

# General Physics: Electricity & Magnetism

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Faraday cage

# Chapter 2: Conductor & dielectrics

1. Conductors
2. Capacitance & capacitors
3. Dielectrics
4. Energy of electrostatic fields

# Chapter 2: Homework

1. 9-2, 9-3, 9-5, 9-11, 9-14
2. Others in the ppt

# 2.1 Conductors

**Equilibrium:** stationary state; time-independent

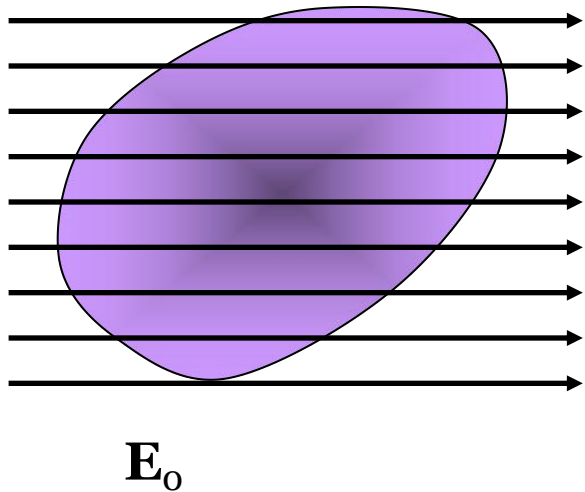


Mountain:  
stationary

Water:  
stationary?

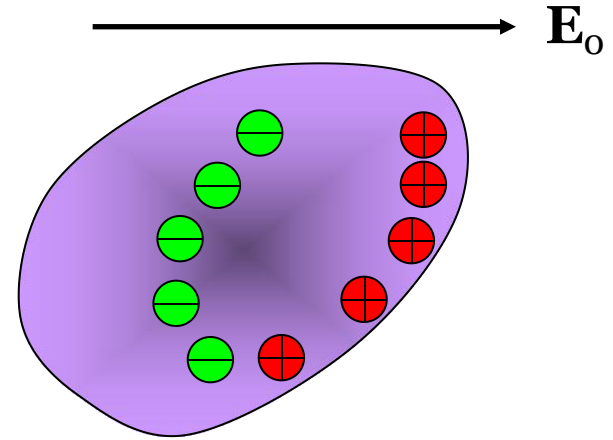
Current:  
stationary?

# 2.1 Conductors



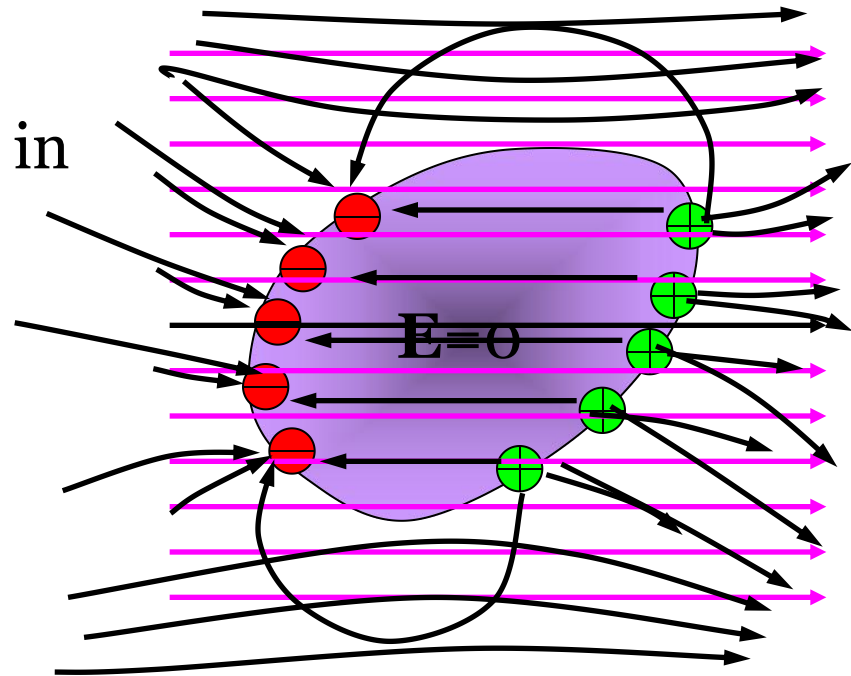
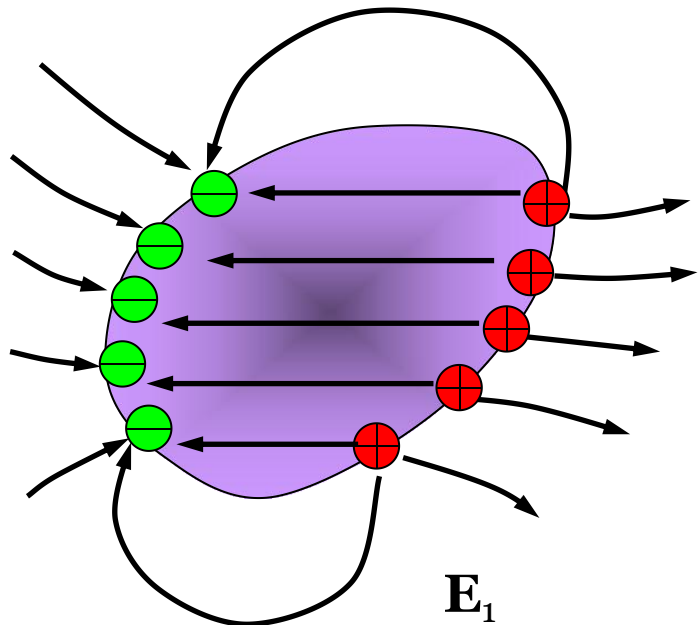
## Conclusion 1:

In an ideal conductor, the electrostatic field



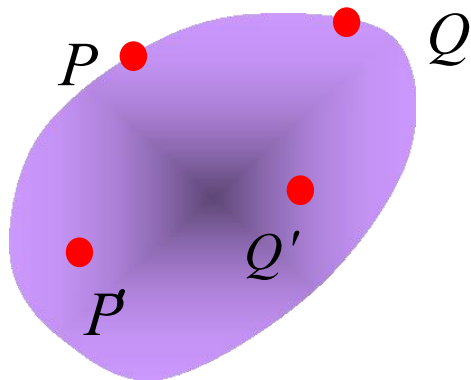
$$\mathbf{E} = \mathbf{E}_0 + \mathbf{E}_1 = 0 \text{ in}$$

the equilibrium condition



# 2.1 Conductors

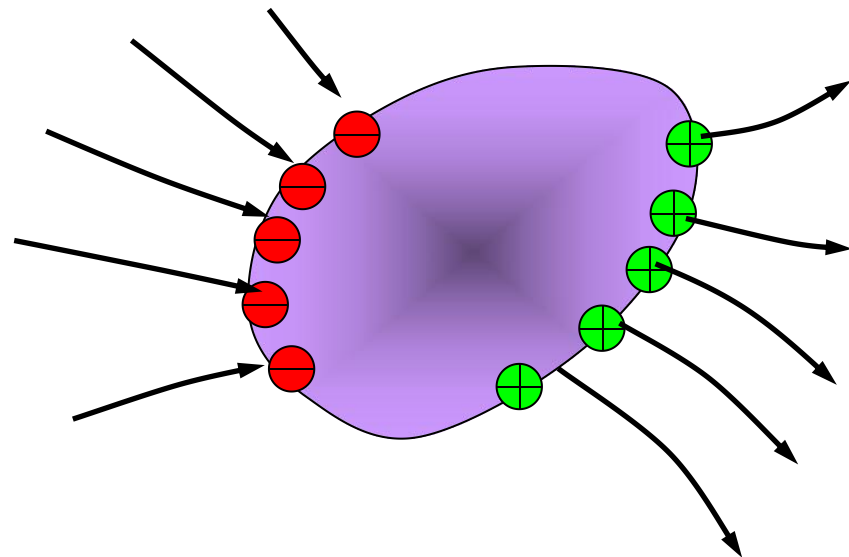
**Conclusion 2:** a conductor is an equipotential (isopotential) body



$$\Delta U_{PQ} = \int_{PQ} \mathbf{E} \cdot d\mathbf{l} = 0$$

**Conclusion 3:**  $\mathbf{E} \parallel \mathbf{n}$  everywhere

**Note:** the pre-condition for electrostatic screening is that there are enough mobile carriers. Is it always satisfied?



