

**EE-517** 

# Bio-Nano-Chip Design

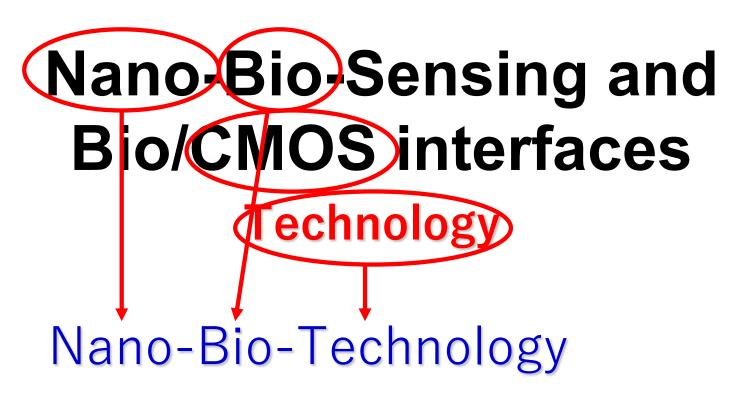


EE-517
Bio-Nano-Chip
Design

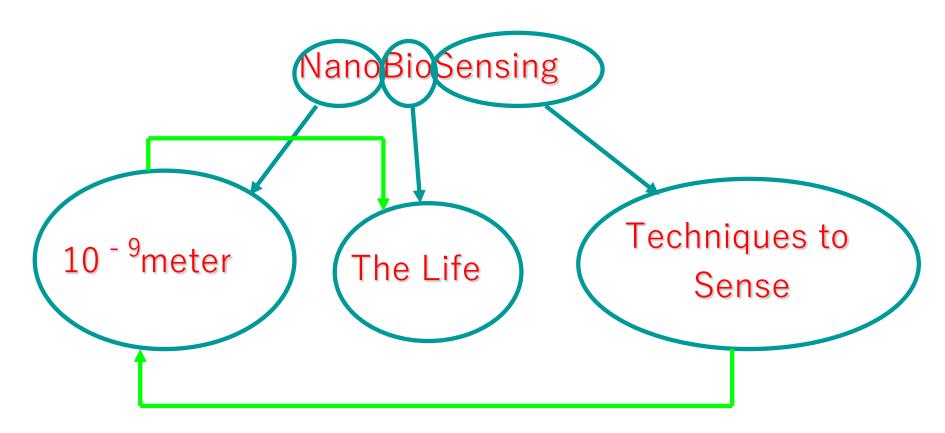
- 1. LECTURES by videos (on SWITCHtube)
- 1. Q/A on Zoom (link by Moodle)

(c) S.Carrara

### What's about The Course?



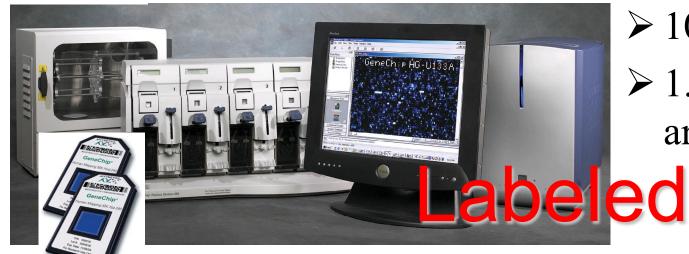
## What's about Nano-Bio-Sensing?



That means "techniques at the nano-metric scale to sense information related to biological processes"



### The Motivation



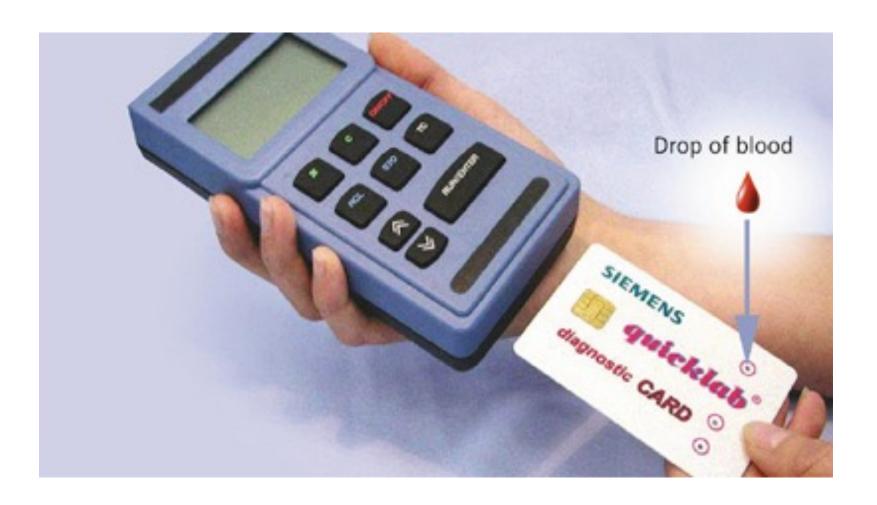
- > 100.000 \$ (machinery)
- > 1.000 \$ the single μ-array

Label-Free

- > 50 \$ (machinery)
- > 0.05 \$ the single strip



# The Quicklab project by Siemens



## Glucometer on iPhone

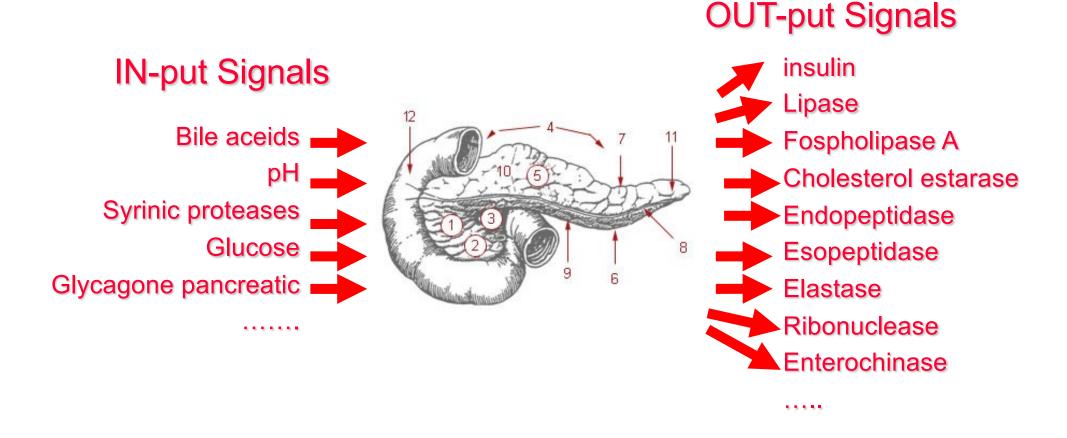


## Next step: the future already begun



Glucose Personal Diagnostics on our Skin

## **The Pancreas Functions**



## What to sense?

```
Simple Molecules | Glucose (Diabetes) |
Cholesterol (heart attack ) |

AFP (Hepato Carcinoma) |
PSA (Prostate) |

DNA | PC-1 gene (prostate cancer) |
psa gene (Hepato Carcinoma)
```

Bio-Markers may be simple molecules, proteins or genes

## What else to sense?

```
Endogenous<br/>MetabolitesInsulin (Diabetes)<br/>β-blockers (heart attack )Anti-cancer<br/>compoundsCyclophosphamide (Breast Cancer)<br/>Docetaxel (Prostate Cancer)AnestheticsPropofol (surgery)<br/>Midazolam (surgery)
```

# Endogenous and Exogenous Metabolites are usually simple molecules

## What to sense?

Simple Molecules Glucose (Diabetes)
Cholesterol (heart attack)

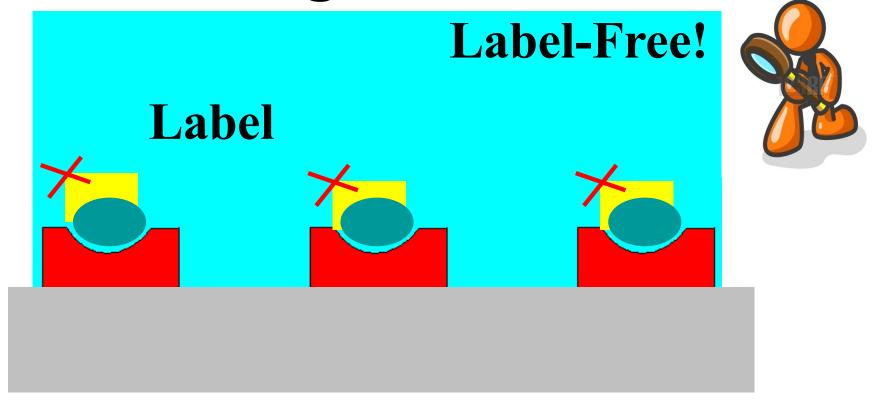
Proteins PSA (Prostate)

DNA PC-1 gene (prostate cancer)
sequences p53 gene (Hepato Carcinoma)

Markers?

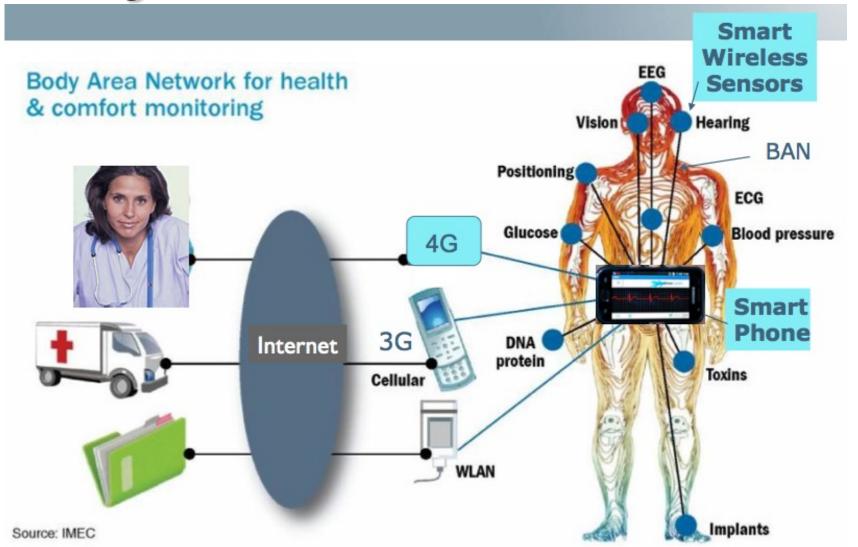
Bio-Markers may be simple molecules, proteins or genes

# Measuring Bio-Markers



The Measure of Bio-markers may be performed in a labeled manner or in label-free mode

# Fully-Connected Human++



Courtesy, Hugo De Man (IMEC)

## The Time' forecast on Human++



[TIME, February 2011]

# Chips under the skin?



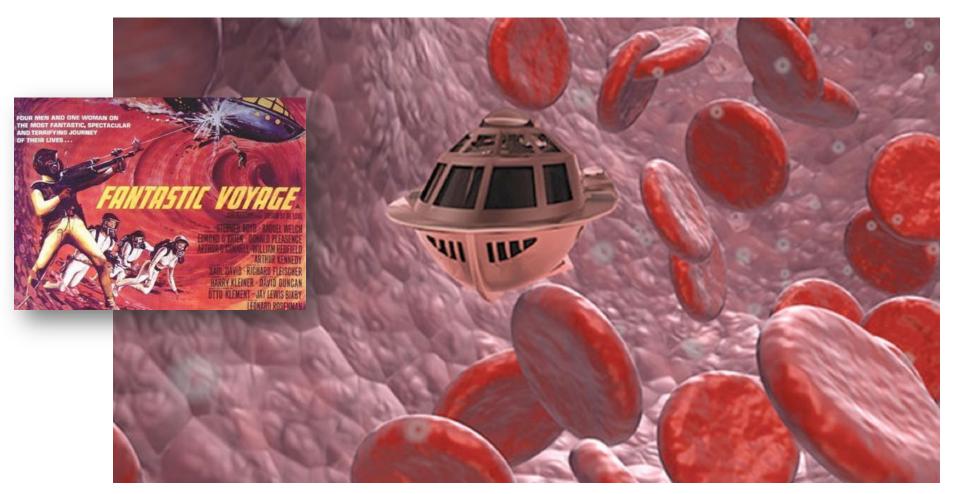
Under the skin: how insertable microchips could unlock the future

Volunteers in Melbourne have had microchips inserted for three months, designed to unlock doors and carry out other tasks. Will they really be any use?

