



Bernhard Ketzer University of Bonn

XIV ICFA School on Instrumentation in Elementary Particle Physics LA HABANA 27 November - 8 December, 2017



universität**bonn**

- Overview: old and new detectors
- Detection principles
- Interaction of charged particles with matter
 - Inelastic collisions with atomic electrons ⇒ ionization energy loss
 - Emission of Cherenkov radiation
 - Emission of Transition radiation
 - ⇒ unified treatment in PAI model
- Mean energy loss: Bethe formula and friends
- Energy loss distributions (straggling functions): Landau et al.
- Multiple scattering
 - important for smaller momenta
 - contribution to momentum resolution





Principle: collection of electrons and ions (holes) produced in detector medium by ionizing radiation

Detector material:

- gas ⇒ fast collection of e⁻ and ions, e.g. Ne, Ar
- liquid ⇒ higher density, e.g. liquid Ar
- solid ⇒ higher density, self-supporting, e.g. semiconductor

Setup:

- vessel with two electrodes and thin entrance window
- filled with active medium
 ⇒ creation of electron-ion (hole) pairs
- electric field between anode and cathode
 - separation of e⁻ and ions (holes), drift and diffusion
 - signal induction
 - collection at anode/cathode



Plan of the Lecture



- 1. Introduction
- 2. Interactions of charged particles with matter
- 3. Drift and diffusion of charges in gases
- 4. Avalanche multiplication of charge
- 5. Signal formation and processing
- 6. Ionization and proportional gaseous detectors
- 7. Track reconstruction and momentum measurement





3 Drift and Diffusion of Charges in Gases

3.1 Drift of charge carriers: equation of motion3.2 Microscopic picture3.3 Diffusion

3.1 Drift of Charge Carriers

Microscopically:

External electric field \Rightarrow acceleration

Collisions ⇒ slowing down

Macroscopically: drift motion with drift velocity u

Ansatz: Langevin equation

$$m\frac{\mathrm{d}\boldsymbol{u}}{\mathrm{d}t} = e\boldsymbol{E} + e(\boldsymbol{u} \times \boldsymbol{B}) - K\boldsymbol{u}$$

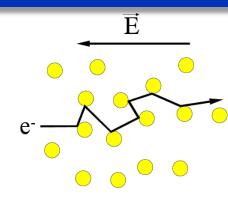
Solution for $t \gg \tau$: steady state for which $\frac{\mathrm{d}\boldsymbol{u}}{\mathrm{d}t} = 0$

Ku = frictional force $\tau = \frac{m}{\nu}$ = characteristic time

 $\omega = \frac{eB}{m}$ = cyclotron frequency

 \Rightarrow Drift velocity *u* dominated by dimensionless parameter $\omega \tau$

 $\boldsymbol{u} = \frac{e}{m} \tau \left| \boldsymbol{E} \right| \frac{1}{1 + \omega^2 \tau^2} \left[\hat{\boldsymbol{E}} + \omega \tau \left(\hat{\boldsymbol{E}} \times \hat{\boldsymbol{B}} \right) + \omega^2 \tau^2 \left(\hat{\boldsymbol{E}} \cdot \hat{\boldsymbol{B}} \right) \hat{\boldsymbol{B}} \right]$



universität**bonn**







$$\boldsymbol{u} = \frac{e}{m}\tau \left|\boldsymbol{E}\right| \frac{1}{1+\omega^{2}\tau^{2}} \left[\hat{\boldsymbol{E}} + \omega\tau\left(\hat{\boldsymbol{E}}\times\hat{\boldsymbol{B}}\right) + \omega^{2}\tau^{2}\left(\hat{\boldsymbol{E}}\cdot\hat{\boldsymbol{B}}\right)\hat{\boldsymbol{B}}\right]$$

Drift velocity u dominated by dimensionless parameter $\omega \tau$ (carries charge sign):

- $\omega \tau = 0 \Rightarrow \mathbf{u} \parallel \mathbf{E}$ $\mathbf{u} = \frac{e}{m} \tau \mathbf{E} = \mu \mathbf{E}, \qquad \mu = \frac{e}{m} \tau$ (scalar mobility, carries charge sign)
- with magnetic field:

 \Rightarrow generalized mobility: $\mu = \frac{e}{m}M^{-1}$ (tensor of 2nd order)

• $\omega \tau \gg 1, \ e > 0$

$$- \hat{\mathbf{E}} \cdot \hat{\mathbf{B}} \neq 0 \Rightarrow \mathbf{u} \parallel \mathbf{B}$$

 $\hat{\mathbf{E}} \cdot \hat{\mathbf{B}} = 0 \Rightarrow \mathbf{u} \parallel (\mathbf{E} \times \mathbf{B})$



Drift of Charge Carriers

Case of *E* orthogonal to *B*:

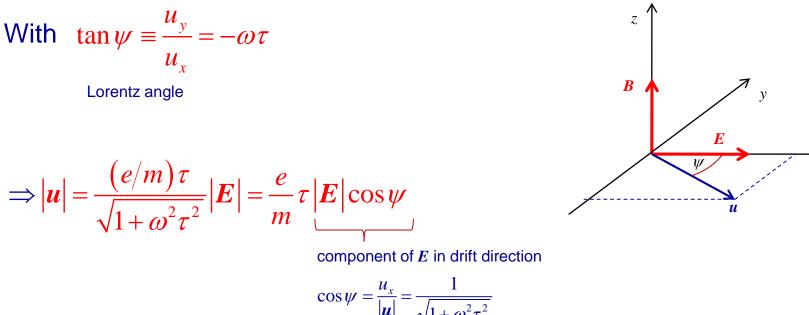
$$\hat{\boldsymbol{E}} \cdot \hat{\boldsymbol{B}} = 0, \quad \boldsymbol{E} = \begin{pmatrix} \boldsymbol{E}_x \\ 0 \\ 0 \end{pmatrix}, \quad \boldsymbol{B} = \begin{pmatrix} 0 \\ 0 \\ B_z \end{pmatrix}$$

$$\Rightarrow u_{x} = \frac{(e/m)\tau}{(1+\omega^{2}\tau^{2})} |\mathbf{E}|$$
$$u_{y} = -\frac{(e/m)\tau}{(1+\omega^{2}\tau^{2})} \omega\tau |\mathbf{E}|$$
$$u_{z} = 0$$

$$\tan\psi \equiv \frac{u_y}{u_x} = -\omega\tau$$

universität**bonn**

Drift of Charge Carriers



i.e. magnitude of drift velocity is determined by component of electric field in drift direction

Gas Detectors

universität**bonn**

х



Numerical example:

- $\mu = 10^4 \text{ cm}^2 \text{V}^{-1} \text{s}^{-1}$ for electrons
 - $\mu = 1 \text{ cm}^2 \text{V}^{-1} \text{s}^{-1} \qquad \text{for ions}$
 - $B = 1T = 10^{-4} \text{ Vs cm}^{-2}$ typical