# 2.1 Conductors

**Example:** to estimate the electrostatic screening length.

Surface density  $\sigma=1 e/a^2$   $a=4$  Angstrom

$$E=\sigma/\varepsilon_0=1.6 \times 10^{-19}/(4 \times 10^{-10})^2/8.85 \times 10^{-12}=1.1 \times 10^{11} \text{ N/C}$$

**case 1:** Typical metal  $\rho=10^{22} e/\text{cm}^3$

$$d=\sigma/\rho=1/(4 \times 10^{-10})^2/[10^{22} / (10^{-2})^3] =6.25 \times 10^{-10} \text{ m}$$

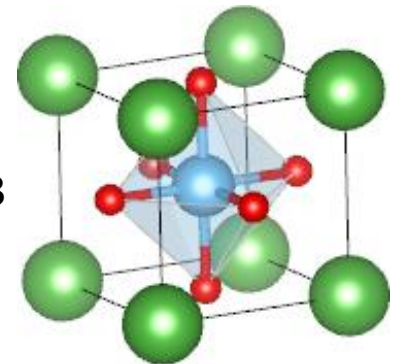
$$=6.25 \text{ \AA}=1.5625 a$$

**case 2:** Intrinsic semiconductor Si  $\rho=1.5 \times 10^{10} e/\text{cm}^3$

$$d=417 \text{ m}$$

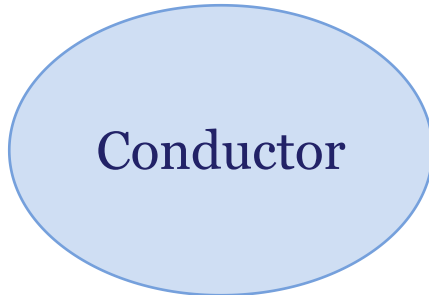
**case 3:** Doped semiconductor Si  $\rho=10^{18} e/\text{cm}^3$

$$d=6.25 \text{ \mu m}=15625 a$$



# 2.1 Conductors

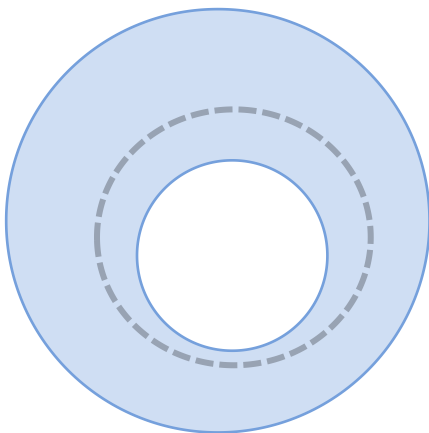
**Conclusion 4:** no net charge in a conductor



$$\Phi_e = \oint_S \mathbf{E} \cdot d\mathbf{S} = \frac{1}{\epsilon_0} \sum_{i=1}^n q_i^{in}$$

$$\nabla \cdot \mathbf{E} = \rho / \epsilon_0 \quad \longrightarrow \quad \rho = 0 \text{ everywhere}$$

Net charge can only appear at the surface of a conductor



How about a conductor with a cavity?

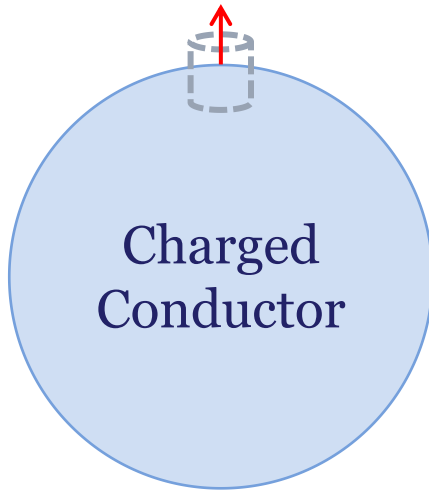
net charge in the cavity = net charge at the inner surface

Applications: 1. [Faraday cylinder](#)

2. [Van de Graaff generator](#)



# 2.1 Conductors



Surface charge density  $\sigma(\mathbf{r})$

$\mathbf{E}$  is perpendicular to the surface

$$\Phi = \int_{dS} \mathbf{E} \cdot d\mathbf{S} + \int_{S-dS} \mathbf{E} \cdot d\mathbf{S}$$

$$= \int_{dS} \mathbf{E} \cdot d\mathbf{S} + 0$$

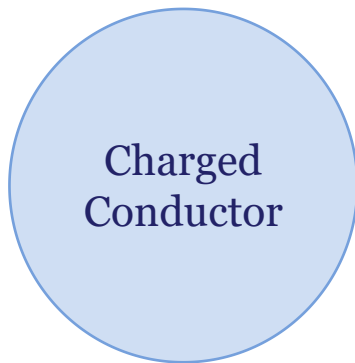
$$= E dS = dq / \epsilon_0 = \sigma dS / \epsilon_0$$

$$\mathbf{E} = \sigma(\mathbf{r}) / \epsilon_0$$

For comparison,  $\mathbf{E} = \sigma(\mathbf{r}) / 2\epsilon_0$  for infinite plate because there are two surfaces.

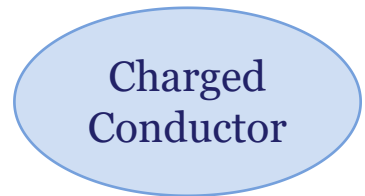
# 2.1 Conductors

For an isolated conductor, the surface charge density may be complex in general, but qualitatively it is in proportional to the local curvature



Sphere: uniform curvature, uniform  $\sigma$

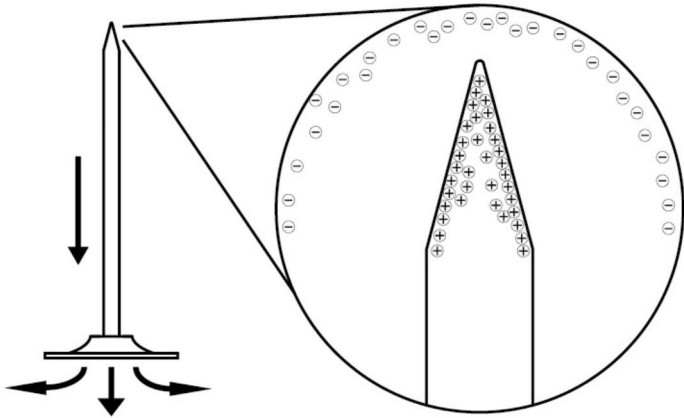
Ellipsoid: larger  $\sigma$  at two ends



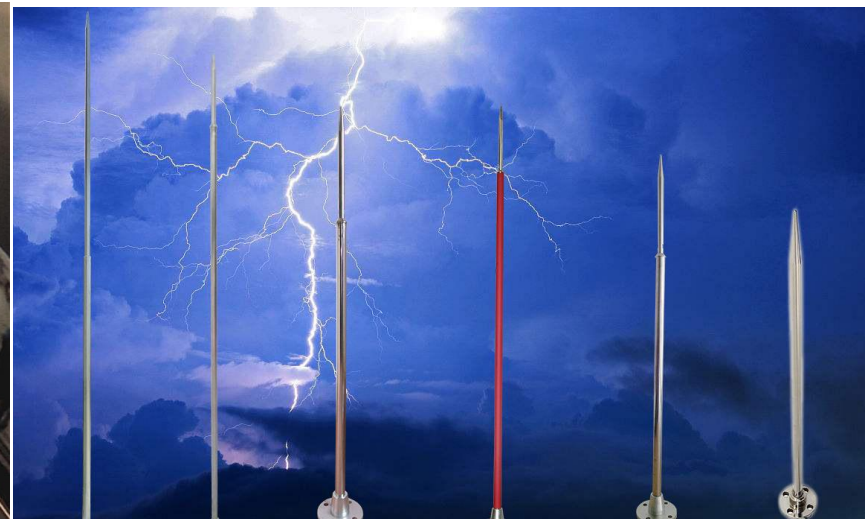
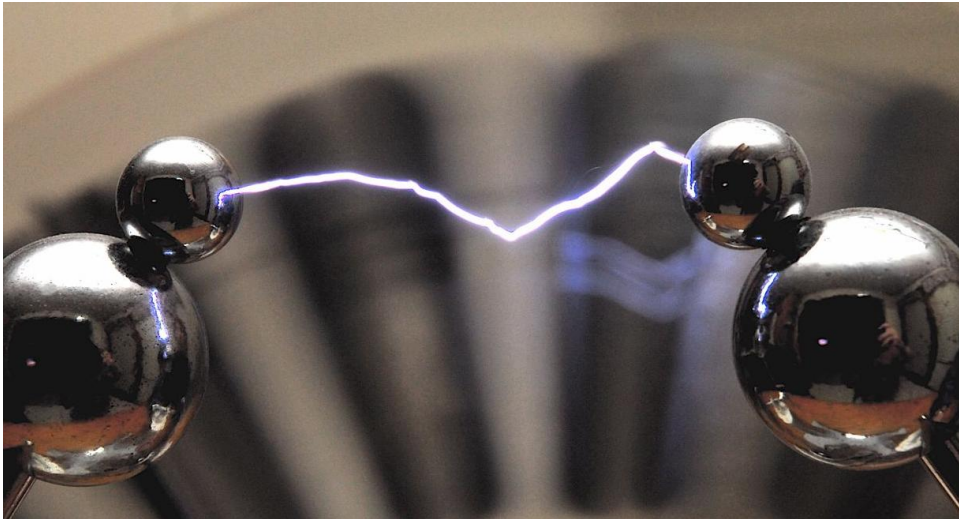
An intuitive understanding (qualitative): With the same distance and the same solid angle, the corresponding surface is large in the side with smaller curvature

Then, how about the negative curvature case?