THEGUARDIAN.COM

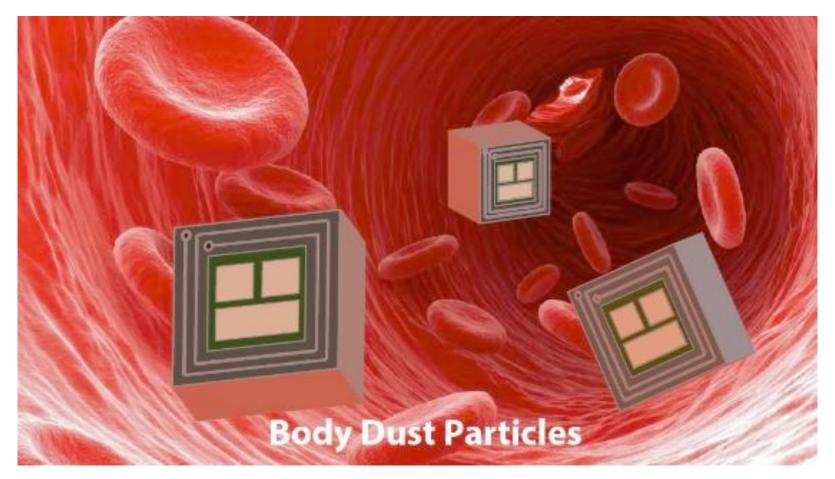
[The Guardian, October 2017]

# How small Chips under the skin?



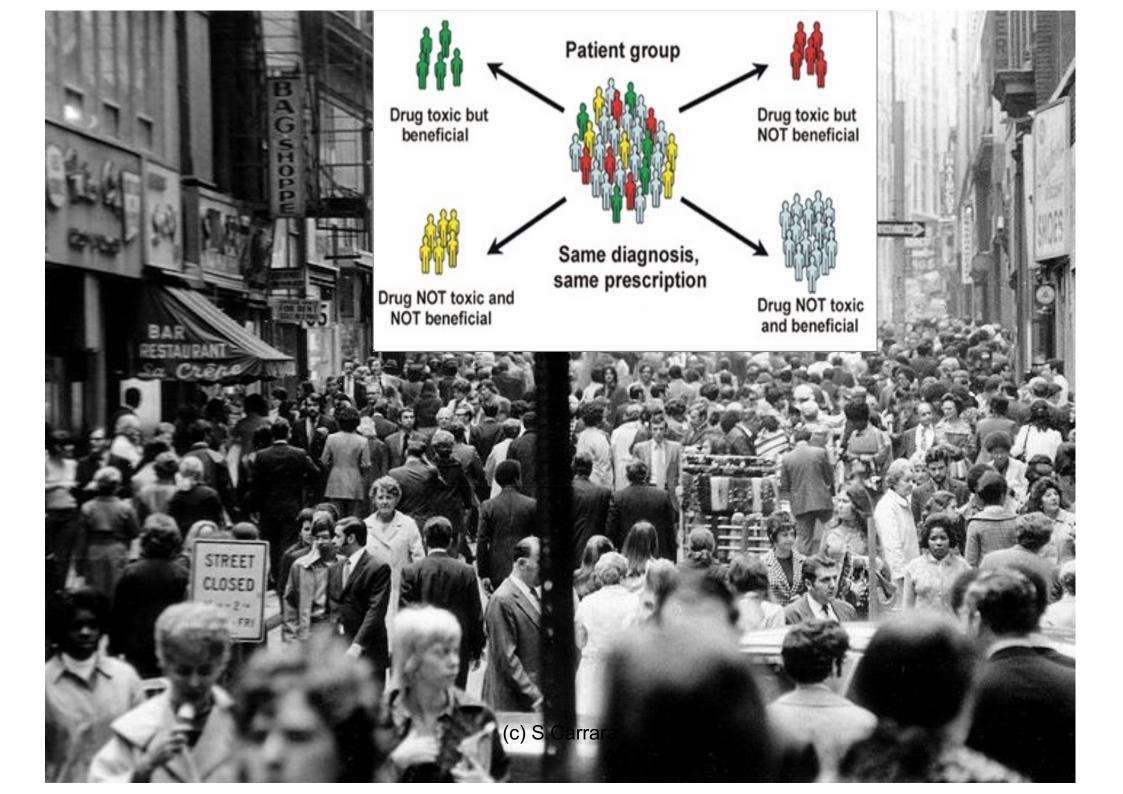
[1966 Sci. Fi. movie titled "Fantastic Voyage"]

# **Body Dust: Drinkable CMOS Bioelectronics**

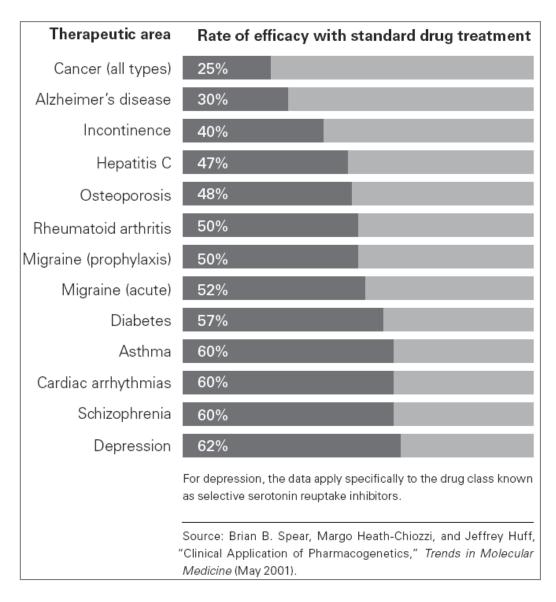


Tracking cancer-cell developing "drinkable" electronic sensors

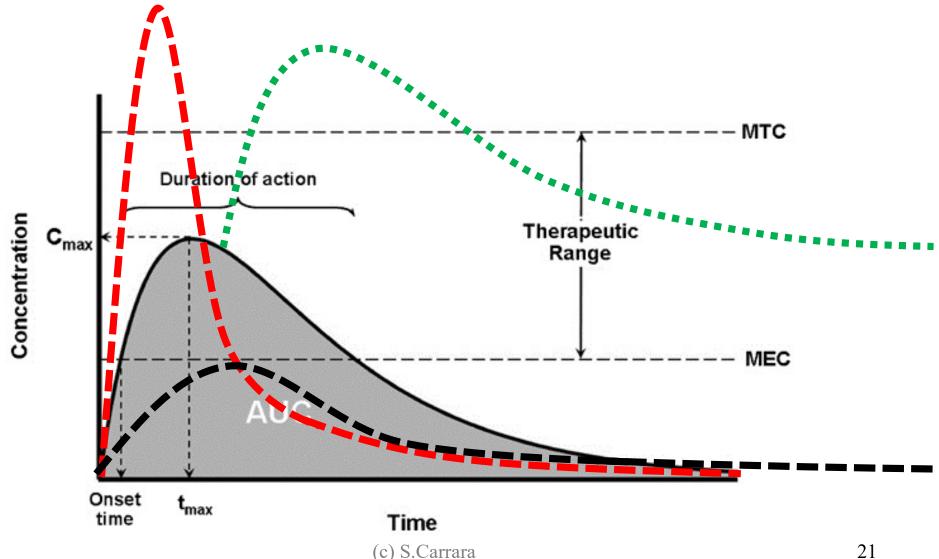
https://actu.epfl.ch/news/tracking-cancer-cell-development-with-drinkable-el/



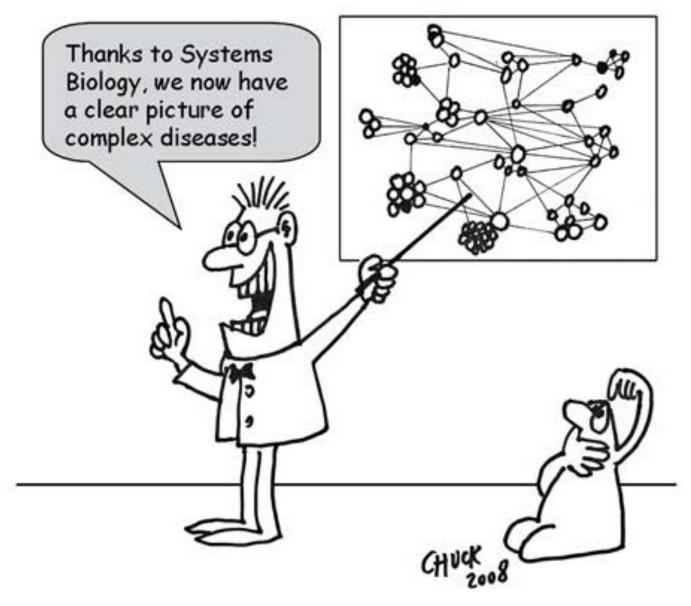
# Low efficacy of used compounds



# Personalized Therapy: the right dose in the right moment!

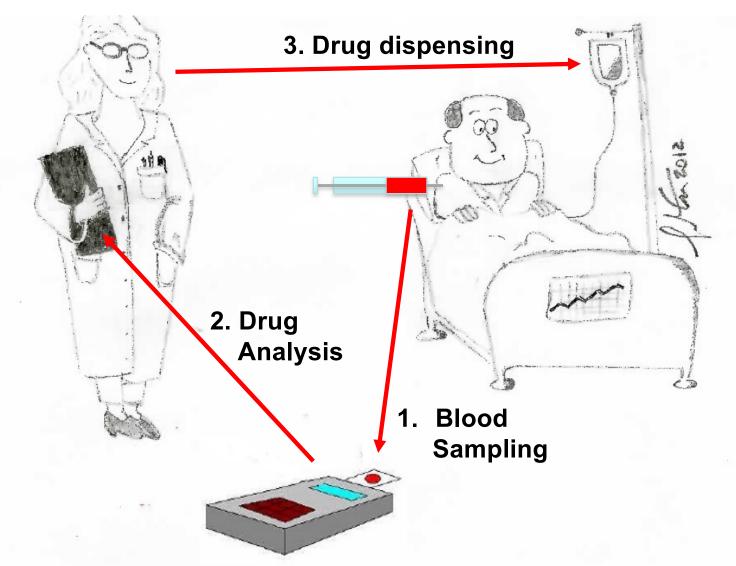


### System Biology is not enough



(c) S.Carrara

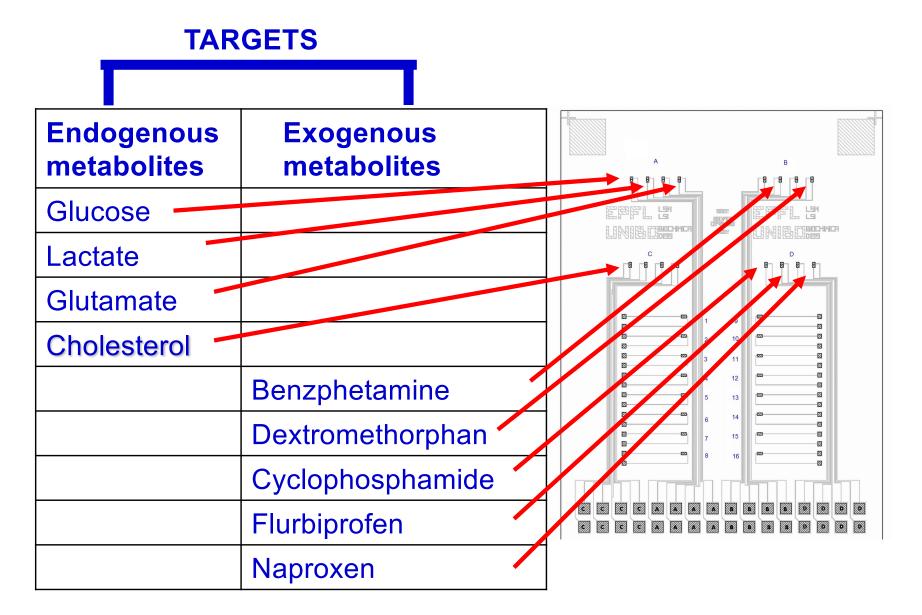
## Personalized Therapy



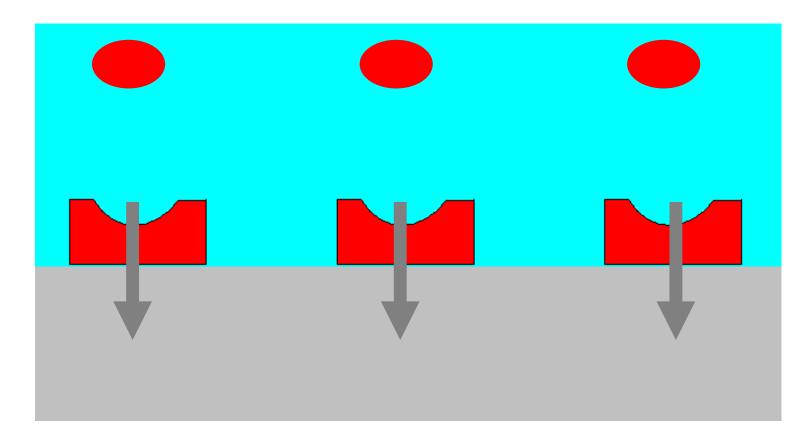
Drugs injection based on patient's pharmacokinetics

(c) S.Carrara

#### The need for new Bio/CMOS systems



# **CMOS/Sample interface**



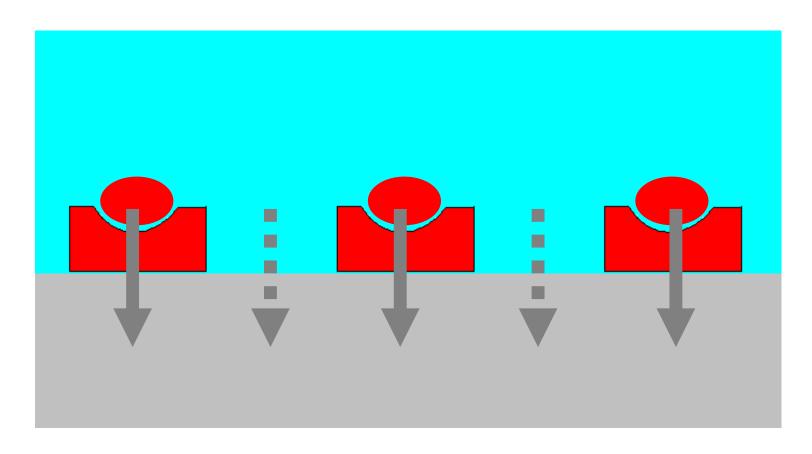
The interface between the CMOS circuit and the biosample needs to be deeply investigated and organized

