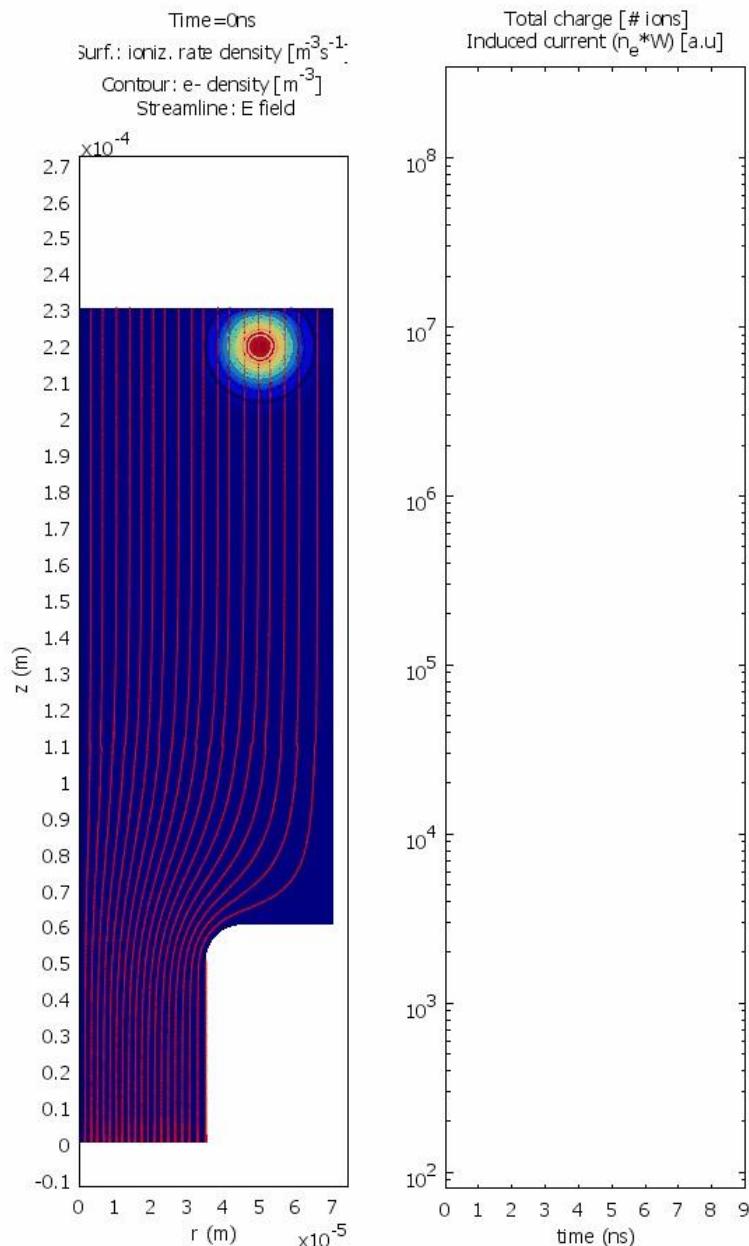
Will the streamer be able to grow out of the hole?



# Cathodeless CAT

Not completely successful, but it sparks at a rather large charge.

Maybe such geometries can be optimized for SQS mode.