# 2.1 Conductors

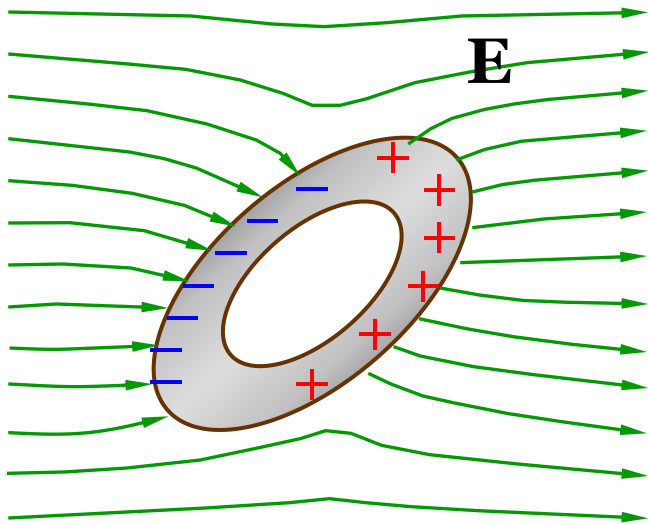


When the local electric field near the tip goes beyond a threshold (breakdown field), the corona discharge occurs.

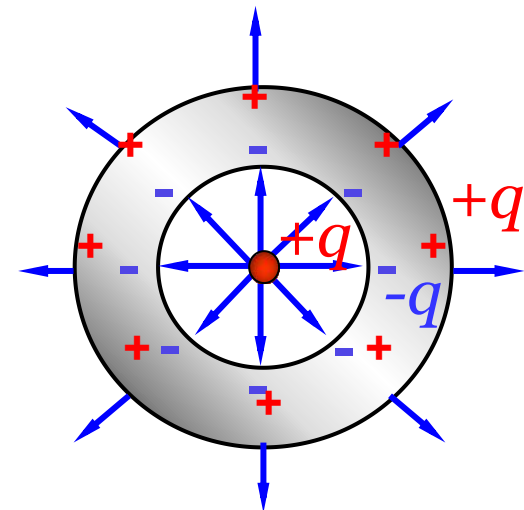


# 2.1 Conductors

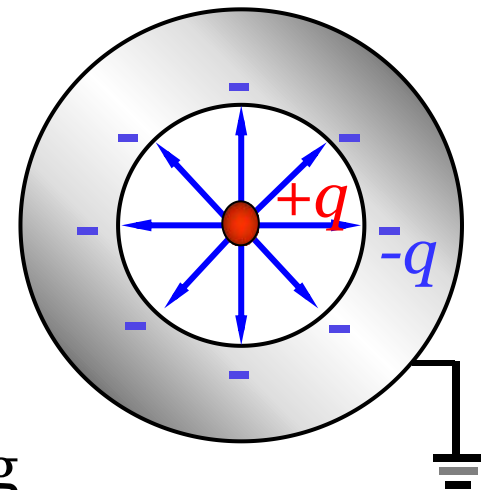
Electrostatic shielding



Outside field ---->  
zero inside



Inner charge ---->  
outside field



Grounding

# 2.1 Conductors

**Example:** an isolated hollow spherical shell ( $R_1$ : outside radius;  $R_2$ : inside radius) + a ball ( $R_3$ : radius), both of which are with charge  $q$ .

Spherical symmetric:  $E(r)$ ,  $U(r)$

1.  $r < R_3$ :  $E=0$  (conductor)

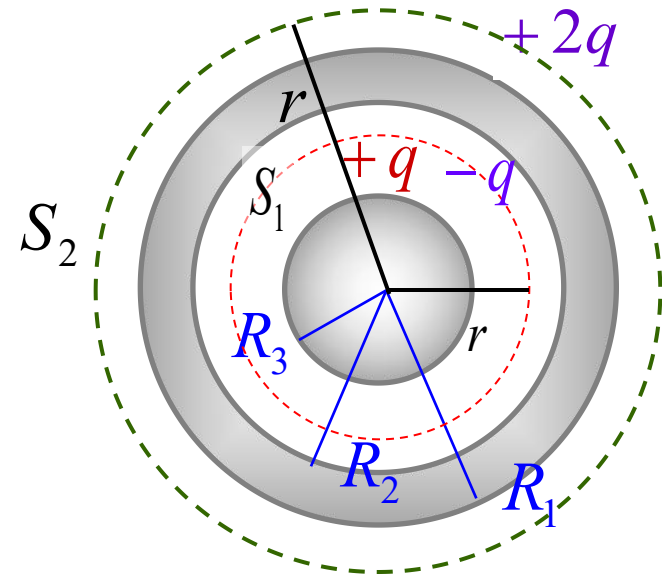
$U = \text{a constant} = q/(4\pi\epsilon_0 R_3)$  ??

2.  $R_3 < r < R_2$   $E(r) = q/(4\pi\epsilon_0 r^2)$

$U(r) = q/(4\pi\epsilon_0 r) + C$

$U(R_3) = q/(4\pi\epsilon_0 R_3) + C = U_3 \rightarrow C = U_3 - q/(4\pi\epsilon_0 R_3)$

$U(R_2) = q/(4\pi\epsilon_0)(1/R_2 - 1/R_3) + U_3$



# 2.1 Conductors

$$U(R_2) = q / (4\pi\epsilon_0) (1/R_2 - 1/R_3) + U_3$$

3.  $R_2 < r < R_1$      $E(r) = 0$  (conductor)

$$U(R_1) = U(R_2)$$

4.  $r > R_1$

$$E(r) = 2q / (4\pi\epsilon_0 r^2)$$

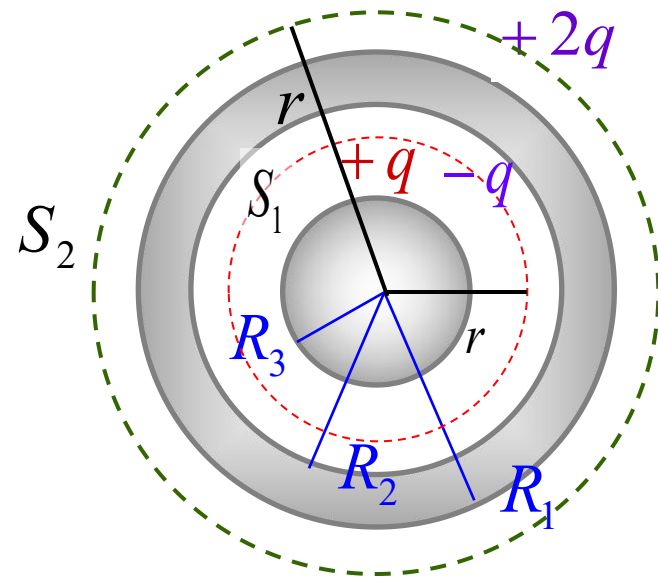
$$U(r) = 2q / (4\pi\epsilon_0 r)$$

$$U(R_1) = 2q / (4\pi\epsilon_0 R_1) = U(R_2)$$

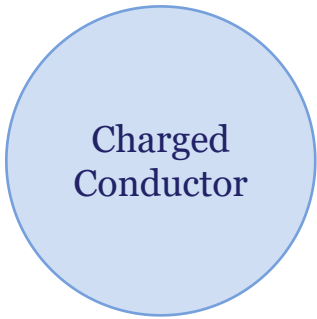
$$2q / (4\pi\epsilon_0 R_1) = q / (4\pi\epsilon_0) (1/R_2 - 1/R_3) + U_3$$

Then  $U_3 = q / (4\pi\epsilon_0) (2/R_1 - 1/R_3 + 1/R_2)$

$$\neq q / (4\pi\epsilon_0 R_3)$$



# 2.2 Capacitance & capacitors



$U \sim Q$  Linear superposition principle

e.g.  $U=Q/(4\pi\epsilon_0 R)$

Capacitance  $C=Q/U=4\pi\epsilon_0 R$



$C$ : the charge needed to increase  
voltage

unit: F     $1 \text{ F} = 1 \text{ C/V}$

For isolated spherical conductor with 1 F

$R=1/4\pi\epsilon_0=9 \times 10^9 \text{ m} = 1400 R_{\text{earth}}$

Thus 1 F is too large in practice

Then 1  $\mu\text{F}$ , 1 nF, 1 pF are more frequently used



Michael Faraday  
1791-1867

# 2.2 Capacitance & capacitors

Two conductors with electrostatic shielding:

Then the electric field is only from the inner charge, independent on external field.