## **Applications in Personalized Therapy**

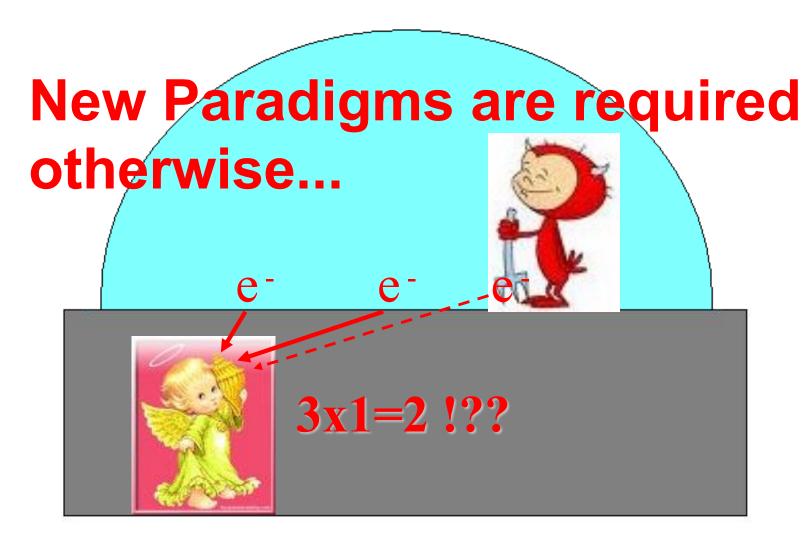
		TAR	GETS
	Probe Enzymes	Endogenous metabolites	Exogenous metabolites
	Glucose Oxidase	Clucose	
	Lactate Oxidase	Lactate	
	Glutamate Oxidase	Glutamate	
	P450 11A1	Cholesterol	
	P450 2B4		Benzphetamine
	P450 3A4		Dextrometorphane
	P450 3A4		Cyclophocphamide
\'	P450 2C9		Flurbiprofene
\	P450 2C9		Naproxene

(c) S.Carrara

# **CMOS/Sample interface**

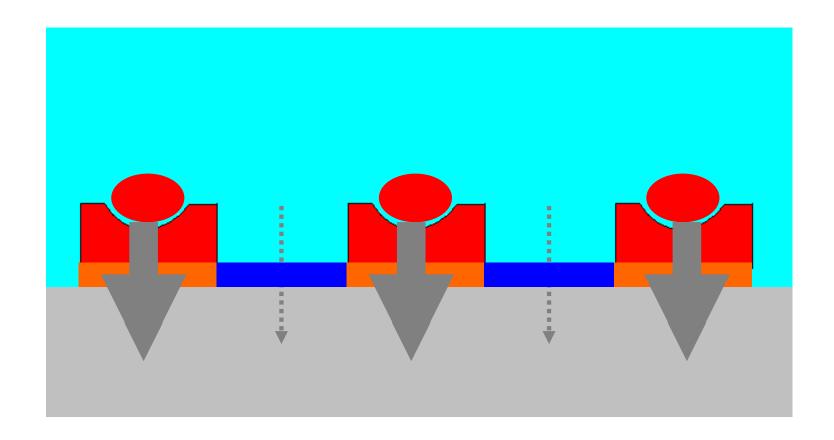


The interface between the CMOS circuit and the biosample needs to be deeply investigated and organized



Excellent CMOS technology is not sufficient if molecules are not doing their own job at the Bio/CMOS interface!

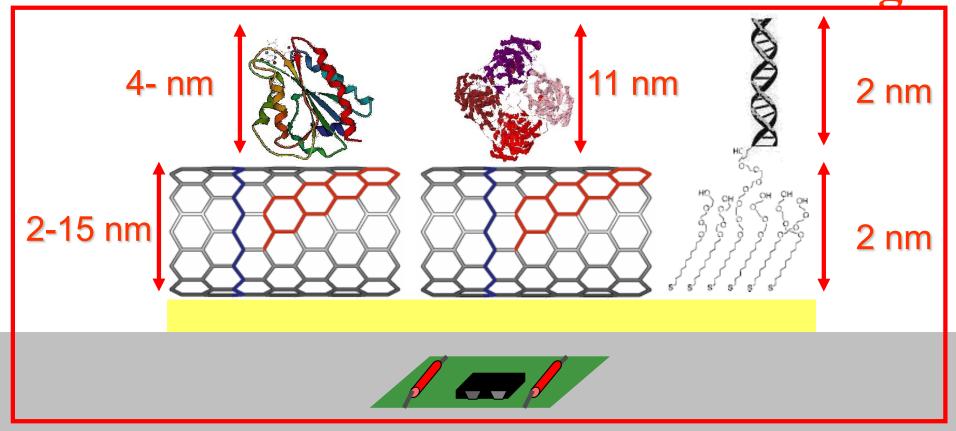
# **CMOS/Sample interface**



The interface between the CMOS circuit and the biosample needs to be deeply investigated and organized

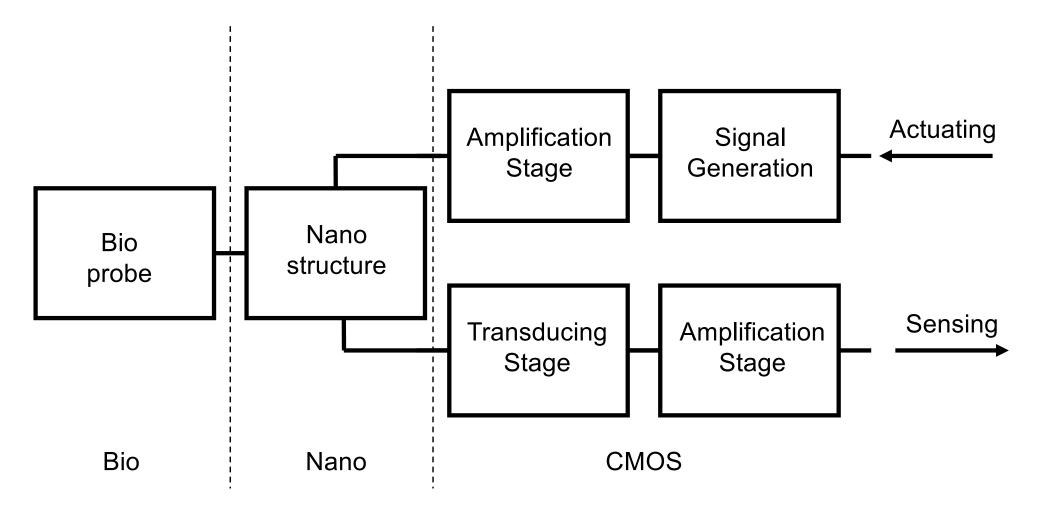
## **New Approach**

Bio/Nano/CMOS Co-Design!



New paradigms for Nano-Bio-CMOS co-design are required to succeed in chip bio-sensing

## Design of the all interfaces



#### The Bio/Nano/CMOS interface



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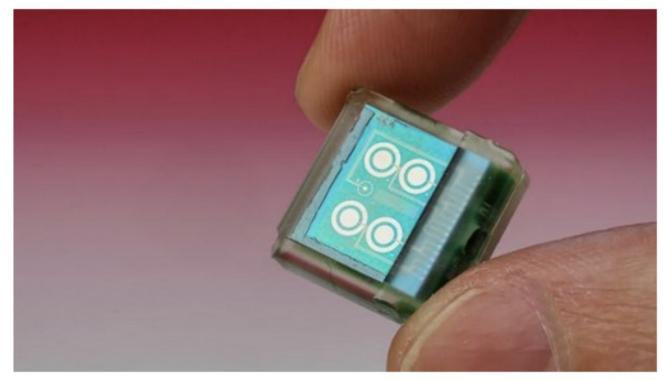
Trending News Charleston shooting Brookfield Zoo Multiple Scierosis Joe Biden

# A subcutaneous biosensor chip to revolutionize tomorrow's medicine

May 29, 2015 9:26 AM Relaxnews











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The microchip implants that let you pay with your hand

By Katherine Latham Business reporter

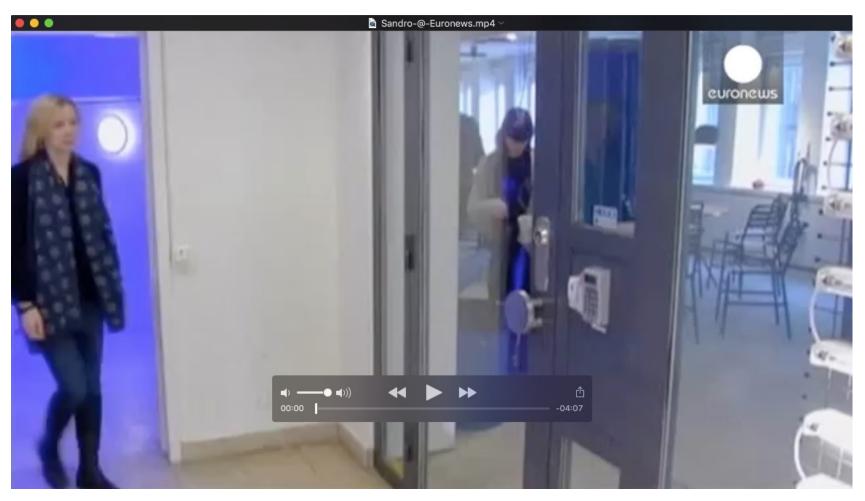
O April, 11th, 2022



An x-ray showing a Walletmor implant, which are injected into a person's hand after a local anaesthetic

For many of us, the idea of having such a chip implanted in our body is an appalling one, but a 2021 survey of more than 4,000 people across the UK and the European Union found that 51% would consider it.

# Chips under the skin?



[EuroNews, June 2015]

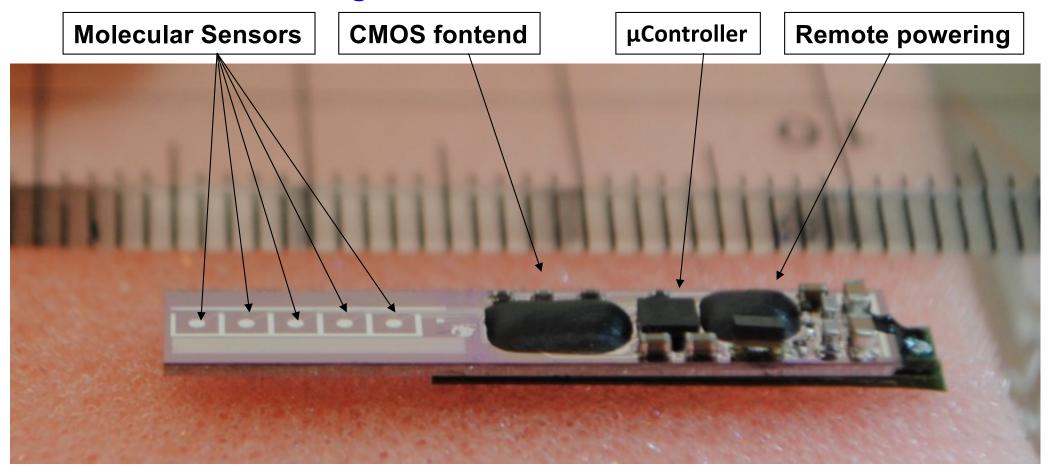
# Under the skin for body sculpting



# **Enhancing human being**



## Size and Shape to be injectable as a Needle?



The IC has been fabricated in UMC 0.18 technology and interfaced to the passive multi-panel platform

## **ECG** monitoring by Medtronic



Mark Phelps by Medtronic, and the Reveal LINQ<sup>TM</sup> system



EE-517: Bio-Nano-Chip Design

Subject of the week	Chapter' paragraphs*
Introduction to Bio-Nano-Chip design, and Conductive Solutions	§1.1-1.5, §2.1-2.7, §2.14-15
Biological molecules: Proteins and DNA building blocks	§3.5-9, §4.13 and §4.17-18
Biological molecules interactions (DNA, Antibodies, Oxidases and Cytochromes)	§4.4-17 and §4.19-23
Biosensors Principle with DNA, Antibodies, and Enzymes	§6.1-4 and §8.2
Biosensors Principle with DNA, Antibodies, and Enzymes  Biosensors Principle by Redox reactions and Faradaic processes	§8.4-8
Nanotechnology for molecular assembly on chip' surfaces (absorption models)	§5.1
Nanotechnology for checking molecular assembly on chip' surfaces (SPR+ AFM)	§5.2
Nanotechnology to prevent electron transfer	§6.3-7
Nanotechnology to enhance electron transfer in redox reactions	§8.4-8, and 8.3 and 8.9
Chip design for electrochemical sensing: basic configurations and equivalent circuits	§9.1-9.2
Amperometric biosensing in constant-bias (Current-to-Voltage & FTCC Methods)	§9.1.2 and 9.3-5
Amperometric biosensing in voltage-scan (VDCM & DDSM Methods)	§10.3-5
Label-free capacitance detection (CBCM & FTCM Methods)	§7.2-6
Review for final exam	

<sup>\*</sup> textbook : Sandro Carrara, Bio/CMOS interfaces and Co-Design, Springer publisher, New York, 2013

