# (Super) Capacitors

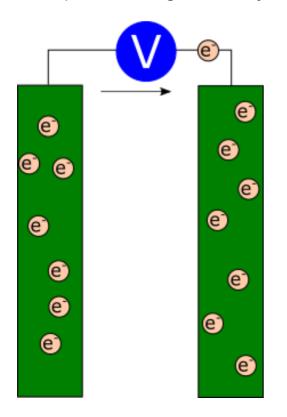


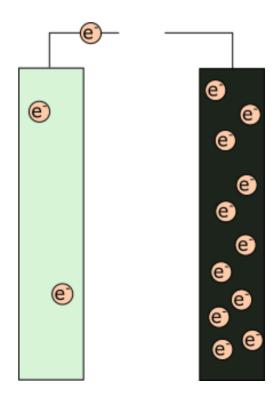
# Faster (dis)charging

Batteries are a good charge-storage solution, but are limited in how quickly they can discharge (safely).

• Ultimately, redox processes can only occur so fast...

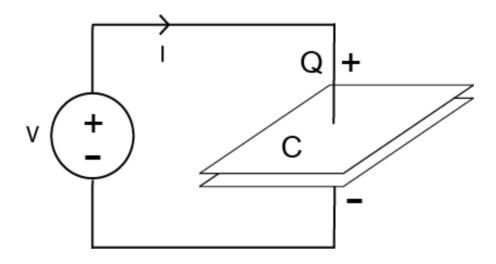
Why not separate charges directly?





#### This is a Capacitor

An arrangement of electrodes of area A, separated by a distance d.



Two electrodes separated by vacuum have a capacitance *C*:

$$C = \frac{\epsilon_0 A}{d}$$

where  $\epsilon_0$  is the permittivity of free space = 8.854 × 10<sup>-12</sup> C<sup>2</sup> J<sup>-1</sup> m<sup>-1</sup>

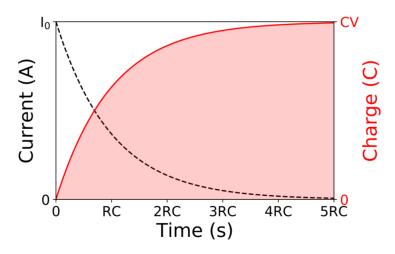
## Charge stored

On charging a capacitor with a constant voltage, current decays with time:

$$I_t = I_0 e^{\left(rac{-t}{RC}
ight)}$$

The charge stored increases with time:

$$Q_t = CV \left[ 1 - e^{\left(rac{-t}{RC}
ight)} 
ight]$$



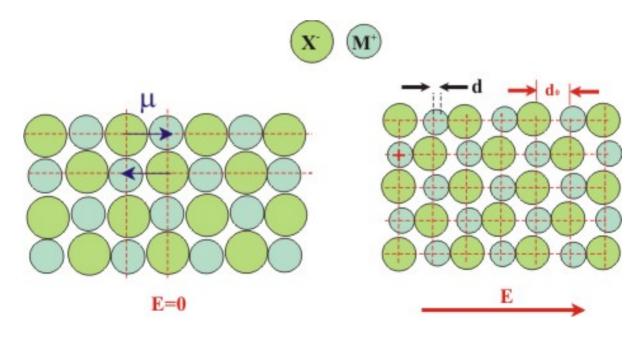
The maximum charge stored, Q = CV

#### Increasing the charge stored

 $C=rac{\epsilon_0 A}{d}$ , so decreasing d or increasing A will increase stored charge. If d gets too small, however, electrons will tunnel from one plate to the other.

#### Alternatively, use a dielectric

• An electrically insulating material in which an applied electric field causes a **displacement** (but not a flow) of charge.