**Case 1:** a plane-parallel capacitor

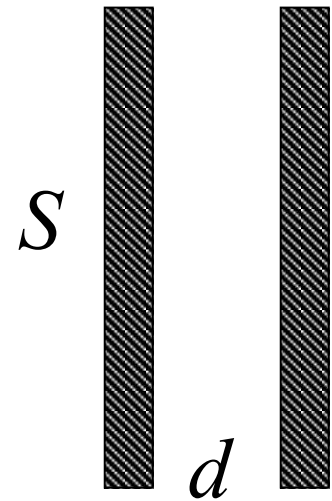
$$E = \sigma / \epsilon_0 \quad V = Ed$$

$$C = Q/V = \sigma S / (\sigma d / \epsilon_0) = \epsilon_0 S / d$$

$$S = 1 \text{ m}^2 \quad d = 1 \text{ mm}$$

$$C = 8.85 \times 10^{-12} \times 1 / 0.001 = 8.85 \times 10^{-9} \text{ F}$$

$$= 8.85 \text{ nF}$$





# 2.2 Capacitance & capacitors

**Case 2:** A coaxial cylindrical capacitor

$$E = \eta / (2\pi\epsilon_0 r)$$

$\eta$ : the line density of inner cylinder

$$\Delta U = \int_{R_1}^{R_2} E dr = \eta / (2\pi\epsilon_0) (\ln R_2 - \ln R_1)$$

$$= \eta \ln(R_2/R_1) / (2\pi\epsilon_0) \quad Q = \eta h$$

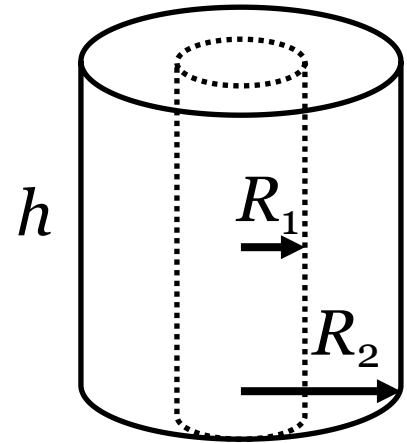
$$C = Q / \Delta U$$

$$= 2\pi\epsilon_0 h / \ln(R_2/R_1)$$

e.g.  $h = 1 \text{ m}$   $R_1 \sim 0.5 R_2$

$$C = 2\pi \times 8.85 \times 10^{-12} / \ln(2)$$

$$= 5.3 \times 10^{-10} \text{ F} = 0.53 \text{ nF}$$



In the limit of  $R_1 \sim R_2$

$$\ln(R_2/R_1) = R_2/R_1 - 1$$

$$C = 2\pi\epsilon_0 h R_1 / (R_2 - R_1)$$

e.g.  $R_1 / (R_2 - R_1) = 1000$

$$C = 55.6 \text{ nF}$$

# 2.2 Capacitance & capacitors

**Case 3:** A homocentric spherical shells

$$E=Q/(4\pi\epsilon_0 r^2)$$

$$\Delta U=\int_{R_1}^{R_2} E dr=Q/(4\pi\epsilon_0) (1/R_1-1/R_2)$$

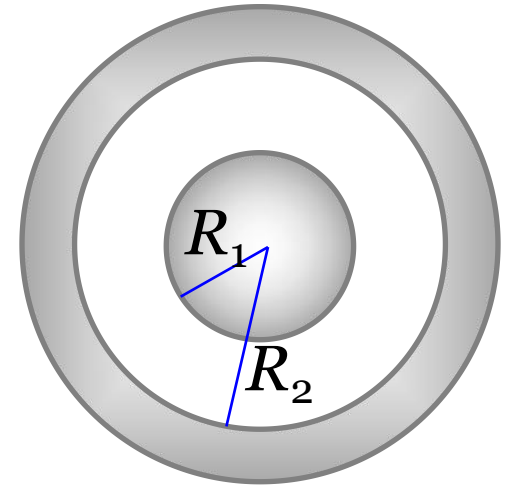
$$C=Q/\Delta U$$

$$=4\pi\epsilon_0 R_1 R_2 / (R_2 - R_1)$$

e.g.  $R_1=1$  m  $R_2\sim 1.1$  m

$$C=4\pi \times 8.85 \times 10^{-12} \times 1.1/0.1$$

$$=1.2 \times 10^{-9} \text{ C}=1.2 \text{ nF}$$



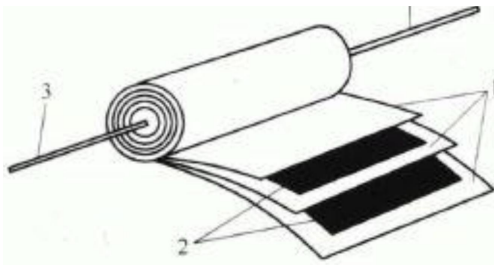
In summary, the general routes to enlarge  $C$ :

1. enlarge the charged surface
2. reduce the distance between charged surfaces

# 2.2 Capacitance & capacitors

## Examples of different kind of capacitors

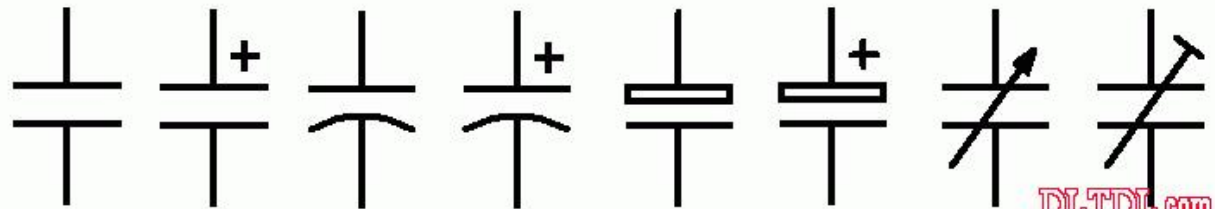
Paper capacitor



Aluminum electrolytic capacitor



Ceramic capacitor



# 2.2 Capacitance & capacitors

- Series connection

$$Q_1=Q_2=Q$$

$$U=U_1+U_2$$

$$C=Q/U=Q/(U_1+U_2)$$

$$1/C=1/C_1+1/C_2 \quad C \text{ is smaller}$$

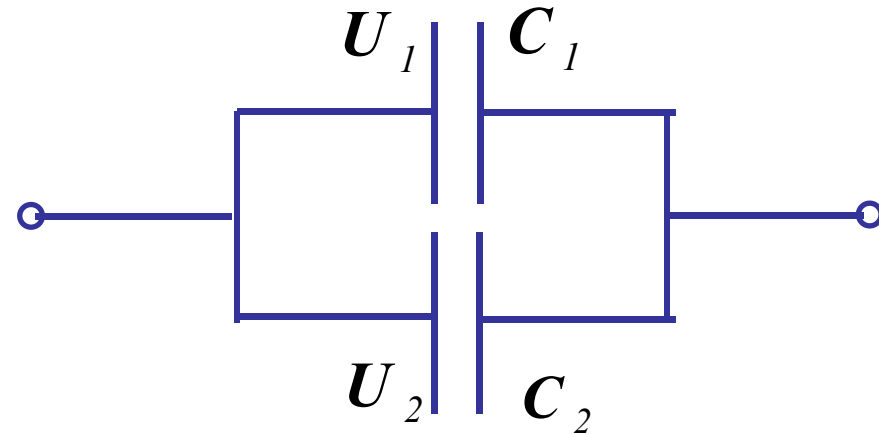
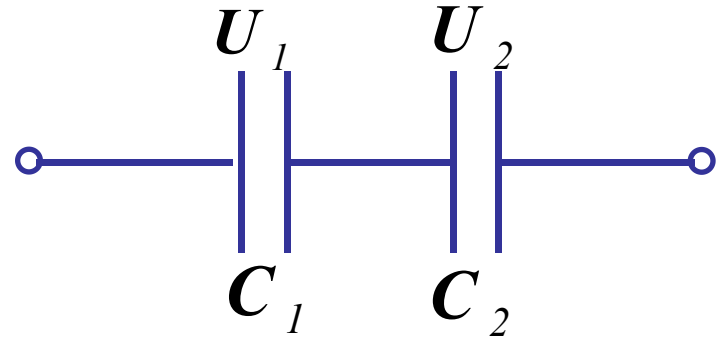
- Parallel connection

$$Q=Q_1+Q_2$$

$$U=U_1=U_2$$

$$C=Q/U=(Q_1+Q_2)/U$$

$$C=C_1+C_2 \quad C \text{ is larger}$$



**Differnt from resistors!**

# 2.2 Capacitance & capacitors

The charging process:

