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PhotoElectroChemistry for Corrosion

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Introduction

- ▶ Photoelectrochemical techniques have been shown to be useful tools for characterizing oxidation layers.
- ▶ Interdisciplinary theoretical underpinnings were built [1–5] such as the Gärtner-Butler model [6, 7] which has been proven to be a simple and robust model for the photocurrent generation.
- ▶ Technical progresses were achieved, allowing to study oxide layers at macroscopic, mesoscopic, and microscopic scales [8, 9], or in-situ in high temperature corrosion conditions [10, 11].

Hypotheses

Several hypotheses are needed in order to apply the theoretical concepts:

▶ semiconductors are considered to be ideal i.e. crystallized and homogeneous

▶ the dielectric constant of the semiconductor is independent of the light wavelength

▶ the capacity of the Helmholtz layer is greater than the capacitance of the space charge capacitance

▶ the potential drop in the Helmholtz layer is independent of the applied potential and is negligible

Warning

The hypotheses are rarely fully respected in the case of oxides or passive films formed on industrial alloys. Nonetheless, the literature shows that the developed models can be applied to non-ideal systems such as oxides and passive films.

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Electronic Band Structure

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Band Model I

- Solids: conductors, semiconductors and insulators.
- Valence and conduction bands correspond to allowed energy states for the electrons.
- E_c is the lowest energy level of the conduction band.
- E_v is the highest energy level of the valence band.
- E_g is the band gap with no allowed energy states.
- E_F is the Fermi Level which describes the distribution of the electrons among both bands.

Fermi Level
The Fermi Level represents the highest energy state that can be occupied level at 0K.
It is equivalent to the electrochemical potential in solid phases.

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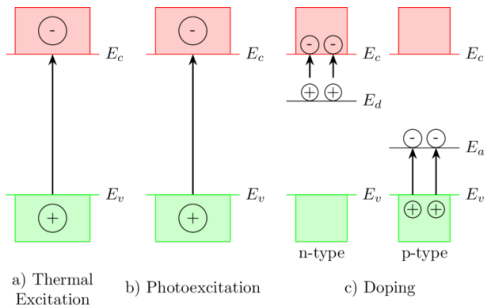
Band Model II

- ▶ The electronic conduction $=$ movement of electrons and/or holes in conduction/valence band.
- ▶ The conduction depends on the number of available charge carriers in the conduction band and in the valence band.
- ▶ In conductors: overlap of the conduction and the valence bands occurs.
- ▶ In semiconductor and insulator: the conduction depends on the band gap and the energy provided by the environment to the electrons from the valence band in order to jump into the conduction band.

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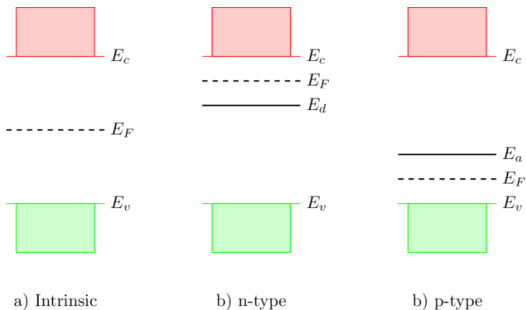
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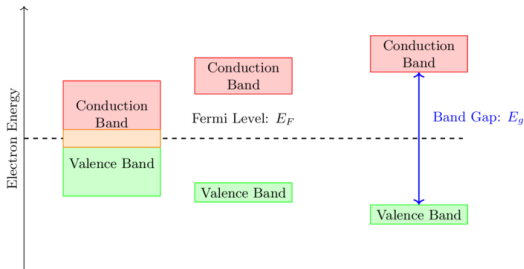
Excitation carrier I

In semiconductors, charge carriers can be generated by three mechanisms:

- ▶ thermal excitation: in the case of very low band gaps, it can be enough in order to eject an electron from E_v to E_c.
- ▶ photoexcitation: ejects electrons from E_v to E_c band when an incident photon ($h\nu > 5eV$) is absorbed.
- ▶ doping: introduces additional energy level located in between E_v to E_c.

a) Thermal Excitation   b) Photoexcitation   c) Doping

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Fermi Level Position

The Fermi level E_F in intrinsic semiconductors is located at the mid-gap. The n-type and p-type doping shift the Fermi level towards band edges E_c and E_v.