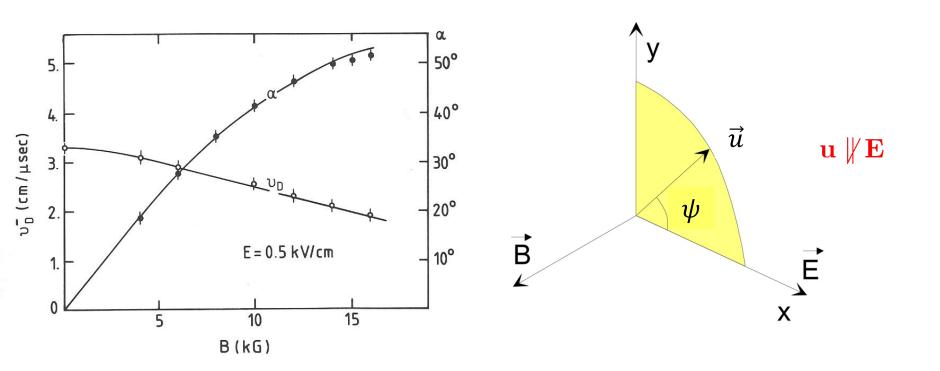
 $\Rightarrow \omega \tau = B \mu \sim \begin{cases} 10^{-4} & \text{for ions} \\ 1 & \text{for electrons, i.e. } \psi = 45^{\circ} \end{cases}$

⇒ effect on ion drift negligible

universität**bonn**



Drift of Charge Carriers



Electron drift velocity and Lorentz angle (E orthogonal to B)

[A. Breskin et al., NIM 124, 189 (1975)]

universität**bonn**



Drifting electrons and ions are scattered on gas molecules
 ⇒ Direction of motion randomized in each collision
 ⇒ "stop & go" motion of individual particles



• constant drift velocity *u* in the direction given by *E* (or *E* and *B*)

Goal: derive basic relations between

- Microscopic quantities of instantaneous velocity c, mean time τ between collisions, and fractional energy loss Λ
 - ⇒ distributed according to distribution functions!
 - ⇒ here: use suitable averages
- Macroscopic quantities of drift velocity *u* and isotropic diffusion coefficient *D* and

universität**bonn**

É





Microscopic picture: electron between two collisions

$$m\frac{\mathrm{d}\boldsymbol{v}}{\mathrm{d}t} = e\boldsymbol{E} \quad \Rightarrow \quad \boldsymbol{v}(t) = \frac{e\boldsymbol{E}}{m}t + \boldsymbol{v}(0)$$

Average in time:

$$\langle \mathbf{v}(t) \rangle \equiv \mathbf{u} = \frac{e\mathbf{E}}{m} \langle t \rangle + \langle \mathbf{v}(0) \rangle$$

 $\langle t \rangle = \tau$ since $\langle t \rangle = \int_{0}^{\infty} t e^{-t/\tau} \frac{\mathrm{d}t}{\tau} = \tau$

average time since last collision = average time between collisions

For $m \ll M \Rightarrow$ electron scatters isotropically, i.e. no preferential direction after collision

- $\Rightarrow \left\langle \boldsymbol{\nu}\left(0\right)\right\rangle = 0$
- $\Rightarrow u = \frac{eE}{m}\tau$
- extra velocity gained by e- in electric field between collisions, in addition to its instantaneous, randomly oriented velocity
- corresponds macroscopically to drift velocity





Next encounter of gas molecule ⇒ extra energy is lost (on average), i.e. there is a balance between energy picked up and collision losses

Drift distance x: average number of collisions $N = \frac{t}{\tau} = \frac{x}{u} \cdot \frac{1}{\tau}$

$$\Rightarrow eEx = \frac{x}{u\tau} \cdot \Lambda \cdot \mathcal{E}_E \qquad \Lambda = \text{average relative energy los}$$

$$\varepsilon_{\text{F}} = \text{equilibrium energy (only particular)}$$

Λ = average relative energy loss per collision $ε_F$ = equilibrium energy (only part due to electric field, not thermal energy)

With
$$\frac{1}{\tau} = n\sigma c$$
 and $\varepsilon = \frac{1}{2}mc^2 = \varepsilon_E + \frac{3}{2}kT \approx \varepsilon_E$ $\varepsilon_E \gg \frac{3}{2}kT$ usually fulfilled for e- in detectors

follows: $u^2 = \frac{eE}{mn\sigma} \sqrt{\frac{\Lambda}{2}}$ drift velocity $\Rightarrow u \propto \sqrt{E}$ if Λ and σ const. $\Rightarrow \mu \propto \frac{1}{\sqrt{E}}$ $c^2 = \frac{eE}{mn\sigma} \sqrt{\frac{2}{\Lambda}}$ average instantaneous velocity





- rapid acceleration in electric field
- small energy loss in elastic collisions with atoms
- e⁻ momentum randomized in collisions
- energy gain in electric field is mainly in random motion >> thermal energy

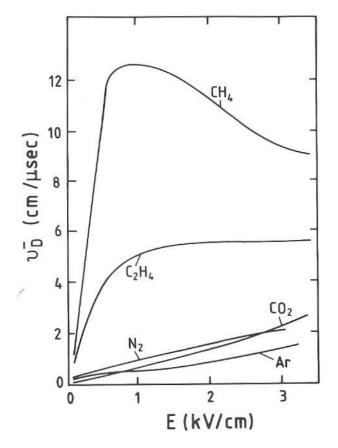
$$\Rightarrow \text{ drift velocity} \qquad u^2 = \frac{eE}{mn\sigma} \sqrt{\frac{\Lambda}{2}} \ll c^2 = \frac{eE}{mn\sigma} \sqrt{\frac{2}{\Lambda}} \qquad \text{average velocity}$$

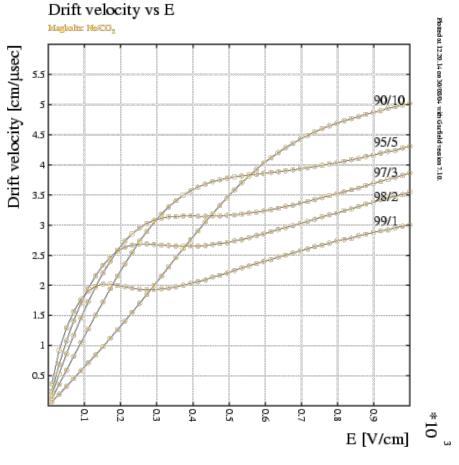
 $\Lambda = \Lambda(\varepsilon)$ average fractional energy loss per collision $\sigma = \sigma(\varepsilon)$ collision cross section 10-1 10-14 λ σ (cm²) Ramsauer minimum (Ar, CH₄, Kr, Xe) 10^{-2} 10-15 CH4 CH, 10-3 AΓ 10-16 10-4 Ar 10-17 10-5 0.001 0.01 0.1 10 0 001 0.01 01 10 (b) [Blum,Rolandi, Springer, 1993] E (eV) (a) ε (eV) B. Ketze

Gas Detectors



Drift Velocity of Electrons





universität**bonn**

Ne/CO₂ Mixtures



3.2.2 Drift of lons



- Only elastic collisions with other gas atoms/molecules
- Two limiting cases:
 - Low field \Rightarrow ion random velocity is thermal (never reached for e-)

$$\Rightarrow u = \left(\frac{1}{m} + \frac{1}{M}\right)^{1/2} \left(\frac{1}{3kT}\right)^{1/2} \frac{eE}{n\sigma} \propto E$$

• High field ⇒ neglect thermal motion (this is the general case for e-)

$$\Rightarrow u = \left(\frac{eE}{mn\sigma}\right)^{1/2} \left[\frac{m}{M}\left(1+\frac{m}{M}\right)\right]^{1/2} \propto \sqrt{E}$$

for ions ($\sigma \sim \text{const.}$, Λ only from elastic collisions)



Field-free gas: quick thermalization of charge carriers in collisions

⇒ Maxwell distribution of velocities (thermal equilibrium):

$$F(c)dc = 4\pi n \left(\frac{m}{2\pi kT}\right)^{3/2} c e^{-mc^2/(2kT)} dc \qquad \overline{c} = \sqrt{\frac{8kT}{\pi m}}$$
$$\overline{E}_{kin} \equiv \varepsilon = \frac{m}{2}\overline{c^2} = \frac{3}{2}kT \qquad \text{Definition of temperature}$$

Point-like charge cloud at t=0 ⇒ Distribution at time t?