$$dW_e = dq(U_+ - U_-) = U dq = q/C dq$$

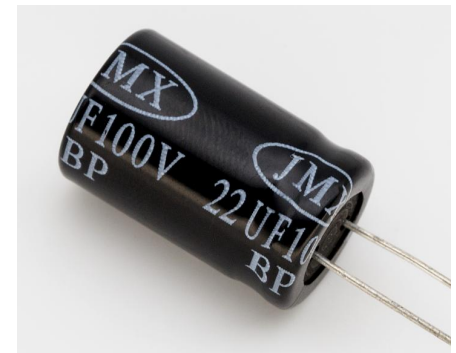
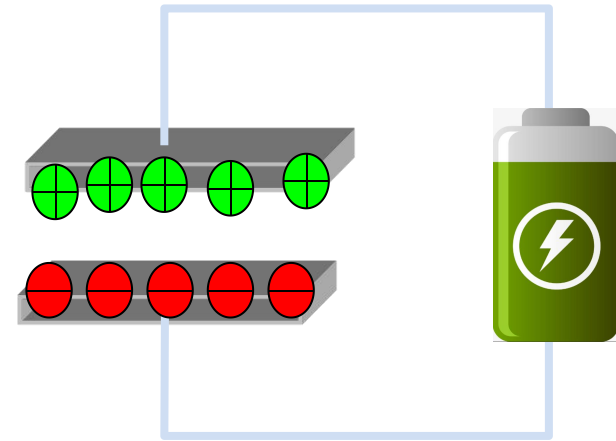
$$W_e = \int dW_e = \int q/C dq$$

$$= Q^2/2C = CU^2/2 = QU/2$$

**Example:** how much energy can a paper capacitor store?

$$W_e = CU^2/2 = 22 \times 10^{-6} \times 100^2/2$$

$$= 0.11 \text{ J}$$



100 V, 22  $\mu$ F

# 2.2 Capacitance & capacitors



AA chargeable battery 600 mAh 1.2 V

$$W_e = UQ = 1.2 \times 0.6 \times 3600 \\ = 2592 \text{ J}$$



AA alkaline battery 2500 mAh 1.5 V

$$W_e = UQ = 1.5 \times 2.5 \times 3600 \\ = 1.35 \times 10^4 \text{ J}$$



Supercapacitor: 3000 F, 3 V

$$W_e = CU^2/2 = 3 \times 10^3 \times 3^2/2 \\ = 1.35 \times 10^4 \text{ J} = 3.75 \text{ Wh}$$

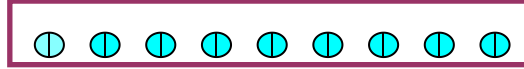
# 2.3 Dielectrics



Vacuum  
 $E = E_0$



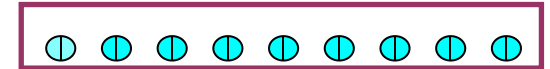
no charge



Dielectrics  
 $E = E_0 + E'$



localized charges  
(dipoles)



Conductors  
 $E = E_0 - E_0 = 0$



mobile charges  
(carriers)



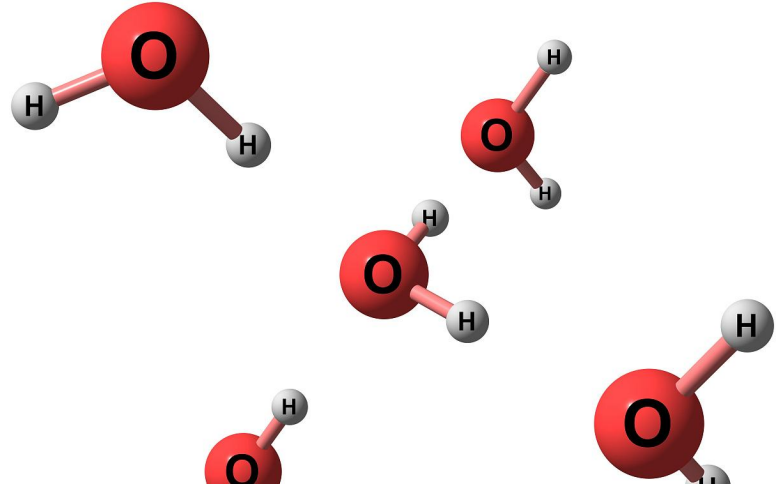
## 2.3 Dielectrics

### a) Polar molecules

$\mathbf{p}_m \neq 0$  for each molecule

$$\mathbf{P} = \sum \mathbf{p}_m = 0 \quad (\mathbf{E} = 0)$$

$$\mathbf{P} = \sum \mathbf{p}_m \neq 0 \quad (\mathbf{E} \neq 0)$$



Electrical poling:

dipoles are aligned by electric field ---->

macroscopic electric polarization

$$\mathbf{P} \sim \mathbf{E}$$



## 2.3 Dielectrics

### b) Nonpolar molecules

$\mathbf{p}_m = 0$  for each molecule

$$\mathbf{P} = \sum \mathbf{p}_m = 0 \quad (\mathbf{E} = 0)$$

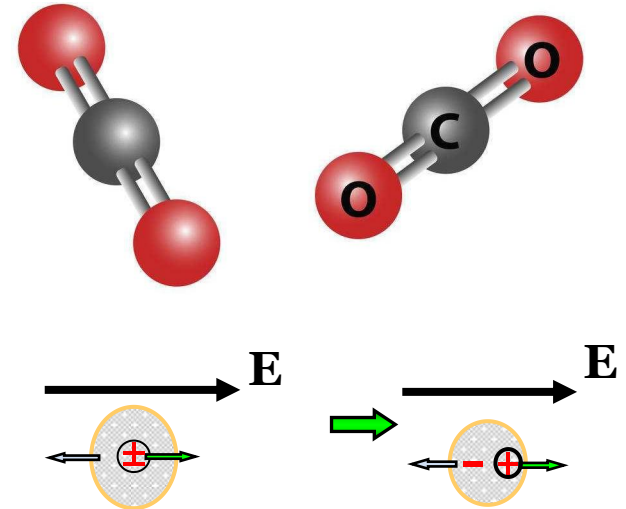
$$\mathbf{P} = \sum \mathbf{p}_m = 0 \quad (\mathbf{E} \neq 0)?$$

$$\mathbf{P} = \sum \mathbf{p}_m \neq 0 \quad (\mathbf{E} \neq 0)$$

Electrical poling:

to create dipoles by electric field ---->  
macroscopic electric polarization

$$\mathbf{P} \sim \mathbf{E}$$



Which  $\mathbf{P}$  is larger for these two cases?

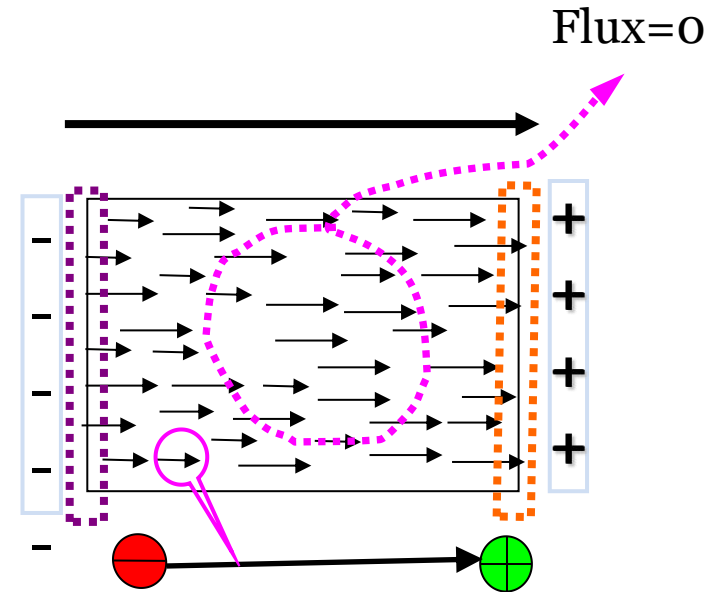
Why?