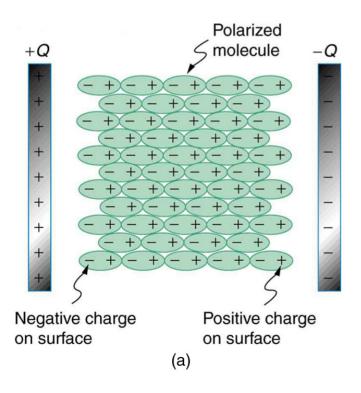


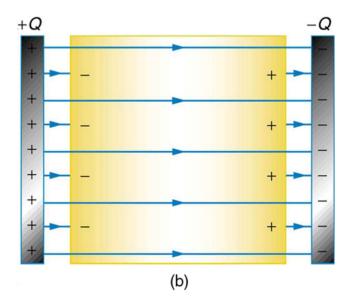
#### Dielectric capacitor

Adding a dielectric between the plates increases the charge capacity.

$$C_{ ext{dielec}} = rac{\epsilon_r \epsilon_0 A}{d}$$

where  $\varepsilon_r$  is the relative permittivity of the dielectric ( $\varepsilon_r$ =  $\varepsilon$  /  $\varepsilon_0$ ) and  $\varepsilon_r$  >  $\varepsilon_0$ 





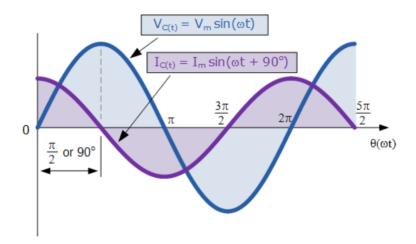
# Example permittivities

Material	Relative Permittivity, $\epsilon_r$
Vacuum	1
Paper	2.0 - 6.0
Polymers	2.0 - 6.0
Silicon oil	2.7 - 2.8
Quartz	3.8 - 4.4
Glass	4 - 15
$Al_2O_3$	10
Ta <sub>2</sub> O <sub>5</sub>	26
TiO <sub>2</sub>	100
CaTiO <sub>3</sub>	130
SrTiO <sub>3</sub>	285
BaTiO <sub>3</sub>	1000 - 10000

## Characterising dielectrics

Apply an alternating (sinusoidal) field, and measure the resulting current *and phase* shift - **Impedance spectroscopy** 

- ullet Applied field,  $E_t=E_0\sin(\omega t)$  , where  $\omega=2\pi f$
- Response current,  $I_t = I_0 \sin(\omega t + \phi)$



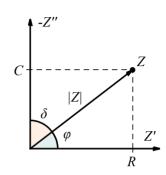
The total impedance  $(Z(\omega)=rac{E_t}{I_t})$  can be represented as a complex number:

- $Z(\omega) = Z_0(\cos\phi + i\sin\phi)$
- In an ideal dielectric, current and voltage should be 90° out-of-phase ( $\varphi = 90^\circ$ ).
- For a resistive material, current and voltage should be in phase ( $\varphi = 0^{\circ}$ ).

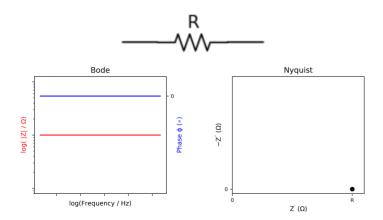
# Impedance analysis

Two standard ways to display data:

- Bode plot: |Z| and φ plotted against frequency
- **Nyquist plot**: -Z" (90° out-of-phase) against Z' (in phase).