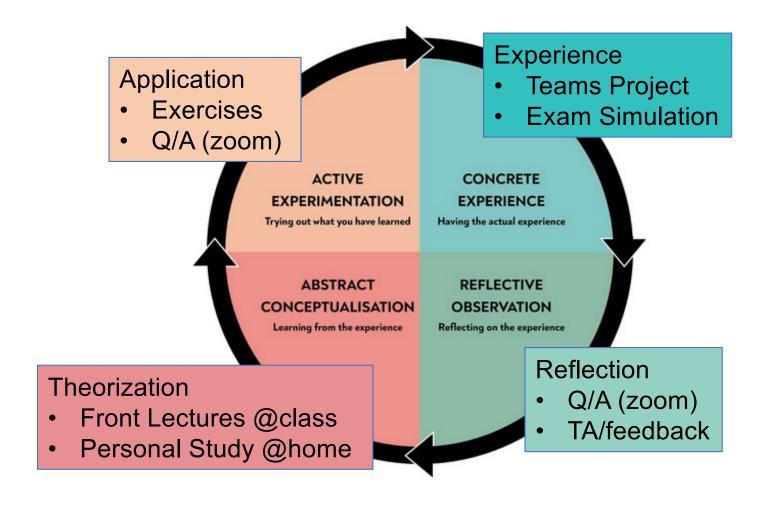


**EE-517:** Bio-Nano-Chip Design





**EE-517:** Bio-Nano-Chip Design



Kolb experimental learning cycle



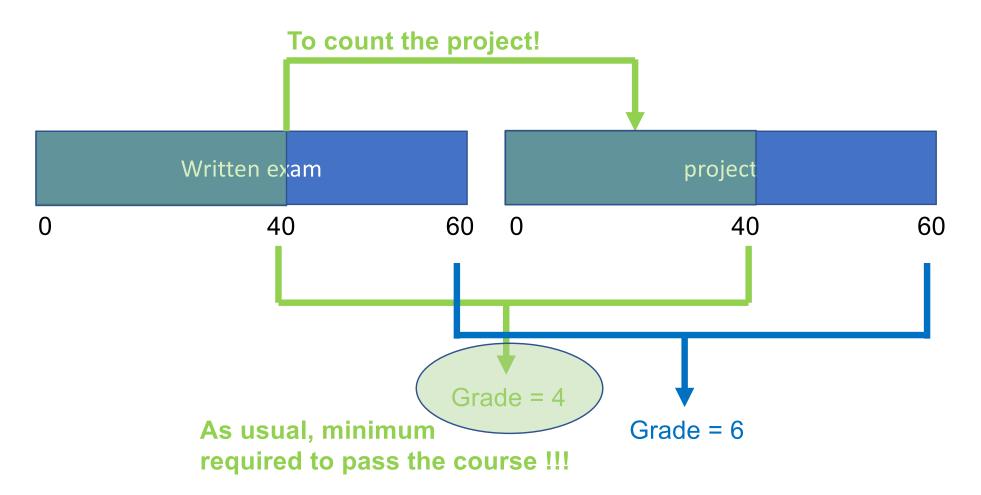
**EE-517:** Bio-Nano-Chip Design

- The project is mandatory.
- Project and written exam count for 60 points each
- Obtaining 40 points out of 60 in the written exam is mandatory for the project to count.
- The final grade is obtained by cumulating points by the written exam and by the project.
- The grade is maximum 6, as usual, obtained by compressing the 120 max points to 6 max grade.

The Exam, the Project, and the Grade



**EE-517:** Bio-Nano-Chip Design



The Exam, the Project, and the Grade



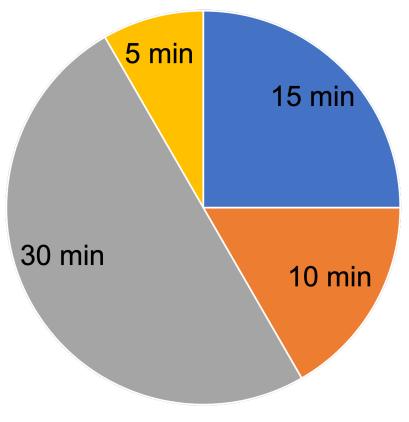
**EE-517:** Bio-Nano-Chip Design

#### **Student Projects - Rules**

- 13. The **Project evaluation** is done based on the following aspects:
  - a. Individual proposal: 15 points (individual mark)
  - b. Progressive report: 15 points (group mark)
  - c. Final report: 15 points (group mark)
  - d. Individual queries during the sessions: 15 points (individual mark)



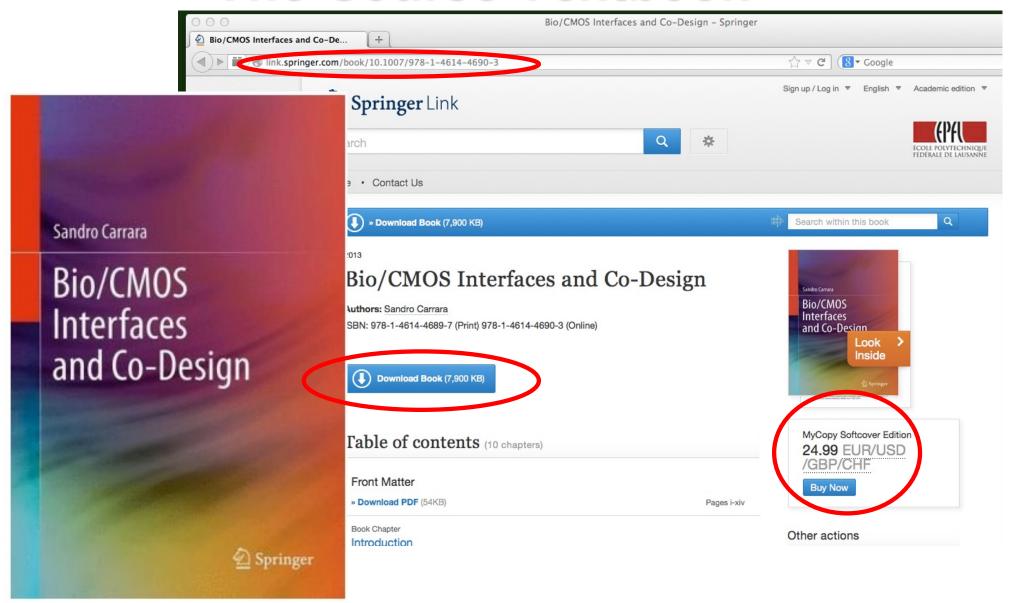
**EE-517:** Bio-Nano-Chip Design



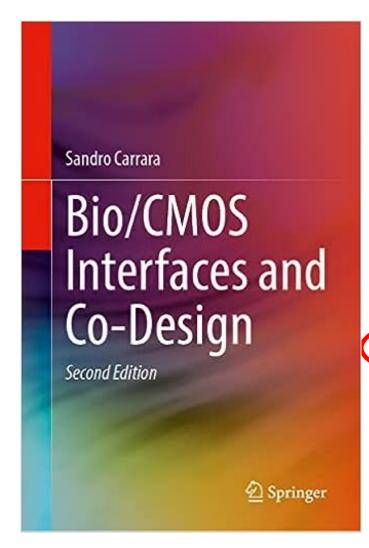
- Coffee/Exercise Q&A TA Intro
- Project Work
  Questions to students

3<sup>rd</sup> hour every week

## **The Course Textbook**

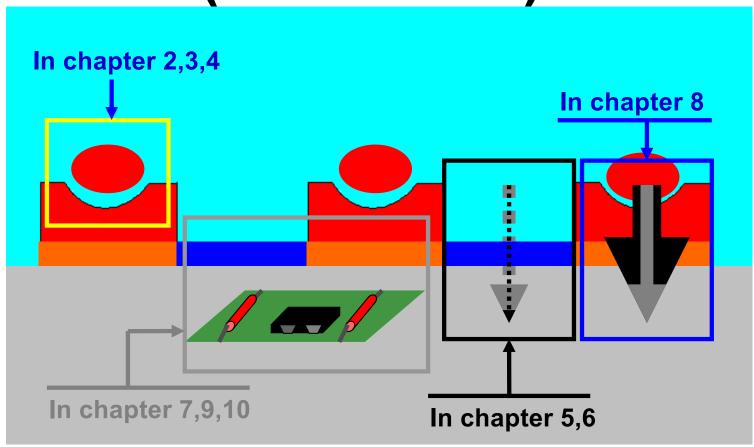


### The Course Textbook: the 2<sup>nd</sup> Edition!



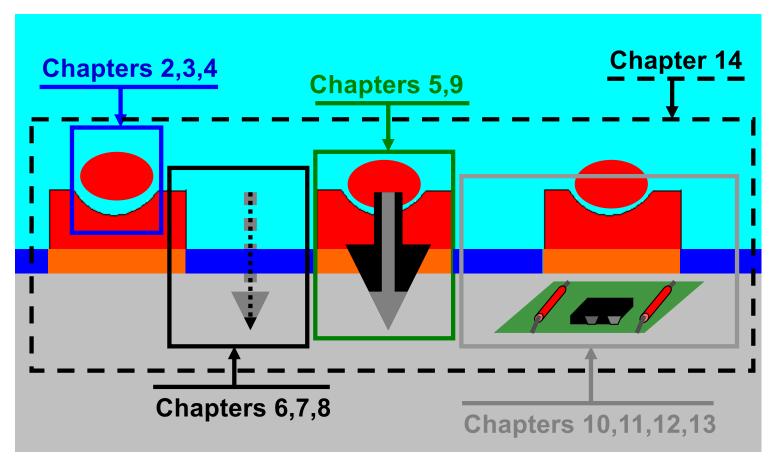
This textbook demonstrates new paradigms for the interface between CMOS circuits and the biological world. A deep theoretical description of such an interface is defined and discussed, while various real applications are demonstrated by also discussing several analog CMOS circuits. Electrochemical techniques are proposed in detail to learn how to design integrated biosensors. Biological materials are described to provide devices selectivity. Nanoscale materials are discussed to provide device sensitivity CMOS circuits are analyzed to provide real applications. Extensive examples with solutions are https://link.springer.com/book/10.1007/978-3-031-31832-0 provided, as well as exercises at the end of each chapter.

# Bio/CMOS interface book (1st Edition)



Introduction to Personal electronics, Distributed Diagnostics, and Bio/CMOS interfaces in Chapter 1

# Bio/CMOS interface book (2<sup>nd</sup> Edition)



Introduction to Personal electronics, Distributed Diagnostics, and Bio/CMOS interfaces in Chapter 1