Ansatz: $\mathbf{J} = -D \nabla n$ \Rightarrow diffusion equation $\frac{\partial n}{\partial t} = D \Delta n$ Solution: Gaussian law $\frac{dN}{N} = \frac{ndz}{N} = \frac{1}{\sqrt{4\pi Dt}} e^{-z^2/4Dt} dz$ $\sqrt{\langle r^2 \rangle} \equiv \sigma_r = \sqrt{6Dt}$





Microscopic picture: consider electron/ion starting at t = 0 with velocity c (1-D) After $N = t/\tau$ collisions (statistically independent) \Rightarrow displacement x

$$x = \sum_{i=1}^{N} \xi_i$$

$$\overline{x} = \sum_{i=1}^{N} \overline{\xi_i} = 0$$

$$\overline{x^2} = N\overline{\xi_i^2}$$

 $\xi_i = x$ -component of *i*th displacement

since $\xi_i = 0$ (isotropic scattering, fulfilled for e- and ions for very small fields)

since probability distribution is equal for each step

With $\xi = c_x t$ follows $\overline{\xi^2} = \overline{c_x^2} \overline{t^2} = \frac{1}{3} \overline{c^2} \overline{t^2}$ symmetry $\overline{t^2} = \int t^2 e^{-t/\tau} \frac{\mathrm{d}t}{\tau} = 2\tau^2 \qquad \Rightarrow \overline{\xi^2} = \frac{2}{3} \overline{c^2} \tau^2$ $\Rightarrow \overline{x^2} = N \overline{\xi^2} = \frac{2}{3} \overline{c^2} \tau t \equiv \sigma_x^2$





Comparison with solution of diffusion equation: $\sigma_x = \sqrt{2Dt}$

$$\Rightarrow D = rac{\overline{c^2} au}{3} = rac{2}{3} rac{arepsilon}{m} au$$
 with $arepsilon = rac{1}{2} m \overline{c^2}$

With $\mu = \frac{e}{m}\tau$ follows $\varepsilon = \frac{3}{2}\frac{De}{\mu}$ (valid for ions and electrons!) \Rightarrow determination of average e⁻ kinetic energy by measurement of D/μ

Diffusing body in thermal equilibrium with environment: $\varepsilon = \frac{3}{2}kT$ (fulfilled for ions at low fields only!)

$$\Rightarrow \frac{D}{\mu} = \frac{kT}{e}$$

Nernst-Townsend formula

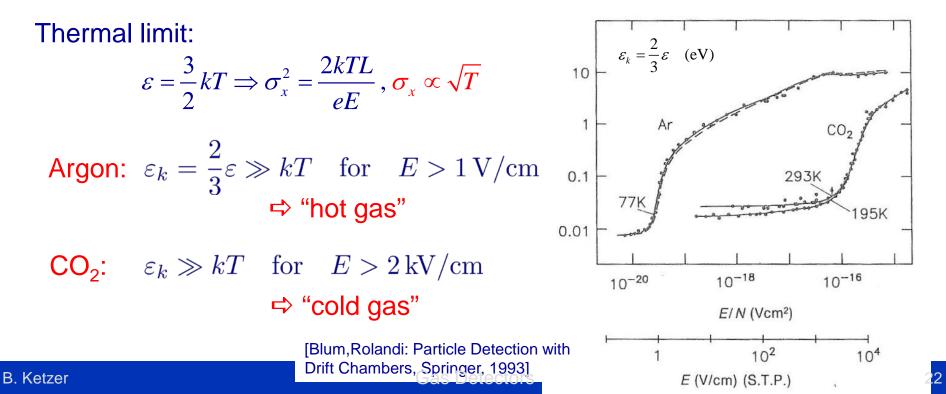


Drift and Diffusion

universität**bonn**

Diffusion depends on average kinetic energy: $\varepsilon = \frac{3}{2} \frac{De}{\mu}$ from kinetic gas theory Width of electron cloud after drifting distance L: $\sigma_x^2 = 2Dt = \frac{2DL}{\mu E} = \frac{4\varepsilon L}{3eE}$

⇒ small e⁻ energies at high drift fields required for small diffusion







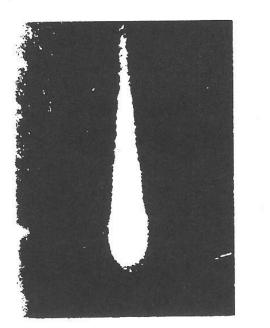
4 Avalanche Multiplication of Charge

4.1 Gas amplification4.2 Choice of detector gas4.3 Statistical fluctuations of gain





Increase electric field in gaseous detector up to several kV/cm ⇒electrons gain sufficient energy between collisions to ionize gas molecules ⇒avalanche generation



Drift velocity $u_e \gg u_{ion}$ Diffusion

drop-like distribution of charges
 (in homogeneous field)

- e- in the front
- tail of positive ions

[L.B. Loeb, Basic Processes of Gaseous Electronics, Univ. Cal. Press (1961)]





Probability of ionization per unit path length: 1. Townsend coefficient

$$\alpha = \frac{1}{\lambda_{\rm ion}} = n\sigma_{\rm ion}$$

 λ_{ion} = mean free path for secondary ionizing collision σ_{ion} = cross section for ionizing collision n = density of gas molecules

Avalanche development: N electrons in avalanche

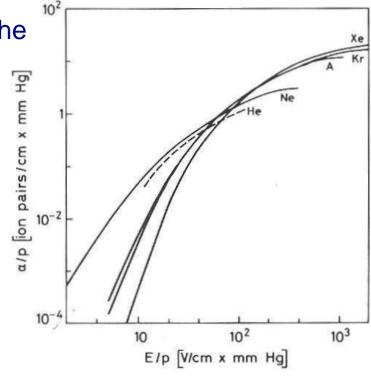
• Homogeneous electric field:

