

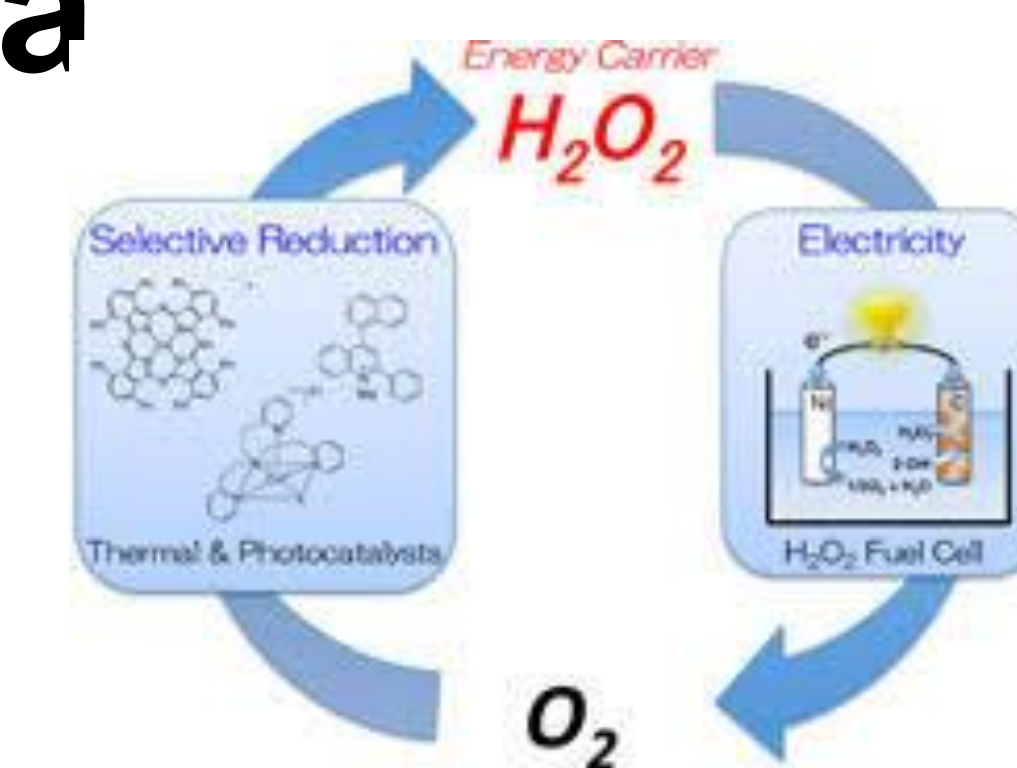


Voltammetric Sensors for Hydrogen Peroxide Detection in Living Cells

Elena Meckaroska, Isidora Ristovska, Milkica Janeva, **Pavlinka Kokoskarova**

Faculty of Medical Sciences, Goce Delcev University, Stip, Macedonia

e-mail: kpavlinka@yahoo.com



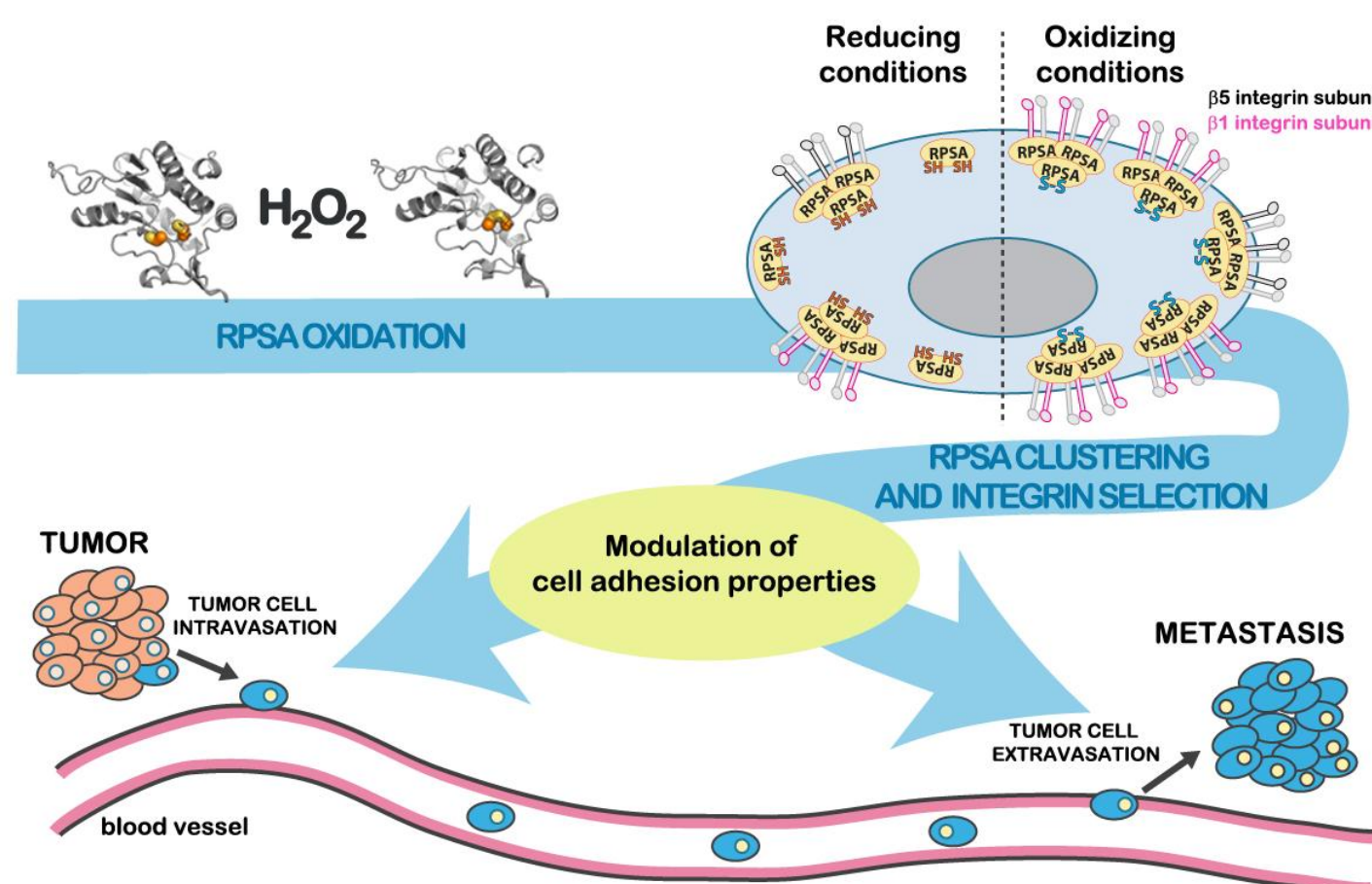
1 Background

Hydrogen Peroxide (H₂O₂) is one of the most important small molecules involved in various signalling processes at all living systems. Hydrogen peroxide is commonly created as a by-product in the respiration processes. There is significant amount of scientific information showing the involvement of H₂O₂ in signalling of stress responses, and in many other redox-signalling related processes. In most of the mechanisms featuring redox-signalling in the cells, it is confirmed that thiol-containing molecules play important role, usually undergoing a redox transformation in reactions with H₂O₂. The redox chemistry of H₂O₂ is quite complex, since it can be involved in variety of reactions. Depending on the pH, H₂O₂ can be seen as a substance with highly oxidative potential, but also as a compound with reductive properties. The two most simple 2-electron scenarios of oxidation and reduction of H₂O₂ can be described by following reactions:

$$\text{H}_2\text{O}_2 + 2\text{H}^+ + 2\text{e}^- \rightarrow 2\text{H}_2\text{O} \quad E^\circ = +1.534 \text{ V vs. SCE (pH of 7.00)}$$

$$2\text{H}^+ + \text{O}_2 + 2\text{e}^- \rightarrow \text{H}_2\text{O}_2 \quad E^\circ = +0.440 \text{ V vs. SCE (pH of 7.00)}$$

In this work, we show several ways of designing voltammetric sensors for hydrogen peroxide quantification. These methods can be designed for direct detection and quantification of hydrogen peroxide, but more efficient are the approaches in the so-called Electrocatalytic-Regenerative (EC') electrode mechanism. With the methods presented here, detection of hydrogen peroxide in micro-to-milimolar range is possible.