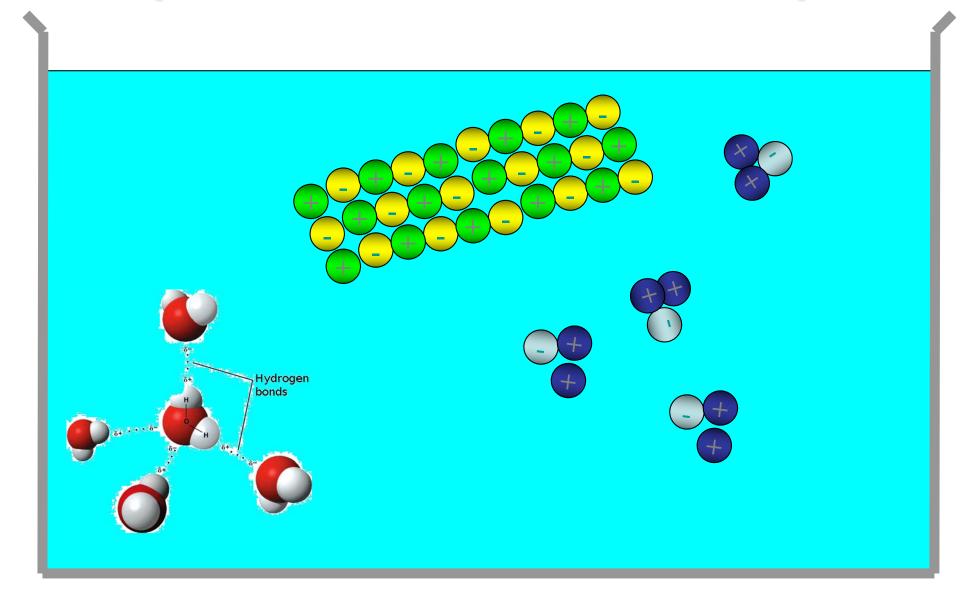
**EE-517:** Bio-Nano-Chip Design

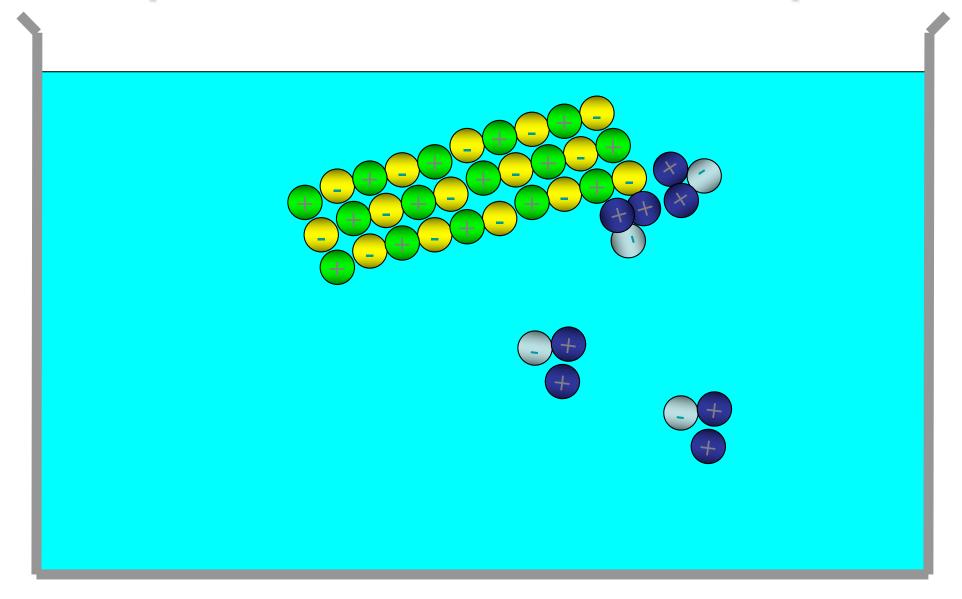
# Lecture #1 Conductive Solutions

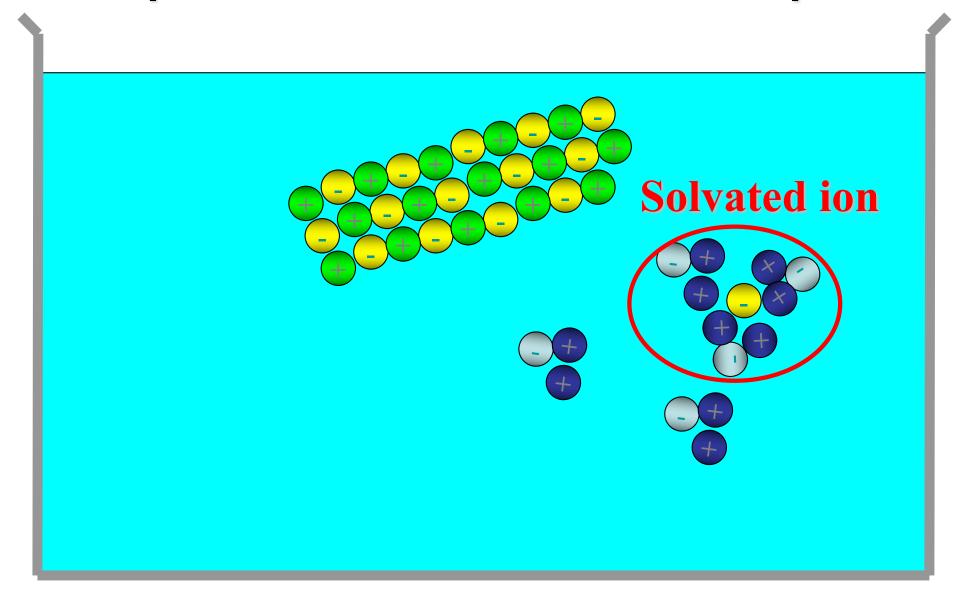
#### **Lecture Outline**

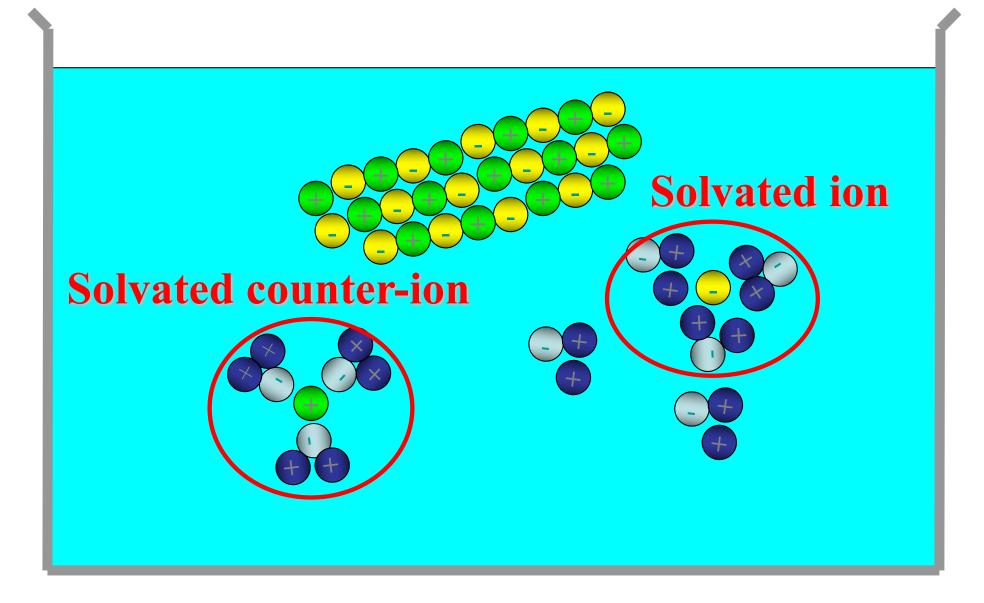
(Book Bio/CMOS: Chapter' paragraphs § 2.1-2.7 and § 2.14-15)

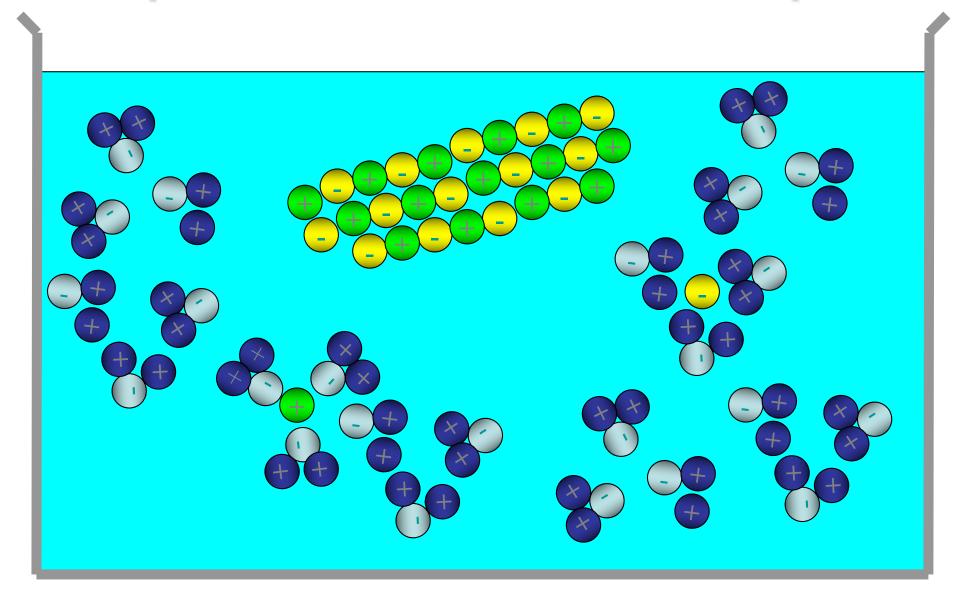
- Solutions of ionic solutes
- Solutions of electrolytes
- Conductive Solutions
- Helmholtz planes
- Redox reactions
- pH