$$\mathrm{d}N = N\alpha\mathrm{d}s \Rightarrow N(s) = N_0 e^{\alpha s}$$

Gain:
$$G = \frac{N}{N_0} = e^{\alpha s}$$

• Inhomogeneous electric field: $\alpha = \alpha(s)$

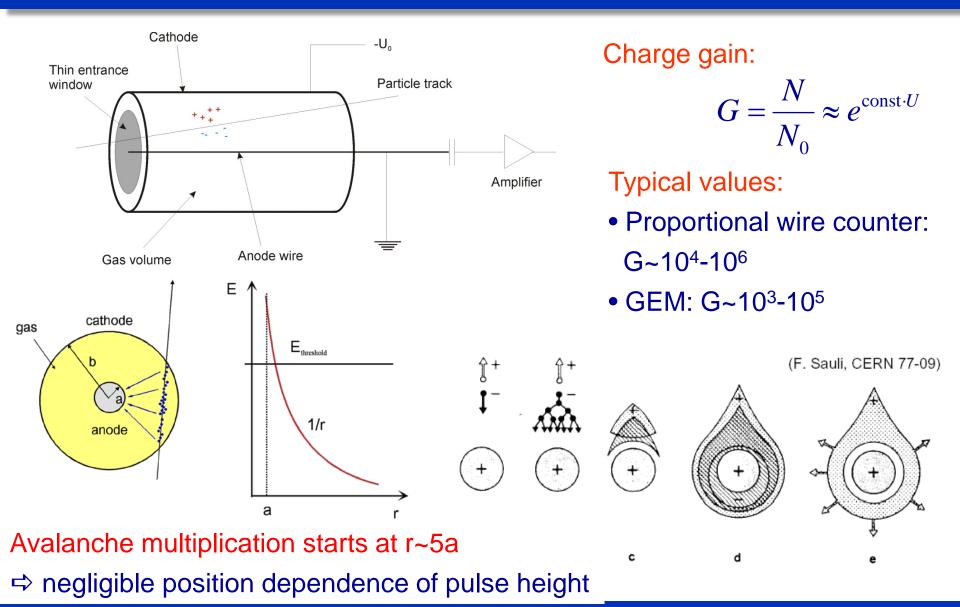
$$G = \exp\left[\int_{s_1}^{s_2} \alpha(s) ds\right] = \exp\left[\int_{E_1}^{E_2} \frac{\alpha(E)}{dE/ds} dE\right]$$



[S.C. Brown, Basic Data of Plasma Physics, MIT Press (1959)]

Proportional Counter





B. Ketzer

Gas Detectors

26





In principle: avalanche formation occurs in every gas, depending on the electric field

Determines substantially the properties of the detector Experimental requirements:

- low operating voltage
- high gain
- good proportionality
- high rate capability
- long lifetime, no aging

⇒ partly contradicting each other

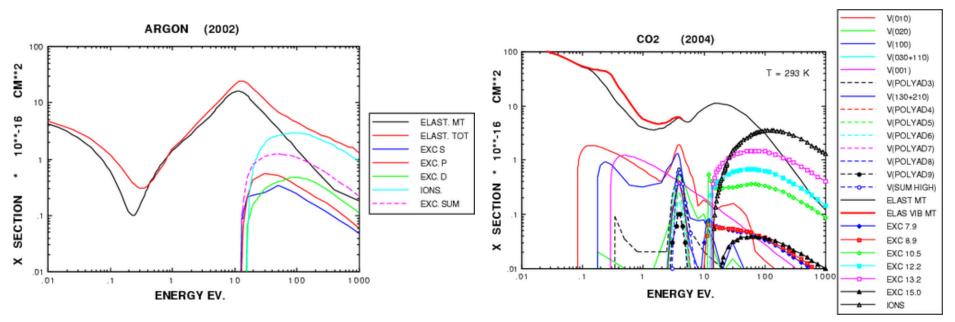
⇒ no ideal gas meeting all requirements





Noble gases: main component

- avalanche multiplication occurs at much lower fields than in complex molecules
- energy dissipation mostly through ionization, in contrast to many non-ionizing energy dissipating modes in polyatomic molecules







Choice of noble gas ⇒ high specific ionization by MIPs (particle physics)

	Хе	Kr	Ar	Ne	Не
Z	54	36	18	10	2
W (eV)	22	24	26	36	41
n _T (ip/cm)	307	192	94	39	7.8

Xe, Kr: expensive, multiple scattering ⇒ deflection of incoming particleHe: high leak rate

Ne: ~8× more expensive than Ar ⇔ closed gas system necessary

Ar: commonly used

But: continuous discharge for $G \ge 10^3 - 10^4$!





Reason: formation of excited and ionized noble gas atoms in avalanche

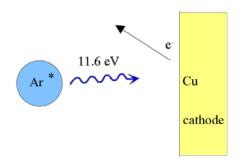
- a) Excited noble gas atoms:
- deexcitation through radiative transitions

 \Rightarrow emission of photons with $hv = E_{ex}$ (11.6 eV for Ar, 16.6 eV for Ne)

• electrodes (metal): $hv > E_w$ (4.4 eV for Cu)

⇒ photons can extract photoelectrons from electrodes

⇒ new avalanches at different locations in detector







- b) Ionized noble gas atoms:
- drift to cathode where they are neutralized by extracting an electron
- conservation of energy
 - emission of photon
 - extraction of another electron from cathode (secondary emission)
 - ⇒ further avalanches, delayed with respect to original event

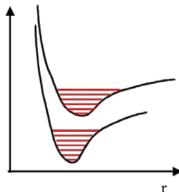




Solution: addition of quench gas, i.e. polyatomic molecules, e.g. CH₄, CO₂, BF₃, ...

a)

- absorption of photons in a wide energy range (7.9 14.5 eV for CH4) through excitation of rotational and vibrational levels
- energy dissipation via dissociation or elastic collisions
 b)
- charge exchange noble gas → quench gas very efficient due to low ionization potential of molecules
- drift of ionized molecule to cathode, neutralisation
- emission of secondary radiation at cathode unlikely
 - dissociation
 - polymerisation (formation of larger molecular complexes)





HISKP

5 Creation of the Signal

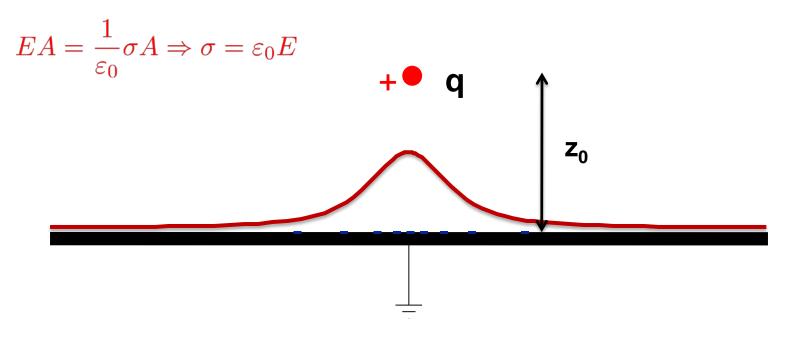
5.1 Signal Formation by Moving Charges
5.2 Ramo-Shockley Theorem
5.3 Planar Detector
5.4 Cylindrical Wire





Consider charge q above a grounded electrode

- electric field is perpendicular to conductor at the surface
- changes take place only on surface
- surface charge density σ and electric field E on the surface are related by Gauss' law





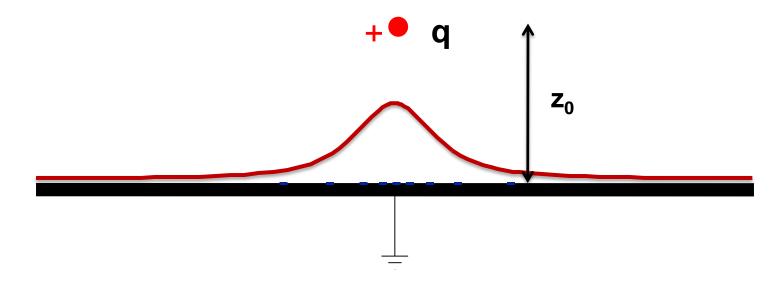


In order to find the charge induced on an electrode, we have to

a) solve the Poisson equation with boundary condition $\phi = 0$ on the conductor surface

$$\Delta \varphi = -\frac{\rho}{\varepsilon_0}, \quad \mathbf{E} = -\nabla \varphi$$