# 2.3 Dielectrics

Dielectrics under  $E$ -field

For a uniform dielectric medium,

- no net charge within the medium (macroscopically)
- net charges exist at the surface/interface (edge state)



Note: for a non-uniform dielectric medium,

- net charge can exist within the medium (macroscopically)

$$\nabla \cdot \mathbf{E} = \rho_e / \epsilon_0$$

## 2.3 Dielectrics

Electric polarization  $\mathbf{P} = \sum_i \mathbf{p}_i / V$

unit:  $\text{C}/\text{m}^2 \sim$  surface charge density  $\sigma$

$\rho$ : volume density of dipole molecules

$q$ : dipole charge

$l$ : dipole distance

surface charge  $Q^+ = \rho q S l$

$\sigma' = Q^+ / S = \rho q l = \rho p = P$

$\sigma' = \mathbf{P} \cdot \mathbf{e}_n$

not free charge, but bound charge



# 2.3 Dielectrics

Depolarizing field in the dielectric region

$$E_0 = \sigma_0 / \epsilon_0$$

$$E' = \sigma' / \epsilon_0 = P / \epsilon_0$$

$$E = E_0 - E' = E_0 - P / \epsilon_0 = (\sigma_0 - \sigma') / \epsilon_0$$

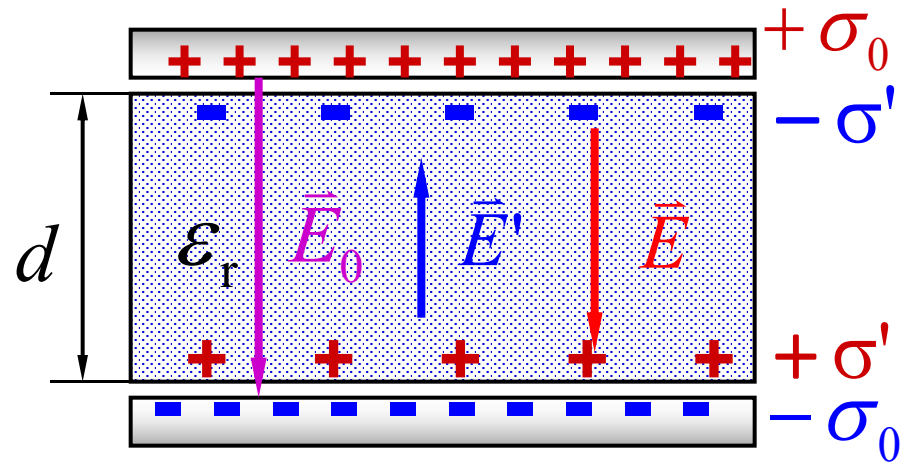
In **linear** dielectric medium

$$P = \chi_e \epsilon_0 E$$

$\chi_e$ : electric susceptibility

$$E = E_0 - E' = E_0 - P / \epsilon_0 = E_0 - \chi_e E$$

$$E = E_0 / (1 + \chi_e)$$

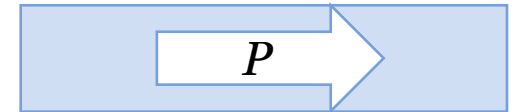


- $\sigma' = \sigma_0 \chi_e / (1 + \chi_e)$
- conducting limit:  
 $\chi_e \rightarrow \infty$
- vacuum limit  
 $\chi_e \rightarrow 0$

## 2.3 Dielectrics

A uniformly polarized rod

$\sigma' = \mathbf{P} \cdot \mathbf{e}_n = P$ , only exist at two ends



if  $l^2 \gg S$ , these two ends can be considered as two point charge  $\pm PS$

Then the depolarizing field at the center of rod

$$E' = 2PS / [4\pi\epsilon_0(l/2)^2] = 2PS / (\pi\epsilon_0 l^2)$$

in the  $l^2 \gg S$  limit

$$E' \rightarrow 0$$

## 2.3 Dielectrics

A uniformly polarized ball

$$\sigma' = \mathbf{P} \cdot \mathbf{e}_n = P \cos \theta$$

$$dS = R^2 \sin \theta d\theta d\varphi$$

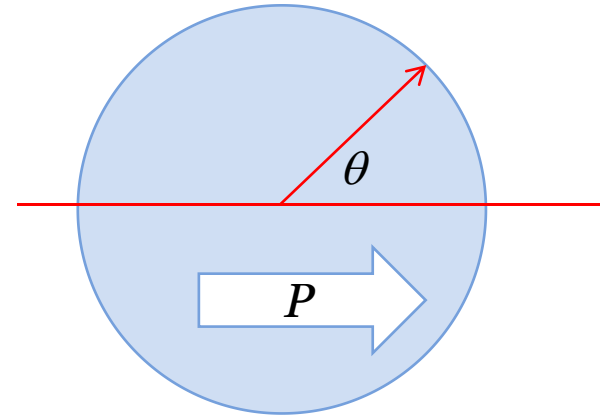
Then the depolarizing field at the center of ball

$$dE' = dq' / (4\pi \epsilon_0 R^2) = \sigma' dS / (4\pi \epsilon_0 R^2) = \sigma' \sin \theta / (4\pi \epsilon_0) d\theta d\varphi$$

$$dE_x' = -dE' \cos \theta = -P \sin \theta \cos^2 \theta / (4\pi \epsilon_0) d\theta d\varphi$$

$$E' = E_x' = \oint dE_x' = P / (2\epsilon_0) \int_0^\pi \cos^2 \theta d\cos \theta = -P / (3\epsilon_0)$$

Depolarizing field depends on the geometry!



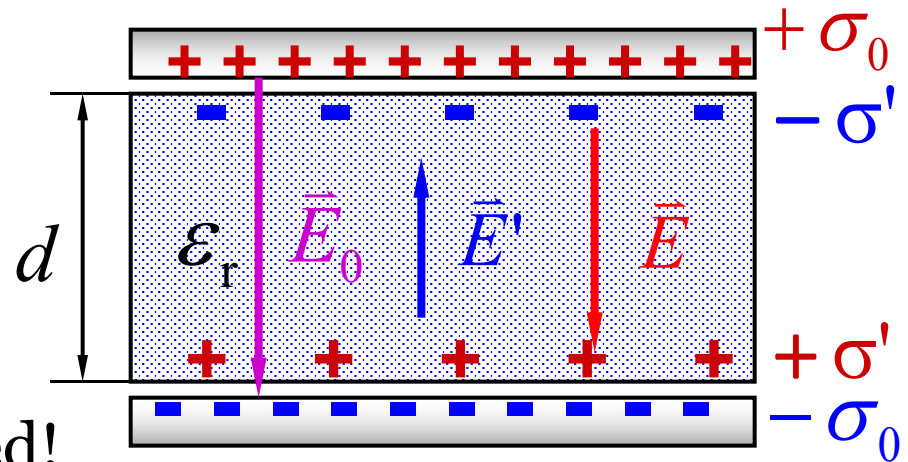
## 2.3 Dielectrics

$$E = E_0 / (1 + \chi_e) < E_0$$

$$U = Ed < E_0 d = U_0$$

$$C = Q/U = (1 + \chi_e) C_0$$

The capacitance is increased!



By defining a **relative dielectric constant**  $\epsilon_r = 1 + \chi_e$

$\epsilon_r, \chi_e$ : dimensionless quantities

Note: the linear isotropic  $P$ - $E$  relation is only an approximation, which works in most cases.

It can be nonlinear and anisotropic, depends on materials and conditions.

## 2.3 Dielectrics

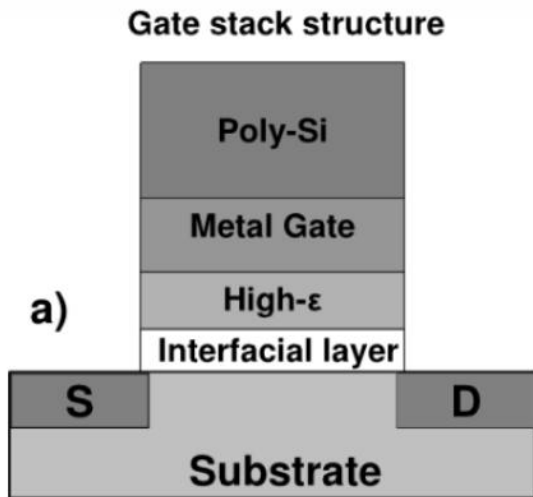
### Dielectric constants of various substances

Substance	Conditions	Dielectric constant ( $\kappa$ )
Air	gas, 0 °C, 1 atm	1.00059
Methane, CH <sub>4</sub>	gas, 0 °C, 1 atm	1.00088
Hydrogen chloride, HCl	gas, 0 °C, 1 atm	1.0046
Water, H <sub>2</sub> O	gas, 110 °C, 1 atm	1.0126
	liquid, 20 °C	80.4
Benzene, C <sub>6</sub> H <sub>6</sub>	liquid, 20 °C	2.28
Methanol, CH <sub>3</sub> OH	liquid, 20 °C	33.6
Ammonia, NH <sub>3</sub>	liquid, -34 °C	22.6
Mineral oil	liquid, 20 °C	2.24
Sodium chloride, NaCl	solid, 20 °C	6.12
Sulfur, S	solid, 20 °C	4.0
Silicon, Si	solid, 20 °C	11.7
Polyethylene	solid, 20 °C	2.25–2.3
Porcelain	solid, 20 °C	6.0-8.0
Paraffin wax	solid, 20 °C	2.1–2.5
Pyrex glass 7070	solid, 20 °C	4.00



# 2.3 Dielectrics

**Extension 1:** high- $k$  dielectric material: a vital key to continue the Moore's law



Under Embargo Until Nov. 4th, 9:00pm PT

## High-k Dielectric reduces leakage substantially

Gate

1.2nm  $\text{SiO}_2$

Silicon substrate

Gate

3.0nm High-k

Silicon substrate

Benefits compared to current process technologies

	High-k vs. $\text{SiO}_2$	Benefit
Capacitance	60% greater	Much faster transistors
Gate dielectric leakage	> 100x reduction	Far cooler

intel.