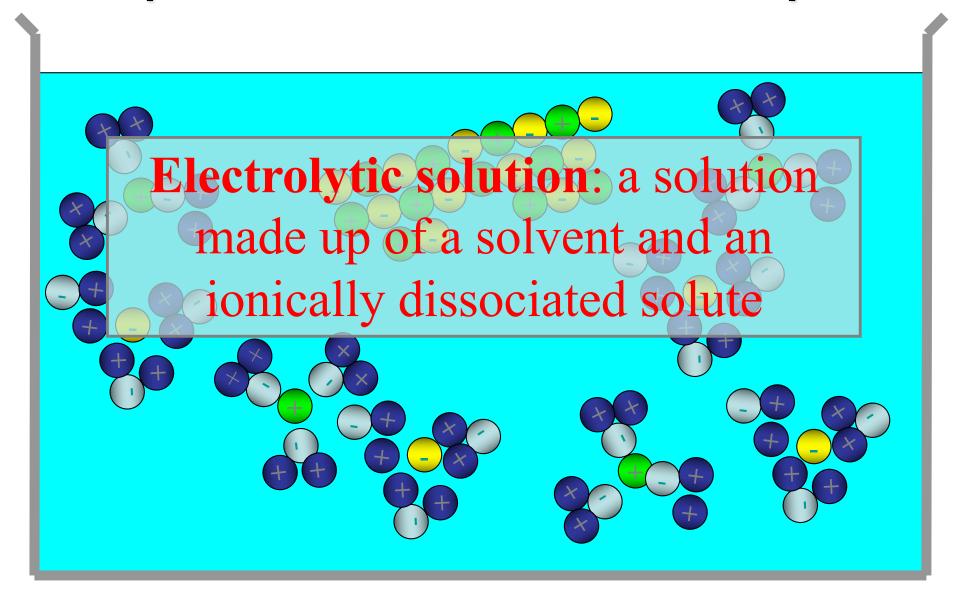


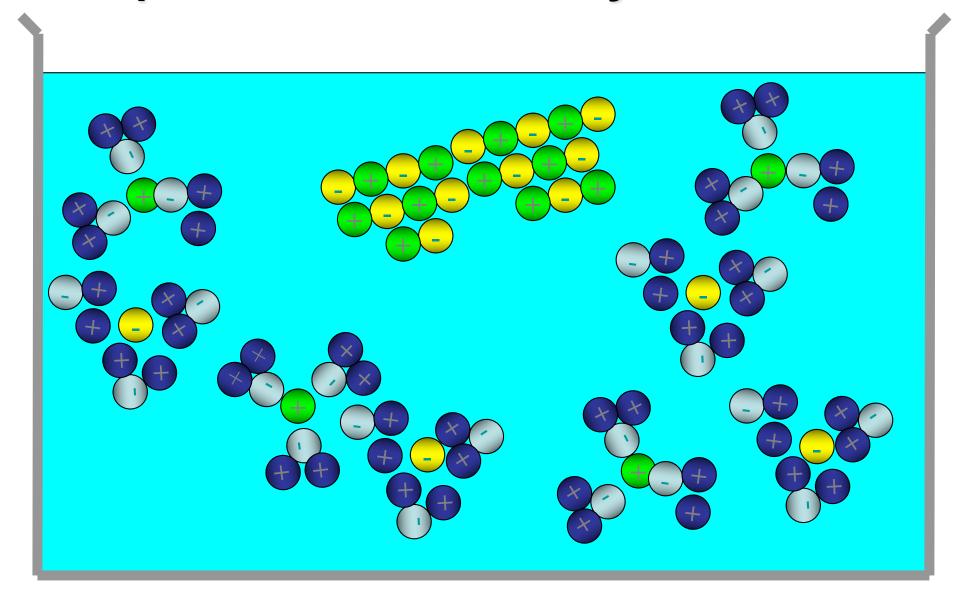


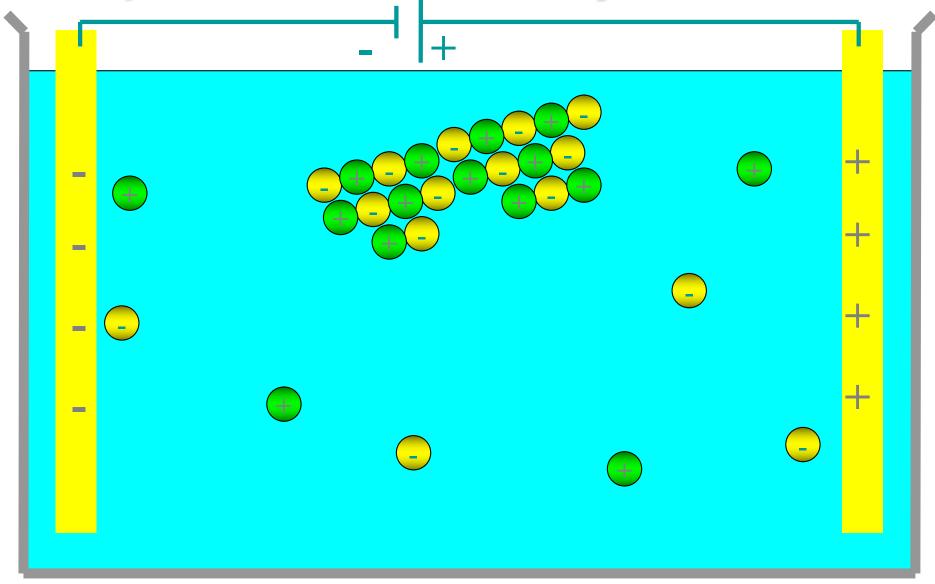


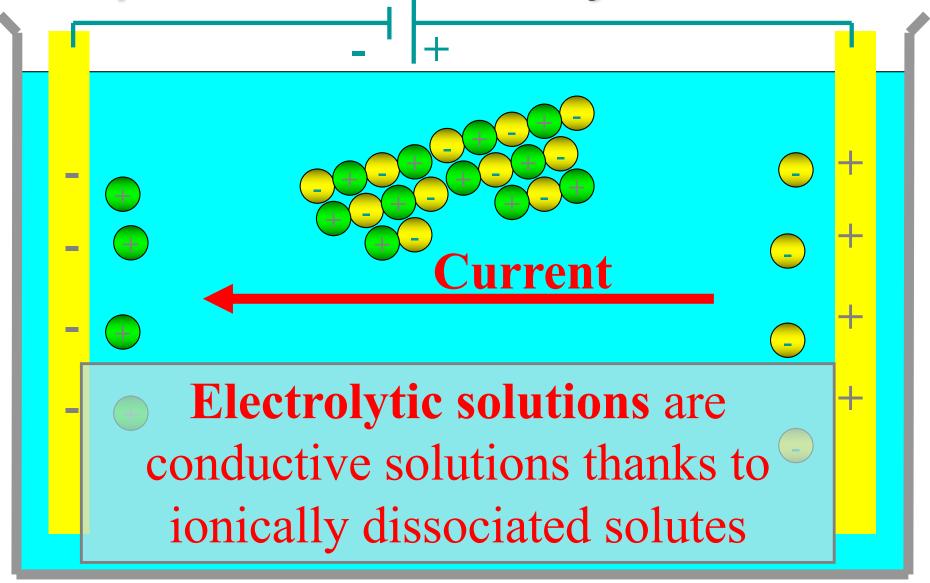












#### **Ionic Solution of HCI**

#### **Dissociation of Chloride Acid**

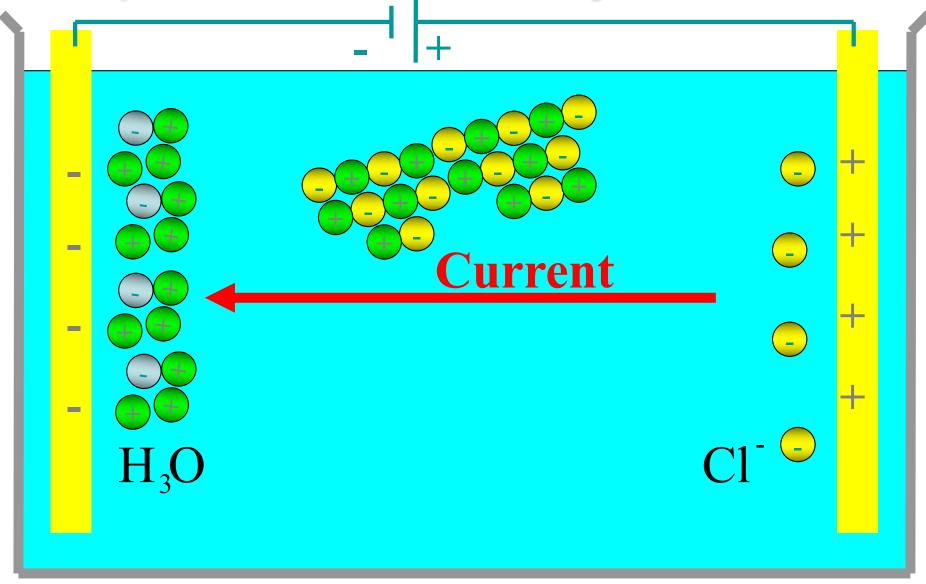
$$2HCl + 2H_2O \rightarrow 2H_3O^+ + 2Cl^-$$

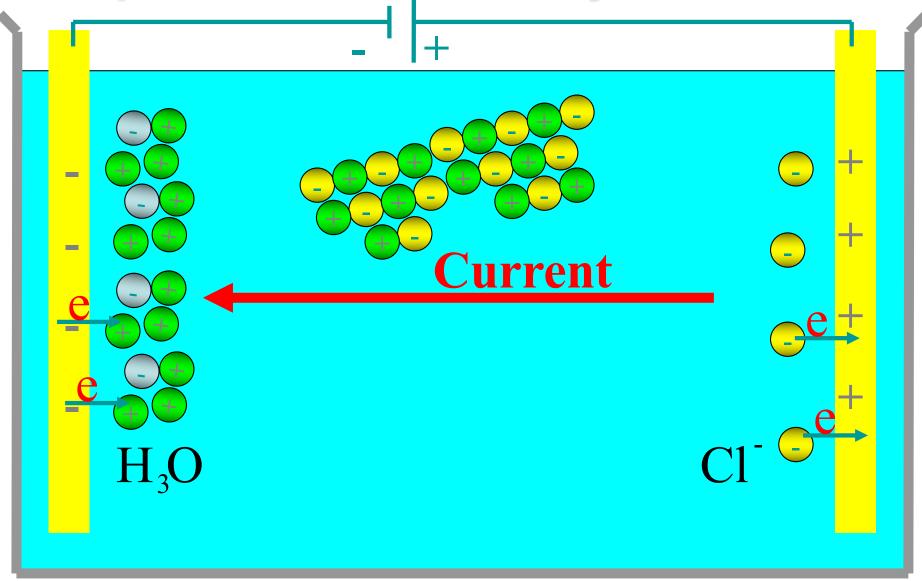
$$2H_3O^+ + 2e^- \rightarrow 2H_2O + H_2 \uparrow$$

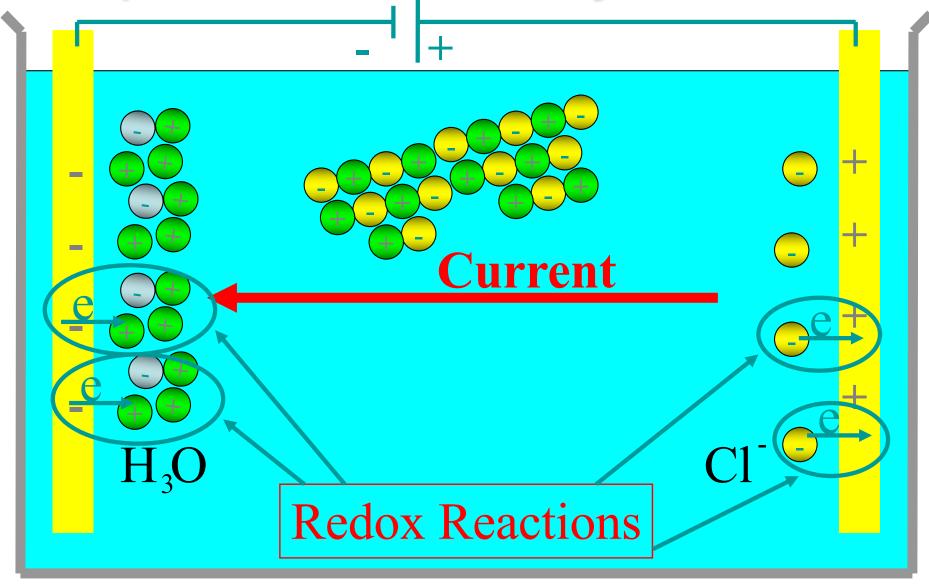
- Electrode (Cathode)

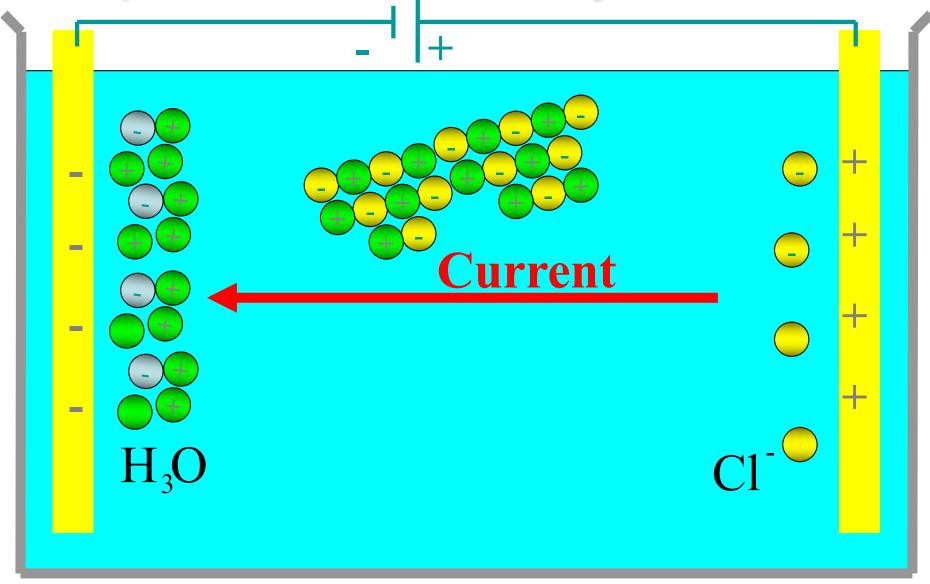
$$2Cl^- \rightarrow Cl_2 \uparrow +2e^-$$

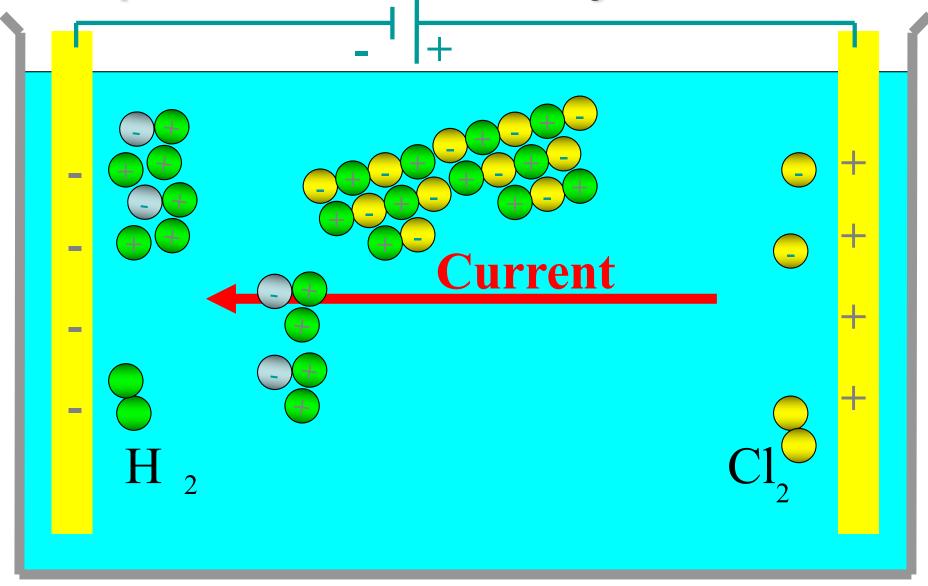
+ Electrode (Anode)