- b) calculate the electric field *E* on the surface of the conductor
- c) integrate $\varepsilon_0 E$ over the surface of the electrode

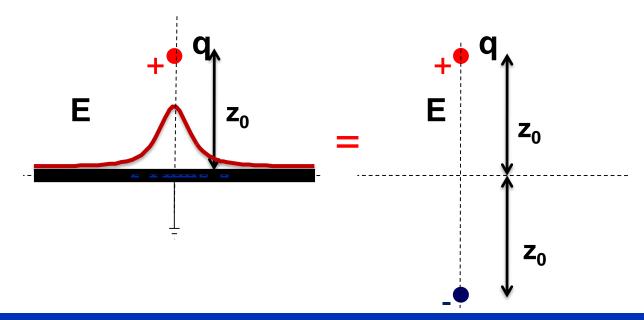




Signal Formation



For this particularly simple setup with one electrode ⇒ use mirror charge

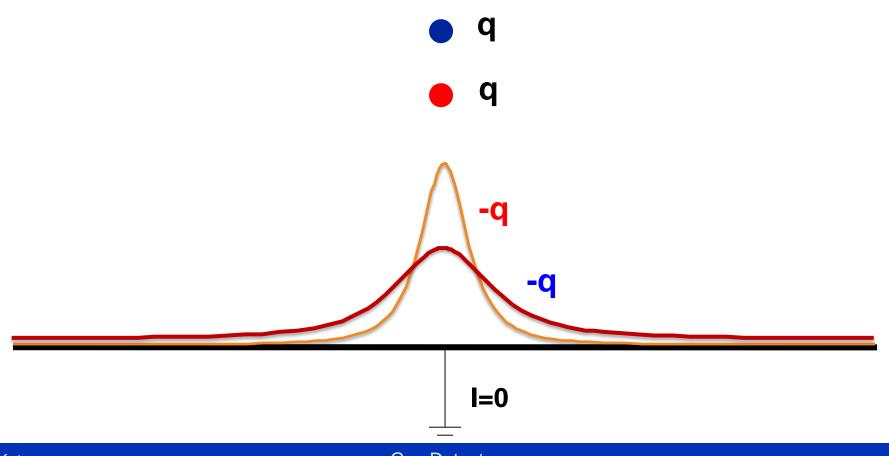






Movement of charge q (no external field needed!)

⇒ change of induced surface charge on electrode

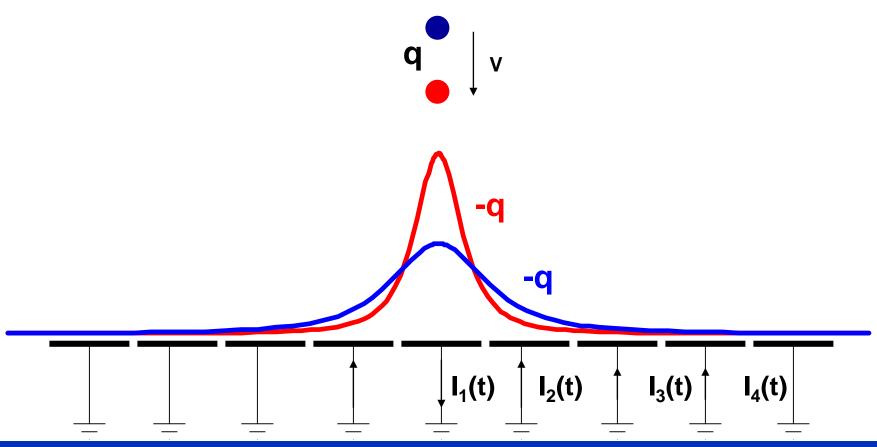






Movement of charge q (no external field needed!)

- ⇒ change of induced surface charge on electrode
- ⇒ current on segmented electrodes

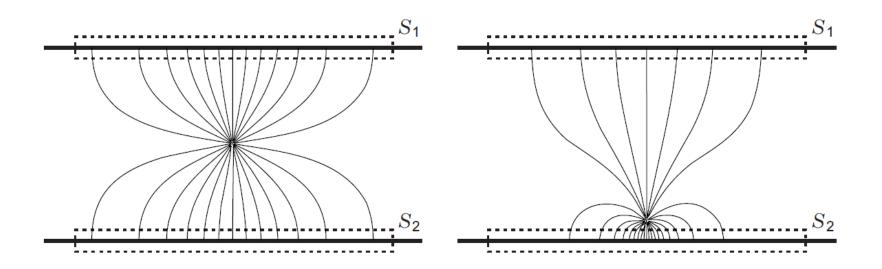






How to calculate the induced signal?

- solve Poisson equation with moving charge
- use mirror charges to get rid of electrodes
- use Ramo-Shockley theorem



[H. Spieler, Semiconductor detector systems, Oxford, 2005]



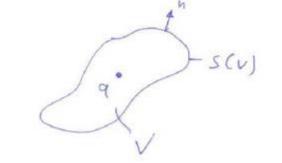
[W. Shockley, J. Appl. Phys. 9, 635 (1938), S. Ramo, Proc. IRE 27, 584 (1939)]

Calculation of signals induced on grounded electrodes:

Gauss' Law: point charge inside closed surface

$$\oint \mathbf{E} \cdot \mathbf{n} \mathrm{d}a = \frac{1}{\varepsilon_0} \int_V \rho(\mathbf{x}) \mathrm{d}^3 x$$

• Green's 2nd theorem:



$$\int_{V} (\phi \Delta \psi - \psi \Delta \phi) \, \mathrm{d}^{3}x = \oint_{S} \left(\phi \frac{\partial \psi}{\partial n} - \psi \frac{\partial \phi}{\partial n} \right) \, \mathrm{d}a$$

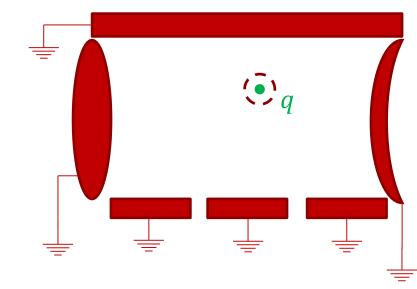
with $\frac{\partial \phi}{\partial n} \equiv \nabla \phi \cdot \mathbf{n} \left[= -\mathbf{E} \cdot \mathbf{n} \right]$





Consider detector volume delimited by grounded electrodes:

Real situation: charge q at position x₀, all electrodes grounded, i.e.
 U_i = 0, i = 1,2, ... ⇒ solution φ(x)





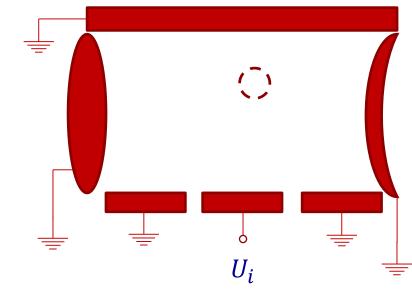


- Real situation: charge q at position x₀, all electrodes grounded, i.e.
 U_i = 0, i = 1,2, ... ⇒ solution φ(x)
- Auxiliary situations: charge q removed, all electrodes grounded except electrode i, i.e.