5 Conclusions

We present several scenarios under conditions of cyclic voltammetry for quantification of hydrogen peroxide in living cells. The methods can be direct, but commonly a so-called „redox enzyme“ mediators are used for getting sensitive quantification of H₂O₂.

References

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2 Results-I

First sensor is based on direct Quantification (reduction of H₂O₂) at metallic electrodes

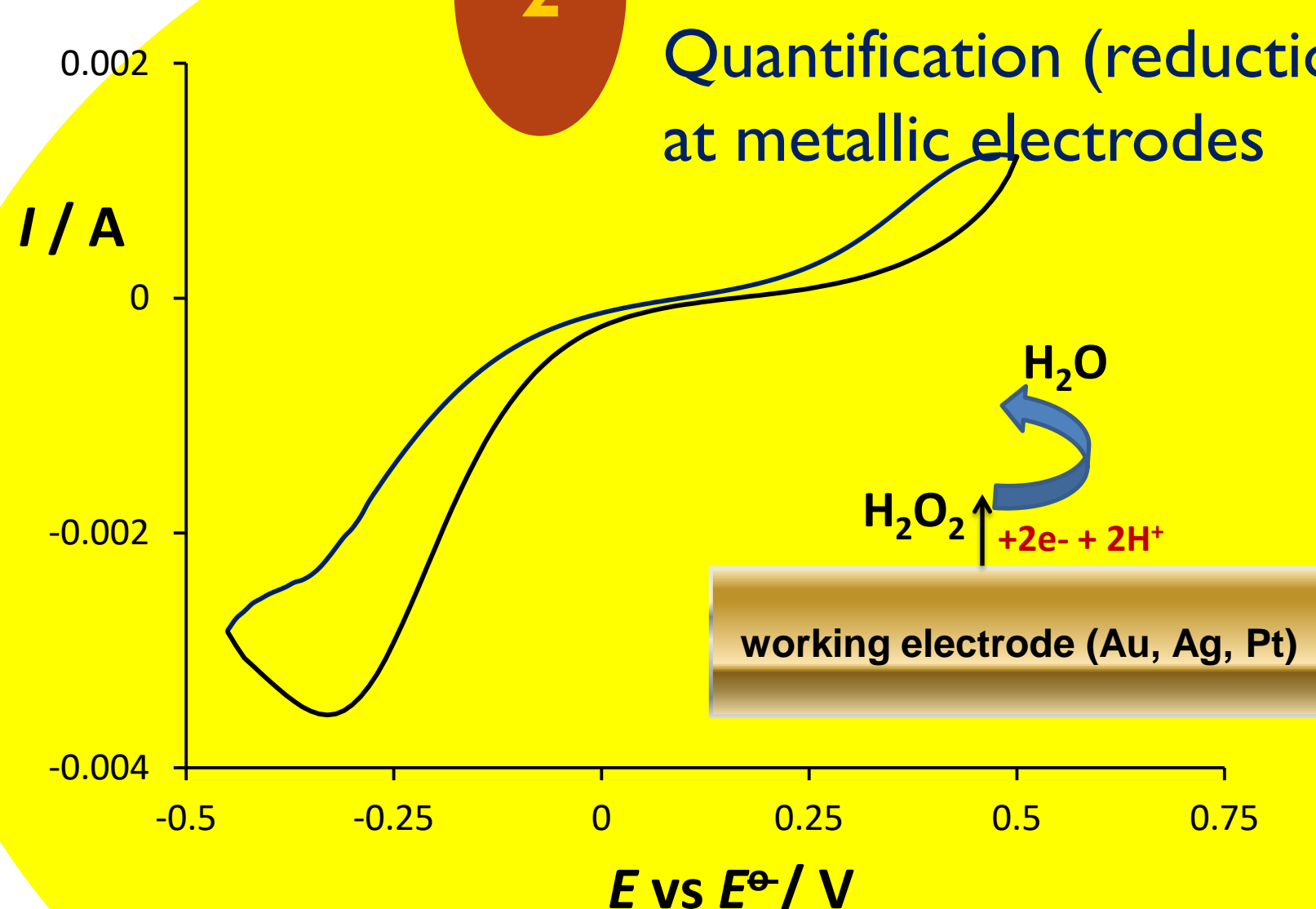


Figure 1. cyclic voltammogram showing reduction of H₂O₂ at bare metallic electrodes