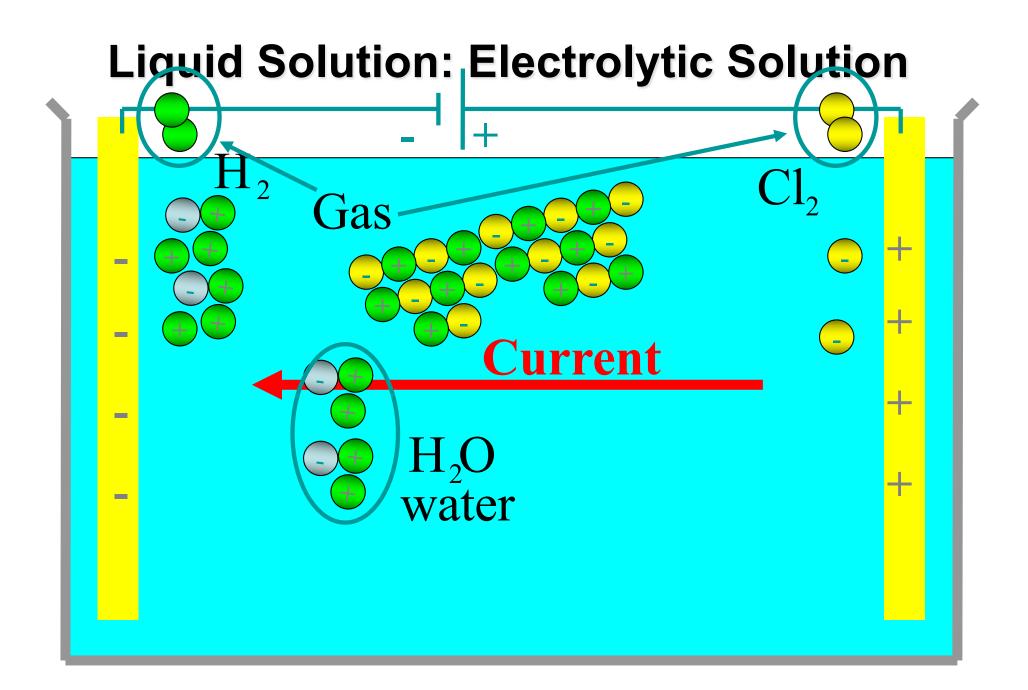


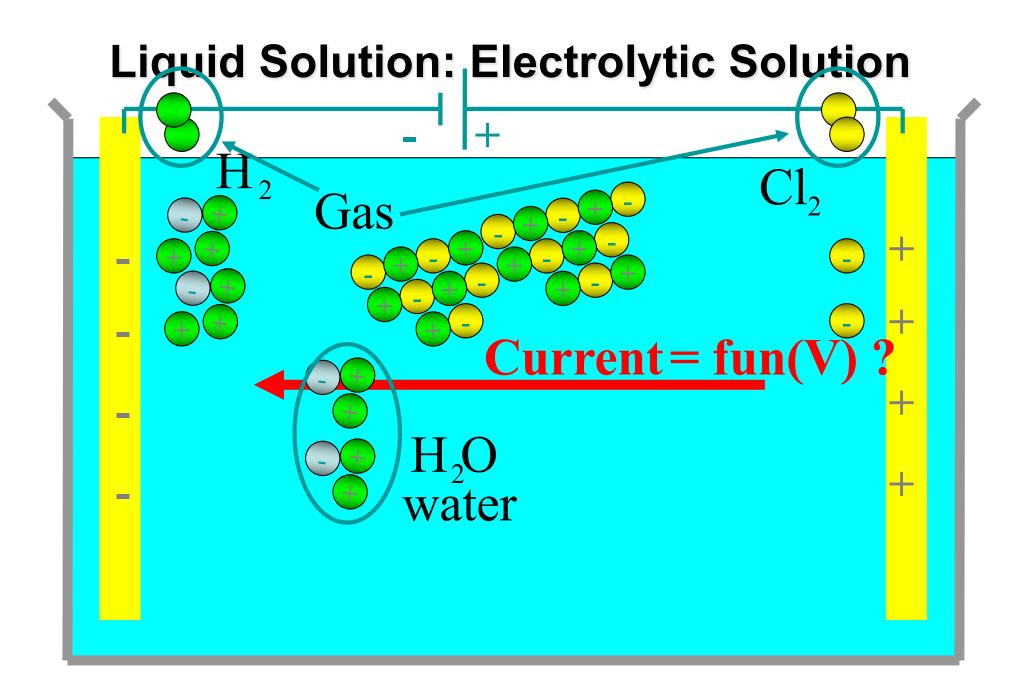


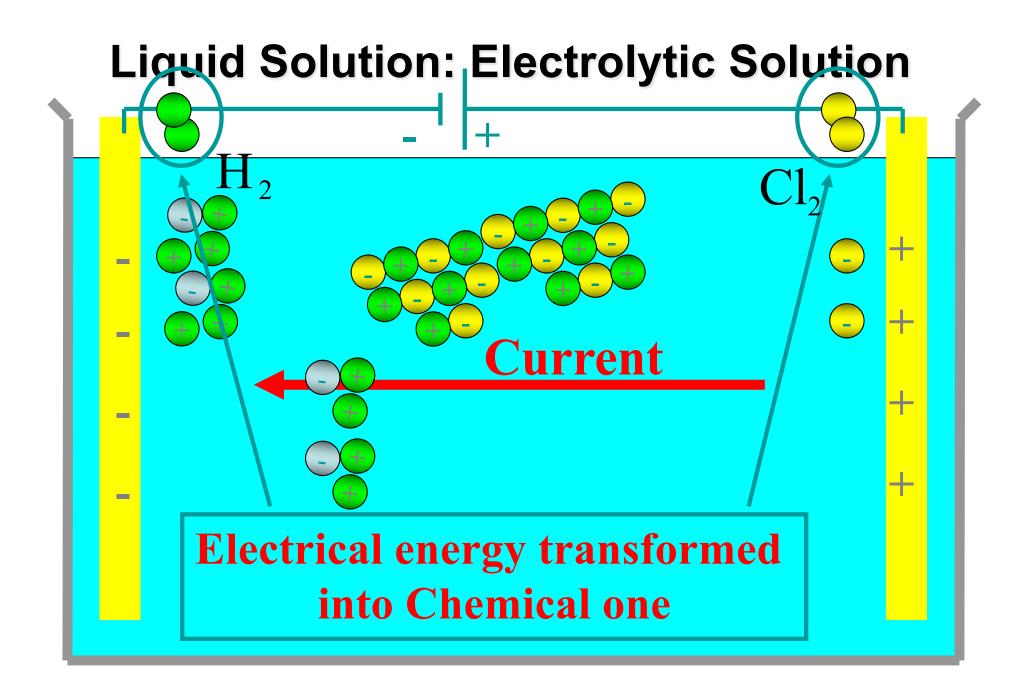


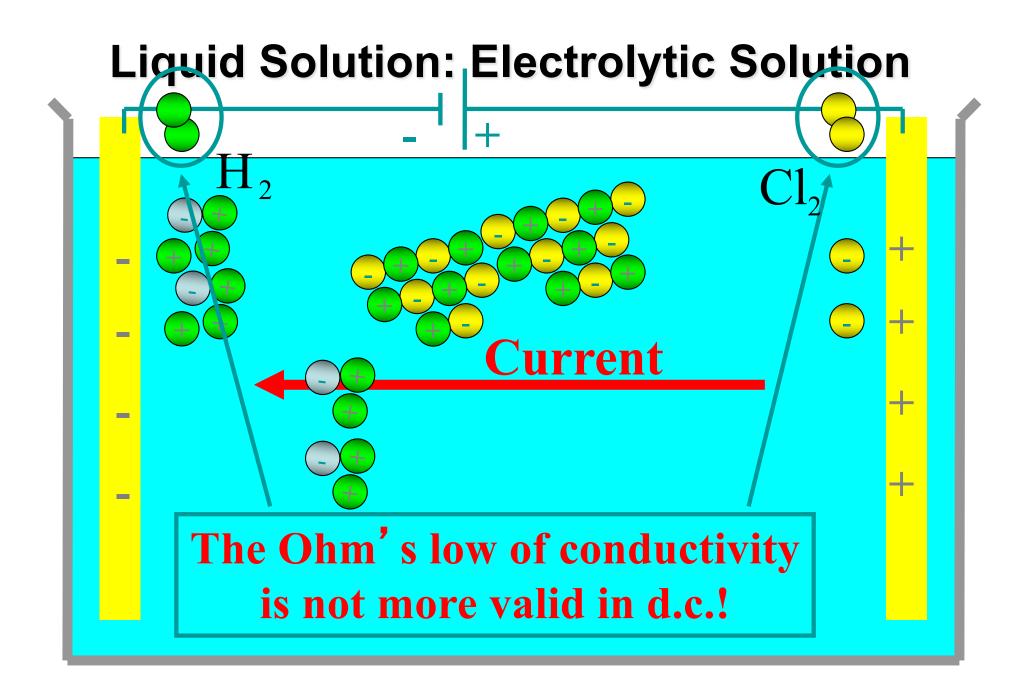


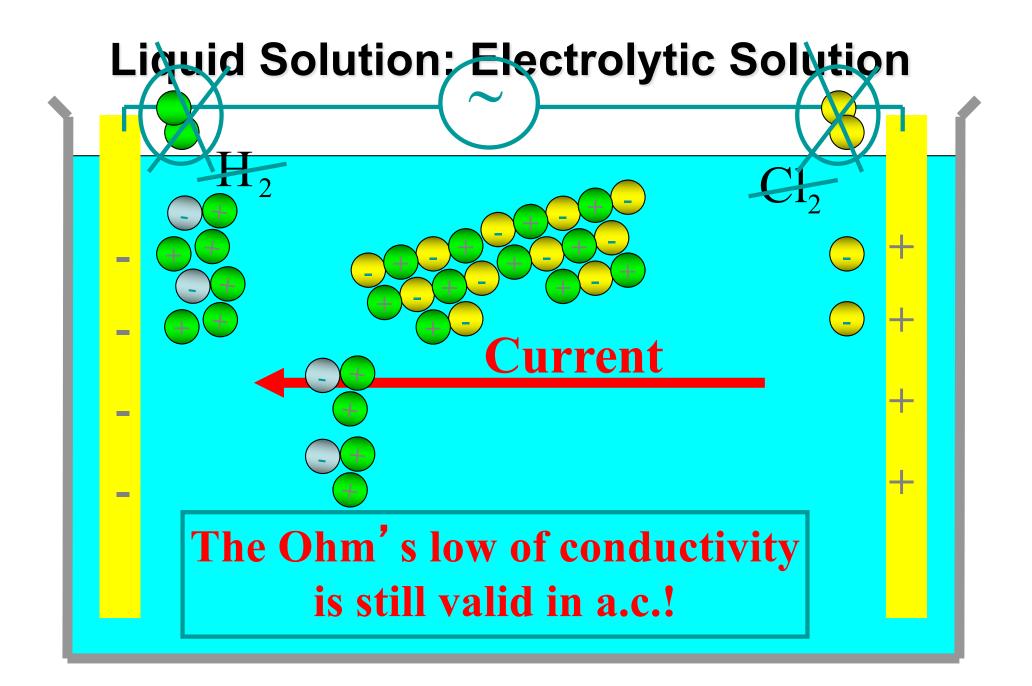


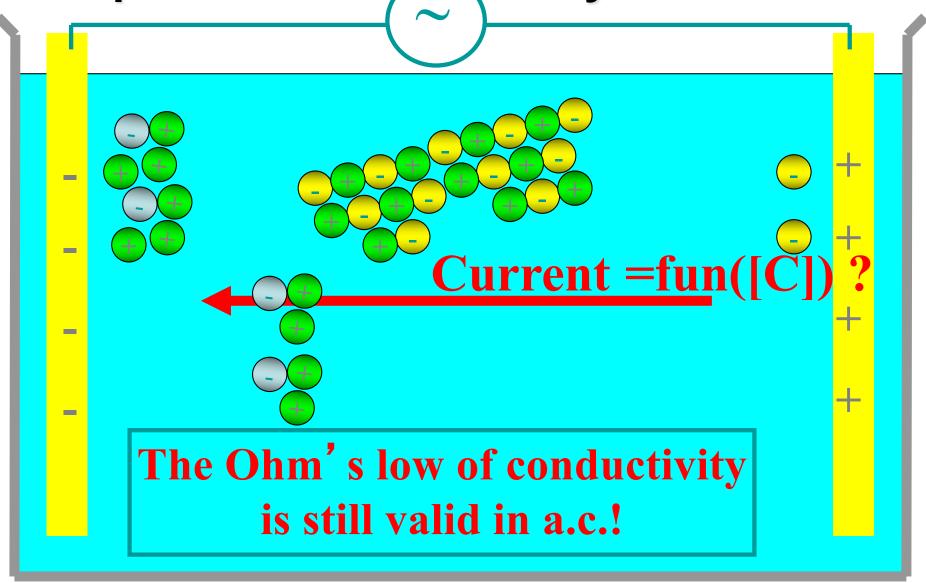






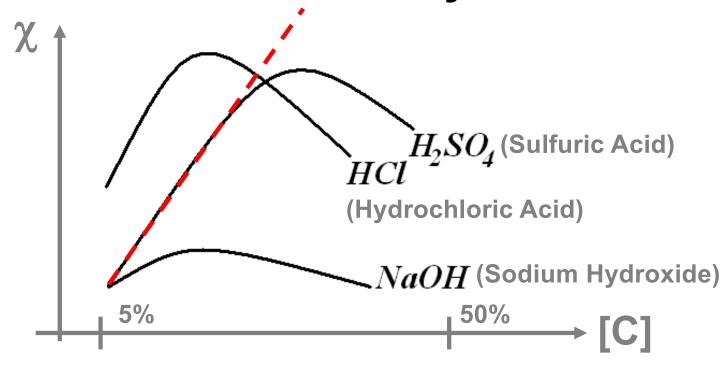




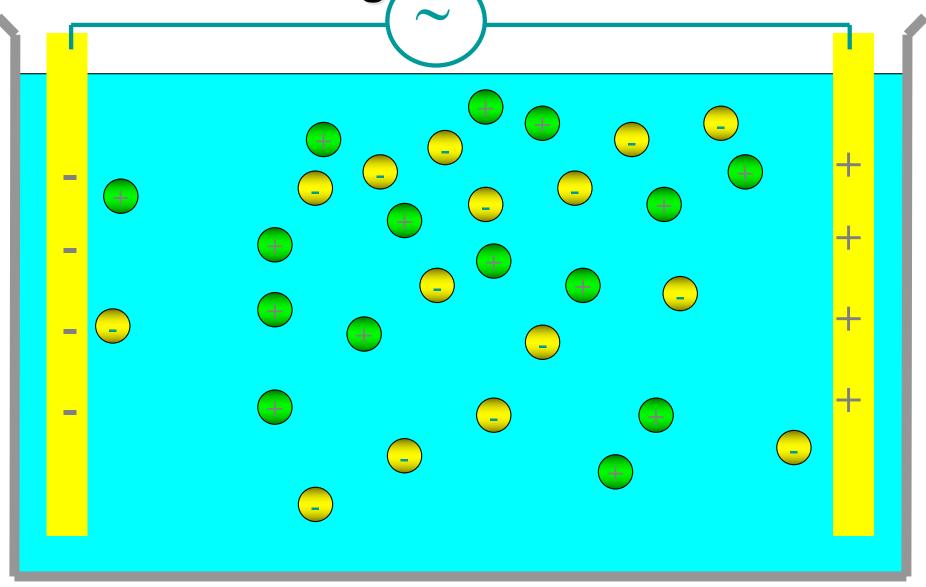


## a.c. Conductivity in Ionic Solutions

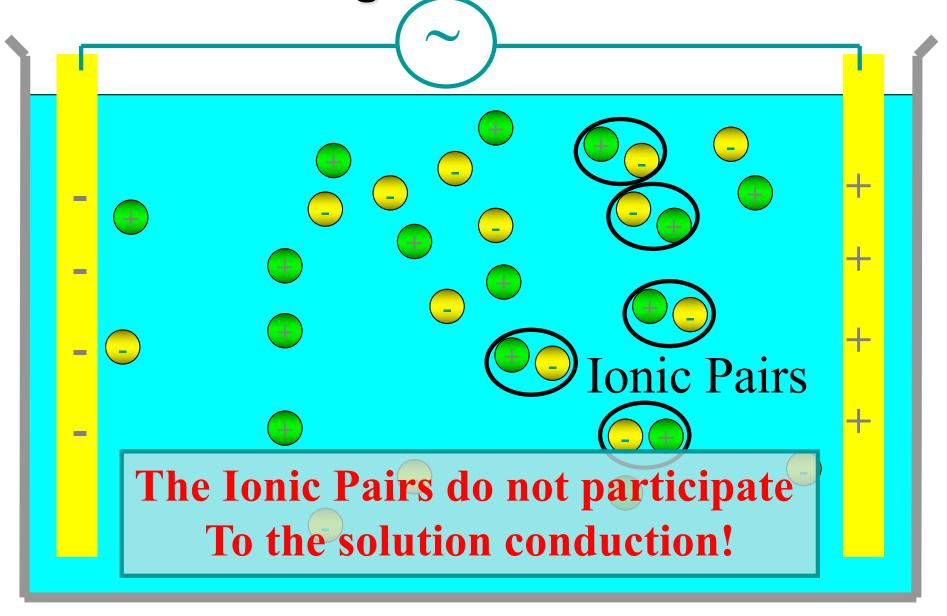
#### **Solution Conductivity**



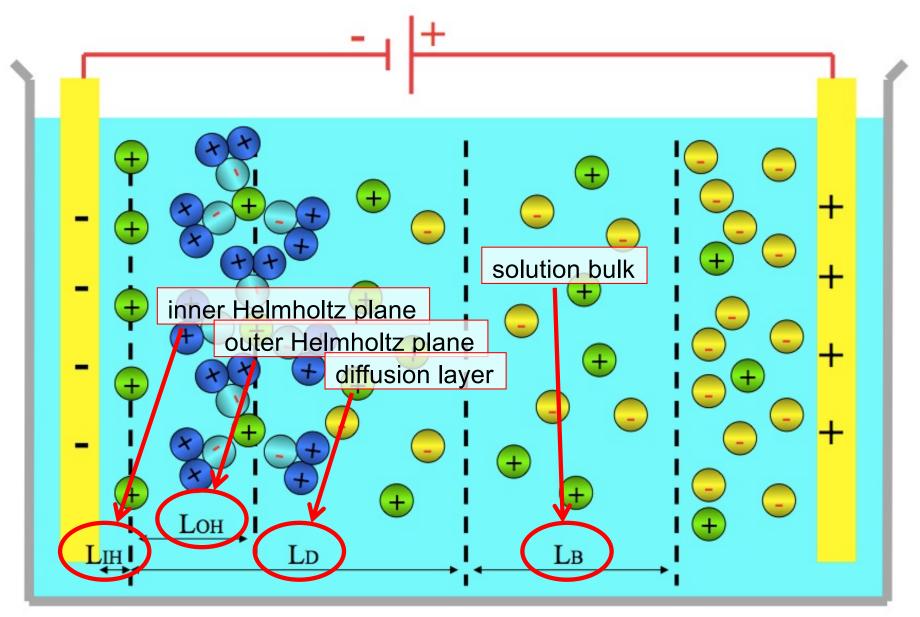
# lons in high concentrations



#### lons in high concentrations



# **Helmholtz Planes**



# Debye Length

Charge density: 
$$\rho_e = \sum_i z_i e n_i$$

 $z_i$  = charge of species i (e.g. +2, -1, etc.)

 $n_i$  = concentration of species *i* (number per volume)

$$abla^2 \phi = 0$$
 In the bulk

$$abla^2 \phi = 0$$
 In the bulk 
$$abla^2 \phi = -\frac{\rho_e}{\kappa \varepsilon_0}$$
 Close to electrodes

For perturbation away from equilibrium at finite temperature

$$\hat{\phi} = \phi - \phi_0 \qquad \rho_e = \sum_i z_i e n_{i0} \exp\left(-\frac{z_i e \hat{\phi}}{k_B T}\right)$$

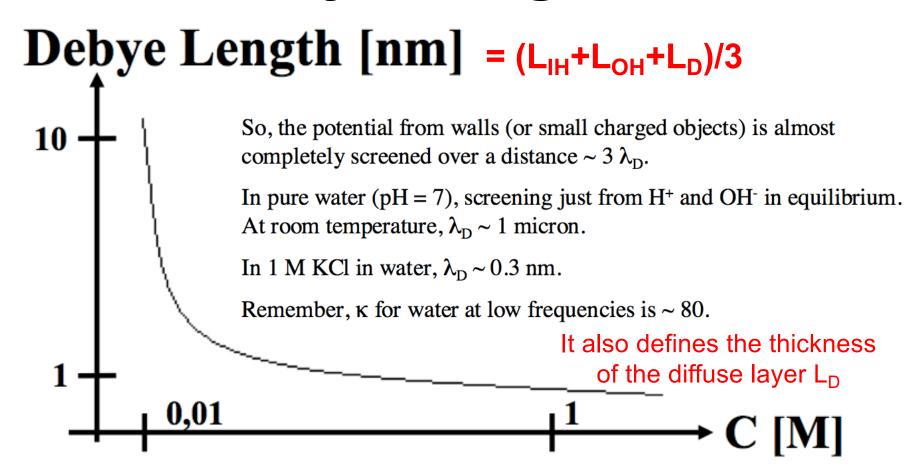
# **Debye Length**

$$\nabla^2 \hat{\phi} = -\frac{1}{\kappa \varepsilon_0} \sum_i z_i e n_{i0} \exp \left( -\frac{z_i e \hat{\phi}}{k_B T} \right) \approx -\frac{1}{\kappa \varepsilon_0} \sum_i z_i e n_{i0} + \frac{e^2}{\kappa \varepsilon_0 k_B T} \sum_i z_i^2 n_{i0} \hat{\phi} \equiv \frac{1}{\lambda_D^2} \hat{\phi}$$

~ 0 for equilibrium neutrality

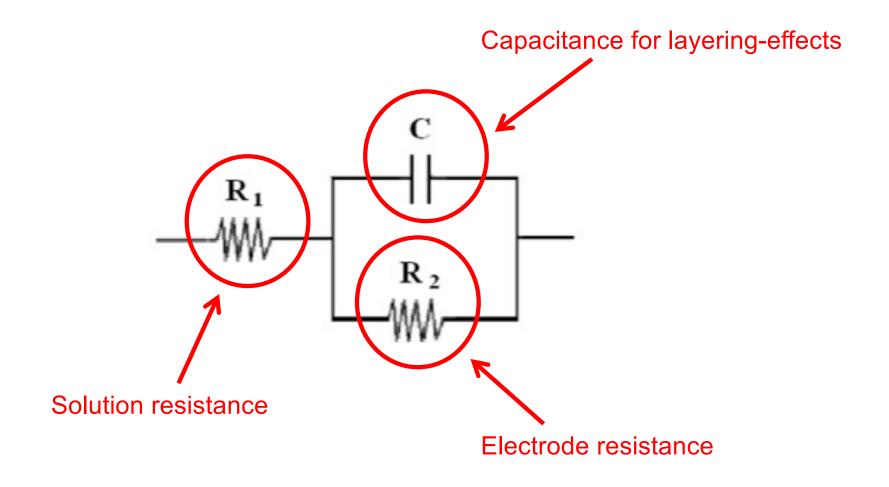
$$\lambda_D = \left(\frac{e^2}{\kappa \varepsilon_0 k_B T} \sum_i z_i^2 n_{i0}\right)^{-1/2}$$

# **Debye Length**



The Bebye Length is defined as the region of charge carrier's net electrostatic effect in solution

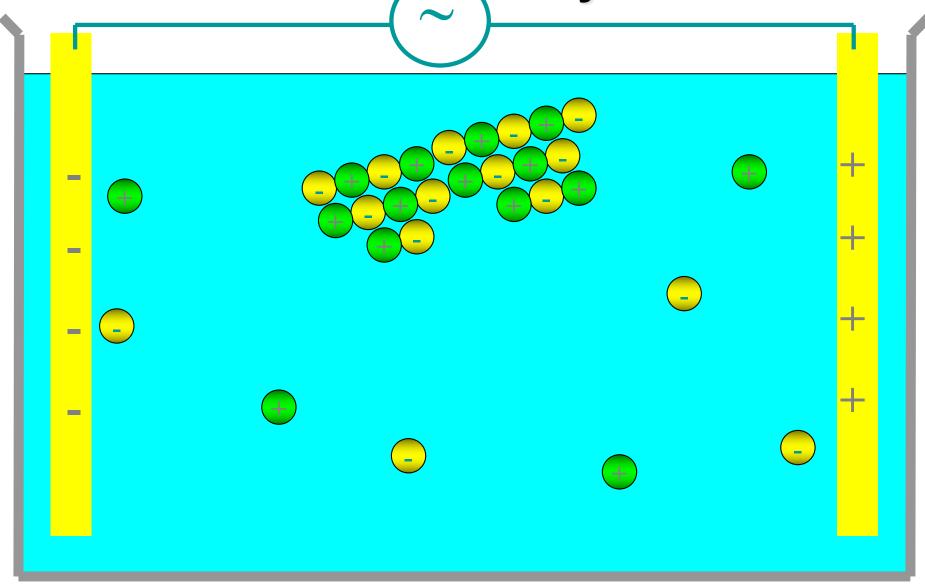
# Equivalent Circuit with Layering effects: Randles Model



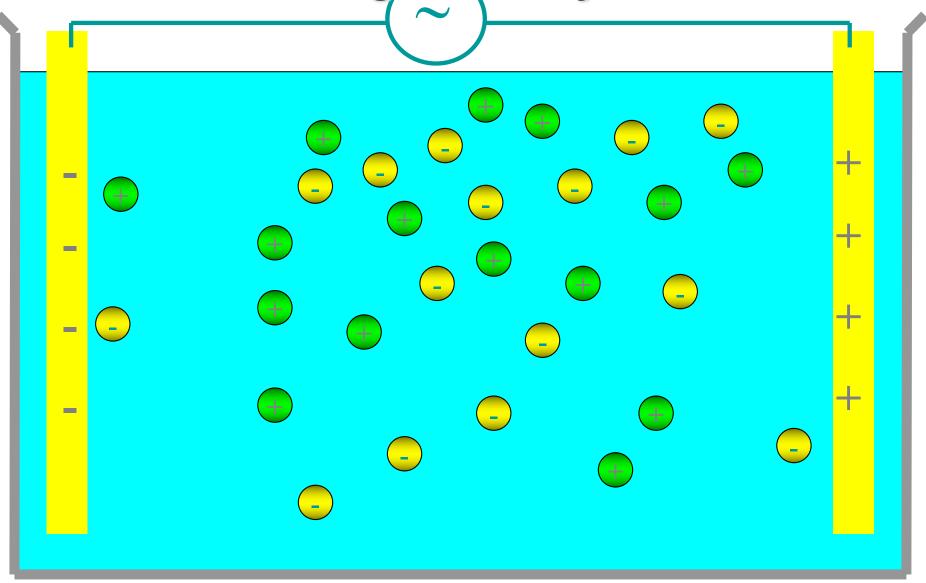
# Kind of Electrolytes

- Strong Electrolytes
- Weak Electrolytes