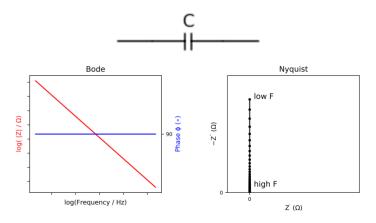
#### Ideal Response



Single Z' value for all frequencies

$$\circ$$
 Z" = 0

• 0° phase shift



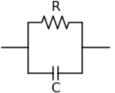
Z" varies with frequency

$$\circ$$
 Z' = 0

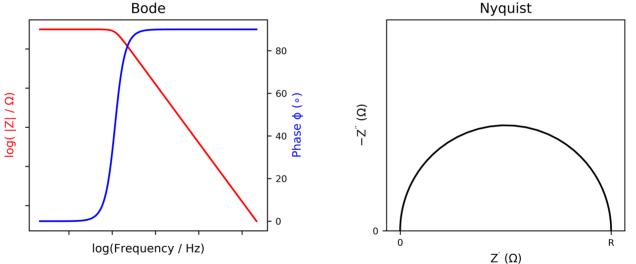
90° phase shift

#### 'Real' Impedance

Many real materials exhibit behaviour like a parallel RC circuit:



<sup>C</sup> Ions flow



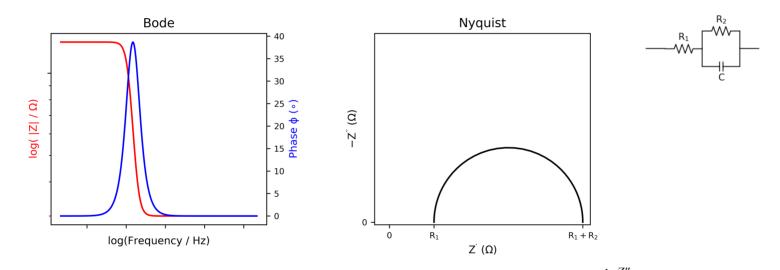
electrode (see later)

• Ionic conduction in a ceramic material building up a charge gradient

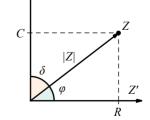
More complex behaviour is often observed, and can be modelled using *equivalent* circuits

#### Real dielectric response

Changing the electric field direction causes the dipoles to rearrange with a characteristic timescale. This timescale means *real* dielectrics show a peak in  $\phi$  with frequency, corresponding to the maximum energy loss.



This *dielectric loss* is often characterised as  $\tan\delta=\frac{Z'}{-Z''}$ , where  $\delta=90^\circ-\phi$ 

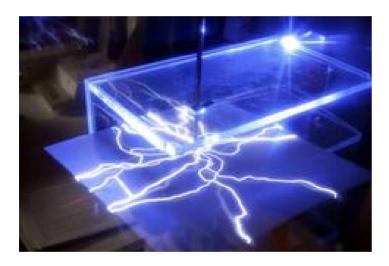


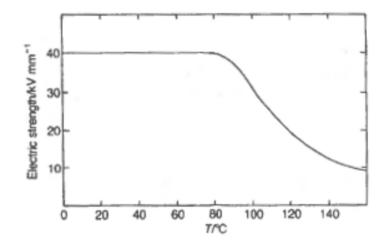
tan δ must be minimised for applications

#### Dielectric breakdown

Dielectrics also break down under high electric fields

- Electrons start to conduct, causing localised heating and breakdown
- This is quantified as the **Dielectric Strength** (in *e.g.* V m<sup>-1</sup>)





# How good is a capacitor for energy storage?

Take e.g. a BaTiO<sub>3</sub>-based capacitor:

- $\varepsilon_r \approx 1000$
- Dielectric strength ≈ 10 MV m<sup>-1</sup>
- Thickness  $(d) \approx 1 \, \mu \text{m}$