a) Intrinsic   b) n-type   b) p-type

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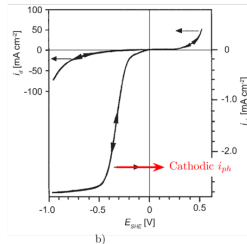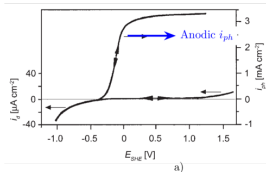
Semiconductor/electrolyte interface under illumination

Electron/hole pairs I

▶ The illumination of the semiconductor/electrolyte interface, with
photons having an energy greater than the band gap, E_g, creates
electron/hole pairs in the semiconductor.

▶ By applying the adequate potential the pairs can be separated.

▶ As a consequence, the majority charge carriers are attracted to the
semiconductor bulk whereas the minority charge carriers are drawn
to the semiconductor/electrolyte interface where they can be
transferred to a RedOx species creating an additional current called
photocurrent.

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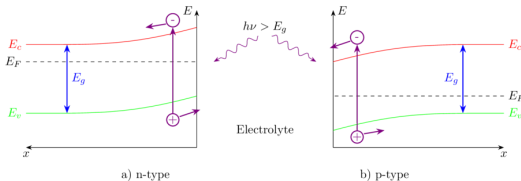
Electron/hole pairs II

- ▶ The photocurrent is significant when the semiconductor/electrolyte junction is in depletion.

- ▶ n-type (p-type) semiconductors generate anodic (cathodic) photocurrents where the electrons (holes) move towards the external circuit whereas the holes (electrons) move towards the interface.

- ▶ The applied potential on n-type (p-type) semiconductors is greater (lower) than the flat band potential.

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Semiconductor/electrolyte interface under illumination

Gärtner-Butler Model I

▶ Gärtner [6] and Butler [7] proposed a simple and robust model for describing the photocurrent (no recombination of the photogenerated electron/hole pairs in the space charge).

▶ The photocurrent depends on
  ▶ w_{sc}: space charge width
  ▶ α: absorption coefficient
  ▶ L_{cc}: the average diffusion length of the minority charge carriers

$$I_{ph} = \phi_0 \left[1 - \frac{\exp(-\alpha_{sc} \cdot w_{sc})}{1 + \alpha_{sc} \cdot L_{cc}} \right] \tag{1}$$

When $\alpha_{sc} \cdot w_{sc} \ll 1$ and $\alpha \cdot L_{cc} \ll 1$, the photocurrent is approximated by the equation 2.

$$I_{ph} = \phi_0 \cdot \alpha \cdot w_{sc} \tag{2}$$

Warning

All absorbed photons generate electron/hole pairs and the minority charge carriers are transferred to the electrolyte and therefore contribute to the photocurrent.

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Gärtner-Butler Model II

Space charge width w_{sc} in depletion depends on:

▶ N_{cc}: number of majority charge carriers ($\sim$ doping)

▶ e: elementary charge of an electron

▶ U: applied potentials

▶ U_{fb}: flat band potential,

▶ ϵ: relative permittivity

▶ ϵ_0: vacuum permittivity

$$w_{sc} = \sqrt{\frac{2\epsilon\epsilon_0}{eN_{cc}}(U - U_{fb} - \frac{kT}{e})} \tag{3}$$

The absorption coefficient α_{sc} depends on:

▶ $h\nu$: light energy $h\nu$

▶ n: band-band transition type where n takes discreet values of 0.5 or 2 when direct or indirect transitions are allowed.

$$\alpha_{sc} = const\frac{(h\nu - E_g)^n}{h\nu} \tag{4}$$

Gärtner-Butler Model III

The complete expression of the photocurrent is therefore given by the equation 5. The latter is obtained by substituting the absorption coefficient α_{sc} and the space charge width w_{sc} from the equation 2 by the equations 3 and 4, respectively.

$$I_{ph} = \phi_0 \cdot const \frac{(h\nu - E_g)^n}{h\nu} \cdot \sqrt{\frac{2\epsilon\epsilon_0}{eN_{cc}}(U - U_{fb} - \frac{kT}{e})} \tag{5}$$

The linear transform with respect to the energy is used for determining the band gaps.

$$\left[\frac{I_{ph} \cdot h\nu}{\phi_0}\right]^{1/n} = const \cdot (h\nu - E_g) \tag{6}$$

The linear transform with respect to the potential is used for determining the semiconducting type, the flat band potential, and the number of majority charge carrier.

$$I_{ph}^2 = const \cdot (U - U_{fb} - \frac{kT}{e}) \tag{7}$$

Identification of minor oxides I

► Benaboud et al. [8] showed that the photoelectrochemical characterization is robust for detecting the presence of minor oxides.