# Weak Electrolyte



# Strong Electrolytes



# Hard and weak Electrolytes

#### Dissociation of the Sodium Chloride

$$NaCl + H_2O \longrightarrow Na^+ + Cl^- + H_2O$$

Dissociation of Hydrogen Cyanide

$$HCN + H_2O \longleftrightarrow H_3O^+ + CN^-$$

 $\alpha$  = Dissociation Degree

# **Equilibrium Constant**

$$AB + H_2O \stackrel{\alpha}{\longleftrightarrow} A^+ + B^- + H_2O$$

$$K = \frac{[A^+][B^-]}{[AB]} = \frac{\alpha [AB]_0 \alpha [AB]_0}{(1-\alpha)[AB]_0} = \frac{\alpha^2}{1-\alpha} [AB]_0$$

Actually true only for weak electrolytes in small concentration!

# Kind of Electrolytes

Strong Electrolytes

Salts
Strong Acids
Strong Bases

Weak Electrolytes

Weak Acids ? Weak Basis

#### **Acid or Basic Solutions**

#### The Acetic Acid

$$CH_3COOH + 2H_2O \rightarrow CH_3COO^- + H^+ + 2H_2O$$

The Sodium Hydroxide
 $NaOH + H_2O \rightarrow Na^+ + OH^- + H_2O$ 

Base

Arrhenius definition

(Not actually true but enough for our aims)

#### Water is neither a base nor an acid

#### Water dissociation

$$2H_2O \leftrightarrow H_3O^+ + OH^-$$

$$K = \frac{[H_3O^+][OH^-]}{[H_2O]^2}$$

$$[H_2O] = 1 \rightarrow K_w = [H_3O^+][OH^-]$$

Water Ionic product

$$\Delta G^{0} = -RT \ln K_{w} = \Delta G^{0}_{H_{3}O^{+}} + \Delta G^{0}_{OH^{-}} - 2\Delta G^{0}_{H_{2}O}$$

# **Definition of pH**

#### Water dissociation

$$\Delta G^{0} = -RT \ln K_{w} = \Delta G^{0}_{H_{3}O^{+}} + \Delta G^{0}_{OH^{-}} - 2\Delta G^{0}_{H_{2}O}$$

$$\ln K_{w} = -\frac{\Delta G^{0}_{H_{3}O^{+}} + \Delta G^{0}_{OH^{-}} - 2\Delta G^{0}_{H_{2}O}}{RT}$$

$$\ln K_{w} = -14$$

$$K_{w} = [H_{3}O^{+}][OH^{-}] = 10^{-14}$$

### pH=7, means "neutral pH"

 $[H_3O^+] = [OH^-] = 10^{-7}M$ 

### Acid, Neutral, or Basic Solutions

$$[H_{3}O^{+}] > 10^{-7}M \rightarrow Acid Solution$$

$$[H_{3}O^{+}] = 10^{-7}M \rightarrow Neutral Solution$$

$$[H_{3}O^{+}] < 10^{-7}M \rightarrow Basic Solution$$

$$pH = Log\left(\frac{1}{[H_{3}O^{+}]}\right); \begin{cases} <7 \rightarrow Acid Solution \\ 7 \rightarrow Neutral Solution \\ >7 \rightarrow Basic Solution \end{cases}$$