3 Results-II

Second sensor is based on Redox properties of a given Enzyme that is **dissolved** in water And whose redox reaction is Coupled with some redox couple Sensitive to H₂O₂

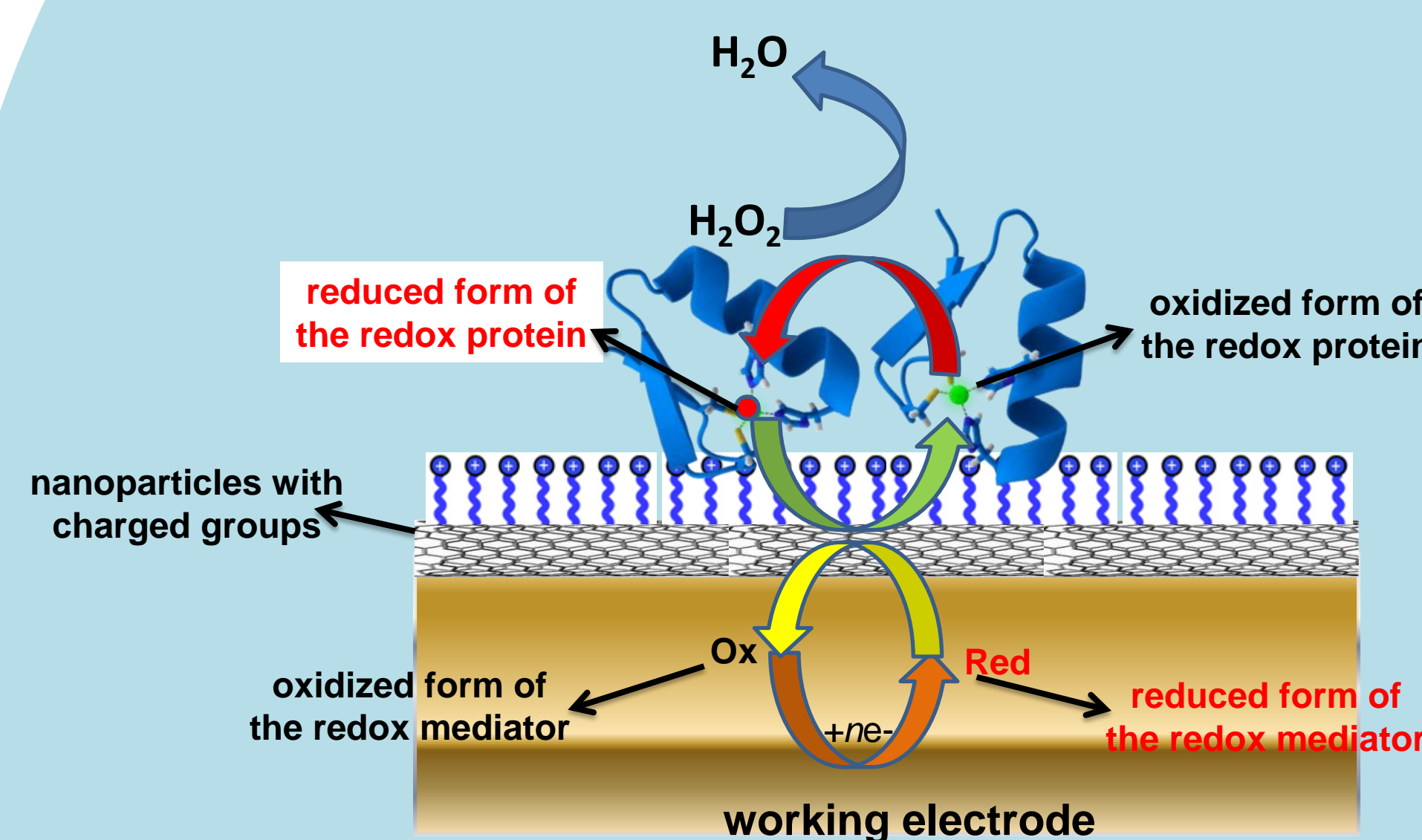
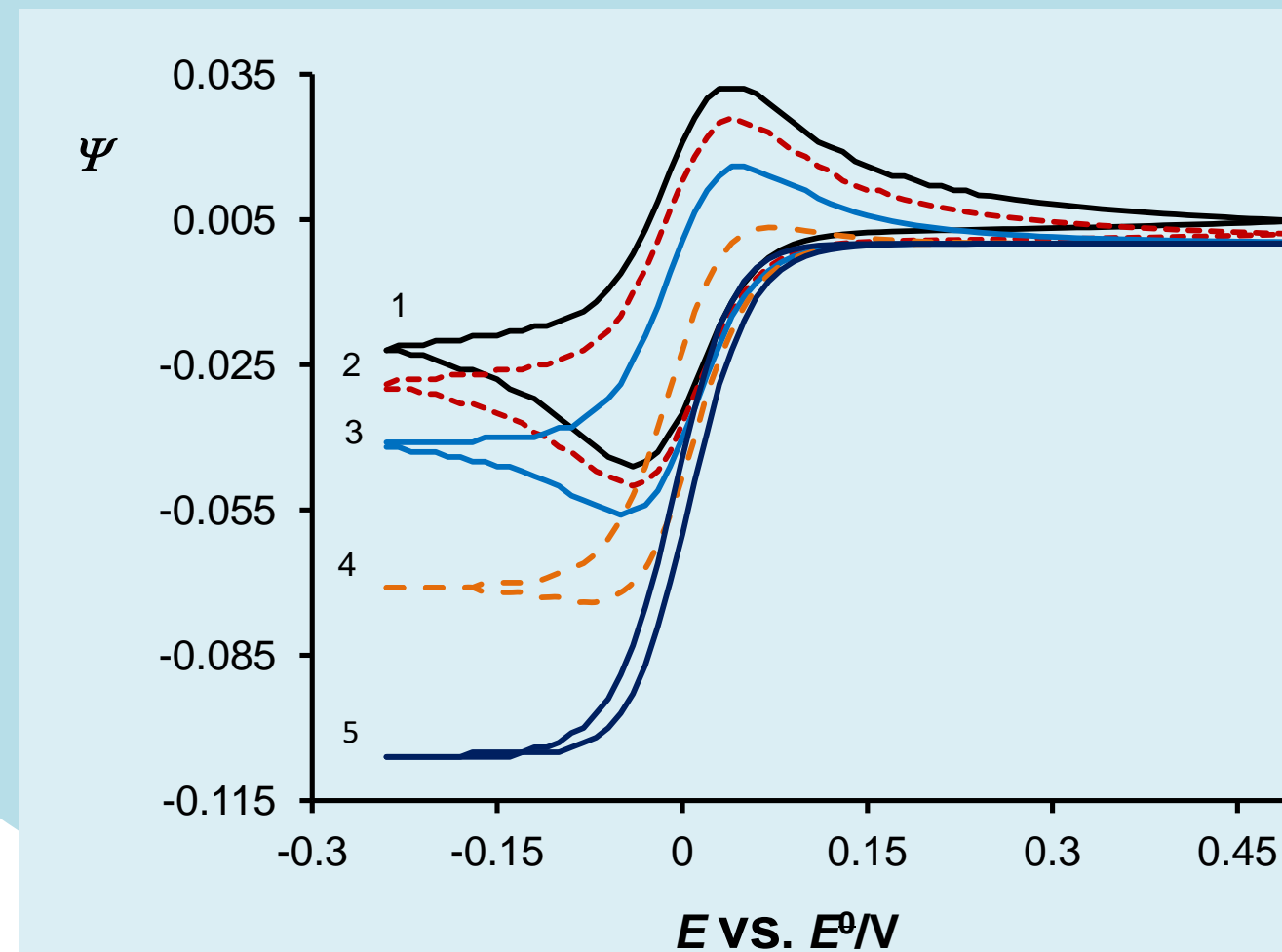


Figure 2. Scheme of voltammetric sensor for H₂O₂ detection based on „dissolved redox enzyme“ with redox mediator Curve 1 is without H₂O₂, and curves 2-5 are with consecutive increase of H₂O₂ concentration. Faradaic currents on voltammograms originate from the reaction Ox + ne- ↔ Red Redox mediator Ox is initially present in the electrochemical cell



4 Results-III

This sensor is based on Redox properties of a given Enzyme that is **adsorbed** on working electrode Surface and whose redox reaction is Coupled with some redox couple Sensitive to H₂O₂