Assuming a total volume of 5 cm<sup>3</sup> (similar to an AA battery):

$$A = rac{5 imes 10^{-6}}{1 imes 10^{-6}} = 5 ext{ m}^2$$

$$C=rac{\epsilon_r\epsilon_0A}{d}=0.04427\,\mathrm{F}$$

Maximum voltage =  $10 imes 10^6/1 imes 10^{-6} = 10 ext{ V}$ , therefore:

$$Q=CV=0.4427 ext{ Coulombs}=0.1229 ext{ mAh}$$

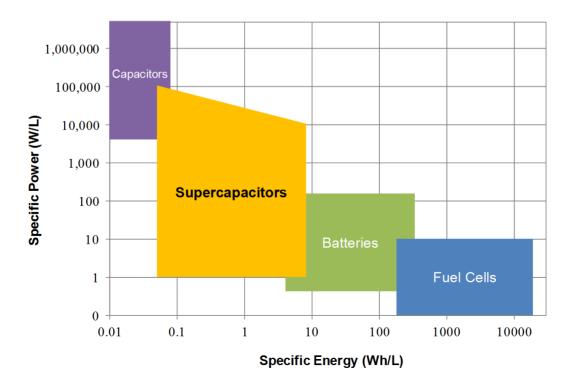
Volumetric charge storage of 24.6 mAh  $L^{-1}$ : Energy capacity = 0.245 Wh  $L^{-1}$ 

## Why are they useful then?

Although energy capacity is worse than for batteries (often by a lot), capacitors can discharge the charge very rapidly

A discharge current of 100 Amps can be easily achieved.

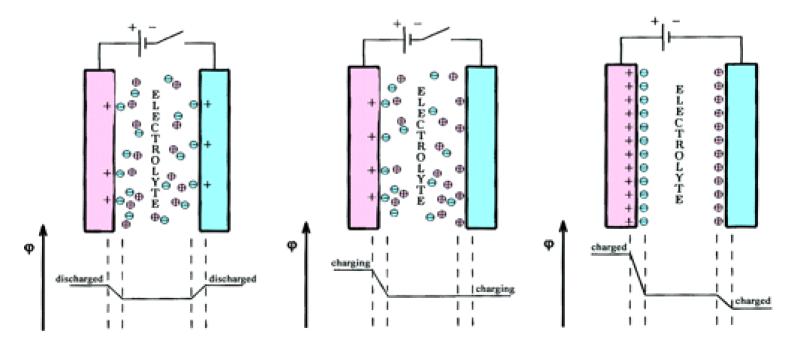
This gives a high specific power. For the BaTiO<sub>3</sub> capacitor considered before, this gives power density approximately 1000 W L<sup>-1</sup>!



#### Supercapacitors

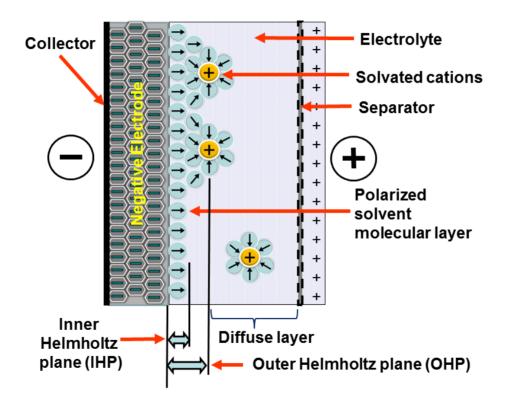
- Higher capacitance (but lower voltage limits) than other capacitors
- Sometimes known as ultracapacitors or electrostatic double layer capacitors (EDLCs)

Rather than a ceramic dielectric, supercapacitors rely on an ionic electrolyte solution, and an ion-permeable membrane to prevent electronic conduction.



#### Supercapacitor operation

Charge is stored in a **Helmholtz double layer** at each electrode:



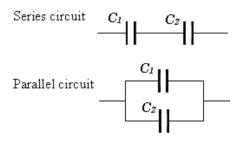
This separation of +ve and -ve charges occurs over a few angstroms

This is effectively a capacitor with very small d

#### Capacitance

Because there are two double-layers, a supercapacitor behaves as two capacitors connected in series:

$$rac{1}{C} = rac{1}{C_a} + rac{1}{C_c}$$



For a symmetric supercapacitor (commonly called ultracapacitor, where anode and cathode are the same material):

$$C=rac{C_A}{2}$$

The total energy stored is:

$$E=rac{CV^2}{2}$$

Charging voltages are typically 1-3 V (depending on electrolyte).