► The strong photocurrent observed at around 5 eV reveals the major oxide i.e. monoclinic zirconia.

► The photocurrent $h\nu < 5eV$ reveals the presence of minor oxides even in "pure" zirconium.

► The slope changes provided an estimation of the band gaps: hematite, chromia and a solid solution of $(Fe_x Cr_{1-x})O_3$.
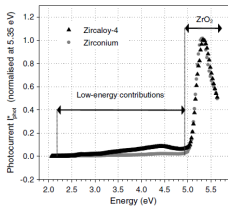

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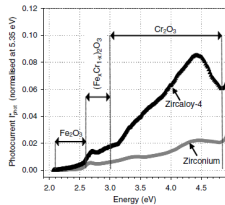
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Semiconduction type I

- ▶ Loucif et al. [12] showed the effect of hydrogen pressure on the semiconduction type on Ni-based alloy 600 oxidized in simulated PWR.

- ▶ The "V-shape" of the normalized photocurrent reveals an isolating behavior of the oxide layer at high hydrogen pressure.

- ▶ The monotonous increase of the normalized photocurrent towards more anodic potentials reveals n-type semiconduction.

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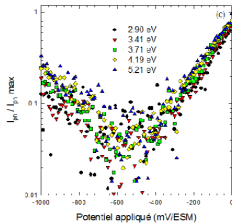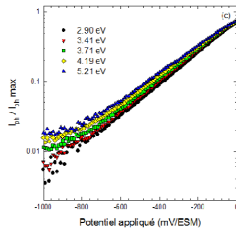
High temperature PEC I

▶ The majority of photoelectrochemical characterizations are performed at room temperature in simple glass/Plexiglas cells where the signal/noise ratio is very good.

▶ High temperature photoelectrochemical characterizations require sophisticated metallic cells and transparent windows able to withstand the arch environment.

▶ Despite the need to improve the signal/noise ratio, the feasibility of the in-situ photoelectrochemical characterizations was demonstrated by Bojinov et al. [10] in 2002 and more recently by Skocic [11] in 2015
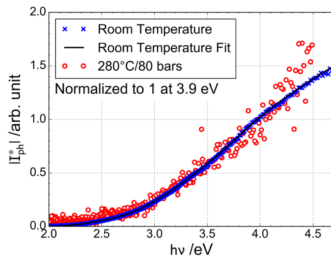
High temperature PEC II

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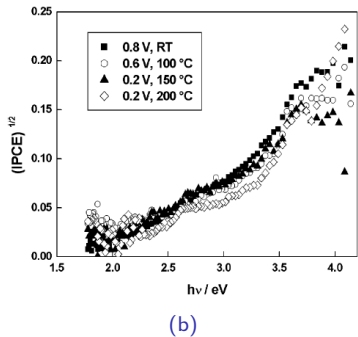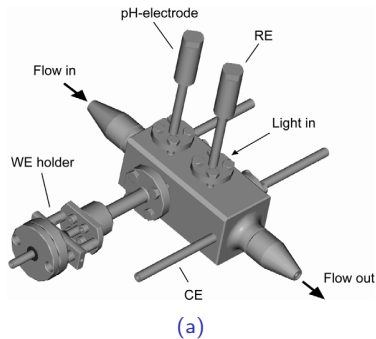
High temperature PEC III
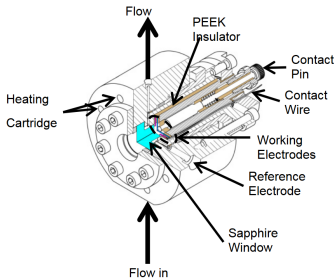
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