10

$\text{SiO}_2$   $\epsilon_r=3.9$

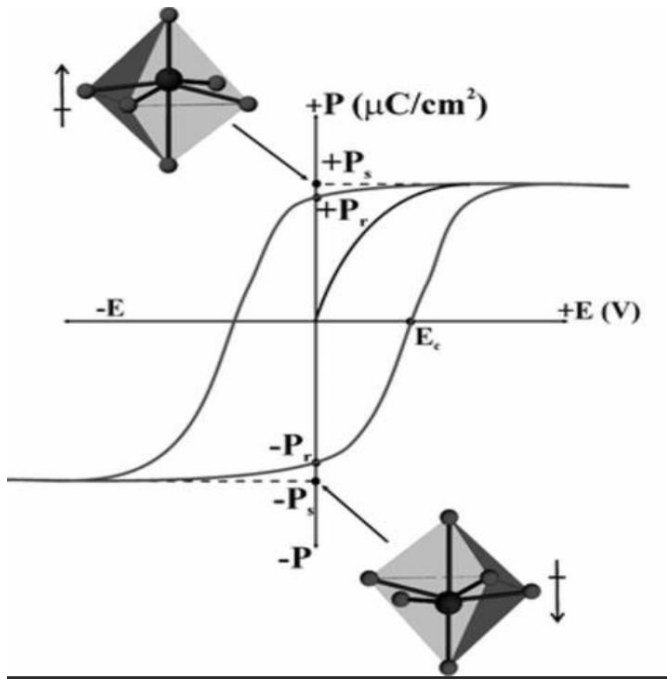
$\text{HfO}_2$   $\epsilon_r=20$

Supplemental reading:

[HKMG\(High-K 栅氧化物层 +Metal Gate\)技术](#)

# 2.3 Dielectrics

## Extension 2: ferroelectrics



Joseph Valasek  
(1897-1993)

《**Physics Today**》：  
Because Valasek was a pioneer whose main work was done well before his field became popular, and because he was very quiet and modest and did not seek recognition and honor for his work, he never achieved the recognition he deserved.”

$P$ : nonlinear to  $E$   
non-volatile states  
discovered in 1920

$\text{BaTiO}_3$ ,  $P_r=20 \mu\text{C}/\text{cm}^2$

$\text{PbTiO}_3$ ,  $P_r=80 \mu\text{C}/\text{cm}^2$

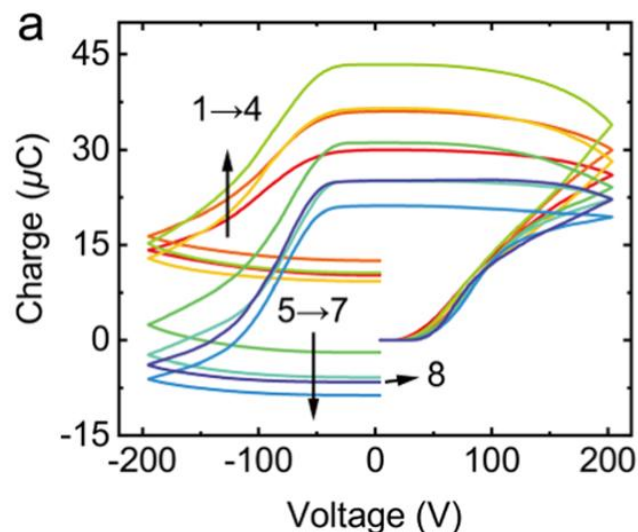
$t\text{-BiFeO}_3$ ,  $P_r=120 \mu\text{C}/\text{cm}^2$

铁电百年诞辰，归来仍是少年

## 2.3 Dielectrics

### High, Multiple, and Nonvolatile Polarizations in Organic–Inorganic Hybrid $[(\text{CH}_3)_3(\text{CH}_2\text{CH}_2\text{Cl})\text{N}]_2\text{InCl}_5 \cdot \text{H}_2\text{O}$ for Memcapacitor

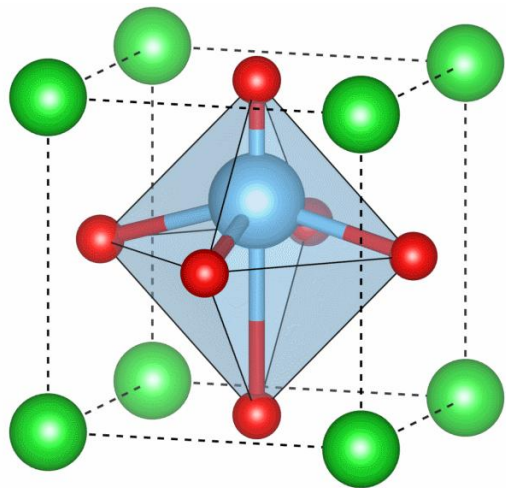
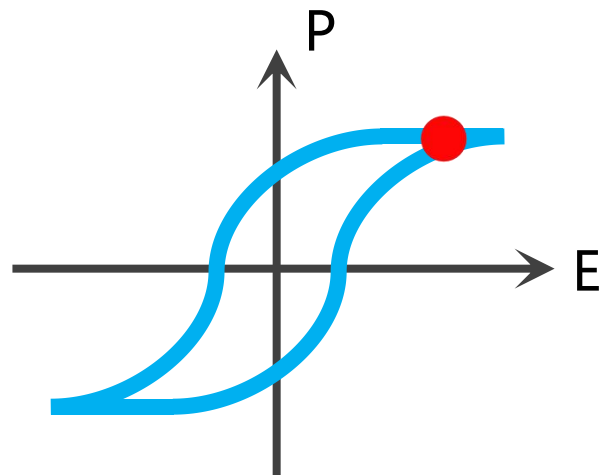
Conventional ferroelectric materials based on displacive and order–disorder types generally have difficulty meeting these requirements due to their low polarization values ( $\sim 150 \mu\text{C}/\text{cm}^2$ ) and persistent electrical hysteresis loops. In this study, we report a novel organic–inorganic hybrid  $(\text{CETM})_2\text{InCl}_5 \cdot \text{H}_2\text{O}$  ( $\text{CETM} = (\text{CH}_3)_3(\text{CH}_2\text{CH}_2\text{Cl})\text{N}$ ) exhibiting an intriguing polarization vs electric field (charge vs voltage) “hysteresis loop” and a record-high nonvolatile polarization over  $30000 \mu\text{C}/\text{cm}^2$  at room temperature.





# 铁电物理

铁电百年仍弥坚，  
量子机制敢为先。  
理论计算调极性，  
磁电对偶续新篇。



**第一代**  
罗息盐等  
分子铁电体

**第二代**  
钛酸钡、锆钛酸铅等  
氧化物铁电体

**第三代**  
磁性铁电体、  
低维铁电体等

1920s

1940-1950s

2000s

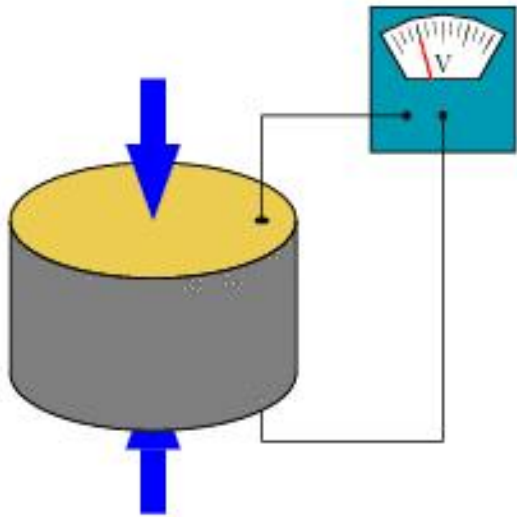
2020s

铁电经典理论

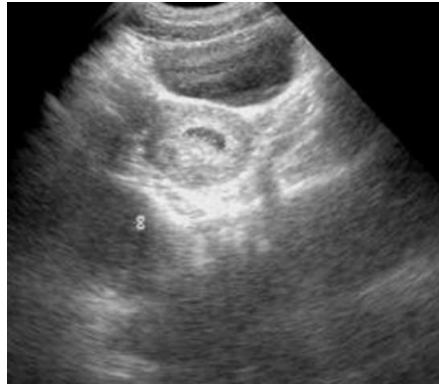
铁电量子理论

# 2.3 Dielectrics

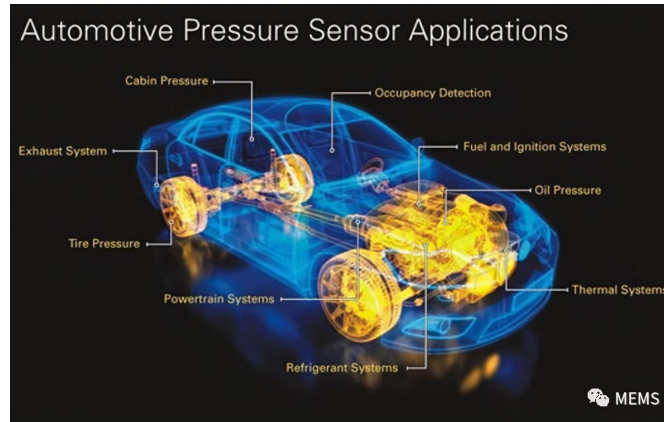
## Extension 3: piezoelectricity



$P$  changes upon pressure  
Microelectromechanical  
Systems (MEMS)