 $U_j = 0, j \neq i \Rightarrow$ solutions $\phi_i(x)$ with $\phi_i(x) = U_i$ at surface of electrode *i*

Space between electrodes and charge q free of charges

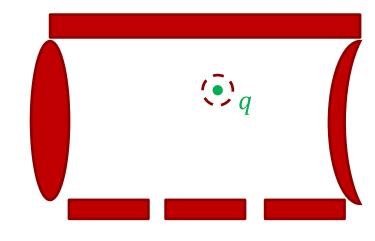
$$\Rightarrow \Delta \phi(\mathbf{x}) = 0$$
$$\Delta \phi_i(\mathbf{x}) = 0$$



universität**bonr**



Apply Green's 2^{nd} theorem to volume *V* delimited by S(V):



universität**bonn**

$$\int_{V} (\phi \Delta \phi_{i} - \phi_{i} \Delta \phi) d^{3}x = \oint_{S(V)} \left(\phi \frac{\partial \phi_{i}}{\partial n} - \phi_{i} \frac{\partial \phi}{\partial n} \right) da \qquad \text{for every } i$$
$$= 0$$

Solve surface integral by splitting it up into 3 parts





$$\Rightarrow Q_i = -q \frac{\phi_i(\mathbf{x}_0)}{U_i}$$

induced charge on electrode i by charge qat position x_0 when all electrodes are grounded

 $\phi_i(x_0) =$ potential at point x_0 when point charge q is removed, electrode i is put to potential U_i and all other electrodes are grounded

weighting potential of electrode *i*

Point charge moving along trajectory $x_0(t)$ \Rightarrow time-dependent induced charge on electrode *i*

$$\Rightarrow$$
 current $I_i(t) = -\frac{\mathrm{d}Q_i(t)}{\mathrm{d}t}$

Sign convention: positive current points away from electrode



10 (1)

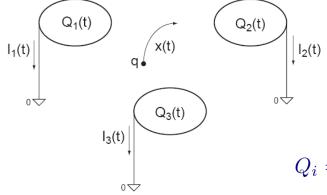
$$I_{i}(t) = -\frac{\mathrm{d}Q_{i}(t)}{\mathrm{d}t} = \frac{q}{U_{i}}\frac{\mathrm{d}}{\mathrm{d}t}\phi\left[\mathbf{x}_{0}(t)\right]$$
$$I_{i}(t) = \frac{q}{U_{i}}\nabla\phi_{i}\left[\mathbf{x}_{0}(t)\right] \cdot \frac{\mathrm{d}\mathbf{x}_{0}(t)}{\mathrm{d}t} = -\frac{q}{U_{i}}\mathbf{E}_{i}\left[\mathbf{x}_{0}(t)\right] \cdot \mathbf{v}(t)$$
$$\begin{array}{c} \mathsf{Ramo-Shockley}\\ \mathsf{Theorem}\end{array}$$

The current induced on a grounded electrode by a point charge qmoving along a trajectory $x_0(t)$ is $I_i(t)$, where $E_i(x_0)$ is the electric field in the case where the charge q is removed, electrode i is set to voltage U_i , and all other electrodes are grounded.

universität**bon**



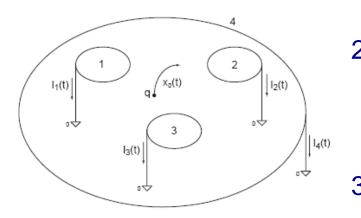




Consequences:

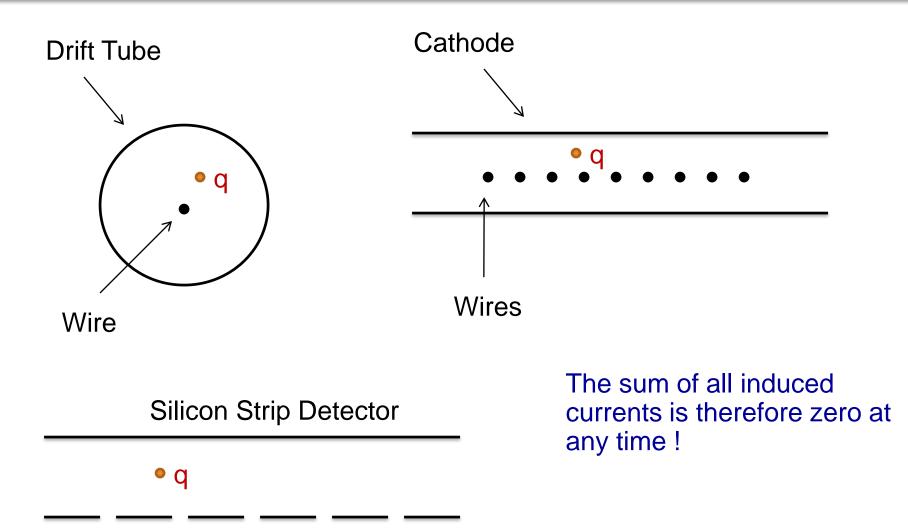
 Charge induced on electrode *i* by a charge *q* moving from point 1 to 2 is

$$Q_{i} = \int_{t_{1}}^{t_{2}} I_{i}(t) dt = -\frac{q}{U_{i}} \int_{t_{1}}^{t_{2}} \boldsymbol{E}_{i}[\boldsymbol{x}(t)] \, \dot{\boldsymbol{x}}(t) dt = \frac{q}{U_{i}} \left[\phi_{i}(\boldsymbol{x}_{1}) - \phi_{i}(\boldsymbol{x}_{2})\right]$$



and is independent of the actual path

- 2. Once all charges have arrived at the electrodes, the total induced charge in a given electrode is equal to the charge that has arrived at this electrode
- 3. In case there is one electrode enclosing all others, the sum of all induced currents is zero at any time



IskP

universität**bonn**





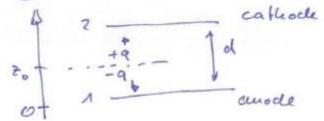


- Planar ionization chamber
- Planar strip detector
- Proportional wire counter



Planar Ionization Chamber

Weighting potential of anode => set anode to potential My and ground catheadle \$ (2=0) = U, $\phi_{1}(z) = \frac{U_{1}}{d}(d-z)$ $E_{1}(z) = U_{1}$ $Ramo: I_{A}(1) = -\frac{q}{u} \cdot E_{A}[2(1)] \cdot 2(1)$ 2 = u = NE $dt = \mu d_1$, $z(t=0) = z_0 = z_0 = z_{(t)} = \mu d_1 \cdot t + z_0$





Planar Ionization Chamber



- Jons and à contribute to signal! , u ion thue Three regions in time:
 - $t < t_e = \frac{z_o}{u_e}$ => $z_{iou}(t) = z_o + u_{iou} t$, $z_{iou} = u_{iou}$ $z_e(t) = z_o - u_e t$, $z_e = -u_e$

->
$$I_{1}(t) - I_{1}^{ion}(t) + I_{2}^{e}(t) = -\frac{q}{u_{1}} \cdot \frac{u_{1}}{d} \cdot u_{iou} - \frac{(-q)}{u_{1}} \cdot \frac{u_{1}}{d} \cdot (-u_{e})$$