# Electrolytes Aqueous

- Acids (e.g. H<sub>2</sub>SO<sub>4</sub>)
- Alkalis (KOH)
- NaClO<sub>4</sub> or LiClO<sub>4</sub>
- LiAsF<sub>6</sub>

#### Organic

*e.g.* acetonitrile, propylene carbonate, tetrahydrofuran with:

- Tetraethylammonium tetrafluoroborate, N(Et)<sub>4</sub>BF<sub>4</sub>
- Triethyl(methyl) tetrafluoroborate, NMe(Et)<sub>3</sub>BF<sub>4</sub>

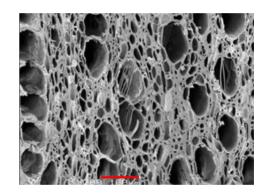
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← Cheaper
Higher voltage →
Wider temperature range →
← Higher conductivity
← Higher specific power
Higher specific energy →
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#### Electrode materials

- C  $\propto$  amount of double-layer  $\propto$  A  $\rightarrow$ 
  - electrodes are designed to have maximum area

#### Porous (activated) carbon

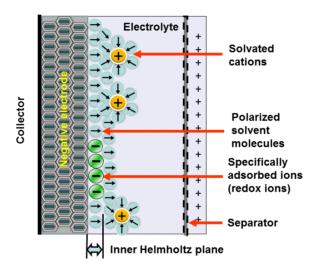
- Surface area exceeding 3000 m<sup>2</sup> g<sup>-1</sup>
- Trade-off between surface area and pore size
  - Smaller pores limit maximum current (power density) but increase energy capacity
- Activated carbon is relatively expensive and potentially unsustainable
  - High temperatures and aggressive chemical activation required
  - Biochar (a by-product of biofuel production) is one alternative





#### Pseudocapacitance

One way to increase energy storage in supercapacitors is to add redox-active species.



- Must be fast, reversible redox processes (so that power density remains high)
- Pseudocapacitance can contribute 100 times the double-layer capacitance
- Redox ions must have affinity for the electrode(s)
  - MnO<sub>2</sub> is commonly used as an electrode

NOTE: In order to be considered pseudocapacitance, charge stored must depend linearly on the applied voltage (otherwise it is behaving like a battery)

## Hybrid technologies

One of the main drawbacks of supercapacitors is that their voltage drops with time

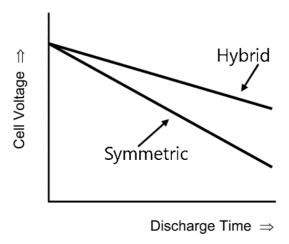
Not ideal for powering devices

Hybrid capacitors combine a battery-electrode with a supercapacitor electrode

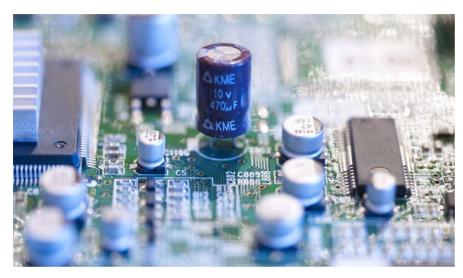
- e.g. replace carbon cathode with NiOOH
- C<sub>battery</sub> ≈ 10 × C<sub>supercap</sub>

$$\circ$$
  $\frac{1}{C} = \frac{1}{C_A} + \frac{1}{10C_A} pprox \frac{1}{C_A}$ 

• In some cases (e.g. thin-film Li electrodes) fast redox kinetics still allow high power applications



# (Super) Capacitor applications



#### Regenerative braking

- Recovers kinetic energy lost when braking
- Large currents generated; supercapacitors required to store charge

#### **Medical devices** *i.e.* pacemakers

 Takes advantage of their lowmaintenance / long life