## Summary

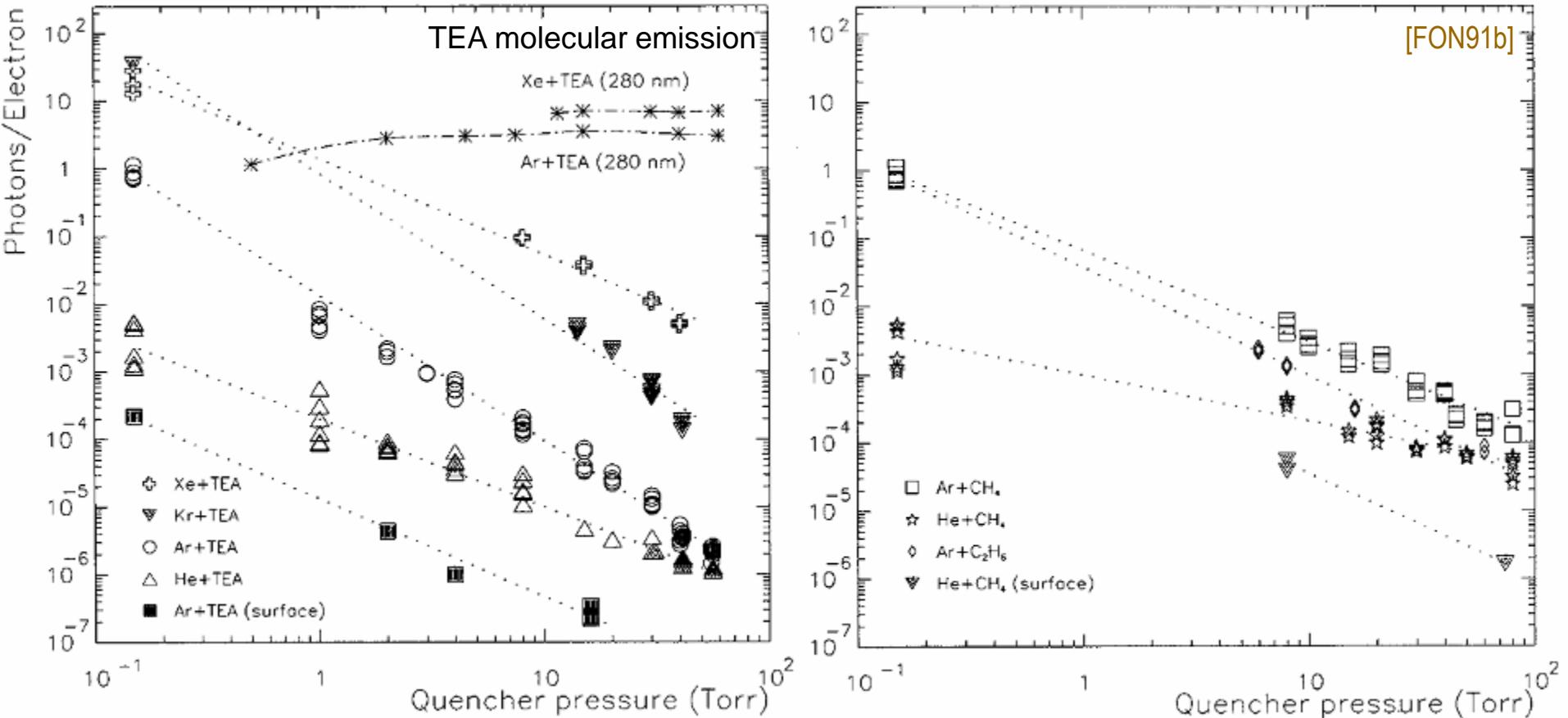
- Streamer breakdown is a recognizable feature (precursor-discharge structure) of several types of gas detectors
- The corresponding hydrodynamic model seems to describe qualitatively fast breakdown also in micropattern detectors
- Streamers can be supported by diffusion alone
- This seems to be qualitatively more in agreement with the empirical observations in detectors than the classical mechanism based on self-photoionization
- Gives correct breakdown limit for GEM
- Seems to reproduce SQS in needles
- Useful tool for detector design and optimization. No full SQS so far...

## References

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# Emission suppression by “quenchers”

Photon yields in PPAC in the band: 120-170nm



There is some evidence that the emission originates mainly from fragments (likely carbon atomic emission lines) at  $\lambda > 140\text{nm}$ .

Photoemission strongly suppressed for quencher concentration 1-10%

If “quenching” was the answer, people ought to have found a sparkless gas, or at least one that would strongly expand the sparking limit.