= $-\frac{q}{d}(u_{ion} + u_{e})$

•
$$t_e < t < t_{ion} = \frac{q_{-2_o}}{u_{ion}} \Rightarrow 2_{ion}(t) = 2_o + u_{ion} t_{12_{ion}} = u_{ion}$$

 $Z_e(t) = 0, \quad Z_e = 0$

$$=$$
 $1_{n}(t) = -d$ uion



universität**bonn**

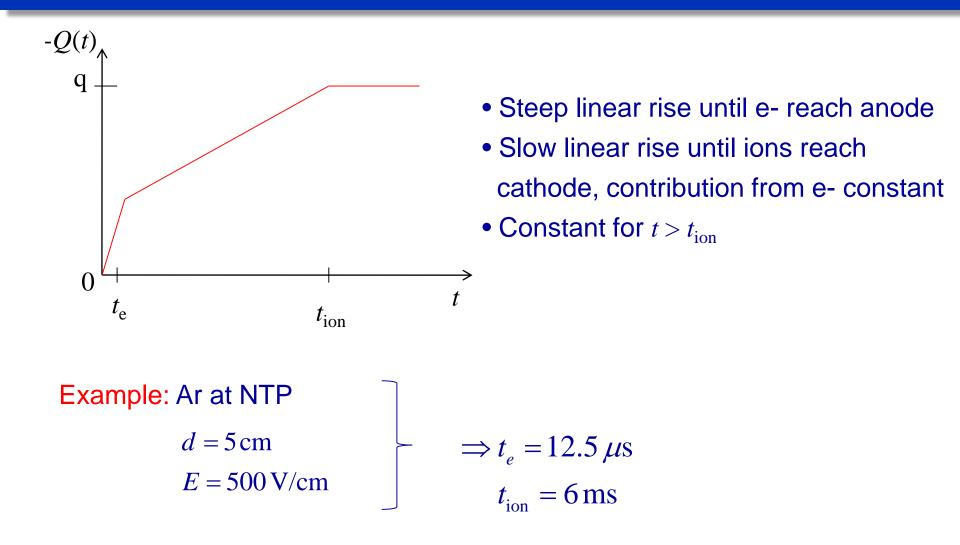
Charge induced on anode :

- Q, (+) SI, (+') d+'
- $t < t_e = 7 \quad Q_1(t) = -\frac{q}{a} (u_{iou} + u_e) \cdot t$
 - · texterion => Q (1) = q (uint + 20)
 - · t > tion => Q, (1) = q (d +0) = q total induced cheve so muscle

HISKP

Planar Ionization Chamber

universität**bonn**





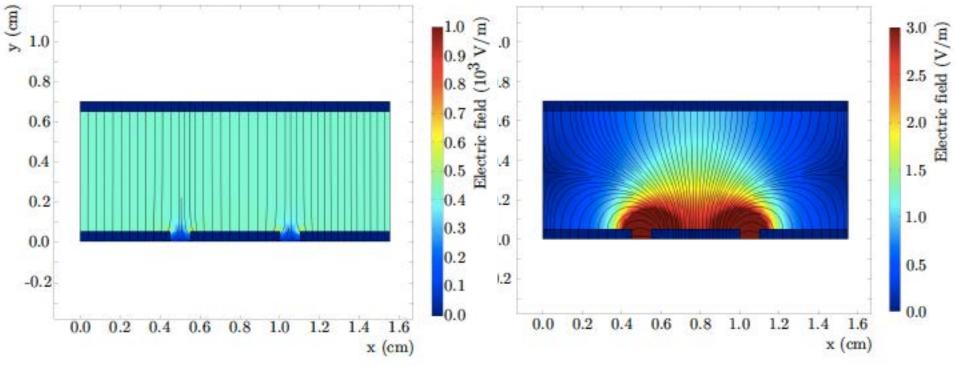


How do the real field and the weighting field look like?



ISKP





M. Berger, PhD, Development Commissioning and Spatial Resolution Studies of a GEM based TPC, Munich, 2015]

Gal beters

Weighting field

Planar Strip Detector

Real field







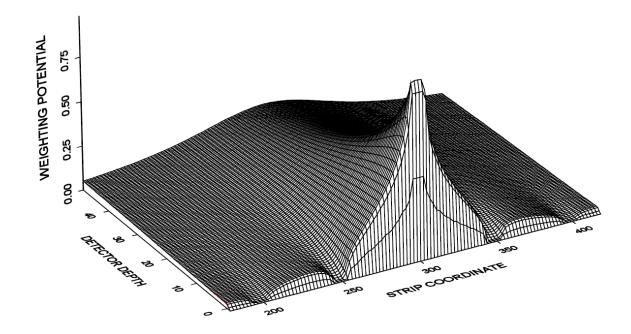


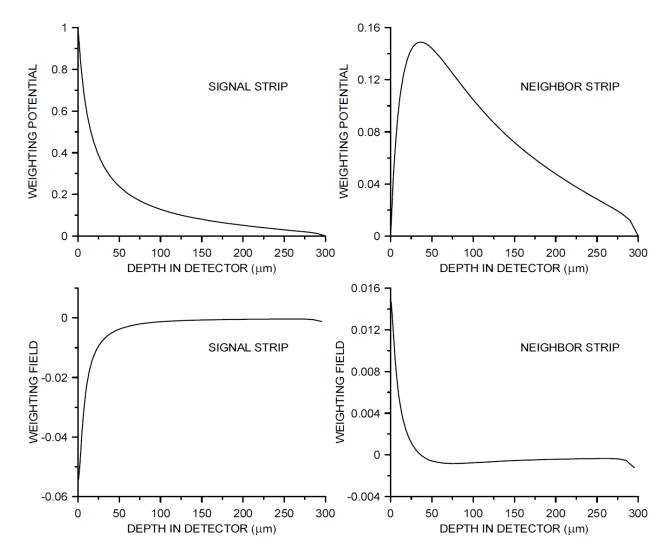
FIG. 2.29. Weighting potential for a 300 μ m thick strip detector with strips on a pitch of 50 μ m. The central strip is at unit potential and the others at zero. Only 50 μ m of depth are shown.

[H. Spieler, Semiconductor detector systems, Oxford, 2005]

universität**bonn**

Planar Strip Detector





[H. Spieler, Semiconductor detector systems, Oxford, 2005]

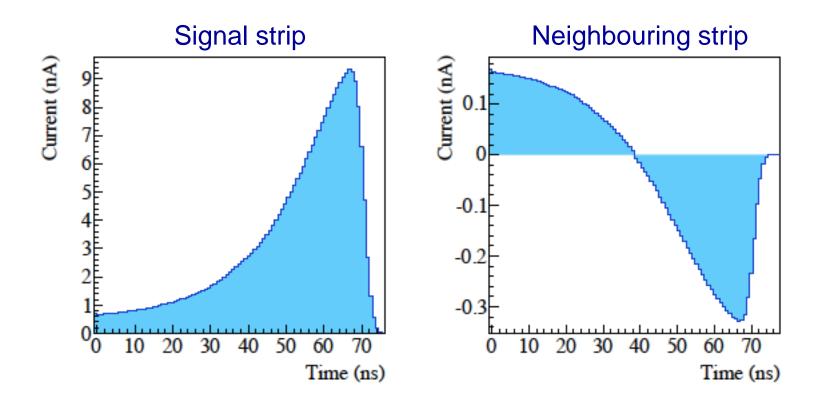
ISKP

Gas Detectors





Currents according to Ramo-Shockley theorem



M. Berger, PhD, Development Commissioning and Spatial Resolution Studies of a GEM based TPC, Munich, 2015]





Analogously to plane ionization chamber: drifting charges induce currents

$$\phi(r) = \frac{\mathcal{C}U}{2\pi\varepsilon_0} \ln\left(\frac{b}{r}\right) \quad \text{for} \quad a < r < b, \ \mathcal{C} = \frac{2\pi\varepsilon_0}{\ln\left(\frac{b}{a}\right)}$$
$$\phi(b) = 0$$