B-scan  
ultrasonography



sensors in  
industry



biosensors



# 2.3 Dielectrics

## Extension 4: electrets



For comparison:

Graphene masks

Inventors:

孙立涛 (SEU)

ultra-large surface

N95 masks

Inventors:

蔡秉燊&刘朝宇(UTK)

electrets

Weakness:

Moisture

sensitive

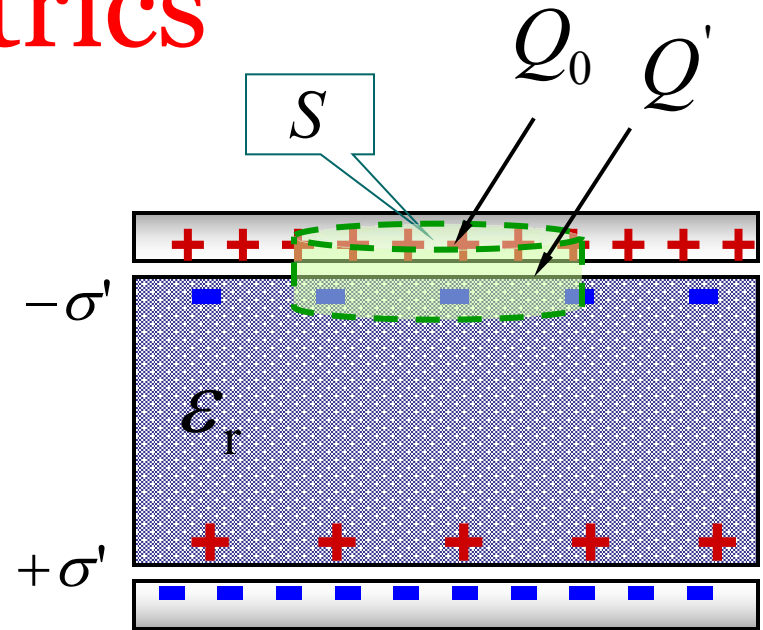
## 2.3 Dielectrics

$$\oint_S \mathbf{E} \cdot d\mathbf{S} = (Q_0 - Q') / \epsilon_0$$

$$Q' = \chi_e / (1 + \chi_e) Q_0 = (1 - 1/\epsilon_r) Q_0$$

$$\oint_S \mathbf{E} \cdot d\mathbf{S} = Q_0 / \epsilon_0 \epsilon_r$$

$$\oint_S \epsilon_0 \epsilon_r \mathbf{E} \cdot d\mathbf{S} = Q_0$$



By defining a new vector field

$$\mathbf{D} = \epsilon_0 \epsilon_r \mathbf{E} = \epsilon \mathbf{E} = \epsilon_0 \mathbf{E} + \mathbf{P}$$

**D**: electric displacement vector;  $\epsilon$ : permittivity of dielectric

Gauss's law in dielectrics:  $\oint_S \mathbf{D} \cdot d\mathbf{S} = Q_0$  or  $\nabla \cdot \mathbf{D} = \rho_0$

Only free carriers are involved.  $\nabla \cdot \mathbf{P} = -\rho'$

## 2.3 Dielectrics

Electric field of a uniformly charged sphere/ball in uniform dielectric medium

outside:

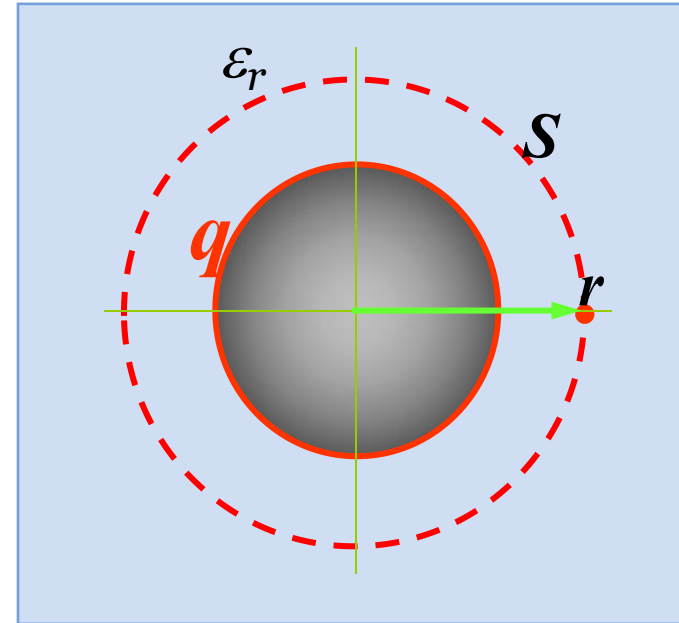
$$\oint_S \mathbf{D} \cdot d\mathbf{S} = Q_0$$

$$4\pi r^2 D = Q_0$$

$$\rightarrow D = Q_0 / (4\pi r^2)$$

$$\mathbf{E} = \mathbf{D} / \epsilon_0 \epsilon_r = Q_0 \mathbf{e}_r / (4\pi \epsilon_0 \epsilon_r r^2)$$

$$= \mathbf{E}_0 / \epsilon_r$$





## 2.3 Dielectrics



**Attention:** whether  $\mathbf{E}=\mathbf{E}_0/\epsilon_r$  generally correct?

**Answer:** no!

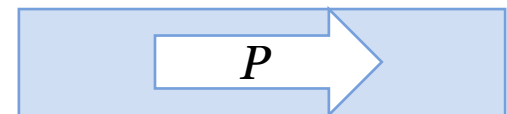
The conditions:

- 1) uniform dielectric medium in the whole space
- 2) uniform dielectric medium with isopotential surface

**Example:** A uniformly polarized rod

$\mathbf{E}' \sim 0$  in the middle point

$\mathbf{E} \approx \mathbf{E}_0 \neq \mathbf{E}_0/\epsilon_r$



# 2.4 Energy of electrostatic fields

The electrostatic energy of a plane-parallel capacitor:

$$W_e = QU/2 = CU^2/2$$

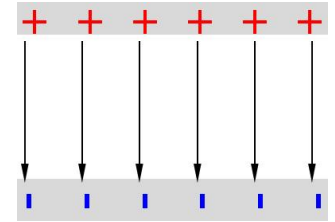
$$= \epsilon S/d (Ed)^2/2 = \epsilon E^2 V/2$$

Energy density

$$w_e = W_e/V = \epsilon E^2/2 = DE/2$$

$W_e = \int w_e dV$  generally works beyond the  
plane-parallel capacitor

**Philosophy:** the energy is carried by the field,  
instead of charge!



## 2.4 Energy of electrostatic fields

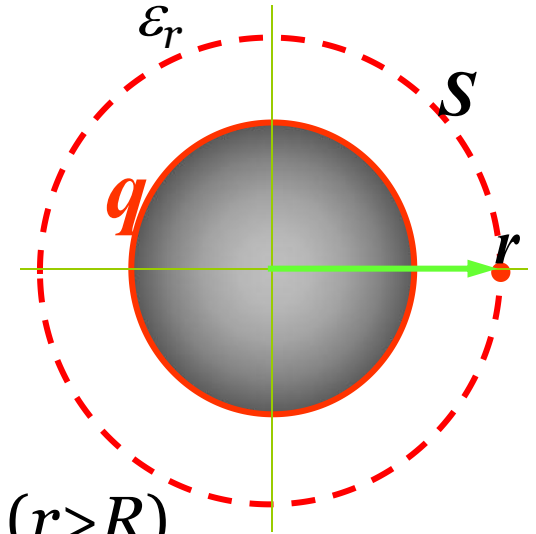
From the viewpoint of field:

**Case 1:** a uniformly charged sphere

$$\begin{aligned}W_e &= \iiint w_e dV = \iiint \varepsilon_0 E^2 dV / 2 \\ &= \varepsilon_0 / 2 \iiint [q / (4\pi\varepsilon_0 r^2)]^2 dV \quad (\text{outside}) \\ &= q^2 / (32\pi^2 \varepsilon_0) \iiint 1/r^4 dV = q^2 / (8\pi\varepsilon_0) \int 1/r^2 dr \quad (r > R) \\ &= q^2 / (8\pi\varepsilon_0 R) \quad \text{equal to the value calculated from charge}\end{aligned}$$

In dielectric medium,

$$W_e = \iiint w_e dV = \iiint DE dV / 2 = q^2 / (8\pi\varepsilon_0 \varepsilon_r R)$$



# Chapter 2: Homework

1. 9-2, 9-3, 9-5, 9-11, 9-14
2. Additional homework: to calculate the field energy of a uniformly charged ball
3. Deadline: Tuesday, May 7