Ramo's theorem:

- weighting potential of wire \Rightarrow set to potential U_1 , ground tube
- here as for parallel plate counter: weighting field ist equal to real field!

$$\phi_1(r) = \frac{U_1 \ln\left(\frac{b}{r}\right)}{\ln\left(\frac{b}{a}\right)} \qquad \qquad E_1(r) = \frac{U_1}{r \ln\left(\frac{b}{a}\right)}$$

Signal Shape of Cylindrical Wire

Contribution of e- and ions: unlfiplication at i'

• electrons:
$$I_{\mu}^{e}(t) = -\frac{(-q)}{U_{\mu}} \cdot \frac{U_{\mu}}{rl_{\mu}(b_{\mu})} \cdot \dot{r}_{e}(t)$$

$$Q_{A}^{e} = \int_{a}^{a} I_{A}^{e}(t) dt = \int_{a+t'}^{a} \frac{q}{r l_{A}(b/a)} dr$$

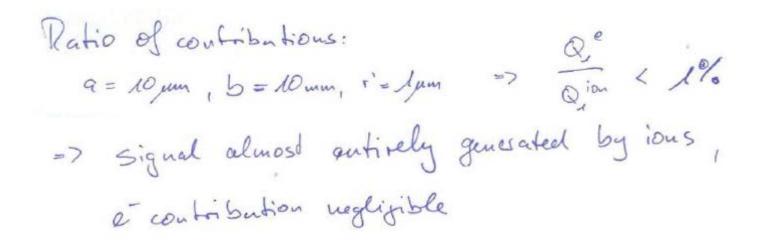
$$= -\frac{q}{\ln(b/a)} \ln\left(\frac{a tr'}{a}\right)$$

· ious :

$$\begin{split} I_{\mu}^{ion}(4) &= -\frac{q}{u_{\mu}} \cdot \frac{u_{\mu}}{\tau lu(b/a)} \cdot \overline{r_{ion}(4)} \\ Q_{\mu}^{ion} &= \int I_{\mu}^{ion}(4) d4 = -\frac{q}{lu(b/a)} lu(\frac{b}{a+r'}) \\ q_{\mu}(r) \end{split}$$

HISKP







Jou trajectory:
$$u = \mu E$$

$$\frac{dr(4)}{dt} = \mu \frac{u}{\tau(4) \ell_{n}(b'_{n})} \implies \tau(4) = \alpha \sqrt{1 + \frac{t}{t_{o}}}, t_{o} = \frac{\alpha^{2} \ell_{u}(b'_{n})}{Z_{\mu}u}$$
with $\tau(0) = \alpha$

$$\tau(4) = \frac{1}{2} \alpha \left(1 + \frac{t}{t_{o}}\right)^{-1/2} \cdot \frac{1}{t_{o}}$$

$$= \sum T_{\mu}(4) = -\frac{\alpha}{u_{\mu}} \cdot \frac{u_{\mu}}{\tau \ell_{u}(b'_{n})} \cdot \tau =$$

$$= -\frac{\alpha}{2\ell_{u}(b'_{n})} \cdot \frac{1}{t_{v}t_{o}}$$

$$= \sum uegative wite Signal$$

B. Keizer perbolic form with characteristic time constant to a ferous

65

Signal Shape of Cylindrical Wire

Juduced charge at time t:

$$Q_{p}(t) = \int_{0}^{t} I_{p}(t') dt' = -\frac{q}{2ln(b_{a})} \cdot ln\left(1 + \frac{t}{t_{o}}\right)$$

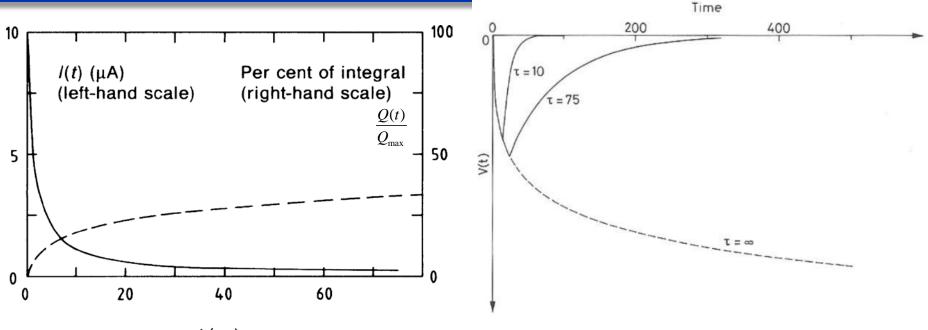
Once ions have arrived at tube wall (at $t = t_{max}$):

 $\Rightarrow Q_1(t_{max}) = -q$

Current induced on cathode: $I_2(t) = -I_1(t)$

Signal Shape





t (ns)

 $t_0 = 1.25 \text{ ns}$ b/a = 500 q = 10⁶

[W. Blum et al., Particle Detection with Drift Chambers, Springer (2008)]

Q(t), U(t) with RC element

[W.R. Leo, Techniques for Nuclear and Particle Physics Experiments, Springer (1994)]

IskP





Many textbooks deduce the signal shape using energy conservation

• Signal is calculated using energy balance:

Energy gained by charge in electric field =

change of energy stored in capacitor

- In some special cases, this argument gives the correct result, e.g. for a 2-electrode system because there the weighting field and the real field are equal.
- But the argument is very misleading:
 - An induced current signal has nothing to do with energy. In a gas detector the electrons are moving at constant speed in a constant electric field, so the energy gained by the electron in the electric field is lost into collisions with the gas, i.e. heating of the gas.
 - In absence of an electric field, the charge can be moved across the gap without using any force and currents are flowing.



- 1. Calculate particle trajectory $x_0(t)$ in the "real" electric field
- 2. Remove all impedance elements, ground the electrodes and calculate currents induced by moving charge on grounded electrodes

$$I_i(t) = \frac{q}{U_i} \nabla \phi_i \left[\boldsymbol{x}_0(t) \right] \cdot \frac{\mathrm{d} \boldsymbol{x}_0(t)}{\mathrm{d} t} = -\frac{q}{U_i} \boldsymbol{E}_i \left[\boldsymbol{x}_0(t) \right] \cdot \boldsymbol{v}(t)$$

3. Place these currents as ideal current sources on a circuit where the electrodes are simple nodes and the mutual electrode capacitances are added between the nodes. They are calculated from the weighting field by

$$c_{nm} = \frac{\varepsilon_0}{V_w} \oint_{\boldsymbol{A}_n} \boldsymbol{E}_m(\boldsymbol{x}) d\boldsymbol{A}$$

$$C_{nn} = \sum_{m} c_{nm} \qquad C_{nm} = -c_{nm} \quad n \neq m$$

